#### Clear Chain and Cable Lube

# **SAFETY DATA SHEET**

SDS PREP DATE: 2/21/2022

SDS REVISION DATE: 2/21/2022

**Product ID:** ACC381

Pro-Line Industrial Products, Inc. Pro-Line Industrial Products, Inc. 723 W University Ave 110-428 Georgetown, TX 78626 800-263-9436

24 Hour Emergency - 1-800-424-9300

This Safety Data Sheet contains environmental, health and toxicology information for your employees. Please make sure this information is given to them. It also contains information to help you meet community Right To Know emergency response reporting requirements under SARA TITLE III and many other laws. If you resell this product, this SDS must be given to the buyer or the information incorporated in your SDS.

# Section 1: Company and Product Identification \_\_\_\_\_\_

Pro-Line Industrial Products, Inc. Pro-Line Industrial Products, Inc. 723 W University Ave 110-428 Georgetown, TX 78626 800-263-9436 24 Hour Emergency - 1-800-424-9300

Product ID: ACC381

Product Name: Clear Chain and Cable

Intended Use: CHAIN AND CABLE LUBRICANT

### Section 2: Hazard(s) Identification

Product Signal Word: DANGER

**Physical Hazard Classification:** Flammable Aerosols, Category 1

Extremely flammable aerosol.

#### **Physical Hazard Precautionary Statements:**

Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.

Do not spray on an open flame or other ignition source.

Do not pierce or burn, even after use.

Protect from sunlight.

Do not expose to temperatures exceeding 50 °C/122°F.

# **Health Hazard Classification(s):**

Acute Toxicity - Oral - Level 4

Acute Toxicity - Dermal - Level 4

Skin Corrosion/Irritation -Level 3

Eye Damage/Irritation -Level 2B

Carcinogenicity - Level 2

Aspiration Hazard - Level 2

#### **Health Hazard Statements:**

Harmful if swallowed.







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May be harmful if swallowed and enters airways.

Harmful in contact with skin.

Causes mild skin irritation.

Causes eye irritation.

Suspected of causing cancer.

#### First-Aid Statement(s):

IF SWALLOWED: Immediately call a POISON CENTER/doctor/physician.

IF SWALLOWED: Call a POISON CENTER/doctor if you feel unwell.

IF ON SKIN: Wash with plenty of water.

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.

Call a POISON CENTER/doctor/physician if you feel unwell.

See Section 12 if specific treatment is applicable.

Do NOT induce vomiting.

If skin irritation occurs: Get medical advice/attention.

If eye irritation persists: Get medical advice/attention.

Wash contaminated clothing before reuse.

#### **Handling and Storage Statement(s):**

Use personal protective equipment as required.

Store locked up.

#### **Exposure Control / Personal Protection Statement(s):**

Obtain special instructions before use.

Do not handle until all safety precautions have been read and understood.

Wash hands and exposed areas thoroughly after handling.

Do no eat, drink or smoke when using this product.

Wear protective gloves/protective clothing/eye protection/face protection.

Use personal protective equipment as required.

#### **Disposal Statement(s):**

Dispose of contents/container in accordance with local regulations.

# **Section 3: Product Composition**

	CAS#	% Range		PEL	TLV
HYDROCARBON PROPELLANT	68476-86-8	5%	10%	NO DATA	NO DATA
ALIPHATIC HYDROCARBON *	110-54-3	1%	10%	500 PPM	50 PPM

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	p-Chlorobenzotrifluoride	98-56-6	1%	10%	NOT ESTABLISHED	NOT ESTABLISHED
	POLYISOBUTYLENE	9003-27-4	1%	10%	NOT ESTABLISHED	NOT ESTABLISHED
	LUBE OIL ADDITIVE CONTAINING	9003-29-6	30%	50%	NOT ESTABLISHED	NOT ESTABLISHED
	BARIUM SULFONATE	CONFIDENTIAL	1%	10%	NOT ESTABLISHED	NOT ESTABLISHED
	PETROLEUM HYDROCARBON BASE	64741-88-4	1%	10%	NOT ESTABLISHED	NOT ESTABLISHED
	ACETONE	67-64-1	10%	20%	TWA 1000 PPM	TWA 750 PPM STEL 1000 PPM
	HYDROTREATED LIGHT DISTILLATE	64742-47-8	1%	10%	NONE ESTABLISHED BY OSHA	NONE ESTABLISHED BY OSHA

Specific chemical identity and exact percentages are withheld as Trade Secret.

#### **Section 4: First-Aid Measures**

IF SWALLOWED: Immediately call a POISON CENTER/doctor/physician.

IF SWALLOWED: Call a POISON CENTER/doctor if you feel unwell.

IF ON SKIN: Wash with plenty of water.

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.

Call a POISON CENTER/doctor/physician if you feel unwell.

See Section 12 if specific treatment is applicable.

Do NOT induce vomiting.

If skin irritation occurs: Get medical advice/attention.

If eye irritation persists: Get medical advice/attention.

Wash contaminated clothing before reuse.

GENERAL: This material is an aspiration hazard and defats the skin. Breathing vapors of high concentrations may cause CNS depression.

EYE CONTACT: Slightly irritating but does not injure eye tissue.

SKIN CONTACT: Low order of toxicity. Frequent or prolonged contact may irritate and cause dermatitis. Skin contact may aggravate an existing dermatitis condition.

INHALATION: High vapor/aerosol concentrations (greater than approximately 100 ppm) are irritating to the eyes and the respiratory tract, may cause headaches, dizziness, anesthesia, drowsiness, unconsciousness, and other central nervous system effects, including death.

INGESTION: Small amounts of this product aspirated into the respiratory system during ingestion or vomiting may cause mild to severe pulmonary injury, possibly minimal toxicity.

#### FIRST AID

EYE CONTACT: Flush eyes with large amounts of water until irritation subsides. If irritation persists, get medical attention.

SKIN CONTACT: Flush with large amounts of water; use soap if available. Remove grossly contaminated clothing, including shoes, and launder before

INHALATION: Using proper respiratory protection, immediately remove the affected victim from exposure. Administer artificial respiration if breathing is stopped. Keep at rest. Call for prompt medical attention

INGESTION: If swallowed, DO NOT induce vomiting. Keep at rest. Get prompt medical attention.

#### **PRECAUTIONS**

SPECIAL PRECAUTIONS: Health studies have shown that many hydrocarbons pose potential human health risks which may vary from person to person. As a precaution, exposure to liquids, vapors, mists or fumes should be minimized.

PERSONAL PROTECTION: For open systems where contact is likely, wear safety glasses with side shields, long sleeves, and chemical resistant gloves.

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Where concentrations in air may exceed the limits, work practice or other means of exposure reduction are not adequate, NIOSH/MSHA approved respirators may be necessary to prevent overexposure by inhalation.

VENTILATION: The use of mechanical dilution ventilation is recommended whenever this product is used in a confined space, is heated above ambient temperatures, or is agitated.

# Section 5: Fire-Fighting Measures \_\_\_\_\_

FIRE AND EXPLOSION HAZARDS: This product releases Flammable Vapors at well below ambient temperatures and readily forms flammable mixtures with air exposed to an ignition source. It will burn in the open or be explosive in confined spaces. Its vapors are heavier than air and may travel long distances to a point of ignition, and then flash back. Alkaline/chlorine gas mixtures have produced explosions.

EXTINGUISHING MEDIA: Dry Chemical. CO2. Halogenated Extinguishing Agent. Stop Gas Flow.

SPECIAL FIREFIGHTING PROCEDURES: Gas fires should not be extinguished unless the gas flow can be stopped immediately. Allow the fire to burn itself out. If the source cannot be shut off immediately, all equipment and surfaces exposed to the fire should be cooled with water to prevent over-heating flash-backs, or explosions. Control fire until gas supply can be shut off. Use proper protective equipment. Use fresh air respirator when exposure to hazardous concentrations of toxic gases is possible.

FIRE FIGHTING: Use water spray to cool fire exposed surfaces and to protect personnel. Isolate "fuel" supply from fire. Use foam, dry chemical, or water spray to extinguish fire. Avoid spraying water directly into storage containers due to danger of boiling over. This liquid is volatile and gives off invisible vapors. Either the liquid or vapor may settle in low areas or travel some distance along the ground or surface to ignition sources where they may ignite or explode.

#### Section 6: Accidental Release Measures

STEPS TO BE TAKEN IN CASE CONTAINER IS PUNCTURED AND MATERIAL IS RELEASED:

Clean up area by mopping or with absorbent materials and place in closed container for disposal. Consult Federal, State, and local disposal authorities.

WASTE DISPOSAL METHOD: Consult local authorities for proper waste disposal procedures. Empty de-pressurized containers can not be reused. Cans which are pressurized or contain liquid must be disposed of in a permitted waste management facility. Consult Federal, State, and local disposal authorities for approved procedures.

# Section 7: Handling and Storage \_\_\_\_\_

Use personal protective equipment as required.

Store locked up.

VENTILATION REQUIREMENT: Use adequate level exhaust ventilation. Note: Where carbon monoxide may be generated, special ventilation may be required. Local exhaust recommended when appropriate to control employee exposure.

RESPIRATORY PROTECTION: Based on contamination level and working limits of the respirator, use a respirator approved by NIOSH/MSHA.

EYES: Face shield and goggles or chemical goggles should be worn.

GLOVES: Impervious gloves should be worn. Gloves contaminated with the product should be discarded. Polyfluorinated polyethylene has been suggested.

OTHER CLOTHING EQUIPMENT: Standard work clothing. Standard work shoes; discard if shoes can not be decontaminated. Store contaminated clothing in well ventilated cabinets or closed containers. Wash contaminated clothing and dry before reuse.

RESPIRATORY PROTECTION: In situations where vapor concentrations exceed the recommended exposure limits, a NIOSH approved organic vapor

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cartridge or air-supplying respirator should be worn.

#### Section 8: Exposure Control / Personal Protection \_\_\_\_\_

Obtain special instructions before use.

Do not handle until all safety precautions have been read and understood.

Wash hands and exposed areas thoroughly after handling.

Do no eat, drink or smoke when using this product.

Wear protective gloves/protective clothing/eye protection/face protection.

Use personal protective equipment as required.

VENTILATION REQUIREMENT: Use adequate level exhaust ventilation. Note: Where carbon monoxide may be generated, special ventilation may be required. Local exhaust recommended when appropriate to control employee exposure.

RESPIRATORY PROTECTION: Based on contamination level and working limits of the respirator, use a respirator approved by NIOSH/MSHA.

EYES: Face shield and goggles or chemical goggles should be worn.

GLOVES: Impervious gloves should be worn. Gloves contaminated with the product should be discarded. Polyfluorinated polyethylene has been suggested.

OTHER CLOTHING EQUIPMENT: Standard work clothing. Standard work shoes; discard if shoes can not be decontaminated. Store contaminated clothing in well ventilated cabinets or closed containers. Wash contaminated clothing and dry before reuse.

RESPIRATORY PROTECTION: In situations where vapor concentrations exceed the recommended exposure limits, a NIOSH approved organic vapor cartridge or air-supplying respirator should be worn.

# **Section 9: Product Properties\_**

Flash Point (CCP): LVL 3 AEROSOL, PROPELLENT: -136°F

Boiling Point for Product: N/D

Vapor Pressure for Product: N/D

Vapor Density for Product: N/D

Specific Gravity: N/D

V.O.C.: <25%

Water Solubility: NIL

Appearance: CLEAR AEROSOL SPRAY

PH: N/D

# **Section 10: Stability and Reactivity**

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STABILITY: Stable

CONDITIONS TO AVOID: Temperatures above 130 degree F.

HAZARDOUS POLYMERIZATION: Will not occur

MATERIALS AND CONDITIONS TO AVOID INCOMPATIBILITY: Strong oxidizing agents

HAZARDOUS DECOMPOSITION PRODUCTS: None

### Section 11: Toxicological Information\_\_\_\_\_

HYDROCARBON PROPELLANT 68476-86-8

ALIPHATIC HYDROCARBON \* 110-54-3

Acute oral toxicity:LD 50 Rat:2,500 mg/kgAcute inhalation toxicity:LC 50 Rat:48,000 ppm, 4 hoursAcute dermal toxicity:LD 50 Rabbit:1,300 mg/kg

Routes of Entry: Inhalation, skin absorption, skin contact

**Acute Exposure Hazards:** 

**INHALATION HAZARD:** Inhalation of vapors irritates the respiratory tract. Overexposure may cause central nervous system depression with lightheadedness, nausea, headache, and blurred vision. Greater exposure may cause muscle weakness, numbness of the extremities, unconsciousness and suffocation. Vapors can displace oxygen, especially in confined spaces.

**INGESTION HAZARD:** May produce gastrointestinal irritation with abdominal pain, nausea, vomiting, and diarrhea. Aspiration into lungs may cause chemical pneumonitis, which may be fatal. May cause central nervous system depression.

**SKIN CONTACT HAZARD:** May cause redness, irritation, dryness, cracking, and pain. Defatting or dermatitis may result from prolonged or repeated exposure. Hexane may be absorbed through the skin with possible systemic effects. There are no reports of skin sensitization through occupational exposure. Sensitization was not observed in a maximization test using 25 volunteers.

**EYE CONTACT HAZARD:** Vapors cause mild irritation. Splashes may cause redness and pain.

**Chronic Exposure Hazards:** Repeated or prolonged skin contact may defat the skin and produce irritation and dermatitis. Prolonged exposure may cause adverse reproductive effects and visual disturbances. Chronic inhalation may cause peripheral nerve disorders and central nervous system effects. Laboratory tests have resulted in mutagenic effects. May affect the developing fetus. Chronic exposure produces peripheral neuropathy with effects including muscular weakness, paresthesia, numbing of the hands, feet, legs, and arms, unsteadiness, and difficulty walking and standing. Repeated exposure may cause nervous system abnormalities with muscle weakness and damage, motor incoordination, and sensation disturbances. Persons with pre-existing skin disorders or eye problems or impaired respiratory function may be more susceptible to the effects of the substance.

p-Chlorobenzotrifluoride 98-56-6

Acute oral toxicity: LD50 Rat: >6,800 mg/kg
Acute inhalation toxicity: LC50 Rat: 4,479 ppm
Acute dermal toxicity: LD50 Rabbit: >2,700 mg/kg
Skin irritation: Rabbit: non-irritating
Eye irritation: Rabbit: non-irritating

A 28-day range-finding inhalation study was conducted in male and female Sprague-Dawley rats exposed to 0, 100, 250, 500, or 1000 ppm for 6 hr/day, 5 days/week. Clinical signs included increased activity at 250 ppm and above. Liver and kidney weights were increased. Microscopic changes in male kidneys stained positive for alpha-2-U globulin and the effects were considered not relevant to humans. Liver cell hypertrophy was seen at all exposures in males. Liver changes were consistent with clinical chemistry and PCBTF-blood level analysis and are believed to be an adaptive response, due to increased liver metabolism. Gavage studies in laboratory rodents for treatment periods of 14, 28, and 90 days have demonstrated significant liver and kidney toxicity at dose levels of 400 - 1000 mg/kg/day. Evidence of target organ toxicity included significant increases in relative liver and kidney weights, clinical chemistry values and histopathological findings. Renal toxicity which occurred only in male rats, was apparently due to "hyaline droplet" nephropathy and is therefore, highly unlikely to develop in man. The NOAEL's for all these studies range from 10 to 100 mg/kg/day. CNS effects were observed in rats exposed to PCBTF at or above 2822 ppm for 4 hours. A 90 day(13 week) rat inhalation toxicity and neurobehavioral study was conducted using exposures of 6 hrs/day, 5 days/week at concentrations of 0, 10, 50 and 250 ppm. There were no PCBTF-related macroscopic observations. Microscopically, PCBTF-related centrilobular hypertrophy was present only in the livers of males and females at the high dose (250 ppm) after 13-weeks of exposure. No centrilobular hypertrophy was observed at any level among recovery animals. There were no PCBTF-related effects on the nervous system as measured by a functional observation battery, muscular activity measurements and neuropathology. A NOEL of 50 ppm was established in this study for liver hepatocyte

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hypertrophy in male and female rats. If the hepatocyte hypertrophy observed is considered to be an adaptive response to PCBTF, the NOAEL for this study is 250ppm.

# Section 12: Ecological Information \_\_\_\_\_

ALIPHATIC HYDROCARBON \* 110-54-3

Ecotoxicity: Experimental studies involving Hexane show acute aquatic toxicity values of 2.1 mg/L and greater than 1000 mg/L. Environmental Fate: Persistence: Volatilization from soil surfaces is expected to be an important fate process. Hexane will be degraded in the atmosphere by reaction with hydroxyl radicals; the half-life of this reaction in air is estimated to be three days. Screening studies suggest that Hexane will undergo biodegradation in soil and water surfaces, but volatilization in expected be the predominant fate process in the environment. Hydrolysis is not expected to be an important environmental fate process. Bioaccumulation: An estimated bioconcentration factor (BCF) of 2300 and log Kow of 3.9 for Hexane suggest the potential for bioconcentration in aquatic organisms is high. Metabolites may partially bioaccumulate in the lipid bilayer of fish tissues. Mobility: Hexane is highly volatile and will partition rapidly in the air. When released into water, Hexane will be lost by volatilization and biodegradation. Hexane is expected to have high mobility in soils/sediments based on a Koc of 150. Volatilization from moist soil surfaces is expected to be an important fate process based on a Henry's law constant of 1.83 atm-m3/mole. Hexane may volatilize from dry surfaces based on its vapor pressure.

p-Chlorobenzotrifluoride 98-56-6

#### **AOUATIC ECOTOX DATA**

Fish:

LC50 (96 hr.) (Rainbow trout) 13.5 mg/L LC50 (96 hr.) (Bluegill sunfish) 12.0 mg/L MATC (31 day) (Fathead minnow) >0.54 <1.4 mg/L\* \*Triethylene glycol used as solvent carrier BCF (48 hr.) (Bluegill sunfish) 121.8 & 202.0 Invertebrates: LC50 (48 hr.) (Water flea) 12.4 mg/L MATC (21 day) (Water flea)  $> 0.03 < 0.05 \text{ mg/L}^*$ \*Acetone used as solvent carrier Plants:

IC50 (72 hr.) (Green & Blue-green algae) 500 mg/L TERRESTRIAL ECOTOX DATA

No data available

#### **ENVIRONMENTAL FATE DATA**

Biotic:

Biodegradation: inconclusive due to volatility

Abiotic:

Atmospheric lifetime: estimated to be 65.9 days for OH radical reaction

Log Kow 3.7 Koc 420 - 530

Water Sol. @ 23 C 29.1

p-Chlorobenzotrifluoride (PCBTF) will preferentially partition to the atmosphere, due to its high volatility. It has been estimated that 99.93% of a 100 Kg spill would end up in the atmosphere, while only 0.06% would partition to water (M. Garlanda, 1990). The aqueous solubility of PCBTF (29.1 mg/L) would also tend to limit its potential impact to exposed aquatic systems. PCBTF has exhibited significant toxicity to aquatic species under laboratory conditions, but is unlikely to exhibit a similar degree of acute toxicity under environmental conditions due to the aforementioned solubility and volatility issues. The moderate level of bioaccumulation measured in laboratory tests will also be subject to environmental mitigation due to PCBTF's physical/chemical properties. PCBTF should rapidly volatilize from dry and moist soils. Volatility, and relative environmental partitioning characteristics, make it unlikely that PCBTF represents a significant threat to aquatic or terrestrial environments.

ACETONE 67-64-1

#### **Acute and Prolonged Toxicity to Fish**

96 h LC 50 Fathead minnow (Pimephales promelas), : 8,733 - 9,482 mg/l Mortality

96 h LC 50 Bluegill (Lepomis macrochirus), : 8,300 mg/l Mortality

96 h LC 50 Rainbow trout, donaldson trout (Oncorhynchus mykiss), : 4,740 - 6,330 mg/l Mortality

#### **Acute Toxicity to Aquatic Invertebrates**

No data

#### **Environmental fate and pathways**

No data

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If applicable, IARL, NPT and OSHA carcinogens and chemicals subject to the reporting requirements of SARA Title III, Section 313 are identified in Section III with an "\*". Additional ecological information is Not Determined.

### Section 13: Disposal Information\_\_\_\_\_

Dispose of contents/container in accordance with local regulations.

WASTE DISPOSAL METHOD: Consult local authorities for proper waste disposal procedures. Empty de-pressurized containers can not be reused. Cans which are pressurized or contain liquid must be disposed of in a permitted waste management facility. Consult Federal, State, and local disposal authorities for approved procedures.

#### Section 14: Transportation Information\_\_\_\_\_

DOT Proper Shipping Name: UN1950

Aerosols, flammable, (each not exceeding 1L

capacity) 2.1

# Section 15: Regulatory Information\_\_\_\_\_

	CAS#	PEL	TLV
HYDROCARBON PROPELLANT	68476-86-8	NO DATA	NO DATA
ALIPHATIC HYDROCARBON *	110-54-3	500 PPM	50 PPM
p-Chlorobenzotrifluoride	98-56-6	NOT ESTABLISHED	NOT ESTABLISHED
POLYISOBUTYLENE	9003-27-4	NOT ESTABLISHED	NOT ESTABLISHED
LUBE OIL ADDITIVE CONTAINING	9003-29-6	NOT ESTABLISHED	NOT ESTABLISHED
BARIUM SULFONATE	CONFIDENTIAL	NOT ESTABLISHED	NOT ESTABLISHED
PETROLEUM HYDROCARBON BASE	64741-88-4	NOT ESTABLISHED	NOT ESTABLISHED
ACETONE	67-64-1	TWA 1000 PPM	TWA 750 PPM STEL 1000 PPM
HYDROTREATED LIGHT DISTILLATE	64742-47-8	NONE ESTABLISHED BY OSHA	NONE ESTABLISHED BY OSHA

State of California SAFE DRINKING WATER AND TOXIC ENFORCEMENT ACT OF 1986.

WARNING: IN ACCORDANCE WITH PROP 65, THIS PRODUCT CONTAINS A CHEMICAL(S) KNOWN TO THE STATE OF CALIFORNIA TO CAUSE CANCER, BIRTH DEFECTS AND OTHER REPRODUCTIVE HARM.

If applicable, IARL, NPT and OSHA carcinogens and chemicals subject to the reporting requirements of SARA Title III, Section 313 are identified above with an "\*"

All material components are listed in the U.S. TSCA inventory.

#### Section 16: Other Information

Consumer Product Safety Act Certification.

This product was evaluated by the Company listed above and is certified to be in compliance with the provisions of the Consumer Product Safety Act, the Federal Hazardous Substances Act, and the Poison Prevention Packaging Act, as applicable. This product was manufactured at the location identified on the SDS. The date of manufacture is stamped on the product container. No testing is required to certify compliance with the above mentioned regulation.

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The above information is based on the data of which we are aware and is believed to be correct as of the date hereof. Since the information contained herein may be applied under conditions beyond our control and with which we may be unfamiliar and since data made available subsequent to the date hereof may suggest modification of the information, we do not assume any responsibility for the results of its use. This information is furnished upon condition that the person receiving it shall make his own determination of the suitability of the material for his particular purpose.