

SAFETY DATA SHEET

1. Identification

Product identifier	SLOW SET ASPHALT EMULSION		
Other means of identification			
SDS number	9586		
Synonyms	APPLICABLE TO ALL CSS-1 D (DILUTE) PRODUCTS * APPLICABLE TO ALL CSS-1H D (DILUTE) PRODUCTS * APPLICABLE TO ALL SS-1 D (DILUTE) PRODUCTS * APPLICABLE TO ALL SS-1H D (DILUTE) PRODUCTS * CQS-1 * CQS-1H * CSS-1 * CSS-1 SPECIAL * CSS-1H * CSS-1H Slurry * CSS-1HP * CSS-1P * SS-1 * SS-1H * CIR-EE(S) * CIR-EE(H) * SB-EE(S) * SB-EE(H)		
Recommended use	Road construction and maintenance		
Recommended restrictions	Other uses are not recommended unless an a that use, which demonstrates that the use will	ssessment is completed, prior to commencement of be controlled.	
Manufacturer/Importer/Supplier/I	Distributor information		
Manufacturer			
Manufacturer	Flint Hills Resources Pine Bend, LLC		
	P.O. Box 2917		
	Wichita, KS		
	67201-2917		
	United States		
Telephone numbers – 24 hour emergency assistance			
Chemtrec	800-424-9300		
Telephone numbers – general assistance			
8-5 (M-F, CST) MSDS Assistance	316-828-7988		
Email:	msdsrequest@fnr.com		
2. Hazard(s) identification			
Physical hazards	Not classified.		
Health hazards	Skin corrosion/irritation	Category 1B	
	Serious eye damage/eye irritation	Category 1	
	Sensitization, skin	Category 1	
	Carcinogenicity	Category 1B	
	Specific target organ toxicity, single exposure	Category 3 narcotic effects	
	Specific target organ toxicity, repeated exposure	Category 2 (liver, thymus, bone marrow)	
	Aspiration hazard	Category 1	
Environmental hazards	Hazardous to the aquatic environment, acute hazard	Category 2	
	Hazardous to the aquatic environment, long-term hazard	Category 2	
OSHA defined hazards	Not classified.		
Label elements			
Signal word	Dangar		
Environmental hazards OSHA defined hazards Label elements Signal word	Aspiration hazard Hazardous to the aquatic environment, acute hazard Hazardous to the aquatic environment, long-term hazard Not classified.	Category 1 Category 2 Category 2	

Hazard statement	Causes severe skin burns and eye damage. May cause an allergic skin reaction. May cause cancer. May cause drowsiness or dizziness. May cause damage to organs (liver, thymus, bone marrow) through prolonged or repeated exposure. May be fatal if swallowed and enters airways. Toxic to aquatic life with long lasting effects.
Precautionary statement	
Prevention	Obtain special instructions before use. Do not handle until all safety precautions have been read and understood.
	Do not breathe mist or vapor. Wash thoroughly after handling. Use only outdoors or in a well-ventilated area. Contaminated work clothing must not be allowed out of the workplace. Avoid release to the environment. Wear protective gloves/protective clothing/eye protection/face protection.
Response	If swallowed: Rinse mouth. Do NOT induce vomiting.
	If on skin (or hair): Take off immediately all contaminated clothing. Rinse skin with water/shower. If inhaled: Remove person to fresh air and keep comfortable for breathing.
	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/doctor. If skin irritation or rash occurs: Get medical advice/attention. Wash contaminated clothing before reuse. Collect spillage.
	Specific treatment (see first aid instructions on this label).
Storage	Store in a well-ventilated place. Keep container tightly closed. Store locked up.
Disposal	Dispose of contents/container in accordance with local/regional/national/international regulations.
Hazard(s) not otherwise classified (HNOC)	Not classified.
Supplemental information Precautionary statement(s)	
Hazard statement	Contains or releases hydrogen sulfide, an extremely flammable and toxic gas. Gas may evolve from this material and accumulate in confined spaces.
	When it is heated, this material may cause thermal burns.
Prevention	Use personal protective equipment as required. Wear protective gloves/eye protection/face protection.

3. Composition/information on ingredients

Mixtures

Chemical name	Common name and synonyms	CAS number	%
WATER		7732-18-5	25 - 93
ASPHALT BINDER		Mixture	7 - 75
CATIONIC EMULSIFIER SALT		Proprietary	≤ 5
OIL DISTILLATES		Proprietary	≤ 5
Additional components	Common name and synonyms	CAS number	%
Chemical name			
PETROLEUM ASPHALT		8052-42-4	≤ 75
POLYMER MODIFIER		Proprietary	≤ 7
HYDROGEN SULFIDE		7783-06-4	≤ 0.1
POLYCYCLIC AROMATIC COM	POUNDS	130498-29-2	< 0.1

Composition comments	Values do not reflect absolute minimums and maximums; these values are typical which may vary from time to time.
	Asphalt component may contain antistrip, vulcanizing agent, and polymer modifier. Asphalt materials can contain hydrogen sulfide, because it is naturally occurring in crude oil from which asphalt is derived. Hydrogen sulfide can also be present as a byproduct of asphalt processing.
	The specific identities of some of the components of this product are being withheld as trade secrets. However, all pertinent hazards are addressed in this SDS.
	This Safety Data Sheet is intended to communicate potential health hazards and potential physical hazards associated with the product(s) covered by this sheet, and is not intended to communicate product specification information. For product specification information, contact your Flint Hills Resources, LP representative.
4. First-aid measures	
Inhalation	Remove to fresh air. If not breathing, institute rescue breathing. If breathing is difficult, ensure airway is clear and give oxygen. If heart has stopped, immediately begin cardiopulmonary resuscitation (CPR).
	Keep affected person warm and at rest. GET IMMEDIATE MEDICAL ATTENTION.
Skin contact	If hot material gets on skin, immediately flush affected area with large amounts of cool water. Do not attempt to remove the material from the skin, or to remove contaminated clothing. Get immediate medical attention.
	For cold material, immediately wash skin with plenty of soap and water after removing contaminated clothing and shoes. Get medical attention if irritation persists.
	Place contaminated clothing in closed container for storage until laundered or discarded. If clothing is to be laundered, inform person performing operation of contaminant's hazardous properties. Discard contaminated leather goods.
Eye contact	Flush immediately with large amounts of water for at least 15 minutes. Eyelids should be held away from the eyeball to ensure thorough rinsing. GET IMMEDIATE MEDICAL ATTENTION.
Ingestion	If spontaneous vomiting occurs, keep head below hips to prevent aspiration and monitor for breathing difficulty.
	Never give anything by mouth to an unconscious person.
	Keep affected person warm and at rest. GET IMMEDIATE MEDICAL ATTENTION.

Most important symptoms/effects, acute and delayed	INHALATION: Fumes, mists, or vapors from the heated material may be irritating to the respiratory tract. Symptoms may include headache, excitation, euphoria, dizziness, incoordination, drowsiness, light-headedness, blurred vision, fatigue, tremors, convulsions, loss of consciousness, coma, respiratory arrest and death, depending on the concentration and duration of exposure.
	Contains hydrogen sulfide gas. Hydrogen sulfide can cause respiratory paralysis and death, depending on the concentration and duration of exposure. Do not rely on ability to smell vapors, since odor fatigue rapidly occurs. Effects of overexposure include irritation of the nose and throat, nausea, vomiting, diarrhea, abdominal pain and signs of nervous system depression (e.g. headache, drowsiness, dizziness, loss of coordination and fatigue), irregular heartbeats, pulmonary edema, weakness and convulsions.
	SKIN: CORROSIVE. Contact can cause skin burns and permanent skin damage. Skin contact may cause harmful effects in other parts of the body.
	Contains a component(s) that may cause allergic skin reactions in some individuals.
	EYES: CORROSIVE. Contact can cause burns and permanent damage to eye tissue. Can cause blindness.
	Vapors may cause eye irritation and sensitivity to light.
	INGESTION: CORROSIVE. Swallowing this material may be harmful or cause death. Harmful effects include burns and permanent damage to the digestive tract, including the mouth, throat, stomach and intestines. Symptoms may include severe abdominal pain and vomiting of blood. Blood loss through damaged tissue can lead to low blood pressure and shock.
	Aspiration into lungs may cause chemical pneumonia and lung damage.
Indication of immediate medical attention and special treatment needed	INHALATION: Inhalation exposure can produce toxic effects. Treat intoxications as hydrogen sulfide exposures. Monitor for respiratory distress. If cough or difficulty in breathing develops, evaluate for upper respiratory tract inflammation, bronchitis, and pneumonitis.
	Treat as corrosive material. Monitor for possible pharyngeal and pulmonary edema. Onset may be delayed up to 24 hours from the time of exposure. Administer supplemental oxygen with assisted ventilation, as required.
	Signs and symptoms of CNS depression, confusion and convulsions should be considered in the assessment and treatment of victims of exposures.
	INGESTION: This material is primarily an irritant and corrosive. As a corrosive, give attention to potential complication of esophagus or stomach perforations if ingested. Use of emetics and lavage are contraindicated. Necrosis and associated inflammatory processes peak at about 48 hours, but may extend up to four days. Initial healing processes occur during the period 4 to 14 days, but the esophageal wall is weakest during this period.
	SKIN: Hot material may cause skin burns. Immerse skin covered with hot material in cool water to limit tissue damage and prevent spread of liquid material. Consider leaving cooled material on skin unless contraindicated by contamination or potential for tattooing. If removal is necessary, mineral oil may be of assistance in minimizing skin loss when removing cool, hardened asphalt.
	EYES: Hot material may cause burns to the eyes. Early ophthalmologic evaluation is recommended.
5. Fire-fighting measures	
Suitable extinguishing media	Use water spray, dry chemical, carbon dioxide or fire-fighting foam for Class B fires to extinguish fire.
Unsuitable extinguishing media	Do not use a solid water stream as it may scatter and spread fire.

Specific hazards arising from the chemical	Combustion may produce COx, SOx, NOx, reactive hydrocarbons, irritating vapors, and other decomposition products in the case of incomplete combustion. Fires involving this product may release hydrogen sulfide.	
	Material will burn in a fire.	
	Hydrogen sulfide can react with the iron in an asphalt storage tank to form iron sulfide. Iron sulfide is pyrophoric. When exposed to air, iron sulfide is capable of igniting spontaneously.	
Special protective equipment and precautions for firefighters	Evacuate area and fight fire from a safe distance.	
	Use water spray to cool adjacent structures and to protect personnel. Shut off source of flow, if possible. Stay away from storage tank ends. Withdraw immediately in case of rising sound from venting safety device or any discoloration of storage tank due to fire. Always stay away from tanks engulfed in flame.	
	Exercise extreme care when using water spray on asphalt tank fires. When water is mixed with hot asphalt, steam may rapidly develop resulting in violent asphalt foaming and possible tank eruptions from increased pressure.	
	Firefighters must wear NIOSH approved positive pressure breathing apparatus (SCBA) with full face mask and full protective equipment.	
6. Accidental release meas	ures	
Personal precautions, protective equipment and emergency procedures	Keep unnecessary people away; isolate hazard area and deny entry. For spills in confined areas, ensure adequate ventilation. For spills outdoors, stay upwind. IF TANK, RAILCAR OR TANK TRUCK IS INVOLVED IN A FIRE, isolate for 800 meters (1/2 mile) in all directions. Evacuate area endangered by release as required. Wear appropriate personal protective equipment. See Exposure Controls/Personal Protection (Section 8).	
Methods and materials for containment and cleaning up	Keep unnecessary people away. Isolate area for at least 50 meters (164 feet) in all directions to preserve public safety. For large spills, if downwind consider initial evacuation for at least 300 meters (1000 feet).	
	For spills on land, scrape up spilled material for disposal. For large spills, dike ahead of spill to contain. For spills on water, contain as much as possible with booms and begin recovery as soon as possible. If material sinks or becomes dispersed, consult with local, state and regional authorities for approved clean up procedures.	
	Use a vapor suppressing foam to reduce vapors. Do not touch or walk through spilled material. Stop leak when safe to do so.	
	See Exposure Controls/Personal Protection (Section 8).	
Environmental precautions	Prevent entry into water ways, sewers, basements or confined areas. Notify local authorities and National Response Center, if required.	
7. Handling and storage		
Precautions for safe handling	Avoid contact with strong oxidizing agents. Prevent small spills to minimize slip hazard or release to the environment. Do not cut, grind, drill, weld (or introduce any other ignition source) on empty containers or reuse containers unless adequate precautions are taken. Avoid extreme temperatures to minimize product degradation.	
	Avoid personal contact with this material. Always observe good personal hygiene measures, such as removing contaminated clothing and protective equipment, washing after handling the material and before entering public areas. Restrict eating, drinking and smoking to designated areas to prevent personal chemical contamination. Routinely wash work clothing and protective equipment to remove contaminants. Do not breathe fumes, vapor or gas. See Section 8 of the SDS for Personal Protective Equipment.	
Conditions for safe storage, including any incompatibilities	Store in tightly closed containers in a cool, dry, isolated, well-ventilated area away from heat, sources of ignition and incompatibles. Avoid contact with strong oxidizing agents. Empty containers may contain material residue. Do not reuse without adequate precautions.	
	Hydrogen sulfide can build up in the head space of storage vessels containing this material. Use appropriate respiratory protection to prevent exposure. See Exposure Controls/Personal Protection (Section 8).	
	When entering a storage vessel that has previously contained this material it is recommended that the atmosphere be monitored for the presence of hydrogen sulfide. See Occupational exposure limits (Section 8) for exposure limits.	

8. Exposure controls/personal protection

Oc	cupational exposure limits					
	US. OSHA Table Z-1 Limits Additional components	s for Air Contan	ninants (29 CFF Type	R 1910.1000)	Value	
	POLYCYCLIC AROMATIC COMPOUNDS (CAS 130498-29-2) US. OSHA Table Z-2 (29 C	FR 1910.1000)	PEL		0.2 mg/m3	
	Additional components	,	Туре		Value	Form
	POLYCYCLIC AROMATIC COMPOUNDS		TWA		0.2 mg/m3	Coal tar pitch volatiles (benzene soluble fraction)
	(CAS 130498-29-2) HYDROGEN SULFIDE (CAS 7783-06-4)		Ceiling		20 ppm	
	US. ACGIH Threshold Lim	it Values	_			-
	Additional components		Туре		Value	Form
	PETROLEUM ASPHALT (CAS 8052-42-4)		TWA		0.5 mg/m3	Inhalable fraction (as benzene-soluble aerosol)
	POLYCYCLIC AROMATIC COMPOUNDS (CAS 130498-29-2)		TWA		0.2 mg/m3	Coal tar pitch volatiles (benzene soluble fraction)
	HYDROGEN SULFIDE		STEL		0.2 mg/m3 5 ppm	Aerosol.
			TWA		1 ppm	
	US. NIOSH: Pocket Guide	to Chemical Ha	zards			
	Additional components		Туре		Value	Form
	PETROLEUM ASPHALT (CAS 8052-42-4)		Ceiling		5 mg/m3	Fume.
	POLYCYCLIC AROMATIC COMPOUNDS (CAS 130498-29-2)		TWA		0.1 mg/m3	Cyclohexane-extractable fraction.
	HYDROGEN SULFIDE (CAS 7783-06-4)		Ceiling		10 ppm	
Bio	logical limit values					
	ACGIH Biological Exposur Additional components	re Indices Value	Specimen	Sampling Time		
	POLYCYCLIC AROMATIC COMPOUNDS (CAS 130498-29-2)	2.5 μg/l	1-Hydroxypy rene in urine	*		
	* - For sampling details, plea	ase see the sour	ce document.			
Exp	oosure guidelines					
	US OSHA Specifically Reg	ulated Substan	ces: Action lev	el and Reference		
	POLYCYCLIC AROMA 130498-29-2)	TIC COMPOUNE	DS (CAS	29 CFR 1910.1002	2	
Ap cor	Appropriate engineering Cons controls equip subst		Consider the following when employing engineering controls and selecting personal protective equipment: potential hazards of the material, applicable exposure limits, job activities, and other substances in the work place.			
		Ventilation are exposures be	nd other forms c elow occupation	of engineering controls al exposure limits and	are the preferred i guidelines.	means for controlling
Ind	ividual protection measure	s, such as pers	onal protective	equipment		
Eye/face protection and/or face shield. Have eye washing facilities readily available where eye contact can			om eyes. Eye o hield. Have eye	contact can be avoide washing facilities rea	d by using chemica dily available where	Il safety glasses, goggles e eye contact can occur.

Skin protection	
Hand protection	Prevent any skin contact with this material. Use appropriate chemical resistant gloves. Glove suitability for a job must be determined by the user for specific use conditions. Contact the glove manufacturer for specific advice on glove selection regarding permeability and breakthrough times for your use conditions. Gloves should be discarded and replaced if there is any indication of degradation or chemical breakthrough.
	When handling hot material, use heat resistant gloves.
Other	Prevent any skin contact with this material. Additional protective clothing may be necessary.
Respiratory protection	The use of air purifying respirators is not recommended where hydrogen sulfide levels may exceed exposure limits. Use a positive pressure air supplied respirator if there is any potential for an uncontrolled release, exposure levels are not known, or any other circumstances where air purifying respirators may not provide adequate protection. See OSHA 29 CFR 1910.134 for more information regarding respiratory protection and Assigned Protection Factors (APFs).
Thermal hazards	Contact with hot material can cause thermal burns which may result in permanent damage. Wear appropriate thermal protective clothing. Additional protection may be necessary to prevent skin contact including use of apron, arm covers, face shield, or boots.

9. Physical and chemical properties

Appearance		
Physical state	Liquid.	
Form	Not applicable	
Color	Dark brown	
Odor	Musty	
Odor threshold	Not available.	
рН	2 - 10.5	
Melting point/freezing point	< 32 °F (< 0 °C)	
Initial boiling point and boiling range	212 °F (100 °C)	
Flash point	>212 °F (>100 °C)	
Evaporation rate	Not available	
Flammability (solid, gas)	Not applicable.	
Upper/lower flammability or exp	losive limits	
Explosive limit - lower (%)	Not available.	
Explosive limit - upper (%)	Not available.	
Vapor pressure	23.76 mmHg at 77 °F (25 °C) (similar to water)	
Vapor density	Not available	
Relative density	0.9 - 1.1 at 60/60 °F (15.6/15.6 ℃)	
Solubility(ies)		
Solubility (water)	Dispersible	
Partition coefficient (n-octanol/water)	Not available	
Auto-ignition temperature	Not available	
Decomposition temperature	Not available.	
Viscosity	10 - 700 SFS at 77 ℉ (25 ℃)	
10. Stability and reactivity		
Reactivity	See statements below.	
Chemical stability	Material is stable under normal conditions.	
Possibility of hazardous reactions	Not anticipated under normal conditions.	
Conditions to avoid	Avoid overheating, emissions generation, unventilated areas, heat, open flames.	
Incompatible materials	Incompatible with strong oxidizing agents. See precautions under Handling & Storage (Section 7).	
Hazardous decomposition products	Not anticipated under normal conditions.	

11. Toxicological information

Information on likely routes of exposure

Components	Species	Test Results	
Acute toxicity	Not classified.		
Information on toxicological ef	fects		
	Aspiration into lungs m	ay cause chemical pneumonia and lung damage.	
	INGESTION: CORROSIVE. Swallow burns and permanent intestines. Symptoms through damaged tissu	ing this material may be harmful or cause death. Harmful effects include damage to the digestive tract, including the mouth, throat, stomach and may include severe abdominal pain and vomiting of blood. Blood loss the can lead to low blood pressure and shock.	
	Vapors may cause eye	irritation and sensitivity to light.	
	EYES: CORROSIVE. Contact blindness.	can cause burns and permanent damage to eye tissue. Can cause	
	Contains a component	(s) that may cause allergic skin reactions in some individuals.	
	SKIN: CORROSIVE. Contact cause harmful effects	can cause skin burns and permanent skin damage. Skin contact may n other parts of the body.	
	Contains hydrogen sul depending on the cond since odor fatigue rapi nausea, vomiting, diari headache, drowsiness pulmonary edema, we	tide gas. Hydrogen sulfide can cause respiratory paralysis and death, entration and duration of exposure. Do not rely on ability to smell vapors, dly occurs. Effects of overexposure include irritation of the nose and throat, hea, abdominal pain and signs of nervous system depression (e.g. dizziness, loss of coordination and fatigue), irregular heartbeats, akness and convulsions.	
Symptoms related to the physical, chemical and toxicological characteristics	INHALATION: Fumes, mists, or vapo Symptoms may include light-headedness, blur respiratory arrest and e	INHALATION: Fumes, mists, or vapors from the heated material may be irritating to the respiratory tract. Symptoms may include headache, excitation, euphoria, dizziness, incoordination, drowsiness, light-headedness, blurred vision, fatigue, tremors, convulsions, loss of consciousness, coma, respiratory arrest and death, depending on the concentration and duration of exposure.	
Ingestion	Likely route of exposur	e	
Eye contact	Likely route of exposur	e	
Skin contact	Likely route of exposur	e	
Inhalation	Likely route of exposur	e	

Acute toxicity	Not oldssined.			
Components	Species	Test Results		
OIL DISTILLATES				
Acute				
Dermal				
LD50	Rat	> 4300 mg/kg		
Inhalation				
Mist				
LC50		4.1 mg/l		
Oral				
LD50	Rat	> 7600 mg/kg		
WATER (CAS 7732-18-5)				
Acute				
Oral				
LD50	Rat	> 89800 mg/kg		
Skin corrosion/irritation	Causes severe skin burns and eye damage.			
Serious eye damage/eye irritation	Causes serious eye damage.			
Respiratory or skin sensitizatio	n			
Respiratory sensitization	Not classified.			
Material name: SLOW SET ASPHAL	T EMULSION	SDS L		

Skin sensitization	May cause an allergic skin reaction.	
Germ cell mutagenicity Not classified.		
Carcinogenicity May cause cancer.		
ACGIH Carcinogens		
ASPHALT (BITUMEN) FUME, AS BENZENE-SOLUBLE AEROSOL, INHALABLE FRACTION (CAS 8052-42-4)		A4 Not classifiable as a human carcinogen.
COAL TAR PITCH VOLAT SOLUBLE AEROSOL (CA	TILES, AS BENZENE IS 130498-29-2)	A1 Confirmed human carcinogen.
IARC Monographs. Overall E	valuation of Carcinogenicity	
PETROLEUM ASPHALT (POLYCYCLIC AROMATIC 130498-29-2)	CAS 8052-42-4) COMPOUNDS (CAS	2B Possibly carcinogenic to humans.1 Carcinogenic to humans.
US. National Toxicology Prog	gram (NTP) Report on Carcinc	ogens
POLYCYCLIC AROMATIC 130498-29-2)	COMPOUNDS (CAS	Reasonably Anticipated to be a Human Carcinogen.
US. OSHA Specifically Regul	ated Substances (29 CFR 191	0.1001-1050)
Not regulated.		
Reproductive toxicity	Not classified.	
Specific target organ toxicity - single exposure	May cause drowsiness or dizzi	ness.
Specific target organ toxicity - repeated exposure	May cause damage to organs exposure.	(liver, thymus, bone marrow) through prolonged or repeated
Aspiration hazard	May be fatal if swallowed and e	enters airways.
Toxicological data		
	HYDROGEN SULFIDE: Hydrogen sulfide causes rapid death due to metabolic asphyxiation. It has an unpleasant "rotten egg" odor that diminishes with increased exposure and is not a reliable indicator for warning of exposure. Eye irritation may occur at levels above 4 ppm. Olfactory fatigue occurs rapidly at levels of 50 ppm or higher. Respiratory effects include irritation with possible pulmonary edema at levels above 50 ppm. At 500 ppm immediate loss of consciousness and death can occur. NIOSH has determined that 100 ppm hydrogen sulfide is immediately dangerous to life and health (IDLH).	
	PROPRIETARY PROCESS OILS: Long-term repeated (lifetime) skin exposure also has been reported to result in an increase in skin tumors in laboratory rodents. The International Agency for Research on Cancer (IARC) has concluded that there is sufficient evidence of carcinogenicity in experimental animals for these types of oils (Group 1B).	
	POLYCYCLIC AROMATIC HY PAHs. Certain PAHs are weak metabolism. Chronic or repeat incidence of tumors of the skin individuals overexposed to cer photosensitivity and eye irritativ respiratory tract irritation, coug precancerous lesions, erythem Oral overexposure to PAHs ha (leukoplakia). Mild nephrotoxic function tests, changes in the i exposed to high levels of PAHs	DROCARBONS (PAHs): Cancer is the most significant endpoint for a carcinogens which become carcinogenic after undergoing ted exposure increases the likelihood of tumor initiation. Increased , bladder, lung and gastrointestinal tract have been described in tain PAHs. Overexposure to PAHs has also been associated with on. Inhalation overexposure of PAHs has been associated with h, and bronchitis. Dermal overexposure has been associated with ha, dermal burns, photosensitivity, acneiform lesions and irritation. s been associated with precancerous growths of the mouth city, congestion and renal cortical hemorrhages and elevated liver mmune system and other effects have been observed in rats s by ingestion.

ASPHALT/ASPHALT LIKE PRODUCTS: Asphalt fumes from heated material have been reported to cause eye, respiratory tract and skin irritation, as well as nausea and headaches. Symptoms may include coughing, wheezing and shortness of breath. An adverse effect on pulmonary function has not been conclusively demonstrated. Studies in humans to determine the potential long-term health effects of asphalt also have had inconsistent results. Epidemiological studies in European paving asphalt worker populations indicated a slight positive association between lung cancer mortality and exposure to asphalt fumes. A case-control examination of these data found no consistent evidence of an association between bitumen and lung cancer risk, possibly due to the confounding effects of potential exposure to coal tar cigarette smoking, and other substances. Additional studies of workers exposed to asphalt emissions during paving with straight-run asphalt showed mutagenic and genotoxic/cytogenetic effects in these workers.

Studies in experimental animals have not established a link between lung cancer and asphalt fume exposure. However, an increase in skin tumors was observed in lifetime studies of laboratory rodents exposed to extracts of asphalt (bitumen) as well as "cutbacks" of asphalt (asphalts that are diluted, dissolved or liquefied in hydrocarbon solvents).

An increased incidence of skin tumors was also observed in lifetime dermal bioassays of laboratory rodents exposed to distillates of fumes generated from roofing flux, an asphalt that is further processed or oxidized. These condensed fumes were collected from an oxidized roofing asphalt at high temperatures (>450 degrees F). Follow up studies suggest that the roofing asphalt distillates act as tumor initiators, involving a genotoxic mechanism. No increases in skin tumors were found in a lifetime study of rodents dermally exposed to distillates of fumes generated from paving asphalt.

The International Agency for Research on Cancer (IARC) recently determined that occupational exposures to oxidized asphalt and their emissions during roofing applications are "probably carcinogenic to humans" (Group 2A). They also determined that occupation exposures to hard asphalts and their emissions during mastic asphalt work and occupational exposures to straight-run asphalts and their emissions during paving operations are "possibly carcinogenic to humans" (Group 2B).

Ecotoxicity	Toxic to aquatic life with long lasting effects.				
Components		Species	Test Results		
OIL DISTILLATES					
Aquatic					
Acute					
Algae	EC50	Pseudokirchnerella subcapitata	10 mg/l, 72 hr		
Crustacea	EC50	Daphnia magna	68 mg/l, 48 hr		
Fish	LC50	Oncorhynchus mykiss	21 mg/l, 96 hr		
Chronic					
Crustacea	NOEC	Daphnia magna	0.2 mg/l, 21 d		
Fish	NOEC	Oncorhynchus mykiss	0.08 mg/l, 14 d		
Persistence and degradability	Not readil	Not readily biodegradable.			
Bioaccumulative potential	May bioad	May bioaccumulate in aquatic organisms.			
Mobility in soil	May partition into soil and water.				
Other adverse effects	No other adverse effects expected.				
13. Disposal consideration	ons				
Disposal instructions	This material, as supplied, when discarded or disposed of, is not a hazardous waste according to Federal Regulations (40 CFR 261).				
Hazardous waste code	The transportation, storage, treatment and disposal of waste material must be conducted in compliance with federal, state, and local regulations. Under RCRA it is the responsibility of the user of the material to determine, at the time of disposal, whether this material meets RCRA criteria for hazardous waste. For additional handling information and protection of employees, see Section 7 (Handling and Storage) and Section 8 (Exposure Controls/Personal Protection). The proper waste code must be evaluated at the time of disposal and should be determined by the user and waste disposal company.				

12. Ecological information

Waste from residues / unused products	Dispose of this material in accordance with all applicable local and national regulations.			
Contaminated packaging	Empty containers should be ta accordance with government re-	ken to an approved waste handling site for recycling or disposal in egulations. Packaging may contain residue that can be hazardous.		
14. Transport information				
General information	In accordance with US DOT, b transportation: *below 212 °F (100 °C), are no *above 212 °F (100 °C), are UI Distillates) 9, III *greater than the flash point, a (Petroleum Distillates) 3, III	ulk and non-bulk shipments of this product, which are offered for t regulated. N3257, Elevated Temperature Liquid, Flammable, NOS (Petroleum re UN3256, Elevated Temperature Liquid, Flammable, NOS		
Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code	Not classified for MARPOL. Pl mode is ship or vessel to deter	ease contact the Transportation Compliance CSO if transportation mine the need for a MARPOL classification.		
15. Regulatory information				
US federal regulations	All ingredients are on the TSCA inventory, or are not required to be listed on the TSCA inventory.			
	This material does not contain concentration) that are subject Superfund Amendments and F	toxic chemicals (in excess of the applicable de minimis to the annual toxic chemical release reporting requirements of the Reauthorization Act (SARA) Section 313 (40 CFR 372).		
	Check local, regional or state/p be more restrictive than federa civil and criminal penalties.	provincial regulations for any additional requirements as these may I laws and regulations. Failure to comply may result in substantial		
US EPCRA (SARA Title III) Se	ection 313 - Toxic Chemical: D	e minimis concentration		
HYDROGEN SULFIDE (C POLYCYCLIC AROMATIC 130498-29-2)	AS 7783-06-4) COMPOUNDS (CAS	1.0 % 0.1 %		
HYDROGEN SULFIDE (C	AS 7783-06-4)	100 LBS		
US EPCRA (SARA Title III) Se	ection 312 - Extremely Hazard	ous: Listed substance		
HYDROGEN SULFIDE (C	AS 7783-06-4)	Listed.		
	AS 7783-06-4)	100 LBS		
US. OSHA Specifically Regul	ated Substances (29 CFR 191	0.1001-1050)		
Not regulated.				
Superfund Amendments and Rea	authorization Act of 1986 (SAF	RA)		
Hazard categories	Immediate Hazard - Yes Delayed Hazard - Yes Fire Hazard - No Pressure Hazard - No Reactivity Hazard - No			
Other federal regulations	Please see Section 2 for OSH	A hazard classification(s) for EPCRA Tier I/Tier II reporting.		
Clean Air Act (CAA) Section 112 Hazardous Air Pollutants (HAPs) List POLYCYCLIC AROMATIC COMPOUNDS (CAS 130498-29-2) Clean Air Act (CAA) Section 112(r) Accidental Belease Prevention (40 CEP 68 130)				
HYDROGEN SULFID	E (CAS 7783-06-4)			
US state regulations				
US. California Proposition 65	5			
WARNING: This product HSC. §25249.5.	contains a chemical known to th	ne State of California to cause cancer. Proposition 65, CAL.		
US - California Proposition 65 - CRT: Listed date/Carcinogenic substance				
PETROLEUM ASPHALT (CAS 8052-42-4) LISTED: January 1, 1990				
16. Other information, inclu	uding date of preparation	n or last revision		
Issue date	04-01-2015			

02-17-2017

Revision date

Version #	04
HMIS® ratings	Health: 2* Flammability: 0 Physical hazard: 0 * Indicates chronic health hazard
NFPA ratings	Health: 2 Flammability: 1 Instability: 0
Disclaimer	THIS SDS HAS BEEN PREPARED TO COMPLY WITH FEDERAL REGULATIONS THAT ARE INTENDED TO QUICKLY PROVIDE USEFUL INFORMATION TO THE USER(S) OF THIS MATERIAL OR PRODUCT - IT IS NOT INTENDED TO SERVE AS A COMPREHENSIVE DISCUSSION OF ALL POSSIBLE RISKS OF HAZARDS, BUT RATHER PROVIDES INFORMATION GENERALLY ACCEPTED IN THE SCIENTIFIC COMMUNITY AS RELEVANT REGARDING THE POTENTIAL HAZARDS OF THIS PRODUCT. ADEQUATE TRAINING, INSTRUCTION, WARNINGS AND SAFE HANDLING PROCEDURES SHOULD BE PROVIDED TO HANDLERS AND USERS. USERS SHOULD REVIEW THE INFORMATION IN THE SDS, AND SATISFY THEMSELVES AS TO ITS SUITABILITY AND COMPLETENESS, INCLUDING ENSURING THAT THIS IS THE MOST CURRENT SDS.
Revision information	Physical & Chemical Properties: Multiple Properties Regulatory Information: United States
Completed by	Flint Hills Resources, LP - Operations EH&S