PREVENTION OF SIGNIFICANT DETERIORATION PRECONSTRUCTION REVIEW AND PRELIMINARY DETERMINATION FOR ADIENT US LLC POLYURETHAN FOAM MANUFACTURING FACILITY IN GILES COUNTY, TENNESSEE

This review was performed by the Tennessee Air Pollution Control Division in accordance with the Rules for Prevention of Significant Deterioration (PSD).

DRAFT, 2022

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Appendix A – PSD Construction Permit 980244

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I. Rule Background

On June 3, 1981, the State of Tennessee adopted Tennessee Air Pollution Control Regulations (TAPCR) 1200-03-09-.01(4), Prevention of Significant Air Quality Deterioration. This Rule has been subsequently amended, with the latest amendments effective April 4, 2018. Under these regulations, a new major stationary source that is included in one of 28 source categories and has the potential or increased potential to emit 100 tons per year or more of any air pollutant regulated in the Clean Air Act must be reviewed with regard to significant deterioration prior to construction. In addition, any source having the potential or increased potential to emit 250 tons per year or more of any of these air pollutants must be reviewed with the same regard.

To comply with the amended PSD regulations, a source with potential emissions greater than significant amounts of a regulated pollutant must meet several criteria. The first criterion is that Best Available Control Technology (BACT) must be applied to all emission points for the applicable PSD pollutant. The second criterion is that the proposed source or modification must not cause or contribute to any violation of the National Ambient Air Quality Standards (NAAQS – see **Table 1**). Finally, increases in ambient concentrations of sulfur dioxide, nitrogen dioxide and particulate matter resulting from emissions discharged by the proposed source must not exceed the increments specified by the PSD regulations (**Table 2**).

	Table 1: National Ambient Air Quality Standards					
Pollutant		Averaging Period	Standard			
Particulate (PM ₁₀)		24-hour	150 μg/m ³			
Matter	(PM _{2.5})	Annual	12.0 µg/m ³ (primary)			
			$15.0 \mu g/m^3$ (secondary)			
		24-hour	35 µg/m ³			
Nitrogen Dio	xide	Annual (primary and secondary)	53 ppb			
(NO ₂)		1-hour (primary)	100 ppb			
Carbon Mono	oxide (CO)	8-hour	9 ppm			
		1-hour	35 ppm			
Sulfur Dioxic	le (SO ₂)	1-hour (primary)	75 ppb			
		3-hour (secondary)	0.5 ppm			
Lead		3-month (primary and secondary)	$0.15 \mu g/m^3$			
Ozone		8-hour (primary and secondary)	0.070 ppm			

Table 2: Maximum Allowable Increases (µg/m3) for Class II Areas			
Pollutant	μg/m ³		
PM10, annual arithmetic mean	17		
PM10, 24-hour maximum	30		
PM2.5, annual arithmetic mean	4		
PM2.5, 24-hour maximum	9		
Sulfur dioxide: Annual arithmetic mean	20		
Sulfur dioxide: 24-hour maximum	91		
Sulfur dioxide: 3-hour maximum	512		
Nitrogen dioxide: Annual arithmetic mean	25		

II. Project Background and Description

On March 30, 2022, Adient US LLC (Adient or Adient Pulaski) submitted an application for a construction permit to increase its emission limits for the production operations at the polyurethane foam manufacturing facility located at 1890 Mines Road, Pulaski, Giles County, Tennessee. This change will increase the Volatile Organic Compound (VOC) allowable emission limit from 308.0 tons per year (tpy) in Title V Operating Permit 569269 to 491.4 tpy.

Adient operates three moisture curing, urethane foam injection lines at its Pulaski, Tennessee facility. The foam lines produce automotive seat cushions and other foam products in clamshell molds. The three molding lines each operate using racetrack-type conveyors, whereby the molds are presented to the various production stations for the foam process to produce a part in its final form. An open mold is presented to the mold release - spray application station where an operator sprays the mold with the mold release agent. The mold release agent is comprised of a wax in a solvent carrier which contains VOC. The mold advances to the pour station where a robot equipped with a urethane component mix head injects the mixed foam components into the open mold. The mold is automatically closed, and the foam reaction occurs in the mold cavity. As the mold advances through the production line, the foam expands, cures and is opened and presented to the extraction station. A worker team removes the molded part, cleans and prepares the mold for another cycle, and the process repeats.

On April 29, 2004, the Technical Secretary issued Title V permit 556316 to the previous owner, Johnson Controls, Inc (Johnson Controls). Condition E4-2 limited VOC emissions from the source to 248.0 tons during all intervals of 12 consecutive months. On May 8, 2006, the Division of Air Pollution Control issued a minor modification to that permit to increase the allowable VOC emissions to 258.0 tons during all intervals of 12 consecutive months.

On February 18, 2010, the Division received an application from Johnson Controls dated February 16, 2010, for a minor modification to Title V permit 556316 to increase the allowable VOC emission limit in

condition E4-2 to 276.7 tons during all intervals of 12 consecutive months. On March 4, 2010, the Division received a letter from Johnson Controls indicating that the baseline emissions used in the February 16, 2010, application were incorrect. The letter stated that "Johnson Controls is requesting an increase in the VOC allowable for the Foam Production Line from 258.0 tons per year to 285.9 tons per year during all intervals of 12 consecutive months." On June 4, 2010, the Division issued Title V renewal permit 562120 which contained condition E4-2 that increased the VOC emission limit for the source to 285.9 tons during all intervals of 12 consecutive months.

On January 15, 2016, the Technical Secretary issued Title V renewal permit 569269. On May 26, 2016, the Division issued an Administrative Amendment to Title V permit 569269 changing the permittee to Adient US LLC (Adient). Adient submitted an application dated July 15, 2016, requesting a Minor Modification for permit 560269 "to increase the VOC limit from 285.9 to 320 tons per year". Adient submitted a revised application dated October 12, 2016, requesting a change to the VOC emission limit to 308 tons per year. On November 21, 2016, the Division issued Minor Modification #1 to Title V permit 569249 which changed the VOC emission limit in condition E4-2 to 308.0 tons during all intervals of 12 consecutive months.

On June 15, 2020, the Division received a revised Title V permit renewal application dated June 11, 2020, from Adient. The application states "During this renewal application process, Adient Pulaski is targeting an increase in Allowable AAP Emissions VOC's from the current 308 tons per AAP to 346 AAP." In June 2021, Adient contacted the Division regarding increasing VOC emissions from the mold release operations to 491.4 tpy¹.

In accordance with 40 CFR 52.21(r)(4), and Division Rule 1200-03-09-.01(4)(a)6. states:

If a particular source or modification becomes a major stationary source or major modification solely by virtue of a relaxation in any enforceable limitation which was established after August 7, 1980, on the capacity of the source or modification otherwise to emit a pollutant, such as a restriction on hours of operation, then the requirements of this paragraph shall apply to the source or modification as though construction had not yet commenced on the source or modification.

The Division determined that the VOC emission limit increase from 248.0 to 258.0 tons during all intervals of 12 consecutive months in 2006 was not the result of a modification to the source, and the facility became a major stationary source solely by relaxation of an enforceable limitation. The Division determined that the VOC emission limit increases to 276.7 tons (during all intervals of 12 consecutive months) in 2010 and to 308.0 tons (during all intervals of 12 consecutive months) in 2016 further relaxed the enforceable emission limitation that kept the facility from being a major stationary source. Therefore, the 2006, 2010, and 2016 emissions increases should have been issued in accordance with paragraph 1200-03-09-.01(4), Prevention of Significant Air Quality Deterioration (PSD).

The proposed modification will result in a significant emission increase for VOC. The project is therefore subject to review under the regulations governing the Prevention of Significant Air Quality Deterioration (PSD).

¹ Projected maximum emissions facility-wide are 502.85 tpy, but Adient will voluntarily accept a limit on facility-wide emissions of 491.40 tpy.

III. Information Used in Analysis

The applicant provided the following information in their March 30, 2022, permit application (Appendix A).

Table 3: Source Description						
Emission Source	Stack (Process Vent) ID	Description	Stack Height (ft)	Stack Exit Flowrate (scfm)	Stack Diameter (ft)	
				1.5.000	2121	
Facility ID	1		80	15,900	3'2"	
(ESRN):	2		90	18,200	3'6"	
28-0076-01	3	Polyurethane Foam Manufacturing Foam line No. 2	96	23,800	3'10"	
	4		61	7,100	2'1"	
All stacks emit at ambient	5		58	5,000	2'	
conditions of	6		10'6"	25,000	5'6"	
70°F, with 1%	7	Polyurethane Foam Manufacturing	10'6"	25,000	5'6"	
moisture from	8	Foam Line No. 1	10'6"	25,000	5'6"	
round discharge	9		10'6"	25,000	5'6"	
points	10		10'6"	25,000	5'6"	
unobstructed in	11	Polyurethane Foam Manufacturing	10'6"	25,000	5'6"	
the upward direction	12	Foam Line No. 3	10'6"	25,000	5'6"	
	13		10'6"	25,000	5'6"	

The proposed modification will affect the emission source listed in Table 3.

Notes: ESRN is the Emission Source Reference Number for the source point on the permit. Stack parameters from **Figure 3** in application dated 2022-03-30

IV. Emissions Analysis

Projected emissions increase from the proposed modification (**Table 4**) were obtained from the information and assumptions given in the March 30, 2022, permit application.

Table 3: Projected Emissions Increases and Permit Emission Limits						
Pollutant	Project Emissions Increase (tons/year)	New Permit Allowable Emission Limits (tons/year)	PSD Significance Threshold (tons/year)	Subject to PSD Review?		
PM/PM ₁₀ /PM _{2.5}	9.45	9.90	25/15/10	No		
VOC	502.85	491.40	40	Yes		

The company's calculated VOC PTE is 502.85 tpy (see calcs in Appendix B of Adient's application dated March 30, 2022), but they chose to request a limit of 491.4 tpy. The facility-wide VOC emission rate is primarily generated by the use and operations associated with the mold release agent, and the balance of the VOC emissions are from other related foam manufacturing and cleaning operations. The VOC emissions from other related foam manufacturing and cleaning operations are relatively low enough that Adient is able to accept a VOC emissions cap of 491.4 tpy. The company provided a letter stating this agreement to the VOC limit of 491.4 tpy.

Since the PSD application submitted by Adient is the result of the 2006 permit modification that made the Adient facility become a major stationary source (by virtue of a relaxation of an enforceable VOC limit), all three foam seating lines are considered new emission units for the purpose of determining baseline actual emissions. Therefore, baseline actual emissions for this source are zero, as required by TAPCR 1200-03-09-.01(4)(a)6.

V. Control Technology Review

V.1 National Emission Standards for Hazardous Air Pollutants (NESHAP)

EPA has promulgated National Emission Standards for Hazardous Air Pollutants (NESHAPs) for various industrial categories. Adient is an area source of hazardous air pollutants (HAPs), which emits less than 10 tons per year of any single HAP and less than 25 tons per year of total HAPs. The application was evaluated to determine the applicability of 40 CFR Part 63 Subpart OOOOOO (National Emission Standards for Hazardous Air Pollutants for Flexible Polyurethane Foam Production and Fabrication Area Sources).

The Adient polyurethane foam operations is an area source of HAPs and an existing source pursuant to 40 CFR §63.11414(c). Accordingly, the operations are subject to 40 CFR Part 63, Subpart OOOOOO: National Emission Standards for Hazardous Air Pollutants for Flexible Polyurethane Foam Production and Fabrication Area Sources. Being subject to 40 CFR Part 63 Subpart OOOOOO, the facility shall not use a material containing methylene chloride as an equipment cleaner to flush the mixhead, use a material containing methylene chloride elsewhere as an equipment cleaner, or use a mold release agent containing methylene chloride flexible polyurethane foam process.

V.2 New Source Performance Standards (NSPS)

Not Applicable

V.3 Other Federal Regulations

The Adient Pulaski facility operates processes which utilize toluene diisocyanate (TDI), Chemical Abstract System no. 26471-62-5. TDI is a regulated hazardous substance under Section 112(r) of the Clean Air Act, Accidental Release Prevention/Risk Management Plan Rule. Adient Pulaski is subject to Program 1 of the rule, which requires preparation and submittal of a Risk Management Plan (RMP) in accordance with 40 CFR §68.12(a) and (b), updated every five years. Program 1 also requires a hazard assessment, which consists of a worst-case release scenario analysis as provided in §68.25 and a five-year accident history as provided in §68.42.

V.4 Best Available Control Technology (BACT) Analysis

Pursuant to TAPCR 1200-03-09-.01(4)(j), this proposed source is required to apply best available control technology for VOC since significant net emission increases are expected from the project as a whole.

Best Available Control Technology means an emission limitation (including a visible emission standard) based on the maximum degree of reduction for each regulated NSR pollutant which would be emitted from any proposed major stationary source or major modification which the Technical Secretary, on a case-by-case basis, taking into account energy, environmental, and economic impacts and other costs, determines is achievable for such source or modification through application of production processes or available methods, systems and techniques, including fuel cleaning or treatment or innovative fuel combustion techniques for control of such pollutant.

In no event shall application of Best Available Control Technology result in emissions of any pollutant which would exceed the emissions allowed by any applicable standard under 40 CFR part 60 or 61. If the Technical Secretary determines that technological or economic limitations on the application of measurement methodology to a particular emissions unit would make the imposition of an emission standard infeasible, a design, equipment, work practice, operational standard or combination thereof, may be prescribed instead to require the application of Best Available Control Technology. Such standard shall, to the degree possible, set forth the emissions reduction achievable by implementation of such design, equipment, work practice or operation, and shall provide for compliance by means which achieve equivalent results.

The EPA policy memorandum dated December 1, 1987, directs applicants and permit reviewers to consider all technically feasible alternatives, including those more stringent than the BACT selection. This is referred to as the "top-down BACT analysis approach." EPA's 1990 New Source Review manual summarizes the top-down BACT analysis in the following steps:

- 1. Identify all control technologies.
- 2. Eliminate technically infeasible options.
- 3. Rank remaining control technologies by control effectiveness.

Table 4: Summary of BACT Analysis				
Emission Source	Pollutant	Emission Limit	Control Technology	
Three Polyurethane Foam Production lines	VOC	491.40 tons of VOC per 12 consecutive months	Utilize good work practice standards to reduce VOC emissions	

4. Evaluate most effective controls and document results.

5. Select BACT.

The results of the BACT analysis are summarized in **Table 5**. Top-down BACT analysis provides that all available control technologies be ranked in descending order of control effectiveness. The most effective control technology is established as BACT unless the applicant demonstrates, and the permitting authority agrees, that technical considerations, or energy, environmental, or economic impacts indicate that the most effective technology is not achievable. If the most stringent technology is eliminated in this fashion, then the next most stringent alternative is considered until a BACT option is selected.

V.4.1 Volatile Organic Compounds (VOC) Emissions – Three Polyurethane Foam Production Lines

BACT applies to each regulated NSR pollutant which would be emitted from any proposed major stationary source or major modification, and BACT analyses are generally performed on each emissions

unit subject to PSD review. Where appropriate, BACT analyses may be performed on groupings of emission units on a case-by-case basis.

Step One: Identify all control technologies: Available control options are those air pollution control technologies or techniques with a practical potential for application to the emissions unit and the regulated pollutant under evaluation. Air pollution control technologies and techniques include the application of production processes or available methods, systems, and techniques, including fuel cleaning or treatment or innovative fuel combustion techniques for control of the affected pollutant. This includes technologies employed outside of the United States. In some circumstances inherently lower-polluting processes are appropriate for consideration as available control alternatives. The control alternatives should include not only existing controls for the source category in question, but also (through technologies. Technologies required under lowest achievable emission rate (LAER) determinations are available for BACT purposes, must also be included as control alternatives, and usually represent the top alternative.

Adient provided the following information in their March 30, 2022, permit application (Appendix A). The application identified the following options to control VOC emissions from the three polyurethane foam production lines:

- Chemical adsorption (carbon and synthetic)
- Recuperative thermal oxidizers coupled with a carbon absorber
- Thermal oxidizer using flare technology
- Scrubber technology
- Refrigeration/condensing VOC control units
- Recuperative thermal oxidizer
- Regenerative thermal oxidizer
- Catalytic thermal oxidizer

The Division reviewed EPA's RACT-BACT-LAER Clearinghouse (RBLC) categories 63.013 and 99.016; guidance documents and found no additional technologies. After further review, the Division proposes Adient utilize good work practice standards to reduce VOC emissions.

Step Two: Eliminate technically infeasible options: In the second step, the technical feasibility of the control options identified in step one is evaluated with respect to the source-specific (or emissions unit-specific) factors. A demonstration of technical infeasibility should be clearly documented and should show, based on physical, chemical, and engineering principles, that technical difficulties would preclude the successful use of the control option on the emissions unit under review. Technically infeasible control options are then eliminated from further consideration in the BACT analysis.

Adient provided the following information in their March 30, 2022, permit application (Appendix A). The application states that the following selected control technologies were rejected as technically infeasible.

Chemical adsorption/Recuperative thermal oxidizers coupled with a carbon adsorber: For this application, large volumes of air are used to collect and transport the mold release solvents and wax overspray to the atmosphere. The wax content in the air will bind on chemically active media surfaces used in adsorber technologies, which makes carbon adsorption and synthetic adsorption incompatible and technically infeasible. For cost projection purposes, a hybrid control system using a carbon concentrator coupled with a [smaller volume] thermal oxidizer was evaluated but is not presented for cost analysis purposes since the concentrator technology is not compatible with the wax materials and thus is not feasible.

Flare technology: Thermal treatment using open or closed flares are typically deployed to control process gases where the exhaust gas has a suitable combustible content and flammability ranges capable of

sustaining an open or closed flame. Since the volatile to air ratio for this application are significantly below the lower flammable limit, flare technology is not suitable and has been determined to be technically infeasible.

Scrubber: The VOCs in this process are not water-soluble. Typical wet and dry scrubbing systems are not capable of collecting and treating VOC airstreams and as a result those technologies are usually reserved or utilized as VOC pretreatment components in a VOC treatment system. Since these technologies do not [substantially] remove or treat VOC laden air, those technologies will not be further evaluated.

Refrigeration/condensing technology: Condensing or refrigeration systems are typically used in low air flow, high VOC content air streams for condensation and collection of the VOC liquid components. For this application, the high-volume air stream and dilute VOC concentration are not compatible or technically feasible control options and are not considered further.

Step Three: Rank remaining technologies by control effectiveness:

Adient provided the following information in their March 30, 2022, permit application (Appendix A).

Since the technologies remaining [not eliminated] are all based on the principle of thermal oxidation (TO), each category will be further evaluated for the economic benefit and cost/control determination. With the desire for a high level of VOC control, those technologies that involve TO have been ranked accordingly by their efficacy and ability to reduce VOCs in relatively dilute airstreams. **Table 6** ranks the remaining control technologies.

Table 5: Ranked Control Options				
Rank	VOC Control Efficiency			
Equal	Recuperative TO	95%		
Equal	Regenerative TO	95%		
Equal	Catalytic TO	95%		
4	Good Work Practice Standards	N/A		

Recuperative Thermal Oxidation: A recuperative TO is a large air-heating device that uses the exhaust temperature to preheat the incoming air using an air-to-air heat exchanger. Relatively low thermal efficiency of the recuperative design results in large fuel gas volume projections, and the relatively short heat of combustion contact time with the process gases (and the targeted VOC compounds) also requires higher treatment (combustion) temperatures to attain the targeted 95% VOC destruction performance.

Regenerative Thermal Oxidization: The regenerative thermal oxidization technology provides operational advantages over recuperative and catalytic technologies in that the thermal efficiency is greatly improved, thereby reducing the quantity of fuel gas needed to attain a specific treatment temperature. The thermal improvement is typically attained using ceramic media and the heat of combustion from the combustion chamber (in this case supplemental fuel burning) is conducted in multiple and sequencing beds in a series of cycles between pre-heat and heat recovery of the treatment beds on a regularly cyclical and frequent basis. The result of this design is fuel gas savings and reduced combustion gas emissions, when compared to other TO technologies.

Catalytic Incineration: This technology allows a reactive catalyst to bring the air stream up to the target VOC oxidation temperature. For the catalyst to work properly, the process gas must be heated to approximately 800°F. Final VOC treatment (destruction) is attained as the reaction of the VOC compounds on the catalytic surface heats the VOC compounds which are then oxidized, and the targeted VOC destruction is complete without the need for additional fuel beyond the preheater section. Catalysts have finite life spans and are subject to catalyst poisoning from compounds in the airstream. Not included in this evaluation are the likely needs for special pretreatment filtration to isolate overspray wax from blinding or contaminating the reactive catalyst surface. Adient is concerned this technology may not be fully compatible with the wax overspray but has presented the US EPA values for comparative and full BACT treatment consideration.

Good Work Practice Standards: A work practice standard is any design, equipment, work practice, operational standard, or combination thereof to reduce air emissions. Good work practice standards may be implemented to satisfy the requirement for the application of BACT when control technology is infeasible or economic limitations exist.

Step Four: Evaluate most effective controls and document results: EPA's 1990 NSR workshop manual states that after technically feasible control options are identified, the energy, environmental, and economic impacts are considered to arrive at the final level of control. If the applicant accepts the highest-ranked control option as BACT, the applicant proceeds to consider whether impacts of unregulated air pollutants or impacts in other media would justify selection of an alternative control option. If there are no outstanding issues regarding collateral environmental impacts, the analysis is ended, and the highest-ranked option is proposed as BACT.

Adient provided the following information in their March 30, 2022, permit application (Appendix A).

The production lines are equipped with existing process ventilation exhaust systems that total approximately 270,000 scfm to meet OSHA spray application requirements. The process exhaust is at room temperature and is comprised of (dilute) VOC concentrations with very low fuel value. The BACT demonstration indicates the VOC concentrations are considered very low for VOC – BACT control treatment considerations. Typical TO systems, treating low concentration VOC streams, require large volumes of fuel gas to maintain proper oxidation temperatures in the combustion chamber of the unit.

The thermal and destruction efficiencies have been summarized and are presented in **Table 7**. The values presented have been calculated using US EPA Cost Control Manual factors. Other factors used in the BACT demonstration are also tabulated for relative comparison of features and benefits. For this BACT demonstration, the three selected TO technologies are believed to be the appropriate types of control for this application.

Table 7: Operating Parameters for VOC Control Equipment							
Gas Preheat of ControlGas Preheat of TreatmentThermal EfficiencyDestructionEstimate							
Control Technology	Treatment Temperature	Efficiency	Efficiency	Fuel Flow	Estimated Heat Input		
Recuperative TO	1, 450 F	70%	95%	3,553 scfm	217.4 MMBtu/hr		
Regenerative TO	1,600 F	95%	95%	431.4 scfm	26.4 MMBtu/hr		
Catalytic	800 F	70%	95%				
Incinerator				1,251.6 scfm	76.6 MMBtu/hr		
Heat inputs are estimated from a conversion factor of 1.020 Btu/scf							

Heat inputs are estimated from a conversion factor of 1,020 Btu/scf

EPA's 1990 New Source Review manual addresses cost effectiveness as follows:

Cost effectiveness (dollars per ton of pollutant reduced) values above the levels experienced by other sources of the same type and pollutant, are taken as an indication that unusual and persuasive differences exist with respect to the source under review. In addition, where the cost of a control alternative for the specific source reviewed is within the range of normal costs for that control alternative, the alternative, in certain limited circumstances, may still be eligible for elimination. To justify elimination of an alternative on these grounds, the applicant should demonstrate to the satisfaction of the permitting agency that costs of pollutant removal for the control alternative are disproportionately high; when compared to the cost of control for that particular pollutant and source in recent BACT determinations. If the circumstances of the differences are adequately documented and explained in the application, and are acceptable to the reviewing agency, they may provide a basis for eliminating the control alternative.

Capital costs include the purchase and installation of equipment items, foundations and supports, piping, insulation, structural steel, and instrumentation. Annual operating costs include utilities, operating labor, and maintenance. The application submitted by Adient on March 30, 2022, states that costs for various control approaches was completed using current costs associated with fuel and electrical utility fees, labor rates, and methods and costs from US EPA's Vatavuk Air Pollution Control Cost Indexes - updated using the U.S Department of Labor Statistics Inflation Calculator. **Table 8** includes the cost per ton of VOC treated for the technically feasible options identified.

Table 8: Cost per Ton of VOC Treated					
Thermal Control Option Dollars per Ton of VOC Treated					
2006 Dollars CPI-Adjusted to 2022 Dollars					
Recuperative Thermal Oxidizer	\$42,261	\$71,032			
Regenerative Thermal Oxidizer	\$10,734	\$18,042			
Catalytic Incinerator	\$19,308	\$32,453			

The application submitted by Adient on March 30, 2022, states that the high projected VOC treatment estimates summarized in **Table 8** result from a variety of factors listed below:

1) The process discharges large volumes of air which are needed to provide a safe working environment for workers in the foam operation area for each of the three foam seating lines.

2) The spray mold release agent and solvent release rates result in low VOC concentrations in the exhaust stream.

3) The high air volumes require extensive and high costs for VOC abatement equipment capable of handling 270,000 scfm and also achieving the needed 95 percent destruction efficiency of the VOC compounds at relatively low concentration.

4) The VOC abatement equipment will require large volumes of fuel in the form of natural gas to heat the large volume, low VOC content, ambient temperature exhaust stream to the target treatment temperatures for each of the targeted and respective TO devices evaluated.

5) The COVID pandemic situation has raised the costs for TO manufacturers and components found in TO equipment. Delays in materials and operating system components have been experienced, along with fabrication and transportation delays.

6) Inflation is at a 40-year high level.

Step Five: Select BACT: The application submitted by Adient on March 30, 2022, concludes that the projected costs per ton of VOC treated are in excess of what would be considered cost effective for VOC controls. On that basis, Adient concluded the current configuration without VOC abatement is BACT for this project and for the existing process.

TDEC-APC Review of Proposed BACT: The Division compared Adient proposed BACT with other control technology reviews for VOC. **Table 9** includes cost information for PSD permits issued in Tennessee in the previous five years. Adient's rejection of add-on controls is consistent with prior determinations issued in Tennessee.

Table 9: Previous BACT Determinations for VOC with Cost Information						
Facility	Permit Number			Selected as BACT?		
Granges Americas Inc.	973712	Thermal Oxidation	\$29,963	No		
Hankook Tire Manufacturing Tennessee,	971720	Regenerative Thermal Oxidation	\$19, 936	No		
LP		Thermal Oxidation	\$61,017	No		
TVA Gleason	975023	Catalytic Oxidation	\$157,000	No		
Domtar Paper	978656		\$10,174	No		
		Catalytic Oxidation	\$21,483			

The RACT-BACT-LAER Clearinghouse (RBLC) was reviewed to identify comparable processes (**Table 10**). The Division identified seven processes for comparison with Adient's proposed BACT.

Table	Table 10: RBLC Search Results – VOC Controls for Polyurethane Foam Manufacturing (Molds)							
RBLC ID	Facility Name	Facility Name Date Process Name		Control Method Description	Efficiency			
IN-0208	NHK Seating of America, Inc.	3/7/2019	Seat Foam Production Line	Regenerative Thermal Oxidizer	95%			
MI-0095	Johnson Controls, Inc.	12/18/2001	MFG Process, Polyurethane Foam	Airless spray gun for mold release/Emission limit	NA			
IN-0137	Foamex Innovations, Inc.	10/12/2012	Polyurethane Foam Coating Line	Best Management Practices/Emission limit	NA			
IN-0137	Foamex Innovations, Inc.	10/12/2012	Polyurethane Foam Production Line	Best Management Practices/Emission limit	NA			
MI-0313	Woodbridge Corporation	11/18/2002	Mold release application	HVLP is used to yield good transfer efficiency. RTO costing \$6500 per ton was not required. (Costs were not verified) / Emission limit	NA			
MI-0163	Steelcase, Inc.	10/9/2002	Mold Release Polyurethane Foam	Water Base Mold Release	100%			
MI-0176	International Foam and Trim	3/25/1994	IN-Mold coating of polyurethane parts.	HVLP coating application equipment used. Dry filters/Emission limit	NA			

NHK Seating of America, Inc. (RBLC ID: IN-0208) uses a regenerative thermal oxidizer for VOC emissions control. After review of the permit in the RBLC, this facility has one automobile seat foam production line, with a capacity of 56 seats per hour, that exhausts through ventilation hoods equipped with dry filters. This line has the capability to use water-based or solvent-based mold release. If solvent-based mold release is used, the VOC emissions are controlled by a regenerative thermal oxidizer. The BACT Analysis (Appendix B of the permit) provided no details for the volume of exhaust air or VOC concentration in the exhaust air stream. An economic impact analysis was not performed as part of BACT. Since no direct comparison of the exhaust stream from the RBLC listing can be made to the Pulaski operation, the control options identified in this BACT demonstration will be used to determine the best available control for this application.

Steelcase, Inc. (RBLC ID: MI-0163) uses water-based mold release agent. There are no details for the polyurethane process in the RBLC for Steelcase, Inc. Adient's attempts to use low emitting alternatives such as co-solvent and water-based mold release agents have failed, resulting in high levels of damaged parts, requiring re-manufacturing and wasted raw materials. Specifically, the use of lower emitting mold release agents does not allow a clean release of the newly formed foam part and frequently causes the foam to stick to the mold, causing damage to the part upon extraction. Therefore, water-based mold release agents cannot be used in Adient's polyurethane foam process.

Pursuant to TAPCR 1200-03-09-.01(4), the following requirements are established as BACT for VOC from source 01, polyurethane foam manufacturing:

• Utilize good work practice standards to reduce VOC emissions

Compliance with this requirement shall be assured by conducting the following daily work practice activities to ensure VOC emissions are minimized and reduced. These activities are outlined in Adient's ISO 14001 Environmental Management System (EMS) Policies as well as internal Standard Work Guidelines:

- All VOC containing mold release containers shall be kept in closed and sealed containers and piping systems at all times until end point usage.
- Inventory storage of all mold release containers shall consist of a fully enclosed locked storage structure complete with self-contained spill containment.
- Mold release material shall be transferred from the tank storage area to the end point discharge via a fully contained and closed loop piping conveyance system.
- In the event of an inadvertent failure of the closed loop conveyance system, flow sensors shall continuously monitor abrupt or out of calibration flow rates to ensure mold release material spillage is kept at a minimum. Spilled material is mitigated immediately by an internal quick reactionary team to ensure VOC containing mold release is contained quickly to not increase potential VOC emissions.
- End point usage of VOC containing mold release material shall continuously be monitored by Line Operators and Line Technicians to ensure proper application at all times.
- Utilizing the ISO14001 EMS Environmental Objectives Form, plant personnel shall participate in monthly mold release usage reduction activities with progress continuously tracked and compared with prior months records.
- Limit of 491.40 tons of VOC per 12 consecutive months.

Compliance with this limit shall be demonstrated by calculating VOC emissions during each calendar month and each period of 12-consecutive months.

VI. Ambient Air Quality Impact Analysis

VI.1 Introduction

On March 30, 2022, Adient submitted an application to expand its operations at 1890 Mines Road in Pulaski, TN by increasing VOC emissions from its currently permitted 308 TPY to 491.4 TPY. The increase in emissions will be due to an increase in the facility's usage of its mold release agent at its three urethane foam injection lines. **Figure 1** shows an aerial photo of the plant and its immediate surroundings which is about 5 km NW of the center of Pulaski. **Figure 2** also shows the near-field within 1 km surrounding the facility.

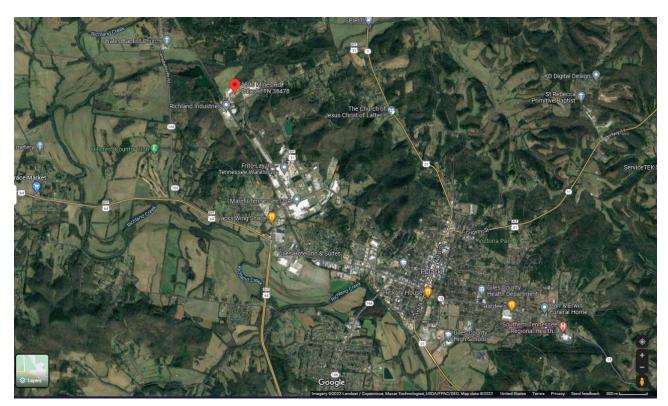
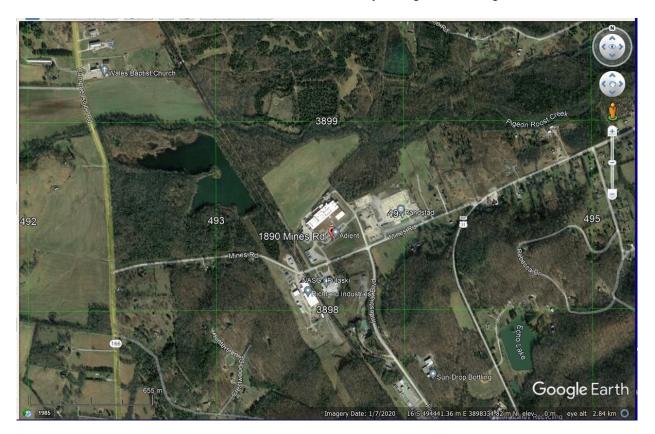


Figure 1 – Adient US, LLC – Pulaski Tennessee Facility in relationship to Pulaski, TN (Google Maps image)

Figure 2 – Adient US, LLC – Pulaski Tennessee Facility Near field with 1 km UTM Grid overlay (Google Earth image)



VI.2 Project Overview

Adient designs, engineers, and manufactures seats for the automotive industry. The manufacturing process includes the production of automotive cushions, backrests, head restraints and other automotive components formulated with polyurethane foam. As part of an overall program to reduce its environmental footprint, Adient has developed a lightweight seating foam and has pioneered the creation of low-emission foams using renewable resources and natural oil polyols. These advancements not only have a direct positive impact on the local community, but also serve to support the automotive industry in lowering the weight of cars, which has a direct effect on improved gas mileage.

Adient - Pulaski operates under Tennessee Division of Air Pollution Control (TDAPC) Title V Permit Number 569269. Federally enforceable emission limits make Adient a major source under Prevention of Significant Deterioration (PSD) regulations [40 CFR 52.21(b)(1)(i)(b)]. Adient proposes to increase annual emissions such that the facility emits more than the PSD Significant Emission Rate (SER) thresholds for the New Source Review (NSR) pollutant category of VOCs. Therefore, increasing the annual emission limitations for this facility will be subject to PSD review (including modeled impact assessment and review) under TAPCR rule 1200-03-09-.01(4)(a)(6).

This section of the PSD Analysis describes the assessment of ambient impacts resulting from the increase in emissions from the proposed permitting action and existing equipment. Section V above discusses the required best available control technology (BACT) analysis. Sections in Volume VI below provide the air quality analysis and analysis of other impacts, respectively. The permit application forms, emission calculations, BACT survey results, and modeling output and plots are contained in the application appendices. With this application, TDAPC proposes to approve Adient's requests for issuance of a PSD Permit for the new operating scenario at their Pulaski facility.

The facility, in Giles County, is located at 1890 Mines Rd in Pulaski, TN 38478, which is about 66 miles south-southwest of downtown Nashville, TN. The area is considered a rural Class II area. The closest Class I areas are: The Sipsey Wilderness Area in north central Alabama (60 mi or 97 km SSW) and Cohutta Wilderness Area in southeastern Tennessee and northern GA (135 mi or 218 km East),

The application describes the process as follows in Section 1.2.2:

"The three molding lines each operate using racetrack-type conveyors, whereby the molds are presented to the various production stations for the foam process to produce a part in its final form. An open mold is presented to the mold release - spray application station where an operator sprays the mold with the mold release agent. The mold advances to the pour station where a robot equipped with a urethane component mix head injects the mixed foam components into the open mold. The mold automatically closes and the foam reaction occurs in the mold cavity. As the mold advances through the production line, the foam expands, cures and is opened and presented to the extraction station. A worker team removes the molded part, cleans and prepares the mold for another cycle, and the process repeats. A process flow diagram is attached in **Figure 2** of the appendix.

Prior to injecting the foam components into the mold, the molds are sprayed with a wax mold release agent to allow removal of the cured foam. The mold release wax is suspended in an aliphatic (non-halogenated) solvent, which contains VOC.

The two-part foam components are moisture-cured producing a polyurethane automotive seat cushion. Halogenated compounds are not used as blowing agents. The foam components are mixed at the gun head, injected/poured into the clamshell molds, the molds are closed and the foam reaction occurs in the mold cavity."

Listed below are the potential emissions from the project compared to the PSD applicability levels for those pollutants emitted at the facility, which require an initial modeling analysis of the facility's projected emissions. Emissions greater than the applicability level necessitate preliminary modeling analyses for those pollutants.

Table 11: Fotential Emissions (Tons/ Fear) Subject to FSD Wodening						
Pollutant	Existing Emissions (tpy) ^[1]	Project Emissions Increases (tpy) ^[2]	PSD Significant Emission Rate (tpy)	PSD Triggered? (Yes/No)		
Filterable Particulate Matter (PM)	113.9 ³	0.0 (9.9 total)	25	No		
Total PM ≤ 10 microns (PM10)	113.9 ³	0.0 (9.9 total)	15	No		
Total PM \leq 2.5 microns (PM2.5)	113.9 ³	0.0 (9.9 total)	10	No		
Sulfur Dioxide (SO ₂)	0.0	0.0	40	No		
Nitrogen Oxides (NOx)	0.0	0.0	40	No		
Volatile Organic Compounds (VOC)	308.0	183.4 (491.4 total)	40	Yes		
Carbon Monoxide (CO)	0.0	0.0	100	No		

Table 11: Potential Emissions (Tons/Year) Subject to PSD Modeling

Notes:

1. From existing Title V Operating Permit 569269, Condition E1.

2. Total VOC and PM is from **Table 2** in **Section 3.1.4** of the permit application.

3. PM existing emissions previously based on the allowable regulatory limit of 0.02 gr/dscf

As required by the PSD regulations, after it is determined that a facility has significant impacts, a typical air quality impact assessment may include some or all of the following steps:

- 1. Determination of the Significant Impact Area (SIA) if any for each pollutant with a Class II SIL &
- 2. monitoring *de minimis* analysis for the proposed emission increase.

Also when proposed new impacts are significant:

- 3. a comprehensive PSD increment consumption analysis for the surrounding Class II area, *and any Class I areas close enough to have significant impacts*,
- 4. a comprehensive Ambient Air Quality Standards impact analysis, and
- 5. an additional airshed impact assessment of the effects on Visibility, Soils, Vegetation, Associated Growth, and Nonattainment Areas, as well as Class I area Air Quality Related Values (AQRV's) if applicable.

The emission rate of PM_{10} is below the significant emission rate (SER) of 15 tpy for PSD applicability, and it is also below the SER threshold for PM (25 tpy) and $PM_{2.5}$ (10 tpy). Also, since the facility process is not heated in a separate curing oven, $PM_{2.5}$ is not an anticipated air pollutant. Hence, all forms of PM maybe considered below the SERs for PM, which makes further PM analysis unnecessary for this permit application.

Since **Table 11** above indicates that this facility is only a major PSD source for VOC, many of the typical ambient PSD analysis steps involving refined modeling with the latest version (v21112) of the refined AERMOD dispersion model were unnecessary for this analysis. For this case only an analysis using Modeled Emission Rates for Precursors (MERPs) was necessary to evaluate the facility's impact on ozone creation from the sources existing VOC emissions.

As a result, VOC emissions were assessed using MERPs and comparing the screening level impact to that of the significant impact level (SIL) associated with the National Ambient Air Quality Standards (NAAQS) for Ozone. Additionally, a breakout of individual source emissions and discharge parameters is unnecessary since MERPs analyses are based on total facility emissions in tons per year.

Finally, representative ozone background data for the MERPs analysis was found, so the preconstruction monitoring requirement was waived based on the availability of representative data from the regional ozone monitor at Fairview Middle School (FMS) in Fairview, Tennessee. Specifically, the regional ozone monitor located at FMS in Fairview, Tennessee was found to be representative of the project site. TDAPC staff agrees with this assessment. The application also describes the monitor as follows:

"The FMS monitor is located approximately 80 kilometers (km) north of Adient's Pulaski facility. Fairview and Pulaski have similar populations of approximately 8,700 and 7,600, respectively. For ozone, a large component of background concentration can be attributed to vehicle use. With similar sized populations, vehicle use can be expected to occur at a similar rate. The two areas also have similar terrain and land use."

Design value data for the FMS monitor is available for the previous ten years (2011 - 2020). FMS monitor information and data for 2020 is provided below.

	Table 12: Representative Ozone Monitor					
Monitor Location	Monitor ID	2020 Design Value (ppb)				
Fairview Middle School,						
Crow Cut Road, Fairview,	471870106	60				
TN 37062						

Table 12: Representative Ozone Monitor

Since VOC is regulated from the facility versus emissions of other pollutants from individual processes and emission points, a summary of the facility-wide emissions for each pollutant and for each modeled scenario, each process and each emission point were unnecessary. Hence **Table 11** emissions above will suffice instead of a detailed description of emission sources and locations which would be pertinent to many other PSD analyses for criterial pollutants.

Similarly, consideration of stack parameters and emissions (based on unit expected maximum capacity) is only pertinent regarding the general release height and annual facility tonnage when comparing them with the general release heights and annual facility tonnages used in EPA's MERPs analysis work. **Table 11** above indicates that the facility's total VOC emissions of 491 TPY is best comparable to the 500 TPY level in EPA's MERPs analyses. Also above, **Table 3: Source Description** indicates that the facility's general emissions release height is less than 100 feet which is most comparable to lower release height of 10 meters, used in EPA's MERPs analyses.

VI.3 CLASS II MODELING: SINGLE-SOURCE IMPACT ANALYSIS

The following sections summarize the methodology used to evaluate the facility's air quality impacts in Class II areas. The dispersion modeling described was performed in accordance with the EPA "Guideline on Air Quality Models" (GAQM, contained in 40 CFR Part 51, Appendix W) (EPA, 2017a), the New Source Review (NSR) Workshop Manual (EPA, 1990), all applicable EPA clarification memorandums and guidance documents, and direction and regulatory guidance provided by the Tennessee Department of Environment and Conservation (TDEC) and EPA Region IV. The modeling analysis focused on demonstrating that the ambient impact of proposed emissions from the Adient project will be in compliance with all applicable NAAQS and PSD Class II increments.

VI.3.1 Dispersion Modeling Methodology

Since VOC emissions are the target of this analysis, EPA's work to define a screening methodology to evaluate precursor emission impacts on ozone formation using EPA's work with photochemical grid modeling (PGM) methods was relied on for this analysis, instead of using the American Meteorological Society / Environmental Protection Agency Regulatory Model (AERMOD) gaussian dispersion model, which is typically used to determine predicted impacts in the Class II area surrounding the facility.

In December 2016, the EPA developed a simple screening methodology to estimate single source impacts on secondary pollutants which they described as: Modeled Emission Rates for Precursors and debuted using the acronym "MERPs". MERPs reflect levels of increased precursor emissions that are not expected to cause a significant contribution to O_3 for PSD applications. A MERP can relate:

 $\Box \Box VOC$ emissions to O₃; and

 \square \square NO_X emissions to O₃.

MERPs modeling methods are intended to conservatively estimate secondary pollutant impacts in what is also termed a Tier 1 screening analysis to demonstrate ambient compliance, before a more refined and resource intensive Tier 2 analysis using detailed photochemical grid modeling is necessary.

VI.3.2 Assessment of Secondary Pollutant Impacts

The SIL for ozone (O_3) which EPA recommended in their April 2018, guidance on SIL's, was used to demonstrate that the proposed potential impacts from Adient do not cause or contribute to a violation of the NAAQS for ozone. The recommended SIL for O_3 was used to assess potential impacts from secondary pollutants emitted from Adient.

The EPA December 2016, guidance memorandum provided a framework on how to develop source-specific or site-specific MERPs. The guidance document did not endorse a specific MERP value, though it did provide illustrative MERPs from the EPA's modeling of two hypothetical sources in various locations across the United States.

EPA's initial 2016 MERPs guidance memorandum was finalized by EPA in April of 2019. Tennessee has also provided more customized MERPs guidance for sources in Tennessee since the 2019 memorandum. According to EPA and Tennessee guidance, sources are required to estimate both the impacts of primarily emitted and secondarily formed pollutants as part of the PSD program. This is normally done using a Tier 1 MERPs analysis first, and if a Tier 1 analysis fails to demonstrate ambient compliance, a Tier 2 analysis using PGM techniques may be used if necessary.

Tennessee's guidance regarding MERPs was used to assess potential impact of secondarily formed O_3 which could be expected from the chemical interaction of Adient's potential emissions of VOCs with nitrogen oxides emitted from off-site combustion sources in the vicinity.

The precursors to ground-level ozone formation are VOC and NO_x . However, since this project only exceeded the SER for VOC and not NO_x , Adient only needed to consider VOC contributions to ozone as a part of this assessment.

VI.3.2.1 Ozone Assessment

TDEC documentation titled "*Tennessee Guidance on the Use of EPA's MERPs to Account for Secondary Ozone and Fine Particulate Formation in Tennessee Under the New Source Review (NSR) Prevention of Significant Deterioration Program (PSD)*" was customized for sources in Tennessee by the Division of Air Pollution Control using the PGM results EPA used in their MERPs guidance. The secondary ozone analysis in this report specifically relied upon the Tennessee Guidance document to conservatively predict impacts resulting from VOC emissions at the Adient- Pulaski facility. Below is a quote from the pertinent part of Tennessee's latest MERPs guidance on page 4.

"The significant Impact Level (SIL) for ozone is 1 ppb.... The units for the Maximum Model Impact are parts per billion (ppb) for ozone.... The most conservative (lowest) MERP values from the six (6) nearby hypothetical sources (in or near Tennessee) by precursor and pollutant are contained in table... These default MERP values can be used for Tier 1 demonstrations in Tennessee without further justification."

The abbreviated **Table 13** from the Tennessee guidance contains the following default MERP values for 8-hour ozone impacts resulting from NOx and VOC emissions.

Precursor	8-hour Ozone
NO _x	156

	VOC	1,542
--	-----	-------

Note: The default values are the lowest (conservative) MERP values for hypothetical sources in and near Tennessee."

In the Tennessee MERPs Guidance on pages 12 and 13 it states:

SILs Analysis

MERPs can be used to determine if a facility's proposed emission increases will result in secondary impacts that are above the SILs. Once either one of the precursor pollutants triggers this analysis because their emissions are above the PSD Significant Emission Rates (SERs), then emissions of the other precursor pollutant must be included in the analysis to determine the synergistic impact that both pollutants have together, even though the other pollutant's emissions may fall below the SER. The analysis is unnecessary only when emissions of both precursor pollutants are below the respective SERs.

For ozone, the following equation should be used:

 $\frac{EMIS_NOx}{MERP_NOx} + \frac{EMIS_VOC}{MERP_VOC} < 1$

EMIS_NOx and EMIS_VOC are the proposed emission increases for NOx and VOC (tpy). MERP_NOx and MEPR_VOC are the MERPs for NOx and VOC (tpy). If the sum of the ratios is less than 1, then the secondary ozone impacts are below the ozone SIL and the applicant does not need to perform a cumulative analysis for ozone. If the sum of the ratios is equal to or greater than 1, the applicant must perform a cumulative analysis for ozone.

VI.3.2.2 Single-Source Impact Modeling Results

Summary results for each significantly emitted pollutant and avg time.

 \underline{O}_3 – The NAAQS for Ozone (O₃) is 70 ppb, which equates to 140 µg/m³, for an 8-hour average. The SIL for Ozone is 1 ppb. Since O₃ is a secondary pollutant formed in the atmosphere by precursor VOC and NOx pollutants, the source was evaluated using single source MERPs methodology below in this section to demonstrate that the source will not cause or contribute to a violation of the NAAQS for O₃.

Resultant details for secondary Ozone resulting from VOC emissions.

The secondary O_3 impact assessment is compared to the established SIL for Ozone of 1 part per billion (ppb). As outlined in **Table 7** of the TDEC November 2019 guidance (seen above), the default MERP values (tpy) for Tennessee PSD applications are 156 tpy of NO_x and 1542 tpy of VOC. Per equations provided above from page 13 of the TDEC guidance

(https://www.tn.gov/content/dam/tn/environment/air/documents/apc-modeling-

page/apc_TN%20Guidance%20on%20the%20Use%20of%20EPAs%20MERPs%20to%20Account%20f or%20Secondary%20Formation%20in%20Tennessee_11222019.pdf), the SIL analysis demonstration for the proposed project at Adient is as follows:

For the Class II significant impact modeling analysis, the maximum predicted impact was compared to the only pertinent PSD Class II SIL, which was the SIL for ozone. The modeled impacts for the Tier 1 secondary pollutant analysis scenario are summarized below.

Since the source does not emit primary ozone and only emits one precursor to secondary ozone formation, the analysis centers around the single precursor VOC. In the Tennessee MERPs equation above, the value for *EMIS_VOC* is 491.4 TPY, while the value for *EMIS_NOx* is zero. The MERP value for VOC related to 8-hour ozone is 1,542 TPY from the MERPs table, so the computed ratio for NOx is zero while the computed ratio for VOC is less than one as seen below.

$$\frac{EMIS_NOx}{MERP_NOx} + \frac{EMIS_VOC}{MERP_VOC} = \frac{0 \ TPY \ of \ NOx}{156 \ NOx \ MERP} + \frac{491.4 \ TPY \ of \ VOC}{1542 \ VOC \ MERP} = 0.32 \ which \ is < 100 \ MERP$$

Hence, an evaluation of the equation indicates that the sum of the computed ratios of emissions to MERPs is also less than 1. Therefore, the 491.4 TPY of VOC emitted by the Adient facility would be expected to have an impact less than the SIL of 1 ppb for ozone. As a result, any further cumulative analysis for VOC is unnecessary to approve the company's ambient assessment for VOC.

Additionally, since the predicted ozone value is less than the threshold value of 1, a cumulative analysis for ozone was unnecessary, and it is not necessary to include the background ozone concentration in a more refined cumulative evaluation for ozone described on page 13 of the Tennessee MERPs guidance.

VI.4 CLASS I AREA AMBIENT AIR QUALITY IMPACT ASSESSMENT

Class I areas are federally protected areas for which more stringent air quality standards apply to protect unique natural, cultural, recreational, and/or historic values. Air quality dispersion modeling analyses to support the PSD application for the Adient-Pulaski Class I analysis include the following assessments:

- 1. Determination of the facility potential pollutant emission quantities relative to PSD significant emission rates (SER) as defined in PSD rules (40 CFR 52.21).
- 2. Determination of the source location and distance within 300 km of any Class I area. Facility impacts at Class I areas located beyond 300 km from the PSD source are considered insignificant.
- 3. Determination of compliance with the Federal Land Managers (FLMs) Air Quality Related Values (AQRVs) in addressing regional haze visibility and acidic deposition.
- 4. Determination of whether facility impacts at Class I areas located within 300 km from the PSD source are considered significant. If so, a determination of compliance with the EPA's NAAQS and PSD increments for those triggered criteria pollutants that have Class I area increments.

The Adient facility submitted separate analyses to assess impacts on AQRVs and on the Class I SILs for the NAAQS and PSD increments.

VI.4.1 Initial Screening Criteria for AQRVS

The Federal Land Managers (FLM) have the authority & responsibility to protect air quality related values (AQRVs) in Class I areas, and to consider in consultation with the permitting authority whether a proposed major emitting facility will have an adverse impact on such values. Class I AQRVs for which PSD modeling is typically conducted include visibility impairment, ozone (O₃) effects on vegetation, and effects of sulfur and nitrogen deposition on soils and surface waters.

The FLMs developed an Initial Screening Criteria, Q/D, to determine if sources greater than 50 km away from a Class I area need to perform any further Class I AQRV impact analyses. The Q/D ratio is calculated by summing the annual VOC, SO₂, NO_x, PM, and H₂SO₄ emissions (in tons per year, based on 24-hour maximum allowable emissions and adjusted as if it were operated for 8,760 hours per year then dividing by the distance (in kilometers) to the nearest Class I area. If the Q/D value is less than or equal to 10, the source is considered to have negligible impacts on AQRVs in the Class I area and no further analyses are needed.

The Initial Screening Criteria for Adient emissions were calculated for the five Class I areas within 300 km of the Adient facility below (with the approximate distance to the facility listed):

- ► Sipsey National Wilderness Area (~ 97 km)
- ► Cohotta Wilderness Area (~218 km)
- ► Mammoth Cave National Park (~ 226 km)
- ► Joyce Kilmer-Slickrock Wilderness Area (~278 km)
- ► Great Smoky Mountain National Park (~281 km)

All other Class I areas are located at distances greater than 300 km from the facility.

A Class I area analysis includes a Class I PSD increment assessment for pollutants subject to PSD review (increasing above the SER) and an AQRV analysis for visibility, ozone, and deposition that could impact a Class I area's resources.

Class I AQRV Analysis

The Class I AQRV analysis was prepared in accordance with the FLM's Air Quality Related Values Work Group (FLAG) *Phase I Report – Revised (2010)*. The FLMs developed a screening-level criteria involving facility emissions and distance to the Class I area (Q/D ratio) for sources greater than 50 km from a Class I area to determine whether adverse impacts could occur to AQRV from a pollutant. The equation used is provided below:

 $Q/D \text{ ratio} = \frac{\text{Facility Emissions (Q)}}{\text{Distance to Class I Area (D)}}$

If the Q/D ratio is below 10, it is presumed that no adverse impact will occur, and no further AQRV analysis is required. An AQRV analysis for the Class I Areas of concern can be found below.

Class I Area	Pollutant	Q (tpy)	D (km)	Q/D	Q/D sum
Sipsey	PM	9.90		0.10	
Wilderness Area	VOC	491.4	97	5.07	5.17
	PM	9.90	218	0.05	2.30

Table 14: Q/D ratios for Class I Areas within 300 km of Adient-Pulaski (From Table 7: Class I AQRV Analysis in permit application)

Cohotta Wilderness Area	VOC	491.4		2.25		
Mammoth Cave	PM	9.90	226	0.04	2.21	
National Park	VOC	491.4	220	2.17	2.21	
Joyce Kilmer-	PM	9.90		0.04		
Slickrock Wilderness Area	VOC	491.4	278	1.77	1.81	
Great Smoky	PM	9.90		0.04		
Mountain National Park	VOC	491.4	281	1.75	1.79	

The Q/D ratios for PM and VOC and their sums for each of the Class I Areas are well below the threshold of 10; therefore, it is presumed there are no adverse impacts from Adient, and no further analysis is required.

Class I Increment Analysis

Adient does not have a significant increase of PM/PM₁₀/PM_{2.5}, above the corresponding SERs, so an analysis of significant PM impacts vs. PM increments or PM NAAQS was not necessary. Additionally, since there is no Class I PSD increment established for VOC, any other increment analysis for this project would not be applicable.

Class I NAAQS Analysis

Finally, since the computed MERP ratio in the Class II area for NOx is zero and the corresponding ratio for VOC is much less than one, then the combined MERP ratio at the much increased distances to the Class I areas is assumed to be even less, making anticipated Class I ozone impacts insignificant as well.

VII. ADDITIONAL IMPACTS ANALYSIS

PSD applies to new major sources or major modifications at existing sources for pollutants where the area the source is located is in attainment or unclassifiable with the NAAQS. Adient is a major source of VOCs, a precursor to ozone. Adient is located in the City of Pulaski, County of Giles in the State of Tennessee, which is designated attainment for ozone.

A PSD major source subject to PSD review is required to conduct an air quality analysis and an additional impacts analysis, among other requirements. Pursuant to 40 CFR §52.21(o), the additional impacts analysis consists of three parts: Growth Analysis, Soils and Vegetation Impacts Analysis and Visibility Impairment Analysis. Each of these analyses is addressed below.

VII.1 GROWTH ANALYSIS

The Adient plant is located in an industrial park setting on the north side of Pulaski, Tennessee. Pulaski is located in the south central portion of the state nearly equidistant between Memphis and Chattanooga. The

general vicinity outside of the industrial development is mainly agricultural pastureland and woodland with some residential development east of the plant (**Figure 1**). The workforce consists of 305 employees living within commuting distance of the plant. The size of the workforce has resulted in little impact on population growth in Giles County, which has a population of 29,503 as of 2018. A review of historical aerial photos dating back to 1998, indicate that there has been little industrial or commercial development in the immediate vicinity of the Adient plant, and no substantive residential growth in the general area.

VII.2 SOIL AND VEGETATION ANALYSIS

Particulate Matter and Volatile Organic Compounds

The criteria for evaluating impacts on soils and vegetation is taken from EPA's, *A Screening Procedure for the Impacts of Air Pollution Sources on Plants, Soils, and Animals*, EPA, 1980. According to US EPA, "...sources more than 10 km from any Class I areas, exemptions provide that no analysis of impairment need be done if emission increases are below specified limits." Specified limits are de minimis values found at 40 CFR 52.21(b)(23)(i).

The Adient operations are located more than 10 km from any Class I areas in the region. The criteria air pollutants emitted by the Adient Pulaski operations include PM and PM_{10} . Emissions of PM and PM_{10} are below the significance levels (de minimis values) found at 40 CFR 52.21(b)(23)(i).

The Adient operations generating particulate matter emissions are not the direct result of combustion or combustion byproducts and therefore, Adient assumes there are no emissions of $PM_{2.5}$, but in any event, all PM emitted by the source is less than the respective thresholds, including the threshold for $PM_{2.5}$.

Adient emits VOC at an annual rate above the de minimis value of 40 tpy, but screening concentrations are not available for VOC. VOC is a precursor to ozone and while ozone is identified as a "Regulated Pollutant", EPA indicates that a screening concentration [for O_3 is] available, and now a simple procedure for estimating the ozone impact of a single source is currently available through the use of MERPs.

Additionally, the secondary NAAQS were established at concentration levels below which no harmful effects to either soil or vegetation is expected (per US EPA, Office of Air Quality Planning and Standards, *New Source Review Workshop Manual (Draft)*, U.S. EPA, Research Triangle Park, NC, October, 1990).⁶ As discussed above, EPA has developed a two-tiered evaluation for secondary ozone formation from VOC. As demonstrated by this application, the VOC emissions from Adient's facility are below the default TDEC MERPs value, indicating that no adverse impact to compliance with the ozone NAAQS is expected. As such, VOC emissions from the facility will not negatively affect soil and vegetation in the surrounding area.

Other than VOC, Adient does not emit any criteria air pollutants above their respective significance thresholds.

VII.3 VISIBILITY IMPAIRMENT ANALYSIS

US EPA prescribes the use of its *Workbook for Plume Visual Screening and Analysis (Revised)*, October 1992 (EPA-454/R92-023), methodologies for purposes of conducting a visibility impairment analysis. A

visibility impairment analysis is generally required to determine the impact on sensitive areas such as state parks, wilderness areas, airports, scenic sites and overlooks. Three levels of screening procedures are outlined by US EPA. If the criteria for the first screening level, the most conservative level, are met, no further analysis is needed.

The VISCREEN model is recommended for the Level 1 screen. The VISCREEN model primarily considers NO_2 and particulate emission increases associated with a project. VISCREEN does not consider or calculate visibility impacts from ozone.

Level 1 Screen Analysis

Level 1 analysis incorporates conservative parameters to determine plume impacts. Default values for particle size and density, and a default of worst-case meteorological condition of F stability and 1.0 meters per second (m/s) wind speed are used for the analysis, while all emissions are assumed to exit the plant from one point. The worst-case meteorological condition is expected to persist for 12 hours with a wind direction that would transport the plume directly to the sensitive area being analyzed.

A maximum particulate matter emission rate of 2.64 lb/hour, based on a maximum emission rate of 9.9 tpy assuming an operating schedule of 7,488 hours/year (6 days per week, 24 hours per day, 52 weeks/year), was input to the model. It was assumed that there were no emissions of NOx, soot, primary nitrogen dioxide and primary sulfate. A background visual range of 25 kilometers was used. All other inputs relied on the default parameters.

While the Pulaski facility is greater than 10 km from any Class I area, a distance of 10 km was assumed in order to run the most conservative analysis of visibility impairment.

Level 1 modeling results indicate that the Adient operations do not adversely impact visibility within or beyond a 10 km radius.

OVERALL RESULTS OF PLUME VISIBILITY SCREENING
SOURCE: Adient US LLC, Pulaski, CLASS I AREA: NA
INSIDE class I area Plume delta E DOES NOT EXCEED screening criterion for SKY background Plume delta E DOES NOT EXCEED screening criterion for TERRAIN background Plume contrast DOES NOT EXCEED screening criterion for SKY background Plume contrast DOES NOT EXCEED screening criterion for TERRAIN background
OUTSIDE class I area Plume delta E DOES NOT EXCEED screening criterion for SKY background Plume delta E DOES NOT EXCEED screening criterion for TERRAIN background Plume contrast DOES NOT EXCEED screening criterion for SKY background Plume contrast DOES NOT EXCEED screening criterion for TERRAIN background
SCREENING CRITERIA: DELTA E = 2.0 GREEN CONTRAST = 0.050

VIII CONCLUSIONS AND CONDITIONS OF APPROVAL FOR AMBIENT AIR QUALITY IMPACT ANALYSIS

Projected emissions of VOC from the proposed modification exceed the PSD significance levels at maximum operating rate and maximum hours of operation. This major modification is subject to review under the regulations for the Prevention of Significant Deterioration contained in 1200-03-09-.01(4). The proposed control technology satisfies the requirement to install BACT, as required by the PSD regulations. The BACT requirements are incorporated into the permit to be issued for the proposed modification. The proposed changes will not result in ambient impacts that would exceed any National Ambient Air Quality Standards or PSD Increments and will not cause or contribute to adverse impacts on Air Quality Related Values in nearby Class I areas.

After review of the information submitted with the PSD application, it is concluded that the proposed modification qualifies for approval, subject to the terms and conditions of the proposed PSD construction permit (Appendix A).

Appendix A –PSD Construction Permit 980244

Appendix B – Emission Summary for Proposed PSD Permit 980244

Emission Summary

Source 01

		Permit Number:	980244
Source Status: New Modificatio	on Expansion Relocation] Permit Status: New	Renewal
PSD⊠ NSPS⊡ NESHAPs⊠	Previous Permit Number:	Construction	Operating 569269

	Pounds/Hour			Tons/Year				
Pollutant	Actual	Uncontrolled Potential	Allowable/ Permitted Potential	Actual	Uncontrolled Potential	Allowable/ Permitted Potential	Date of Data	Applicable Standard TAPCR 1200-03-
VOC [1]					502.85	491.4	3/30/2022	0901(4) 0707(2)
PM ^[2]	2.16	2.16	3.00	6.22	9.46	9.90	3/30/2022	0701(5)

1. VOC emissions are uncontrolled and are based on the data in Page 36 of the application (3/30/22). The facility has agreed to limit (through recordkeeping and good work practice standards) VOC emissions to 491.4 tons per any period of 12-consecutive months. The emission limit is based on an agreement letter dated July 18, 2022, from the permittee.

2. PM emissions are uncontrolled and are based on the data in Page 37 of the application (3/30/22). The regulatory allowable limit (TAPCR 1200-03-07-.03(1)) for PM is 46.29 lb/hr. The facility has agreed to limit (through recordkeeping) PM emissions to 3.00 lb/hr and 9.90 tons per year based on an agreement letter dated July 18, 2022, from the permittee.

The actual TPY is based on the facility's operating hours of 5,760 hr/yr from APC 10 form in the application (3/30/22).

Appendix C – Application for Proposed PSD Permit 980244

Aemilia Hamel

From:	Air.Pollution Control
Sent:	Wednesday, 30 March, 2022 20:43
То:	APC Permitting
Subject:	FW: Email 1 of 2 - Construction Permit Application_Source ID 28-0076 Adient US, LLC
Attachments:	Adient_Air Quality Construction Permit Application_March 2022_final_PART 1 OF 2.pdf

From: O'Brien, Ann <AOBrien@scsengineers.com>
Sent: Wednesday, March 30, 2022 5:02 PM
To: Air.Pollution Control <Air.Pollution.Control@tn.gov>
Cc: Kris Patrick Foster <kris.patrick.foster@adient.com>
Subject: [EXTERNAL] Email 1 of 2 - Construction Permit Application_Source ID 28-0076 Adient US, LLC

*** This is an EXTERNAL email. Please exercise caution. DO NOT open attachments or click links from unknown senders or unexpected email - STS-Security. ***

To Whom It May Concern, on behalf of Adient US, LLC, Source ID No. 28-0076, attached please find a construction permit application, which includes all sections except for the Safety Data Sheets. Given the size of the application, which exceeds 25 MB, we are sending it in two separate emails. Please let me know if you have any questions. Or, you can contact Kris P. Foster, EHS Analyst with Adient US, LLC. He can be reached at kris.patrick.foster@adient.com.

Attached to this email are all application documents EXCEPT the Safety Data Sheets. They will be sent as an attachment to a second email (Email 2 of 2 – Construction Permit Application_Source ID 28-0076 Adient US, LLC).

An original application is being sent to the Department by way of certified mail. Best regards, Ann O'Brien

Ann P. O'Brien Project Manager SCS Engineers 6215 North Lundy Avenue Chicago, IL 60646 USA 773-775-6362 (W) 773-540-5199 (C) aobrien@scsengineers.com

Driven by Client Success www.scsengineers.com 1890 Mines Road Pulaski, Tennessee 38478 Tel 931.363.5666 Fax 931.424-6722



March 30, 2022

Michelle Owenby, Technical Secretary State of Tennessee, Department of Environment and Conservation Division of Air Pollution Control William R. Snodgrass Tennessee Tower 312 Rosa L. Parks Avenue, 15th Floor Nashville, TN 37243

SUBJECT: Air Construction Permit Application, Prevention of Significant Deterioration Adient US, LLC 1890 Mines Road, Pulaski, TN 38478

To Whom It May Concern,

Enclosed please find an air construction permit application for Adient US, LLC (Adient) located at 1890 Mines Road, Pulaski (Giles County), Tennessee. Adient designs, engineers and manufactures seats for the automotive industry. The manufacturing process includes the production of automotive cushions, backrests, head restraints and other automotive components formulated with polyurethane foam. This air quality construction permit application has been prepared in accordance with the requirements of the Tennessee Department of Environment & Conservation (TDEC), Air Pollution Control Regulations, Tennessee Code: Title 68: Chapter 201. Adient US, LLC (Adient) is a major source of air pollution, specifically as to volatile organic compounds (VOC) and is seeking authority to expand its foam manufacturing operations located in Pulaski, Tennessee.

By way of background, in June 2021, Adient approached TDEC regarding increasing VOC emissions from the facility to 491.4 tpy. Discussions regarding this increase continued during the remainder of 2021 and into the spring of 2022. In October 2021, TDEC alleged that Adient had not met certain permit requirements. Adient denied these allegations and discussions between the parties ensued. As a result of these discussions, the parties entered into a Consent Order on February 24, 2022. Pursuant to the discussions, Adient is submitting this PSD permit application.

Should you have any questions or need additional information, please contact Mr. Kris Patrick Foster, EHS Analyst at (931) 363 – 5666 or <u>kris.patrick.foster@adient.com</u>. You can also contact Ms. Stephanie Taylor, Project Professional with SCS Engineers. Stephanie can be reached at (913) 749 – 0733 or <u>staylor@scsengineers.com</u>.

Thank you for your assistance in processing this permit application. We look forward to meeting and working with you on this project.

Sincerely,

Ryan Speck Plant Manager

Ms. Michelle Owenby, Technical Secretary March 30, 2022

Encl: Air Construction Permit Application, Prevention of Significant Deterioration

Cc: Mr. Kris Patrick Foster, EHS Analyst Mr. Rick H. Palmer, Regional EHS Manager Ms. Stephanie Taylor, SCS Engineers Mr. Jeff Pfost, Environmental Partners, Inc.

Air Quality Construction Permit Application

Prevention of Significant Deterioration

Adient US, LLC 1890 Mines Road Pulaski, Tennessee 38478

March 30, 2022



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1.0 INTRODUCTION

This air quality construction permit application has been prepared in accordance with the requirements of the Tennessee Department of Environment & Conservation (TDEC), Air Pollution Control Regulations, Tennessee Code: Title 68: Chapter 201. Adient US, LLC (Adient) is a major source of air pollution, specifically as to volatile organic compounds (VOC) and is seeking authority to expand its operations located in Pulaski, Tennessee (**Figure 1**).

1.1 SITE LOCATION AND CONTACTS

Company name:	Adient US, LLC
Street address:	1890 Mines Road
City, State, Zip Code:	Pulaski, Tennessee 38478
Telephone number:	(931) 363 - 5666
Website:	https://www.adient.com
Responsible official:	Mr. Ryan Speck
Job title:	Plant Manager
Electronic mailing address:	Ryan.speck@adient.com
Environmental contact:	Mr. Kris Patrick Foster
Job title:	EHS Analyst
Electronic mailing address:	kris.patrick.foster@adient.com
Corporate environmental contact:	Mr. Rick H. Palmer
Job title:	Regional EHS Manager
Electronic mailing address:	ricki.h.palmer@adient.com

1.2 PROCESS OVERVIEW

Adient designs, engineers and manufactures seats for the automotive industry. The manufacturing process includes the production of automotive cushions, backrests, head restraints and other automotive components formulated with polyurethane foam. As part of an overall program to reduce its environmental footprint, Adient has developed a lightweight seating foam and has pioneered the creation of low-emission foams using renewable resources and natural oil polyols. These advancements not only have a direct positive impact on the local community, but also serve to support the automotive industry in lowering the weight of cars, which has a direct effect on improved gas mileage.

Standard Industrial Classification:	2821: Plastics Materials, Synthetic Resins and Nonvulcanizable Elastomers	
North American Industrial Classification System:	326150: Urethane and Other Foam Product (except Polystyrene) Manufacturing	
Establishments primarily engaged in manufacturing syn nonvulcanizable elastomers. Important products of this phenolic and other tar acid resins; urea and melamine acrylic resins; polyethylene resins; polypropylene resins; petroleum polymer resins; miscellaneous resins, includin polyesters, polycarbonate resins, acetal resins, and fluo	industry include: cellulose plastics materials; resins; vinyl resins; styrene resins; alkyd resins; rosin modified resins; coumarone-indene and g polyamide resins, silicones, polyisobutylenes,	

1.2.1 Process Description, Pulaski Operations

Adient operates three moisture curing, urethane foam injection lines at its Pulaski, Tennessee facility. The foam lines produce automotive seat cushions and other foam products in clamshell molds.

The three molding lines each operate using racetrack-type conveyors, whereby the molds are presented to the various production stations for the foam process to produce a part in its final form. An open mold is presented to the mold release - spray application station where an operator sprays the mold with the mold release agent. The mold advances to the pour station where a robot equipped with a urethane component mix head injects the mixed foam components into the open mold. The mold is automatically closed and the foam reaction occurs in the mold cavity. As the mold advances through the production line, the foam expands, cures and is opened and presented to the extraction station. A worker team removes the molded part, cleans and prepares the mold for another cycle, and the process repeats. A process flow diagram is attached (**Figure 2**).

Prior to injecting the foam components into the mold, the molds are sprayed with a wax mold release agent to allow removal of the cured foam. The mold release wax is suspended in an aliphatic (non-halogenated) solvent, which contains VOC.

The two-part foam components are moisture-cured producing a polyurethane automotive seat cushion. Halogenated compounds **are not used** as blowing agents. The foam components are mixed at the gun head, injected/poured into the clamshell molds, the molds are closed and the foam reaction occurs in the mold cavity.

Copies of the Safety Data Sheets (SDSs) for the wax mold release agent and other raw materials containing regulated air pollutants (**Appendix A**) are attached.

2.0 AIR PERMIT HISTORY

April 29, 2004: TDEC issued Title V operating permit number 556316 ("Permit 556316"), to Johnson Controls, Inc. (28-0076) for a polyurethane foam manufacturing operation. Condition E4-2 of Permit 556316 limited the VOC emissions to 248.0 tons during any 12-consecutive months.

May 8, 2006: TDEC issued Minor Modification #2 to Permit 556316, to Johnson Controls, Inc. to increase the allowable VOC emissions from 248.0 to 258.0 tons during any 12-consecutive months. According to TDEC, documentation attached to Minor Modification #2 indicated that the increase was based on a letter dated January 23, 2006 from Kathy Arnold. The Division has no copy of the January 23, 2006 letter.

February 18, 2010: TDEC received an application from Johnson Controls, Inc. dated February 16, 2010, for a minor modification to Permit 556316, to increase the allowable VOC emissions from 258.0 to 276.7 tons per 12-consecutive months.

March 4, 2010: According to TDEC, it received a letter from Johnson Controls, Inc. that indicated the baseline emissions used in the February 16, 2010 application was incorrect. The letter stated, "Johnson Controls is requesting an increase in the VOC allowable for the Foam Production."

June 4, 2010: TDEC issued renewal Title V operating permit number 562120 to Johnson Controls, Inc. which contained condition E4-2 that increased the VOC emission limit to 285.9 tons during any 12-consecutive months.

January 15, 2016: TDEC issued renewal Title V operating permit number 569269 ("Permit 569269"), to Johnson Controls, Inc. for the polyurethane foam manufacturing operation.

May 26, 2016: TDEC issued Administrative Amendment #1 to Permit 569269, for the ownership change of the polyurethane foam manufacturing operation from Johnson Controls, Inc. to Adient US, LLC, which occurred on May 1, 2016.

November 21, 2016: TDEC issued Minor Modification #1 to Permit 569269, to Adient US, LLC, to increase the VOC emissions from 285.9 tons during all intervals of 12-consecutive months to 308 tpy during all intervals of 12-consecutive months.

June 15, 2020: TDEC received a revised Title V permit renewal application dated June 11, 2020 from Adient US, LLC. The application stated, "During this renewal application process, Adient Pulaski is targeting an increase in Allowable AAP Emissions VOC's from the current 308 tpy per AAP to 346 AAP."

In June 2021, Adient approached TDEC regarding increasing VOC emissions from the mold release operations to 491.4 tpy¹. Discussions regarding this increase continued during the remainder of 2021 and into the spring of 2022. In October 2021, TDEC alleged that Adient had not met certain permit requirements. Adient denied these allegations and discussions between the parties ensued. As a result of these discussions, the parties entered into a Consent Order on February 24, 2022. Pursuant to the discussions, Adient is submitting this PSD permit application.

3.0 CONSTRUCTION PERMIT APPLICATION

As a result of increased business related to a new automotive program, Adient is seeking a construction air permit for the existing Pulaski foam operations. Adient is a major source of VOC and accordingly, this permit application is being submitted under the Prevention of Significant Deterioration (PSD) permitting program. Pursuant to 1200-03-09-.01(4), Prevention of Significant Air Quality Deterioration (PSD), the following elements are addressed:

- 1. Emission calculations for applicable regulated air pollutants, namely criteria air pollutants and hazardous air pollutants (HAPs);
- 2. Applicable federal and state air pollution control regulations and rules and standards;
- 3. Top-down Best Available Control Technology (BACT) analysis;
- 4. Air quality assessment;
- 5. Additional impact analysis including potential growth impacts, soil and water impacts and visibility impacts;
- 6. Additional information and data provided on TDEC permit application forms.

Adient understands that this construction permit application is subject to public participation pursuant to 1200-03-09-.01(4). Adient further understands that the Department accomplishes the requirement associated with public participation by electronic notice on the Department's website on a monthly basis including applicants seeking authority to construct, modify and/or operate air pollution emission sources.

3.1 FACILITY-WIDE EMISSION RATES

3.1.1 Volatile Organic Compounds (VOCs)

As described above, projected VOC emissions are the result of in-mold release agents used to produce polyurethane foam seating systems for the automotive industry. The mold release agents may be sprayed or

¹ Projected maximum emissions facility-wide are 502 tpy, but Adient will voluntarily accept a limit on facility-wide emissions of 491.4 tpy.

manually applied to the mold surface, prior to foam component application. The mold release agent is comprised of wax in a solvent carrier.

Attempts to use low emitting alternatives such as co-solvent mold release agents have failed, resulting in high levels of damaged parts, requiring re-manufacturing and wasted raw materials. Specifically, the use of lower emitting mold release agents does not allow a clean release of the newly formed foam part and frequently causes the foam to stick to the mold, causing damage to the part upon extraction.

Adient is proposing to increase the allowable facility-wide VOC emission rate to 491.4 tons per year (tpy). The facility-wide VOC emission rate is primarily generated by the use and operations associated with the mold release agent, and the balance of the VOC emissions are from other related foam manufacturing and cleaning operations. The VOC emissions from other related foam manufacturing and cleaning operations are relatively low enough that Adient is able to accept a VOC emissions cap of 491.4 tpy.

The project net emissions increases and decreases are summarized below. Pursuant to TDEC's request, baseline emissions in this application are 0.0 tpy.

VOC Emissions - Netting Exercise				
Baseline VOC emissions: 0.0 tpy (past actual)				
Proposed VOC emission increases:	491.4 tpy			
Contemporaneous VOC emission decreases:	0.0 tpy			
Net VOC emission increases:	491.4 tpy			

 Table 1.
 Contemporaneous VOC Emission Decreases and Increases

Detailed VOC emissions calculations are included with this application (**Appendix B**). Because the existing operation is considered a PSD major source of VOC emissions (greater than 250 tons), the requested VOC increases (the project) are greater than 250 tpy, and the emissions increases and net emissions increases are also greater than the PSD significance level of 40 tons, this request requires a major source PSD review.

3.1.2 Particulate Matter Emissions

The wax mold release agents (subject of the VOC increases) contain wax solids suspended in a solvent carrier. The solids portion of the mold release agents are sprayed or manually applied.

Particulate emissions result from overspray of the mold release agent. The purpose of the mold release application is to apply the release wax to the mold which prevents the foam from sticking to the mold surface. The process uses high volume, low pressure (HVLP) spray applicators for improved transfer efficiency. Overspray in the form of total suspended particulate (TSP) has been equated to particulate matter 10 microns and smaller (PM₁₀ filterable) particulates. The spray wax particles are highly cohesive such that they stick to the mold surface as larger agglomerated overspray particles. Since the process is not heated in a separate curing oven and does not involve combustion or by-products of incomplete combustion, it is assumed PM_{2.5} is negligible, if emitted at all.

HVLP applicators are typically equated to a transfer efficiency of 65 percent. For purposes of determining worst case/highest particle emissions, a transfer efficiency of 55 percent has been assumed in the calculation of particle emissions.

Adient has calculated a maximum PM/PM_{10} (and $PM_{2.5}$) emission rate of 9.455 tpy, but is seeking authority to emit up to 9.90 tpy of particulate matter emissions for which emission calculations are attached (**Appendix B**).²

The emission rate of PM_{10} is below the significant emission rate (SER) of 15 tpy for PSD applicability, and it is also below the SER threshold for PM (25 tpy) and $PM_{2.5}$ (10 tpy). Reference is made to $PM_{2.5}$ although as mentioned above, the process is not heated in a separate curing oven and accordingly, $PM_{2.5}$ is not an anticipated air pollutant.

No other criteria pollutants are associated with the process so potential emissions for criteria pollutants other than VOC and PM and PM_{10} are considered negligible. Detailed criteria air pollutant emission calculations, including emission rates for all processed raw materials are included in **Appendix B**.

3.1.3 Hazardous Air Pollutant (HAP) Emissions

Some raw materials contain low concentrations of federal hazardous air pollutants (HAPs) and consequently, Adient is an area source of HAPs. The calculation of HAP emission rates is included in **Appendix B**.

3.1.4 Criteria Air Pollutant and HAP Emissions – Summary

Total facility-wide criteria air pollutant and HAP emission rates are summarized below.

Criteria Air Pollutant and HAP Emissions					
Regulated	Baseline Emissions	Emission Increases (Decreases)	Proposed Allowable Emissions	PSD Significance Threshold	
Air Pollutant ³		Tons p	er Year		Comments
SO ₂	NA	0.00	0.00	40	<ser< td=""></ser<>
NOx	NA	0.00	0.00	40	<ser< td=""></ser<>
СО	NA	0.00	0.00	100	<ser< td=""></ser<>
PM total	0.00	0.00	9.90	25	<ser< td=""></ser<>
PM10	NA	0.00	9.90	15	<ser< td=""></ser<>
PM _{2.5}	NA	0.00	9.90	10	<ser< td=""></ser<>
VOC ⁴	0.00	0.00	491.4	40	>SER
Pb	NA	0.00	0.00	0.6	<ser< td=""></ser<>

Table 2.	Facility-wide	Emissions	Summary

 $^{^2}$ The current maximum allowed particulate matter (TSP) for the mold release operation is 113.9 tpy. However, this particulate emission rate is much higher than currently sought as part of this permit application on the belief that at the time the existing PM emissions were established, a different PM estimating method and/assumptions of emissions factors were used. Careful reconsideration of the materials using mass balance with inclusion of EPA established spray gun transfer efficiency guidelines results in much lower worst-case maximum potential particulate emissions projections. Further, equating total particulate matter to PM_{10} is believed to also present worst case PM_{10} projected/potential emissions.

³ The Pulaski foam manufacturing facility is currently permitted to emit 113.90 tpy of PM (total) and 308.00 tpy of VOC. ⁴ Projected maximum VOC emission rates are estimated to be 502 tpy, but Adient will accept a voluntary limit on emissions of 491.40.

Criteria Air Pollutant and HAP Emissions					
Regulated	Baseline Emissions	Emission Increases (Decreases)	Proposed Allowable Emissions	PSD Significance Threshold	
Air Pollutant ³		Comments			
CO ₂ e	NA	0.00	0.00	75,000	<ser< td=""></ser<>
HF	NA	0.00	0.00	3	<ser< td=""></ser<>
H ₂ SO ₄	NA	0.00	0.00	7	<ser< td=""></ser<>
H ₂ S	NA	0.00	0.00	10	<ser< td=""></ser<>
Highest single HAP	NA	NA	<10 tpy	NA	NA
Total HAPs	NA	NA	<25 tpy	NA	NA

4.0 **REGULATORY APPLICABILITY**

The Pulaski foam manufacturing facility is a major source of VOC emissions and an area source of HAPs. The following federal and state regulatory programs are applicable to these operations.

4.1 NEW SOURCE PERFORMANCE STANDARDS

There are no New Source Performance Standards (NSPS) pursuant to 40 CFR Part 60 that are applicable to the molded foam operations conducted at the Adient Pulaski facility.

Adient does not operate any fuel combustion equipment except small air makeup units that provide space heating by way of natural gas combustion and consequently, 40 CFR 60 subpart Dc is not applicable.

Adient Pulaski does not operate any new stationary compression ignition (CI) internal combustion engines (ICE), as defined (40 CFR 60 subpart IIII).

Adient Pulaski does not operate any new stationary spark ignition (SI) internal combustion engines (ICE), as defined (40 CFR 60 subpart JJJJ).

4.2 NATIONAL EMISSION STANDARDS FOR HAZARDOUS AIR POLLUTANTS

The Adient Pulaski foam operations are an area source of HAPs and an existing source pursuant to 40 CFR §63.11414(c) accordingly, the operations are subject to 40 CFR 63 subpart 000000 National Emission Standards for Hazardous Air Pollutants for Flexible Polyurethane Foam Production and Fabrication Area Sources. The following standards will apply to this source:

- The permittee must not use a material containing methylene chloride as an equipment cleaner to flush the mix head or use a material containing methylene chloride elsewhere as an equipment cleaner in a molded flexible polyurethane foam process.
- The permittee must not use a mold release agent containing methylene chloride in a molded flexible polyurethane foam process.

Reference: 40 CFR 63.11416(c) and 1200-03-09-.03(8) TAPCR

- The permittee shall keep a certification on file at the plant site that contains the following statements, and must be signed by a responsible official:
 - This facility does not use any equipment cleaner to flush the mix head which contains methylene chloride, or any other equipment cleaner containing methylene chloride in a molded flexible polyurethane foam process in accordance with § 63.11416(c)(1).
 - This facility does not use any mold release agent containing methylene chloride in a molded flexible polyurethane foam process in accordance with § 63.11416(c)(2).

Reference: 40 CFR 63.11417(c)

The Adient Pulaski operations are an area source of HAPs and accordingly, are not subject to the boiler MACT (40 CFR 63 DDDDD).

The Adient Pulaski facility does not operate any stationary reciprocating internal combustion engines (RICE) and accordingly, is not subject to 40 CFR 63 subpart ZZZZ.

4.3 OTHER FEDERAL REGULATIONS

The Adient Pulaski operates processes which utilize toluene diisocyanate (TDI), Chemical Abstract System no. 26471-62-5. TDI is a regulated hazardous substance under Section 112(r) of the Clean Air Act, Accidental Release Prevention/Risk Management Plan Rule.

Adient Pulaski is subject to Program 1 of the rule, which requires preparation and submittal of a Risk Management Plan (RMP) in accordance with 40 CFR §68.12(a) and (b), updated every 5 years. Program 1 also requires a hazard assessment, which consists of a worst-case release scenario analysis as provided in §68.25 and a five-year accident history as provided in §68.42.

Reference: 40 CFR 68

4.4 FEDERAL AND STATE RECORDKEEPING, REPORTING AND MONITORING REQUIREMENTS AND EMISSION STANDARDS

Regulatory Reference	Requirement	Comments
Recordkeep	oing, Reporting and Monitoring Re	equirements
40 CFR 82	Operation, maintenance and disposal of regulated refrigerants	
1200-03-0902(11)(e)1(iii)	Monitoring and associated recordkeeping requirements	
1200-03-09.02(11)(e)1(iii)(II)II	Records retention	
1200-03-0902(11)(e)1(iii)	Monitoring and records reporting	
200-03-0902(11)(e)3(v)(IV)	Annual compliance certifications	Submittal to both EPA Region 4 and TDEC
1200-03-0501(1)		
1200-03-0503(6)	Visible emissions standards	
1200-03-0502(1)		
1200-03-08	Fugitive dust prohibition and control standards	

Regulatory Reference	Requirement	Comments				
Recordkeeping, Reporting and Monitoring Requirements						
1200-03-2602(2)(i)	Annual Accounting Period and Associated Reporting	Annual emissions reporting and payment of emission fees				
1200-03-0902(11)(e)1.(iii)	Semi-annual reporting	Monitoring and recordkeeping reporting				
1200-03-1002(2)(a)	Documentation of all VOC and HAP-containing materials	Maintain SDSs, records of VOC and HAP content and maintain these records for five years				
1200-03-3203(3)	Accidental release plan	Maintain the plan and on an annual basis, submittal a certification of compliance with the plan				
1200-03-1002(2)(a)	Recordkeeping of logs	Mandates time schedule for routine, ongoing records entries				
1200-03-0904.	VOC and HAP records	Maintain records of VOC and HAP usages and emissions				
	Emission Standards					
1200-03-0704(1).	Particulate matter (PM) emitted from this source shall not exceed 0.02 gr/dscf) of exhaust gases					
1200-03-0707(2)	VOC emissions limit	Currently 308 tpy; proposed 491.4 tpy				

5.0 BEST AVAILABLE CONTROL TECHNOLOGY (BACT) ANALYSIS

PSD regulations require that a PSD major modification incorporate and use the best available control technology (BACT). This application includes a top-down BACT demonstration for VOC emissions using the 5-step top-down BACT demonstration methods as follows:

- Step 1 Identify all control technologies
- Step 2 Eliminate technically infeasible options
- Step 3 Rank remaining control technologies by control effectiveness
- Step 4 Evaluate most effective controls and document results
- Step 5 Findings and Conclusions

5.1 IDENTIFICATION OF ALL CONTROL TECHNOLOGIES

The feasibility of various VOC control options are first determined based upon conventional pollutant specific control technologies. For this application, the control of volatile organic compounds (VOCs) results from the use of mold release agent on the three foam molding lines. The mold release agents are comprised of mold release wax consisting of approximately 4.1 percent, by weight, solids suspended in a VOC solvent. The evaporation of the mold release solvent (VOC carrier) results in VOC emissions which is the subject of the BACT control demonstration.

The considerations have focused on air treatment systems and technologies capable of treating relatively dilute VOC containing air that also contains wax overspray. Water scrubbing devices and dry filtration methods would be considered highly effective at removing the wax solids from the airstream but limited or

ineffective at removing VOC compounds. The technologies capable of reducing VOC content in air have been identified as follows:

- Chemical adsorption (carbon or synthetic)
- Recuperative thermal oxidizers coupled with a carbon absorber.
- Thermal oxidizer using flare technology
- Scrubber technology
- Refrigeration/condensing VOC control units
- Recuperative thermal oxidizers
- Regenerative thermal oxidizers
- Catalytic thermal oxidizers

5.2 ELIMINATION OF TECHNICALLY INFEASIBLE OPTIONS

Chemical adsorption: For this application, large volumes of air are used to collect and transport the mold release solvents and wax overspray to the atmosphere. The wax content in the air will bind on chemically active media surfaces used in adsorber technologies, which makes carbon adsorption and synthetic adsorption incompatible and technically infeasible. For cost projection purposes, a hybrid control system using a carbon concentrator coupled with a [smaller volume] thermal oxidizer was evaluated but is not presented for cost analysis purposes since the concentrator technology is not compatible with the wax materials and thus is not feasible. For this reason, oxidation of the entire air stream (without concentration) is technically the most viable VOC control strategy for the BACT demonstration.

Flare technology: Thermal treatment using open or closed flares are typically deployed to control process gases where the exhaust gas has a suitable combustible content and flammability ranges capable of sustaining an open or closed flame. Since the volatile to air ratio for this application are significantly below the lower flammable limit, flare technology is not suitable and has been determined to be technically infeasible.

Scrubber: Typical wet and dry scrubbing system are not capable of collecting and treating VOC airstreams and as a result those technologies are usually reserved or utilized as VOC pretreatment components in a VOC treatment system. Since these technologies do not [substantially] remove or treat VOC laden air, those technologies will not be further evaluated.

Refrigeration/condensing technology: Condensing or refrigeration systems are typically used in low air flow, high VOC content air streams for condensation and collection of the VOC liquid components. For this application, the high-volume air stream and dilute VOC concentration are not compatible or technically feasible control options and are not considered further.

5.3 RANKING OF REMAINING CONTROL TECHNOLOGIES

Oxidation by way of heating air streams above the thermal destruction temperature are highly effective at treating VOC air streams of varying concentrations. Wax particles from overspray and accompanying VOCladen exhaust is compatible with high temperatures found in most oxidizing equipment, although wax solids build up on heat exchangers and solid catalysts may present operational and maintenance issues. Regardless of the wax content concerns, the remaining oxidizer base treatment technologies may be feasible and are further evaluated in the BACT demonstration.

Ranking of the remaining control technologies has been compiled based on the anticipated effectiveness of the control equipment (thermal oxidization). All of the remaining technologies (not eliminated) are oxidizers that are highly effective in treating (reducing) VOCs from the foam line mold release agent. While carbon

monoxide and oxides of nitrogen are also components of thermal oxidization by-products, the benefits of VOC controls typically outweigh the relatively small adverse impacts from the by-product gases produced.

Since the technologies remaining [not eliminated] are all based on the principle of thermal oxidation (TO), each category will be further evaluated for the economic benefit and cost/control determination. With the desire for high level of VOC control, those technologies that involve TO have been ranked accordingly by their efficacy and ability to reduce VOCs in relatively dilute airstreams.

The identification and ranking of TO technologies include:

- Recuperative thermal oxidation
- Regenerative thermal oxidation
- Catalytic thermal oxidation.

The production lines are currently equipped with existing process ventilation exhaust systems that total approximately 270,000 scfm (**Figure 3**). The process exhaust is at room temperature and is comprised of (dilute) VOC concentrations with very low fuel value. The concentration of VOC in the exhaust stream at the proposed (full potential) concentration rates is calculated at 6.93 x 10⁻⁶ pounds of VOC per cfm of exhaust or the equivalent of 0.0069 pounds of VOC per 1,000 scfm. Roughly and on average, the concentration of the targeted VOC is slightly less than 96 parts per million (ppm) by weight. Typical lower flammable limits (LFL) concentrations for VOC compounds are measured in percentages and to be combustible would require VOC concentrations that are several orders of magnitude higher than this process air to sustain combustion or even to have measurable fuel value. This is the reason the BACT demonstration indicates the VOC concentrations. Typically TO systems treating low concentration VOC streams require large volumes of fuel gas to maintain proper oxidation temperatures in the combustion chamber of the unit.

The three molding lines each operate using racetrack-type conveyors, whereby the molds are presented to the various production stations for the foam process to produce a part in its final form. The nature and positioning of the personnel working on the respective molding line places the de-mold, mold cleaning, and spray wax stations in close proximity. Hooded spray exhaust is provided for a relatively large area of the human occupied stations. For worker safety and health reasons, and to meet OSHA spray application requirements, the exhaust rate from each of the lines is high. At the same time, the spray rate of VOC's in the spray wax material is relatively low compared to other types of industrial VOC sources, resulting in very high air flow rates with relatively low VOC concentrations.

Spray application areas are also regulated by [safe] building code requirements under the guidance and restrictions of the National Fire Protection Association (NFPA). These regulations are intended to minimize fire hazards and provide a safe working environment for workers using flammable and combustible materials. The mold release agent is comprised of solvent carriers that are considered flammable agents. NFPA requires solvent concentrations be maintained well below their respective flammable limits. Safe limits are maintained in the process air system as a result of adequate ventilation rates and proper system design components and elements. The exhaust volumes presented in the BACT analysis are the accumulation of measured process exhaust volumes. They are considered necessary for personnel safety reasons and believed accurate to the BACT demonstration.

Because the VOC laden airstream is dilute and is also at room temperature, it has little or no heat value, requiring thermal oxidation (TO) treatment to bring the gases to their final and desired treatment temperature, for example to 1,600°F. With little or no fuel value, the TO control unit will be required to supplement the exhaust gas with large quantities of fuel. The large volume of process exhaust also requires very large volumes of fuel gas, whereby the amount of fuel gas is directly proportional to the thermal

efficiency of the TO technology option evaluated. For example, the greater the thermal efficiency, the less fuel is required to attain and maintain proper treatment temperature; and as a result, lower operating costs are realized and lower combustion emissions are also possible with higher thermal efficiency TO units/technologies.

Those VOC treatment systems identified in the BACT analysis capable of high thermal efficiency reduce the quantity (and cost) of supplemental fuel which reduces the overall (annual) operating cost of the treatment system. Of the systems ranked above, the regenerative thermal oxidizer has the highest thermal efficiency while maintaining a high degree of VOC removal/treatment efficacy. Catalytic systems require lower air preheat temperatures and appropriately sized catalytic exchanger sections. Catalytic costs and size limitations may restrict the viability of this treatment technology, however catalytic oxidation is further evaluated for cost/benefit determination.

The thermal and destruction efficiencies have been summarized and are presented in **Table 3**. The values presented have been calculated using US EPA Cost Control Manual factors. Other factors used in the BACT demonstration are also tabulated for relative comparison of features and benefits. For this BACT demonstration, the three selected TO technologies are believed to be the appropriate types of control for this application and each TO type is discussed further.

Table 3.	Operating Parameters for VOC Control Equipment
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Control Technology	Gas Preheat of Treatment Temperature	Thermal Efficiency	Destruction Efficiency	Fuel Flow	Estimated Heat Input	
Recuperative TO	1,450° F	70%	95%	3,553 scfm	217.4 MMBtu/hr	
Regenerative TO	1,600° F	95%	95%	431.4 scfm	26.4 MMBtu/hr	
Catalytic TO	800° F	70%	95%	1251.6 scfm	76.6 MMBtu/hr	
Heat inputs are estimated from a conversion factor of 1,020 Btu/scf						

Recuperative TO technology has a low thermal efficiency and as a result very high fuel operating cost as evidenced in **Table 3** above. This TO technology is further evaluated for cost/benefit determination. As set forth above, all three TO technologies have equivalent destruction efficiencies and are therefore ranked equally from a VOC control and viability perspective.

5.4 EVALUATE MOST EFFECTIVE CONTROLS AND DOCUMENT RESULTS

Adient completed evaluations of various VOC treatment and control options identified above (treatment and control type as well as configuration options). Costs for various control approaches was completed using *current* costs associated with fuel and electrical utility fees, labor rates, and methods and costs from US EPA's Vatavuk *Air Pollution Control Cost Indexes* - updated using the U.S Department of Labor Statistics Inflation Calculator. TDEC has directed Adient to perform the BACT cost/benefit exercise without adding retrofit costs, and as a result retrofit costs are not considered in the analysis, as Adient believes they should be.

5.4.1 Recuperative Thermal Oxidation

A recuperative TO is a large air-heating device that uses the exhaust temperature to preheat the incoming air using an air-to-air heat exchanger. This technology recovers and recuperates a portion of the spent (combusted) gas energy, providing the technology name. Air-to-air heat exchangers have limited efficiencies and a value of 70% thermal efficiency was used by the EPA Cost Control demonstration tool. Relatively low thermal efficiency of the recuperative design results in large fuel gas volume projections, and the relatively

short heat of combustion contact time with the process gases (and the targeted VOC compounds) also requires higher treatment (combustion) temperatures to attain the targeted 95% VOC destruction performance. These factors combine to yield the EPA Cost Control projected fuel gas rates that are the highest for the three TO technologies evaluated.

5.4.2 Regenerative Thermal Oxidization

The regenerative thermal oxidization technology provides operational advantages over recuperative and catalytic technologies, in that the thermal efficiency is greatly improved, thereby reducing the quantity of fuel gas needed to attain a specific treatment temperature. The thermal improvement is typically attained using ceramic media and the heat of combustion from the combustion chamber (in this case supplemental fuel burning) is conducted in multiple and sequencing beds in a series of cycles between pre-heat and heat-recovery of the treatment beds on a regularly cyclical and frequent basis; for example, every couple of minutes. The cycle times are field adjusted for actual conditions and loadings to minimize the output VOC concentrations to the extent possible.

Because of the long contact time with the heat transfer media, the targeted VOC compound destruction is increased, allowing the combustion temperature to be lower than that of the recuperative or catalytic TO counterparts. For example, the 95% destruction can be achieved at a temperature of 1,450°F and with a higher thermal efficiency (typically 95% thermal recovery). The result of this design is fuel gas savings and reduced combustion gas emissions, when compared to other TO technologies. As stated, the fuel needs and typical thermal efficiency are as calculated using the US EPA BACT Cost Control Manual and are also typical of field observations for this technology.

5.4.3 Catalytic Incineration

This technology allows a reactive catalyst to bring the air stream up to the target VOC oxidation temperature. For the catalyst to work properly, the process gas must first be raised to approximately 800°F. Final VOC treatment (destruction) is attained as the reaction of the VOC compounds on the catalytic surface heats the VOC compounds which are then oxidized and the targeted VOC destruction is complete without the need for additional fuel beyond the preheater section. Catalytic TO technologies are evaluated in the US EPA Cost Control tool at 70% thermal efficiency. Not included in this evaluation are the likely needs for special pretreatment filtration to isolate overspray wax from blinding or contaminating the reactive catalyst surface.

Since lower temperatures needed on the front end of the control unit are realized with this technology, fuel gas consumption for this technology are less than that of comparable recuperative TO technologies. To their detriment, catalyst have finite life spans, are subject to catalyst poising from compounds in the airstream, and require periodic replacement and additional maintenance, adding to their higher initial and operating/maintenance costs. The US EPA BACT Cost Control tool accounts for these factors including the initial costs, maintenance costs, catalyst replacement costs and fuel costs for comparative purposes in the BACT demonstration, but does not account for special pre-filtration equipment costs. The Company is concerned this technology may not be fully compatible with the wax overspray but has presented the US EPA values for comparative and full BACT treatment consideration.

5.5 COST/BENEFIT EVALUATION

The specific details of the cost/benefit evaluations conducted are included in **Appendix C**. The lowest cost/benefit VOC control options were tabulated for regenerative thermal oxidizer technology, due to the high thermal efficiency and high rate of VOC treatment efficacy. As described in above, three treatment types were evaluated for cost/benefit determination.

	Dollars per Ton of VOC Treated			
		CPI-Adjusted to 2022		
Thermal Control Option	2006 Dollars	Dollars		
Recuperative Thermal Oxidizer	\$42,261	\$71,032		
Regenerative Thermal Oxidizer	\$10,734	\$18,042		
Catalytic Incinerator	\$19,308	\$32,453		

Table 4. BACT Cost/Benefit Evaluation

Even absent any escalation using 2006 costs, the overall treatment costs results in treatment costs that are excessively high and not economically feasible. The high projected VOC treatment estimates summarized in **Table 4** result from a variety of factors listed below.

- 1) The process discharges large volumes of air which are needed to provide a safe working environment for workers in the foam operation area for each of the three foam seating lines.
- 2) The spray mold release agent and solvent release rates result in low VOC concentrations in the exhaust stream.
- 3) The high air volumes require extensive and high costs for VOC abatement equipment capable of handling 270,000 scfm and also achieving the needed 95 percent destruction efficiency of the VOC compounds at relatively low concentration.
- 4) The VOC abatement equipment will require large volumes of fuel in the form of natural gas to heat the large volume, low VOC content, ambient temperature exhaust stream to the target treatment temperatures for each of the targeted and respective TO devices evaluated.
- 5) The COVID pandemic situation has raised the costs for TO manufacturers and components found in TO equipment. Delays in materials and operating system components have been experienced, along with fabrication and transportation delays.
- 6) Inflation is at a 40-year high level.

The impact and effect of some of the factors listed above have not been included in the BACT evaluation provided, but will result in higher costs than the evaluation presented in the economic portion of the BACT. Since as directed by TDEC, a baseline of zero emissions was used, the costs would be even higher if actual emissions were considered.

5.6 OTHER CONSIDERATIONS

The Pulaski site is located in an area that is designated as attainment for ozone, and VOC is a precursor to ozone. This should reduce the sensitivity or perceived need for VOC abatement which typically raises the "reasonable range" used by the permitting authority to require the use of controls for new VOC projects. It appears the cost/benefit consideration remains higher than reasonable for each of the technologies at each of the adjusted or escalated values used in this demonstration.

The cost projections do not include special retrofit costs since the air permit application seeks the allowed rate of 491 tons per year with a baseline of zero. As a result, actual costs incurred would be higher than projected in the BACT demonstration if the baseline were higher.

The US EPA tools were stretched beyond the reasonable maximum flow rates of 100,000 scfm as if the costs projections and escalations were linear. Experience with large demand systems dictate that replicate and multiple units are needed for such applications due to unit engineering, production, and transportation limitations for the air pollution control unit manufacturer/supplier.

Because the regenerative TO has the lowest cost benefit ratio, a regenerative TO manufacturer was contacted for verification of commercial availability, size and maximum design flow rates. For a system capable of processing 270,000 scfm would require the deployment of four of their largest units which have the individual capability of 70,000 scfm each. This replicated installation would require the costs be calculated on a ¹/₄ basis (one for each regenerative TO unit) and as a result of the replicated fabrication and construction costs, would result in a much higher cost/benefit factor.

The environmental impact of burning natural gas is not put forth in the US EPA tool or into the evaluation and discussion. Oxides of nitrogen, carbon monoxide, and large volumes of carbon dioxide (among other products of combustion) would result if a regenerative TO was required. Adient is very sensitive to its greenhouse gas (GHG) impacts and such a system would increase both the consumption of fossil fuel (in the form of natural gas), increase the release of other air pollutants that are products of combustion, and result in the release of large quantities of GHGs.

5.7 FINDINGS AND CONCLUSIONS

A query from the RACT/BACT/LAER Clearinghouse (RBLC) was conducted for flexible foam production and only one record was returned for a facility in Indiana (see **Appendix D**), which used regenerative thermal oxidization controls. No details were provided in the RBLC listing for the volume of air, the number of lines or for the dollars per ton of treatment. Adient was able to confirm from the RBLC listing that the technology used for the Indiana site is the same technology (regenerative thermal oxidation) that yielded the lowest treatment costs summarized in Table 3 above. Since no direct comparison of process specific details from the RBLC listing could be made to the Pulaski operation, Adient is relying on the control options identified in its BACT demonstration to determine the best available control for this application.

Typically, VOC control strategies considered BACT have a cost/benefit in the range of \$5,000 to \$6,000 per ton of treatment. The best available control demonstrations for this application yield costs that are far in excess of this range, with the lowest cost technology being the regenerative thermal oxidizer. Projected costs and processing equipment yields annual operating and control benefits far in excess of what would otherwise be considered cost effective for VOC controls. Further, these figures also do not include retrofitting costs which would make the costs per ton of treatment even higher. On that basis, Adient concludes and asserts the current configuration without VOC abatement is BACT for this project and for the existing process.

6.0 AIR QUALITY ANALYSIS

An air quality analysis for VOC is required for the proposed project because the net emission increase exceeds the PSD SER for VOC (40 tpy). The air quality analysis involves demonstrating compliance with National Ambient Air Quality Standards (NAAQS) and PSD increment standards, as well as a Class I Area impact analysis. There are no established National Ambient Air Quality Standards (NAAQS) or PSD increment standards for VOC; however, VOC is considered a precursor emission to ozone. As such, a secondary impact analysis is required for VOC emissions in relation to secondary ozone formation.

6.1 OZONE ANALYSIS

6.1.1 Preconstruction Monitoring

Since the project's net emission increase is greater than 100 tpy for VOC, preconstruction monitoring must be addressed for ozone. Per US EPA's *Ambient Monitoring Guidelines for PSD*, data from an existing monitor may be used if it is considered representative of the facility location. When determining if data is representative of the site, location, quality of data, and the currentness of the data must be considered.

Ozone is considered a regional pollutant; therefore, possible chemical reactions that could influence ambient concentrations must also be considered.

Adient believes that the regional ozone monitor located at the Fairview Middle School (FMS) in Fairview, Tennessee is representative of the project site. The FMS monitor is located approximately 80 kilometers (km) north of Adient's Pulaski facility. Fairview and Pulaski have similar populations of approximately 8,700 and 7,600, respectively. For ozone, a large component of background concentration can be attributed to vehicle use. With similar sized populations, vehicle use can be expected to occur at a similar rate. The two areas also have similar terrain and land use.

Design value data for the FMS monitor is available for the previous ten years (2011 – 2020). FMS monitor information and data for 2020 is provided below. Adient is requesting that TDEC waive the preconstruction monitoring requirement based on the availability of representative data from the FMS ozone monitor.

Monitor Location	Monitor ID	2020 Design Value (ppb)
Fairview Middle School, Crow Cut Road, Fairview,	471870106	60

Table 5. Representative Ozone Monitor

6.1.2 Secondary Impact Analysis

TN 37062

There is no established NAAQS for VOC; however, VOC is a precursor to ozone, which does have an established NAAQS. US EPA has developed a two-tiered evaluation for secondary ozone formation. The Tier I analysis allows for a qualitative assessment based on existing modeling studies performed by US EPA. The Tier I approach involves comparison of facility emissions to a Modeled Emission Rate for Precursors (MERPs) developed by US EPA. TDEC's guidance document, *Tennessee Guidance on the Use of EPA's MERPs to Account for Secondary Ozone and Fine Particulate in Tennessee Under the New Source Review PSD Program (TDEC MERPs Guidance),* was utilized to assess if further analysis is required for precursor emissions.

To evaluate compliance, a Tier 1 demonstration using the *TDEC MERPs Guidance* and suggested default values was completed. A comparison of Adient's VOC emissions to TDEC's default MERPs for VOC is shown in the table below.

Pollutant	Facility Emissions⁵ (tpy)	TDEC Default MERPs Value (tpy)	Emissions Below MERPs?
VOC	491.4	1,542	Yes

Table 6. MERPs /

In accordance with the *TDEC MERPs Guidance*, a Tier I demonstration is successful if the facility emissions are less than the default MERPs value. As demonstrated above, Adient's emissions are below the TDEC

⁵ Adient has voluntarily agreed to limit facility-wide VOC emissions to 491.4 tpy.

default MERPs value for ozone; therefore, no further analysis is required, and no adverse secondary impacts for ozone are expected to occur.

6.2 CLASS I AREA ANALYSIS

Class I areas are provided the most protection under the PSD program, ensuring protection to natural, scenic, recreational, and/or historical areas. TDEC requires sources to evaluate impacts to Class I areas located within 300 km of the project site. There are five Class I areas within 300km of Adient – Sipsey National Wilderness Area (~ 97 km), Cohotta Wilderness Area (~218 km), Mammoth Cave National Park (~ 226 km), Joyce Kilmer-Slickrock Wilderness Area (~278 km), and Great Smoky Mountain National Park (~281 km).

A Class I area analysis includes a Class I PSD increment assessment for pollutants subject to PSD review (increasing above the SER) and an Air Quality Related Values (AQRV) analysis for visibility, ozone, and deposition that could impact a Class I area's resources.

6.2.1 Class I Increment Analysis

Adient does not have a significant increase of $PM/PM_{10}/PM_{2.5}$, and there is no Class I PSD increment established for VOC. As such, a Class I increment analysis is not required.

6.2.2 Class I AQRV Analysis

The Class I AQRV analysis was prepared in accordance with the Federal Land Manager's (FLM's) Air Quality Related Values Work Group (FLAG) *Phase I Report – Revised (2010)*. The FLMs developed a screening criteria involving facility emissions and distance to the Class I area (Q/D ratio) for sources greater than 50 km from a Class I area to determine whether adverse impacts could occur to AQRV from a pollutant. The equation used is provided below:

Facility Emissions (Q) Distance to Class I Area (D)

If the Q/D ratio is below 10, it is presumed that no adverse impact will occur, and no further AQRV analysis is required. An AQRV analysis for the Class I Areas of concern can be found below.

Class I Area	Pollutant	Q (tpy)	D (km)	Q/D
	PM	9.90	97	0.10
Sipsey Wilderness Area	VOC	491.4	9/	5.07
Cohotta Wilderness	PM	9.90	010	0.05
Area	VOC	491.4	218	2.25
Mammoth Cave	PM	9.90	224	0.04
National Park	VOC	491.4	226	2.17
Joyce Kilmer-Slickrock	PM	9.90	070	0.04
Wilderness Area	VOC	491.4	278	1.77
Great Smoky Mountain	PM	9.90	001	0.04
National Park	VOC	491.4	281	1.75

Table 7. Class I AQRV Analysis

The Q/D ratio for PM and VOC in each of the Class I Areas are below the threshold of 10; therefore, it is presumed there are no adverse impacts from Adient, and no further analysis is required. .

7.0 ADDITIONAL IMPACTS ANALYSIS

PSD applies to new major sources or major modifications at existing sources for pollutants where the area the source is located is in attainment or unclassifiable with the NAAQS. Adient is a major source of VOCs, a precursor to ozone. Adient is located in the City of Pulaski, County of Giles in the State of Tennessee, which is designated attainment for ozone.

A PSD major source subject to PSD review is required to conduct an air quality analysis and an additional impacts analysis, among other requirements. Pursuant to 40 CFR §52.21(o), the additional impacts analysis consists of three parts: Growth Analysis, Soils and Vegetation Impacts Analysis and Visibility Impairment Analysis. Each of these analyses is addressed below.

7.1 GROWTH ANALYSIS

The Adient plant is located in an industrial park setting on the north side of Pulaski, Tennessee. Pulaski is located in the south central portion of the state nearly equidistant between Memphis and Chattanooga. The general vicinity outside of the industrial development is mainly agricultural pastureland and woodland with some residential development east of the plant (**Figure 4**). The workforce consists of 305 employees living within commuting distance of the plant. The size of the workforce has resulted in little impact on population growth in Giles County, which has a population of 29,503 as of 2018. A review of historical aerial photos dating back to 1998, indicate that there has been little industrial or commercial development in the immediate vicinity of the Adient plant, and no substantive residential growth in the general area.

7.2 SOIL AND VEGETATION ANALYSIS

7.2.1 Particulate Matter and Volatile Organic Compounds

The criteria for evaluating impacts on soils and vegetation is taken from EPA's, A Screening Procedure for the Impacts of Air Pollution Sources on Plants, Soils, and Animals, EPA, 1980. According to US EPA, "...sources more than 10 km from any Class I areas, exemptions provide that no analysis of impairment need be done if emission increases are below specified limits." Specified limits are de minimis values found at 40 CFR 52.21(b)(23)(i).

The Adient operations are located more than 10 km from any Class I areas in the region. The criteria air pollutants emitted by the Adient Pulaski operations include particulate matter (PM) and PM_{10} . Emissions of PM and PM_{10} are below the significance levels (de minimis values) found at 40 CFR 52.21(b)(23)(i). The Adient operations generating particulate matter emissions are not the direct result of combustion or combustion byproducts and therefore, Adient assumes there are no emissions of $PM_{2.5}$, but in any event, all PM emitted by the source is less than the respective thresholds, including the threshold for $PM_{2.5}$.

Adient emits VOC at an annual rate above the de minimis value of 40 tpy, but screening concentrations are not available for VOC. VOC is a precursor to ozone and while ozone is identified in Table 2.1 as a "Regulated Pollutant", the EPA screening guide indicates that "a screening concentration [for O_3 is] available, but no simple procedure for estimating the ozone impact of a single source is currently available." Additionally, the secondary NAAQS were established at concentration levels below which no harmful effects to either soil or vegetation is expected.⁶ As discussed in Section 6.1.2, EPA has developed a two-tiered evaluation for

⁶ US EPA, Office of Air Quality Planning and Standards, *New Source Review Workshop Manual (Draft)*, U.S. EPA, Research Triangle Park, NC, October, 1990.

secondary ozone formation from VOC. As demonstrated by this application, the VOC emissions from Adient's facility are below the default TDEC MERPs value, indicating that no adverse impact to compliance with the ozone NAAQS is expected. As such, VOC emissions from the facility will not negatively affect soil and vegetation in the surrounding area.

Other than VOC, Adient does not emit any criteria air pollutants above their respective significance thresholds.

7.3 VISIBILITY IMPAIRMENT ANALYSIS

US EPA prescribes the use of its *Workbook for Plume Visual Screening and Analysis (Revised)*, October 1992 (EPA-454/R92-023), methodologies for purposes of conducting a visibility impairment analysis. A visibility impairment analysis is generally required to determine the impact on sensitive areas such as state parks, wilderness areas, airports, scenic sites and overlooks. Three levels of screening procedures are outlined by US EPA. If the criteria for the first screening level, the most conservative level, are met, no further analysis is needed.

The VISCREEN model is recommended for the Level 1 screen. The VISCREEN model primarily considers NO2 and particulate emission increases associated with a project. VISCREEN does not consider or calculate visibility impacts from ozone.

7.3.1 Level 1 Screen Analysis

Level 1 analysis incorporates conservative parameters to determine plume impacts. Default values for particle size and density, and a default of worst-case meteorological condition of F stability and 1.0 meters per second (m/s) wind speed are used for the analysis, while all emissions are assumed to exit the plant from one point. The worst-case meteorological condition is expected to persist for 12 hours with a wind direction that would transport the plume directly to the sensitive area being analyzed.

A maximum particulate matter emission rate of 2.64 lb/hour, based on a maximum emission rate of 9.9 tpy assuming an operating schedule of 7,488 hours/year (6 days per week, 24 hours per day, 52 weeks/year), was input to the model. It was assumed that there were no emissions of NOx, soot, primary nitrogen dioxide and primary sulfate. A background visual range of 25 kilometers was used. All other inputs relied on the default parameters.

While the Pulaski facility is greater than 10 km from any Class I area (**Figure 5**), a distance of 10 km was assumed in order to run the most conservative analysis of visibility impairment.

Level 1 modeling results indicate that the Adient operations do not adversely impact visibility within or beyond a 10 km radius.

OVERALL RESULTS OF PLUME VISIBILITY SCREENING
SOURCE: Adient US LLC, Pulaski, CLASS I ARFA: NA
INSIDE class I area
Plume delta E DOES NOT EXCEED screening criterion for SKY background Plume delta E DOES NOT EXCEED screening criterion for TERRAIN background
Plume contrast DOES NOT EXCEED screening criterion for SKY background Plume contrast DOES NOT EXCEED screening criterion for TERRAIN background
OUTSIDE class I area
Plume delta E DOES NOT EXCEED screening criterion for SKY background Plume delta E DOES NOT EXCEED screening criterion for TERRAIN background
Plume contrast DOES NOT EXCEED screening criterion for SKY background Plume contrast DOES NOT EXCEED screening criterion for TERRAIN background
SCREENING CRITERIA: DELTA E = 2.0
GREEN CONTRAST = 0.050

8.0 TDEC CONSTRUCTION PERMIT APPLICATION FORMS

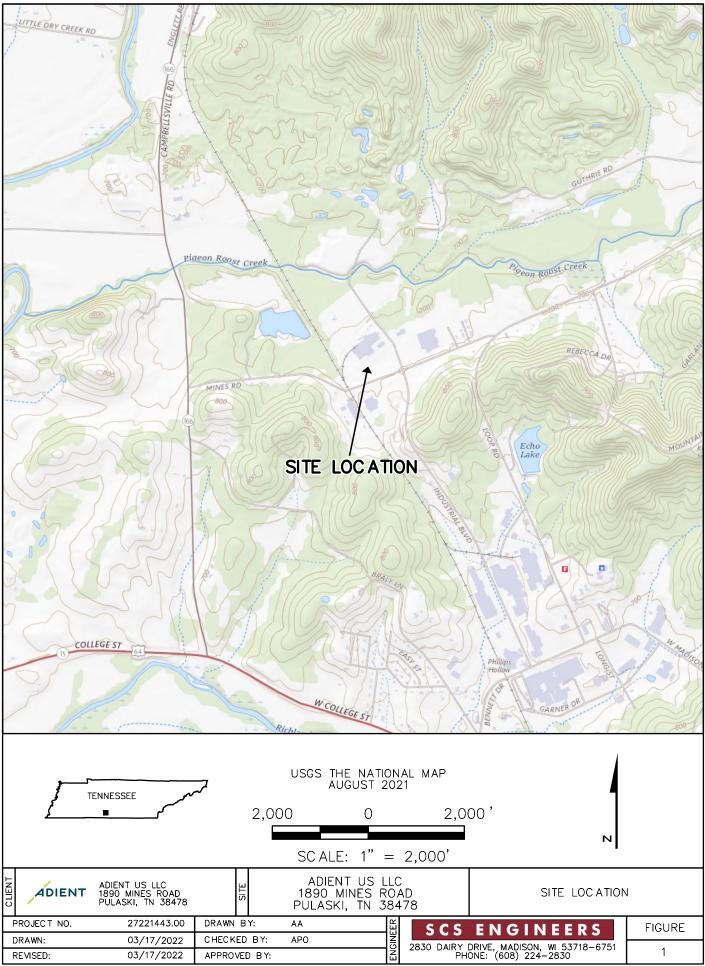
TDEC construction permit application forms are included in Appendix E.

- APC 1 Identification
- APC 2 Operations and Flow Diagrams
- APC 3 Stack Identification
- APC 10 Miscellaneous Processes
- APC 19 Compliance Certification Monitoring and Reporting
- APC 22 Compliance Demonstration by Monitoring Control System Parameters or Operating Parameters of a Process
- APC 26 Compliance Demonstration by Recordkeeping
- APC 29 Emission Summary for the Facility
- APC 30 Current Emissions Requirements and Status
- APC 31 Compliance Plan and Compliance Certification

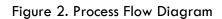
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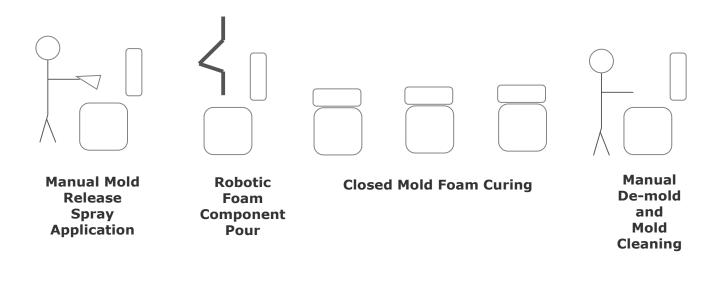
Figures

- Site Location Map 1
- 2
- Process Flow Diagram Stack Exhaust Flow Rates 3
- 4 Aerial View
- Map Visibility Impairment Analysis 5

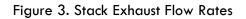


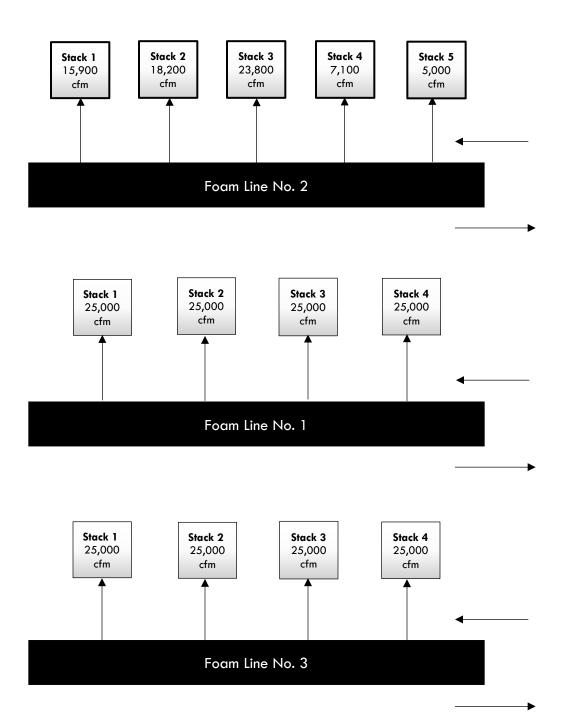
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Foam Line Processing Step Diagram

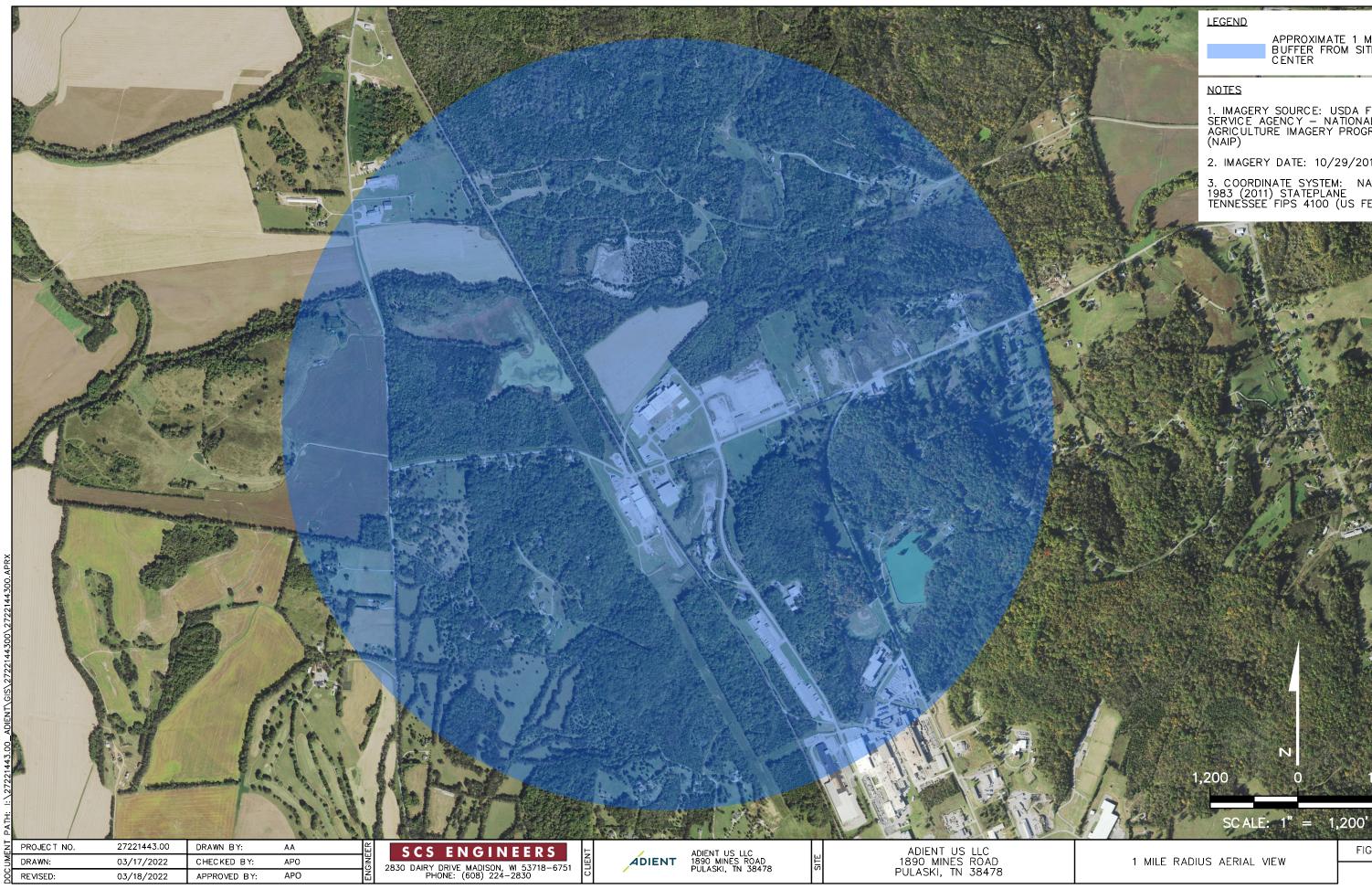




	Exhaust Flow Rate	Height ¹	Diameter	Temperature	Moisture Content	Configuration	Direction
Stack						Round,	Up, Down,
ID No.	scfm	ft	ft	°F	%	Rectangular	Horizontal
1	15,900	80'	3' 2"	70	<1%	Round	Up
2	18,200	90'	3' 6"	70	<1%	Round	Up
3	23,800	96'	3' 10"	70	<1%	Round	Up
4	7,100	61'	2' 1"	70	<1%	Round	Up
5	5,000	58'	2'	70	<1%	Round	Up
6	25,000	10' 6"	5' 6"	70	<1%	Round	Up
7	25,000	10' 6"	5' 6"	70	<1%	Round	Up
8	25,000	10' 6"	5' 6"	70	<1%	Round	Up
9	25,000	10' 6"	5' 6"	70	<1%	Round	Up
10	25,000	10' 6"	5' 6"	70	<1%	Round	Up
11	25,000	10' 6"	5' 6"	70	<1%	Round	Up
12	25,000	10' 6"	5' 6"	70	<1%	Round	Up
13	25,000	10' 6"	5' 6"	70	<1%	Round	Up

Figure 3. Stack Parameters (Cont'd)

¹ Height above roof; roof height is 35'.



APPROXIMATE 1 MILE BUFFER FROM SITE CENTER

1. IMAGERY SOURCE: USDA FARM SERVICE AGENCY – NATIONAL AGRICULTURE IMAGERY PROGRAM (NAIP)

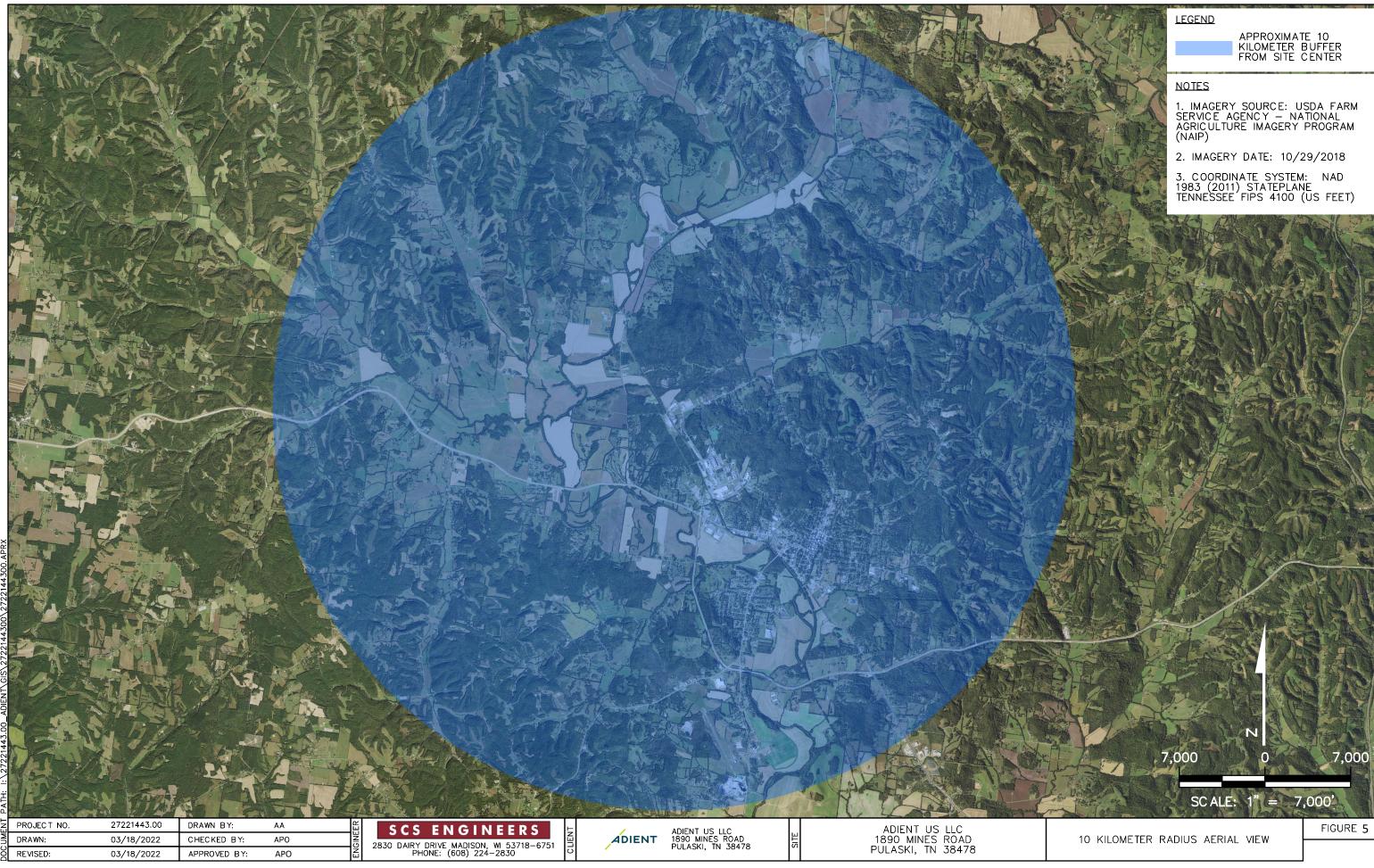
2. IMAGERY DATE: 10/29/2018

3. COORDINATE SYSTEM: NAD 1983 (2011) STATEPLANE TENNESSEE FIPS 4100 (US FEET)

FIGURE 4

1,200

N





Appendix A

Safety Data Sheets

Appendix B

Emission Calculations

Table 1. Potential to Emit - Volatile Organic Compounds (VOC) Emissions Adient US LLC

FID 28-0076-01

Source:	Foam	Production -	General	Process
SCC:	3080	1005		

			Potential to Emit ²									
			Max	cimum Raw Ma	aterial Usage	Rate ³	Volatile Organic Compounds (VOC)					
	-	Material Usage Ite ¹	Density	Usage		Usage	Content				Emissions	
Raw Material	gal/mo	gal/yr	lb/gal	gal/mo	mo/yr	gal/yr	lb VOC/gal	E mission Factor	Unit	lb/hr	lb/yr	tpy
Mold Release Agent ⁴	8,389	100,668	6.30	13,295	12	159,540	6.15	6.16	lb VOC/gal	112	982,766	491.38
Paste Wax	65	780	6.84	104	12	1,248	5.15	5.15	lb VOC/gal	0.73	6,427	3.21
Tegostab B 8737 LF (Silicon)	603	7,236	8.16	965	12	11,578	-	1.5%	% by wt	0.16	1,417	0.71
Tegostab B 8724 LF2 (Surfactant)	1,206	14,472	7.90	1,930	12	23,155	0.00	0.00	lb VOC/gal	0	0	0.00
Glycerine	1,541	18,492	10.52	2,466	12	29,587	0.50	0.50	lb VOC/gal	1.69	14,794	7.40
DL-02596 Black	563	6,756	9.82	901	12	10,810	0.00	0.00	lb VOC/gal	0	0	0.00
Jeffcat -2F20	240	2,880	7.09	384	12	4,608	0.00	0.00	lb VOC/gal	0	0	0.00
Dabco 33 LX	506	6,072	9.31	810	12	9,715	0.00	0.00	lb VOC/gal	0	0	0.00
YUKOL - 8331	505	6,060	8.72	808	12	9,696	0.00	0.00	lb VOC/gal	0	0	0.00
BIOH 2828	3,053	36,636	8.40	4,885	12	58,618	0.00	0.00	lb VOC/gal	0	0	0.00
Multranol 3901	9,605	115,260	8.42	15,368	12	184,416	0.00	0.00	lb VOC/gal	0	0	0.00
Polyol (E833)	73,188	878,256	8.55	117,101	12	1,405,210	0.00	0.00	lb VOC/gal	0	0	0.00
Polyol (E855)	89,843	1,078,116	8.84	143,749	12	1,724,986	0.00	0.00	lb VOC/gal	0	0	0.00
Voranol 360 Polyol DA	7,038	84,456	9.04	11,261	12	135,130	0.00	0.00	lb VOC/gal	0	0	0.00
Voranate TDI T-80 (TDI)	45,385	544,620	10.17	72,616	12	871,392	10.09	3.29E-05	lb/lb ⁵	0.0333	291.70	0.1458
Diphenylmethane diisocyanate (MDI)	4,080	48,960	10.09	6,528	12	78,336	10.00	9.39E-06	lb/lb ^{5,6}	0.0008	7.42	0.0037
Diethanolamine (DEOA)	1,232	14,784	9.09	1,971	12	23,654	7.78	7.94E-06	lb/lb ⁷	0.0002	1.71	0.0009
Total	247,042	2,964,504		395,140		4,741,678				114.81	1,005,705	502.85

8,760 hr/yr

Footnotes

1. Average raw material usage rates are based on recent semi-annual emissions reporting data (October 2020 through September 2021).

2. Potential to emit is based on worst-case conservative assumptions; Adient will voluntarily accept a facility-wide VOC limit of 491.4 tpy.

3. With the exception of the mold release agent, maximum usage rates are based on highest monthly usage rate of each respective raw material for the period of October 2020 through September 2021 multiplied by a factor of 1.60 (491 tpy proposed VOC emission rate/308 tpy current allowable VOC emission rate).

4. Maximum usage rate of mold release agent is based on a worst-case conservative assumption allowing for potential growth in production rates from customer requirements.

5. TDI and MDI emission factors are based on a source test conducted on October 1, 1997 (see Title V permit renewal application, 2014).

6. All diphenylmethane diisocyanate is assumed to be methylene diphenyl diisocyanate (MDI); historically assumed.

7. Emission factors are based on emission factors used in the 2014 Title V permit renewal application, and semi-annual/12-month emission reports.

Source

Safety Data Sheets

Table 2. Particulate Matter Emissions Adient US LLC

FID 28-0076-01

Source: Foam Production - General Process SCC: 30801005

	Potential to Emit							
	Maximum Product Usage Rate Particulate Matter Emissio				issions ¹			
					Transfer			
				Content ²	Efficiency ³	Emissions		
Process	lb/mo	mo/yr	lb/yr	wt %	%	lb/hr	lb/yr	tpy
Mold release	85,400	12	1,024,800	4.10%	55.00%	2.16	18,908	9.45

Operation

8,760 hr/yr

Footnotes

1. Assume PM is equivalent to PM10; PM2.5 is not an anticipated air pollutant emission, but assumed under conservative worst-case PTE calculations.

2. Safety Data Sheet: Mold Release Agent_Chem Trend_PU-11434

3. Transfer efficiency is a conservative assumption; industry standards for spray gun efficiency is 65%.

Table 3. Potential to Emit - Hazardous Air Pollutant (HAP) Emissions Adient US LLC FID 28-0076-01

Source: Foam Production - General Process SCC: 30801005

		Potential to Emit								
	N	Naximum Prod	uct Usage Rat	e		Hazardous Air Pollutants (HAPs)				
	Density	Usage		Usage	Content ²				Emissions	
Raw Material	lb/gal	gal/mo	mo/yr	gal/yr	lb VOC/gal	Emission Factor	Unit	lb/hr	lb/yr	tpy
TD-80 (TDI)	10.17	72,616	12	871,392	10.09	3.29E-05	lb/lb ¹	0.0333	291.70	0.1458
Diphenylmethane diisocyanate (MDI)	10.09	6,528	12	78,336	10.00	9.39E-06	lb/lb ^{1,2}	0.0008	7.42	0.0037
Diethanolamine (DEOA)	9.09	1,971	12	23,654	7.78	7.94E-06	lb/lb ³	0.0002	1.71	0.0009
Total								0.03	301	0.15



Footnotes

1. TDI and MDI emission factors are based on a source test conducted on October 1, 1997 (see Title V permit renewal application, 2014).

2. All diphenylmethane diisocyanate is assumed to be methylene diphenyl diisocyanate (MDI); historically assumed.

3. Emission factors are based on emission factors used in the 2014 Title V permit renewal application, and semi-annual/12-month emission reports.

Source

Safety Data Sheets

Appendix C

BACT Analysis – Backup Calculations

DATA INPUTS

ATA INPUTS		
Company Name	Adient	
Location	Pulaski, TN	
Process Name	Spray Mold Release 3 Foan	n Lines
Annual Hours of Operation	8,760	hrs/yr
Process Exhaust Flow Rate	270,000	scfm
VOC Capture Efficiency	80	%
VOC Control Efficiency	95	%
Annual VOC Emissions (before controls)	491.4	tons/yr
Natural Gas Rate	7.80	\$/1000ft ³
Electricity Rate	0.083	\$/kwh
VOC Heating Value	21,095	BTU/lb
Process Exhaust Temperature	80	°F
2006 Dollars CPI Recuperative Oxidizer	146.4	
2006 Dollars CPI Regenerative Oxidizer	139.0	
2006 Dollar CPI Catalytic Oxidizer	217.4	
Date of Vatavuk Cost Index	3rd Quarter 2006	
Operating Labor Rate	75	\$/hr
Annual Interest Rate	7	%
Control System Life	20	years
Cost of Permanent Total Enclosure (\$)	0	

DATA OUTPUTS

Process Exhaust Gas Heat Content Annual VOC Emissions Controlled		0.146 BTU/scf 373.464 tons			
Annual Costs of Control Device	vice 2006 dollars				PI adjusted to 22 dollars
Recuperative	\$	42,261	\$/ton VOC Controlled	\$	71,032
Regenerative	\$	10,734	\$/ton VOC Controlled	\$	18,042
Catalytic	\$	19,308	\$/ton VOC Controlled	\$	32,453
Minimum	\$	10,734	\$/ton VOC Controlled	\$	18,042

BACT Cost-Benefit Evaluation

Company Name: Adient Location: Pulaski, TN Process: Spray Mold Release 3 Foam Lines

TOTAL ANNUAL COST SPREADSHEET PROGRAM -- RECUPERATIVE THERMAL OXIDIZERS

Describes the annual operating costs for purchasing, installing and operating a recuperative thermal oxidizer to control the above process.

COST BASE DATE: April 1988 [1]

VAPCCI	[2]	3rd Quarter 2006	146.4
--------	-----	------------------	-------

INPUT PARAMETERS

Gas flowrate (scfm):	270000
Reference temperature (oF):	77
Inlet gas temperature (oF):	80
Inlet gas density (lb/scf):	0.0739
Primary heat recovery (fraction):	0.70
Waste gas heat content (BTU/scf):	0.146
Waste gas heat content (BTU/lb):	1.98
Gas heat capacity (BTU/lb-oF):	0.255
Combustion temperature (oF):	1600
Preheat temperature (oF):	1144
Fuel heat of combustion (BTU/lb):	21502
Fuel density (lb/ft3):	0.0408

DESIGN PARAMETERS

Auxiliary Fuel Reqrmnt (lb/min):	145.275
(scfm):	3560.7
Total Gas Flowrate (scfm):	273561

3/28/2022

CAPITAL COSTS

Equipment Costs (\$): Incinerator:	
<pre> Incinerator: @ 0 % heat recovery:</pre>	0
@ 35 % heat recovery:	0
0 50 % heat recovery:	0
@ 70 % heat recovery:	488,088
PTE Containment or other capital costs	100,000
Total Equipment Costbase:	488,088
' 'escalated:	893,766
Instrumentation:	0
Sales Tax:	26,813
Freight:	44,688
Purchased Equipment Cost (\$):	1,054,644
Direct Installation Costs:	
Foundations & Supports:	84,372
Handling & Erection:	147,650
Electrical:	42,186
Piping:	21,093
Ductwork and Insulation:	10,546
Painting:	10,546
Direct Installation Cost:	316,393
Site Preparation:	0
Buildings or PTE:	0
Total Direct Cost:	1,371,038

BACT Cost-Benefit Evaluation

Indirect Installation Costs: Engineering:	105,464	
Field Expenses:	52,732	
Contractor Fees:	105,464	
Start-Up:	21,093	
Performance Test:	10,546	
Contingencies:	31,639	
2	·	
Total Indirect Cost:	326,940	
Total Capital Investment (\$):	1,697,977	
ANNUAL COST INPUTS		
Operating factor (hr/yr):	8760	
Operating labor rate (\$/hr):	75.00	
Maintenance labor rate (\$/hr):	82.50	
Operating labor factor (hr/sh):	0.5	
Maintenance labor factor (hr/sh):	0.5	
Electricity price (\$/kwh):	0.083	
Natural gas price (\$/mscf):	7.80	
Annual interest rate (fraction):	0.070	
Control system life (years):	20	
Capital recovery factor:	0.0944	
Taxes, insurance, admin. factor:	0.04	
Pressure drop (in. w.c.):	19.0	

ANNUAL COSTS

Item	Cost (\$/yr)	Wt. Factor	W.F.(cond.)
Operating labor	41,063	0.003	
Supervisory labor	6,159	0.000	
Maintenance labor	45,169	0.003	
Maintenance materials	45,169	0.003	
Natural gas	14,597,540	0.925	
Electricity	736,981	0.047	
Overhead	82,536	0.005	0.014
Taxes, insurance, administrative	67,919	0.004	
Capital recovery	160,277	0.010	0.014
Total Annual Cost	15,782,811	1.000	1.000

[1] Original equipment costs reflect this date. [2] VAPCCI = Vatavuk Air Pollution Control Cost Index (for thermal incinerators) corresponding to year and quarter shown. Original equipment cost, purchased equipment cost, and total capital investment have been escalated to this data via the VAPCCI and control equipment vendor data. Latest indexes included herein.

RACT Cost Summary Table	2006 Costs for 270000 scfm system		U.S. Bureau of Labor Statistics Correction Factor - 2006 to 2022 CPI Inflation Calculator
-		2022 dollars	1.6808
1 Purchased Equipment Cost (PEC)	1,054,644	1,772,646	
2 Total Direct Cost (includes PEC)	1,371,038		
3 Total Indirect Cost	326,940		
4 Total Capital Investment (= 2+3)	1,697,977	2,853,960	
5 Annual Direct Operating Costs	15,472,080	26,005,471	
6 Annual Indirect Operating Costs	150,455	252,884	
7 Annual Capital Recovery Costs	<u>160,277</u>	269,394	
8 Total Annual Costs (= 5+6+7)	15,782,811	26,527,749	
	2006 dollars		
Oxidizer VOC Control Efficiency	95 %		
Annual VOC Input to the Control Device	393.12 tons		
Annual VOC Emissions Controlled	373.464		
Annual VOC Emissions after Controls	19.656	2022 dolars	
Annual Cost of Control Device	\$ 42,261 \$/ton Controlle	e 71,031.61	

BACT Cost-Benefit Evaluation

Company Name: Adient Location: Pulaski, TN Process: Spray Mold Release 3 Foam Lines

TOTAL ANNUAL COST SPREADSHEET PROGRAM--REGENERATIVE THERMAL OXIDIZER (RTO)

Describes the annual operating costs for purchasing, installing and operating a regenerative thermal oxidizer to control the above process.

COST BASE DATE: December 1988 [1]

VAPCCI [2]	3rd Quarter 2006	139.0	

INPUT PARAMETERS

Gas flowrate (scfm):	270000
Reference temperature (oF):	77
Inlet gas temperature (oF):	80
Inlet gas density (lb/scf):	0.0739
Primary heat recovery (fraction):	0.95
Waste gas heat content (BTU/scf):	0.146
Waste gas heat content (BTU/lb):	1.98
Gas heat capacity (BTU/lb-oF):	0.255
Combustion temperature (oF):	1450
Heat loss (fraction):	0.01
Exit temperature (oF):	149
Fuel heat of combustion (BTU/lb):	21502
Fuel density (lb/ft3):	0.0408

DESIGN PARAMETERS

Auxiliary Fuel Requirement (lb/min):		17.642
	(scfm):	432.4
Total Gas Flowrate (scfm):		270432

TOTAL CAPITAL INVESTMENT (\$) [3] (Cost correlations range: 5000 to 500,000 scfm)

PTE Containment or other capital costs	0
@ 85 % heat recoverybase:	0
' 'escalated:	0
@ 95 % heat recoverybase:	6,755,111
' 'escalated:	10,269,247

ANNUAL COST INPUTS

Operating factor (hr/yr):	8760
Operating labor rate (\$/hr):	75.00
Maintenance labor rate (\$/hr):	82.50
Operating labor factor (hr/sh):	0.50
Maintenance labor factor (hr/wk):	1.00
Electricity price (\$/kwh):	0.083
Natural gas price (\$/mscf):	7.80
Annual interest rate (fraction):	0.070
Control system life (years):	20
Capital recovery factor:	0.0944
Taxes, insurance, admin. factor:	0.04
Pressure drop (in. w.c.):	20.0

ANNUAL COSTS

Item	Cost (\$/yr)	Wt. Factor	W.F.(co	nd.)
Operating labor	41,063	0.010		
Supervisory labor	6,159	0.002		
Maintenance labor	4,290	0.001		
Maintenance materials	4,290	0.001		
Natural gas	1,772,684	0.442		
Electricity	766,841	0.191		
Overhead	33,481	0.008		0.022
Taxes, insurance, administrative	410,770	0.102		
Capital recovery	969,344	0.242		0.344
Total Annual Cost	4,008,922	1.000		1.000

BACT Cost-Benefit Evaluation

[1] Base total capital investment reflects this date.
[2] VAPCCI = Vatavuk Air Pollution Control Cost Index (for regenerative thermal oxidizers) corresponding to year and quarter shown. Base total capital investment has been escalated to this date via VAPCCI and control equipment vendor data. Latest indexes included herein.
[3] Source: Vatavuk, William M. ESTIMATING COSTS OF AIR POLLUTION CONTROL. Boca Raton, FL: Lewis Publishers, 1990.

COMPARISON OF REECO/DUPONT, CO\$T-AIR, AND MANUAL RTO COSTS: (1st Qtr. '91 \$)

Flow (scfm)	REECo (\$)	Manual (\$) [a]	Manual/REECo	CO\$T-AIR (\$) [b	C-A/REECo
2,000	340,000	371,061	1.09	640,305	1.88
5,000	425,000	423,946	1.00	713,363	1.68
10,000	500,000	512,087	1.02	835,125	1.67
25,000	850,000	776,511	0.91	1,200,413	1.41
50,000	1,500,000	1,217,217	0.81	1,809,225	1.21
100,000	2,850,000	2,098,629	0.74	3,026,850	1.06

[a] Escalated from April. '88 to 1st quarter '91 and multiplied by installation factor of 1.416 (1.2*1.18). Range of correlation: 10,000 to 100,000 scfm.

[b] Escalated from Dec. '88 to 1st quarter '91. Costs pertain to 95% heat recovery units. Range of correlation: 5,000 to 500,000 scfm.

	Costs for 270000		U.S. Bureau of Labor Statistics Correction Factor - 2006 to 2022 CPI Inflation
RACT Cost Summary Table for RTO	<u>scfm system</u>	<u>2022 Costs</u>	Calculator
1 Total Capital Investment	#########	17,260,550	1.6808
2 Annual Direct Operating Costs 3 Annual Indirect Operating Costs 4 Annual Capital Recovery Costs 5 Total Annual Costs (= 2+3+4)	2,595,327 444,251 <u>969,344</u> 4,008,922	4,362,226 746,697 1,629,274 6,738,197	
Oxidizer VOC Control Efficiency Annual VOC Input to the Control Device Annual VOC Emissions Controlled Annual VOC Emissions after Controls	95 % 393.12 tons 373.464 19.656		
Cost of Control Device	\$ 10,734 \$/ton Controlled	\$18,042]

Company Name: Adient Location: Pulaski, TN Process: Spray Mold Release 3 Foam Lines

TOTAL ANNUAL COST SPREADSHEET PROGRAM--CATALYTIC INCINERATORS (FIXED)

Describes the annual operating costs for purchasing, installing and operating a Catalytic Oxidizer to control the above process.

COST REFERENCE DATE: April 1988 [1]

VAPCCI [2]	3rd Quarter 2006		217.4
	IN	IPUT PARAMETERS	
Waste gas hea Waste gas hea Gas heat capa Combustion te Preheat tempe	nperature (oF): erature (oF): ity (lb/scf): ecovery (fraction): at content (BTU/scf): at content (BTU/lb): acity (BTU/lb-oF): emperature (oF): erature (oF): ombustion (BTU/lb):		270000 77 80 0.0739 0.70 0.146 1.98 0.248 650 479 21502 0.0408

DESIGN PARAMETERS

Auxiliary Fuel Reqrmnt (lb/min):		51.173
	(scfm):	1254.3
Total Gas Flowrate (scfm):		271254
Catalyst Volume (ft3):		525.3

CAPITAL COSTS

Equipment	: Costs (\$):
-----------	-----------	------

-- Incinerator:

@ 0 % heat recovery:

0

	@ 35 % heat recovery:@ 50 % heat recovery:	0 0
	@ 70 % heat recovery:	1,453,093
Other (auxiliary		0
Total Equipment C		1,453,093
' 'esca	llated:	3,414,836
	Instrumentation	341,484
	Sales Tax	102,445
	Freight	170,742
Purchased Equ	ipment Cost (\$):	4,029,506
Direct Installation	Costs:	
	Foundation & Supports	322,361
	Handling & Erection	564,131
	Electrical	161,180
	Piping	80,590
	Ductwork & Insulation	40,295
	Painting	40,295
	Buildings or PTE:	0
	Total Direct Cost:	5,238,358
Indirect Installation	n Costs:	
	Engineering	402,951
	Field Expenses	201,475
	Contractor Fees	402,951
	Start-Up	80,590
	Performance Test	40,295
	Contingencies	120,885
	Total Indirect Costs:	1,249,147
Total Capital	Investment (\$):	6,487,505

ANNUAL COST INPUTS

Operating factor (hr/yr):

8760

Operating labor rate (\$/hr):	75.00
Maintenance labor rate (\$/hr):	82.50
Operating labor factor (hr/sh):	0.5
Maintenance labor factor (hr/sh):	0.5
Electricity price (\$/kwh):	0.083
Catalyst price (\$/ft3):	650
Natural gas price (\$/mscf):	7.80
Annual interest rate (fraction):	0.07
Control system life (years):	20
Catalyst life (years):	2
Capital recovery factor (system):	0.0944
Capital recovery factor (catalyst):	0.5531
Taxes, insurance, admin. factor:	0.04
Pressure drop (in. w.c.):	21.0

ANNUAL COSTS

Item	Cost (\$/yr)	Wt. Factor	W.F.(co	nd.)
Operating labor	41,063	0.006		
Supervisory labor	6,159	0.001		
Maintenance labor	45,169	0.006		
Maintenance materials	45,169	0.006		
Natural gas	5,142,028	0.713		
Electricity	807,684	0.112		
Catalyst replacement	203,972	0.028		
Overhead	82,536	0.011		0.031
Taxes, insurance, administrative	259,500	0.036		
Capital recovery	577,564	0.080		0.116
Total Annual Cost	7,210,843	1.000		1.000

[1] Original equipment costs reflect this date.

[2] VAPCCI = Vatavuk Air Pollution Control Cost Index (for catalytic incinerators) corresponding to year and quarter shown. Original equipment cost, purchased equipment cost, and total capital investment have been escalated to this date via the VAPCCI and control equipment vendor data.

BACT Cost-Benefit Evaluation

RACT Cost Summary Table

U.S. Bureau of Labor Statistics Correction Factor - 2006 to 2022 CPI Inflation Calculator 1.6808

	2006 dollars	2022 dollars
1 Purchased Equipment Cost (PEC)	4,029,506	6,772,794
2 Total Direct Cost (includes PEC)	5,238,358	8,804,632
3 Total Indirect Cost	1,249,147	2,099,566
4 Total Capital Investment (= 2+3)	6,487,505	10,904,199
5 Annual Direct Operating Costs	6,291,244	10,574,322
Annual Indirect Operating Costs	342,036	574,894
7 Annual Capital Recovery Costs	577,564	970,769
8 Total Annual Costs (= 5+6+7)	7,210,843	12,119,985

Oxidizer VOC Control Efficiency (%)	95 %
Annual VOC Input to the Control Device	393.12 tons
Annual VOC Emissions Controlled	373.464 tons
Annual VOC Emissions after Controls	19.656 tons
Annual Cost of Control Device	\$ 19,308 per ton controlled \$32,453

				u of Labor Statistics Correction 5 to 2022 CPI Inflation Calculator
		3rd Qtr. 2006	2022	1.6808
	Regen	139.0	233.6	
	Recup	146.4	246.1	
	CatOX	217.4	365.4	

Appendix D

RACT/BACT/LAER Clearinghouse Search Results

T

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COMPREHENSIVE REPORT Report Date:11/18/2021

RBLC ID:	IN-0208 (final)	Date Determination Last Updated:	03/07/2019
Corporate/Company Name:	NHK SEATING OF AMERICA, INC.	Permit Number:	023-34689-00035
Facility Name:	NHK SEATING OF AMERICA, INC.	Permit Date:	11/25/2014 (actual)
Facility Contact:	DEAN HILL 7656597824 DHILL@NHKSEATING.COM	FRS Number:	110055569718
Facility Description:	AUTOMOBILE SEAT MANUFACTURING OPERATION	SIC Code:	2531
Permit Type:	D: Both B (Add new process to existing facility) &C (Modify process at existing facility)	NAICS Code:	336360
Permit URL:	HTTP://PERMITS.AIR.IDEM.IN.GOV/34689F.PDF		
EPA Region:	5	COUNTRY:	USA
Facility County:	CLINTON		
Facility State:	IN		
Facility ZIP Code:	46041		
Permit Issued By:	INDIANA DEPT OF ENV MGMT, OFC OF AIR (Agency Name) MR. MATT STUCKEY(Agency Contact) (317) 233-0203 mstuckey@idem.in.j	gov	
Other Agency Contact Info:	PERMIT WRITER: BRIAN WRIGHT BWRIGHT1@IDEM.IN.GOV		
	317-234-6544		
	SECTION CHIEF: NATHAN BELL 317-233-5670		

Permit Notes:

Process/Pollutant Information

PROCESS NAME:	SEAT FOAM PRODU	SEAT FOAM PRODUCTION LINE			
Process Typ	e: 63.013 (Flexible Poly	urethane Foam Production)			
Primary Fu	el:				
Throughpu	t: 56.00 SEATS				
Process Not	es:				
Р	OLLUTANT NAME:	Volatile Organic Compounds (VOC)			
CA	AS Number:	VOC			
Те	st Method:	Unspecified			
Po	llutant Group(s):	(Volatile Organic Compounds (VOC))			
En	nission Limit 1:	95.0000 % OVERALL CONTROL EF 3 HOURS			
En	nission Limit 2:	10.0000 PPMV 3 HOURS			
Sta	andard Emission:				
Die	d factors, other then air pollut	ion technology considerations influence the BACT decisions: N			
Ca	se-by-Case Basis:	OTHER CASE-BY-CASE			
Ot	her Applicable Requirements:				
Co	ntrol Method:	(A) REGENERATIVE THERMAL OXIDIZER			
Es	t. % Efficiency:	95.000			
Co	st Effectiveness:	0 \$/ton			
Ine	cremental Cost Effectiveness:	0 \$/ton			
Co	mpliance Verified:	No			
Po	Pollutant/Compliance Notes: STATE BACT				

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Appendix E

TDEC Construction Permit Application Forms



TITLE V PERMIT APPLICATION INDEX OF AIR POLLUTION PERMIT APPLICATION FORMS

Section 1: Identification and Diagrams				
This application contains the	APC Form 1, Facility Identification 1			
following forms:	APC Form 2, Operations and Flow Diagrams 1			

Section 2: Emission Source Description Forms				
Total number of this form				
	APC Form 3, Stack Identification	1		
	APC Form 4, Fuel Burning Non-Process Equipment			
	APC Form 5, Stationary Gas Turbines or Internal Combustion Engines			
	APC Form 6, Storage Tanks			
This application contains the following forms (one form for each incinerator, printing	APC Form 7, Incinerators			
operation, fuel burning installation, etc.):	APC Form 8, Printing Operations			
	APC Form 9, Painting and Coating Operations			
	APC Form 10, Miscellaneous Processes	1		
	APC Form 33, Stage I and Stage II Vapor Recovery Equipment			
	APC Form 34, Open Burning			

Section 3: Air Pollution Control System Forms					
Total number of this form					
	APC Form 11, Control Equipment - Miscellaneous				
	APC Form 13, Adsorbers				
This application contains the following forms (one form for each control system in use at the	APC Form 14, Catalytic or Thermal Oxidation Equipment				
facility):	APC Form 15, Cyclones/Settling Chambers				
	APC Form 17, Wet Collection Systems				
	APC Form 18, Baghouse/Fabric Filters				

APC Index

		Total number of this form	
	APC Form 19, Compliance Certification - Monitoring and Reporting - Description of Methods for Determining Compliance	1	
	APC Form 20, Continuous Emissions Monitoring	an a	
	APC Form 21, Portable Monitors		
	APC Form 22, Control System Parameters or Operating Parameters of a Process	1	
	APC Form 23, Monitoring Maintenance Procedures		
	APC Form 24, Stack Testing	l.	
This application contains the following forms (one form for each incinerator, printing operation, fuel burning installation, etc.);	APC Form 25, Fuel Sampling and Analysis		
inition, fuel burning instantion, etc.).	APC Form 26, Record Keeping	1	
	APC Form 27, Other Methods		
	APC Form 28, Emissions from Process Emissions Sources / Fuel Burning Installations / Incinerators	n na martina 1	
	APC Form 29, Emissions Summary for the Facility or for the Source Contained in This Application	1	
	APC Form 30, Current Emissions Requirements and Status	1	
	APC Form 31, Compliance Plan and Compliance Certification	1	
	APC Form 32, Air Monitoring Network		

Section 5: Statement of Completeness and Certification of Compliance

I have reviewed this application in its entirety and to the best of my knowledge, and based on information and belief formed after reasonable inquiry, the statements and information contained in this application are true, accurate, and complete. I have provided all the information that is necessary for compliance purposes and this application consists of 11 pages and they are numbered from page 1 to 11. The status of this facility's compliance with all applicable air pollution control requirements, including the enhanced monitoring and compliance certification requirements of the Federal Clean Air Act, is reported in this application along with the methods to be used for compliance demonstration.

Name and Title of Responsible Official

Ryan Speck, Plant Manager

Signature of Responsible Official

Telephone Number with Area Code

3 30 22

931-363-5666

Date of Application

(For definition of responsible official, see instructions for APC Form 1)



TITLE V PERMIT APPLICATION FACILITY IDENTIFICATION

		SIT	E INFO	RMATION				
1. Organization's legal name		For	APC company point no.					
Adient US, LLC			APC					
2. Site name (if different from legal name)			Use	APC Log/Permit no.				
					Only			
3. Site address (St./Rd./Hwy.)					NAICS o	r SIC Code		
1890 Mines Road					326150			
City or distance to nearest tow	vn		Zip co	de	County r	County name		
Pulaski			38478		Giles	Giles		
4. Site location (in Lat./Long)	Latitude				-	Longitude		
	36 13' 46.41" N				87 04' 1	4.41" W		
	CONTACT	INFORM	ATION	(RESPONS	IBLE OFFIC	IAL)		
5. Responsible official contact						umber with area code		
Ryan Speck					931-363	-5666		
6. Mailing address (St./Rd./Hwy	.)				Fax num	ber with area code		
1890 Mines Road					931-363	-5787		
City		State		Zip code		Email address		
Pulaski		TN	3	38478	ryan.spe	eck@adient.com		
	CON	TACT IN	FORMA	ATION (TEC	CHNICAL)			
7. Principal technical contact			Phonem	umber with area code				
Kris P. Foster			931-363	-5666				
8. Mailing address (St./Rd./Hwy.)			Fax num	ber with area code				
1890 Mines Road					931-363	-5787		
City State Zip code		Email ad	dress					
Pulaski		TN	3	38478	kris.patr	kris.patrick.foster@adient.com		
	CO	ONTACT I	INFORM	MATION (B	ILLING)			
11. Billing contact					Phonem	umber with area code		
Rhonda Gardner					931-363	-5666		
12. Mailing address (St./Rd./Hwy	.)				Fax num	Fax number with area code		
1890 Mines Road					931-363	931-363-5787		
City State Zip code		Email ad	Email address					
Pulaski		TN	3	38478	rhonda.	o.garner@adient.com		
		TYPE OF	F PERM	IT REQUES	TED			
13. Permit requested for:								
Initial application to operate :			Minor perm	nit modification :				
Permit rene	wal to operate :				Significat	nt modification :		
Administrative permit amendment :				Cor	nstruction permit :			

(OVER)

A	PC	1

HAZARDOUS AIR POLLUTANTS, DESIGNATIONS, AND OTHER PERMITS ASSOCIATED WITH FACILITY					
14. Is this facility subject to the provisions governing Tennessee Air Pollution Control regulations?	g prevention of accidental releases of hazar	dous air contaminants contained in Chapter 1200-03-32 of the Yes No			
If the answer is Yes, are you in compliance with	the provisions of Chapter 1200-03-32 of the	ne Tennessee Air Pollution Control regulations?			
15. If facility is located in an area designated as "No	n-Attainment" or "Additional Control", inc	licate the pollutant(s) for the designation.			
Attainment for all pollutants.					
 List all valid Air Pollution permits issued to the s reference numbers listed on the permit(s)]. 	sources contained in this application [identi	fy all permits with most recent permit numbers and emission source			
569269 Title V Operating Permit (28-0076)					
17. Page number :	Revision number:	Date of revision:			
2					



TITLE V PERMIT APPLICATION OPERATIONS AND FLOW DIAGRAMS

1. Please list, identify, and describe briefly process emission sources, fuel burning installations, and incinerators that are contained in this application. Please attach a flow diagram for this application.

 List the <u>states</u> that are within 50 miles of your facility Alabama 	y.
3. Are there any storage piles?	YES NO
Tank Farm (Closed Loop System) Foam Crushing Mold Cleaning	ecause of size or production rate and cite the applicable regulations. 1200-03-0904(5)(a)4(i) 1200-03-0904(5)(a)4(i) 1200-03-0904(5)(a)4(i) 1200-03-0904(5)(f)76
	I (poured) into the open mold, the mold is closed, the foam d and the foam cures in the closed mold. Once cured, the foam ed and the cycle is repeated.
cushions. Foam cushions or buns a with mold release, and/or with hand to the mold. VOC emissions from the	nufacturing - production of polyurethane foam automotive seat are produced on one of three production lines. Molds are sprayed applied application techniques to prevent the foam from sticking ne process are mostly from the use of solvent and water based old release materials may be emitted as total suspended

RDA 1298



TITLE V PERMIT APPLICATION STACK IDENTIFICATION

GENERAL IDENTIFICATION AND DESCRIPTION					
1. Facility name:					
Adient US, LLC					
2. Emission source (identify): 28-0076-01 Polyurethane Foam Manufac	cturing				
	STACK DESCRIPTION				
3. Stack ID (or flow diagram point identification):					
F1 through F13 (please see Figure 3 for s	stack parameters)				
4. Stack height above grade in feet:					
5. Velocity (data at exit conditions):	6. Inside dimensions at outlet in feet:				
(Actual feet per second)					
7. Exhaust flow rate at exit conditions (ACFM):	8. Flow rate at standard conditions (DSCFM):				
9. Exhaust temperature:	10. Moisture content (data at exit conditions):				
	Grains per dry				
Degrees Fahrenheit (°F)	standard cubic Percent foot (gr./dscf.)				
11. Exhaust temperature that is equaled or exceeded during ninet	y (90) percent or more of the operating time (<u>for stacks subject to diffusion equation only</u>):				
NA					
	(°F)				
12 If this stack is equipped with continuous pollutant monitoring	equipment required for compliance, what pollutant(s) does this equipment monitor (e.g., Opacity,				
$SO_2, NO_x, etc.)?$	· Jt				
NA					
Complete the appropriate APC form(s) 4,5,7,8,9, or 10 for	each source exhausting through this stack.				
BYPASS STACK DESCRIPTION					
13. Do you have a bypass stack?	v				
Yes No					
If yes, describe the conditions which require its use & complete APC form 4 for the bypass stack. Please identify the stack number(s) of flow diagram point					
number(s) exhausting through this bypass stack.					
4 ^{4.} Page number: Revision N	Tumber: Date of Revision:				
4					



TITLE V PERMIT APPLICATION MISCELLANEOUS PROCESSES

GENERAL IDENTIFICATION AND DESCRIPTION

1. Facility name: Adient US, LLC							
2. Process emission source (identify):							
3. Stack ID or flow diagram p	28-0076-01 Polyurethane Foam Manufacturing 3. Stack ID or flow diagram point identification(s): 4. Year of construction or last modification:						
F1 through F13			ation was in 2010				
-	lled for compliance, attach an appropriate Air Pol	llution Control s	ystem form.				
5. Normal operating schedule	: 24 Hrs./Day 5 Days/Wk. 24	.0 Days/Y	ſr.				
6. Location of this process em	nission source in UTM coordinates: UT	M Vertical : <u>3</u>	898.504 UTMHorizontal: 4	93.569			
7. Describe this process (Plea	se attach a flow diagram of this process) and chee	ck one of the fol	llowing:				
Batch_	Continuous A detailed description of the pro-	ocess is include	d in the narrative portion of the app	olication.			
	PROCESS MATERIA	L INPUT AN	ND OUTPUT				
8. List the types and amounts	of raw materials input to this process:						
Material	Storage/Material handling proces	SS	Average usage (units)	Maximum usage (units)			
Foam chemicals	Chemical storage/delivered directly	to process	2,965,000 gal/yr	4,742,000 gal/yr			
			See attached emission	See attached emission			
			calculations for average	calculations for maximum			
	usage by raw material usage by raw material						
9. List the types and amounts	of primary products produced by this process:						
Material	Storage/Material handling proces	SS	Average usage (units)	Maximum usage (units)			
Auto foam products	Foam manufacturing		3,178,091 FBE	9,833,712 FBE			
			Full Bench Equivalent	Full Bench Equivalent			
10. Process fuel usage:							
Type of fuelMax heat input (10° BTU/Hr.)Average usage (units)Maximum usage (units)							
NA							
11. List any solvents, cleaners,	etc., associated with this process:						
Mold maintenance and cleaning materials.							
If the emissions and/or operations of this process are monitored for compliance, please attach the appropriate Compliance Demonstration form.							
12. Describe any fugitive emissions associated with this process, such as outdoor storage piles, open conveyors, open air sand blasting, material handling operations,							
etc. (please attach a separate sheet if necessary).							
All emissions assumed to be a point source.							
13. Page number:	13. Page number: Date of Revision:						
5	iconsistent calibor.		Date of Revision.				



TITLE V PERMIT APPLICATION COMPLIANCE CERTIFICATION - MONITORING AND REPORTING DESCRIPTION OF METHODS USED FOR DETERMINING COMPLIANCE

requ mon durii	All sources that are subject to 1200-03-0902(11) of the Tennessee Air Pollution Control Regulations are required to certify compliance with all applicable requirements by including a statement within the permit application of the methods used for determining compliance. This statement must include a description of the monitoring, recordkeeping, and reporting requirements and test methods. In addition, the application must include a schedule for compliance certification submittals during the permit term. These submittals must be no less frequent than annually and may need to be more frequent if specified by the underlying applicable requirement or the Technical Secretary.				
	GENERAL IDENTIFICATION AND DESCRIPTION				
1.	Facility name: Adient US, LLC				
2.	Process emission source, fuel burning installation, or incinerator (identify): 28-0076-01 Polyurethane Foam Manufa	cturing			
3.	Stack ID or flow diagram point identification(s): Figure 2 Process Flow Diagram and Figure 3 Stack Paramete	rs			
	METHODS OF DETERMINING COMPLIANCE				
4.	This source as described under Item #2 of this application will use the following method(s) for determining compliance with a (and special operating conditions from an existing permit). Check all that apply and attach the appropriate form(s)	applicable requirements			
	Continuous Emission Monitoring (CEM) - APC 20 Pollutant(s):				
	Emission Monitoring Using Portable Monitors - APC 21 Pollutant(s):				
	Monitoring Control System Parameters or Operating Parameters of a Process - APC22 Pollutant(s): Opacity - Visual Observation				
	Monitoring Maintenance Procedures - APC 23 Pollutant(s):				
	Stack Testing - APC 24 Pollutant(s):				
	Fuel Sampling & Analysis (FSA) - APC 25 Pollutant(s):				
	Recordkeeping - APC 26 Pollutant(s): PM and VOC				
	Other (please describe) - APC 27 Pollutant(s):				
5.	Compliance certification reports will be submitted to the Division according to the following schedule:				
	Start date: Per Title V Permit Requirements				
	And every 365 days thereafter.				
6.	Compliance monitoring reports will be submitted to the Division according to the following schedule:				
	Start date: Per Title V Permit Requirements				
	And every <u>365</u> days thereafter.				
7. 6	Page number:Revision number:Date of revision:				



TITLE V PERMIT APPLICATION - COMPLIANCE DEMONSTRATION BY MONITORING CONTROL SYSTEM PARAMETERS OR OPERATING PARAMETERS OF A PROCESS

The monitoring of a control system parameter or a process parameter shall be acceptable as a compliance demonstration method provided that a correlation between the parameter value and the emission rate of a particular pollutant is established.

GENERAL IDENTIFICATION AND DESCRIPTION

1. Facility name: Adient US, LLC Stack ID or flow diagram point identification(s)
 F1 through F13

3. Emission source:

28-0076-01 Polyurethane Foam Manufacturing

MONITORING DESCRIPTION

4. Pollutant(s) being monitored:

Opacity

5. Description of the method of monitoring and establishment of correlation between the parameter value and the emission rate of a particular pollutant: Visible emissions reading pursuant to EPA Method 9.

6. Compliance demonstration frequency (specify the frequency with which compliance will be demonstrated):

Annually per the Opacity Decision Tree dated June 18, 1996, amended September 11, 2013.

7. Page number:

Revision number:

Date of revision:

7



TITLE V PERMIT APPLICATION

COMPLIANCE DEMONSTRATION BY RECORDKEEPING

requirement is established.	rovided that a correlation between the parameter value recorded and the applicable
GENERAL IDENTIFIC	CATION AND DESCRIPTION
1. Facility name:	2. Stack ID or flow diagram point identification(s):
Adient US, LLC	F1 through F13
3. Emission source (identify):	
28-0076-01 Polyurethane Foam Manufacturing	
MONITORING AND REC	CORDKEEPING DESCRIPTION
4. Pollutant(s) or parameter being monitored:	
PM and VOC	
5. Material or parameter being monitored and recorded:	
Material Usage and Material Formulation Data	
6. Method of monitoring and recording:	
Monthly record of the actual usage of Polyurethane Foam Manufa	acturing materials.
Emissions are estimated as follows:	
1. PM - (Material Usage) x (PM content) x (1-55% Transfer Effici	ency)
2. VOC - (Material Usage) x (VOC content)	
3. TDI, MDI, DEOA - (Material Usage) x (HAP content) x (Emissi	ions Factor)
Note: TDI and MDI are reactants in the foam materials and are la emission factors are from testing performed on October 1, 1997.	argely consumed or locked into the matrix of the foam. The compound
 Compliance demonstration frequency (specify the frequency with which co Monthly Calculation 	ompliance will be demonstrated):
8. Page number: Revision number: 8	Date of revision:



TITLE V PERMIT APPLICATION EMISSION SUMMARY FOR THE FACILITY OR FOR THE SOURCES CONTAINED IN THIS APPLICATION

GENERAL IDENTIFICATION AND DESCRIPTION

1. Facility name: Adient US, LLC

EMISSIONS SUMMARY TABLE - CRITERIA AND SELECTED POLLUTANTS

2. Complete the following emissions summary for regulated air pollutants at this facility or for the sources contained in this application.

	Summary of Maximum Allowable Emissions		Summary of Actual Emissions		
Air Pollutant	Tons per Year	Reserved for State use (Pounds per Hour- Item 4, APC 28)	Tons per Year	Reserved for State use (Pounds per Hour- Item 4, APC 28)	
Part iculate Matter (TSP)	9.9	NA	<9.9	NA	
Sulfur Dioxide					
Volatile Organic Compounds	491.4	NA	305	NA	
Carbon Monoxide					
Lead					
Nitrogen Oxides					
Total Reduced Sulfur					
Mercury					
Asbestos					
Beryllium					
Vinyl Chlorides					
Fluorides					
Gaseous Fluorides					
Greenhouse Gases in CO ₂ Equivalents					

(Continued from previous page) EMISSIONS SUMMARY TABLE – HAZARDOUS AIR POLLUTANTS

3. Complete the following emissions summary for regulated air pollutants that are hazardous air pollutant(s) at this facility or for the sources contained in this application.

	Summary of Maximum Allowable Emissions		Summary of Actual Emissions		
Air Pollutant & CAS	Tons per Year	Reserved for State use (Pounds per Hour- Item 5, APC 28)	Tons per Year	Reserved for State use (Pounds per Hour- Item 5, APC 28)	
TDI, 26471-62-5	9.9	NA	<1	NA	
MDI, 101-68-8	9.9	NA	<1	NA	
Diethanolamine, 111-42-2	9.9	NA	<1	NA	
Page number:	Revision nur	nber:	Date of revision:		



TITLE V PERMIT APPLICATION CURRENT EMISSIONS REQUIREMENTS AND STATUS

GENERAL IDENTIFICATION AND DESCRIPTION					
1. Facility name:	2. Emission source number				
Adient US, LLC	28-0076-01				
3. Describe the process emission source / fuel burning installation / incinerator.					
Polyurethane Foam Manufacturing					
	SSIONS AND REQUIREMENTS				
4. Identify if only a part of the source is subject to this requirement 5. Pollutant 6. Applicable requirement(s): Regulations, 40 CFR, perm air quality based standards					
Please refer to the application narrative for a list of applicable					
	process and accordingly, the PSD application includes the most current				
	Regarding compliance status associated with applicable requirements,				
please refer to TDEC application form APC 31 below, Note 5	5.				
10. Other applicable requirements (new requirements that apply to this source during the term of this permit)					
11. Page number:Revision number:10	Date of revision:				



TITLE V PERMIT APPLICATION COMPLIANCE PLAN AND COMPLIANCE CERTIFICATION GENERAL IDENTIFICATION AND DESCRIPTION

Adient Ú	1. Facility name: Adient US LLC Pulaski			
2. List all	he process emission source(s) or fuel burning installation(s) or	rincinerator(s) that are part of this application.		
28-0076	01 Polyurethane Foam Manufacturing			
	COMPLIANCE F	PLAN AND CERTIFICATION		
3. Indicate	that source(s) which are contained in this application are pres	ently in compliance with all applicable requirements, by checking the following:		
_X	A. Attached is a statement of identification of the source to assure compliance with all the applicable requirem	(s) currently in compliance. We will continue to operate and maintain the source(s) ents for the duration of the permit.		
	B APC 30 form(s) includes new requirements that apply requirements on a timely basis.	y or will apply to the source(s) during the term of the permit. We will meet such		
4. Indicate	that there are source(s) that are contained in this application w	hich are not presently in full compliance, by check ing both of the following:		
	A. Attached is a statement of identification of the source and the proposed solution.	(s) not in compliance, non-complying requirement(s), brief description of the problem,		
	B. We will achieve compliance according to the following	g schedule:		
	Action	Deadline		
Progres	s reports will be submitted:			
Start da		0 days thereafter until compliance is achieved.		
5. State th under se	e compliance status with any applicable compliance assurance ction 114(a)(3) of the Clean Air Act as of the date of submitta	monitoring and compliance certification requirements that have been promulgated l of this APC 31.		
requirem	NOTE: As set forth is the attached Permit Narrative, TDEC alleged non-compliance with some permit requirements and Adient disputed those allegations. On February 24, 2022, Adient and TDEC entered into a Consent Order to resolve those concerns.			
6. Page nu	mber: Revision number:	Date of revision:		

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SAFETY DATA SHEET

Chem-Trend® PU-11434



Section 1. Identification

Product name :

*.

γ. i

: Chem-Trend® PU-11434

Relevant identified uses of the substance or mixture and uses advised against Not available.

Supplier's details	: Chem-Trend LP 1445 W McPherson Park Dr PO Box 860, Howell MI 48844-0860 517-546-4520
Emergency telephone number and Telephone number	: +1 517 546 4520

Section 2. Hazard	s identification
OSHA/HCS status	: This material is considered hazardous by the OSHA Hazard Communication Standard (29 CFR 1910.1200).
Classification of the substance or mixture	: FLAMMABLE LIQUIDS - Category 3 ACUTE TOXICITY (inhalation) - Category 4 SKIN IRRITATION - Category 2 EYE IRRITATION - Category 2A SPECIFIC TARGET ORGAN TOXICITY (SINGLE EXPOSURE) (Narcotic effects) - Category 3 ASPIRATION HAZARD - Category 1
GHS label elements	
Hazard pictograms	
Signal word	: Danger
Hazard statements	 Flammable liquid and vapor. Harmful if inhaled. Causes serious eye irritation. Causes skin irritation. May be fatal if swallowed and enters airways. May cause drowsiness or dizziness.
Precautionary statements	
Prevention	: Wear protective gloves. Wear eye or face protection. Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking. Use explosion-proof electrical, ventilating, lighting and all material-handling equipment. Use only non-sparking tools. Take precautionary measures against static discharge. Keep container tightly closed. Use only outdoors or in a well-ventilated area. Avoid breathing vapor. Wash hands thoroughly after handling.

Section 2. Hazards identification

Response	: IF INHALED: Remove person to fresh air and keep comfortable for breathing. Call a POISON CENTER or physician if you feel unwell. IF SWALLOWED: Immediately call a POISON CENTER or physician. Do NOT induce vomiting. IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water or shower. IF ON SKIN: Wash with plenty of soap and water. Take off contaminated clothing and wash it before reuse. If skin irritation occurs: Get medical attention. IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. If eye irritation persists: Get medical attention.		
Storage	: Store locked up. Store in a well-ventilated place. Keep cool.		
Disposal	 Dispose of contents and container in accordance with all local, regional, national and international regulations. 		
Hazards not otherwise classified	: None known.		

Section 3. Composition/information on ingredients

Substance/mixture : Mixture			
Ingredient name	%	CAS number	
nonane Hydrocarbon naphthas	≥50 - ≤75 ≥25 - ≤50	111-84-2	

Section 4. First aid measures

Description of necessary first aid measures

Eye contact	 Immediately flush eyes with plenty of water, occasionally lifting the upper and lower eyelids. Check for and remove any contact lenses. Continue to rinse for at least 10 minutes. Get medical attention. 			
Inhalation	: Remove victim to fresh air and keep at rest in a position comfortable for breathing. If it is suspected that fumes are still present, the rescuer should wear an appropriate mask or self-contained breathing apparatus. If not breathing, if breathing is irregular or if respiratory arrest occurs, provide artificial respiration or oxygen by trained personnel. It may be dangerous to the person providing aid to give mouth-to-mouth resuscitation. Get medical attention. If necessary, call a poison center or physician. If unconscious, place in recovery position and get medical attention immediately. Maintain an open airway. Loosen tight clothing such as a collar, tie, belt or waistband.			
Skin contact	: Flush contaminated skin with plenty of water. Remove contaminated clothing and shoes. Continue to rinse for at least 10 minutes. Get medical attention. Wash clothing before reuse. Clean shoes thoroughly before reuse.			
Ingestion	: Get medical attention immediately. Call a poison center or physician. Wash out mouth with water. Remove dentures if any. Remove victim to fresh air and keep at rest in a position comfortable for breathing. If material has been swallowed and the exposed person is conscious, give small quantities of water to drink. Stop if the exposed person feels sick as vomiting may be dangerous. Aspiration hazard if swallowed. Can enter lungs and cause damage. Do not induce vomiting. If vomiting occurs, the head should be kept low so that vomit does not enter the lungs. Never give anything by mouth to an unconscious person. If unconscious, place in recovery position and get medical attention immediately. Maintain an open airway. Loosen tight clothing such as a collar, tie, belt or waistband.			
Most important symptoms/effects, acute and delayed				
Potential acute health effec	ts			

Eye contact	:	Causes	serious e	ye irritation.

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Section 4. First aid measures					
Inhalation	: Harmful if inhaled. Can cause central nervous system (CNS) depression. May cause drowsiness or dizziness.				
Skin contact	: Causes skin irritation.				
Ingestion	 Can cause central nervous system (CNS) depression. May be fatal if swallowed and enters airways. 				
<u>Over-exposure signs/sym</u>	ptoms				
Eye contact	: Adverse symptoms may include the following: pain or irritation watering redness				
Inhalation	: Adverse symptoms may include the following: nausea or vomiting headache drowsiness/fatigue dizziness/vertigo unconsciousness				
Skin contact	: Adverse symptoms may include the following: irritation redness				
Ingestion	: Adverse symptoms may include the following: nausea or vomiting				
Indication of immediate me	dical attention and special treatment needed, if necessary				
Notes to physician	: Treat symptomatically. Contact poison treatment specialist immediately if large quantities have been ingested or inhaled.				
Specific treatments	: No specific treatment.				
Protection of first-aiders	: No action shall be taken involving any personal risk or without suitable training. If it is suspected that fumes are still present, the rescuer should wear an appropriate mask or self-contained breathing apparatus. It may be dangerous to the person providing aid to give mouth-to-mouth resuscitation.				

See toxicological information (Section 11)

Section 5. Fire-fighting measures

Extinguishing media	
Suitable extinguishing media	: Use dry chemical, CO ₂ , water spray (fog) or foam.
Unsuitable extinguishing media	: Do not use water jet.
Specific hazards arising from the chemical	: Flammable liquid and vapor. Runoff to sewer may create fire or explosion hazard. In a fire or if heated, a pressure increase will occur and the container may burst, with the risk of a subsequent explosion. The vapor/gas is heavier than air and will spread along the ground. Vapors may accumulate in low or confined areas or travel a considerable distance to a source of ignition and flash back. Fire water contaminated with this material must be contained and prevented from being discharged to any waterway, sewer or drain.
Hazardous thermal decomposition products	: Decomposition products may include the following materials: carbon dioxide carbon monoxide

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Section 5. Fire-fighting measures

Special protective actions for fire-fighters	: Promptly isolate the scene by removing all persons from the vicinity of the incident if there is a fire. No action shall be taken involving any personal risk or without suitable training. Move containers from fire area if this can be done without risk. Use water spray to keep fire-exposed containers cool.
Special protective equipment for fire-fighters	: Fire-fighters should wear appropriate protective equipment and self-contained breathing apparatus (SCBA) with a full face-piece operated in positive pressure mode.

Section 6. Accidental release measures

Personal precautions, protective equipment and emergency procedures						
For non-emergency personnel	: No action shall be taken involving any personal risk or without suitable training. Evacuate surrounding areas. Keep unnecessary and unprotected personnel from entering. Do not touch or walk through spilled material. Shut off all ignition sources. No flares, smoking or flames in hazard area. Avoid breathing vapor or mist. Provide adequate ventilation. Wear appropriate respirator when ventilation is inadequate. Put on appropriate personal protective equipment.					
For emergency responders	: If specialized clothing is required to deal with the spillage, take note of any information in Section 8 on suitable and unsuitable materials. See also the information in "For non-emergency personnel".					
Environmental precautions	: Avoid dispersal of spilled material and runoff and contact with soil, waterways, drains and sewers. Inform the relevant authorities if the product has caused environmental pollution (sewers, waterways, soil or air). Water polluting material. May be harmful to the environment if released in large quantities.					

Methods and materials for containment and cleaning up

Stop leak if without risk. Move containers from spill area. Use spark-proof tools and explosion-proof equipment. Approach release from upwind. Prevent entry into sewers, water courses, basements or confined areas. Wash spillages into an effluent treatment plant or proceed as follows. Contain and collect spillage with non-combustible, absorbent material e.g. sand, earth, vermiculite or diatomaceous earth and place in container for disposal according to local regulations (see Section 13). Dispose of via a licensed waste disposal contractor. Contaminated absorbent material may pose the same hazard as the spilled product. Note: see Section 1 for emergency contact information and Section 13 for waste disposal.

Section 7. Handling and storage

Precautions for safe handling

Protective measures: Put on appropriate personal protective equipment (see Section 8). Do not swallow.
Avoid contact with eyes, skin and clothing. Avoid breathing vapor or mist. Avoid
release to the environment. Use only with adequate ventilation. Wear appropriate
respirator when ventilation is inadequate. Do not enter storage areas and confined
spaces unless adequately ventilated. Keep in the original container or an approved
alternative made from a compatible material, kept tightly closed when not in use. Store
and use away from heat, sparks, open flame or any other ignition source. Use
explosion-proof electrical (ventilating, lighting and material handling) equipment. Use
only non-sparking tools. Take precautionary measures against electrostatic discharges.
Empty containers retain product residue and can be hazardous. Do not reuse container.

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Section 7. Handling and storage

Conditions for safe storage,	: Store in accordance with local regulations. Store in a segregated and approved area.
including any	Store in original container protected from direct sunlight in a dry, cool and well-ventilated
incompatibilities	area, away from incompatible materials (see Section 10) and food and drink. Store
·	locked up. Eliminate all ignition sources. Separate from oxidizing materials. Keep
	container tightly closed and sealed until ready for use. Containers that have been
	opened must be carefully resealed and kept upright to prevent leakage. Do not store in
	unlabeled containers. Use appropriate containment to avoid environmental
	contamination. See Section 10 for incompatible materials before handling or use.

Section 8. Exposure controls/personal protection

Control parameters

Occupational exposure limits

Ingredient name			Exposure limits				
nonane		ACGIH TLV (United States, 3/2017). TWA: 200 ppm 8 hours. TWA: 1050 mg/m ³ 8 hours.					
Hydrocarbon naphthas			ACGIH TLV (United S TWA: 200 mg/m³, (a	States, 3/2017). A			
Appropriate engineering controls	:	other engineeri recommended	Idequate ventilation. U Ing controls to keep wo or statutory limits. The oncentrations below ar pment.	rker exposure to a engineering cont	airborne contami rols also need to	nants below keep gas,	
Environmental exposure controls	:	they comply wit cases, fume sc	n ventilation or work pro th the requirements of e rubbers, filters or engir ry to reduce emissions	environmental pro neering modification	tection legislation	n. In some	
Individual protection meas							
Eye/face protection		assessment inc gases or dusts.	r complying with an app dicates this is necessar . If contact is possible, t indicates a higher deg	y to avoid exposu the following prot	re to liquid splas ection should be	hes, mists, worn, unles:	S
Hand protection	:	worn at all time necessary. Co during use that noted that the t glove manufact	tant, impervious gloves s when handling chem nsidering the paramete the gloves are still reta ime to breakthrough fo turers. In the case of n of the gloves cannot be	ical products if a r ers specified by the ining their protect r any glove materi nixtures, consisting	isk assessment i e glove manufact ive properties. It al may be differe g of several subs	indicates this turer, check should be ent for differe	s is
Body protection	:	performed and handling this pr static protective	ctive equipment for the the risks involved and oduct. When there is a clothing. For the grea anti-static overalls, boo	should be approve a risk of ignition fro itest protection fro	ed by a specialist om static electric	t before ity, wear ant	i-
Other skin protection	:	based on the ta	otwear and any addition ask being performed an e handling this product	d the risks involve			
Respiratory protection	:	appropriate star	azard and potential for ndard or certification. I ection program to ensu	Respirators must I	be used accordir	ng to a	
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Section 8. Exposure controls/personal protection

Section 9. Physical and chemical properties

Physical state	Liquid.	Color	Off-white.
Odor	Hydrocarbon.	Odor threshold	Not available.
рН	Not applicable.	Melting point	Not available.
Boiling point	154.4°C (309.9°F)	Flash point	Closed cup: 39.4°C (102.9°F) [Setaflash.]
Burning time	Not applicable.	Burning rate	Not applicable.
Evaporation rate	<1 (butyl acetate = 1)	Flammability (solid, gas)	Not available.
Lower and upper explosive (flammable) limits	Not available.	Vapor pressure	Not available.
Vapor density	>1 [Air = 1]	Relative density	0.784
Solubility	Insoluble in the following materials: cold water.	Solubility in water	Not available.
Partition coefficient: n- octanol/water	Not available.	Auto-ignition temperature	Not available.
Decomposition temperature	Not available.	SADT	Not available.
Viscosity	Kinematic (40°C (104°F)): <0.2 cm²/s (<20 cSt)	Volatility	95.9

Lower and upper explosive (flammable) limits Distillates (petroleum), hydrotreated light nonane

Lower: 0.6% Upper: 5.5% Lower: 0.8% Upper: 2.9%

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Aerosol product

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Section 10. Stability and reactivity				
Reactivity	:	No specific test data related to reactivity available for this product or its ingredients.		
Chemical stability	:	The product is stable.		
Possibility of hazardous reactions	;	Under normal conditions of storage and use, hazardous reactions will not occur.		

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Section 10. Stability and reactivity

Conditions to avoid		Avoid all possible sources of ignition (spark or flame). Do not pressurize, cut, weld, braze, solder, drill, grind or expose containers to heat or sources of ignition. Do not allow vapor to accumulate in low or confined areas.
Incompatible materials	:	Reactive or incompatible with the following materials: oxidizing materials
Hazardous decomposition products	:	Under normal conditions of storage and use, hazardous decomposition products should not be produced.

Section 11. Toxicological information

Information on toxicological effects

Acute toxicity

Product/ingredient name	Result	Species	Dose	Exposure
nonane	LC50 Inhalation Gas. LC50 Inhalation Vapor	Rat Rat	3200 ppm 17000 mg/m³	4 hours 4 hours
Irritation/Corrosion	: Causes serious eye irritatio	n. Causes skin irri	tation.	
Sensitization	: No known significant effects or critical hazards.			
Mutagenicity	: No known significant effects or critical hazards.			
Carcinogenicity	: No known significant effects or critical hazards.			
Reproductive toxicity	: No known significant effects or critical hazards.			
Teratogenicity	: No known significant effects or critical hazards.			

Specific target organ toxicity (single exposure)

Name	Target organs
Hydrocarbon naphthas	Narcotic effects

Specific target organ toxicity (repeated exposure)

Not available.

Aspiration hazard

Name	Result
Chem-Trend® PU-11434	ASPIRATION HAZARD - Category 1
nonane	ASPIRATION HAZARD - Category 1
Hydrocarbon naphthas	ASPIRATION HAZARD - Category 1

Information on the likely : Not available. routes of exposure

100100 01	expected
Potential	acute health effects

Eye contact	: Causes serious eye irritation.
Inhalation	: Harmful if inhaled. Can cause central nervous system (CNS) depression. May cause drowsiness or dizziness.
Skin contact	: Causes skin irritation.
Ingestion	: Can cause central nervous system (CNS) depression. May be fatal if swallowed and enters airways.

Symptoms related to the physical, chemical and toxicological characteristics

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Section 11. Toxicological information

Eye contact	Skin contact
Adverse symptoms may include the following: pain or irritation watering redness	Adverse symptoms may include the following: irritation redness
Inhalation	Ingestion
Adverse symptoms may include the following: nausea or vomiting headache drowsiness/fatigue dizziness/vertigo unconsciousness	Adverse symptoms may include the following: nausea or vomiting

Delayed and immediate effects and also chronic effects from short and long term exposure

Short term exposure	
Potential immediate effects	: Not available.
Potential delayed effects	: Not available.
Long term exposure	
Potential immediate effects	: Not available.
Potential delayed effects	: Not available.

Numerical measures of toxicity

Acute toxicity estimates

Route	ATE value	
Inhalation (gases) Inhalation (vapors)	5679.4 ppm 14.48 mg/l	

Section 12. Ecological information

No known significant effects or critical hazards.

Section 13. Disposal considerations

Disposal methods : The generation of waste should be avoided or minimized wherever possible. Disposal of this product, solutions and any by-products should at all times comply with the requirements of environmental protection and waste disposal legislation and any regional local authority requirements. Dispose of surplus and non-recyclable products via a licensed waste disposal contractor. Waste should not be disposed of untreated to the sewer unless fully compliant with the requirements of all authorities with jurisdiction. Waste packaging should be recycled. Incineration or landfill should only be considered when recycling is not feasible. This material and its container must be disposed of in a safe way. Care should be taken when handling emptied containers that have not been cleaned or rinsed out. Empty containers or liners may retain some product residues. Vapor from product residues may create a highly flammable or explosive atmosphere inside the container. Do not cut, weld or grind used containers unless they have been cleaned thoroughly internally. Avoid dispersal of spilled material and runoff and contact with soil, waterways, drains and sewers.

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Section 13. Disposal considerations

RCRA classification

: D001 Because of its ignitability if the product is disposed of in its original form.

Section 14. Transport information

	DOT Classification	Bulk	TDG Classification	ΙΑΤΑ	IMDG
UN number	Not regulated.	NA1993	UN1993	UN1993	UN1993
UN proper shipping name	-	Combustible liquid, n.o.s.(nonane, Hydrocarbon naphthas)	FLAMMABLE LIQUID, N.O.S. (nonane, Hydrocarbon naphthas)	Flammable liquid, n.o.s. (nonane, Hydrocarbon naphthas)	FLAMMABLE LIQUID, N.O.S. (nonane, Hydrocarbon naphthas)
Transport hazard class(es)	-		3	3	3
Packing group	-	111	111	111	111
Environmental hazards	No.	Yes.	No.	No.	Yes.

Emergency Response Guidebook (ERG): 128

Additional information

DOT Classification	:	-
TDG Classification		Product classified as per the following sections of the Transportation of Dangerous Goods Regulations: 2.18-2.19 (Class 3). <u>Explosive Limit and Limited Quantity Index</u> 5 <u>Passenger Carrying Road or Rail Index</u> 60 <u>Special provisions</u> 16
IMDG		The marine pollutant mark is not required when transported in sizes of ≤5 L or ≤5 kg. <u>Emergency schedules</u> F-E, _S-E_ <u>Special provisions</u> 223, 274, 955
ΙΑΤΑ	-	The environmentally hazardous substance mark may appear if required by other transportation regulations. <u>Quantity limitation</u> Passenger and Cargo Aircraft: 60 L. Packaging instructions: 355. Cargo Aircraft Only: 220 L. Packaging instructions: 366. Limited Quantities - Passenger Aircraft: 10 L. Packaging instructions: Y344. <u>Special provisions</u> A3
Special precautions for user	l	Transport within user's premises: always transport in closed containers that are upright and secure. Ensure that persons transporting the product know what to do in the event of an accident or spillage.

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Section 15. Regulatory information

Inventory list	
Australia	: All components are listed or exempted.
Canada	: At least one component is not listed in DSL but all such components are listed in NDSL.
China	: All components are listed or exempted.
Europe	: Contact local supplier or distributor.
Japan	: Japan inventory (ENCS): At least one component is not listed. Japan inventory (ISHL): At least one component is not listed.
New Zealand	: All components are listed or exempted.
Philippines	: All components are listed or exempted.
Republic of Korea	: All components are listed or exempted.
Taiwan	: All components are listed or exempted.
United States	: All components are listed or exempted.
Clean Air Act Section 112	2(b) Hazardous Air Pollutants (HAPs)

Not applicable.

SARA 302/304	
Composition/information	n on ingredients
No products were found.	
SARA 304 RQ	: Not applicable.
<u>SARA 311/312</u>	
Classification	 FLAMMABLE LIQUIDS - Category 3 ACUTE TOXICITY (inhalation) - Category 4 SKIN IRRITATION - Category 2 EYE IRRITATION - Category 2A SPECIFIC TARGET ORGAN TOXICITY (SINGLE EXPOSURE) (Narcotic effects) - Category 3 ASPIRATION HAZARD - Category 1
State regulations	
Massachusetts	: The following components are listed: NONANE; Waxy solid1
New York	: None of the components are listed.
New Jersey	: The following components are listed: NONANE; Waxy solid1
Pennsylvania	: The following components are listed: NONANE; Waxy solid1
<u>California Prop. 65</u>	

▲ WARNING: This product can expose you to benzene, which is known to the State of California to cause cancer and birth defects or other reproductive harm. This product can expose you to chemicals including naphthalene, ethylbenzene, cumene, which are known to the State of California to cause cancer, and Toluene, which is known to the State of California to cause birth defects or other reproductive harm. For more information go to www. P65Warnings.ca.gov.

U.S. Federal regulations : TSCA 12(b) one-time export: nonane

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Chem-Trend® PU-11434

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Section 16. Other information

Hazardous Material Inf	ormation System (U.S.A.)
Health: 3 /	Flammability : 2 Physical hazards : 0 Personal protection Code : H
National Fire Protectio	n Association (U.S.A.)
Health : 1	Flammability : 2 Instability/Reactivity : 0 Special : -
<u>History</u>	
Date of issue/Date of revision	: 4/22/2018
Date of previous issu	e : 6/27/2017
Version	: 1.1
Prepared by	: Chem-Trend Regulatory Affairs Department.
Key to abbreviations	: ATE = Acute Toxicity Estimate BCF = Bioconcentration Factor GHS = Globally Harmonized System of Classification and Labelling of Chemicals IATA = International Air Transport Association IBC = International Air Transport Association IMDG = International Maritime Dangerous Goods LogPow = logarithm of the octanol/water partition coefficient MARPOL = International Convention for the Prevention of Pollution From Ships, 1973 as modified by the Protocol of 1978. ("Marpol" = marine pollution) UN = United Nations

✓ Indicates information that has changed from previously issued version.

Notice to reader

Information presented herein has been compiled from information provided to us by our suppliers and other sources considered to be dependable and is accurate and reliable to the best of our knowledge and belief but is not guaranteed to be so. Nothing herein is to be construed as recommending any practice or the use of any product in violation of any patent or in violation of any law or regulation. It is the users' responsibility to determine the suitability of any material for a specific purpose and to adopt such safety precautions as may be necessary. We make no warranty as to the results to be obtained in using any material and, since conditions of use are not under our control, we must necessarily disclaim all liability with respect to the use of any material supplied by us.



SAFETY DATA SHEET

Product Name: Rikeizai H-179K-7

SECTION 1) CHEMICAL PRODUCT AND SUPPLIER'S IDENTIFICATION

Product Name:	Rikeizai H-179K-7		
SDS Number:	CRSH-179K-7		
Revision Date:	Jan 08, 2019	Date Printed:	Sep 09, 2019
Version:	2.2	Supersedes Date:	Jan 30, 2018
Manufacturer's Name:	Cytech Products		
Address:	906 Peterson Drive Elizabethtown, KY, U	JS, 42701	
Emergency Phone:	1-800-424-9300 (CHEMTREC)		
Information Phone Numbe	r: 1-270-769-1046		
Fax:			

Product/Recommended Uses: Mold Release Prep and Touchup Paste Wax for Polyurethane Molding

SECTION 2) HAZARDS IDENTIFICATION

Classification

Acute toxicity Inhalation - Category 4

Aspiration Hazard - Category 1

Eye Irritation - Category 2B

Flammable Liquids - Category 3

Skin Irritation - Category 2

Specific Target Organ Toxicity -Single Exposure (Narcotic Effects) - Category 3

Pictograms



Signal Word

Danger

Hazardous Statements - Health

H332 - Harmful if inhaled

H304 - May be fatal if swallowed and enters airways

H320 - Causes eye irritation

H315 - Causes skin irritation

H336 - May cause drowsiness or dizziness

Hazardous Statements - Physical

H226 - Flammable liquid and vapor

Precautionary Statements - General

P101 - If medical advice is needed, have product container or label at hand.

P102 - Keep out of reach of children.

P103 - Read label before use.

Precautionary Statements - Prevention

- P261 Avoid breathing dust/fume/gas/mist/vapors/spray.
- P271 Use only outdoors or in a well-ventilated area.
- P264 Wash thoroughly/ hands thoroughly after handling.
- P210 Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.
- P233 Keep container tightly closed.
- P240 Ground/bond container and receiving equipment.
- P241 Use explosion-proof electrical, ventilating, lighting equipment.
- P242 Use only non-sparking tools.
- P243 Take action to prevent static discharges.
- P280 Wear protective gloves/protective clothing/eye protection/face protection.

Precautionary Statements - Response

- P304 + P340 IF INHALED: Remove person to fresh air and keep comfortable for breathing.
- P312 Call a POISON CENTER or doctor, if you feel unwell.
- P301 + P310 IF SWALLOWED: Immediately call a POISON CENTER or doctor.
- P331 Do NOT induce vomiting.

P305 + P351 + P338 - IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.

- P337 + P313 If eye irritation persists: Get medical advice/attention.
- P303 + P361 + P353 IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water or shower.
- P370 + P378 In case of fire: Use carbon-dixoxide, alcohol foam, water spray or dry chemical to extinguish.
- P302 + P352 IF ON SKIN: Wash with plenty of water.
- P321 Specific treatment (see first-aid on this label).
- P332 + P313 If skin irritation occurs: Get medical advice/attention.
- P362 + P364 Take off contaminated clothing. And wash it before reuse.

Precautionary Statements - Storage

P405 - Store locked up.

P403 + P235 - Store in a well-ventilated place. Keep cool.

P403 + P405 - Store in a well-ventilated place. Store locked up.

Precautionary Statements - Disposal

P501 - Dispose of contents/container in accordance with local/national/international regulation. Under RCRA it is the responsibility of the user of the products to determine at the time of disposal whether the product meets RCRA criteria for hazardous waste. Waste management should be in full compliance with federal, state and local laws.

Hazards Not Otherwise Classified (HNOC)

None.

Acute toxicity of less than one percent of the mixture is unknown

SECTION 3) COMPOSITION / INFORMATION ON INGREDIENTS

CAS	Chemical Name	% By Weight
		70 by weight
0064742-47-8	ISOPARAFFINIC PETROLEUM DIS	TILLATE 72% - 76%

Specific chemical identity and/or exact percentage (concentration) of the composition has been withheld to protect confidentiality.

SECTION 4) FIRST-AID MEASURES

Inhalation

Remove source of exposure or move person to fresh air and keep comfortable for breathing. If experiencing respiratory symptoms: Call a POISON CENTER/doctor. If breathing is difficult, trained personnel should administer emergency oxygen if advised to do so by the POISON CENTER/doctor. If exposed/lf you feel unwell/lf concerned: Call a POISON CENTER/doctor.

Skin Contact

Take off contaminated clothing, shoes and leather goods (e.g. watchbands, belts). Wash with plenty of lukewarm, gently flowing water for a duration of 15-20 minutes or until medical aid is available. IF exposed or concerned: Get medical advice/attention. Wash contaminated clothing before re-use.

Eye Contact

If irritation occurs, cautiously rinse eyes with lukewarm, gently flowing water for 5 minutes, while holding the eyelids open. If eye irritation persists: Get medical advice/attention.

Ingestion

Rinse mouth. Drink 2-3 glasses of water. Immediately call a POISON CENTER/doctor. Do NOT induce vomiting. If vomiting occurs naturally, lie on your side, in the recovery position.

Most Important Symptoms/Effects, Acute and Delayed

Skin Contact: Prolonged skin contact may cause irritation or dermatitis.

Inhalation: Vapor concentrations above the TLV may cause nasal and respiratory irritation, dizziness, weakness, fatigue, nausea, headache, and, at extreme levels, asphyxiation.

Indication of Immediate Medical Attention and Special Treatment Needed

No data available.

SECTION 5) FIRE-FIGHTING MEASURES

Suitable Extinguishing Media

Small Fire : Dry chemical, carbon dioxide, water-spray or alcohol-resistant foam.

Large Fire : Water spray, fog or alcohol-resistant foam.

Carbon dioxide can displace oxygen. Use caution when applying carbon dioxide in confined spaces.

Unsuitable Extinguishing Media

Do not use straight stream of water.

Specific Hazards in Case of Fire

Fire will produce irritating gases.

Containers may explode in fire.

Runoff from fire control may cause pollution.

Fire-Fighting Procedures

Isolate immediate hazard area and keep unauthorized personnel out. Stop spill/release if it can be done safely. Move undamaged containers from immediate hazard area if it can be done safely. Cool containers with flooding quantities of water until well after fire is out. Caution should be exercised when using water or foam as frothing may occur, especially if sprayed into containers of hot, burning liquid. Dispose of fire debris and contaminated extinguishing water in accordance with official regulations. Do not allow contaminated extinguishing water or surface waters.

Large Fire: Dike fire-control water for later disposal; do not scatter the material.

Special Protective Actions

Wear positive pressure self-contained breathing apparatus (SCBA).

Structural firefighters' protective clothing provides limited protection in fire situations ONLY; it is not effective in spill situations where direct contact with the substance is possible.

SECTION 6) ACCIDENTAL RELEASE MEASURES

Emergency Procedure

Evacuate and isolate hazard area and keep unauthorized personnel away. Stay uphill and/or upstream. Ventilate closed spaces before entering. Do not touch damaged containers or spilled material unless wearing appropriate protective clothing. ELIMINATE all ignition sources (no smoking, flares, sparks or flames in immediate area). All equipment used when handling the product must be grounded. A vapor-suppressing foam may be used to reduce vapors.

Recommended Equipment

Wear liquid tight chemical protective clothing in combination with positive pressure self-contained breathing apparatus (SCBA).

Personal Precautions

DO NOT breathe vapor or mist.

DO NOT get on skin, eyes or clothing.

Environmental Precautions

Stop spill/release if it can be done safely. Prevent spilled material from entering sewers, storm drains, other unauthorized drainage systems and natural waterways by using sand, earth, or other appropriate barriers. Inform the relevant authorities if the product has caused environmental pollution (sewers, waterways, soil or air).

Methods and Materials for Containment and Cleaning up

Absorb or cover with dry earth, sand or other non-combustible material and transfer to containers. Ventilate area after clean-up is complete.

SECTION 7) HANDLING AND STORAGE

General

Wash hands after use.

Do not get in eyes, on skin or on clothing.

Do not breathe vapors or mists.

Use good personal hygiene practices.

Eating, drinking and smoking in work areas is prohibited.

This product is not intended for human or animal consumption.

Remove contaminated clothing and protective equipment before entering eating areas.

All containers must be properly labelled.

Eyewash stations and showers should be available in areas where this material is used and stored.

Ventilation Requirements

Use only with adequate ventilation to control air contaminants to their exposure limits. The use of local ventilation is recommended to control emissions near the source. Report ventilation failures immediately.

Storage Room Requirements

Segregate from other hazard classes and store in a cool, dry, well ventilated area, away from sources of ignition and incompatibilities.

Protect containers against banging or other physical damage when storing, transferring, or using them.

Provide secondary containment for toxic materials.

Keep containers securely sealed when not in use.

Keep the smallest amount of material in work areas.

Procedures must be conducted in a fume hood, glove box, or other suitable containment device.

Empty container can retain residue and may be dangerous.

Label cabinets with "TOXIC CHEMICALS" or similar warning.

Store between 40-100°F.

SECTION 8) EXPOSURE CONTROLS/PERSONAL PROTECTION

Eye protection

Wear indirect-vent, impact and splash resistant goggles when working with liquids.

Skin protection

Use of gloves approved from relevant standards that meet or are equivalent to OSHA 29 CFR 1910.132. Suitability and durability of a glove is dependent on usage, e.g. frequency and duration of contact, chemical resistance of glove material, glove thickness, dexterity. Always seek advice from glove suppliers. Contaminated gloves should be replaced. Use of an apron and over- boots of chemically impervious materials such as neoprene or nitrile rubber. The type of protective equipment must be selected according to the concentration and amount of the dangerous substance at the specific workplace.

Respiratory protection

If engineering controls do not maintain airborne concentrations to a level which is adequate to protect worker, a respiratory protection program that meets or is equivalent to OSHA 29 CFR 1910.134 should be followed. Check with respiratory protective equipment suppliers.

Appropriate Engineering Controls

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Provide exhaust ventilation or other engineering controls to keep the airborne concentrations of vapors below their respective threshold limit value.

Chemical Name	OSHA STEL (mg/m3)	OSHA STEL (ppm)	OSHA TWA (mg/m3)	OSHA TWA (ppm)	OSHA Carcinogen	OSHA Skin designation	OSHA Tables (Z1, Z2, Z3)	NIOSH STEL (mg/m3)
ISOPARAFFINI C			2000	500			1 1 1	- ATT
PETROLEUM DISTILLATE								

Chemical Name	NIOSH TWA (mg/m3)	NIOSH STEL (ppm)	NIOSH TWA (ppm)	NIOSH Carcinogen	ACGIH STEL (mg/m3)	ACGIH STEL (ppm)	ACGIH TWA (mg/m3)	ACGIH TWA (ppm)
ISOPARAFFINI C							[(L)[N159](L) [N800]]; [5 (I)	(L)[N159](L) [N800]
PETROLEUM							[N159]5 (I)	[]
DISTILLATE							[N800]];	

Chemical	ACGIH	ACGIH	ACGIH
Name	TLV Basis	Carcinogen	Notations
ISOPARAFFINI C PETROLEUM DISTILLATE	URT irr [N159]URT irr [N800]	[A2[N159]A2 [N800]]; [A4 [N159]A4 [N800]];	[A2[N159]A2 [N800]]; [A4 [N159]A4 [N800]];

URT - Upper respiratory tract

SECTION 9) PHYSICAL AND CHEMICAL PROPERTIES

Physical and Chemical Properties

Non-Volatiles wt%	25 +/- 5
Density	6.84000 lb/gal
Density VOC	5.15000 lb/gal
Specific Gravity	0.82000
Appearance	White Paste
Odor Threshold	N/A
Odor Description	Slight hydrocarbon solvent odor
рН	N/A
Water Solubility	Negligible
Flammability	Flash point at or above 100°F/38°C and less than 200°F/93°C
Flash Point Symbol	N/A
Flash Point	42 °C
Viscosity (cps @ 77°F)	Less than 1000
Lower Explosion Level	0.6
Upper Explosion Level	6
Vapor Pressure	0.37
Vapor Density	5.3000000000
Freezing Point	N/A
Melting Point	N/A
Low Boiling Point	318 °F
High Boiling Point	400 °F
Auto Ignition Temp	230 °C
Decomposition Pt	N/A

SECTION 10) STABILITY AND REACTIVITY

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N/A

Stability

Stable under normal conditions.

Conditions to Avoid

Avoid heat, spark, flame, direct sunlight and incompatible materials.

Hazardous Reactions/Polymerization

Will not occur.

Incompatible Materials

Avoid materials incompatible with water.

Hazardous Decomposition Products

No data available.

SECTION 11) TOXICOLOGICAL INFORMATION

Likely Route of Exposure

Inhalation, ingestion, skin absorption, eye contact.

Aspiration Hazard

May be fatal if swallowed and enters airways

Carcinogenicity

No data available.

Germ Cell Mutagenicity

No data available.

Reproductive Toxicity

No data available.

Respiratory/Skin Sensitization

0064742-47-8 ISOPARAFFINIC PETROLEUM DISTILLATE

The substance defats the skin, which may cause dryness or cracking.

Serious Eye Damage/Irritation

Causes eye irritation

0064742-47-8 ISOPARAFFINIC PETROLEUM DISTILLATE

The vapour is mildly irritating to the eyes.

Skin Corrosion/Irritation

Causes skin irritation

Specific Target Organ Toxicity - Repeated Exposure

No data available.

Specific Target Organ Toxicity - Single Exposure

May cause drowsiness or dizziness

0064742-47-8 ISOPARAFFINIC PETROLEUM DISTILLATE

May cause effects on the central nervous system.

Acute Toxicity

Harmful if inhaled

0064742-47-8 ISOPARAFFINIC PETROLEUM DISTILLATE

If swallowed, can easily enter the airways and could result in aspiration pneumonitis.

If swallowed, can easily enter the airways and could result in aspiration pneumonitis. Inhalation of high concentrations may cause dizziness, anesthesia, unconsciousness.

Likely Routes of Exposure

0064742-47-8 ISOPARAFFINIC PETROLEUM DISTILLATE

The substance can be absorbed into the body by inhalation of its vapour and by ingestion.

SECTION 12) ECOLOGICAL INFORMATION

Toxicity

No data available.

Other Adverse Effects

No data available.

Persistence and Degradability

0064742-47-8 ISOPARAFFINIC PETROLEUM DISTILLATE

Expected to be inherently biodegradable. The volatile constituents will oxidize rapidly by photochemical reactions in air.

Mobility in Soil

0064742-47-8 ISOPARAFFINIC PETROLEUM DISTILLATE

Floats on water. Contains volatile constituents. Evaporates within a day from water or soil surfaces. Large volumes may penetrate soil and could contaminate groundwater.

SECTION 13) DISPOSAL CONSIDERATIONS

Waste Disposal

It is the responsibility of the user of the product to determine at the time of disposal whether the product meets local criteria for hazardous waste. Waste management should be in full compliance with national, state and local laws.

Empty Containers retain product residue which may exhibit hazards of material, therefore do not pressurize, cut, glaze, weld or use for any other purposes. Return drums to reclamation centers for proper cleaning and reuse.

SECTION 14) TRANSPORT INFORMATION

U.S. DOT Information

UN number: UN 1268 Proper shipping name: Petroleum Distillates, N.O.S. (ISOPARAFFINIC PETROLEUM DISTILLATE) Hazard class: Combustible liquid Packaging group: III Hazardous substance (RQ): No Data Available Toxic-Inhalation Hazard: No Data Available Marine Pollutant: No Data Available Note / Special Provision: No Data Available

IMDG Information

UN number: UN 1268 Proper shipping name: Petroleum Distillates, N.O.S. (ISOPARAFFINIC PETROLEUM DISTILLATE) Hazard class: 3 Packaging group: III Marine Pollutant: No Data Available Note / Special Provision: No Data Available

IATA Information

UN number: UN 1268 Proper shipping name: Petroleum Distillates, N.O.S. (ISOPARAFFINIC PETROLEUM DISTILLATE) Hazard class: 3 Packaging group: III Marine Pollutant: No Data Available Note / Special Provision: No Data Available

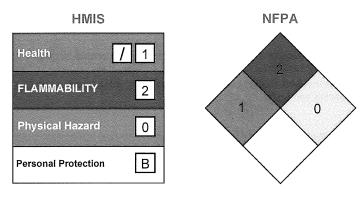
SECTION 15) REGULATORY INFORMATION

CAS	Chemical Name	% By Weight	Regulation List
0064742-47-8	ISOPARAFFINIC PETROLEUM DISTILLATE	72% - 76%	SARA312,VOC,IARCCarcinogen,TS CA,TSCA_UVCB - CHEMICAL SUBSTANCES OF UNKNOWN OR VARIABLE COMPOSITION, COMPLEX REACTION PRODUCTS AND BIOLOGICAL MATERIALS
0009002-88-4	POLYETHYLENE	20% - 22%	SARA312,IARCCarcinogen,TSCA
0009010-79-1	Polypropylene	1.5% - 4%	SARA312,TSCA

SECTION 16) OTHER INFORMATION INCLUDING INFORMATION ON PREPARATION AND REVISION OF THE SDS

Glossary

ACGIH- American Conference of Governmental Industrial Hygienists; ANSI- American National Standards Institute; Canadian TDG Canadian Transportation of Dangerous Goods; CAS- Chemical Abstract Service; Chemtrec- Chemical Transportation Emergency Center (US); CHIP- Chemical Hazard Information and Packaging; DSL- Domestic Substances List; EC- Equivalent Concentration; EH40 (UK)-HSE Guidance Note EH40 Occupational Exposure Limits; EPCRA- mergency Planning and Community Right-To-Know Act; SLEffects screening levels; HMIS- Hazardous Material Information Service; LC- Lethal Concentration; LD- Lethal Dose; NFPA- National Fire Protection Association; OEL- Occupational Exposure Limits; OSHA- Occupational Safety and Health Administration, US Department of Labor; PEL- Permissible Exposure Limit; ARA (Title III)- Superfund Amendments and eauthorization Act; SARA 313- Superfund Amendments and Reauthorization on Environmental Quality; TLV- Threshold Limit Value; TSCA- Toxic Substances Control Act Public Law 94-469; TWA - Time Weighted Value; US DOT- US Department of Transportation; WHMIS- Workplace Hazardous Materials Information System.



(*) - Chronic effects

Caution: HMIS® ratings are based on a 0-4 rating scale, with 0 representing minimal hazards or risks, and 4 representing significant hazards or risks

DISCLAIMER

To the best of our knowledge, the information contained herein is accurate. However, neither the above named supplier nor any of its subsidiaries assumes any liability whatsoever for the accuracy or completeness of the information contained herein. Final determination of suitability of any material is the sole responsibility of the user. All materials may present unknown hazards and should be used with caution. Although certain hazards are described herein, we cannot guarantee that these are the only hazards that exist. The above information pertains to this product as currently formulated, and is based on the information available at this time. Addition of reducers or other additives to this product may substantially alter the composition and hazards of the product. Since conditions of use are outside our control, we make no warranties, express or implied, and assume no liability in connection with any use of this information.

Version 2.1: Revision Date: Jan 08, 2019



Cytech Products Inc. 906 Peterson Drive Elizabethtown, KY 42701

Rikeizai H-179K-7

Release Agent

<u>Volume</u>

78.69%

Acoustical / Carpet / High Resilience Foam

The composition given below is the composition of the Product as Formulated. Slight variation may occur in individual batches.

Weight/Gallon Non-Volatiles (% by weight) Non-Volatiles (% by volume) Volatiles (% by weight) Volatiles (% by volume) Density of Blend (kg/liter) Volatile Organic Compounds Minus Exempt Solvents.

24.51% 21.31% 75.49% 78.69% 0.8179

6.8221 Lbs/Gal

75.49% by wt.; 5.150 lbs/gal; 617.11 grams/liter

For Environmental Reporting and Permitting use Actual Batch Data and Processing Information. For Further Assistance, Contact Your Sales Representative.

Density Of Solvent Blend 6.5202 Pounds/Gallon

This product is **not** photochemically reactive as per SCAQMD rule 102.

Volatile Composition:

CAS#

64742-47-8

<u>Composition</u> Hydrocarbon Naphtha <u>Weight</u> 75.49%

Percent of Volatile

Solvent Blend NFPA Hazard Rating

 $H = 1 \qquad F = 2 \qquad R = 0$

Hazardous and Safety Information

This product is in solvent base form. The product should always be kept away from excessive heat or open flames. Gloves should be worn when handling the product. If spills occur, dispose of them in accordance with local, federal, state or international regulations. Hands and arms should be washed thoroughly with soap and water after contact. The Safety Data Sheet (SDS) for this product contains detailed information regarding all hazardous and safety information. Always read the SDS that accompanies this product before use.

The information, statements, and data contained herein are to the best of Cytech Products' knowledge and belief, accurate and reliable as of the date issued. Cytech Products does not warrant or guarantee their accuracy or reliability, and Cytech Products shall not be liable for any loss or damage arising out of the use thereof. This information is provided as a guide, with the understanding that it is the customer's responsibility to safely and correctly use our products based on the customer's actual conditions. Therefore, Cytech Products Inc. assumes no warranty, expressed or implied, regarding the actual use, storage, handling or merchantability of this product for a particular purpose. The information provided herein should not be assumed as recommendations to infringe on any patents or licenses.

AQD H-179K-7 1.0 05/09/17

Phone: 270-769-1046 Fax: 270-769-1048 Email: sales@cytechproducts.com



SAFETY DATA SHEET

Classified in accordance 29 CFR 1910.1200

1. Identification

Product identifier: TEGOSTAB B 8737 LF 2

Chemical name: Preparation of organomodified polysiloxanes

Other means of identification

Recommended restrictions

Recommended use: Industrial Use Restrictions on use: None known.

Manufacturer/Importer/Distributor Information

Company Name	: Evonik Corporation Nutrition & Care PO Box 34628 Richmond, VA 23234 USA

Telephone : +1 804 727 0700

Fax : +1 804 727 0845

E-mail : us-productsafety-cs@evonik.com

Emergency telephone number:

24-Hour Health	: +1 800 424 9300 (CHEMTREC - US & CANADA)
Emergency	+1 800 681 9531 (CHEMTREC MEXICO)
0, 1	+1 703 527 3887 (CHEMTREC WORLD)

2. Hazard(s) identification

Hazard Classification

Physical Hazards

Flammable liquids

Category 4

Health Hazards

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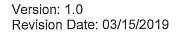
Toxic to reproduction

Category 2

Label Elements

Hazard Symbol:





Signal Word: Hazard Statement:	Warning Combustible liquid. Suspected of damaging fertility or the unborn child.
Precautionary Statements	
Prevention:	Obtain special instructions before use. Do not handle until all safety precautions have been read and understood. Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking. Wear protective gloves/eye protection/face protection.
Response:	IF exposed or concerned: Get medical advice/attention.
Storage:	Store in a well-ventilated place. Keep cool.
Hazard(s) not otherwise classified (HNOC):	None.

3. Composition/information on ingredients	

Chemical name:

Preparation of organomodified polysiloxanes

ΟΠΙΚ

POWER TO CREATE

Mixtures

Chemical Identity	CAS number	Content in percent (%)*
Hexamethyl disiloxane	107-46-0	0.5 - 1%
propane-1,2-diol	57-55-6	<0.5%
2-Propanol, 1,1',1"-nitrilotris-	122-20-3	<0.005%
Phenol, 2,6-bis(1,1-dimethylethyl)-4-	128-37-0	<0.001%

* All concentrations are percent by weight unless ingredient is a gas. Gas concentrations are in percent by volume.

The exact concentration has been withheld as a trade secret.

4. First-aid measures

Description of necessary first-aid measures

General information:	Remove soiled or soaked clothing immediately
Inhalation:	Ensure supply of fresh air. In the event of symptoms seek medical advice.
Skin Contact:	In case of contact with skin wash off with soap and water. In the event of symptoms seek medical advice.
Eye contact:	In case of contact with eyes rinse thoroughly with water. In the event of symptoms seek medical advice.
Ingestion:	Thoroughly clean the mouth with water In the event of symptoms seek medical advice.

Version: 1.0 Revision Date: 03/15/2019

	Version: 1.0 Revision Date: 03/15/2019
Personal Protection for First- aid Responders:	Do not inhale explosion and/or combustion gases, Self-contained breathing apparatus.
Most important symptoms/effec	cts, acute and delayed
Symptoms:	No special hints.
Hazards:	No data available.
Indication of immediate medica	l attention and special treatment needed
Treatment:	Treat symptomatically.
5. Fire-fighting measures	
Suitable (and unsuitable) exting	guishing media
Suitable extinguishing media:	foam, carbon dioxide, dry powder, water spray.
Unsuitable extinguishing media:	Full water jet
Specific hazards arising from the chemical:	In the event of fire the following can be released: - Carbon monoxide, carbon dioxide, silicon dioxide Under certain conditions of combustion traces of other toxic substances cannot be excluded
Special protective equipment a	and precautions for firefighters
Special fire fighting procedures:	Take action to prevent static discharges. Keep away from sources of ignition - no smoking. Vapours may form explosive mixtures with air Cool endangered containers by water spray
Special protective equipment for fire-fighters:	Do not inhale explosion and/or combustion gases Self-contained breathing apparatus.
6. Accidental release measures	
Personal precautions, protective equipment and emergency procedures:	Use personal protective equipment. Ensure adequate ventilation. Keep away sources of ignition.
Methods and material for containment and cleaning up:	Take up with absorbent material (eg sand, kieselguhr, universal binder) Dispose of absorbed material in accordance with the regulations.
Environmental Precautions:	Do not allow to enter drains or waterways Do not discharge into the subsoil/soil.
7. Handling and storage	
Handling	
Technical measures (e.g. Loca and general ventilation):	
Safe handling advice:	Avoid contact with skin and eyes. Do not inhale gases/vapours/aerosols.No special measures necessary if stored and handled as prescribed.

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special measures necessary if stored and handled as prescribed.



Hygiene measures:	Do not eat, drink or smoke when working. Wash hands before breaks and immediately after handling the product. Remove soiled or soaked clothing immediately.
Storage	
Safe storage conditions:	Keep container tightly closed and in a well-ventilated place.
Safe packaging materials:	No data available.
8. Exposure controls/personal	protection
Control Parameters	
Occupational Exposure Li	mits
	None of the components have assigned exposure limits. Hazardous components without workplace control parameters
Appropriate Engineering Controls	No data available.
Individual protection measure	s, such as personal protective equipment
Eye/face protection:	This product is not classified as a hazardous substance. Any necessity for eye protection must be determined within the scope of a risk assessment.
Skin Protection Hand Protection:	Material: gloves made of nitril (NBR) Break-through time: > 480 min Material: gloves made of chloroprene (CR, e.g. Neoprene) Break-through time: > 480 min Material: protective gloves made of fluorinated rubber (FKM, e.g. Viton) Break-through time: > 480 min
Skin and Body Protection:	protective clothing
Respiratory Protection:	in case of formation of vapours/aerosols: Short term: filter apparatus, combination filter A-P2
Hygiene measures:	Do not eat, drink or smoke when working. Wash hands before breaks and immediately after handling the product. Remove soiled or soaked clothing immediately.
9. Physical and chemical prope	rties
Annoaranco	

liquid
liquid
colourless to yellowish
Characteristic
not measured
not determined
not measured
not measured
147.20 °F (DIN EN ISO 2719)
not measured



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Explosive limit - upper (%):	not measured	
Explosive limit - lower (%):	not measured	
Vapor pressure:	not measured	
Vapor density (air=1):	not measured	
Density:	0.958 - 0.978 g/cm3 (25 °C) (DIN 51757)	
Relative density:	No data available.	
Solubility(ies)		
Solubility in Water:	Insoluble	
Solubility (other):	not measured	
Partition coefficient (n-octano	l/water): not measured	
Self Ignition Temperature:	not measured	
Decomposition Temperature:	not measured	
Kinematic viscosity:	No data available.	
Dynamic viscosity:	20 - 40 mPa.s (25 °C, DIN 53019)	
Other information		
Explosive properties:	not measured	
Oxidizing properties:	not measured	
Minimum ignition temperature		
Metal Corrosion:	not measured	
10. Stability and reactivity		
	see section "Possibility of hazardous reactions"	
Reactivity:	SEE SECTOR 1 Dasibility of hazardoud rodations	
Chemical Stability:	The product is stable under normal conditions.	
Possibility of hazardous reactions:	Reactions with acids. Reactions with alkalies.	

Conditions to avoid:	Unknown
Incompatible Materials:	Unknown
Hazardous Decomposition Products:	None with proper storage and handling.

11. Toxicological information

Inf	ormation on likely routes of ex Inhalation:	xposure If handled correctly, not a relevant route of exposure. Information on effects are given below.
	Skin Contact:	Relevant route of exposure. Information on effects are given below.
	Eye contact:	Relevant route of exposure. Information on effects are given below.
- Autor	Ingestion:	If handled correctly, not a relevant route of exposure. Information on effects are given below.

Symptoms related to the physical, chemical and toxicological characteristics

Inhalation:	No data available.
minalation	no data aranasier



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Eye contact:	No data available.	
Ingestion:	No data available.	
Information on toxicological effects		
Acute toxicity (list all possible	routes of exposure)	
Oral Product:	no data available	
Dermal Product:	no data available	
Inhalation Product:	no data available	
Repeated dose toxicity Product:	no data available	
Skin Corrosion/Irritation Product:	no data available	
Serious Eye Damage/Eye Irritati Product:	on no data available	
Respiratory or Skin Sensitizatio Product:	n no data available	
Carcinogenicity Product:	No data available.	
IARC Monographs on the Evaluation No carcinogens present or n	ation of Carcinogenic Risks to Humans: one present in regulated quantities	
US. National Toxicology Program (NTP) Report on Carcinogens: No carcinogens present or none present in regulated quantities		
US. OSHA Specifically Regulated Substances (29 CFR 1910.1001-1050): No carcinogens present or none present in regulated quantities		
Germ Cell Mutagenicity		
In vitro Product:	No data available.	
Components: Hexamethyl disiloxane	Ames test (OECD 471): negative	
In vivo Product:	No data available.	
Components: Hexamethyl disiloxane	Chromosomal aberration (OECD 475): negative	
Reproductive toxicity Product:	No data available.	



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Specific Target Organ	Toxicity - Repeated Exposure
Product:	no data available

Acute hazards to the aquatic environment:

Aspiration Hazard Product:	Not classified
	Proper use provided, no adverse health effects have been observed or have been come to our knowledge.

12. Ecological information

Ecotoxicity:

Fish Product:	No data available.	
Aquatic Invertebrates Product:	No data available.	
Chronic hazards to the aquati	c environment:	
Fish Product:	No data available.	
Aquatic Invertebrates Product:	No data available.	
Toxicity to Aquatic Plants Product:	No data available.	
Persistence and Degradability		
Biodegradation Product:	No data available.	
BOD/COD Ratio Product:	No data available.	
Bioaccumulative potential Bioconcentration Factor (BCF) Product: No data available.		
Partition Coefficient n-octanol / v Product:	water (log Kow) Log Kow: not measured	
Mobility in soil:	No data available.	

Components:



Version: 1.0 Revision Date: 03/15/2019

Hexamethyl disiloxane propane-1,2-diol 2-Propanol, 1,1',1''- nitrilotris- Phenol, 2,6-bis(1,1- dimethylethyl)-4-methyl-	No data available. No data available. No data available. No data available.
Other adverse effects:	Do not allow to enter soil, waterways or waste water canal.
13. Disposal considerations	
Disposal methods:	In accordance with local authority regulations, take to special waste incineration plant
Contaminated Packaging:	If empty contaminated containers are recycled or disposed of, the receiver must be informed about possible hazards.

14. Transport information

Domestic regulation

49 CFR

Not regulated as a dangerous good

International Regulations

UNRTDG

Not regulated as a dangerous good

IATA-DGR

Not regulated as a dangerous good

IMDG-Code

Not regulated as a dangerous good

Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code

Not applicable for product as supplied.

15. Regulatory information

US Federal Regulations

TSCA Section 12(b) Export Notification (40 CFR 707, Subpt. D) None present or none present in regulated quantities.

US. Toxic Substances Control Act (TSCA) Section 5(a)(2) Final Significant New Use Rules (SNURs) (40 CFR 721, Subpt E)

None present or none present in regulated quantities.

US. OSHA Specifically Regulated Substances (29 CFR 1910.1001-1050)

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CERCLA Hazardous Substance List (40 CFR 302.4):

None present or none present in regulated quantities.

Superfund Amendments and Reauthorization Act of 1986 (SARA)

Hazard categories

Flammable (gases, aerosols, liquids, or solids), Reproductive toxicity

SARA 302 Extremely Hazardous Substance

None present or none present in regulated quantities.

US. EPCRA (SARA Title III) Section 304 Extremely Hazardous Substances Reporting Quantities and the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) Hazardous Substances

SARA 311/312 Hazardous Chemical Chemical Identity Threshold Planning Quantity

SARA 313 (TRI Reporting)

None present or none present in regulated quantities.

Clean Air Act (CAA) Section 112(r) Accidental Release Prevention (40 CFR 68.130): None present or none present in regulated quantities.

Clean Water Act Section 311 Hazardous Substances (40 CFR 117.3)

None present or none present in regulated quantities. **US State Regulations**

> **US. California Proposition 65** No ingredient requiring a warning under CA Prop 65.

- US. New Jersey Worker and Community Right-to-Know Act No ingredient regulated by NJ Right-to-Know Law present.
- US. Massachusetts RTK Substance List No ingredient regulated by MA Right-to-Know Law present.
- US. Pennsylvania RTK Hazardous Substances No ingredient regulated by PA Right-to-Know Law present.

US. Rhode Island RTK

No ingredient regulated by RI Right-to-Know Law present.

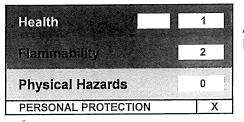
Inventory Status:

US TSCA Inventory: Canada DSL Inventory List: Included on Inventory. Included on Inventory.



16.Other information, including date of preparation or last revision

HMIS Hazard ID



Ask supervisor or safety specialist for handling instructions

Hazard rating: 0 - Minimal; 1 - Slight; 2 - Moderate; 3 - Serious; 4 - Severe; RNP - Rating not possible; *Chronic health effect

Issue Date:	03/15/2019
Version #:	1.0
Further Information:	none
Revision Information:	Changes since the last version are highlighted in the margin. This version replaces all previous versions.
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SAFETY DATA SHEET

Classified in accordance 29 CFR 1910.1200

1. Identification

Product identifier: TEGOSTAB B 8724 LF 2

Chemical name:

Preparation of organomodified polysiloxanes

Other means of identification

Recommended restrictions

Recommended use: Industrial Use **Restrictions on use:** None known.

Manufacturer/Importer/Distributor Information

Company Name	: Evonik Corporation Nutrition & Care PO Box 34628 Richmond, VA 23234 USA	
Telephone	: +1 804 727 0700	
Fax	: +1 804 727 0845	
E-mail	: us-productsafety-cs@evonik.com	
Emergency telephone number:		

24-Hour Health : +1 800 424 9300 (CHEMTREC - US & CANADA) Emergency +1 800 681 9531 (CHEMTREC MEXICO)

+1	703 527	3887 (0	CHEMTREC	WORLD)

2. Hazard(s) identification

Hazard Classification

Label Elements

Hazard Symbol:	No symbol
Signal Word:	No signal word.
Hazard Statement:	Not applicable
Precautionary Statements	Not a hazardous substance or mixture according to 29 CFR 1910.1200.

None.



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3. Composition/information on in	ngredients		
Chemical name: Preparation of organomodified poly	Chemical name: Preparation of organomodified polysiloxanes		
Mixtures			
Composition Comments:	No hazardous ingredients.		
4. First-aid measures			
Description of necessary first-a	id measures		
General information:	Remove soiled or soaked clothing immediately		
Inhalation:	Ensure supply of fresh air.		
Skin Contact:	In case of contact with skin wash off with soap and water. Consult a doctor if skin irritation persists.		
Eye contact:	In case of contact with eyes rinse thoroughly with plenty of water. If symptoms persist, seek medical advice.		
Ingestion:	Medical treatment		
Personal Protection for First- aid Responders:	Do not inhale explosion and/or combustion gases, Use self-contained breathing apparatus and wear full protective suit.		
Most important symptoms/effec	cts, acute and delayed		
Symptoms:	No information is on file to date regarding acute and/or delayed post- exposure symptoms and effects.		
Hazards:	No data available.		
Indication of immediate medica	I attention and special treatment needed		
Treatment:	Treat symptomatically.		
5. Fire-fighting measures			
Suitable (and unsuitable) exting	guishing media		
Suitable extinguishing media:	foam, carbon dioxide, dry powder, water spray.		
Unsuitable extinguishing media:	Full water jet		
Specific hazards arising from the chemical:	In the event of fire the following can be released: - Carbon monoxide, carbon dioxide, silicon dioxide Under certain conditions of combustion traces of other toxic substances cannot be excluded		
Special protective equipment and precautions for firefighters			
Special fire fighting procedures:	No specific precautions.		
Special protective equipment for fire-fighters:	Do not inhale explosion and/or combustion gases Use self-contained breathing apparatus and wear full protective suit.		



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6. Accidental release measures	
Personal precautions, protective equipment and emergency procedures:	Use personal protective equipment.
Methods and material for containment and cleaning up:	Take up with absorbent material (eg sand, kieselguhr, universal binder) Dispose of absorbed material in accordance with the regulations.
Environmental Precautions:	Do not allow to enter drains or waterways Do not discharge into the subsoil/soil.
7. Handling and storage	
landling	
Technical measures (e.g. Local and general ventilation):	No data available.
Safe handling advice:	Avoid contact with skin and eyes. Ensure adequate ventilation.
Contact avoidance measures:	No data available.
Hygiene measures:	Do not eat, drink or smoke when working. Wash hands before breaks and immediately after handling the product. Remove soiled or soaked clothing immediately.
Storage	
Safe storage conditions:	Keep container tightly closed and in a well-ventilated place.
Safe packaging materials:	No data available.
3. Exposure controls/personal pro	otection
Control Parameters Occupational Exposure Limit	is
	None of the components have assigned exposure limits.
Appropriate Engineering Controls	No data available.
Individual protection measures, s	such as personal protective equipment
Eye/face protection:	goggles with side pieces
Skin Protection Hand Protection:	Additional Information: gloves made of nitril (NBR)
Skin and Body Protection:	light protective clothing
Respiratory Protection:	in case of formation of vapours/aerosols: Short term: filter apparatus, Filter A-P2
Hygiene measures:	Do not eat, drink or smoke when working. Wash hands before breaks and

Hygiene measures: Do not eat, drink or smoke when working. Wash hands before breaks and immediately after handling the product. Remove soiled or soaked clothing immediately.



Version: 1.1 Revision Date: 04/30/2019

9. Physical and chemical p	roperties
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of Thysical and chemical properties		
Appearance		
Physical state:	liquid	
Form:	liquid	
Color:	colourless to yellowish	
Odor:	Characteristic	
Odor Threshold:	not measured	
pH:	not determined	
Freezing point:	not measured	
Boiling Point:	not measured	
Flash Point:	363.20 °F (DIN EN ISO 2719)	
Evaporation Rate:	not measured	
Flammability (solid, gas):	no data available	
Explosive limit - upper (%):	not measured	
Explosive limit - lower (%):	not measured	
Vapor pressure:	not measured	
Vapor density (air=1):	not measured	
Density:	0.959 - 0.979 g/cm3 (25 °C) (DIN 51757)	
Relative density:	No data available.	
Solubility(ies)		
Solubility in Water:	Insoluble	
Solubility (other):	not measured	
Partition coefficient (n-octano	I/water): not measured	
Self Ignition Temperature:	, not measured	
Decomposition Temperature:	not measured	
Kinematic viscosity:	No data available.	
Dynamic viscosity:	20 - 40 mPa.s (25 °C, DIN 53019)	
-		
Other information	not measured	
Explosive properties:		
Oxidizing properties:	not measured	
Minimum ignition temperature	e: not measured not measured	
Metal Corrosion:	not measured	
10. Stability and reactivity		
Reactivity:	see section "Possibility of hazardous reactions"	
Chemical Stability:	The product is stable under normal conditions.	
Possibility of hazardous reactions:	Reactions with acids. Reactions with alkalies.	
Conditions to avoid:	Unknown	
Incompatible Materials:	Unknown	
Hazardous Decomposition Products:	None with proper storage and handling.	
11. Toxicological information		



Information on likely route Inhalation:	s of exposure If handled correctly, not a relevant route of exposure. Information on effects are given below.
Skin Contact:	Relevant route of exposure. Information on effects are given below.
Eye contact:	Relevant route of exposure. Information on effects are given below.
Ingestion:	If handled correctly, not a relevant route of exposure. Information on effects are given below.

Symptoms related to the physical, chemical and toxicological characteristics

Inhalation:	No data available.
Skin Contact:	No data available.
Eye contact:	No data available.
Ingestion:	No data available.

Information on toxicological effects

Acute toxicity (list all possible routes of exposure)

Oral Product:	no data available
Dermal Product:	no data available
Inhalation Product:	no data available

Repeated dose toxicity	
Product:	no data available

Skin Corrosion/Irritation Product: no data available

- Serious Eye Damage/Eye Irritation Product: no data available
- Respiratory or Skin Sensitization Product: no data available
- Carcinogenicity Product: No data available.
- IARC Monographs on the Evaluation of Carcinogenic Risks to Humans: No carcinogens present or none present in regulated quantities
- US. National Toxicology Program (NTP) Report on Carcinogens: No carcinogens present or none present in regulated quantities
- US. OSHA Specifically Regulated Substances (29 CFR 1910.1001-1050): No carcinogens present or none present in regulated quantities



Germ Cell Mutagenicity

A - A	
In vitro Product:	No data available.
In vivo Product:	No data available.
Reproductive toxicity Product:	No data available.
Specific Target Organ Toxicity - Product:	Single Exposure no data available
Specific Target Organ Toxicity - Product:	Repeated Exposure no data available
Aspiration Hazard Product:	Not classified
Other effects:	Proper use provided, no adverse health effects have been observed or have been come to our knowledge. Due to the composition of the product it cannot be excluded: Irritates eyes and skin

12. Ecological information

Ecotoxicity:

Acute hazards to the aquatic o	environment:
Fish Product:	No data available.
Aquatic Invertebrates Product:	No data available.
Chronic hazards to the aquat	ic environment:
Fish Product:	No data available.
Aquatic Invertebrates Product:	No data available.
Toxicity to Aquatic Plants Product:	No data available.
Persistence and Degradability	
Biodegradation Product:	No data available.
BOD/COD Ratio Product:	No data available.
Bioaccumulative potential	

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Bioconcentration Factor (B Product:	SCF) No data available.				
Partition Coefficient n-octanol / Product:	water (log Kow) Log Kow: not measured				
Mobility in soil:	No data available.				
Other adverse effects:	Do not allow to enter soil, waterways or waste water canal.				
13. Disposal considerations					
Disposal methods:	In accordance with local authority regulations, take to special waste incineration plant				
Contaminated Packaging:	If empty contaminated containers are recycled or disposed of, the receiver must be informed about possible hazards.				
14. Transport information					

Domestic regulation

49 CFR

Not regulated as a dangerous good

International Regulations

UNRTDG

Not regulated as a dangerous good

IATA-DGR

Not regulated as a dangerous good

IMDG-Code

Not regulated as a dangerous good

Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code

Not applicable for product as supplied.

15. Regulatory information

US Federal Regulations

TSCA Section 12(b) Export Notification (40 CFR 707, Subpt. D) None present or none present in regulated quantities.

US. Toxic Substances Control Act (TSCA) Section 5(a)(2) Final Significant New Use Rules (SNURs) (40 CFR 721, Subpt E)

None present or none present in regulated quantities.

US. OSHA Specifically Regulated Substances (29 CFR 1910.1001-1050) None present or none present in regulated quantities.



CERCLA Hazardous Substance List (40 CFR 302.4):

None present or none present in regulated quantities.

Superfund Amendments and Reauthorization Act of 1986 (SARA)

Hazard categories

Not classified

SARA 302 Extremely Hazardous Substance

None present or none present in regulated quantities.

US. EPCRA (SARA Title III) Section 304 Extremely Hazardous Substances Reporting Quantities and the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) Hazardous Substances

SARA 311/312 Hazardous Chemical

None present or none present in regulated quantities.

SARA 313 (TRI Reporting)

None present or none present in regulated quantities.

Clean Air Act (CAA) Section 112(r) Accidental Release Prevention (40 CFR 68.130): None present or none present in regulated quantities.

Clean Water Act Section 311 Hazardous Substances (40 CFR 117.3)

None present or none present in regulated quantities. **US State Regulations**

US. California Proposition 65

No ingredient requiring a warning under CA Prop 65.

US. New Jersey Worker and Community Right-to-Know Act No ingredient regulated by NJ Right-to-Know Law present.

US. Massachusetts RTK - Substance List

No ingredient regulated by MA Right-to-Know Law present.

US. Pennsylvania RTK - Hazardous Substances No ingredient regulated by PA Right-to-Know Law present.

US. Rhode Island RTK

No ingredient regulated by RI Right-to-Know Law present.

Inventory Status:

US TSCA Inventory: In Canada DSL Inventory List: In

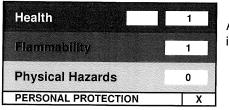
Included on Inventory. Included on Inventory.



16.Other information, including date of preparation or last revision

HMIS Hazard ID

 $\frac{i}{\lambda}n$



Ask supervisor or safety specialist for handling instructions

Hazard rating: 0 - Minimal; 1 - Slight; 2 - Moderate; 3 - Serious; 4 - Severe; RNP - Rating not possible; *Chronic health effect

Issue Date:	04/30/2019
Version #:	1.1
Further Information:	none
Revision Information:	Changes since the last version are highlighted in the margin. This version replaces all previous versions.
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Glycerine 99.7% USP Kosher

Safety Data Sheetaccording to Federal Register / Vol. 77, No. 58 / Monday, March 26, 2012 / Rules and RegulationsRevision Date: 12/3/2018Supersedes: 10/21/2013

Version: 2.0

SECTION 1: IDENTIFICATION OF THE SUBSTANCE/MIXTURE AND OF THE COMPANY

1.1. Product Identifier

A division of JACOB STERN & SONS, INC. - SINCE 1857-

Product Form: Substance Product Name: Glycerine 99.7% USP Kosher

CAS No: 56-81-5 Formula: C3H8O3/C3H5(OH)3 Synonyms: 1,2,3-Propanetriol; Glycerol; Glycerin

1.2. Intended Use of the Product

Use of the substance/mixture: NF Excipient

1.3. Name, Address, and Telephone of the Responsible Party

Company

Acme-Hardesty Co. 450 Sentry Parkway Blue Bell, PA 19422 T 866-226-3834 T 215-591-3610 www.acme-hardesty.com

1.4. Emergency Telephone Number

Emergency Number : 800-424-9300

For Chemical Emergency, Spill, Leak, Fire, Exposure, or Accident, call CHEMTREC - Day or Night

SECTION 2: HAZARDS IDENTIFICATION

2.1. Classification of the Substance or Mixture

Classification (GHS-US) Not classified

2.2. Label Elements

GHS-US Labeling Not applicable

2.3. Other Hazards No additional information available

No additional information available

2.4. <u>Unknown Acute Toxicity (GHS-US)</u>:

No data available

SECTION 3: COMPOSITION/INFORMATION ON INGREDIENTS

3.1. Substances

Name	Product Identifier	%	Classification (GHS-US)	
Glycerin	(CAS No) 56-81-5	99.7	Not classified	

Full text of H-phrases: see section 16

3.2. Mixtures

Not applicable

Full text of H-phrases: see section 16

SECTION 4: FIRST AID MEASURES

4.1. Description of First Aid Measures

First-aid Measures General: Never give anything by mouth to an unconscious person. If you feel unwell, seek medical advice (show the label where possible).

First-aid Measures After Inhalation: When symptoms occur: go into open air and ventilate suspected area.

First-aid Measures After Skin Contact: Remove contaminated clothing. Drench affected area with water for at least 15 minutes.

First-aid Measures After Eye Contact: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.

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First-aid Measures After Ingestion: Rinse mouth. Do NOT induce vomiting.

4.2. Most important symptoms and effects, both acute and delayed

Symptoms/Injuries: None expected under normal conditions of use.

Symptoms/Injuries After Inhalation: Not expected to present a significant inhalation hazard under anticipated conditions of normal use.

Symptoms/Injuries After Skin Contact: None under normal conditions.

Symptoms/Injuries After Eye Contact: Direct contact with the eyes is likely irritating.

Symptoms/Injuries After Ingestion: If a large quantity has been ingested: May cause nausea, vomiting, and diarrhea.

4.3. Indication of Any Immediate Medical Attention and Special Treatment Needed

If medical advice is needed, have product container or label at hand.

SECTION 5: FIREFIGHTING MEASURES

5.1. Extinguishing Media

Suitable Extinguishing Media: Use extinguishing media appropriate for surrounding fire. Water spray or fog, dry chemical powder, alcohol-resistant foam, carbon dioxide (CO2).

Unsuitable Extinguishing Media: None known.

5.2. Special Hazards Arising From the Substance or Mixture

Fire Hazard: Not flammable.

Explosion Hazard: Product is not explosive.

Reactivity: Stable at ambient temperature and under normal conditions of use.

5.3. Advice for Firefighters

Firefighting Instructions: Exercise caution when fighting any chemical fire.

Protection During Firefighting: Do not enter fire area without proper protective equipment, including respiratory protection.

SECTION 6: ACCIDENTAL RELEASE MEASURES

6.1. Personal Precautions, Protective Equipment and Emergency Procedures

General Measures: Handle in accordance with good industrial hygiene and safety practice.

6.1.1. For Non-emergency Personnel

Protective Equipment: Use appropriate personal protection equipment (PPE).

Emergency Procedures: Evacuate unnecessary personnel.

6.1.2. For Emergency Responders

Protective Equipment: Equip cleanup crew with proper protection.

Emergency Procedures: Ventilate area.

6.2. Environmental Precautions

Prevent entry to sewers and public waters.

6.3. Methods and Material for Containment and Cleaning Up

For Containment: Absorb and/or contain spill with inert material, then place in suitable container. **Methods for Cleaning Up:** Clear up spills immediately and dispose of waste safely.

6.4. Reference to Other Sections

See heading 8, Exposure Controls and Personal Protection.

SECTION 7: HANDLING AND STORAGE

7.1. Precautions for Safe Handling

Hygiene Measures: Handle in accordance with good industrial hygiene and safety procedures. Wash hands and other exposed areas with mild soap and water before eating, drinking, or smoking and again when leaving work. Do no eat, drink or smoke when using this product.

7.2. Conditions for Safe Storage, Including Any Incompatibilities

Storage Conditions: Store in a dry, cool and well-ventilated place. Keep container closed when not in use. **Incompatible Products:** Strong acids. Strong bases. Strong oxidizers.

7.3. Specific End Use(s) NF Excipient

SECTION 8: EXPOSURE CONTROLS/PERSONAL PROTECTION

8.1. Control Parameters

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Glycerin (56-81-5) USA OSHA OSHA PEL (TWA) (mg/m3) 5 mg/m³			
8.2. Exposure Controls	5 lig/iii		
Personal Protective Equipment	: Gloves. Safety glasses.		
Hand Protection	: Wear chemically resistant protective gloves. The breakthrough time of the selected		
	gloves must be greater than the intended use period.		
Eye Protection	: Chemical goggles or safety glasses.		
Respiratory Protection Other Information	 In case of inadequate ventilation wear respiratory protection. When using, do not eat, drink or smoke. 		
SECTION 9: PHYSICAL AND CHI			
9.1. Information on Basic Physical a	•		
Physical State	: Liquid		
Appearance Odor	: Colorless. Viscous. : Characteristic.		
Odor Threshold	: No data available		
pH	: No data available		
Relative Evaporation Rate (butylacetate=1			
Melting Point	: 18 °C (64.4°C)		
Freezing Point	: No data available		
Boiling Point	: 290 °C (554°F) (decomposition)		
Flash Point	: 177 °C (350.6°F) Cleveland Open Cup		
Auto-ignition Temperature	: 400 °C (752°F)		
Decomposition Temperature	: No data available		
Flammability (solid, gas)	: No data available		
Vapor Pressure	: < 0.01 hPa		
Relative Vapor Density at 20 °C	: No data available		
Relative Density	: No data available		
Specific Gravity	: 1.26 g/cm ³		
Solubility	: Fully miscible.		
Log Pow	: No data available		
Log Kow	: -1.76		
Viscosity, Kinematic	: 1300 mPas @ 20 °C		
Viscosity, Dynamic	: No data available		
Explosive Properties	: No data available		
Oxidizing Properties	: No data available		
Explosive Limits : Not applicable			
9.2. Other Information			

VOC content

: ≤0.5 %

SECTION 10: STABILITY AND REACTIVITY

10.1 Reactivity: Stable at ambient temperature and under normal conditions of use.

10.2 Chemical Stability: Product is stable.

10.3 Possibility of Hazardous Reactions: Hazardous polymerization will not occur.

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10.4 Conditions to Avoid: Direct sunlight.Extremely high or low temperatures.

10.5 Incompatible Materials: Strong acids. Strong bases. Strong oxidizers.

10.6 Hazardous Decomposition Products: Under fire conditions this material may produce hazardous carbon dioxide (CO2), carbon monoxide (CO), various low molecular weight hydrocarbons, and smoke.

SECTION 11: TOXICOLOGICAL INFORMATION

11.1. Information On Toxicological Effects

Acute Toxicity : Not classified

Glycerin (56-81-5)	
LD50 Dermal Rabbit	> 10 g/kg
LC50 Inhalation Rat (mg/l)	> 570 mg/m³ (Exposure time: 1 h)
ATE (Oral)	12600.000 mg/kg

Skin Corrosion/Irritation: Not classified Serious Eye Damage/Irritation: Not classified

Respiratory or Skin Sensitization: Not classified Germ Cell Mutagenicity: Not classified Carcinogenicity: Not classified

Reproductive Toxicity: Not classified Specific Target Organ Toxicity (Single Exposure): Not classified

Specific Target Organ Toxicity (Repeated Exposure): Not classified

Aspiration Hazard: Not classified

Symptoms/Injuries After Inhalation: Not expected to present a significant inhalation hazard under anticipated conditions of normal use.

Symptoms/Injuries After Skin Contact: None under normal conditions.

Symptoms/Injuries After Eye Contact: Direct contact with the eyes is likely irritating.

Symptoms/Injuries After Ingestion: If a large quantity has been ingested: May cause nausea, vomiting, and diarrhea.

SECTION 12: ECOLOGICAL INFORMATION

12.1. Toxicity

Glycerin (56-81-5)	
LC50 Fish 1	51 (51 - 57) ml/l (Exposure time: 96 h - Species: Oncorhynchus mykiss [static])
EC50 Daphnia 1	> 500 mg/l (Exposure time: 24 h - Species: Daphnia magna)

12.2. Persistence and Degradability

Glycerine 99.7% USP Kosher		
Persistence and Degradability	The substance is biodegradable. Unlikely to persist.	
12.3. Bioaccumulative Potential		

12.3. Bioaccumulative Potential

Glycerine 99.7% USP Kosher		
Log Pow	-1.76 Based on the n-octanol/water partition coefficient accumulation in organisms is not expected.	
Bioaccumulative Potential		
Glycerin (56-81-5)		
BCF fish 1	(no bioaccumulation)	

Log Pow

12.4. Mobility in Soil No additional information available

12.5. Other Adverse Effects

Other Information

: Avoid release to the environment.

SECTION 13: DISPOSAL CONSIDERATIONS

13.1. Waste treatment methods

Waste Disposal Recommendations: Dispose of waste material in accordance with all local, regional, national, and international regulations.

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SECTION 14: TRANSPORT INFORMATION

In Accordance With ICAO/IATA/DOT/TDG

14.1. UN Number Not applicable

14.2. UN Proper Shipping Name Not regulated for transport.

14.3. Additional Information

Other information

: No supplementary information available.

Transport by Sea Not regulated for transport.

Air Transport Not regulated for transport.

SECTION 15: REGULATORY INFORMATION

15.1 US Federal Regulations

Glycerin (56-81-5)	
Listed on the United States TSCA (Toxic Substan	nces Control Act) inventory
EPA TSCA Regulatory Flag	Y2 - Y2 - indicates an exempt polymer that is a polyester and is made only from reactants included in a specified list of low concern reactants that comprises one of the eligibility criteria for the exemption rule.

15.2 US State Regulations

Glycerin (56-81-5)

U.S Connecticut - Hazardous Air Pollutants - HLVs (30 min)	
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- U.S. Connecticut Hazardous Air Pollutants HLVs (8 hr)
- U.S. Idaho Occupational Exposure Limits TWAs
- U.S. Massachusetts Right To Know List
- U.S. Michigan Occupational Exposure Limits TWAs
- U.S. Minnesota Hazardous Substance List
- U.S. Minnesota Permissible Exposure Limits TWAs
- U.S. New Hampshire Regulated Toxic Air Pollutants Ambient Air Levels (AALs) 24-Hour
- U.S. New Hampshire Regulated Toxic Air Pollutants Ambient Air Levels (AALs) Annual
- U.S. New Jersey Right to Know Hazardous Substance List
- U.S. North Dakota Air Pollutants Guideline Concentrations 8-Hour
- U.S. Oregon Permissible Exposure Limits TWAs
- U.S. Pennsylvania RTK (Right to Know) List
- U.S. Tennessee Occupational Exposure Limits TWAs
- U.S. Texas Effects Screening Levels Long Term
- U.S. Texas Effects Screening Levels Short Term
- U.S. Vermont Permissible Exposure Limits TWAs
- U.S. Washington Permissible Exposure Limits STELs
- U.S. Washington Permissible Exposure Limits TWAs

Global Inventories

TSCA	Listed - CAS # 56-81-5
DSL/NDSL	Listed.
EINECS/ELINCS	Listed EINECS 200-289-5
ENCS	Listed. Nikkaji Number - J1.916J
KECL	Listed. KE-29297
PICCS	Listed.
AICS	Listed.

Legend

TSCA - United States Toxic Substances Control Act Section 8(b) Inventory DSL/NDSL - Canadian Domestic Substances List/Non-Domestic Substances List EINECS/ELINCS - European Inventory of Existing Chemical Substances/European List of Notified Chemical Substances ENCS - Japan Existing and New Chemical Substances

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- KECL Korean Existing and Evaluated Chemical Substances
- PICCS Philippines Inventory of Chemicals and Chemical Substances
- AICS Australian Inventory of Chemical Substances

SECTION 16: OTHER INFORMATION

Other Information : This document has been prepared in accordance with the SDS requirements of the OSHA Hazard Communication Standard 29 CFR 1910.1200.

The data herein are based on our current knowledge and believed to be reliable. Acme-Hardesty Co., provides this information without any representation or warranty, expressed or implied, regarding its accuracy or correctness.

Users must make their own determination that handling, storage, use and disposal of the product in the anticipated manner is safe and appropriate. Because these actions of the user are out of our control, and may be beyond our knowledge, we do not assume responsibility and expressly disclaim liability for loss, damage, expense or any other claim arising out of or in any way connected with the handling, storage, use or disposal of the product or container.

SDS US (GHS HazCom) - US Only

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DL-02596 Black LV Specification: 000000038190

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Version Number: 02



Revision Date: 08-21-2015

	1. Identification	
	Product Identifier	DL-02596 Black LV
	Other means of identification	
	SAP Specification	00000038190
	Recommended use	Not available.
	Recommended restrictions	None known.
	Manufacturer/Importer/Supplier	/Distributor information
	Manufacturer	
	Company	Chromaflo Technologies Corporation 2600 Michigan Avenue Ashtabula,OH 44005-0816 USA
	Telephone	440-997-5137
	Telefax	440-992-3613
	NA; EMERGENCY NUMBER	866-519-4752
	GLOBAL: EMERGENCY NUMBER	(+1) 760-476-3962
	CANADA: CANUTEC EMERGENCY NUMBER	613-996-6666
·	Product Regulatory Services	ehs_americas@chromaflo.com
	2. Hazard(s) identification	
	Physical hazards	Not classified,
	Health hazards	Not classified.
	OSHA defined hazards	Not classified.
	Label elements	
	Hazard symbol	None.
	Signal word	None.
	Hazard statement	The mixture does not meet the criteria for classification.
•	Precautionary statement	
	Prevention	Observe good industrial hygiene practices.
	Response	Wash hands after handling.
	Storage	Store away from incompatible materials.
	Disposal	Dispose of waste and residues in accordance with local author

 Storage
 Store away from incompatible materials.

 Disposal
 Dispose of waste and residues in accordance with local authority requirements.

 Hazard(s) not otherwise classified (HNOC)
 None known.

 Supplemental information
 If product is in liquid or paste form, physical or health hazards listed related to dust are a

If product is in liquid or paste form, physical or health hazards listed related to dust are not considered significant. However, product may contain substances that could be potential hazards if caused to become airborne due to grinding, sanding or other abrasive processes.

3. Composition/information on ingredients

Mixtures

Chemical name	Common name and synonyms	CAS number	%
Carbon Black		1333-86-4	20 - 40
Oligomeric Hydrocarbons*		Proprietary*	2.5 - 10
Other components below reportable levels			60 - 80

*Designates that a specific chemical identity and/or percentage of composition has been withheld as a trade secret.

4. First-aid measures

 $\mathbb{E}[\frac{n}{2}]$

	in not una medoures	
	Inhalation	Move to fresh air, Call a physician if symptoms develop or persist.
and the second second	Skin contact	Wash off with soap and water. Get medical attention if irritation develops and persists.
	Eye contact	Rinse with water. Get medical attention if irritation develops and persists.
	Ingestion	Rinse mouth. Get medical attention if symptoms occur,
	Most important symptoms/effects, acute and delayed	Direct contact with eyes may cause temporary irritation.
	Indication of immediate medical attention and special treatment needed	Treat symptomatically.
	General Information	Ensure that medical personnel are aware of the material(s) involved, and take precautions to protect themselves.
	5. Fire-fighting measures	
	Suitable extinguishing media	Water fog. Foam. Dry chemical powder. Carbon dioxide (CO2),
	Unsuitable extinguishing media	Do not use water jet as an extinguisher, as this will spread the fire.
	Specific hazards arising from the chemical	During fire, gases hazardous to health may be formed.
	Special protective equipment and precautions for firefighters	Self-contained breathing apparatus and full protective clothing must be worn in case of fire,
	Fire fighting	Move containers from fire area if you can do so without risk.

Move containers from fire area if you can do so without risk.

Use standard firefighting procedures and consider the hazards of other involved materials. No unusual fire or explosion hazards noted.

6. Accidental release measures

equipment/instructions **Specific methods**

General fire hazards

1000000	Personal precautions, protective equipment and emergency procedures	Keep unnecessary personnel away. Keep people away from and upwind of spill/leak. Keep out of low areas. Do not touch damaged containers or spilled material unless wearing appropriate protective clothing. For personal protection, see section 8 of the SDS,
	Methods and materials for containment and cleaning up	Large Spills: Stop the flow of material, if this is without risk. Dike the spilled material, where this is possible. Cover with plastic sheet to prevent spreading. Absorb in vermiculite, dry sand or earth and place into containers. Use water spray to reduce vapors or divert vapor cloud drift. Prevent entry into waterways, sewer, basements or confined areas. Following product recovery, flush area with water.
		Small Spills: Wipe up with absorbent material (e.g. cloth, fleece). Clean surface thoroughly to remove residual contamination.
	Environmental precautions	Never return spills to original containers for re-use. For waste disposal, see section 13 of the SDS. Avoid discharge into drains, water courses or onto the ground.
	7. Handling and storage	Avoid protoprod protoproto

Precautions for safe handling Avoid prolonged exposure. Observe good industrial hygiene practices, Store in original tightly closed container. Store away from incompatible materials (see Section 10 Conditions for safe storage, including any incompatibilities of the SDS).

8. Exposure controls/personal protection

cupational exposure limits US. OSHA Table Z-1 Limits for	Air Contaminants (29 CFR 1910.	4000	
Components	Type	Value	
Carbon Black (CAS 1333-86-4)	PEL	3.5 mg/m3	
US. ACGIH Threshold Limit Val Components	ues Type	Value	Form
Carbon Black (CAS 1333-86-4)	TWA	3 mg/m3	Inhalable fraction,

US. NIOSH: Pocket Guide to Chemical Hazards

Components	Туре	Value
Carbon Black (CAS 1333-86-4)	TWA	0.1 mg/m3
Biological limit values	No biological exposure limits noted for	or the ingredient(s).
Appropriate engineering controls	Good general ventilation should be u applicable, use process enclosures.	sed. Ventilation rates should be matched to conditions. If ocal exhaust ventilation, or other engineering controls to mended exposure limits, if exposure limits have not been
Individual protection measures,	, such as personal protective equipm	
Eye/face protection	If contact is likely, safety glasses with	side shields are recommended.
Skin protection		
Hand protection	For prolonged or repeated skin conta	ct use sultable protective gloves.
Other	Wear suitable protective clothing.	· · · ·
Respiratory protection	When workers are facing concentration certified respirators. Use a NIOSH/MS vapor/mist at levels exceeding the exp	ons above the exposure limit they must use appropriate SHA approved respirator if there is a risk of exposure to posure limits.
Thermal hazards	Wear appropriate thermal protective of	lothing, when necessary.
General hygiene considerations	Always observe good personal hygier	ne measures, such as washing after handling the material noking. Routinely wash work clothing and protective

9. Physical and chemical properties

Appearance

· · · · · · · · · · · · · · · · · · ·	
Physical state	Liquid,
Form	Liquid. Viscous Liquid
Color	Black,
Odor	slight to none
Odor threshold	Not available.
pH	Not available.
Melting point/freezing point	Not available.
Initial boiling point and boiling range	7592 °F (4200 °C) estimated
Flash point	333.7 °F (167.6 °C) estimated
Evaporation rate	Not available.
Flammability (solid, gas)	Not available.
Upper/lower flammability or exp	losive limits
Flammability limit - lower (%)	Not available.
Flammability limit - upper (%)	Not available.
Explosive limit - lower (%)	Not available.
Explosive limit - upper (%)	Not available.
Vapor pressure	Not available.
Vapor density	Not available.
Relative density	Not available.
Solubility(ies)	
Solubility (water)	Not available.
Partition coefficient (n-octanol/water)	Not available.
Auto-ignition temperature	Not available.
Decomposition temperature	Not available.
Viscosity	Not available.

 Other information

 Density
 9,821 lb/gai

 Flammability class
 Combustible [IIB estimated]

10. Stability and reactivity

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Reactivity	The product is stable and non-reactive under normal conditions of use, storage and transport.
Chemical stability	Material is stable under normal conditions,
Possibility of hazardous reactions	No dangerous reaction known under conditions of normal use.
Conditions to avoid	Avoid temperatures exceeding the flash point. Contact with incompatible materials.
Incompatible materials	Strong oxidizing agents.
Hazardous decomposition products	No hazardous decomposition products are known.

11. Toxicological information

Information on likely routes of exposure

Inhalation	Prolonged inhalation may be harmful.
Skin contact	No adverse effects due to skin contact are expected.
Eye contact	Direct contact with eyes may cause temporary irritation.
Ingestion	Expected to be a low ingestion hazard.
Symptoms related to the physical, chemical and toxicological characteristics	Direct contact with eyes may cause temporary irritation.

Information on toxicological effects

Acute toxicity

Product	Species	Test Results
DL-02596 Black LV		
<u>Acute</u>		
Dermal		
LD50	Rabbit	1373.1575 ml/kg estimated
Oral		• · · · · · · · ·
LD50	Rat	22929.7773 mg/kg estimated
Components	Species	Test Results
Carbon Black (CAS 1333-86-4)		
Acute		
Oral		
LD50	Rat	> 8000 mg/kg
* Estimates for product may	be based on additional component	data not shown.
Skin corrosion/irritation	Prolonged skin contact may cause	
Serious eye damage/eye Irritation	Direct contact with eyes may cau	
Respiratory or skin sensitizati	on	
Respiratory sensitization	Not available.	
Skin sensitization	This product is not expected to c	ause skin sensitization.
Germ cell mutagenicity		duct or any components present at greater than 0.1% are
Carcinogenicity	This product is not considered to	be a carcinogen by IARC, ACGIH, NTP, or OSHA.
IARC Monographs. Overal	Evaluation of Carcinogenicity	- , ,,
Carbon Black (CAS 133	· · ·	B Possibly carcinogenic to humans. 1001-1050)
US. OSHA Specifically Reg Not listed.		

Specific target organ toxicity - single exposure	Not classified.
Single exposure Specific target organ toxicity - repeated exposure	Not classified,
Aspiration hazard	Not available.
Chronic effects	Prolonged inhalation may be harmful.
12. Ecological information	I
Ecotoxicity	The product is not classified as environmentally hazardous. However, this does not exclude the

	possibility that large or frequent spills can have a harmful or damaging effect on the environment,
Persistence and degradability	No data is available on the degradability of this product.
Bioaccumulative potential	No data available.
Mobility in soil	No data available.
Other adverse effects	No other adverse environmental effects (e.g. ozone depletion, photochemical ozone creation potential, endocrine disruption, global warming potential) are expected from this component.

13. Disposal considerations

Disposal instructions	Collect and reclaim or dispose in sealed containers at licensed waste disposal site.
Local disposal regulations \rightarrow	Dispose in accordance with all applicable regulations.
Hazardous waste code	The waste code should be assigned in discussion between the user, the producer and the waste disposal company.
Waste from residues / unused products	Dispose of in accordance with local regulations. Empty containers or liners may retain some product residues. This material and its container must be disposed of in a safe manner (see: Disposal instructions).
Contaminated packaging	Empty containers should be taken to an approved waste handling site for recycling or disposal. Since emptied containers may retain product residue, follow label warnings even after container is emptied.

14. Transport information

DOT

Not regulated as dangerous goods.

ΙΑΤΑ

Not regulated as dangerous goods.

IMDG

Not regulated as dangerous goods.

Transport in bulk according to Not available. Annex II of MARPOL 73/78 and the IBC Code

15. Regulatory information

All components are on the U.S. EPA TSCA Inventory List,

TSCA Section 12(b) Export Notification (40 CFR 707, Subpt. D)

Not regulated.

CERCLA Hazardous Substance List (40 CFR 302.4)

Not listed.

US federal regulations

SARA 304 Emergency release notification

Not regulated,

US. OSHA Specifically Regulated Substances (29 CFR 1910.1001-1050) Not listed.

Not listed.

Superfund Amendments and Reauthorization Act of 1986 (SARA) Hazard categories Immediate Hazard - No

Immediate Hazard - No Delayed Hazard - No Fire Hazard - No Pressure Hazard - No Reactivity Hazard - No

SARA 302 Extremely hazardous substance

Not listed.

1			
SARA 311/312 Hazardous chemical	No		
SARA 313 (TRI reporting) Not regulated.			
Other federal regulations			
Clean Air Act (CAA) Sectio	n 112 Hazardous Air F	Pollutants (HAPs) List	
Not regulated.			
Clean Air Act (CAA) Sectio Not regulated.	n 112(r) Accidental Re	elease Prevention (40 CFR 68.130)	
Safe Drinking Water Act (SDWA)	Not regulated.		
US state regulations			
US - New Jersey RTK - Sub	stances: Listed subst	tance	
		tment of Justice (California Health and S	afety Code Section 11100)
Not listed. US. California. Candidate C (a))	hemicals List. Safer C	Consumer Products Regulations (Cal. Co	de Regs, tit. 22, 69502.3, subd.
Carbon Black (CAS 1333 US. Massachusetts RTK - S			
Carbon Black (CAS 1333 US. New Jersey Worker and		-Know Act	
Not regulated.			
US. Pennsylvania RTK - Ha			
Carbon Black (CAS 1333 US. Pennsylvania Worker a		to-Know Law	
Carbon Black (CAS 1333 US. Rhode Island RTK			
Not regulated.			,
US. California Proposition 6 WARNING: This product		own to the State of California to cause canc	er.
		late/Carcinogenic substance	
Carbon Black (CAS		Listed: February 21, 2003	
International Inventories	·		
Country(s) or region	Inventory name		On inventory (yes/no)*
Australia	-	of Chemical Substances (AICS)	Yes
Canada	Domestic Substances		Yes
Canada	Non-Domestic Substa	ances List (NDSL)	No
China	Inventory of Existing (Chemical Substances in China (IECSC)	Yes
Europe		of Existing Commercial Chemical	No
Europe	European List of Notif	fied Chemical Substances (ELINCS)	No
Japan		and New Chemical Substances (ENCS)	Yes
Korea	Existing Chemicals Li	- ,	Yes
New Zealand	New Zealand Inventor	ſŷ	Yes
Philippines	Philippine Inventory of (PICCS)	f Chemicals and Chemical Substances	Yes
United States & Puerto Rico		ntrol Act (TSCA) Inventory	Yes
Taiwan		als Substances Control Act	No
*A "Yes" indicates that all compon A "No" indicates that any or more	ents of this product compl	ly wilh the inventory requirements administered b of are not listed or exempt from listing on the inve	w the governing country(r)

16. Other information, including date of preparation or last revision

Issue date

ji di u

04-23-2015

Revision date Version # Disclaimer

08-21-2015

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The information contained herein is based on data believed to be reliable and the manufacturer disclaims any liability incurred from the use or reliance upon the same. The information given is designed only as a guidance for safe handling, use, processing, storage, transportation, disposal and release and is not to be considered a warranty or quality specification. The information relates only to the specific material designated and may not be valid for such material used in combination with any other materials or in any process, unless specified in the text. This safety information is not a license to use this material as claimed by any patents of third parties. The user alone must finally determine whether a contemplated use of this material will infringe any such patents, and for obtaining any required licenses. The information in the sheet was written based on the best knowledge and experience currently available.

Revision Information

Physical & Chemical Properties: Multiple Properties

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Version	Revision Date:	SDS Number:	Date of last issue: 06/27/2017	
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SECTION 1. IDENTIFICATION

Product name

: JEFFCAT® ZF-20

Manufacturer or supplier's details

Company name of supplier Address Telephone	:	Huntsman International LLC P.O. Box 4980 The Woodlands, TX 77387 United States of America (USA) TechInfo: (281) 719-7780
E-mail address of person responsible for the SDS	:	SDS@huntsman.com
Emergency telephone number	:	Chemtrec: (800) 424-9300 or (703) 527-3887
Recommended use of the che Recommended use	m i :	ical and restrictions on use Component of a Polyurethane System.
Restrictions on use	:	For industrial use only.

SECTION 2. HAZARDS IDENTIFICATION

GHS classification in accord Flammable liquids	lance with 29 CFR 1910.1200 : Category 4
Acute toxicity (Oral)	: Category 4
Acute toxicity (Inhalation)	: Category 4
Acute toxicity (Dermal)	: Category 3
Skin corrosion	: Category 1B
Serious eye damage	: Category 1
Short-term (acute) aquatic hazard	: Category 3
GHS label elements Hazard pictograms	

Signal word

: Danger

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Haza	rd statements	H311 Toxic in c	armful if swallowed or if inhaled. ontact with skin. evere skin burns and eye damage.
Preca	autionary statements	No smoking. P261 Avoid bre P264 Wash skin P270 Do not ea P271 Use only P273 Avoid rele P280 Wear prot face protection. Response: P301 + P312 IF if you feel unwe P301 + P330 + induce vomiting P303 + P361 + all contaminate P304 + P340 + and keep comfc CENTER/docto P305 + P351 + water for severa and easy to do. CENTER/docto P362 Take off c P370 + P378 In alcohol-resistan Storage: P403 + P235 Si P405 Store lock Disposal: P501 Dispose c	 SWALLOWED: Call a POISON CENTER/doctor II. P331 IF SWALLOWED: Rinse mouth. Do NOT P353 IF ON SKIN (or hair): Take off immediately d clothing. Rinse skin with water/shower. P310 IF INHALED: Remove person to fresh air ortable for breathing. Immediately call a POISON r. P338 + P310 IF IN EYES: Rinse cautiously with al minutes. Remove contact lenses, if present Continue rinsing. Immediately call a POISON r. contaminated clothing and wash before reuse. case of fire: Use dry sand, dry chemical or it foam to extinguish.
Othe	r hazards		

None known.

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

Substance / Mixture	:	Substance

Chemical nature : Amines

Hazardous components

Chemical name	CAS-No,	Concentration (% w/w)
N,N,N',N'-tetramethyl-2,2'-	3033-62-3	90 - 100
oxybis(ethylamine)		

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The specific chemical identity and/or exact percentage (concentration) of composition may be withheld as a trade secret.

SECTION 4. FIRST AID MEASURES

General advice	:	Move out of dangerous area. Consult a physician. Show this safety data sheet to the doctor in attendance. Symptoms of poisoning may appear several hours later. Treat symptomatically. Get medical attention if symptoms occur.
If inhaled	:	If inhaled, remove to fresh air. Get medical attention if symptoms occur.
In case of skin contact	:	Immediate medical treatment is necessary as untreated wounds from corrosion of the skin heal slowly and with difficulty. Take victim immediately to hospital. If on skin, rinse well with water. If on clothes, remove clothes.
In case of eye contact	:	Small amounts splashed into eyes can cause irreversible tissue damage and blindness. In the case of contact with eyes, rinse immediately with plenty of water and seek medical advice. Continue rinsing eyes during transport to hospital. Remove contact lenses. Keep eye wide open while rinsing. If eye irritation persists, consult a specialist.
If swallowed	:	Clean mouth with water and drink afterwards plenty of water. Keep respiratory tract clear. Do NOT induce vomiting. Never give anything by mouth to an unconscious person. If symptoms persist, call a physician. Take victim immediately to hospital.
Most important symptoms and effects, both acute and delayed	:	Blurred vision
Notes to physician	:	Treat symptomatically.

SECTION 5. FIREFIGHTING MEASURES

Suitable extinguishing media	:	Use extinguishing measures that are appropriate to local circumstances and the surrounding environment.
		Carbon dioxide (CO2)

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Date of last issue: 06/27/2017 Revision Date: SDS Number: Version Date of first issue: 03/14/2016 02/26/2019 400001000079 1.3 : High volume water jet Unsuitable extinguishing media Do not allow run-off from fire fighting to enter drains or water Specific hazards during • courses. firefighting No hazardous combustion products are known Hazardous combustion : products : No data is available on the product itself. Specific extinguishing methods : Collect contaminated fire extinguishing water separately. This Further information must not be discharged into drains. Fire residues and contaminated fire extinguishing water must be disposed of in accordance with local regulations. For safety reasons in case of fire, cans should be stored separately in closed containments. Use a water spray to cool fully closed containers. : In the event of fire, wear self-contained breathing apparatus. Special protective equipment for firefighters

SECTION 6. ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures	Use personal protective equipment. Refer to protective measures listed in sections 7 and 8.	
Environmental precautions	Prevent product from entering drains. Prevent further leakage or spillage if safe to do so. If the product contaminates rivers and lakes or drains info respective authorities.	۰rm
Methods and materials for containment and cleaning up	Contain spillage, and then collect with non-combustible absorbent material, (e.g. sand, earth, diatomaceous earth vermiculite) and place in container for disposal according local / national regulations (see section 13). Keep in suitable, closed containers for disposal.	ı, to

SECTION 7. HANDLING AND STORAGE

Advice on protection against fire and explosion	:	Do not spray on a naked flame or any incandescent material. Keep away from open flames, hot surfaces and sources of ignition.
Advice on safe handling	:	Avoid formation of aerosol. Do not breathe vapours/dust. Avoid contact with skin and eyes. For personal protection see section 8. Smoking, eating and drinking should be prohibited in the application area. Provide sufficient air exchange and/or exhaust in work rooms.

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			ls during handling keep bottle on a metal tray. nse water in accordance with local and national
Cond	litions for safe storage	Containers wh upright to prev Observe label	-ventilated place. ich are opened must be carefully resealed and kept vent leakage.
Mate	erials to avoid	: For incompation SDS.	tible materials please refer to Section 10 of this
	ner information on age stability	: Stable under	normal conditions.

SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Components	CAS-No.	Value type (Form of exposure)	Control parameters / Permissible concentration	Basis
N,N,N',N'-tetramethyl-2,2'- oxybis(ethylamine)	3033-62-3	TWA	0.05 ppm	ACGIH
		STEL	0.15 ppm	ACGIH

Components with workplace control parameters

Personal protective equipment

	O want and least ask and ventilation is recommanded	d to
Respiratory protection	General and local exhaust ventilation is recommended maintain vapor exposures below recommended limits. concentrations are above recommended limits or are unknown, appropriate respiratory protection should be Follow OSHA respirator regulations (29 CFR 1910.134 use NIOSH/MSHA approved respirators. Protection pr by air purifying respirators against exposure to any hazardous chemical is limited. Use a positive pressure supplied respirator if there is any potential for uncontro release, exposure levels are unknown, or any other circumstance where air purifying respirators may not p adequate protection.	Where worn. 4) and rovided e air blled
Hand protection		
Remarks	The suitability for a specific workplace should be discu with the producers of the protective gloves.	issed
Eye protection	Eye wash bottle with pure water Tightly fitting safety goggles Wear face-shield and protective suit for abnormal proc problems.	cessing
Skin and body protection	Impervious clothing	

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Hygie	ene measure s	concentration of : Avoid contact v When using do When using do	protection according to the amount and of the dangerous substance at the work place. with skin, eyes and clothing. o not eat or drink. o not smoke. efore breaks and immediately after handling

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance	: liquid
Colour	: light vellow
Odour	: ammoniacal
Odour Threshold	: No data is available on the product itself.
рН	: 11.6
Melting point	: < -4 °F / < -20 °C
Boiling point	: 373.8 °F / 189.9 °C (1,013 hPa)
Flash point	: 154 °F / 68 °C Method: closed cup
Evaporation rate	: No data is available on the product itself.
Flammability (solid, gas)	: No data is available on the product itself.
Flammability (liquids)	: No data is available on the product itself.
Upper explosion limit / Upper flammability limit	: No data is available on the product itself.
Lower explosion limit / Lower flammability limit	: No data is available on the product itself.
Vapour pressure	: 0.49 hPa (68 °F / 20 °C)
Relative vapour density	: 1
Relative density	: 0.85 (68 °F / 20 °C)
Density	: 0.85 g/cm3 (68 °F / 20 °C)
Solubility(ies) Water solubility	: 1,000 g/l completely miscible (68 °F / 20 °C)
Solubility in other solvents	: Solvent: Methanol Description: soluble

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octar Auto-	ion coefficient: n- iol/water ignition temperature mal decomposition	: 316 °F / 158	34 (68 °F / 20 °C) °C vailable on the product itself.	
	Accelerating mposition temperature IT)	: No data is a	vailable on the product itself.	
Visco Vi	osity scosity, dynamic	: <20 mPa.s	(68 °F / 20 °C)	
Explo	osive properties	: No data is a	vailable on the product itself.	
Oxidi	zing properties	: No data is a	vailable on the product itself.	
Mole	cular weight	: 160.3 g/mol		
Meta	l corrosion rate	: Not corrosiv	e to metals	
Partie	cle size	: No data is a	vailable on the product itself.	

SECTION 10. STABILITY AND REACTIVITY

Reactivity Chemical stability Possibility of hazardous	 No dangerous reaction known under conditions of normal use. Stable under normal conditions. Vapours may form explosive mixture with air.
reactions Conditions to avoid	: Heat, flames and sparks.
Incompatible materials	: Strong acids and oxidizing agents
Hazardous decomposition products	: No hazardous decomposition products are known.

SECTION 11. TOXICOLOGICAL INFORMATION

Information on likely routes of exposure	: No data is available on the product itself.
Acute toxicity Acute oral toxicity - Product	: Acute toxicity estimate : 681.09 mg/kg Method: Calculation method
Acute inhalation toxicity - Product	: Acute toxicity estimate: 4.02 mg/l Exposure time: 4 h Test atmosphere: dust/mist Method: Calculation method

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Acute dermal toxicity -Product : Acute toxicity estimate : 442.66 mg/kg Method: Calculation method

Acute toxicity (other routes of : No data available administration)

Skin corrosion/irritation

Components:

N,N,N',N'-tetramethyl-2,2'-oxybis(ethylamine): Species: Rabbit Method: OECD Test Guideline 404 Result: Causes burns

Serious eye damage/eye irritation

Components:

N,N,N',N'-tetramethyl-2,2'-oxybis(ethylamine): Species: Rabbit Result: Risk of serious damage to eyes. Method: OECD Test Guideline 405

Respiratory or skin sensitisation

Components:

N,N,N',N'-tetramethyl-2,2'-oxybis(ethylamine): Exposure routes: Skin Species: Guinea pig Method: OECD Test Guideline 406 Result: Does not cause skin sensitisation.

Assessment:

No data available

Germ cell mutagenicity

Components:

N,N,N',N'-tetramethyl-2,2'-oxybis(ethylamine): Genotoxicity in vitro : Concentration: .08 - .18 mg/ml Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 476 Result: negative

> Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 471 Result: negative

> Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 479 Result: Not classified due to inconclusive data.

Metabolic activation: negative Method: OECD Test Guideline 482 Result: negative

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Components:

N,N,N',N'-tetramethyl-2,2'-oxyl Genotoxicity in vivo	 bis(ethylamine): Application Route: Intraperitoneal injection Dose: 45 - 145 mg/kg Method: OECD Test Guideline 474 Result: negative
Carcinogenicity No data available	
Carcinogenicity - Assessment	: No data available
IARC	No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC.
ACGIH	No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen or potential carcinogen by ACGIH.
OSHA	No component of this product present at levels greater than or equal to 0.1% is on OSHA's list of regulated carcinogens.
ΝΤΡ	No component of this product present at levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen by NTP.
Reproductive toxicity Effects on fertility	: No data available
Components: N,N,N',N'-tetramethyl-2,2'-oxyl Effects on foetal development	bis(ethylamine): Species: Rabbit Application Route: Dermal General Toxicity Maternal: No observed adverse effect level: 2.5 mg/kg body weight Embryo-foetal toxicity: No observed adverse effect level: 12 mg/kg body weight Method: OECD Test Guideline 414 Result: Embryotoxic effects and adverse effects on the offspring were detected only at high maternally toxic doses
Reproductive toxicity - Assessment	: No data available
STOT - single exposure No data available	

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STOT - repeated exposure

No data available

Repeated dose toxicity

Components:

N,N,N',N'-tetramethyl-2,2'-oxybis(ethylamine): Species: Rat, male and female : 8.2 mg/m3 Application Route: Ingestion Test atmosphere: vapour Exposure time: 336 h Number of exposures: 6 h Method: Subacute toxicity

Repeated dose toxicity - : No data available Assessment

Aspiration toxicity No data available

Experience with human exposure

General Information: No	data available
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Inhalation: No data available

Skin contact: No data available

Components:

N,N,N',N'-tetramethyl-2,2'-oxybis(ethylamine): Eye contact : Symptoms: Blurred vision

Ingestion: No data available

Toxicology, Metabolism, Distribution No data available

Neurological effects No data available

Further information Ingestion:

No data available

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SECTION 12. ECOLOGICAL INFORMATION

Ecotoxicity

Components:

N,N,N',N'-tetramethyl-2,2'-oxybis(ethylamine): Toxicity to fish : LC50 (Brachydanio rerio (zebrafish)): ca. 131.2 mg/l Exposure time: 96 h Test Type: semi-static test Test substance: Fresh water Method: OECD Test Guideline 203

Components:

bis(ethylamine):
: EC50 (Daphnia magna (Water flea)): 102 mg/l
Exposure time: 48 h
Test Type: static test
Test substance: Fresh water
Method: OECD Test Guideline 202

Components:

N,N,N',N'-tetramethyl-2,2'-oxybis(ethylamine):

N,N,N',N'-tetramethyl-2,2'-oxyl Toxicity to algae/aquatic plants	 ErC50 (Selenastrum capricornutum (green algae)): 24 mg/l Exposure time: 72 h Test Type: static test Test substance: Fresh water Method: OECD Test Guideline 201
M-Factor (Acute aquatic toxicity)	: No data available
Toxicity to fish (Chronic toxicity)	: No data available
Toxicity to daphnia and other aquatic invertebrates (Chronic toxicity)	: No data available
M-Factor (Chronic aquatic toxicity)	: No data available
Toxicity to microorganisms	: No data available
Toxicity to soil dwelling organisms	: No data available
Plant toxicity	: No data available
Sediment toxicity	: No data available
Toxicity to terrestrial organisms	: No data available
Ecotoxicology Assessment	

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Versio	on Revision Date: 02/26/2019		DS Number: 0001000079	Date of last issue: 06/27/2017 Date of first issue: 03/14/2016
A	Acute aquatic toxicity	:	No data availab	le
N	Components: N,N,N',N'-tetramethyl-2,2'-			
			·	s no known ecotoxicological effects.
	oxicity Data on Soil		No data availab	
	Other organisms relevant the environment	to :	No data availab	le
P	ersistence and degrada	bility		
N	Components: I,N,N',N'-tetramethyl-2,2'- Biodegradability		Inoculum: activa Result: Inherent Biodegradation: Exposure time: Method: OECD Inoculum: activa Result: Not reac Biodegradation: Exposure time:	ly biodegradable. < 10 % 28 d Test Guideline 302B lily biodegradable. 2 % 28 d
	liochemical Oxygen Demand (BOD)	:	Method: OECD No data availabl	Test Guideline 301F le
	hemical Oxygen Demano COD)	i :	No data availabl	e
В	OD/COD	:	No data availabl	e
т	hOD	:	No data availabl	e
В	OD/ThOD	:	No data availabl	е
	issolved organic carbon DOC)	:	No data availabl	e
	hysico-chemical emovability	:	No data availabl	e
S	tability in water	:	No data availabl	e
P	hotodegradation	:	No data availabl	e
	npact on Sewage reatment	:	No data availabl	e

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ersion 3	Revision Date: 02/26/2019	SDS Number: 400001000079	Date of last issue: 06/27/2017 Date of first issue: 03/14/2016	
Bioac	cumulative potential			
Comp	onents:			
N,N,N	,N'-tetramethyl-2,2'-oxy			
Bioaco	cumulation	: Remarks: Bio	accumulation is unlikely.	
	on coefficient: n- l/water - Product	: log Pow: -0.34	4 (68 °F / 20 °C)	
Mobili	ty in soil			
Mobilit	у	: No data availa	able	
	ution among nmental compartments	: No data availa	able	
Stabili	ty in soil	: No data availa	ble	
Other	adverse effects			
	nmental fate and	: No data availa	ble	
Result assess	s of PBT and vPvB sment	: No data availa	ble	
Endoc potent	rine disrupting al	: No data availa	ble	
	oed organic bound ns (AOX)	: No data availa	ble	
Hazar	dous to the ozone laye)r		
Ozone	-Depletion Potential	Protection of S Substances Remarks: This manufactured) CFR Protection of Environment; Part 82 Stratospheric Ozone - CAA Section 602 Cla s product neither contains, nor was with a Class I or Class II ODS as defined b Act Section 602 (40 CFR 82, Subpt. A, Ap	oy tl
	nal ecological ation - Product		ntal hazard cannot be excluded in the ever I handling or disposal. Jatic life.	nt of
Global (GWP)	warming potential	: No data availa	ble	

SECTION 13. DISPOSAL CONSIDERATIONS

Disposal methods	
Waste from residues	 The product should not be allowed to enter drains, water courses or the soil. Do not contaminate ponds, waterways or ditches with

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		Dispose of as h national regula	sed waste management company. nazardous waste in compliance with local and	
Contaminated packaging		 Empty remaining contents. Dispose of as unused product. Do not re-use empty containers. Do not burn, or use a cutting torch on, the empty drum. 		

SECTION 14. TRANSPORT INFORMATION

ΙΑΤΑ		
UN/ID No.	: UN 2922	
Proper shipping name	: Corrosive liquid, toxic, n.o.s. (BIS(DIMETHYLAMINOETHYL)ETHER)	
Class	: 8	
Subsidiary risk	: 6.1	
Packing group	: 11	
Labels	: Corrosive, Toxic	
Packing instruction (cargo aircraft)	: 855	
Packing instruction (passenger aircraft)	: 851	
IMDG		
UN number	: UN 2922	
Proper shipping name	: CORROSIVE LIQUID, TOXIC, N.O.S. (BIS(DIMETHYLAMINOETHYL)ETHER)	
Class	; 8	
Subsidiary risk	: 6.1	
Packing group	: 11	
Labels	: 8 (6.1)	
EmS Code	: F-A, S-B	
Marine pollutant	: no	
Transport in bulk according	g to Annex II of MARPOL 73/78 and the IBC Code	

Not applicable for product as supplied.

National Regulations

DOT Classification UN/ID/NA number	-	UN 2922
Proper shipping name	:	CORROSIVE LIQUIDS, TOXIC, N.O.S. (BIS(DIMETHYLAMINOETHYL)ETHER)
Class	:	8
Subsidiary risk	:	6.1

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Packing group	: 11
Labels	: CORROSIVE, POISON
ERG Code	: 154
Marine pollutant	: no

Special precautions for user

The transport classification(s) provided herein are for informational purposes only, and solely based upon the properties of the unpackaged material as it is described within this Safety Data Sheet. Transportation classifications may vary by mode of transportation, package sizes, and variations in regional or country regulations.

SECTION 15. REGULATORY INFORMATION

EPCRA - Emergency Planning and Community Right-to-Know Act

CERCLA Reportable Quantity

This material does not contain any components with a CERCLA RQ.

SARA 311/312 Hazards	;	Flammable (gases, aerosols, liquids, or solids) Acute toxicity (any route of exposure) Skin corrosion or irritation Serious eye damage or eye irritation
SARA 313	:	This material does not contain any chemical components with known CAS numbers that exceed the threshold (De Minimis) reporting levels established by SARA Title III, Section 313.

This product does not contain any hazardous air pollutants (HAP), as defined by the U.S. Clean Air Act Section 112 (40 CFR 61).

California Prop. 65

This product does not contain any chemicals known to State of California to cause cancer, birth defects, or any other reproductive harm.

The components of this product are reported in the following inventories:CH INV: On the inventory, or in compliance with the inventory			
DSL	: All components of this product are on the Canadian DSL		
AICS	: On the inventory, or in compliance with the inventory		
NZIoC	: On the inventory, or in compliance with the inventory		
ENCS	: On the inventory, or in compliance with the inventory		
KECI	: On the inventory, or in compliance with the inventory		
PICCS	: On the inventory, or in compliance with the inventory		
IECSC	: On the inventory, or in compliance with the inventory		
TCSI	: On the inventory, or in compliance with the inventory		
TSCA	: On the inventory, or in compliance with the inventory		

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Inventories

AICS (Australia), DSL (Canada), IECSC (China), REACH (European Union), ENCS (Japan), ISHL (Japan), KECI (Korea), NZIoC (New Zealand), PICCS (Philippines), TCSI (Taiwan), TSCA (USA)

TSCA - 5(a) Significant New Use Rule List of Chemicals No substances are subject to a Significant New Use Rule.

US. Toxic Substances Control Act (TSCA) Section 12(b) Export Notification (40 CFR 707, Subpt D)

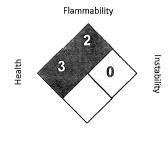
No substances are subject to TSCA 12(b) export notification requirements.

SECTION 16. OTHER INFORMATION

Further information

NFPA 704:

Dovision Data



Special hazard.

HMIS® IV:

HEALTH	3
FLAMMABILITY	2
PHYSICAL HAZARD	0

HMIS® ratings are based on a 0-4 rating scale, with 0 representing minimal hazards or risks, and 4 representing significant hazards or risks. The "*" represents a chronic hazard, while the "/" represents the absence of a chronic hazard

Revision Date	. 02/20/2019
ACGIH ACGIH / TWA ACGIH / STEL	 USA. ACGIH Threshold Limit Values (TLV) 8-hour, time-weighted average Short-term exposure limit

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The information and recommendations in this publication are to the best of our knowledge, information and belief accurate at the date of publication, NOTHING HEREIN IS TO BE CONSTRUED AS A WARRANTY, EXPRESS OR OTHERWISE.

IN ALL CASES, IT IS THE RESPONSIBILITY OF THE USER TO DETERMINE THE APPLICABILITY OF SUCH INFORMATION AND RECOMMENDATIONS AND THE SUITABILITY OF ANY PRODUCT FOR ITS OWN PARTICULAR PURPOSE.

THE PRODUCT MAY PRESENT HAZARDS AND SHOULD BE USED WITH CAUTION. WHILE CERTAIN HAZARDS ARE DESCRIBED IN THIS PUBLICATION, NO GUARANTEE IS MADE THAT THESE ARE THE ONLY HAZARDS THAT EXIST.

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Hazards, toxicity and behaviour of the products may differ when used with other materials and are dependent upon the manufacturing circumstances or other processes. Such hazards, toxicity and behaviour should be determined by the user and made known to handlers, processors and end users.

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NO PERSON OR ORGANIZATION EXCEPT A DULY AUTHORIZED HUNTSMAN EMPLOYEE IS AUTHORIZED TO PROVIDE OR MAKE AVAILABLE DATA SHEETS FOR HUNTSMAN PRODUCTS. DATA SHEETS FROM UNAUTHORIZED SOURCES MAY CONTAIN INFORMATION THAT IS NO LONGER CURRENT OR ACCURATE.

SAFETY DATA SHEET (SDS-US) DABCO 33 LX VA-No. 07139598 Version 2.0 / US EVOI Revision date 06/07/2018 POWER TO CREATE Print Date 01/09/2019 Page 1/11

JIK

1. Identification 1.1. Product ident	n of the substance/mixture and of the company/undertaking						
Trade name							
Hade Hame	. DADGO 33 EX						
.2. Recommende	d use of the chemical and restrictions on use						
Recommende	d use : Polyurethane Foam Catalyst						
Non-recomme use(s)	nded : None known.						
.3. Details of the	supplier of the safety data sheet						
Company	: Evonik Corporation Nutrition & Care PO Box 34628 Richmond VA 23234 USA						
Telephone	: +1 (0)804-727-0700						
Telefax	: +1 (0)804-727-0845						
E-mail	: us-productsafety-cs@evonik.com						
Contact Canada							
Company	: Evonik Canada Inc. 3380 South Service Burlington ON L7N 3J5 Canada						
Telephone	: +1 (0)905-336-3423-						
Telefax	: +1 (0)905-332-5632						
E-mail	: us-productsafety-cs@evonik.com						
4. Emergency tel	ephone number						
Emergency information	: Non-Emergency Phone Number : (800) 732-5616						
CHEMTREC - CHEMTREC - I	RGENCY TELEPHONE NUMBERS: JS & CANADA toll free: +1-800-424-9300 MEXICO toll free: 01-800-681-9531 LOBAL - Collect calls accepted: +1-703-527-3887						
Hazards iden	tification						
1. Classification	of the substance or mixture						

GHS classification in accordance with 29 C	FR 1910.1200	
Skin irritation	Category 2	H315
Eye irritation	Category 2A	H319

2.2. Label elements

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GHS cl	assification in a	ccordance with 29 CFR 1910	0.1200	
Symbo	l(s)			
Signal	word	: Warning		
hazard	statement	: H315 - Causes skin irritatio H319 - Causes serious eye		
Precau Statem (Preve		: P264 - Wash skin thorough P280 - Wear protective glo		protection.
Precau Statem (Respo		: P302 + P352 - IF ON SKIN P305 + P351 + P338 - IF IN Remove contact lenses, if p P332 + P313 - If skin irritati P337 + P313 - If eye irritati P362 + P364 - Take off cor	VEYES: Rinse cautiously present and easy to do. C on occurs: Get medical a on persists: Get medical a	v with water for several minutes. Continue rinsing. advice/ attention. advice/ attention.

2.3. Other hazards

None known.

3. Composition/information on ingredients

3.1. Substances

-3.2. Mixtures

Hazardous components

Chemical Name	NJ Trade secrets CAS-No.	Concentration	Classification
1,4-Diazabicyclooctane	- 280-57-9	Ø 33%	H228, 1, Flam. Sol. H302, 4, Acute Tox., Oral H315, 2, Skin Irrit. H318, 1, Eye Dam.

Texts of H phrases, see in Chapter 16

4. First aid measures

4.1. Description of first aid measures

General advice	Seek medical advice. If breathing has stopped or is labored, give assisted respirations. Supplemental oxygen may be indicated. If the heart has stopped, trained personnel should begin cardiopulmonary resuscitation immediately.
Inhalation	: Move to fresh air.
Skin contact	: Wash off immediately with soap and plenty of water.Wash off immediately with plenty of water for at least 15 minutes. Immediately remove contaminated clothing, and any extraneous chemical, if possible to do so without delay.
Eye contact	: Rinse immediately with plenty of water for at least 15 minutes.Rinse immediately with plenty of water, also under the eyelids, for at least 15 minutes. Remove contact lenses.

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	Ingestion		ent aspiration of vomit. th to an unconscious per	Turn victim's head to the si son.	de.Never give anything by
4.2.	Most important sym	ptoms and	d effects, both acute ar	nd delayed	
	Symptoms	: Up to	o now no symptoms are	known.	
4.3.	Indication of any im	mediate m	edical attention and s	pecial treatment needed	
	Treat symptomatically	<i>י</i> .			
5.	Fire-fighting meas	ures			
5.1.	Extinguishing media	l			
	Suitable extinguishing media	Dry c Dry s Lime	on dioxide (CO2) themical and stone powder nol-resistant foam		
	Unsuitable extinguishing media	: no da	ata available		
5.2.	Special hazards arisi	ing from t	he substance or mixtu	re	
	Incomplete combustic May generate ammor May generate toxic ni Burning produces nox Downwind personnel	ia gas. trogen oxic tious and t	de gases. oxic fumes.		
5.3.	Advice for firefighter	s			
	Use personal protectiv Wear self-contained b		ent. pparatus for firefighting i	f necessary.	
6.	Accidental release	measure	es		
6.1.	Personal precautions	s, protecti	ve equipment and eme	rgency procedures	
			s.Wear suitable protecti paratus and chemically p	ve clothing, gloves and eye protective clothing.	/face protection.
6.2.	Environmental preca	utions			
	Construct a dike to pre	event sprea	ading.		
6.3.	Methods and materia	l for conta	ainment and cleaning u	ıp	
	Place in appropriate cl Approach suspected le			gency Response number f	or advice.
	If possible, stop flow o	f product.			
7.	Handling and stora	ge			
7.1.	Precautions for safe	handling			
	Advice on safe handling	: Use p	ersonal protective equip	ment.	
		_			

Emergency showers and eye wash stations should be readily accessible. Adhere to work practice rules established by government regulations. Avoid contact with eyes.

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	Hygiene measures	:	Provide readily accessible e	ve wash stations and safety	showers.
	General protective measures	:	Discard contaminated leather Wash hands at the end of ea toilet.		ting, smoking or using the
7.2.	Conditions for safe s	stora	ge, including any incompat	ibilities	
	Prevention of fire an	d ex	plosion		
	Information	:	No special measures require	ed.	
	Storage				
	Information	:	Do not store near acids. Keep containers tightly close	d in a dry, cool and well-ve	ntilated place.
	Advice on common storage	:	Do not store in reactive meta	l containers.	

8. Exposure controls/personal protection

8.1. Control parameters

Exposure limit(s)

Ingredients	CAS-No.	Statutory	Value type	Value	Short-term
		basis/list	(Form of exposure;		
		(Update)	Expressed as)		

Contains no substances with occupational exposure limit values. Hazardous components without workplace control parameters

8.2. Exposure controls

Engineering controls

Appropriate	: Provide readily accessible eye wash stations and safety showers.
engineering controls	Provide natural or explosion-proof ventilation adequate to ensure concentrations are
	kept below exposure limits.

Personal protective equipment

Eye protection	: Chemical resistant goggles must be worn.
Hand protection	 Chemical-resistant, impervious gloves complying with an approved standard should be worn at all times when handling chemical products if a risk assessment indicates this is necessary. butyl-rubber Nitrile rubber Neoprene gloves Impervious gloves
Body Protection	: No specific recommendation.Long sleeve shirts and trousers without cuffs.
Respiratory protection	: Not required for properly ventilated areas.No personal respiratory protective equipment normally required.

9. Physical and chemical properties

9.1. Information on basic physical and chemical properties

Physical state	:	liquid
Form	:	liquid
Colour	:	colourless

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Odour	: ammoniacal		· · ·
Odour Threshold	: no data available		
рН	: 10.7		
Melting point	: no data available		
Boiling point	: Boiling point/range 218 °C		
Flash point	: >110 °C		
Evaporation rate	: no data available		
Flammability	: no data available		
Upper Explosion/Ignition Limit	: no data available		
Lower explosion limit	: no data available		
Vapour pressure	: 4.799 hPa (21 °C)		
Relative vapour density	: no data available		
Relative density	: 1.019 (21 °C)		
Solubility(ies)	: no data available		
Water solubility	: completely soluble		
Partition coefficient: n- octanol/water	: no data available		
Autoignition temperature	: no data available		
Thermal decomposition	: no data available		
Viscosity, kinematic	: no data available		
Viscosity, dynamic	: 305 mPa.s (21.1 °C)		
	223 mPa.s (25 °C)		
	43.8 mPa.s (54.4 °C)		

Explosive properties : no data available

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Oxidising prop	erties	: n	o data available			
9.2. Other in	formation					
Density		: n	o data available			
Metal corrosion	n	: n	o data available			
Ignition temper	rature	: n	o data available			
volatile organic compounds : This product does not contain any chemicals listed under the Section 111 SOCMI Intermediate or Final VOC's (40 CFR 60. This product does not contain any VOC exemptions listed under Section 450.			duct does not contain any chemicals listed under the U.S. Clean Air Act 111 SOCMI Intermediate or Final VOC's (40 CFR 60.489).			
		nder the U.S. Clean Air				

10. Stability and reactivity

10.1. Reactivity

see section "Possibility of hazardous reactions"

10.2. Chemical stability

Stable under normal conditions.

10.3. Possibility of hazardous reactions

no data available

10.4. Conditions to avoid

no data available

10.5. Incompatible materials

Reactive metals (e.g. sodium, calcium, zinc etc.). Materials reactive with hydroxyl compounds. Dehydrating Agents. Organic acids (i.e. acetic acid, citric acid etc.). Mineral acids. sodium hypochlorite Product slowly corrodes copper, aluminum, zinc and galvanized surfaces. Reaction with peroxides may result in violent decomposition of peroxide possibly creating an explosion. Oxidizing agents

10.6. Hazardous decomposition products

Nitric acid Ammonia Nitrogen oxides (NOx) Nitrogen oxide can react with water vapors to form corrosive nitric acid. Carbon monoxide Carbon dioxide (CO2) Aldehydes Flammable hydrocarbon fragments. Heating above 65 °C in the presence of strong base can liberate flammable hydrocarbon fragments. Carbon oxides

11. Toxicological information

11.1. Information on toxicological effects

Acute toxicity (oral) : LD50

Species: Rat Dose: 3,200 mg/kg Method: estimated

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Acute toxic (inhalation		No data is available o	n the product itself.	
Acute toxic (dermal)	ity	No data is available o	n the product itself.	
		Acute toxicity estimate Dose: > 5,000 mg/kg Method: Calculation m		
Irritation/co the skin	prrosion of	Method: see user defi Remarks: Mild skin irr The data are derived t products (conclusion b	itation rom the evaluations or test re	sults achieved with similar
Serious ey eye irritatio		Result: Irritating to eye Remarks: Moderate e Data from a comparat	ye irritation	
		of the eye from the atr Corneal edema may	nosphere.	na) when absorbed into the tissue ue haze" or "fog" around lights. effect.
Respirator sensitizatio		no data available		
Repeated of toxicity	dose	no data available		
CMR asses	sment			
Carcinoger	nicity :	no data available		
Mutagenici	ty :	No data is available or	n the product itself.	
Teratogeni	city :	No data is available or	n the product itself.	
Toxicity to reproductic		No data is available on	o data is available on the product itself.	

No component of this product present at levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen by NTP.

US. IARC Monographs on Occupational Exposures to Chemical Agents

No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC.

US. OSHA Specifically Regulated Substances (29 CFR 1910.1001-1050)

No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen or potential carcinogen by OSHA.

Specific Target Organ Toxicity - Single exposure	: no data available
Specific Target Organ Toxicity - Repeated exposure	: no data available
Aspiration hazard	: no data available

12. Ecological information

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	Ecotoxicology Asses	sme	ent				
	Acute aquatic toxicity	:	no data available				
	Chronic aquatic toxicity	:	no data available				
12.1.	Toxicity						
	Aquatoxicity, fish	:	No data is available on the p	roduct itself.			
	Aquatoxicity, invertebrates	:	No data is available on the p	roduct itself.			
	Aquatoxicity, algae / aquatic plants	:	No data is available on the p	roduct itself.			
	Toxicity in microorganisms	:	no data available				
	chronic toxicity in fish	:	no data available				
	Chronic toxicity in aquatic Invertebrates	;	no data available				
12.2.	Persistence and degradability						
	Photodegradation	:	no data available				
	Biological degradability	:	no data available				
12.3.	Bioaccumulative pote	ntia	I				
	Bioaccumulation	:	no data available				
12.4.	Mobility in soil						
	Environmental distribution	:	no data available				
12.5.	Results of PBT and v	₽vB	assessment				
	PBT and vPvB assessment	:	No data available				
12.6.	Other adverse effects						
	General Information		Do not allow to enter soil, wa The product is classified as s Regulation on the Classificat	lightly hazardous to wa	aters (according to the German		
13.	Disposal considerat	tion	S				
13.1.	Waste treatment meth						
	Product	:	Contact supplier if guidance i	s required.			
	Contaminated	:	Dispose of container and uni	used contents in accord	lance with federal, state, and loca		

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packaging

requirements.

Transport information 14.

Not dangerous according to transport regulations.

14.1. UN number: 14.2. UN proper shipping name: Transport hazard class(es): 14.3. 14.4. Packing group: 14.5. Environmental hazards: 14.6. Special precautions for user: Yes Not dangerous according to transport regulations.

15. **Regulatory information**

US Federal Regulations

SARA Title III Section 311/312 Hazard Categories - Acute Health Hazard

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 302 Extremely Hazardous Substance (40 CFR 355, Appendix A)

- This material does not contain any components with a SARA 302 RQ.

SARA 304 - Emergency Release Notification

- This material does not contain any components with a section 304 EHS RQ.

US. EPA CERCLA Hazardous Substances (40 CFR 302)

- This material does not contain any components with a CERCLA RQ.

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 313 Toxic Chemicals (40 CFR 372.65) - Supplier Notification Required

- This material does not contain any chemical components with known CAS numbers that exceed the threshold (De Minimis) reporting levels established by SARA Title III, Section 313.

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 302 Extremely Hazardous Substance (40 CFR 355, Appendix A)

- No chemicals in this material are subject to the reporting requirements of SARA Title III, Section 302.

US. Clean Air Act - Hazardous Air Pollutants (HAP)

- This product does not contain any hazardous air pollutants (HAP), as defined by the U.S. Clean Air Act Section 112 (40 CFR 61).

US. Clean Air Act Section 112(r); Regulated toxic and flammable substances for Accidental Release Prevention - 40 CFR 68.130 (subpart F)

- This product does not contain any chemicals listed under the U.S. Clean Air Act Section 112(r) for Accidental Release Prevention (40 CFR 68.130, Subpart F).

USA. Clean Water Act - Section 311: Designation of Hazardous Substances. CFR Title 40, Part 116 - This product does not contain any Hazardous Substances listed under the U.S. CleanWater Act, Section 311, Table 116.4Å.

USA. Clean Water Act - Section 311, Determination of Reportable Quantities for Hazardous Chemicals. CFR Title 40, Part 117

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- This product does not contain any Hazardous Chemicals listed under the U.S. CleanWater Act, Section 311, Table 117.3.

USA. Clean Water Act - Section 307: Toxic Pollutants

- This product does not contain any toxic pollutants listed under the U.S. Clean Water Act Section 307

State Regulations

US. California Safe Drinking Water & Toxic Enforcement Act (Proposition 65) This product does not contain any chemicals known to State of California to cause cancer, birth defects, or any other reproductive harm.

9	Health:	2
	Flammability:	1
	Reactivity:	0

Notification status

USA (TSCA)	:	listed/registered or exempted
Canada (DSL)	:	listed/registered or exempted

16. Other information

List of references

Revision date : 06/07/2018

Relevant H phrases from chapter 3

H228 :	Flammable solid.
H302 :	Harmful if swallowed.
H315 :	Causes skin irritation.
H318 :	Causes serious eye damage.

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SAFETY DATA SHEET (SDS-US)

DARCO	33	LX	
VA-No.			

07139598



Legend

ADR	European Agreement concerning the International Conviews of Departments Conde by Depart
ADR	European Agreement concerning the International Carriage of Dangerous Goods by Road European Agreement concerning the International Carriage of Dangerous Goods by Inland
	Waterways
ADNR	European agreement concerning the international carriage of dangerous goods by inland
<i>ABRIC</i>	waterways (ADN)
ASTM	American Society for Testing and Materials
ATP	Adaptation to Technical Progress
BCF	Bioconcentration factor
BetrSichV	German Ordinance on Industrial Safety and Health
C.C.	closed cup
CAS	Chemical Abstract Services
CESIO	European Committee of Organic Surfactants and their Intermediates
ChemG	German Chemicals Act
CMR	carcinogenic-mutagenic-toxic for reproduction
DIN	German Institute for Standardization
DMEL	Derived minimum effect level
DNEL	Derived no effect level
EINECS	European Inventory of Existing Commercial Chemical Substances
EC50	half maximal effective concentration
GefStoffV	German Ordinance on Hazardous Substances
GGVSEB	German ordinance for road, rail and inland waterway transportation of dangerous goods
GGVSee	German ordinance for sea transportation of dangerous goods
GLP GMO	Good Laboratory Practice
IATA	Genetic Modified Organism
ICAO	International Air Transport Association International Civil Aviation Organization
IMDG	International Maritime Dangerous Goods
ISO	International Organization For Standardization
LOAEL	Lowest observed adverse effect level
LOEL	Lowest observed effect level
NOAEL	No observed adverse effect level
NOEC	no observed effect concentration
NOEL	no observed effect level
O. C.	open cup
OECD	Organisation for Economic Cooperation and Development
OEL	Occupational Exposure Limit
PBT	Persistent, bioaccumulative, toxic
PEC	Predicted effect concentration
PNEC	Predicted no effect concentration
REACH	REACH registration
RID	Convention concerning International Carriage by Rail
STOT	Specific Target Organ Toxicity
SVHC	Substances of Very High Concern
TA TPR	Technical Instructions
TRGS	Third Party Representative (Art. 4)
VCI	Technical Rules for Hazardous Substances German chemical industry association
vPvB	very persistent, very bioaccumulative
VOC	volatile organic compounds
VwVwS	German Administrative Regulation on the Classification of Substances Hazardous to Waters
	into Water Hazard Classes
WGK	Water Hazard Class
WHO	World Health Organization
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SAFETY DATA SHEET

Y-8331

Date of issue: 2017-03-08

Revision date: Not applicable

Version: R0001.0001

1. IDENTIFICATION

A. Product name

- Y-8331 [US-P-181]

B. Recommended use and restriction on use

- General use - Restriction on use : Polyol Components for the production of polyurethane : Not available

- C. Manufacturer / Supplier / Distributor information - Company name : Mitsui Chemicals & SKC Polyurethanes Inc. - Address : 255, Yongjam-ro, Nam-gu, Ulsan, 44782, Korea - Telephone number : +82-52-278-5242 - Emergency telephone : +82-52-278-5242
- 2. HAZARD IDENTIFICATION

A. GHS Classification - Not applicable

number

B. GHS label elements

- Hazard symbols
- Not applicable
- o Signal words - Not applicable
- o Hazard statements
- Not applicable

• Precautionary statements

- 1) Prevention
- Not applicable
 - 2) Response
 - Not applicable
- 3) Storage
- Not applicable
- 4) Disposal
 - Not applicable

C. Other hazards which do not result in classification : (NFPA Classification)

• NFPA grade (0 ~ 4 level)

- Health : 1 , Flammability : 1, Reactivity : 0

Chemical Name	Trade names and Synonyms	CAS No.	Content(%)
Polyether Polyol Resin	- ·	9082-00-2	100

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4. FIRST AID MEASURES

A. Eye contact

- Do not rub your eyes.
- Immediately flush eyes with plenty of water for at least 15minutes and call a doctor/physician.

B. Skin contact

- Flush skin with plenty of wter for at least 15 minutes while removing contaminated clothing and shoes.
- Laundering enough contaminated clothing before reuse.

C. Inhalation contact

- When exposed to large amounts of steam and mist, move to fresh air.
- Take specific treatment if needed.

D. Ingestion contact

- About whether I should induce vomiting Take the advice of a doctor.
- Rinse your mouth with water immediately.

E. Delayed and immediate effects and also chronic effects from short and long term exposure

- Not available

F. Notes to physician

- Notify medical personnel of contaminated situations and have them take appropriate protective measures.

5. FIREFIGHTING MEASURES

A. Suitable (Unsuitable) extinguishing media

- Dry chemical, carbon dioxide, regular foam extinguishing agent, spray
- Avoid use of water jet for extinguishing

B. Specific hazards arising from the chemical

- Not available

C. Special protective actions for firefighters

- Cool containers with water until well after fire is out.
- Avoid inhalation of materials or combustion by-products.
- Use appropriate extinguishing measure suitable for surrounding fire.
- Wear appropriate protective equipment.
- Keep containers cool with water spray.
- Vapor or gas is burned at distant ignition sources can be spread quickly.

6. ACCIDENTAL RELEASE MEASURES

A. Personal precautions, protective equipment and emergency procedures

- Ventilate closed spaces before entering.
- Must work against the wind, let the upwind people to evacuate.
- Move container to safe area from the leak area.
- Remove all sources of ignition.
- Do not direct water at spill or source of leak.

B. Environmental precautions

- Prevent runoff and contact with waterways, drains or sewers.
- If large amounts have been spilled, inform the relevant authorities.

C. Methods and materials for containment and cleaning up

- Large spill : Stay upwind and keep out of low areas. Dike for later disposal.
- Notification to central government, local government. When emissions at least of the standard amount

- Appropriate container for disposal of spilled material collected.
- Small leak: sand or other non-combustible material, please let use absorption.
- Wipe off the solvent.
- Dike for later disposal.

7. HANDLING AND STORAGE

A. Precautions for safe handling

- Comply with all applicable laws and regulations for handling
- Get the manual before use.
- Dealing only with a well-ventilated place.
- Do not inhale the steam prolonged or repeated.

B. Conditions for safe storage, including any incompatibilities

- Do not apply direct heat.
- Save applicable laws and regulations.
- Keep in the original container.
- Keep sealed when not in use.
- Prevent static electricity and keep away from combustible materials or heat sources.

8. EXPOSURE CONTROLS/PERSONAL PROTECTION

A. Exposure limits

- ACGIH TLV
- Not available

\circ OSHA PEL

- Not available

B. Engineering controls

- A system of local and/or general exhaust is recommended to keep employee exposures above the Exposure Limits. Local exhaust ventilation is generally preferred because it can control the emissions of the contaminant at its source, preventing dispersion of it into the general work area. The use of local exhaust ventilation is recommended to control emissions near the source.

C. Individual protection measures, such as personal protective equipment

Respiratory protection

- Under conditions of frequent use or heavy exposure, Respiratory protection may be needed.
- Respiratory protection is ranked in order from minimum to maximum.
- Consider warning properties before use.
- Any chemical cartridge respirator with organic vapor cartridge(s).
- Any chemical cartridge respirator with a full facepiece and organic vaporcartridge(s).
- Any air-purifying respirator with a full facepiece and an organic vapor canister.

- For Unknown Concentration or Immediately Dangerous to Life or Health : Any supplied-air respirator with full facepiece and operated in a pressure-demand or other positive-pressure mode in combination with a separate escape supply. Any self-contained breathing apparatus with a full facepiece.

• Eye protection

- Wear primary eye protection such as splash resistant safety goggles with a secondary protection face shield.

- Provide an emergency eye wash station and quick drench shower in the immediate work area.

Hand protection

- Wear appropriate glove.

o Skin protection

- Wear appropriate clothing.

o Others

- Not available

9. PHYSICAL AND CHEMICAL PROPERTIES

A. Appearance

- Appearance	Liquid(Viscous liquid)
- Color	Colorless
B. Odor	Almost Odorless
C. Odor threshold	Not available
D. pH	5.5~7.5
E. Melting point/Freezing point	-30~-50 ℃
F. Initial Boiling Point/Boiling Ranges	Not available
G. Flash point	225 °C ~ 235 °C
H. Evaporation rate	Not available
I. Flammability(solid, gas)	Not available
J. Upper/Lower Flammability or explosive limits	Not available
K. Vapour pressure	Not available
L. Solubility	Not available
M. Vapour density	Not available
N. Specific gravity(Relative density)	Not available
O. Partition coefficient of n-octanol/water	Not available
P. Autoignition temperature	Not available
Q. Decomposition temperature	Not available
R. Viscosity	Not available
S. Molecular weight	Not available

10. STABILITY AND REACTIVITY

A. Chemical Stability

- This material is stable under recommended storage and handling conditions.

B. Possibility of hazardous reactions

- Hazardous Polymerization will not occur.

C. Conditions to avoid

- Avoid contact with incompatible materials and condition.
- Avoid : Accumulation of electrostatic charges, Heating, Flames and hot surfaces

D. Incompatible materials

- Not available

E. Hazardous decomposition products

- May emit flammable vapour if involved in fire.

11. TOXICOLOGICAL INFORMATION

A. Information on the likely routes of exposure

(Respiratory tracts)

- Not available

o (Oral)

- Not available
- (Eye·Skin)

- Not available

B. Delayed and immediate effects and also chronic effects from short and long term exposure

Acute toxicity

- * Oral
 - [Polyether Polyol Resin] : LD50 > 10000 mg/kg Rat
 - * Dermal
 - [Polyether Polyol Resin] : LD50 > 5000 mg/kg Rabbit
 - * Inhalation
 - Not available
- Skin corrosion/irritation

- Not available

 \circ Serious eye damage/irritation

Not available
 Respiratory sensitization

- Not available

Skin sensitization

- Not available

Carcinogenicity

* IARC

- Not available

* OSHA - Not available

* ACGIH

- Not available

* NTP

- Not available

* EU CLP

- Not available

 \circ Germ cell mutagenicity

- Not available

• Reproductive toxicity - Not available

• STOT-single exposure

- Not available

• STOT-repeated exposure

- Not available

Aspiration hazard

- Not available

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3	24	风气日日	OGICA	911 N H K U K 4	リング自己に行	N

A. Ecotoxicity

0 Fish

- Not available

• Crustaceans

- Not available

Algae

- Not available

B. Persistence and degradability

• Persistence

- Not available

Degradability

- Not available

C. Bioaccumulative potential

 \circ Bioaccumulative potential

- Not available

Biodegration

- Not available

D. Mobility in soil

- Not available

E. Other adverse effects

- Not available

A. Disposal methods

- Since more than two kinds of designaed waste is mixed, it is difficult to treat seperatly, then can be reduction or stabilization by incineration or similar process.

- If water separation is possible, pre-process with Water separation process.
- Dispose by incineration.

B. Special precautions for disposal

- The user of this product must disposal by oneself or entrust to waste disposer or person who other's waste recycle and dispose, person who establish and operate waste disposal facilities.

- Dispose of waste in accordance with all applicable laws and regulations.

14. TRANSPORT INFORMATION

A. UN No. (IMDG)

- Not applicable

B. Proper shipping name

- Not applicable

C. Hazard Class

- Not applicable

D. IMDG Packing group

- Not applicable

E. Marine pollutant

- Not applicable

F. Special precautions for user related to transport or transportation measures

- Local transport follows in accordance with Dangerous goods Safety Management Law.

- Package and transport follow in accordance with Department of Transportation (DOT) and other regulatory agency requirements.
- Air transport(IATA): Not subject to IATA regulations.

- EmS FIRE SCHEDULE : Not available

- EmS SPILLAGE SCHEDULE : Not available

15. REGULATORY INFORMATION

A. National and/or international regulatory information

- POPs Management Law
 - Not applicable
- Information of EU Classification
 - * Classification
 - Not applicable
 - * Risk Phrases
 - Not applicable
 - * Safety Phrase
 - Not applicable
- U.S. Federal regulations
- * OSHA PROCESS SAFETY (29CFR1910.119)
 - Not applicable
- * CERCLA Section 103 (40CFR302.4)
- Not applicable
- * EPCRA Section 302 (40CFR355.30)
 - Not applicable
- * EPCRA Section 304 (40CFR355.40)
 - Not applicable
- * EPCRA Section 313 (40CFR372.65)
 - Not applicable
- o Rotterdam Convention listed ingredients

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Not applicable
Stockholm Convention listed ingredients

Not applicable

Montreal Protocol listed ingredients

Not applicable
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16. OTHER INFORMATION

A. Reference

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The information contained herein is believed to be accurate. It is provided independently of any sale of the product for purpose of hazard communication. It is not intended to constitute performance information concerning the product. No express warranty, or implied warranty of merchantability or fitness for a particular purpose is made with respect to the product or the information contained herein.
This Safety Data Sheet was compiled with data and information from the following sources: KOSHA, NITE, ESIS, NLM, SIDS, IPCS

B. Issue date

- 2017-03-08

C. Revision number and Last date revised

- Not applicable

D. Other

- This SDS is prepared according to the Globally Harmonized System (GHS).

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MATERIAL SAFETY DATA SHEET



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1. Product and Company Identification

Material name.	.BiOH 2828
Document #	
Revision date	12-5-2013
Product use	For research and development purposes only.
Synonym(s)	Modified oligomeric vegetable oil
Manufacturer information	Cargill Biobased Polyurethanes 9320 Excelsior Blvd. Hopkins, Minnesota 55343 United States CustomerService_Polyols@cargill.com General Information: 1-877-Poly BiOH (765-9246) 24 Hour Emergency: 1-800-424-9300
2. Hazards Identification	
Physical state	Liquid.
Emergency overview	CAUTION
	Prolonged or repeated skin contact may cause drying, cracking, or irritation.
OSHA regulatory status	This product is not hazardous according to OSHA 29CFR 1910.1200.
Potential health effects Routes of exposure	Not applicable.
Eyes	Direct contact with eyes may cause temporary irritation.
Skin	Prolonged or repeated contact may cause itching, redness, and rash in some individuals.
Inhalation	Under normal conditions of intended use, this material is not expected to be an inhalation hazard.
Ingestion	No harmful effects expected in amounts likely to be ingested by accident.

3. Composition / Information on Ingredients

Components		CAS #	Percent
Modified oligomeric vegetable oil		Not Applicable	> 99
Composition comments	All concentrations are in percent by weight unle percent by volume.	ess ingredient is a gas. Ga	s concentrations are in
4. First Aid Measures			
First aid procedures			
Eye contact	Any material that contacts the eye should be w remove contact lenses. Get medical attention p		
Skin contact	Wash skin with soap and water. Get medical at	ttention promptly if sympto	ms occur after washing.
Inhalation	If symptomatic, move to fresh air. Get medical	attention if symptoms pers	ist.
Ingestion	First aid is normally not required. However, if g attention.	reater than 1/2 liter (pint) ir	ngested, seek medical
5. Fire Fighting Measures			
Flammable properties	No unusual fire or explosion hazards noted.		
Extinguishing media			
Suitable extinguishing media	Water. Water fog. Foam. Dry chemical powder.	. Carbon dioxide (CO2).	
Fire fighting equipment/instructions	Self-contained breathing apparatus and full pro	tective clothing must be w	orn in case of fire.
Hazardous combustion products	Carbon oxides.		
BiOH 2828			MSDS N/

6. Accidental Release Measures

Personal precautions Methods for cleaning up	Wear appropriate personal protective equipment (See Section 8). Absorb spill with vermiculite or other inert material, then place in a container for chemical waste.	
	Large Spills: Flush area with water. Prevent runoff from entering drains, sewers, or streams. Dike for later disposal.	
7. Handling and Storage		
Handling	Observe good industrial hygiene practices.	
Storage	Keep container closed. Store totes and drums below 38°C (100°F); shelf life: 12 months. Store bulk containers below 46°C (114°F); shelf life: 6 months.	

8. Exposure Controls / Personal Protection

Occupational exposure limits Engineering controls	No exposure limits noted for ingredient(s). Ensure adequate ventilation, especially in confined areas.
Personal protective equipment	
Eye / face protection	Wear safety glasses with side shields (or goggles).
Skin protection	Wear chemical-resistant gloves, footwear and protective clothing appropriate for risk of exposure. Contact glove manufacturer for specific information.
Respiratory protection	If engineering controls do not maintain airborne concentrations below recommended exposure limits (where applicable) or to an acceptable level (in countries where exposure limits have not been established), an approved respirator must be worn. In the United States of America, if respirators are used, a program should be instituted to assure compliance with OSHA Standard 63 FR 1152, January 8, 1998. Respirator type: Air-purifying respirator with an appropriate, government approved (where applicable), air-purifying filter, cartridge or canister.
General hygiene considerations	Always observe good personal hygiene measures, such as washing after handling the material and before eating, drinking, and/or smoking. Routinely wash work clothing and protective equipment to remove contaminants.

9. Physical & Chemical Properties

•	•
Appearance	Not available.
Color	Pink / pale yellow.
Odor	Alcohol.
Odor threshold	Not available.
Physical state	Liquid.
Form	Viscous liquid.
рН	Not available.
Melting point	Not available.
Freezing point	Not available.
Boiling point	Decomposes.
Flash point	> 266 °F (> 130 °C) Closed Cup
Evaporation rate	Not available.
Flammability limits in air, upper, % by volume	Not available.
Flammability limits in air, lower, % by volume	Not available.
Vapor pressure	Not available.
Vapor density	Not available.
Specific gravity	1.01 (25°C)
Solubility (water)	Insoluble.
Partition coefficient (n-octanol/water)	Not available.
Auto-ignition temperature	867.2 °F (464 °C)
Decomposition temperature	> 302 °F (> 150 °C)

Viscosity	3200 - 10000 cP
Density	8.4 lb/gal

10. Chemical Stability & Reactivity Information

Chemical stability	Material is stable under normal conditions.
Conditions to avoid	Excessive heat.
Incompatible materials	Strong oxidizing agents.
Hazardous decomposition products	No hazardous decomposition products are known.
Possibility of hazardous reactions	Hazardous polymerization does not occur.

11. Toxicological Information

Toxicological data Product	Test Results
Experimental Products A	Acute Oral LD50 Rat: > 2000 mg/kg
Sensitization	No sensitizing effects known.
Carcinogenicity	This product is not considered to be a carcinogen by IARC, ACGIH, NTP, or OSHA.
Mutagenicity	Ames test: Negative.
12. Ecological Informat	ion

12. Ecological Information

Ecotoxicity	Not expected to be harmful to aquatic organisms.
Persistence and degradability	No data available.
Bioaccumulation / Accumulation	No data available.
Partition coefficient (n-octanol/water)	Not available.
Mobility in environmental media	No data available.

13. Disposal Considerations

Disposal instructions	Dispose of contents/container in accordance with local/regional/national/international regulations. When this product as supplied is to be discarded as waste, it does not meet the definition of a RCRA waste under 40 CFR 261.
Contaminated packaging	Since emptied containers may retain product residue, follow label warnings even after container is emptied

14. Transport Information

DOT Not regulated as dangerous goods. ΙΑΤΑ Not regulated as dangerous goods. IMDG Not regulated as dangerous goods. TDG Not regulated as dangerous goods.

15. Regulatory Information

US federal regulations This product is not hazardous according to OSHA 29CFR 1910.1200.

CERCLA (Superfund) reportable quantity (lbs)

None

Superfund Amendments and Reauthorization Act of 1986 (SARA)

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Hazard categories	Immediate Hazard - No Delayed Hazard - No Fire Hazard - No Pressure Hazard - No Reactivity Hazard - No	
Section 302 extremely hazardous substance	No	
Section 311 hazardous chemical	No	
Drug Enforcement Agency (DEA)	Not controlled	
WHMIS status	Non-controlled	
Inventory status		
Country(s) or region	Inventory name	On inventory (yes/no)*
Canada	Domestic Substances List (DSL)	Yes
Canada	Non-Domestic Substances List (NDSL)	No
United States & Puerto Rico	Toxic Substances Control Act (TSCA) Inventory	Yes
*A "Yes" indicates that all compo	nents of this product comply with the inventory requirements administered	by the governing country(s)
State regulations	ulations This product does not contain a chemical known to the State of California to cause cancer, birth defects or other reproductive harm.	
16. Other Information		

Further information HMIS® ratings	HMIS® is a registered trade and service mark of the NPCA. Health: 1 Flammability: 1 Physical hazard: 0
NFPA ratings	Health: 1 Flammability: 1 Instability: 0
Disclaimer	To the best of our knowledge, the information contained herein is accurate. However, neither the above named supplier nor any of its subsidiaries assumes any liability whatsoever for completeness of the information contained herein. Final determination of suitability of any material is the sole responsibility of the user. All materials may present unknown hazards and should be used with caution. Although certain hazards are described herein, we cannot guarantee that these are the only hazards that exist.
Issue date	10-14-2010

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SAFETY DATA SHEET



1. Identification

Covestro LLC 1 Covestro Circle Pittsburgh, PA 15205 USA

TRANSPORTATION EMERGENCY CALL CHEMTREC:

INTERNATIONAL:

(800) 424-9300 (703) 527-3887

NON-TRANSPORTATION Emergency Phone: Information Phone:

Call Chemtrec (844) 646-0545

Product Name: Material Number: Use: MULTRANOL 3901 02916227 Polyol components for the production of polyurethanes

2. Hazards Identification

This product is not classified as hazardous according to OSHA HazCom 2012 (29 CFR 1910.1200).

3. Composition/Information on Ingredients

Hazardous Components

There are no hazardous components above the relevant concentration limits according to OSHA HazCom 2012.

4. First Aid Measures

Most Important Symptom(s)/Effect(s)

Acute: Not expected to cause adverse acute health effects.

Eye Contact

In case of contact, flush eyes with plenty of lukewarm water. Get medical attention if irritation develops.

Skin Contact

In case of skin contact, wash affected areas with soap and water. Thoroughly clean shoes before reuse. Wash clothing before reuse. Get medical attention if irritation develops and persists.

Inhalation

If inhaled, remove to fresh air. If breathing is difficult, give oxygen. Get medical attention if irritation

Material Name: MULTRANOL 3901

Material Number: 02916227

Page: 1 of 6

develops.

Ingestion

If ingested, do not induce vomiting unless directed to do so by medical personnel. Get medical attention.

5. Firefighting Measures	
Suitable Extinguishing Media:	Carbon dioxide (CO2), Dry chemical, Foam, water spray for large fires.
Unsuitable Extinguishing Media	No Data Available

Fire Fighting Procedure

Firefighters should be equipped with self-contained breathing apparatus to protect against potentially toxic and irritating fumes. Toxic and irritating gases/fumes may be given off during burning or thermal decomposition. Use cold water spray to cool fire-exposed containers to minimize the risk of rupture.

Hazardous Decomposition Products

By Fire and Thermal Decomposition: Carbon dioxide (CO2), carbon monoxide (CO), oxides of nitrogen (NOx), dense black smoke., Other undetermined compounds

6. Accidental Release Measures

Spill and Leak Procedures

Dike or dam spilled material and control further spillage, if possible. Cover spill with inert material (e. g., dry sand or earth) and collect for proper disposal. Collect and place in appropriately marked sealable containers for disposal. Wash spill area with soap and water.

7. Handling and Storage

Handling/Storage Precautions

Handle in accordance with good industrial hygiene and safety practices. Wash thoroughly after handling. Keep container closed when not in use. Material is hygroscopic and may absorb small amounts of atmospheric moisture. If contamination with isocyanates is suspected, do not reseal containers. Avoid inhalation of vapour or mist.

Storage Temperature	
Minimum:	15 °C (59 °F)
Maximum:	48 °C (118.4 °F)

Storage Conditions

Store separate from food products.

Employee education and training in the safe use and handling of this product are required under the OSHA Hazard Communication Standard 29 CFR 1910.1200.

Substances to Avoid

Oxidizing agents, Isocyanates

_Material Name: MULTRANOL 3901

8. Exposure Controls/Personal Protection

The recommendations in this section should not be a substitute for a personal protective equipment (PPE) assessment performed by the employer as required by 29 CFR 1910 Subpart I.

Exposure Limits

Any component which is listed in section 3 and is not listed in this section does not have a known ACGIH TLV, OSHA PEL or supplier recommended occupational exposure limit.

Industrial Hygiene/Ventilation Measures

Use local and general exhaust ventilation to control levels of exposure.

Respiratory Protection

None required under normal conditions of use.

Hand Protection

Ensure gloves remain in good condition during use and replace if any deterioration is observed.

Permeation resistant gloves., Butyl rubber gloves., Nitrile rubber gloves., Neoprene gloves

Eye Protection

Safety glasses with side-shields

Skin Protection

No special skin protection requirements during normal handling and use.

Additional Protective Measures

Employees should wash their hands and face before eating, drinking, or using tobacco products. Educate and train employees in the safe use and handling of this product.

9. Physical and Chemical Properties

State of Matter:	liquid	
Color:	Colorless	
Odor:	slight	
Odor Threshold:	No Data Available	
pH:	Not Established	
Freezing Point:	Not Established	
Boiling Point:	not established	
Flash Point:	193 °C (379.4 °F)	
Evaporation Rate:	No Data Available	
Lower Explosion Limit:	Not Established	
Upper Explosion Limit:	Not Established	
Vapor Pressure:	Not Established	
Vapor Density:	No Data Available	
Density:	1.01 g/cm^3 (<i>i</i>) $25 \text{ °C} (77 \text{ °F})$	
Relative Vapor Density:	No Data Available	
Specific Gravity:	1.01 @ 25 °C (77 °F)	
Solubility in Water:	completely soluble	
Partition Coefficient: n-	No Data Available	
octanol/water:		
Auto-ignition Temperature:	Not Established	
Matarial Names MIII TRANIOL 2001		
Material Name: MULTRANOL 3901	Material Number: 02916227	

Page: 3 of 6

Decomposition Temperature: Unblocking Temperature: Dynamic Viscosity: Kinematic Viscosity: Bulk Density: Molecular Weight: Hygroscopicity: Self Ignition: Not established No Data Available 1,020 - 1,220 mPa.s @ 25 °C (77 °F) No Data Available 1,010 kg/m3 No Data Available hygroscopic not applicable

10. Stability and Reactivity

Hazardous Reactions

Hazardous polymerisation does not occur.

Stability

Stable

Materials to Avoid

Oxidizing agents, Isocyanates

Conditions to Avoid None known.

Hazardous Decomposition Products

By Fire and Thermal Decomposition: Carbon dioxide (CO2), carbon monoxide (CO), oxides of nitrogen (NOx), dense black smoke., Other undetermined compounds

11. Toxicological Information	가지 않는 것을 가지 않는 것을 가지 않는 것을 가지 않는 것을 했다. 그는 것은 것은 가지 가지 않는 것은 가지 않는 것을 같다. 또한 것을 하는 것을 가지 않는 것을 것을 하는 것을 하는 것을 하는 것을 하는 것을 하는 것을 하는 것을 것을 했다. 것을 것을 하는
Likely Routes of Exposure:	Skin Contact Eye Contact

Health Effects and Symptoms

Acute: Not expected to cause adverse acute health effects. Chronic: Not expected to cause adverse chronic health effects.

Toxicity Data for: MULTRANOL 3901

No data available for this product.

Carcinogenicity:

No carcinogenic substances as defined by IARC, NTP and/or OSHA

12. Ecological Information

Ecological Data for: MULTRANOL 3901

No data available for this component.

Material Name: MULTRANOL 3901

Material Number: 02916227 Page: 4 of 6

13. Disposal Considerations

Waste Disposal Method

Waste disposal should be in accordance with existing federal, state and local environmental control laws.

Empty Container Precautions

Recondition or dispose of empty container in accordance with governmental regulations. Empty containers retain product residue; observe all precautions for product. Do not heat or cut container with electric or gas torch.

14. Transportation Information

Land transport (DOT) Non-Regulated

<u>Sea transport (IMDG)</u> Non-Regulated

<u>Air transport (ICAO/IATA)</u> Non-Regulated

15. Regulatory Information

United States Federal Regulations

US. Toxic Substances Control Act: Listed on the Active Portion of the TSCA Inventory.

No substances are subject to TSCA 12(b) export notification requirements.

US. EPA CERCLA Hazardous Substances (40 CFR 302) Components: None

SARA Section 311/312 Hazard Categories:

Refer to hazard classification information in Section 2.

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 302 Extremely Hazardous Substance (40 CFR 355, Appendix A) Components: None

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 313 Toxic Chemicals (40 CFR 372.65) - Supplier Notification Required Components: None

US. EPA Resource Conservation and Recovery Act (RCRA) Composite List of Hazardous Wastes and Appendix VIII Hazardous Constituents (40 CFR 261):

Under RCRA, it is the responsibility of the person who generates a solid waste, as defined in 40 CFR 261.2, to determine if that waste is a hazardous waste.

State Right-To-Know Information

The following chemicals are specifically listed by individual states; other product specific health and safety data in other sections of the SDS may also be applicable for state requirements. For details on your regulatory requirements you should contact the appropriate agency in your state.

Material Name: MULTRANOL 3901

Massachusetts, New Jersey or Pennsylvania Right to Know Substance Lists:ConcentrationComponents>=1%Polyether Polyol9082-00-2

California Proposition 65 List: None.

CFATS (Chemical Facility Anti-Terrorism Standards) Chemicals

To the best of our knowledge, this product does not contain Appendix A Chemicals of Interest (COI), at or above the Screening Threshold Quantity (STQ), as defined by the Department of Homeland Security Chemical Facility Anti-terrorism Standard (CFATS, 6 CFR Part 27).

Based on information provided by our suppliers, this product is considered "DRC Conflict Free" as defined by the SEC Conflict Minerals Final Rule (Release No. 34-67716; File No. S7-40-10; Date: 2012-08-22).

16. Other Information

The method of hazard communication for Covestro LLC is comprised of product labels and safety data sheets. Safety data sheets for all of our products and general product declarations are available for download at www.productsafetyfirst.covestro.com.

Contact:	Product Safety Department
Telephone:	(412) 413-2835
Version Date:	05/07/2021
SDS Version:	2.8

Information contained in this SDS is believed to be accurate but is furnished without warranty, express or implied, including warranties of merchantability or fitness for a particular purpose. The information relates only to the specific material designated herein. Covestro LLC. assumes no legal responsibility for use of or reliance upon the information in this SDS and such information shall in no case be considered a part of our terms and conditions of sale. The user is responsible for determining whether the Covestro product is suitable for user's method of use or application. Covestro is not liable for any failure to observe the precautionary measures described in this SDS or for any misuse of the product.

Changes since the last version are highlighted in the margin. This version replaces all previous versions.

Material Name: MULTRANOL 3901

Page: 6 of 6

SAFETY DATA SHEET



1. Identification

Covestro LLC 1 Covestro Circle Pittsburgh, PA 15205 USA TRANSPORTATION EMERGENCY CALL CHEMTREC: INTERNATIONAL:

(800) 424-9300 (703) 527-3887

NON-TRANSPORTATION Emergency Phone: Information Phone:

Call Chemtrec (844) 646-0545

Product Name: Material Number: Use: HYPERLITE® POLYOL E-833 04216435 Polyol components for the production of polyurethanes

2. Hazards Identification

This product is not classified as hazardous according to OSHA HazCom 2012 (29 CFR 1910.1200).

3. Composition/Information on Ingredients

Hazardous Components

There are no hazardous components above the relevant concentration limits according to OSHA HazCom 2012.

4. First Aid Measures

Most Important Symptom(s)/Effect(s) Acute: Not expected to cause adverse acute health effects.

Eye Contact

In case of contact, flush eyes with plenty of lukewarm water. Get medical attention if irritation develops.

Skin Contact

In case of skin contact, wash affected areas with soap and water. Thoroughly clean shoes before reuse. Wash clothing before reuse. Get medical attention if irritation develops and persists.

Inhalation

If inhaled, remove to fresh air. If breathing is difficult, give oxygen. Get medical attention if irritation

Material Name: HYPERLITE® POLYOL E-833

Material Number: 04216435

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develops.

Ingestion

If ingested, do not induce vomiting unless directed to do so by medical personnel. Get medical attention.

5. Firefighting Measures	
Suitable Extinguishing Media:	Carbon dioxide (CO2), Dry chemical, Foam, water spray for large fires.
Unsuitable Extinguishing Media	No Data Available

Fire Fighting Procedure

Firefighters should be equipped with self-contained breathing apparatus to protect against potentially toxic and irritating fumes. Toxic and irritating gases/fumes may be given off during burning or thermal decomposition. Use cold water spray to cool fire-exposed containers to minimize the risk of rupture.

Hazardous Decomposition Products

By Fire and Thermal Decomposition: Carbon dioxide (CO2), carbon monoxide (CO), oxides of nitrogen (NOx), dense black smoke., Other undetermined compounds

6. Accidental Release Measures

Spill and Leak Procedures

Dike or dam spilled material and control further spillage, if possible. Cover spill with inert material (e. g., dry sand or earth) and collect for proper disposal. Collect and place in appropriately marked sealable containers for disposal. Wash spill area with soap and water.

7. Handling and Storage

Handling/Storage Precautions

Handle in accordance with good industrial hygiene and safety practices. Wash thoroughly after handling. Keep container closed when not in use. Material is hygroscopic and may absorb small amounts of atmospheric moisture. If contamination with isocyanates is suspected, do not reseal containers. Avoid inhalation of vapour or mist.

Storage Temperature	
Minimum:	20 °C (68 °F)
Maximum:	60 °C (140 °F)

Storage Conditions Store separate from food products.

Employee education and training in the safe use and handling of this product are required under the OSHA Hazard Communication Standard 29 CFR 1910.1200.

Substances to Avoid Oxidizing agents, Isocyanates

Material Name: HYPERLITE® POLYOL E-833

Material Number: 04216435

Page: 2 of 6

8. Exposure Controls/Personal Protection

The recommendations in this section should not be a substitute for a personal protective equipment (PPE) assessment performed by the employer as required by 29 CFR 1910 Subpart I.

Exposure Limits

Country specific exposure limits have not been established or are not applicable

Any component which is listed in section 3 and is not listed in this section does not have a known ACGIH TLV, OSHA PEL or supplier recommended occupational exposure limit.

Industrial Hygiene/Ventilation Measures

Use local and general exhaust ventilation to control levels of exposure.

Respiratory Protection

None required under normal conditions of use.

Hand Protection

Ensure gloves remain in good condition during use and replace if any deterioration is observed.

Permeation resistant gloves., Butyl rubber gloves., Nitrile rubber gloves., Neoprene gloves

Eye Protection

Safety glasses with side-shields

Skin Protection

No special skin protection requirements during normal handling and use.

Additional Protective Measures

Employees should wash their hands and face before eating, drinking, or using tobacco products. Educate and train employees in the safe use and handling of this product.

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State of Matter:	liquid		
Appearance:	viscous		
Color:	Colorless		
Odor:	mild		
Odor Threshold:	No Data Available		
pH:	no data available		
Boiling Point:	> 200 °C (> 392 °F)		
Flash Point:	> 93.33 °C (199.99 °F)		
Evaporation Rate:	No Data Available		
Lower explosion limit:	No Data Available		
Upper Explosion Limit:	No Data Available		
Vapor Pressure:	< 0.01 mmHg @ 20 °C (68 °F)		
Vapor Density:	No Data Available		
Density:	1.025 g/cm ³ @ 20 °C (68 °F) (DIN 51757)		
Relative Vapor Density:	No Data Available		
Specific Gravity:	1.025 @ 20 °C (68 °F)		
Solubility in Water:	slightly soluble		
Partition Coefficient: n-	No Data Available		
P			
Material Name: HYPERLITE® POLYOL	E-833 Material Number: 04216435		

Page: 3 of 6

octanol/water: Auto-ignition Temperature: Decomposition Temperature: Unblocking Temperature: Dynamic Viscosity: Kinematic Viscosity: Bulk Density: Molecular Weight: Pour point: Hygroscopicity:

No Data Available Not established No Data Available No Data Available 1,028 - 1,030 kg/m3 No Data Available -12.22 - -8.89 °C (10 - 16 °F) hygroscopic

10. Stability and Reactivity

Hazardous Reactions

Hazardous polymerisation does not occur.

Stability Stable

Materials to Avoid Oxidizing agents, Isocyanates

Conditions to Avoid None known.

Hazardous Decomposition Products

By Fire and Thermal Decomposition: Carbon dioxide (CO2), carbon monoxide (CO), oxides of nitrogen (NOx), dense black smoke., Other undetermined compounds

11.	T	oxicol	logical	Info	rmation	

Likely Routes of Exposure:

Skin Contact Eye Contact

Health Effects and Symptoms

Acute: Not expected to cause adverse acute health effects. Chronic: Not expected to cause adverse chronic health effects.

Toxicity Data for: HYPERLITE® POLYOL E-833

The components in this product are either not classified, below the relevant concentration limits, or do not have any toxicity data associated with them.

Carcinogenicity:

No carcinogenic substances as defined by IARC, NTP and/or OSHA

12. Ecological Information

Ecological Data for: HYPERLITE® POLYOL E-833

Material Name: HYPERLITE® POLYOL E-833

Material Number: 04216435

Page: 4 of 6

No data available for this product.

13. Disposal Considerations

Waste Disposal Method

Waste disposal should be in accordance with existing federal, state and local environmental control laws.

Empty Container Precautions

Recondition or dispose of empty container in accordance with governmental regulations. Empty containers retain product residue; observe all precautions for product. Do not heat or cut container with electric or gas torch.

14. Transportation Information

Land transport (DOT) Non-Regulated

<u>Sea transport (IMDG)</u> Non-Regulated

<u>Air transport (ICAO/IATA)</u> Non-Regulated

15. Regulatory Information

United States Federal Regulations

US. Toxic Substances Control Act: Listed on the Active Portion of the TSCA Inventory.

No substances are subject to TSCA 12(b) export notification requirements.

US. EPA CERCLA Hazardous Substances (40 CFR 302) Components: None

SARA Section 311/312 Hazard Categories:

Refer to hazard classification information in Section 2.

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 302 Extremely Hazardous Substance (40 CFR 355, Appendix A) Components: None

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 313 Toxic Chemicals (40 CFR 372.65) - Supplier Notification Required Components: None

US. EPA Resource Conservation and Recovery Act (RCRA) Composite List of Hazardous Wastes and Appendix VIII Hazardous Constituents (40 CFR 261):

Under RCRA, it is the responsibility of the person who generates a solid waste, as defined in 40 CFR 261.2, to determine if that waste is a hazardous waste.

State Right-To-Know Information

Material Name: HYPERLITE® POLYOL E-833

Material Number: 04216435

Page: 5 of 6

The following chemicals are specifically listed by individual states; other product specific health and safety data in other sections of the SDS may also be applicable for state requirements. For details on your regulatory requirements you should contact the appropriate agency in your state.

Massachusetts, New Jersey or Pennsylvania Right to Know Substance Lists:

Concentration	Components	CAS-No.
>=1%	Polyether Polyol	CAS# is a trade secret
>=1%	Polyether Polyol	CAS# is a trade secret
>=1%	Polyether Polyol	CAS# is a trade secret
>=1%	Polyether Polyol	CAS# is a trade secret

California Proposition 65 List:

None.

CFATS (Chemical Facility Anti-Terrorism Standards) Chemicals

To the best of our knowledge, this product does not contain Appendix A Chemicals of Interest (COI), at or above the Screening Threshold Quantity (STQ), as defined by the Department of Homeland Security Chemical Facility Anti-terrorism Standard (CFATS, 6 CFR Part 27).

Based on information provided by our suppliers, this product is considered "DRC Conflict Free" as defined by the SEC Conflict Minerals Final Rule (Release No. 34-67716; File No. S7-40-10; Date: 2012-08-22).

16. Other Information

The method of hazard communication for Covestro LLC is comprised of product labels and safety data sheets. Safety data sheets for all of our products and general product declarations are available for download at www.productsafetyfirst.covestro.com.

Contact:	Product Safety Department
Telephone:	(412) 413-2835
Version Date:	05/06/2021
SDS Version:	2.11

Information contained in this SDS is believed to be accurate but is furnished without warranty, express or implied, including warranties of merchantability or fitness for a particular purpose. The information relates only to the specific material designated herein. Covestro LLC. assumes no legal responsibility for use of or reliance upon the information in this SDS and such information shall in no case be considered a part of our terms and conditions of sale. The user is responsible for determining whether the Covestro product is suitable for user's method of use or application. Covestro is not liable for any failure to observe the precautionary measures described in this SDS or for any misuse of the product.

Changes since the last version are highlighted in the margin. This version replaces all previous versions.

SAFETY DATA SHEET



1. Identification

Covestro LLC 1 Covestro Circle Pittsburgh, PA 15205 USA

TRANSPORTATION EMERGENCY CALL CHEMTREC:

(800) 424-9300 (703) 527-3887

NON-TRANSPORTATION Emergency Phone: Information Phone:

INTERNATIONAL:

Call Chemtrec (844) 646-0545

Product Name: Material Number: Chemical Family: Use: HYPERLITE® POLYOL E-855 83470772 Polymer Polyol Polyol components for the production of polyurethanes

2. Hazards Identification

This product is not classified as hazardous according to OSHA HazCom 2012 (29 CFR 1910.1200).

3. Composition/Information on Ingredients

Hazardous Components

There are no hazardous components above the relevant concentration limits according to OSHA HazCom 2012.

4. First Aid Measures

Most Important Symptom(s)/Effect(s)

Acute: Not expected to cause adverse acute health effects.

Eye Contact

In case of contact, flush eyes with plenty of lukewarm water. Get medical attention if irritation develops.

Skin Contact

In case of skin contact, wash affected areas with soap and water. Thoroughly clean shoes before reuse. Wash clothing before reuse. Get medical attention if irritation develops and persists.

Inhalation

Material Name: HYPERLITE® POLYOL E-855

Page: 1 of 7

Material Number: 83470772

If inhaled, remove to fresh air. If breathing is difficult, give oxygen. Get medical attention if irritation develops.

Ingestion

If ingested, do not induce vomiting unless directed to do so by medical personnel. Get medical attention.

5. Firefighting Measures	
Suitable Extinguishing Media:	Carbon dioxide (CO2), Dry chemical, Foam, water spray for large fires.
Unsuitable Extinguishing Media	No Data Available

Fire Fighting Procedure

Firefighters should be equipped with self-contained breathing apparatus to protect against potentially toxic and irritating fumes. Toxic and irritating gases/fumes may be given off during burning or thermal decomposition. Use cold water spray to cool fire-exposed containers to minimize the risk of rupture.

Hazardous Decomposition Products

By Fire and Thermal Decomposition: Carbon dioxide (CO2), carbon monoxide (CO), oxides of nitrogen (NOx), dense black smoke., Other undetermined compounds

6. Accidental Release Measures

Spill and Leak Procedures

Dike or dam spilled material and control further spillage, if possible. Cover spill with inert material (e. g., dry sand or earth) and collect for proper disposal. Collect and place in appropriately marked sealable containers for disposal. Wash spill area with soap and water.

7. Handling and Storage

Handling/Storage Precautions

Handle in accordance with good industrial hygiene and safety practices. Wash thoroughly after handling. Keep container closed when not in use. Material is hygroscopic and may absorb small amounts of atmospheric moisture. If contamination with isocyanates is suspected, do not reseal containers. Avoid inhalation of vapour or mist.

Storage Temperature	
Minimum:	20 °C (68 °F)
Maximum:	60 °C (140 °F)

Storage Conditions Store separate from food products.

Employee education and training in the safe use and handling of this product are required under the OSHA Hazard Communication Standard 29 CFR 1910.1200.

Substances to Avoid

Oxidizing agents, Isocyanates

Material Name: HYPERLITE® POLYOL E-855

Material Number: 83470772

8. Exposure Controls/Personal Protection

The recommendations in this section should not be a substitute for a personal protective equipment (PPE) assessment performed by the employer as required by 29 CFR 1910 Subpart I.

Exposure Limits

Country specific exposure limits have not been established or are not applicable

Any component which is listed in section 3 and is not listed in this section does not have a known ACGIH TLV, OSHA PEL or supplier recommended occupational exposure limit.

Industrial Hygiene/Ventilation Measures

Use local and general exhaust ventilation to control levels of exposure.

Respiratory Protection

None required under normal conditions of use., NIOSH approved air-supplied respirator during die cleaning, high temperature processing or when thermal decomposition is suspected.

Hand Protection

Ensure gloves remain in good condition during use and replace if any deterioration is observed.

Permeation resistant gloves., butyl-rubber, Nitrile rubber, Neoprene gloves

Eye Protection

Safety glasses with side-shields

Skin Protection

Wear as appropriate:, Impervious protective clothing.

Additional Protective Measures

Employees should wash their hands and face before eating, drinking, or using tobacco products. Educate and train employees in the safe use and handling of this product.

9. Physical and Chemical Properties

State of Matter:	liquid	
Appearance:	viscous	
Color:	White	
Odor:	mild	
Odor Threshold:	No Data Available	
pH:	4 - 8	
Boiling Point:	> 200 °C (> 392 °F) (DIN 53171)	
Flash Point:	213 - 232 °C (415.4 - 449.6 °F) (DIN EN 22719)	
Evaporation Rate:	No Data Available	
Lower explosion limit:	No Data Available	
Upper Explosion Limit:	No Data Available	
Vapor Pressure:	< 0.001 mmHg @ 20 °C (68 °F)	
Vapor Density:	No Data Available	
Density:	1,055 kg/m3 @ 20 °C (68 °F) (DIN 51757)	
Relative Vapor Density:	No Data Available	
Specific Gravity:	1.06 @ 25 °C (77 °F)	
Material Name: HYPERLITE® POLY	OL E-855 Material Number: 83470772	
Page: 3 of 7		

Solubility in Water: slightly soluble Partition Coefficient: n-No Data Available octanol/water: Auto-ignition Temperature: No Data Available **Decomposition Temperature:** Not established Dynamic Viscosity: 4,119 - 6,024 cps @ 25 °C (77 °F) (DIN 53019) Kinematic Viscosity: No Data Available < 130,000 Molecular Weight: -21 - -13 °C (-5.8 - 8.6 °F) (ISO 3016) **Pour point:** hygroscopic Hygroscopicity:

10. Stability and Reactivity

Hazardous Reactions

Hazardous polymerisation does not occur.

Stability Stable

Materials to Avoid

Oxidizing agents, Isocyanates

Conditions to Avoid

None known.

Hazardous Decomposition Products

By Fire and Thermal Decomposition: Carbon dioxide (CO2), carbon monoxide (CO), oxides of nitrogen (NOx), dense black smoke., Other undetermined compounds

11. Toxicological Information

Likely Routes of Exposure:

Skin Contact Eye Contact

Health Effects and Symptoms

Acute: Not expected to cause adverse acute health effects. **Chronic:** Not expected to cause adverse chronic health effects.

Toxicity Data for: HYPERLITE® POLYOL E-855

The components in this product are either not classified, below the relevant concentration limits, or do not have any toxicity data associated with them.

Acute Oral Toxicity

Acute toxicity estimate: 4,249 mg/kg (Calculation method)

Carcinogenicity:

No carcinogenic substances as defined by IARC, NTP and/or OSHA

12. Ecological Information

Material Name: HYPERLITE® POLYOL E-855

Page: 4 of 7

Ecological Data for: HYPERLITE® POLYOL E-855

No data available for this product. The components in this product are either not classified, below the relevant concentration limits, or do not have any ecotoxicity data.

13. Disposal Considerations

Waste Disposal Method

Waste disposal should be in accordance with existing federal, state and local environmental control laws.

Empty Container Precautions

Recondition or dispose of empty container in accordance with governmental regulations. Empty containers retain product residue; observe all precautions for product. Do not heat or cut container with electric or gas torch.

14. Transportation Information

Land transport (DOT) Non-Regulated

<u>Sea transport (IMDG)</u> Non-Regulated

<u>Air transport (ICAO/IATA)</u> Non-Regulated

15. Regulatory Information

United States Federal Regulations

US. Toxic Substances Control Act: Listed on the Active Portion of the TSCA Inventory.

No substances are subject to TSCA 12(b) export notification requirements.

US. EPA CERCLA Hazardous Substances (40 CFR 302) Components:

None

SARA Section 311/312 Hazard Categories:

Refer to hazard classification information in Section 2.

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 302 Extremely Hazardous Substance (40 CFR 355, Appendix A) Components: None

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 313 Toxic Chemicals (40 CFR 372.65) - Supplier Notification Required Components: None

US. EPA Resource Conservation and Recovery Act (RCRA) Composite List of Hazardous Wastes and Appendix VIII Hazardous Constituents (40 CFR 261):

Under RCRA, it is the responsibility of the person who generates a solid waste, as defined in 40 CFR 261.2, to determine if that waste is a hazardous waste.

Material Name: HYPERLITE® POLYOL E-855	
	-

State Right-To-Know Information

The following chemicals are specifically listed by individual states; other product specific health and safety data in other sections of the SDS may also be applicable for state requirements. For details on your regulatory requirements you should contact the appropriate agency in your state.

Massachusetts, New Jersey or Pennsylvania Right to Know Substance Lists:

Concentration	Components	CAS-No.
>=1%	Polyether Polyol	CAS# is a trade secret
>=1%	Polymer	CAS# is a trade secret
>=1%	Polymer	CAS# is a trade secret
>=1%	Polymer	CAS# is a trade secret

New Jersey Environmental Hazardous Substances List and/or New Jersey RTK Special Hazardous Substances Lists:

Concentration	<u>Components</u>	CAS-No.
>=1%	Polymer	CAS# is a trade secret

Massachusetts Right to Know Extraordinarily Hazardous Substance List:

Concentration	Components	CAS-No.
<=25 ppm	Acrylonitrile	107-13-1
<=50 ppm	Styrene	100-42-5

California Proposition 65 List:

Concentration	Components	CAS-No.
<=25 ppm	Acrylonitrile	107-13-1
<=50 ppm	Styrene	100-42-5

CFATS (Chemical Facility Anti-Terrorism Standards) Chemicals

To the best of our knowledge, this product does not contain Appendix A Chemicals of Interest (COI), at or above the Screening Threshold Quantity (STQ), as defined by the Department of Homeland Security Chemical Facility Anti-terrorism Standard (CFATS, 6 CFR Part 27).

CFATS (Chemical Facility Anti-Terrorism Standards) Chemicals

To the best of our knowledge, this product does not contain Appendix A Chemicals of Interest (COI), at or above the Screening Threshold Quantity (STQ), as defined by the Department of Homeland Security Chemical Facility Anti-terrorism Standard (CFATS, 6 CFR Part 27).

Based on information provided by our suppliers, this product is considered "DRC Conflict Free" as defined by the SEC Conflict Minerals Final Rule (Release No. 34-67716; File No. S7-40-10; Date: 2012-08-22).

16. Other Information

The method of hazard communication for Covestro LLC is comprised of product labels and safety data sheets. Safety data sheets for all of our products and general product declarations are available for download at www.productsafetyfirst.covestro.com.

Contact: Telephone:	Product Safety Department (412) 413-2835
Version Date:	09/24/2019
SDS Version:	3.5

Material Name: HYPERLITE® POLYOL E-855

Information contained in this SDS is believed to be accurate but is furnished without warranty, express or implied, including warranties of merchantability or fitness for a particular purpose. The information relates only to the specific material designated herein. Covestro LLC. assumes no legal responsibility for use of or reliance upon the information in this SDS and such information shall in no case be considered a part of our terms and conditions of sale. The user is responsible for determining whether the Covestro product is suitable for user's method of use or application. Covestro is not liable for any failure to observe the precautionary measures described in this SDS or for any misuse of the product.

Changes since the last version are highlighted in the margin. This version replaces all previous versions.

Material Name: HYPERLITE® POLYOL E-855



THE DOW CHEMICAL COMPANY

Product name: VORANOL™ 360 Polyol DA

Issue Date: 01/11/2019 **Print Date:** 01/15/2020

THE DOW CHEMICAL COMPANY encourages and expects you to read and understand the entire (M)SDS, as there is important information throughout the document. We expect you to follow the precautions identified in this document unless your use conditions would necessitate other appropriate methods or actions.

1. IDENTIFICATION

Product name: VORANOL[™] 360 Polyol DA

Recommended use of the chemical and restrictions on use

Identified uses: For industrial use. Component(s) for the manufacture of urethane polymers. We recommend that you use this product in a manner consistent with the listed use. If your intended use is not consistent with the stated use, please contact your sales or technical service representative.

COMPANY IDENTIFICATION

THE DOW CHEMICAL COMPANY 2030 DOW CENTER MIDLAND MI 48674-0000 UNITED STATES

Customer Information Number:

800-258-2436 SDSQuestion@dow.com

EMERGENCY TELEPHONE NUMBER

24-Hour Emergency Contact: CHEMTREC +1 800-424-9300 Local Emergency Contact: 800-424-9300

2. HAZARDS IDENTIFICATION

Hazard classification

GHS classification in accordance with 29 CFR 1910.1200 Not a hazardous substance or mixture.

Other hazards

No data available

3. COMPOSITION/INFORMATION ON INGREDIENTS

Synonyms: Sucrose. propoxylated This product is a substance.

Component

CASRN

Concentration

Sucrose, propylene oxide	9049-71-2	>= 65.0 - <= 75.0 %
Glycerol propylene oxide	25791-96-2	>= 25.0 - <= 35.0 %

4. FIRST AID MEASURES

Description of first aid measures

General advice:

First Aid responders should pay attention to self-protection and use the recommended protective clothing (chemical resistant gloves, splash protection). If potential for exposure exists refer to Section 8 for specific personal protective equipment.

Inhalation: Move person to fresh air; if effects occur, consult a physician.

Skin contact: Wash off with plenty of water. Suitable emergency safety shower facility should be immediately available.

Eye contact: Flush eyes thoroughly with water for several minutes. Remove contact lenses after the initial 1-2 minutes and continue flushing for several additional minutes. If effects occur, consult a physician, preferably an ophthalmologist.

Ingestion: If swallowed, seek medical attention. Do not induce vomiting unless directed to do so by medical personnel.

Most important symptoms and effects, both acute and delayed:

Aside from the information found under Description of first aid measures (above) and Indication of immediate medical attention and special treatment needed (below), any additional important symptoms and effects are described in Section 11: Toxicology Information.

Indication of any immediate medical attention and special treatment needed

Notes to physician: If burn is present, treat as any thermal burn, after decontamination. No specific antidote. Treatment of exposure should be directed at the control of symptoms and the clinical condition of the patient.

5. FIREFIGHTING MEASURES

Suitable extinguishing media: Water fog or fine spray. Dry chemical fire extinguishers. Carbon dioxide fire extinguishers. Foam. Alcohol resistant foams (ATC type) are preferred. General purpose synthetic foams (including AFFF) or protein foams may function, but will be less effective.

Unsuitable extinguishing media: Do not use direct water stream. May spread fire.

Special hazards arising from the substance or mixture

Hazardous combustion products: During a fire, smoke may contain the original material in addition to combustion products of varying composition which may be toxic and/or irritating. Combustion products may include and are not limited to: Carbon monoxide. Carbon dioxide.

Unusual Fire and Explosion Hazards: Container may rupture from gas generation in a fire situation. Violent steam generation or eruption may occur upon application of direct water stream to hot liquids.

Advice for firefighters

Fire Fighting Procedures: Keep people away. Isolate fire and deny unnecessary entry. Use water spray to cool fire exposed containers and fire affected zone until fire is out and danger of reignition has passed. Fight fire from protected location or safe distance. Consider the use of unmanned hose holders or monitor nozzles. Immediately withdraw all personnel from the area in case of rising sound from venting safety device or discoloration of the container. Do not use direct water stream. May spread fire. Move container from fire area if this is possible without hazard. Burning liquids may be moved by flushing with water to protect personnel and minimize property damage. Contain fire water run-off if possible. Fire water run-off, if not contained, may cause environmental damage. Review the "Accidental Release Measures" and the "Ecological Information" sections of this (M)SDS.

Special protective equipment for firefighters: Wear positive-pressure self-contained breathing apparatus (SCBA) and protective fire fighting clothing (includes fire fighting helmet, coat, trousers, boots, and gloves). Avoid contact with this material during fire fighting operations. If contact is likely, change to full chemical resistant fire fighting clothing with self-contained breathing apparatus. If this is not available, wear full chemical resistant clothing with self-contained breathing apparatus and fight fire from a remote location. For protective equipment in post-fire or non-fire clean-up situations, refer to the relevant sections.

6. ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures: Isolate area. Refer to section 7, Handling, for additional precautionary measures. Keep unnecessary and unprotected personnel from entering the area. Spilled material may cause a slipping hazard. Use appropriate safety equipment. For additional information, refer to Section 8, Exposure Controls and Personal Protection.

Environmental precautions: Prevent from entering into soil, ditches, sewers, waterways and/or groundwater. See Section 12, Ecological Information.

Methods and materials for containment and cleaning up: Contain spilled material if possible. Absorb with materials such as: Dirt. Sand. Sawdust. Collect in suitable and properly labeled containers. Wash the spill site with water. See Section 13, Disposal Considerations, for additional information.

7. HANDLING AND STORAGE

Precautions for safe handling: Wash thoroughly after handling. Keep container closed. This material is hygroscopic in nature. Product shipped/handled hot can cause thermal burns. See Section 8, EXPOSURE CONTROLS AND PERSONAL PROTECTION. Spills of these organic materials on hot fibrous insulations may lead to lowering of the autoignition

temperatures possibly resulting in spontaneous combustion.

Conditions for safe storage: Protect from atmospheric moisture. Store in a dry place. Avoid prolonged exposure to heat and air. Store in the following material(s): Carbon steel. Stainless steel. Polypropylene. Polyethylene-lined container. Teflon. Glass-lined container. Aluminum. Plasite 3066

lined container. Plasite 3070 lined container. 316 stainless steel. See Section 10 for more specific information.

Storage stability

Storage temperature:Storage Period:15 - 35 °C (59 - 95 °F)24 Month

8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Control parameters

If exposure limits exist, they are listed below. If no exposure limits are displayed, then no values are applicable.

Exposure controls

Engineering controls: Use local exhaust ventilation, or other engineering controls to maintain airborne levels below exposure limit requirements or guidelines. If there are no applicable exposure limit requirements or guidelines, general ventilation should be sufficient for most operations. Local exhaust ventilation may be necessary for some operations.

Individual protection measures

Eye/face protection: Use safety glasses (with side shields). When handling hot material: Use chemical goggles. Wear a face-shield which allows use of chemical goggles, or wear a full-face respirator, to protect face and eyes when there is any likelihood of splashes. **Skin protection**

Hand protection: Chemical protective gloves should not be needed when handling this material. Consistent with general hygienic practice for any material, skin contact should be minimized. Use gloves with insulation for thermal protection, when needed. **Other protection:** No precautions other than clean body-covering clothing should be needed. When handling hot material, protect skin from thermal burns. Selection of specific items will depend on the operation.

Respiratory protection: Respiratory protection should be worn when there is a potential to exceed the exposure limit requirements or guidelines. If there are no applicable exposure limit requirements or guidelines, wear respiratory protection when adverse effects, such as respiratory irritation or discomfort have been experienced, or where indicated by your risk assessment process. For most conditions, no respiratory protection should be needed; however, if material is heated or sprayed, use an approved air-purifying respirator. The following should be effective types of air-purifying respirators: Organic vapor cartridge with a particulate pre-filter.

9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance

Physical state Color Odor Odor Threshold pH Melting point/range Freezing point Liquid. Colorless to yellow Mild No test data available > 7 - < 12 *Estimated.* No test data available No test data available ۰.

Boiling point (760 mmHg) Flash point Evaporation Rate (Butyl Acetate = 1)	Decomposes at elevated temperature closed cup >163 °C(325 °F) <i>ASTM D 93</i> No test data available
Flammability (solid, gas)	Not Applicable
Lower explosion limit	No test data available
Upper explosion limit	No test data available
Vapor Pressure	Literature negligible at ambient temperature
Relative Vapor Density (air = 1)	>1 Literature
Relative Density (water = 1)	> 1.00 at 25 °C (77 °F) / 25 °C <i>Literature</i>
Water solubility	soluble
Partition coefficient: n- octanol/water	No data available
Auto-ignition temperature	No test data available
Decomposition temperature	No data available
Dynamic Viscosity	1,100 mPa.s at 25 °C(77 °F) <i>Estimated</i> .
Kinematic Viscosity	No test data available
Explosive properties	Not explosive
Oxidizing properties	No
Liquid Density	1.0840 g/cm3 at 25 °C (77 °F) <i>ASTM D4669</i>
Molecular weight	No data available

NOTE: The physical data presented above are typical values and should not be construed as a specification.

10. STABILITY AND REACTIVITY

Reactivity: No data available

Chemical stability: Stable under recommended storage conditions. See Storage, Section 7.

Possibility of hazardous reactions: Will not occur by itself.

Conditions to avoid: Product can oxidize at elevated temperatures. Generation of gas during decomposition can cause pressure in closed systems.

Incompatible materials: Avoid contact with oxidizing materials. Avoid contact with: Strong acids. Strong bases. Avoid unintended contact with isocyanates. The reaction of polyols and isocyanates generates heat.

Hazardous decomposition products: Decomposition products depend upon temperature, air supply and the presence of other materials. Decomposition products can include and are not limited to: Carbon dioxide. Alcohols. Ethers. Hydrocarbons. Ketones. Polymer fragments.

11. TOXICOLOGICAL INFORMATION

Toxicological information appears in this section when such data is available.

Acute toxicity

Acute oral toxicity

Low toxicity if swallowed. Small amounts swallowed incidentally as a result of normal handling operations are not likely to cause injury; however, swallowing larger amounts may cause injury.

Typical for this family of materials.

LD50, Rat, > 2,000 mg/kg No deaths occurred at this concentration.

Acute dermal toxicity

Prolonged skin contact is unlikely to result in absorption of harmful amounts.

Typical for this family of materials.

LD50, Rabbit, > 2,000 mg/kg No deaths occurred at this concentration.

Acute inhalation toxicity

At room temperature, exposure to vapor is minimal due to low volatility; single exposure is not likely to be hazardous. Vapor from heated material or mist may cause respiratory irritation. The LC50 has not been determined.

Skin corrosion/irritation

Prolonged exposure not likely to cause significant skin irritation. Material may be handled at elevated temperatures; contact with heated material may cause thermal burns.

Serious eye damage/eye irritation

May cause pain disproportionate to the level of irritation to eye tissues. May cause slight temporary eye irritation. Corneal injury is unlikely.

Sensitization

Contains component(s) which have not demonstrated the potential for contact allergy in mice. Based on information for component(s): Did not cause allergic skin reactions when tested in guinea pigs.

For respiratory sensitization: No relevant data found.

Specific Target Organ Systemic Toxicity (Single Exposure)

Evaluation of available data suggests that this material is not an STOT-SE toxicant.

Specific Target Organ Systemic Toxicity (Repeated Exposure) Based on available data, repeated exposures are not anticipated to cause significant adverse effects.

Carcinogenicity

Similar polyols did not cause cancer in long-term animal studies.

Teratogenicity

Based on information for component(s): Did not cause birth defects or any other fetal effects in laboratory animals.

Reproductive toxicity

Contains component(s) which did not interfere with reproduction in animal studies.

Mutagenicity

Based on information for component(s): In vitro genetic toxicity studies were negative.

Aspiration Hazard

Based on physical properties, not likely to be an aspiration hazard.

COMPONENTS INFLUENCING TOXICOLOGY:

Sucrose, propylene oxide

Acute inhalation toxicity

Typical for this family of materials. No deaths occurred following exposure to a saturated atmosphere.

Glycerol propylene oxide

Acute inhalation toxicity The LC50 has not been determined.

12. ECOLOGICAL INFORMATION

Ecotoxicological information appears in this section when such data is available.

Toxicity

Sucrose, propylene oxide

Acute toxicity to fish

Material is practically non-toxic to aquatic organisms on an acute basis (LC50/EC50/EL50/LL50 >100 mg/L in the most sensitive species tested). LC50, Danio rerio (zebra fish), static test, 96 Hour, 6,310 mg/l, OECD Test Guideline 203 or Equivalent

Acute toxicity to aquatic invertebrates

EC50, Daphnia magna (Water flea), static test, 48 Hour, 9,890 mg/l, OECD Test Guideline 202 or Equivalent

Chronic toxicity to aquatic invertebrates

NOEC, Daphnia magna (Water flea), semi-static test, 21 d, mortality, >= 10 mg/l

Glycerol propylene oxide

Acute toxicity to fish

Material is practically non-toxic to aquatic organisms on an acute basis (LC50/EC50/EL50/LL50 >100 mg/L in the most sensitive species tested). For this family of materials: LC50, Leuciscus idus (Golden orfe), semi-static test, 96 Hour, > 1,000 mg/l, OECD Test Guideline 203 or Equivalent

Acute toxicity to aquatic invertebrates

For this family of materials: EC50, Daphnia magna (Water flea), static test, 48 Hour, > 100 mg/l, OECD Test Guideline 202 or Equivalent

Acute toxicity to algae/aquatic plants

For this family of materials: EC50, Desmodesmus subspicatus (green algae), static test, 72 Hour, Growth rate inhibition, > 100 mg/l, OECD Test Guideline 201 or Equivalent

Chronic toxicity to aquatic invertebrates

NOEC, Daphnia magna (Water flea), semi-static test, 21 d, mortality, >= 10 mg/l

Persistence and degradability

Sucrose, propylene oxide

Biodegradability: Based on information for a similar material: Material is inherently biodegradable (reaches > 20% biodegradation in OECD test(s) for inherent biodegradability).

Glycerol propylene oxide

Biodegradability: For this family of materials: Based on stringent OECD test guidelines, this material cannot be considered as readily biodegradable; however, these results do not necessarily mean that the material is not biodegradable under environmental conditions. Material is ultimately biodegradable (reaches > 70% mineralization in OECD test(s) for inherent biodegradability).

10-day Window: Fail

Biodegradation: 40 % Exposure time: 28 d Method: OECD Test Guideline 301B or Equivalent 10-day Window: Not applicable Biodegradation: 99 % Exposure time: 28 d Method: OECD Test Guideline 302B or Equivalent

Bioaccumulative potential

Sucrose, propylene oxide

Bioaccumulation: Bioconcentration potential is low (BCF < 100 or Log Pow < 3). **Partition coefficient:** n-octanol/water(log Pow): -3.38 - -3.25 Estimated.

Glycerol propylene oxide

Bioaccumulation: No bioconcentration is expected because of the relatively high water solubility.

Mobility in soil

Sucrose, propylene oxide

No relevant data found.

Glycerol propylene oxide

No relevant data found.

13. DISPOSAL CONSIDERATIONS

Disposal methods: DO NOT DUMP INTO ANY SEWERS, ON THE GROUND, OR INTO ANY BODY OF WATER. All disposal practices must be in compliance with all Federal, State/Provincial and local

Product name: VORANOL™ 360 Polyol DA

laws and regulations. Regulations may vary in different locations. Waste characterizations and compliance with applicable laws are the responsibility solely of the waste generator. AS YOUR SUPPLIER, WE HAVE NO CONTROL OVER THE MANAGEMENT PRACTICES OR MANUFACTURING PROCESSES OF PARTIES HANDLING OR USING THIS MATERIAL. THE INFORMATION PRESENTED HERE PERTAINS ONLY TO THE PRODUCT AS SHIPPED IN ITS INTENDED CONDITION AS DESCRIBED IN MSDS SECTION: Composition Information. FOR UNUSED & UNCONTAMINATED PRODUCT, the preferred options include sending to a licensed, permitted: Recycler. Reclaimer. Incinerator or other thermal destruction device. For additional information, refer to: Handling & Storage Information, MSDS Section 7 Stability & Reactivity Information, MSDS Section 10 Regulatory Information, MSDS Section 15

14. TRANSPORT INFORMATION

DOT

Not regulated for transport

Classification for SEA transport (IMO-IMDG):

Transport in bulk according to Annex I or II of MARPOL 73/78 and the IBC or IGC Code Not regulated for transport Consult IMO regulations before transporting ocean bulk

Classification for AIR transport (IATA/ICAO):

Not regulated for transport

This information is not intended to convey all specific regulatory or operational requirements/information relating to this product. Transportation classifications may vary by container volume and may be influenced by regional or country variations in regulations. Additional transportation system information can be obtained through an authorized sales or customer service representative. It is the responsibility of the transporting organization to follow all applicable laws, regulations and rules relating to the transportation of the material.

15. REGULATORY INFORMATION

Superfund Amendments and Reauthorization Act of 1986 Title III (Emergency Planning and Community Right-to-Know Act of 1986) Sections 311 and 312 No SARA Hazards

Superfund Amendments and Reauthorization Act of 1986 Title III (Emergency Planning and Community Right-to-Know Act of 1986) Section 313

This material does not contain any chemical components with known CAS numbers that exceed the threshold (De Minimis) reporting levels established by SARA Title III, Section 313.

Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA) Section 103

This material does not contain any components with a CERCLA RQ.

Pennsylvania Worker and Community Right-To-Know Act:

To the best of our knowledge, this product does not contain chemicals at levels which require reporting under this statute.

California Prop. 65

This product does not contain any chemicals known to State of California to cause cancer, birth defects, or any other reproductive harm.

United States TSCA Inventory (TSCA)

All components of this product are in compliance with the inventory listing requirements of the U.S. Toxic Substances Control Act (TSCA) Chemical Substance Inventory.

16. OTHER INFORMATION

Revision

Identification Number: 377078 / A001 / Issue Date: 01/11/2019 / Version: 6.0 Most recent revision(s) are noted by the bold, double bars in left-hand margin throughout this document.

Full text of other abbreviations

AICS - Australian Inventory of Chemical Substances: ASTM - American Society for the Testing of Materials: bw - Body weight: CERCLA - Comprehensive Environmental Response, Compensation. and Liability Act; CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DOT - Department of Transportation; DSL - Domestic Substances List (Canada); ECx - Concentration associated with x% response; EHS - Extremely Hazardous Substance: ELx - Loading rate associated with x% response: EmS - Emergency Schedule: ENCS - Existing and New Chemical Substances (Japan); ErCx - Concentration associated with x% growth rate response; ERG - Emergency Response Guide; GHS - Globally Harmonized System; GLP - Good Laboratory Practice: HMIS - Hazardous Materials Identification System: IARC - International Agency for Research on Cancer; IATA - International Air Transport Association; IBC - International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk; IC50 - Half maximal inhibitory concentration; ICAO - International Civil Aviation Organization; IECSC - Inventory of Existing Chemical Substances in China; IMDG - International Maritime Dangerous Goods; IMO -International Maritime Organization; ISHL - Industrial Safety and Health Law (Japan); ISO -International Organisation for Standardization; KECI - Korea Existing Chemicals Inventory; LC50 -Lethal Concentration to 50 % of a test population; LD50 - Lethal Dose to 50% of a test population (Median Lethal Dose); MARPOL - International Convention for the Prevention of Pollution from Ships; MSHA - Mine Safety and Health Administration; n.o.s. - Not Otherwise Specified; NFPA - National Fire Protection Association; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NTP - National Toxicology Program; NZIoC - New Zealand Inventory of Chemicals; OECD - Organization for Economic Co-operation and Development; OPPTS - Office of Chemical Safety and Pollution Prevention; PBT - Persistent, Bioaccumulative and Toxic substance; PICCS - Philippines Inventory of Chemicals and Chemical Substances; (Q)SAR - (Quantitative) Structure Activity Relationship; RCRA -Resource Conservation and Recovery Act; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals; RQ - Reportable Quantity; SADT - Self-Accelerating Decomposition Temperature; SARA -

Superfund Amendments and Reauthorization Act; SDS - Safety Data Sheet; TCSI - Taiwan Chemical Substance Inventory; TSCA - Toxic Substances Control Act (United States); UN - United Nations; UNRTDG - United Nations Recommendations on the Transport of Dangerous Goods; vPvB - Very Persistent and Very Bioaccumulative

Information Source and References

This SDS is prepared by Product Regulatory Services and Hazard Communications Groups from information supplied by internal references within our company.

THE DOW CHEMICAL COMPANY urges each customer or recipient of this (M)SDS to study it carefully and consult appropriate expertise, as necessary or appropriate, to become aware of and understand the data contained in this (M)SDS and any hazards associated with the product. The information herein is provided in good faith and believed to be accurate as of the effective date shown above. However, no warranty, express or implied, is given. Regulatory requirements are subject to change and may differ between various locations. It is the buyer's/user's responsibility to ensure that his activities comply with all federal, state, provincial or local laws. The information presented here pertains only to the product as shipped. Since conditions for use of the product are not under the control of the manufacturer, it is the buyer's/user's duty to determine the conditions necessary for the safe use of this product. Due to the proliferation of sources for information such as manufacturer-specific (M)SDSs, we are not and cannot be responsible for (M)SDS obtained from any source other than ourselves. If you have obtained an (M)SDS from another source or if you are not sure that the (M)SDS you have is current, please contact us for the most current version. US



THE DOW CHEMICAL COMPANY

Product name: VORANATE™ T-80 Type I TDI DA

Issue Date: 07/26/2019 **Print Date:** 01/15/2020

THE DOW CHEMICAL COMPANY encourages and expects you to read and understand the entire (M)SDS, as there is important information throughout the document. We expect you to follow the precautions identified in this document unless your use conditions would necessitate other appropriate methods or actions.

1. IDENTIFICATION

Product name: VORANATE™ T-80 Type I TDI DA

Recommended use of the chemical and restrictions on use

Identified uses: Component(s) for the manufacture of urethane polymers. We recommend that you use this product in a manner consistent with the listed use. If your intended use is not consistent with the stated use, please contact your sales or technical service representative. For industrial use only.

COMPANY IDENTIFICATION

THE DOW CHEMICAL COMPANY 2030 DOW CENTER MIDLAND MI 48674-0000 UNITED STATES

Customer Information Number:

800-258-2436 SDSQuestion@dow.com

EMERGENCY TELEPHONE NUMBER

24-Hour Emergency Contact: CHEMTREC +1 800-424-9300 Local Emergency Contact: 800-424-9300

2. HAZARDS IDENTIFICATION

Hazard classification

GHS classification in accordance with 29 CFR 1910.1200 Acute toxicity - Category 1 - Inhalation Skin irritation - Category 2 Eye irritation - Category 2A Respiratory sensitisation - Category 1 Skin sensitisation - Category 1 Carcinogenicity - Category 2 Specific target organ toxicity - single exposure - Category 3

Label elements Hazard pictograms



Signal word: DANGER!

Hazards

Causes skin irritation. May cause an allergic skin reaction. Causes serious eye irritation. Fatal if inhaled. May cause allergy or asthma symptoms or breathing difficulties if inhaled. May cause respiratory irritation. Suspected of causing cancer.

Precautionary statements

Prevention

Obtain special instructions before use. Do not handle until all safety precautions have been read and understood. Do not breathe dust/ fume/ gas/ mist/ vapours/ spray. Wash skin thoroughly after handling. Use only outdoors or in a well-ventilated area. Contaminated work clothing must not be allowed out of the workplace. Wear protective gloves/ protective clothing/ eye protection/ face protection. In case of inadequate ventilation wear respiratory protection.

Response

IF ON SKIN: Wash with plenty of soap and water. IF INHALED: Remove person to fresh air and keep comfortable for breathing. Immediately call a POISON CENTER/doctor. IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing, IF exposed or concerned: Get medical advice/ attention. If skin irritation or rash occurs: Get medical advice/ attention. If eye irritation persists: Get medical advice/ attention. Take off contaminated clothing and wash before reuse.

Storage

Store in a well-ventilated place. Keep container tightly closed. Store locked up.

Disposal

Dispose of contents/ container to an approved waste disposal plant.

Other hazards

No data available

3. COMPOSITION/INFORMATION ON INGREDIENTS

Synonyms: Toluene Diisocyanate This product is a substance.		
Component	CASRN	Concentration
Toluene-diisocyanate, mixture of toluene-2,4- di-isocyanate and toluene-2,6-di-isocyanate	26471-62-5	100.0%

Note

Toluene-diisocyanate (TDI) with CAS# 26471-62-5 describes a mixture of toluene-2,4-di-isocyanate and toluene-2,6-di-isocyanate.

4. FIRST AID MEASURES

Description of first aid measures

General advice:

First Aid responders should pay attention to self-protection and use the recommended protective clothing (chemical resistant gloves, splash protection). If potential for exposure exists refer to Section 8 for specific personal protective equipment.

Inhalation: Move person to fresh air. If not breathing, give artificial respiration; if by mouth to mouth use rescuer protection (pocket mask, etc). If breathing is difficult, oxygen should be administered by qualified personnel. Call a physician or transport to a medical facility.

Skin contact: Remove material from skin immediately by washing with soap and plenty of water. Remove contaminated clothing and shoes while washing. Seek medical attention if irritation persists. Wash clothing before reuse. An MDI skin decontamination study demonstrated that cleaning very soon after exposure is important, and that a polyglycol-based skin cleanser or corn oil may be more effective than soap and water. This may also apply to other isocyanates. Discard items which cannot be decontaminated, including leather articles such as shoes, belts and watchbands. Suitable emergency safety shower facility should be available in work area.

Eye contact: Immediately flush eyes with water; remove contact lenses, if present, after the first 5 minutes, then continue flushing eyes for at least 15 minutes. Obtain medical attention without delay, preferably from an ophthalmologist. Suitable emergency eye wash facility should be immediately available.

Ingestion: Do not induce vomiting. Give one cup (8 ounces or 240 ml) of water or milk if available and transport to a medical facility. Do not give anything by mouth unless the person is fully conscious.

Most important symptoms and effects, both acute and delayed:

Aside from the information found under Description of first aid measures (above) and Indication of immediate medical attention and special treatment needed (below), any additional important symptoms and effects are described in Section 11: Toxicology Information.

Indication of any immediate medical attention and special treatment needed

Notes to physician: Maintain adequate ventilation and oxygenation of the patient. May cause respiratory sensitization or asthma-like symptoms. Bronchodilators, expectorants and antitussives may be of help. Treat bronchospasm with inhaled beta2 agonist and oral or parenteral corticosteroids. Respiratory symptoms, including pulmonary edema, may be delayed. Persons receiving significant exposure should be observed 24-48 hours for signs of respiratory distress. If you are sensitized to

diisocyanates, consult your physician regarding working with other respiratory irritants or sensitizers. Due to irritant properties, swallowing may result in burns/ulceration of mouth, stomach and lower gastrointestinal tract with subsequent stricture. Aspiration of vomitus may cause lung injury. Suggest endotracheal/esophageal control if lavage is done. Cholinesterase inhibition has been noted in human exposure but is not of benefit in determining exposure and is not correlated with signs of exposure. Treatment of exposure should be directed at the control of symptoms and the clinical condition of the patient. Excessive exposure may aggravate preexisting asthma and other respiratory disorders (e.g. emphysema, bronchitis, reactive airways dysfunction syndrome).

5. FIREFIGHTING MEASURES

Extinguishing media

Suitable extinguishing media: Water fog or fine spray.. Dry chemical fire extinguishers.. Carbon dioxide fire extinguishers.. Foam.. Alcohol resistant foams (ATC type) are preferred. General purpose synthetic foams (including AFFF) or protein foams may function, but will be less effective..

Unsuitable extinguishing media: Do not use direct water stream.. May spread fire..

Special hazards arising from the substance or mixture

Hazardous combustion products: During a fire, smoke may contain the original material in addition to combustion products of varying composition which may be toxic and/or irritating.. Combustion products may include and are not limited to:. Nitrogen oxides.. Isocyanates.. Hydrogen cyanide.. Carbon monoxide.. Carbon dioxide..

Unusual Fire and Explosion Hazards: Material reacts slowly with water, releasing carbon dioxide which can cause pressure buildup and rupture of closed containers. Elevated temperatures accelerate this reaction.. Container may rupture from gas generation in a fire situation.. Violent steam generation or eruption may occur upon application of direct water stream to hot liquids.. Dense smoke is produced when product burns..

Advice for firefighters

Fire Fighting Procedures: Keep people away. Isolate fire and deny unnecessary entry.. Stay upwind. Keep out of low areas where gases (fumes) can accumulate.. Water is not recommended, but may be applied in large quantities as a fine spray when other extinguishing agents are not available.. Do not use direct water stream. May spread fire.. Fight fire from protected location or safe distance. Consider the use of unmanned hose holders or monitor nozzles.. Immediately withdraw all personnel from the area in case of rising sound from venting safety device or discoloration of the container.. Move container from fire area if this is possible without hazard.. Use water spray to cool fire-exposed containers and fire-affected zone until fire is out.. Contain fire water run-off if possible. Fire water run-off, if not contained, may cause environmental damage.. Review the "Accidental Release Measures" and the "Ecological Information" sections of this (M)SDS..

Special protective equipment for firefighters: Wear positive-pressure self-contained breathing apparatus (SCBA) and protective fire fighting clothing (includes fire fighting helmet, coat, trousers, boots, and gloves). Avoid contact with this material during fire fighting operations. If contact is likely, change to full chemical resistant fire fighting clothing with self-contained breathing apparatus. If this is not available, wear full chemical resistant clothing

with self-contained breathing apparatus and fight fire from a remote location. For protective equipment in post-fire or non-fire clean-up situations, refer to the relevant sections.

6. ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures: Isolate area. Keep unnecessary and unprotected personnel from entering the area. Keep personnel out of low areas. Spilled material may cause a slipping hazard. Keep upwind of spill. Ventilate area of leak or spill. If available, use foam to smother or suppress. Refer to section 7, Handling, for additional precautionary measures. See Section 10 for more specific information. Use appropriate safety equipment. For additional information, refer to Section 8, Exposure Controls and Personal Protection.

Environmental precautions: Prevent from entering into soil, ditches, sewers, waterways and/or groundwater. See Section 12, Ecological Information.

Methods and materials for containment and cleaning up: Contain spilled material if possible. Absorb with materials such as: Dirt. Vermiculite. Sand. Clay. Do NOT use absorbent materials such as: Cement powder (Note: may generate heat). Collect in suitable and properly labeled open containers. Do not place in sealed containers. Suitable containers include: Metal drums. Plastic drums. Polylined fiber pacs. Wash the spill site with large quantities of water. Attempt to neutralize by adding suitable decontaminant solution: Formulation 1: sodium carbonate 5 - 10%; liquid detergent 0.2 - 2%; water to make up to 100%, OR Formulation 2: concentrated ammonia solution 3 - 8%; liquid detergent 0.2 - 2%; water to make up to 100%. If ammonia is used, use good ventilation to prevent vapor exposure. Contact your supplier for clean-up assistance. See Section 13, Disposal Considerations, for additional information.

7. HANDLING AND STORAGE

Precautions for safe handling: Do not breathe vapors, mist or gas. Avoid contact with eyes, skin, and clothing. Avoid prolonged or repeated contact with skin. Wash thoroughly after handling. Use with adequate ventilation. This material is hygroscopic in nature. Keep container tightly closed. See Section 8, EXPOSURE CONTROLS AND PERSONAL PROTECTION. Spills of these organic materials on hot fibrous insulations may lead to lowering of the autoignition temperatures possibly resulting in spontaneous combustion.

Conditions for safe storage: Store in a dry place. Protect from atmospheric moisture. Do not store product contaminated with water to prevent potential hazardous reaction. See Section 10 for more specific information.

Storage stability

Storage temperature: 18 - 40 °C (64 - 104 °F) Storage Period: 12 Month

8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Control parameters

If exposure limits exist, they are listed below. If no exposure limits are displayed, then no values are applicable.

Component	Regulation	Type of listing	Value

Toluene-diisocyanate, mixture of toluene-2,4-di- isocyanate and toluene-2,6- di-isocyanate	Dow IHG	TWA Inhalable fraction and vapor	0.005 ppm
	Further information: SKIN, Respiratory sensitizer	DSEN, RSEN: Absorbed via	Skin, Skin Sensitizer,
	Dow IHG	C Inhalable fraction and vapor	0.02 ppm
	Further information: SKIN, Respiratory sensitizer	DSEN, RSEN: Absorbed via	Skin, Skin Sensitizer,
	OSHA Z-1	С	0.14 mg/m3 0.02 ppm
	Further information: (b): Th determined from breathing-	e value in mg/m3 is approxim zone air samples.	ate.; [©] : Ceiling limit is to be
	ACGIH	TWA Inhalable	0.001 ppm
		fraction and vapor	
	pulm func: Pulmonary funct	Dermal Sensitization; RSEN ion; eye irr: Eye irritation; as nown relevance to humans;	sthma: Asthma; A3: Confirmed
	ACGIH	STEL Inhalable fraction and vapor	0.005 ppm
Dialogical accuration of	pulm func: Pulmonary funct animal carcinogen with unk absorption	Dermal Sensitization; RSEN	thma: Asthma; A3: Confirmed
Biological occupational exp	osure limits		

Components CAS-No. Control Biological Sampling Permissible Basis parameters specimen time concentration Toluene-diisocyanate, 26471-62toluene Urine End of 5 µg/g ACGIH mixture of toluene-2,4-5 diamine shift creatinine BEI di-isocyanate and toluene-2,6-di-

isocyanate

Exposure controls

Engineering controls: Use only with adequate ventilation. Local exhaust ventilation may be necessary for some operations. Provide general and/or local exhaust ventilation to control airborne levels below the exposure guidelines. Exhaust systems should be designed to move the air away from the source of vapor/aerosol generation and people working at this point. The odor and irritancy of this material are inadequate to warn of excessive exposure.

Individual protection measures

Eye/face protection: Use chemical goggles. If exposure causes eye discomfort, use a full-face respirator.

Skin protection

Hand protection: Use gloves chemically resistant to this material. Examples of preferred glove barrier materials include: Butyl rubber. Polyethylene. Chlorinated polyethylene. Ethyl vinyl alcohol laminate ("EVAL"). Examples of acceptable glove barrier materials include: Viton. Neoprene. Polyvinyl chloride ("PVC" or "vinyl"). Nitrile/butadiene rubber ("nitrile" or "NBR"). NOTICE: The selection of a specific glove for a particular application and duration of use in a workplace should also take into account all relevant workplace factors such as, but not limited to: Other chemicals which may be handled, physical requirements (cut/puncture protection, dexterity, thermal protection), potential body reactions to glove materials, as well as the instructions/specifications provided by the glove supplier.

Other protection: Use protective clothing chemically resistant to this material. Selection of specific items such as face shield, boots, apron, or full body suit will depend on the task. Remove contaminated clothing immediately, wash skin area with soap and water. Contaminated clothing should be disposed of properly or decontaminated and laundered before reuse. Items which cannot be decontaminated, such as shoes, belts and watchbands, should be removed and disposed of properly.

Respiratory protection: Atmospheric levels should be maintained below the exposure guideline. When atmospheric levels may exceed the exposure guideline, use an approved air-purifying respirator equipped with an organic vapor sorbent and a particle filter. For situations where the atmospheric levels may exceed the level for which an air-purifying respirator is effective, use a positive-pressure air-supplying respirator (air line or self-contained breathing apparatus). For emergency response or for situations where the atmospheric level is unknown, use an approved positive-pressure self-contained breathing apparatus or positive-pressure air line with auxiliary self-contained air supply. The following should be effective types of air-purifying respirators: Organic vapor cartridge with a particulate pre-filter.

9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance	
Physical state	Liquid.
Color	Colorless to yellow
Odor	Sharp
Odor Threshold	0.05 - 0.2 ppm <i>Based on Literature for TDI</i> . Odor is inadequate warning of excessive exposure.
рН	Not applicable
Melting point/range	9.5 - 10 °C(49.1 - 50 °F) EU Method A.1 (Melting / Freezing Temperature)
Freezing point	9.5 - 10 °C(49.1 - 50 °F) EU Method A.1 (Melting / Freezing Temperature)
Boiling point (760 mmHg)	252 - 254 °C(486 - 489 °F) at 1,013 hPa <i>EU Method A.2</i> <i>(Boiling Temperature)</i>
Flash point	closed cup 126 °C (259 °F) <i>ASTM D 93</i>
Evaporation Rate (Butyl Acetate = 1)	<1 Literature
Flammability (solid, gas)	Not Applicable
Lower explosion limit	0.9 % vol Literature (2,4 isomer of TDI)
Upper explosion limit	9.5 % vol Literature
Vapor Pressure	0.015 hPa at 20 °C (68 °F) <i>Calculated.</i> 0.01 mmHg at 20 °C (68 °F) <i>Literature</i>
Relative Vapor Density (air = 1)	6.0 Literature
Relative Density (water = 1)	1.22 at 20 °C(68 °F)/ 20 °C EU Method A.3 (Relative Density)
Water solubility	insoluble
Partition coefficient: n- octanol/water	No data available
Auto-ignition temperature	> 595 °C (> 1,103 °F) EC Method A15
Decomposition temperature	No test data available

Dynamic Viscosity
Kinematic Viscosity
Explosive properties
Oxidizing properties
Liquid Density
Molecular weight
Percent volatility

3 mPa.s at 25 °C (77 °F) *Literature* 2.221 mm2/s at 20 °C (68 °F) *Literature* Not explosive No 1.22 g/cm3 at 20 °C (68 °F) *Literature* No test data available No data available

NOTE: The physical data presented above are typical values and should not be construed as a specification.

10. STABILITY AND REACTIVITY

Reactivity: Diisocyanates react with many materials and the rate of reaction increases with temperature as well as increased contact; these reactions can become violent. Contact is increased by stirring or if the other material mixes with the diisocyanate. Diisocyanates are not soluble in water and sink to the bottom, but react slowly at the interface. The reaction forms carbon dioxide gas and a layer of solid polyurea. Reaction with water will generate carbon dioxide and heat.

Chemical stability: Stable under recommended storage conditions. See Storage, Section 7.

Possibility of hazardous reactions: Can occur. Exposure to elevated temperatures can cause product to decompose and generate gas. This can cause pressure build-up and/or rupturing of closed containers. Polymerization can be catalyzed by: Strong bases. Water.

Conditions to avoid: Exposure to elevated temperatures can cause product to decompose. Generation of gas during decomposition can cause pressure in closed systems. Pressure build-up can be rapid. Avoid moisture. Material reacts slowly with water, releasing carbon dioxide which can cause pressure buildup and rupture of closed containers. Elevated temperatures accelerate this reaction.

Incompatible materials: Avoid contact with: Acids. Alcohols. Amines. Water. Ammonia. Bases. Metal compounds. Moist air. Strong oxidizers. Diisocyanates react with many materials and the rate of reaction increases with temperature as well as increased contact; these reactions can become violent. Contact is increased by stirring or if the other material mixes with the diisocyanate. Diisocyanates are not soluble in water and sink to the bottom, but react slowly at the interface. The reaction forms carbon dioxide gas and a layer of solid polyurea. Reaction with water will generate carbon dioxide and heat. Avoid contact with metals such as: Aluminum. Zinc. Brass. Tin. Copper. Galvanized metals. Avoid contact with absorbent materials such as: Moist organic absorbents. Avoid unintended contact with polyols. The reaction of polyols and isocyanates generate heat.

Hazardous decomposition products: Decomposition products depend upon temperature, air supply and the presence of other materials. Gases are released during decomposition.

11. TOXICOLOGICAL INFORMATION

Toxicological information appears in this section when such data is available.

Acute toxicity

Acute oral toxicity

Low toxicity if swallowed. Small amounts swallowed incidentally as a result of normal handling operations are not likely to cause injury; however, swallowing larger amounts may cause injury. Swallowing may result in gastrointestinal irritation or ulceration.

LD50, Rat, female, 4,130 mg/kg

Acute dermal toxicity

Prolonged skin contact is unlikely to result in absorption of harmful amounts.

LD50, Rabbit, male and female, > 9,400 mg/kg

Acute inhalation toxicity

Easily attainable vapor concentrations may cause serious adverse effects, even death. Excessive exposure to TDI may cause severe irritation of the upper respiratory tract and lungs, fluid in the lungs, permanent decrease of lung function, neurologic disorders, cholinesterase depression and gastrointestinal distress.

LC50, Rat, 1 Hour, vapour, 0.48 mg/l LC50, Mouse, 6 Hour, dust/mist, 0.1 mg/l

Skin corrosion/irritation

Prolonged contact may cause severe skin irritation with local redness and discomfort.

Serious eye damage/eye irritation

May cause severe eye irritation. May cause moderate corneal injury. Vapor may cause eye irritation experienced as mild discomfort and redness.

Sensitization

Skin contact may cause an allergic skin reaction.

Animal studies have shown that skin contact with isocyanates may play a role in respiratory sensitization.

May cause allergic respiratory reaction.

Reexposure to extremely low isocyanate concentrations may cause allergic respiratory reactions in individuals already sensitized.

Asthma-like symptoms may include coughing, difficult breathing and a feeling of tightness in the chest. Occasionally, breathing difficulties may be life threatening. Effects may be delayed.

Specific Target Organ Systemic Toxicity (Single Exposure)

May cause respiratory irritation. Route of Exposure: Inhalation Target Organs: Respiratory Tract

Specific Target Organ Systemic Toxicity (Repeated Exposure)

Based on available data, repeated exposures are not anticipated to cause additional significant adverse effects.

Carcinogenicity

An oral study in which high doses of TDI were reported to cause cancer in animals has been found to contain numerous deficiencies which compromise the validity of the study. TDI did not cause cancer in laboratory animals exposed by inhalation, the most likely route of exposure.

Teratogenicity

TDI did not cause birth defects in laboratory animals. Slight effects were observed in the fetus but only at doses which caused toxic effects to the mother.

Reproductive toxicity

In animal studies, TDI has been shown not to interfere with reproduction.

Mutagenicity

In vitro genetic toxicity studies were negative in some cases and positive in other cases. Animal genetic toxicity studies were negative. Results of a Drosophila study were reported to be weakly positive; however, these positive findings are believed to be due to degradation of TDI in the solvent delivery vehicle.

Aspiration Hazard

Consinensulation

Based on physical properties, not likely to be an aspiration hazard.

Carcinogenicity		
Component	List	Classification
Toluene-diisocyanate, mixture	IARC	Group 2B: Possibly carcinogenic to
of toluene-2,4-di-isocyanate		humans
and toluene-2,6-di-isocyanate		
	US NTP	Reasonably anticipated to be a human
		carcinogen
	ACGIH	A3: Confirmed animal carcinogen with
		unknown relevance to humans.

12. ECOLOGICAL INFORMATION

Ecotoxicological information appears in this section when such data is available.

Toxicity

Acute toxicity to fish

Material is slightly toxic to aquatic organisms on an acute basis (LC50/EC50 between 10 and 100 mg/L in the most sensitive species tested).

LC50, Oncorhynchus mykiss (rainbow trout), static test, 96 Hour, 133 mg/l, OECD Test Guideline 203 or Equivalent

Acute toxicity to aquatic invertebrates

EC50, Daphnia magna (Water flea), static test, 48 Hour, 12.5 mg/l, OECD Test Guideline 202 or Equivalent

Acute toxicity to algae/aquatic plants

EC50, Skeletonema costatum (marine diatom), static test, 96 Hour, 3,230 mg/l, OECD Test Guideline 201 or Equivalent

EC50, Chlorella vulgaris (Fresh water algae), static test, 96 Hour, 4,300 mg/l, OECD Test Guideline 201 or Equivalent

Toxicity to bacteria

EC50, activated sludge, Respiration inhibition, 3 Hour, > 100 mg/l, OECD 209 Test

Long-term (chronic) aquatic hazard Chronic toxicity to aquatic invertebrates NOEC, Daphnia magna (Water flea), static test, 21 d, number of offspring, 1.1 mg/l

LOEC, Daphnia magna (Water flea), static test, 21 d, number of offspring, 2.2 mg/l

Persistence and degradability

Biodegradability: In the aquatic and terrestrial environment, material reacts with water forming predominantly insoluble polyureas which appear to be stable. In the atmospheric environment, material is expected to have a short tropospheric half-life, based on calculations and by analogy with related diisocyanates.

Bioaccumulative potential

Bioaccumulation: No data available.

Mobility in soil

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In the aquatic and terrestrial environment, movement is expected to be limited by its reaction with water forming predominantly insoluble polyureas.

13. DISPOSAL CONSIDERATIONS

Disposal methods: DO NOT DUMP INTO ANY SEWERS, ON THE GROUND, OR INTO ANY BODY OF WATER. All disposal practices must be in compliance with all Federal, State/Provincial and local laws and regulations. Regulations may vary in different locations. Waste characterizations and compliance with applicable laws are the responsibility solely of the waste generator. AS YOUR SUPPLIER, WE HAVE NO CONTROL OVER THE MANAGEMENT PRACTICES OR MANUFACTURING PROCESSES OF PARTIES HANDLING OR USING THIS MATERIAL. THE INFORMATION PRESENTED HERE PERTAINS ONLY TO THE PRODUCT AS SHIPPED IN ITS INTENDED CONDITION AS DESCRIBED IN MSDS SECTION: Composition Information. FOR UNUSED & UNCONTAMINATED PRODUCT, the preferred options include sending to a licensed, permitted: Recycler. Reclaimer. Incinerator or other thermal destruction device. For additional information, refer to: Handling & Storage Information, MSDS Section 7 Stability & Reactivity Information, MSDS Section10 Regulatory Information, MSDS Section 15

14. TRANSPORT INFORMATION

DOT

Proper shipping name	Toluene diisocyanate
UN number	UN 2078
Class	6.1
Packing group	
Reportable Quantity	Toluene diisocyanate

Classification for SEA transport (IMO-IMDG):

Proper shipping name	TOLUENE DIISOCYANATE
UN number	UN 2078
Class	6.1
Packing group	II

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Marine pollutant Transport in bulk according to Annex I or II of MARPOL 73/78 and the IBC or IGC Code	No Consult IMO regulations before transporting ocean bulk
Classification for AIR transport (I <i>A</i>	TA/ICAO):
Proper shipping name	Toluene diisocyanate

Proper shipping name	Toluene diisocyan
UN number	UN 2078
Class	6.1
Packing group][

This information is not intended to convey all specific regulatory or operational requirements/information relating to this product. Transportation classifications may vary by container volume and may be influenced by regional or country variations in regulations. Additional transportation system information can be obtained through an authorized sales or customer service representative. It is the responsibility of the transporting organization to follow all applicable laws, regulations and rules relating to the transportation of the material.

15. REGULATORY INFORMATION

Superfund Amendments and Reauthorization Act of 1986 Title III (Emergency Planning and Community Right-to-Know Act of 1986) Sections 311 and 312 Acute toxicity (any route of exposure) Skin corrosion or irritation Serious eye damage or eye irritation Respiratory or skin sensitisation Carcinogenicity Specific target organ toxicity (single or repeated exposure)

Superfund Amendments and Reauthorization Act of 1986 Title III (Emergency Planning and Community Right-to-Know Act of 1986) Section 313

This product contains the following substances which are subject to the reporting requirements of Section 313 of Title III of the Superfund Amendments and Reauthorization Act of 1986 and which are listed in 40 CFR 372.

Componenta

components	CASRN
2,4-Toluene diisocyanate	584-84-9
2,6-Toluene diisocyanate	91-08-7

Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA) Section 103

Components	CASRN	RQ (RCRA Code)
Toluene-diisocyanate, mixture of toluene-2,4-	26471-62-5	100 lbs RQ
di-isocyanate and toluene-2,6-di-isocyanate		

Pennsylvania Worker and Community Right-To-Know Act:

To the best of our knowledge, this product does not contain chemicals at levels which require reporting under this statute.

California Prop. 65

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WARNING: This product can expose you to chemicals including 2,4-Toluene diisocyanate, 2,6-Toluene diisocyanate, which is/are known to the State of California to cause cancer. For more information go to www.P65Warnings.ca.gov.

United States TSCA Inventory (TSCA)

All components of this product are in compliance with the inventory listing requirements of the U.S. Toxic Substances Control Act (TSCA) Chemical Substance Inventory.

16. OTHER INFORMATION

Product Literature

Additional information on this product may be obtained by calling your sales or customer service contact.

Revision

Identification Number: 148691 / A001 / Issue Date: 07/26/2019 / Version: 15.0 Most recent revision(s) are noted by the bold, double bars in left-hand margin throughout this document.

Legend

USA. ACGIH Threshold Limit Values (TLV)	
Ceiling limit	
Dow Industrial Hygiene Guideline	
USA. Occupational Exposure Limits (OSHA) - Table Z-1 Limits for Air Contaminants	
Short-term exposure limit	
8-hour, time-weighted average	
	Dow Industrial Hygiene Guideline USA. Occupational Exposure Limits (OSHA) - Table Z-1 Limits for Air Contaminants Short-term exposure limit

Full text of other abbreviations

AICS - Australian Inventory of Chemical Substances; ASTM - American Society for the Testing of Materials; bw - Body weight; CERCLA - Comprehensive Environmental Response, Compensation, and Liability Act; CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DOT - Department of Transportation; DSL - Domestic Substances List (Canada); ECx - Concentration associated with x% response; EHS - Extremely Hazardous Substance; ELx - Loading rate associated with x% response; EmS - Emergency Schedule; ENCS - Existing and New Chemical Substances (Japan); ErCx - Concentration associated with x% growth rate response; ERG - Emergency Response Guide; GHS - Globally Harmonized System; GLP - Good Laboratory Practice; HMIS - Hazardous Materials Identification System; IARC - International Agency for Research on Cancer; IATA - International Air Transport Association; IBC - International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk; IC50 - Half maximal inhibitory concentration; ICAO - International Civil Aviation Organization; IECSC - Inventory of Existing Chemical Substances in China; IMDG - International Maritime Dangerous Goods; IMO -International Maritime Organization; ISHL - Industrial Safety and Health Law (Japan); ISO -International Organisation for Standardization; KECI - Korea Existing Chemicals Inventory; LC50 -Lethal Concentration to 50 % of a test population; LD50 - Lethal Dose to 50% of a test population (Median Lethal Dose); MARPOL - International Convention for the Prevention of Pollution from Ships; MSHA - Mine Safety and Health Administration; n.o.s. - Not Otherwise Specified; NFPA - National Fire Protection Association; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No

Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NTP - National Toxicology Program; NZIoC - New Zealand Inventory of Chemicals; OECD - Organization for Economic Co-operation and Development; OPPTS - Office of Chemical Safety and Pollution Prevention; PBT - Persistent, Bioaccumulative and Toxic substance; PICCS - Philippines Inventory of Chemicals and Chemical Substances; (Q)SAR - (Quantitative) Structure Activity Relationship; RCRA -Resource Conservation and Recovery Act; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals; RQ - Reportable Quantity; SADT - Self-Accelerating Decomposition Temperature; SARA -Superfund Amendments and Reauthorization Act; SDS - Safety Data Sheet; TCSI - Taiwan Chemical Substance Inventory; TSCA - Toxic Substances Control Act (United States); UN - United Nations; UNRTDG - United Nations Recommendations on the Transport of Dangerous Goods; vPvB - Very Persistent and Very Bioaccumulative

Information Source and References

This SDS is prepared by Product Regulatory Services and Hazard Communications Groups from information supplied by internal references within our company.

THE DOW CHEMICAL COMPANY urges each customer or recipient of this (M)SDS to study it carefully and consult appropriate expertise, as necessary or appropriate, to become aware of and understand the data contained in this (M)SDS and any hazards associated with the product. The information herein is provided in good faith and believed to be accurate as of the effective date shown above. However, no warranty, express or implied, is given. Regulatory requirements are subject to change and may differ between various locations. It is the buyer's/user's responsibility to ensure that his activities comply with all federal, state, provincial or local laws. The information presented here pertains only to the product as shipped. Since conditions for use of the product are not under the control of the manufacturer, it is the buyer's/user's duty to determine the conditions necessary for the safe use of this product. Due to the proliferation of sources for information such as manufacturer-specific (M)SDSs, we are not and cannot be responsible for (M)SDS obtained from any source other than ourselves. If you have obtained an (M)SDS from another source or if you are not sure that the (M)SDS you have is current, please contact us for the most current version.

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Version 1.2	Revision Date: 10/27/2016	SDS Number: 400001000139	Date of last issue: 05/20/2016 Date of first issue: 05/19/2016	
SECTION	1. IDENTIFICATION		-	
Produ	ct name	: RUBINATE®	7304	
Manu	facturer or supplier'	s details		
Comp Addre	any name of supplier ss	 Huntsman Polyurethanes P.O. Box 4980 The Woodlands, TX 77387 United States of America 		
Telepl	none	: Tech Info:(800		

: MSDS@huntsman.com

Emergency telephone number : Chemtrec: (800) 424-9300 or (703) 527-3887

Recommended use of the chemical and restrictions on use

Recommended use	:	Component of a Polyurethane System.
Restrictions on use	:	For industrial use only.

SECTION 2. HAZARDS IDENTIFICATION

E-mail address of person

responsible for the SDS

GHS Classification Acute toxicity (Inhalation)	: Category 4
Skin irritation	: Category 2
Eye irritation	: Category 2B
Respiratory sensitisation	: Category 1
Skin sensitisation	: Category 1
Specific target organ toxicity - single exposure	: Category 3 (Respiratory system)
GHS label elements Hazard pictograms	

Signal word

: Danger

Hazard statements

: H315 + H320 Causes skin and eye irritation. H317 May cause an allergic skin reaction.

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/ersion 1.2	Revision Date: 10/27/2016	SDS Number: 400001000139	Date of last issue: 05/20/2016 Date of first issue: 05/19/2016
		difficulties if inh	se allergy or asthma symptoms or breathing
Preca	autionary statements	P264 Wash ski P271 Use only P272 Contamir the workplace. P280 Wear pro P285 In case o protection. Response: P302 + P352 IF P304 + P340 + and keep comff CENTER/docto P305 + P351 + for several minit to do. Continue P333 + P313 If attention. P337 + P313 If attention. P342 + P311 If POISON CENT P362 Take off o Storage: P403 + P233 S tightly closed. P405 Store locl Disposal:	f inadequate ventilation wear respiratory F ON SKIN: Wash with plenty of soap and wate P312 IF INHALED: Remove person to fresh ai ortable for breathing. Call a POISON or if you feel unwell. P338 IF IN EYES: Rinse cautiously with water utes. Remove contact lenses, if present and ea e rinsing. skin irritation or rash occurs: Get medical advice eye irritation persists: Get medical advice/ experiencing respiratory symptoms: Call a 'ER/doctor. contaminated clothing and wash before reuse. tore in a well-ventilated place. Keep container

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

Substance / Mixture : Mixture

Hazardous components

Chemical name	CAS-No.	Concentration (% w/w)
4,4'-methylenediphenyl diisocyanate	101-68-8	60 - 100
Diphenylmethanediisocyanate	9016-87-9	13 - 30
Diphenylmethane-2,4'- diisocyanate	5873-54-1	13 - 30

The specific chemical identity and/or exact percentage (concentration) of composition has been withheld as a trade secret.

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SECTION 4. FIRST AID MEASURES

and effects, both acute and

General advice	:	Move out of dangerous area. Do not leave the victim unattended. Consult a physician. Show this safety data sheet to the doctor in attendance.
If inhaled	:	If breathed in, move person into fresh air. Call a physician or poison control centre immediately. Keep patient warm and at rest. Keep respiratory tract clear. If breathing is difficult, give oxygen. If breathing is irregular or stopped, administer artificial respiration. If unconscious place in recovery position and seek medical advice. Consult a physician immediately if symptoms such as shortness of breath or asthma are observed. A hyper-reactive response to even minimal concentrations of diisocyanates may develop in sensitised persons. LC50 (rat) : ca. 490 mg/m ³ (4 hours) : using experimentally produced respirable aerosol having aerodynamic diameter <5microns.
In case of skin contact	:	In case of contact, immediately flush skin with soap and plenty of water. Take off contaminated clothing and shoes immediately. Wash contaminated clothing before reuse. Thoroughly clean shoes before reuse. Call a physician if irritation develops or persists. An MDI study has demonstrated that a polyglycol-based skin cleanser (such as D-TamTM, PEG-400) or corn oil may be more effective than soap and water.
In case of eye contact	:	In case of eye contact, remove contact lens and rinse immediately with plenty of water, also under the eyelids, for at least 15 minutes. Protect unharmed eye. Keep eye wide open while rinsing. If eye irritation persists, consult a specialist.
If swallowed	:	Gently wipe or rinse the inside of the mouth with water. DO NOT induce vomiting unless directed to do so by a physician or poison control center. Keep respiratory tract clear. Keep at rest. Do not give milk or alcoholic beverages. If a person vomits when lying on his back, place him in the recovery position. Never give anything by mouth to an unconscious person. If symptoms persist, call a physician. Take victim immediately to hospital.
Most important symptoms	:	None known.

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delay	ved		
Prote	ection of first-aiders	suitable training. It may be danger mouth-to-mouth If potential for ex personal protecti First Aid respond	posure exists refer to Section 8 for specific
Note	s to physician		d supportive therapy as needed. Following medical follow-up should be monitored for at
			edure should be established in consultation esponsible for industrial medicine.

SECTION 5. FIREFIGHTING MEASURES

Suitable extinguishing media	;	Use extinguishing measures that are appropriate to local circumstances and the surrounding environment. Foam Carbon dioxide (CO2) Dry powder
Unsuitable extinguishing media	:	High volume water jet
Specific hazards during firefighting	:	Do not allow run-off from fire fighting to enter drains or water courses. The pressure in sealed containers can increase under the influence of heat. Exposure to decomposition products may be a hazard to health.
Hazardous combustion products	:	Carbon monoxide, carbon dioxide and unburned hydrocarbons (smoke). Nitrogen oxides (NOx) Hydrogen cyanide (hydrocyanic acid)
Specific extinguishing methods	:	Cool containers/tanks with water spray.
Further information	:	Standard procedure for chemical fires. Due to reaction with water producing CO2-gas, a hazardous build-up of pressure could result if contaminated containers are re-sealed. Use extinguishing measures that are appropriate to local circumstances and the surrounding environment. Collect contaminated fire extinguishing water separately. This must not be discharged into drains. Prevent fire extinguishing water from contaminating surface

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/ersion 1.2	Revision Date: 10/27/2016	SDS Number: 40000100013	Date of last issue: 05/20/2016 Date of first issue: 05/19/2016
		Fire residu	e ground water system. es and contaminated fire extinguishing water must ed of in accordance with local regulations.
	ial protective equipment efighters		pproved positive pressure self-contained breathing in addition to standard fire fighting gear.
SECTION	6. ACCIDENTAL RELE	ASE MEASUR	ES
prote	onal precautions, ctive equipment and gency procedures	Immediate Ensure add Keep peop Only qualif equipment Never retu Treat recov considerat For dispos Make sure absorbent The dange	nal protective equipment. ly evacuate personnel to safe areas. equate ventilation. ble away from and upwind of spill/leak. ied personnel equipped with suitable protective may intervene. rn spills in original containers for re-use. vered material as described in the section "Disposa ions". al considerations see section 13. that there is a sufficient amount of neutralizing/ material near the storage area. r areas must be delimited and identified using arning and safety signs.
Envir	onmental precautions	environme Do not allo Prevent pro Prevent fur Local autho cannot be o If the produ	w material to contaminate ground water system. oduct from entering drains. ther leakage or spillage if safe to do so. orities should be advised if significant spillages
	ods and materials for inment and cleaning up	Dilute with Contain sp material, (e and transfe national rea Clean cont Sweep up container fo Neutralize The compo Section 16. Remove ar Clean-up m If the produ Spilled MD The area s dust particl If the produ Soak up wi	nethods - small spillage plenty of water. illage, soak up with non-combustible absorbent a.g. sand, earth, diatomaceous earth, vermiculite) er to a container for disposal according to local / gulations (see section 13). aminated surface thoroughly. or vacuum up spillage and collect in suitable or disposal. small spillages with decontaminant. ositions of liquid decontaminants are given in d dispose of residues. nethods - large spillage uct is in its solid form: I flakes should be picked up carefully. hould be vacuum cleaned to remove remaining es completely. uct is in its liquid form: th inert absorbent material (e.g. sand, silica gel, universal binder_sawdust)

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/ersion 1.2	Revision Date: 10/27/2016	SDS Number: 400001000139	Date of last issue: 05/20/2016 Date of first issue: 05/19/2016
		Shovel into o Wash the sp Test atmosp	ct for at least 30 minutes. open-top drums for further decontamination. illage area with water. here for MDI vapour. able, closed containers for disposal.
SECTION	7. HANDLING AND ST	ORAGE	
Techn	ical measures	: Ensure that the workstat	eyewash stations and safety showers are close to ion location.
Local/	Total ventilation	: Use only wit	h adequate ventilation.
	e on protection against d explosion	: Normal mea	sures for preventive fire protection.
Advice	e on safe handling	Avoid forma Do not breat Avoid expos Avoid contac Smoking, ea application a Provide suffi Open drum o Dispose of r regulations. Persons sus allergies, chi	I protection see section 8. tion of aerosol. he vapours/dust. ure - obtain special instructions before use. et with skin and eyes. ting and drinking should be prohibited in the irea. cient air exchange and/or exhaust in work rooms. carefully as content may be under pressure. nse water in accordance with local and national ceptible to skin sensitisation problems or asthma, ronic or recurrent respiratory disease should not d in any process in which this mixture is being
Condi	tions for safe storage	Observe label Electrical inst technological Keep containe Observe label Electrical inst	allations / working materials must comply with the
Recon	nmended storage	: 20 - 35 °C	safety standards.
tempe		: 6 Months	

SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Components with workplace control parameters

Components	CAS-No.	Value type	Control	Basis
		(Form of	parameters /	
		exposure)	Permissible	
			concentration	

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	sion Date: 7/2016	SDS Number: 400001000139		[*] last issue: 05/20/201 [*] first issue: 05/19/201	
4,4'-methylen diisocyanate	ediphenyl	101-68-8	TWA	0.005 ppm	ACGIH
			C	0.02 ppm 0.2 mg/m3	OSHA Z-
Personal pro	tective equip	oment			
Respiratory protection: Use a properly fitted, air-purifying or air-fed respirator complying with an approved standard if a risk assessme indicates this is necessary. Respirator selection must be based on known or anticipa exposure levels, the hazards of the product and the safe working limits of the selected respirator.					ssessment or anticipated
Hand protecti Remarks	on	Protective g made polyu	loves should b rethane produc	l contact use protectiv be worn when handlin cts to avoid contact w nay be hazardous in c	ig freshly vith trace
		EN374: prof microorgani provide suit polyethylene laminated (" Nitrile/butad	ective gloves a sms. Example able protection e, Polyethylene EVAL"), Polyc iene rubber ("r	oves classified under against chemicals an s of glove materials t n include: Butyl rubbe e, Ethyl vinyl alcohol hloroprene (Neopren nitrile" or "NBR"), Pol lastomer (Viton*).	d hat might r, Chlorinated copolymers e*),
		glove with p	rotection class 240 minutes	ently repeated contac of 5 or higher (breal according to EN374)	kthrough time
		class of 3 or minutes acc	higher (break ording to EN3 ed gloves shoเ	expected, a glove wi through time greater 74) is recommended. uld be decontaminate	than 60
		application a take into ac not limited to requirement	and duration o count all requis o : other chem s (cut/puncture as well as inst	specific glove for a p f use in a workplace s site workplace factors icals that may be har e protection, dexterity ructions/specification	should also s such as, but ndled, physical v, thermal
Eye protection	ı	be used whe to avoid exp Chemical sp Always wea	en a risk asses osure to liquid blash goggles. r eye protectio	y with an approved sta ssment indicates this I splashes, mists or d on when the potential	is necessary usts. for inadvertent

eye contact with the product cannot be excluded. Please follow all applicable local/national requirements when

RUBINATE® 7304

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			ctive measures for a specific workplace. ewash stations and safety showers are close ion location.		
Skin a	and body protection	Choose body p	: Impervious clothing Choose body protection according to the amount and concentration of the dangerous substance at the work place.		
Protective measures		gloves, safety g The type of pro to the concentr at the specific v Ensure that eye	ctive equipment comprising: suitable protective goggles and protective clothing betective equipment must be selected according ation and amount of the dangerous substance workplace. e flushing systems and safety showers are to the working place.		
Hygiene measures		practice. Wash face, har handling. Remove contar before entering When using do When using do Contaminated v workplace. Wash hands ar handling the pro	not eat or drink. not smoke. work clothing should not be allowed out of the nd face before breaks and immediately after		

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance	: liquid
Colour	: brown
Odour	: No data is available on the product itself.
Odour Threshold	: No data is available on the product itself.
рН	: No data is available on the product itself.
Boiling point	: > 300 °C No information available.
Flash point	: > 110 °CMethod: closed cup
Evaporation rate	: No data is available on the product itself.
Flammability (solid, gas)	: No data is available on the product itself.
Flammability (liquids)	: No data is available on the product itself.
Upper explosion limit	: No data is available on the product itself.
Lower explosion limit	: No data is available on the product itself.

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	Vapour	pressure	:	No dàta is availat	ple on the product itself.
	Relative	e vapour density	:	No data is availat	ole on the product itself.
	Relative	e density	:	1.2 (25 °C)	
	Density		:	1.21 g/cm3 (25 °(C)
	Solubilit Wate	ty(ies) er solubility	:	No data is availat	ble on the product itself.
	Solut	pility in other solvents	:	No data is availat	ble on the product itself.
	Partitior octanol/	n coefficient: n-	:	No data is availat	ble on the product itself.
		nition temperature	:	No data is availat	ble on the product itself.
	Therma	l decomposition	:	No data is availat	ble on the product itself.
		celerating osition temperature	:	No data is availat	ble on the product itself.
	Viscosit Visco	y sity, dynamic	:	18 - 32 mPa.s (25	5 °C)

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SECTION 10. STABILITY AND REACTIVITY

Reactivity Chemical stability Possibility of hazardous reactions	 Stable under recommended storage conditions. Stable under normal conditions. Reaction with water (moisture) produces CO2-gas. Exothermic reaction with materials containing active hydroge groups. The reaction becomes progressively more vigorous and can be violent at higher temperatures if the miscibility of the reaction partners is good or is supported by stirring or by the presence of solvents. MDI is insoluble with, and heavier than water and sinks to the bottom but reacts slowly at the interface. A solid water-insoluble layer of polyurea is formed at the interface by liberating carbon dioxide gas. No decomposition if stored and applied as directed. 	
Conditions to avoid	: Extremes of temperature and direct sunlight. Exposure to air or moisture over prolonged periods.	
Incompatible materials	: Acids Bases Amines Steam	
Hazardous decomposition	: Carbon dioxide (CO2), carbon monoxide (CO), oxides of nitrogen (NOx), dense black smoke	

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			nide (hydrocyanic acid) ces noxious and toxic fumes.
CTION	11. TOXICOLOGICA	L INFORMATION	
Inforn expos	•	of : No data is avai	lable on the product itself.
Acute	e toxicity		
	oonents:		
Acute	nethylenediphenyl diis oral yComponents	: LD50 (Rat, mal	e): > 10,000 mg/kg Test Guideline 401
Acute	nylmethanediisocyan oral yComponents	: LD50 (Rat, mal	e): > 10,000 mg/kg ' Test Guideline 401
Acute Produ	inhalation toxicity - ct	: Acute toxicity e Exposure time: Test atmosphe Method: Calcul	re: dust/mist
4,4'-m	ponents: hethylenediphenyl diis dermal toxicity	: LD50 (Rabbit, r	nale and female): > 9,400 mg/kg Test Guideline 402
	nylmethanediisocyan dermal toxicity	: LD50 (Rabbit, r	nale and female): > 9,400 mg/kg Test Guideline 402
	nylmethane-2,4'- diiso dermal toxicity	: LD50 (Rabbit, r	nale and female): > 9,400 mg/kg Test Guideline 402
	toxicity (other routes istration)	of : No data availab	le
Skin o	orrosion/irritation		
Comp	<u>onents:</u>		
Specie	ethylenediphenyl diis es: Rabbit d: OECD Test Guidel		

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Diphenylmethanediisocyanate: Species: Rabbit Assessment: Irritating to skin. Method: OECD Test Guideline 404 Result: Skin irritation

Diphenylmethane-2,4'- diisocyanate: Species: Rabbit Assessment: Irritant Method: OECD Test Guideline 404 Result: Irritating to skin.

Serious eye damage/eye irritation

Components:

4,4'-methylenediphenyl diisocyanate: Species: Rabbit Result: Mild eye irritation

Diphenylmethanediisocyanate: Species: Rabbit Result: Irritation to eyes, reversing within 7 days Assessment: Mild eye irritant Method: OECD Test Guideline 405

Diphenylmethane-2,4'- diisocyanate: Species: Humans Result: Irritation to eyes, reversing within 7 days Assessment: Mild eye irritant Method: OECD Test Guideline 405 Remarks: Mild eye irritation

Respiratory or skin sensitisation

Components:

4,4'-methylenediphenyl diisocyanate: Exposure routes: Skin Species: Mouse Method: OECD Test Guideline 429 Result: May cause sensitisation by skin contact.

Exposure routes: Respiratory Tract Species: Guinea pig Result: May cause sensitisation by inhalation.

Diphenylmethanediisocyanate: Exposure routes: Skin Species: Guinea pig Method: OECD Test Guideline 406 Result: May cause sensitisation by skin contact.

Exposure routes: Respiratory Tract Species: Rat Result: May cause sensitisation by inhalation.

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Sp As	posure routes: Skin pecies: Mouse sessment: May cause ser psult: Causes sensitisation		act.
Sp As	posure routes: Respirator pecies: Guinea pig sessment: May cause ser psult: Causes sensitisation	nsitisation by inhalatior	
4,4	omponents: 1'-methylenediphenyl diiso sessment:	•	isation by inhalation and skin contact.
	ohenylmethanediisocyana sessment:	May cause an alle	ergic skin reaction., May cause allergy or s or breathing difficulties if inhaled.
	ohenylmethane-2,4'- diisoo sessment:	cyanate: Mild eye irritation	
	erm cell mutagenicity emponents:		
	l'-methylenediphenyl diiso enotoxicity in vitro	: Concentration: 2 Metabolic activa	00 ug/plate tion: with and without metabolic activation e 67/548/EEC, Annex, B.13/14
	phenylmethanediisocyana notoxicity in vitro	: Concentration: 2 Metabolic activa	00 ug/plate tion: with and without metabolic activation e 67/548/EEC, Annex, B.13/14
	ohenylmethane-2,4'- diisoo notoxicity in vitro	: Metabolic activa	tion: with and without metabolic activation Fest Guideline 471
4,4	mponents: '-methylenediphenyl diiso notoxicity in vivo	: Application Rout Exposure time: 3 Dose: 118 mg/m	Weeks
	henylmethanediisocyanat notoxicity in vivo	: Application Rout	e: Inhalation ified due to inconclusive data.

Application Route: Inhalation



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1.1	بر ب	11 .1		() IN	UT	

Assessment

	ylmethane-2,4'- diiso oxicity in vivo	Result: negativ cyanate: : Application Ro	/m3 D Test Guideline 474
	-	: Application Ro	
		Exposure time Dose: 118 mg, Method: OECI Result: negativ	: 3 Weeks /m3 D Test Guideline 474
Diphen	onents: ylmethanediisocyana cell mutagenicity- ment		erial or mammalian cell cultures did not show acts.
Germ o Assess	cell mutagenicity- ment	: No data availa	ble
Carcin	ogenicity		
Compo	nents'		
Species Applica Exposu Dose: 1 Freque Method Result:	ethylenediphenyl diisc s: Rat, (male and fem ition Route: Inhalation irre time: 24 month(s) I mg/m ³ ncy of Treatment: 5 c I: OECD Test Guideli positive Organs: Lungs	nale) n łaily	
Species Applica Exposu Dose: 1 Freque Method	ylmethanediisocyana s: Rat, (male and fem tion Route: Inhalatior re time: 24 month(s) I mg/m ³ ncy of Treatment: 5 c I: OECD Test Guideli positive	nale) ı laily	
Species Applica Exposu Dose: 1 Frequen Method Result:	ylmethane-2,4'- diiso s: Rat, (male and fem tion Route: Inhalatior re time: 24 month(s) mg/m³ ncy of Treatment: 5 c : OECD Test Guideli positive Organs: Lungs	laily	

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ersion 2	Revision Date: 10/27/2016	SDS Number: 400001000139	Date of last issue: 05/20/2016 Date of first issue: 05/19/2016	
IARC	:		this product present at levels greater than or lentified as probable, possible or confirmed n by IARC.	
OSH	A	No component of this product present at levels greater thar equal to 0.1% is identified as a carcinogen or potential carcinogen by OSHA.		
NTP			this product present at levels greater than or lentified as a known or anticipated carcinogen	
Repro	oductive toxicity			
<u>Comp</u>	<u>oonents:</u>			
	nylmethanediisocyar s on fertility	: Species: Rat, n Application Rou Method: OECD	nale and female ite: Inhalation Test Guideline 414 gnificant adverse effects were reported	
Diphe	nylmethane-2,4'- diis	ocyanate:		
·		Species: Rat, fe Application Rou Method: OECD		
		Application Rou Method: OECD	nale and female ite: Inhalation Test Guideline 414 testing did not show any effects on fertility.	
	<u>ponents:</u> nethylenediphenyl diis	socyanate:		
Effect	s on foetal opment	: Species: Rat, fe Application Rou General Toxicit mg/m ³	ite: Inhalation y Maternal: No observed adverse effect level: Test Guideline 414	
Diphe	nylmethanediisocyan	ate:		
		Application Rou General Toxicit	y Maternal: 4 mg/m³ Test Guideline 414	
Diphe	nylmethane-2,4'- diis	ocyanate:		
	,,	Species: Rat, m Application Rou General Toxicit mg/m ³		

Method: OECD Test Guideline 414



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Result: No teratogenic effects

Components:

Diphenylmethanediisocyanate:Reproductive toxicity -AssessmentNo toxicity to reproductionNo evidence of adverse effects on sexual function and fertility,
or on development, based on animal experiments.

STOT - single exposure

Components:

4,4'-methylenediphenyl diisocyanate: Exposure routes: Inhalation Target Organs: Respiratory Tract Assessment: May cause respiratory irritation.

Diphenylmethanediisocyanate: Exposure routes: Inhalation Target Organs: Respiratory Tract Assessment: May cause respiratory irritation.

Diphenylmethane-2,4'- diisocyanate: Exposure routes: Inhalation Target Organs: Respiratory system Assessment: The substance or mixture is classified as specific target organ toxicant, single exposure, category 3 with respiratory tract irritation.

STOT - repeated exposure

No data available

Repeated dose toxicity

Components:

4,4'-methylenediphenyl diisocyanate: Species: Rat, male and female NOEC: 0.2 mg/m3 Exposure time: 2 yr Number of exposures: 5 d Method: OECD Test Guideline 453

Diphenylmethanediisocyanate: Species: Rat, male and female NOEC: 0.2 mg/m3 Test atmosphere: dust/mist Exposure time: 2 yr Number of exposures: 5 d Method: OECD Test Guideline 453

Diphenylmethane-2,4'- diisocvanate:

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Species: Rat, male and female NOEC: 0.2 mg/m3 Exposure time: 2 yr Number of exposures: 5 d Method: OECD Test Guideline 453

Components:

Diphenylmethane-2,4'- diisocyanate: Repeated dose toxicity - : Mild eye irritation Assessment

Aspiration toxicity

No data available

Experience with human exposure

General Information:	No data available
Inhalation:	No data available
Skin contact:	No data available
Eye contact:	No data available
Ingestion:	No data available

Toxicology, Metabolism, Distribution

No data available

Neurological effects

No data available

Further information

Ingestion: No data available

SECTION 12. ECOLOGICAL INFORMATION

Ecotoxicity

Components:

4,4'-methylenediphenyl diisocyanate: Toxicity to fish : LC50 (Brachydanio rerio (zebrafish)): > 1,000 mg/l Exposure time: 96 h Test Type: static test Method: OECD Test Guideline 203

Diphenvlmethanediisocvanate:

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ersion 2	Revision Date: 10/27/2016	SDS Num 40000100		Date of last issue: 05/20/2016 Date of first issue: 05/19/2016	
,Toxicity to fish		Expos Test 1 Test s	sure time: 9 Гуре: static substance:		
			> 1,000 mg sure time: 9		
	enylmethane-2,4'- diisc ity to fish	: LC50 Expos Test T Test s	sure time: 9 Type: static substance:		
4,4'-m Toxici	conents: nethylenediphenyl diiso ity to daphnia and othe ic invertebrates	r : EC50 Expos Test T Test s	sure time: 2 Type: static substance: 1		
Toxici	nylmethanediisocyana ity to daphnia and othe ic invertebrates	r : EC50 Expos Test T Test s	sure time: 2 ype: static substance: 1		
Toxici	nylmethane-2,4'- diiso ty to daphnia and othe ic invertebrates	r : EC50 Expos Test T Test s	ure time: 2 ype: static ubstance: l		
Diphe	<u>ponents:</u> nylmethanediisocyana ty to algae	: EC50 subsp Expos Test T Test s	icatus)): > ure time: 7 ype: static ubstance: I	2 h	
M-Fac toxicit	ctor (Acute aquatic y)	: No da	ta available	9	
Toxici toxicit	ty to fish (Chronic	: No dat	ta available	9	

Components:

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	Toxicity aquatic	hylenediphenyl diisocy to daphnia and other invertebrates toxicity)			tatic test resh water
	Toxicity aquatic	Imethanediisocyanate to daphnia and other invertebrates toxicity)	:	NOEC (Daphnia n Exposure time: 21 Test Type: semi-s Test substance: F Method: OECD Te	tatic test resh water
	Toxicity aquatic	Imethane-2,4'- diisocy to daphnia and other invertebrates toxicity)	rana :	ite: NOEC (Daphnia n Exposure time: 21 Test Type: semi-s Test substance: F Method: OECD Te	tatic test resh water
	M-Facto toxicity)	r (Chronic aquatic	:	No data available	
		<u>nents:</u> Imethanediisocyanate to bacteria	:	EC50 (activated s Exposure time: 3 l Test Type: static t Test substance: F Method: OECD Te	est resh water
		Imethane-2,4'- diisocy to bacteria			n est resh water
	Compo	<u>nents:</u>			
		hylenediphenyl diisocy to soil dwelling ns	yan: :	ate: NOEC (Eisenia fe Exposure time: 33 Method: OECD Te	
		Imethanediisocyanate to soil dwelling ns		EC50 (Eisenia feti Exposure time: 33 Method: OECD Te	
		Imethane-2,4'- diisocy to soil dwelling ns	ana :	ite : NOEC (Eisenia fe Exposure time: 33 Method: OECD Te	

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Plant	toxicity	: No data avail	able
Sedin	nent toxicity	: No data avail	able
Toxic organ	ity to terrestrial isms	: No data avai	able
	xicology Assessment aquatic toxicity	: No data avai	able
Chror	nic aquatic toxicity	: No data avai	able
Toxic	ity Data on Soil	: No data avai	able
	organisms relevant to nvironment	: No data avail	able
	er information: Ita available		
<u>Com</u> 4,4'-m	s tence and degradabil <u>ponents:</u> nethylenediphenyl diisoc gradability	yanate: : Inoculum: Do Concentratio Result: Not b Biodegradatio Exposure tim	n: 30 mg/l iodegradable on: 0 %
	nylmethanediisocyanate gradability	: Inoculum: Do Concentratio Result: Not b Biodegradatio Exposure tim	iodegradable on: 0 %
	nylmethane-2,4'- diisoc gradability	: Inoculum: Do Concentratio Result: Not b Biodegradatio Exposure tim	n: 30 mg/l iodegradable on: 0 %
	emical Oxygen and (BOD)	: No data avail	able
Chem (COD	ical Oxygen Demand)	: No data avail	able

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BOD	/COD	: No data availab	ble
ThO	C	: No data availab	ble
BOD	/ThOD	: No data availab	ble
Disso (DOC	blved organic carbon	: No data availab	ble
	ico-chemical vability	: No data availab	ble
<u>Com</u>	ponents:		
	nethylenediphenyl diisoo lity in water	: Degradation ha	lf life(DT50): 20 hrs (25 °C) ormation available. n water
•	enylmethanediisocyanat lity in water	: Degradation ha	lf life(DT50): 0.8 d (25 °C) prmation available. n water
Photo	odegradation	: No data availab	le
Impa Treat	ct on Sewage ment	: No data availab	le
Bioad	ccumulative potential		
4,4'-n	ponents: nethylenediphenyl diisoc cumulation	: Species: Cyprin Bioconcentratio	nus carpio (Carp) n factor (BCF): 200 ccumulation is unlikely.
•	enylmethanediisocyanate cumulation	: Species: Cyprin Bioconcentratio	nus carpio (Carp) n factor (BCF): 200 ccumulation is unlikely.
	enylmethane-2,4'- diisoc cumulation	: Species: Cyprin Bioconcentratio	us carpio (Carp) n factor (BCF): 200 ccumulation is unlikely.
<u>Com</u>	<u>oonents:</u>		
Partiti	nethylenediphenyl diisoc on coefficient: n- ol/water	: log Pow: 4.51 (2 pH: 7	20 °C) Test Guideline 117
	enylmethane-2,4'- diisocy on coefficient: n-	anate : : log Pow: 4.51 (2	20 °C)

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octar	nol/water	pH: 7 Method: OECD	Test Guideline 117
Mob Mobi	ility in soil ility	: No data availat	ble
	ibution among onmental compartments	: No data availat	ble
Stab	ility in soil	: No data availat	le
	er adverse effects ronmental fate and ways	: No data availat	le
	ilts of PBT and vPvB ssment	: No data availat	le
Endo poter	ocrine disrupting ntial	: No data availat	le
	orbed organic bound gens (AOX)	: No data availab	le
Haza	rdous to the ozone laye	er	
Ozon	e-Depletion Potential	Protection of St Substances Remarks: This manufactured w	CFR Protection of Environment; Part 82 ratospheric Ozone - CAA Section 602 Class I product neither contains, nor was vith a Class I or Class II ODS as defined by the Act Section 602 (40 CFR 82, Subpt. A, App.A +
	ional ecological nation	: No data availab	le
Globa (GWI	al warming potential P)	: No data availab	le

SECTION 13. DISPOSAL CONSIDERATIONS

Disposal methods	
Waste from residues	 Do not dispose of waste into sewer. Do not contaminate ponds, waterways or ditches with chemical or used container. Send to a licensed waste management company.
Contaminated packaging	: Empty remaining contents. Dispose of as unused product. Do not re-use empty containers.



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SECTION 14. TRANSPORT INFORMATION

Date:

International Regulation

IATA

Not regulated as dangerous goods

IMDG

Not regulated as dangerous goods

Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code Not applicable for product as supplied.

National Regulations

Packing group

Labels

ERG Code Marine pollutant

DOT Classification	
UN/ID/NA number	: NA 3082
Proper shipping name	: OTHER REGULAT (Methylene Diphen
Class	: 9

OTHER REGULATED SUBSTANCES, LIQUID, N.O.S. (Methylene Diphenyl Diisocyanate)
9
III
CLASS 9
171
no

SECTION 15. REGULATORY INFORMATION

EPCRA - Emergency Planning and Community Right-to-Know Act

CERCLA Reportable Quantity

Components	CAS-No.	Component RQ (lbs)	Calculated product RQ (lbs)
4,4'-methylenediphenyl diisocyanate	101-68-8	5000	8308
chlorobenzene	108-90-7	100	*

*: Calculated RQ exceeds reasonably attainable upper limit.

SARA 311/312 Hazards	No SARA Hazards		
SARA 313	: The following components are subject to reporting level established by SARA Title III, Section 313:		rting levels
	l,4'-methylenedip liisocyanate	bhenyl 101-68-8	60.1825 %
	Diphenylmethane ate	diisocyan 9016-87-9	23.1 %

The following chemical(s) are listed as HAP under the U.S. Clean Air Act, Section 12 (40 CFR 61):

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		4,4'-methylenediphe diisocyanate	nyl 101-68-8	60.1825 %			
	Califo	ornia Prop. 65	•	es not contain any chemicals known to State cause cancer, birth defects, or any other rm.			
	The c	omponents of this prod	uct are reported ir	the following inventories:			
	CH IN	IV		contains substances listed on the Swiss e inventory, or in compliance with the			
	TSCA	ι.	: On the inventory	ν, or in compliance with the inventory			
	DSL AICS			All components of this product are on the Canadian DSL On the inventory, or in compliance with the inventory			
	NZIOC		: On the inventory	, or in compliance with the inventory			
	ENCS KECI	j		 ν, or in compliance with the inventory ν, or in compliance with the inventory 			
	PICCS		: On the inventory	, or in compliance with the inventory			
	IECS0 TCSI	5		 γ, or in compliance with the inventory γ, or in compliance with the inventory 			
			· · · · · · · · · · · · · · · · · · ·	, <u>1</u>			

Inventories

AICS (Australia), DSL (Canada), IECSC (China), REACH (European Union), ENCS (Japan); ISHL (Japan), KECI (Korea), NZIoC (New Zealand), PICCS (Philippines), TCSI (Taiwan), TSCA (USA)

TSCA - 5(a) Significant New Use Rule List of Chemicals

No substances are subject to a Significant New Use Rule.

US. Toxic Substances Control Act (TSCA) Section 12(b) Export Notification (40 CFR 707, Subpt D)

No substances are subject to TSCA 12(b) export notification requirements.



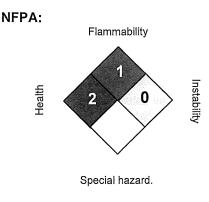
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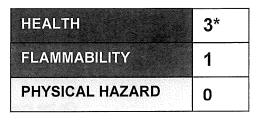
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SECTION 16. OTHER INFORMATION

Further information



HMIS III:



0 = not significant, 1 =Slight,



4 = Extreme, * = Chronic

Liquid decontaminants (percentages by weight or volume) :

Decontaminant 1 : *- sodium carbonate : 5 - 10 % *- liquid detergent : 0.2 - 2 % *- water : to make up to 100 %

Decontaminant 2 : *- concentrated ammonia solution : 3 - 8 % *- liquid detergent : 0.2 - 2 % *- water : to make up to 100 %

Decontaminant 1 reacts slower with diisocyanates but is more environmentally friendly than decontaminant 2.

Decontaminant 2 contains ammonia. Ammonia presents health hazards. (See supplier safety information.)

Revision Date

: 10/27/2016

The information and recommendations in this publication are to the best of our knowledge, information and belief accurate at the date of publication, NOTHING HEREIN IS TO BE CONSTRUED AS A WARRANTY, EXPRESS OR OTHERWISE.

IN ALL CASES, IT IS THE RESPONSIBILITY OF THE USER TO DETERMINE THE APPLICABILITY OF SUCH INFORMATION AND RECOMMENDATIONS AND THE SUITABILITY OF ANY PRODUCT FOR ITS OWN PARTICULAR PURPOSE.

THE PRODUCT MAY PRESENT HAZARDS AND SHOULD BE USED WITH CAUTION. WHILE CERTAIN HAZARDS ARE DESCRIBED IN THIS PUBLICATION, NO GUARANTEE IS MADE THAT THESE ARE THE ONLY HAZARDS THAT EXIST.

Hazards, toxicity and behaviour of the products may differ when used with other materials and are dependent upon the manufacturing circumstances or other processes. Such hazards, toxicity and behaviour should be determined by the user and made known to handlers, processors and end users.

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Section 1: Identification

1.1 Product identifier:

Diethanolamine LFG 85

Other means of identification:

01456

Chemical names: DEA LFG 85; Diethanolamine Low Freezing Grade 85

1.2 Recommended use:

Chemical raw material for industrial and professional uses.

Restrictions on use: Not for food or drug use. Keep out of reach of children.

1.3 Details of the supplier of the Safety Data Sheet:

INEOS Oxide Block 5501 21255 A Louisiana Hwy. 1 South Plaquemine, LA 70764 (866) 865-4767 www.ineosoxide.com

1.4 Emergency Phone No.

CHEMTREC 1-800-424-9300, 24-hours

Section 2: Hazard Identification

2.1 Classification:

Serious eye damage – Cat. 1; H318 Acute toxicity (Oral) – Cat. 4; H302 Skin irritation – Cat. 2; H315 Reproductive toxicity – Cat. 2; H361 Specific target organ toxicity (repeated exposure) – Cat. 2; H373

2.2 Label elements:



Danger. Causes serious eye damage. Harmful if swallowed. Causes skin irritation. Suspected of damaging fertility or the unborn child. May cause damage to organs (blood, kidneys, liver) through prolonged or repeated exposure if swallowed.

Prevention:

Obtain special instructions before use. Do not handle until all safety precautions have been read and understood. Wear eye protection, face protection, protective gloves and protective clothing. Do not breathe fume, mist, vapors or spray. Wash hands and exposed skin thoroughly after handling. Do not eat, drink or smoke when using this product.

Response:

IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. Immediately call a POISON CENTER or doctor.

IF SWALLOWED: Call a POISON CENTER or doctor if you feel unwell. Rinse mouth.

IF ON SKIN: Wash with plenty of water and soap.

If skin irritation occurs: Get medical attention. Take off contaminated clothing and wash it before reuse.

If exposed or concerned: Get medical advice.

Storage Store locked up.

Disposal:

Dispose of contents and container in accordance with local, regional, national and international regulations.

SAFETY DATA SHEET

2.3 Other hazards:

May be corrosive to copper and copper alloys (e.g. brass), some aluminum alloys, zinc, zinc alloys and galvanized surfaces.

Harmful to aquatic life with long lasting effects.

Section 3: Composition/Information on Ingredients

3.1 Substances:

Chemical Name	CAS No.	<u>EC #</u>	<u>Wt.%</u>	GHS Classification
Diethanolamine Other identifiers: DEA; 2,2'-iminodiethanol	111-42-2	203-868-0	85	Eye dam. 1; H318 Acute tox. 4; H302 Skin irrit. 2; H315 Repr. tox. 2; H361 STOT RE 2; H373 Aquatic chronic 3; H412
Water	7732-18-5	231-791-2	15	Not classified

Section 4: First-Aid Measures

4.1 First-aid measures:

Inhalation: Remove source of exposure or move to fresh air. Get medical advice or attention if you feel unwell or are concerned.

Eye Contact: Rinse cautiously with water for several minutes. Remove contact lenses if present and easy to do. Continue rinsing. Take care not to rinse contaminated water into the unaffected eye or onto the face. Immediately call a Poison Centre or doctor. Immediate treatment is required.

Skin Contact: Take off immediately contaminated clothing, shoes and leather goods (e.g. watchbands, belts). Immediately wash gently and thoroughly with lukewarm, gently flowing water and mild soap for 15-20 minutes. If skin irritation occurs, get medical advice or attention. Thoroughly clean clothing, shoes and leather goods before reuse or dispose of safely.

Ingestion: Immediately call a Poison Centre or doctor. Do not induce vomiting. If vomiting occurs naturally, lie on your side in the recovery position. Rinse mouth with water.

4.2 Most important symptoms and effects, acute and delayed:

See Section 11 of this SDS where additional symptoms and important health effects are described.

Inhalation: Symptoms of exposure may include coughing, wheezing, sore throat, chest pain, difficult breathing, nausea and vomiting.

Eye Contact: Causes severe irritation and eye damage. Serious damage, even blindness, may result if treatment is delayed. Symptoms include redness, pain and tearing. If DEA is rinsed out of the eye immediately, eye injury may be prevented.

Skin Contact: Causes severe skin irritation. Symptoms include local pain, redness and swelling. Prolonged contact may cause chemical burns, blister formation and possible tissue destruction.

Ingestion: Harmful if swallowed; oral toxicity in rats ranged from 680 to1820 mg/kg (LD₅₀). Swallowing can cause severe irritation and/or burns of the digestive tract. Symptoms include abdominal and chest pain, nausea, vomiting and diarrhea. Exposure by ingestion may cause reproductive toxicity.

Repeated exposure by ingestion may cause adverse effects to the kidney and liver and may cause anemia. Aspiration into the lungs during ingestion or vomiting may cause lung injury.

4.3 Immediate medical attention and special treatment:

If in eyes or if swallowed, call emergency medical services.

Section 5: Fire-fighting Measures

5.1 Suitable extinguishing media:

Water fog or fine spray, alcohol-resistant foam or dry chemical. Use water spray to cool fire-exposed containers.

Unsuitable extinguishing media: High pressure water streams may scatter hot liquid and may spread the fire. Violent steam generation or eruption may occur upon application of direct water stream to hot liquids.

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5.2 Specific hazards arising from the chemical:

Product can burn if heated or if involved in a fire [Flash point = 163°C (325°F)].

Heat from a fire can cause a rapid build-up of pressure inside containers, which may cause explosive rupture.

During a fire, smoke may contain vaporized DEA in addition to unidentified toxic and/or irritating compounds.

Thermal decomposition and combustion products may include toxic nitrogen oxide, hydrogen cyanide, formaldehyde carbon monoxide, carbon dioxide and ammonia gases.

5.3 Special protective equipment and precautions for fire-fighters:

Evacuate the area and fight fire from a safe distance or a protected location.

Thermal decomposition products such as nitrogen oxides and hydrogen cyanide are hazardous to health.

Do not enter without specialized protective equipment suitable for the situation.

Approach the fire from upwind to avoid hazardous vapors.

Burning liquids may be extinguished by dilution with water. Water spray may be used to flush spills away from ignition sources. Avoid all contact with this material during fire-fighting operations. Wear chemical resistant clothing (chemical splash suit) and positive-pressure self-contained breathing apparatus. Contain water run-off if possible.

Section 6: Accidental Release Measures

6.1 Personal precautions, protective equipment and emergency procedures:

Isolate the area; keep all unprotected people away from the spill area. Ventilate the area.

Extinguish or remove all ignition sources.

Prevent inhalation exposures, skin and possible eye contact.

Wear protective gloves, protective clothing and face protection (See Section 8).

Ensure clean-up is conducted by trained personnel only.

Do not touch or walk through the spilled material.

Spilled material may pose a slipping hazard.

6.2 Environmental precautions:

Prevent material from contaminating soil and from entering sewers or waterways.

6.3 Methods and material for containment and cleaning up:

Stop or reduce leak if safe to do so. Contain the spill with earth, sand or other suitable non-combustible absorbent. Keep materials which can burn away from spilled product. Do not absorb with sawdust, woodchips or other cellulose materials.

Clean up spills immediately.

Scoop up spilled product and any contaminated absorbents into appropriate, labeled containers. Contaminated absorbent may pose the same hazards as the spilled product. Flush the area with water and collect wash-water for proper disposal.

Section 7: Handling and Storage

7.1 Precautions for safe handling:

Do not eat, drink or smoke when using this product.

Wear eye/face protection and protective gloves and other equipment required for the workplace.

Wash hands and exposed skin after handling.

Do not breathe fume/mist/vapors/spray.

Avoid generating airborne fumes/vapors/mist from this product.

Handle this product with adequate ventilation.

Keep away from flames and hot surfaces.

Prevent handling with incompatible materials such as sodium nitrite, strong acids and oxidizing agents (see Section 10).

Prevent release of this material to the environment; prevent spills and keep away from drains.

Never perform any welding, cutting, soldering, drilling or other hot work on an empty vessel, container or piping until all liquid and vapors have been cleared.

Inspect containers for leaks before handling. Prevent damage to containers. Assume that empty containers contain residues which are hazardous.

Remove contaminated clothing promptly. Keep contaminated clothing in closed containers; discard or launder before rewearing. Maintain good housekeeping. Inform laundry personnel of contaminant's hazards. Do not take working clothes home.



7.2 Conditions for safe storage:

Keep containers tightly closed when not in use.

Store in a cool, dry and well-ventilated place. Store away from sunlight, heat and ignition sources.

Keep storage area away from work areas.

Store away from strong oxidants, strong acids and other incompatible materials (see Section 10).

Do not store in containers made of aluminum, copper, brass or other copper alloys, zinc, zinc alloys or galvanized steel. Store separated from food and feedstuffs.

Section 8: Exposure Controls / Personal Protection

8.1 Control parameters

Occupational Exposure Limits: Consult the local jurisdiction (e.g. province/territory) for their occupational exposure limits. Legislation for Canadian OH&S is available at : http://www.ccohs.ca/oshanswers/information/govt.html

Ingredient	ACGIH® TLV®	U.S. OSHA PEL	Other exposure limits
Diethanolamine	1 mg/m ³ Skin	15 mg/m ³ (3 ppm)	NIOSH REL: 3 ppm (15 mg/m ³)
(Inhalable fraction and vapor)		io ing/in (o ppin)	Quebec (Canada) VEMP: 3 ppm (13 mg/m ³)

8.2 Engineering controls

Engineering controls: Use only in a well-ventilated area. Use local exhaust ventilation for operations involving heating or spraying. Use local exhaust ventilation in workplaces where general ventilation is not adequate to control the amount in air. Ventilation system should be made of corrosion-resistant material.

If engineering controls and work practices are not effective in controlling exposure to this material or if adverse health symptoms are experienced, then wear suitable personal protection equipment including approved respiratory protection. Have appropriate equipment available for use in emergencies such as spills or fire.

8.3 Individual protection measures (PPE)

Eye/Face protection: Wear chemical safety goggles. Wear a face-shield or full-face respirator when needed to prevent exposure to liquid, vapour or fume.

Skin protection: Wear chemical protective gloves. Wear clean, body-covering, protective coveralls to prevent skin exposure. If spill or splashing is possible, wear chemical protective apron and boots.

Recommended materials for protective gloves and clothing include butyl rubber, neoprene rubber. Resistance of specific materials can vary from product to product; evaluate resistance under conditions of use and maintain clothing carefully.

Respiratory protection: If concentrations in air exceed the occupational exposure limits, then wear respiratory protection. Respiratory protection should not be necessary unless the product is heated to release vapours or a mist is created. If airborne vapour or mist exposure is likely wear a chemical cartridge respirator with cartridges to protect against ethanolamine, or a powered air-purifying respirator with cartridges to protect against ethanolamine, or a full-face self-contained breathing apparatus. For spills or uncontrolled releases, wear a supplied-air respirator.

If respiratory protection is required, institute a complete respiratory protection program including selection, fit testing, training, maintenance and inspection A respiratory protection program that meets the regulatory requirement, such as OSHA's 29 CFR 1910.134, ANSI Z88.2 or Canadian Standards Association (CSA) Standard Z94.4, must be followed whenever workplace conditions warrant a respirator's use.

Other protection: Workplaces should have a safety shower, hand-wash station and eye-wash fountain available.

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Section 9: Physical and Chemical Properties

asic physical and chemical properties:	
Appearance:	Liquid at temperatures above freezing point. Colorless, viscous.
	Solid at temperatures below freezing point.
Odor:	Ammonia-like, fish-like disagreeable odor.
Odor threshold:	0.27 ppm
pH:	11 (10% solution)
Melting point/freezing point:	27 °C (81 °F) (melting); -2 °C (28 °F) (freezing)
Initial boiling point and boiling range:	268°C (514°F)
Flash point:	163°C (325°F) PMCC; ASTM D93
Evaporation Rate:	< 0.01 (n-Butyl Acetate = 1)
Flammability (solid, gas):	Not applicable
Upper/lower flammability or explosive limits:	Not available
Vapor pressure:	0.000037 kPa at 25 °C (77 °F)
Vapor density:	3.7 (air=1)
Relative density:	1.09 (water = 1)
Solubility (ies):	95.4 g/L @ 20°C. Soluble in water
Partition coefficient (n-octanol/water, log Kow):	-2.46 25°C; OECD 107
Auto-ignition temperature:	> 662 °C (1224 °F)
Decomposition temperature:	200 °C (392 °F)
Viscosity:	190 - 270 mPa.s at 20 °C (dynamic)
Absolute density:	1090.7 kg/m ³ @ 25°C

Section 10: Stability and Reactivity

10.1 Reactivity:

Not reactive under normal conditions of use.

10.2 Chemical stability:

Unstable under certain conditions - see Conditions to Avoid.

Alkanolamine substances are decomposed by light and slowly oxidized by air, turning yellow and then brown. This reaction is accelerated by heat and the presence of metals.

Alkanolamine substances are oxidized by air slowly with evolution of heat. This reaction may lead to spontaneous combustion if the substance is on an adsorbent or on a high surface area material (e.g. absorbent material or thermal insulation).

10.3 Possibility of hazardous reactions:

Heating increases the risk of fire.

Heating above 60°C in aluminum can result in corrosion and generation of flammable hydrogen gas.

Reacts with cellulose nitrate causing fire and explosion hazard.

Reacts violently with strong acids and strong oxidants (e.g. nitric acid, hydrogen peroxide).

Contact with nitrosating agents, under acidic conditions such as nitrous acid, sodium nitrite or nitrogen oxides, can form nitrosamines some of which are potent carcinogens.

10.4 Conditions to avoid:

Avoid high temperatures and contact with sources of ignition. Avoid exposing product to air, light and moisture. Avoid direct sunlight.

10.5 Incompatible materials:

Contact with strong acids, strong oxidizing agents, halogenated hydrocarbons, nitrating agents may increase risk of vigorous or violent reaction.

Contact with alkali metals, metal hydrides and aluminum may generate flammable hydrogen gas.

Contact with Isocyanates and Isothiocyanates: reaction may be rapid, evolving heat.

Product may be corrosive to aluminum alloys at elevated temperatures, many 400 series stainless steel alloys, copper, zinc, and aluminum bronze.

In combination with water, the product may be corrosive to copper and copper alloys (e.g. brass), some aluminum alloys, zinc, zinc alloys, and galvanized surfaces.

Diethanolamine attacks some polymers including polyvinylchloride, polyurethane, polyamide imide, polyvinylidene fluoride and high-density polyethylene at elevated temperatures.



10.6 Hazardous decomposition products:

Decomposes at temperatures above 200°C; hazardous decomposition products may include nitrogen oxides, ammonia, hydrogen cyanide, formaldehyde. Hazardous decomposition products depend upon temperature, air supply and the presence of other materials.

Oxidation in air may form transient, organic peroxides or thermally unstable N-oxides such as hydroxylamines and carbamates form as well as nitrosamines, which are suspected cancer causing chemicals. Oxidation of Diethanolamine and decomposition of products is accelerated by light, heat, and/or presence of metals or metal oxides.

Section 11: Toxicological Information

11.1 Information on toxicological effects

Likely routes of exposure

Ingestion, Eye contact, Skin contact, Inhalation.

Acute toxicity

Inhalation: LC₅₀ of pure DEA was greater than 0.2 mg/L (greater than saturated vapor concentration and no mortalities). Inhalation test equivalent to OECD test guideline 403 and was for an 8 hour exposure time. Calculated 4-hour exposure LC₅₀ = 0.4 mg/L.

Based on information from animal tests, breathing vapors, spray or fume may cause nose and throat irritation, lung injury. Symptoms may include coughing, shortness of breath, difficult breathing and tightness in the chest. Symptoms may develop hours after exposure and are made worse by physical effort.

Ingestion: Harmful if swallowed. Acute toxicity estimate for the mixture is 1886 mg/kg (rat). Based on information from animal tests swallowing may cause liver and kidney damage, brain damage and blood changes.

Skin: Acute toxicity estimate for the mixture is 10,000 mg/kg (rabbit). Dermal exposures caused dose-related anemia and toxicity to the liver and kidneys in rats and mice.

11.2 Acute toxicity data:

Ingredient	LD ₅₀ Oral	LD ₅₀ Dermal	<u>LC₅₀ Inhalation</u> (4 hrs.)
Diethanolamine (DEA)	1600 mg/kg (rat)	8180 mg/kg (rabbit)	> 0.4 mg/L (rat)

Skin corrosion / irritation:

Human experience and animal studies caused moderate or severe skin irritation. Irritating to skin in an animal study according to OECD test guideline 404 (24, 48, 72 hours) in rabbit.

Serious eye damage / irritation:

Human experience and animal tests caused serious eye damage.

Highly irritating, causing serious eye damage in an animal study according to OECD test guideline 405 (24, 48, 72 hours) in rabbit.

STOT (Specific Target Organ Toxicity) Single Exposure:

Inhalation: Short-term inhalation of 858-6000 mg/m³ (cited as 200-1400 ppm) Diethanolamine aerosol or vapor caused breathing difficulties and some deaths in male rats. Continuous inhalation of 107 mg/m³ (cited as 25 ppm) for 216 hours (9 days) resulted in increased liver and kidney weights and altered kidney and liver function.

Ingestion: Ingestion of large quantities may cause liver and kidney damage, brain damage and blood changes. Skin and ingestion exposures have caused dose-related anemia and toxicity to the liver and kidneys in rats and mice.

Skin absorption: In male rats, a significant increase in relative kidney and liver weights occurred following a single oral dose of 200 mg/kg and higher. Signs of anemia were present in female rats ingesting 79 mg/kg/day and higher for 2 weeks and in male rats ingesting 162 mg/kg/day and higher for 2 weeks.

Aspiration hazard:

Due to the alkaline property of DEA, any aspiration during ingestion or vomiting could result in lung injury. Does not meet the criteria for classification in the class: Aspiration hazard.

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11.3 Chronic toxicity:

STOT (Specific Target Organ Toxicity) Repeated Exposure:

In tests with animals, long-term ingestion, inhalation and skin contact exposures to high doses caused anemia and damage to the liver and kidney.

A NOAEC for systemic effects in rats was 15 mg/m³, exposure by inhalation.

A NOAEC for upper respiratory tract irritation was 3 mg/m³.

A LOAEL for systemic effects by dermal application was 32 mg/kg bw/day in rats. Effects included dermal hyperkeratosis, anemia, liver toxicity and nephropathy.

In female rats, repeated oral doses, a LOAEL of 14 mg/kg bw/day was reported for anemia.

Respiratory and / or skin sensitization:

Not known to be a skin sensitizer. Not known to be a respiratory sensitizer.

Not sensitizing in a study conducted according to OECD guideline 406, in Guinea pig.

Germ cell mutagenicity:

Evidence from animal studies, cultured mammalian cells, and bacterial studies does not indicate that DEA is a mutagen. Negative results from in vitro tests according to OECD guidelines 471 (E. coli) and 476 (S. typhimurium). Negative results from in vivo tests in mice according to OECD guideline 474.

Reproductive effects:

Suspected of damaging fertility or the unborn child. For DEA, an extended One-Generation Reproductive Toxicity Study according to OECD TG 443 was performed. The NOAEL (no observed adverse effect level) for fertility and reproductive performance in P0 and F1 rats was 300 ppm, oral in drinking water based on lower number of implants and effects to reproductive organs in both males and females at the LOAEL (lowest observed adverse effect level) of 1000 ppm.

Developmental effects:

Suspected of damaging fertility or the unborn child. For DEA, an extended One-Generation Reproductive Toxicity Study according to OECD TG 443 was performed. The NOAEL for developmental toxicity in F1 rats was 100 ppm, oral in drinking water based on impaired pup survival at the LOAEL of 300 ppm.

Effects on or via lactation:

No information was located.

Carcinogenicity:

NTP Report: Under the conditions of 2 year dermal studies, there was no evidence of carcinogenic activity of DEA in F344/N rats administered 16, 32 or 64 mg/kg DEA or in female F344/N rats administered 8, 16 or 32 mg/kg. There was clear evidence of carcinogenic activity of DEA in male and female B6C3F1 mice based on increased incidences of liver neoplasms in males and females and increased incidences of renal tubule neoplasms in males. IARC Monograph Volume101 reports a mechanism for liver tumor induction in mice exposed to DEA that involves the inhibition of choline uptake in the liver. As humans are less susceptible to choline deficiency than rats or mice, the results may not be predictive of induction of cancer in humans.

IARC lists DEA in Group 2B – Possibly carcinogenic to humans. ACGIH designates DEA as A3 – confirmed animal carcinogen with unknown relevance to humans. DEA is not listed on the NTP Report On Carcinogens.

Interactions with other chemicals:

Diethanolamine may react chemically with nitrosating compounds under certain conditions to form nitrosamines, which are mutagenic and possibly carcinogenic.

Section 12: Ecological Information

12.1 Ecotoxicity:

Data for pure Diethanolamine:

 $\begin{array}{l} LC_{50} \hspace{0.2cm} fish = 1460 \hspace{0.2cm} mg/L \hspace{0.2cm} (\textit{Pimephales promelas; 96-hour; fresh water; static)} \\ EC_{50} \hspace{0.2cm} crustacea = 55 \hspace{0.2cm} mg/L \hspace{0.2cm} (\textit{Daphnia magna; 48-hour; fresh water; static)} \\ ErC_{50} \hspace{0.2cm} algae = 2.2 \hspace{0.2cm} mg/L \hspace{0.2cm} (\textit{Pseudokirchneriella subcapitata; 96-hour; fresh water; semi-static)} \\ NOEC \hspace{0.2cm} crustacea = 0.78 \hspace{0.2cm} mg/L \hspace{0.2cm} (\textit{Daphnia magna; 21-day; fresh water; semi-static)} \end{array}$

12.2 Persistence and degradability:

Degrades rapidly based on quantitative tests. Biodegradation in water: 93% in 28 days, test according to OECD 301F.

12.3 Bioaccumulative potential:

Low potential to bioaccumulate based on log Kow -2.46 @ 20°C, test data according to OECD 107.



12.4 Mobility in soil:

Volatilization of DEA from water is very slow (Henry's Law Constant (H) is 5.35E-14 atm m³/mol). Potential for mobility in soil is very high (Koc between 0 and 50). Log soil organic carbon partition coefficient (log Koc) is estimated to be 0.60.

12.5 Other information:

Not dangerous for the ozone layer (According to EU Council Regulation No 1005/2009) For detailed Ecological data, write to the address in Section 1 or email INEOS Oxide Technical Services at oxide.us.techservices@ineos.com.

Section 13: Disposal Considerations

13.1 Disposal methods

Do NOT discard into any sewers, on the ground or into any body of water. Store material for disposal as indicated in Section 7 Handling and Storage.

Dispose of in accordance with local/regional/national/ international regulations.

For unused, uncontaminated product, the preferred options include sending to a licensed, permitted recycler, reclaimer incinerator or other thermal destruction device.

USA: Under RCRA, it is the responsibility of the user of the product to determine, at the time of disposal, whether the product meets RCRA criteria for hazardous waste.

Other information:

Waste characterizations and compliance with applicable laws are the responsibility solely of the waste generator. INEOS Oxide HAS NO CONTROL OVER THE MANAGEMENT PRACTICES OR MANUFACTURING PROCESSES OF PARTIES HANDLING OR USING THIS MATERIAL. THE INFORMATION PRESENTED HERE PERTAINS ONLY TO THE PRODUCT AS SHIPPED IN ITS INTENDED CONDITION AS DESCRIBED IN SDS SECTION 2.

As a service to its customers, INEOS Oxide can provide names of information resources to help identify waste management companies and other facilities which recycle, reprocess or manage chemicals or plastics, and that manage used drums. Email INEOS Oxide Technical Services at oxide.us.techservices@ineos.com.

Section 14: Transport Information

14.1 U.S. Hazardous Materials Regulation (DOT 49CFR):

U.S. Bulk shipments exceeding >117 lbs (>53.4 kg) ship as: UN3082 ENVIRONMENTALLY HAZARDOUS SUBSTANCE, LIQUID, N.O.S. (DIETHANOLAMINE), Class 9, PG III

Reportable Quantity (RQ) for U.S. Shipments: 100 lbs (45.4 kg) for Diethanolamine

14.2 IMO classification:

Not regulated

Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code: Diethanolamine: Category Y

14.3 ICAO/IATA classification:

Not regulated

14.4 Canadian Transportation of Dangerous Goods (TDG):

Not regulated

For regulatory information regarding transportation, if required, consult product shipping papers, or your INEOS Oxide representative.

Section 15: Regulatory Information

15.1 Safety, health and environmental regulations:

USA:

OSHA: This material is considered a hazardous chemical by the OSHA Hazard Communication Standard 29 CFR 1910.1200 (2012).

Toxic Substances Control Act (TSCA) Section 8(b): All ingredients are listed on the TSCA Inventory.

Additional USA regulatory lists: Clean Air Act -Diethanolamine.

SARA Title III - Section 313. Diethanolamine

California Safe Drinking Water and Toxic Enforcement Act, Proposition 65: Diethanolamine (CAS No. 111-42-2) is on the list of chemicals known to the State to cause cancer.

State Right To Know - Diethanolamine: Massachusetts. New Jersey. Pennsylvania.

Canada:

DSL status: Listed on the DSL (Domestic Substances List).

NPRI Substances: Diethanolamine (and its salts) NPRI Part (Threshold Category): 1A, Reportable to NPRI if manufactured, processed, or otherwise used at quantities greater than: 10 tonnes.

European Union:

European Inventories: 2',2"-iminodiethanol listed in EINECS # 203-868-0.

Other international inventories

Australia: Present on the Inventory of Chemical Substances (AICS).

China: Present on the Chinese inventory (IECSC), 11481.

Japan: Present on ENCS ; Diethanolamine (2)-354; (2)-302.

Korea: Present on the Inventory of Existing and Evaluated Chemical Substances; Diethanolamine KE-20959.

Mexico: Present on the inventory (INSQ).

New Zealand: Present on the inventory (NZIoC) HSNO Approval: HSR002962.

Philippines: Present on the inventory (PICCS).

Taiwan: Present on the inventory (TCSI).

Thailand: Present on the inventory (TCSI / TECI) 55-1-00670.

Turkey: Present on the inventory. EC# 203-868-0

Vietnam: Present on the national chemicals inventory NCI 0623.

Section 16: Other Information

Revision date:

February 20, 2020

Revision summary:

Replaces previous version June 2018 with the following revisions: Section 1.2: Recommended use and restrictions on use Section 2.1, 2.2: new classification and labeling for reproductive toxicity hazard Section 3.1: new GHS classification for DEA Reproductive toxicity category 2; H361 Section 9: new data for Physical and Chemical properties

References and sources for data:

CCOHS – ChemInfo www.ccohs.ca/oshanswers/information/govt.html ECHA - Information on Chemicals, Registered Substances HSDB – Hazardous Substances Data Bank® IARC Monograph Volume101 National Toxicology Program (NTP) – Report on Carcinogens. NIOSH – National Institute for Occupational Safety and Health RTECS® - Registry of Toxic Effects of Chemical Substances

Additional information:

The information provided on this SDS is correct to the best of our knowledge, information and belief at the date of its publication. The information given is designed only as a guide for safe handling, use, processing, storage, transportation, disposal and release and is not to be considered as a warranty or quality specification. The information relates only to the specific material designated and may not be valid for such material used in combination with any other material or in any process, unless specified in the text.

Appendix D - Public Notice

PUBLIC NOTICE

Adient US LLC has applied to the Tennessee Department of Environment & Conservation, Division of Air Pollution Control (Division identification number 28-0076/980244) for approval to expand its polyurethane foam manufacturing process at the facility in Pulaski, Giles County, Tennessee. Adient US LLC proposes to increase production of the three polyurethane foam production lines. The project is subject to review under the State rule for Prevention of Significant Deterioration of Air Quality (PSD), Paragraph 1200-03-09-.01(4) of the Tennessee Air Pollution Control Regulations, which requires a public notification and 30-day public comment period.

The Division of Air Pollution Control has reviewed the application with respect to the above-mentioned PSD regulations and has determined that construction can be approved if certain conditions are met. A copy of the PSD application materials, a copy of the PSD preliminary determination, and a copy of the draft construction permit are available for public inspection during normal business hours at the following locations:

Columbia Environmental Field Office	Tennessee Department of Environment & Conservation
Division of Air Pollution Control	Division of Air Pollution Control
1421 Hampshire Pike	William R. Snodgrass Tennessee Tower
Columbia, TN 38401	312 Rosa L. Parks Avenue, 15th Floor
	Nashville, Tennessee 37243

Electronic copies of the draft permits are available by accessing the TDEC internet site located at:

https://www.tn.gov/environment/ppo-public-participation/ppo-public-participation/ppo-air.html

Questions concerning the source(s) may be addressed to Tracy Kefauver at (615) 532-0536 or by e-mail at <u>Tracy.Kefauver@tn.gov</u>.

Interested parties are invited to review these materials and comment. In addition, a public hearing may be requested at which written or oral presentations may be made. To be considered, written comments or requests for a public hearing must be received no later than 4:30 PM on **August 19, 2022.** To assure that written comments are received and addressed in a timely manner, written comments must be submitted using one of the following methods:

- Mail, private carrier, or hand delivery: Address written comments to Ms. Michelle W. Owenby, Director, Division of Air Pollution Control, William R. Snodgrass Tennessee Tower, 312 Rosa L. Parks Avenue 15th Floor, Nashville, Tennessee 37243.
- 2. E-mail: Submit electronic comments to <u>air.pollution.control@tn.gov</u>.

A final determination will be made after weighing all relevant comments.

Individuals with disabilities who wish to review information maintained at the above-mentioned depositories should contact the Tennessee Department of Environment and Conservation to discuss any auxiliary aids or services needed to facilitate such review. Such contact may be in person, by writing, telephone, or other means, and should be made no less than ten days prior to the end of the public comment period to allow time to provide such aid or services. Contact the Tennessee Department of Environment and Conservation ADA Coordinator, William R. Snodgrass Tennessee Tower, 312 Rosa L. Parks Avenue 22nd Floor, Nashville, TN 37243, 1-(866)-253-5827. Hearing impaired callers may use the Tennessee Relay Service, 1-(800)-848-0298.

Air Pollution Control

DATE: August 19, 2022

Assigned to - Tracy Kefauver

Appendix E – Correspondence

The following are EPA Region IV modeling correspondence (dated April 19, 2022) on the Adient Pulaski air quality analysis.

Sent: To: Cc: Subject:	Tuesday, April 19, 2022 1:43 PM Haidar Alrawi; Gillam, Rick; Shepherd, Lorinda
Ca	
Subject:	Paul LaRock; Lacey Hardin; Richard Smrz; Tracy Kefauver; Travis Blake; Lusky, Katy
500,000	[EXTERNAL] RE: Adient -Pulaski, TN - PSD Application
*** This is an EXTERNAL senders or unexpected e	. email. Please exercise caution. DO NOT open attachments or click links from unknown email - STS-Security. ***
Haidar,	
I hope you are doing we	20.
	e VOC-only air quality analysis for Adient-Pulaski to us for review. We have reviewed n Sections 6 and 7 of the application and we have no comments regarding the
-Chris	
Christopher M. Howard	
Regional Meteorologist	
US EPA Region 4 - Atlan	
404/562-9036	
Howard.chris@epa.gov	

From: Haidar Alrawi <Haidar.Alrawi@tn.gov> Sent: Wednesday, April 13, 2022 3:03 PM To: Howard, Chris <Howard.Chris@epa.gov>; Gillam, Rick <Gillam.Rick@epa.gov>; Rinck, Todd <Rinck.Todd@epa.gov>; Shepherd, Lorinda <Shepherd.Lorinda@epa.gov> Cc: Paul LaRock <Paul.LaRock@tn.gov>; lacey.hardin@tn.gov; Richard Smrz <Richard.Smrz@tn.gov>; Tracy Kefauver <Tracy.Kefauver@tn.gov>; Travis Blake <travis.blake@tn.gov> Subject: RE: Adient -Pulaski, TN - PSD Application

Good afternoon,

Attached, please find a PSD construction permit application that was received on 3/30 for the Adient facility in Pulaski, TN. This source manufactures cushion seats for the automotive industry and only VOC emissions are subject to PSD review at a potential-to-emit emission rate of about 500 tons per year. The modeling analyses are addressed in sections 6 and 7 of the application.

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Please let me know if you have any questions or need additional information.

The following are responses to questions from EPA Region IV (dated August 8, 2022) after review of Adient's construction permit application

Tracy Kefauver

From:	Kris Patrick Foster <kris.patrick.foster@adient.com></kris.patrick.foster@adient.com>
Sent:	Monday, August 8, 2022 5:24 PM
То:	Tracy Kefauver
Cc:	Tawanna Reid
Subject:	[EXTERNAL] RE: Adient PSD Application
Attachments:	220808_TDEC_PSD Construction Permit Application_EPA Region 4
	Comments_Response_final.pdf

Tracy,

Please see attached responses to the below EPA questions. If you have any further inquiries, please don't hesitate to contact me!

Thank you!



Kris P. Foster Environmental, Health and Safety Lead 1890 Mines Road Pulaski, TN 38478 Cell: 931-638-5918 Office: 931-424-7848

Adient – INTERNAL

From: Tracy Kefauver <Tracy.Kefauver@tn.gov>
Sent: Tuesday, August 2, 2022 2:39 PM
To: Kris Patrick Foster <kris.patrick.foster@adient.com>
Cc: Tawanna Reid <Tawanna.Reid@tn.gov>
Subject: FW: Adient PSD Application

CAUTION: This email originated from outside of the organization. <u>Do not click links or open attachments</u> unless you recognize the sender and know the content is safe.

Hi Kris,

Please see the bulleted items in the below email from EPA regarding Adient's PSD construction permit application. The first and last bullet items have already been included in the permit and/or preliminary determination (my comment in red below). Please respond to us regarding bullets 2-4. We will evaluate and include if necessary, in the preliminary determination and/or permit.

We plan to have a draft permit and preliminary determination document to you for your review by end of this week.

Also of note, we submitted this PSD construction application to EPA on April 14, 2022.

Please contact me with any questions.

Thanks,



Tracy Kefauver | TDEC-Environmental Protection Specialist 2 Division of Air Pollution Control William R. Snodgrass Tennessee Tower, 15th Floor 312 Rosa L. Parks Avenue, Nashville, TN 37243 p. 615-532-0536 Tracy.Kefauver@tn.gov https://www.tn.gov/environment/permit-permits/permit-air-home.html

We value your opinion. Please take a few minutes to <u>complete our customer service survey</u>. Internal Customers please complete our <u>customer satisfaction survey</u>.

From: Ferrando, Emily <Ferrando.Emily@epa.gov>
Sent: Friday, July 29, 2022 12:45 PM
To: Tracy Kefauver <Tracy.Kefauver@tn.gov>
Cc: Tawanna Reid <Tawanna.Reid@tn.gov>
Subject: [EXTERNAL] Adient PSD Application

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Hi Tracy,

I hope that you are doing well! I have been looking over the Adient PSD Application and there were a few things that I saw and wanted to touch base with you on. I know you are working on the draft permit so you may have already addressed these items but I just wanted to share what we found when looking through the application. I would be happy to further discuss any of these items with you, feel free to give me a call or I can set up a Teams meeting for us to go over them.

- It appears as though a baseline of 0 in the emission calculations was used because Adient became a major PSD source by relaxing enforceable limits, in accordance with TDEC rule 1200-03-09-.01(4)(a)6, and did not obtain a PSD permit at that time as included in Adient's February 2022 consent decree. Because this reasoning is not very clear in the application, it is recommended to include it in the preliminary and final determinations so that it is present in the permit record. This is included in the preliminary determination.
- The emission rates used in the calculations in Appendix B of the application are noted to be from a source test conducted on October 1, 1997. It is recommended to use emission factors from a more recent source test.
- The applications included discussion of filterable PM emissions but does not address condensible PM emissions. It is recommended to include any condensible PM emissions and further explain how no PM2.5 emissions are expected.
- The application states that 2006 dollars were used to calculate the control costs in the BACT analysis and Appendix C lists the cost base date as April 1988. Per Section 2.5.3 of EPA's Control Cost Manual, "It should be

noted that the accuracy associated with escalation (and its reverse, de-escalation) declines the longer the time period over which this is done. Escalation with a time horizon of more than five years is typically not considered appropriate as such escalation does not yield a reasonably accurate estimate." It is recommended to perform the cost analysis with more recent data than that of either 1988 or 2006. Additionally, the inclusion of more detailed cost calculations used to estimate the equipment costs or a quote, if a quote was used to determine the equipment costs, would be helpful in order to better understand the cost of add on controls.

• In the conclusion paragraph of the BACT analysis, the application states "Adient concludes and asserts the current configuration without VOC abatement is BACT for this project and for the existing process" and does not provide a numerical BACT limit. Even if no add on control technology is chosen as BACT, it is recommended to include a numerical BACT limit in the construction permit. The numerical limit is provided in the preliminary determination and the construction permit 980244.

Thank you, Emily

Emily Ferrando Air Permits Section EPA Region 4, Atlanta GA (404) 562-9042 Pronouns: <u>she/her</u> The following is the revised response requested by EPA Region IV (dated August 16, 2022) to clarify TDI emission factor

Tracy Kefauver

From:
Sent:
To:
Subject:

Kris Patrick Foster <kris.patrick.foster@adient.com> Tuesday, August 16, 2022 1:12 PM Tracy Kefauver [EXTERNAL] Requested clarification

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Tracy,

Please see clarification of TDI emissions factors:

The testing conducted on October 1, 1997 was specific to TDI and MDI from foam manufacturing. This is the most recent testing event representative of the foam types used by the Company. A more recent testing event at the company sites is not available for use.

Further, current formulations and process knowledge would indicate the emissions factors from the October 1997 testing, used to predict TDI and MDI emissions, are believed to provide the worst case emissions rates resulting in an overstatement and estimate of TDI and MDI emissions. Since the projected emissions for TDI and MDI are worst case and possibly an overstate of emission, the very low emission rates are believed a reasonable and conservatively high estimate for permitting purposes and provides an adequate margin of compliance with applicable regulations.



Kris P. Foster Environmental, Health and Safety Lead 1890 Mines Road Pulaski, TN 38478 Cell: 931-638-5918 Office: 931-424-7848

Adient – INTERNAL

Appendix F – Email transmittals to EPA and Affected States

Appendix G – Response to Comments