

## 2-Aminobenzotrifluoride

Revision Date: 20-06-22 Revision Number: 5.1

#### SECTION 1 Identification of the substance / mixture and of the company / undertaking

**Product Name** 2-AMINOBENZOTRIFLUORIDE **Chemical Name** Not Available **Synonyms** C7-H6-F3-N; F3CC6H4NH2; o-toluidine, alpha, alpha, alpha-trifluoride; o-ABTF; o-aminobenzal fluoride; o-aminobenzotrifluoride; toluene, 2-amino-alpha,alpha,alpha-trifluoro-; o-(trifluoromethyl)aniline; o-(trifluoromethyl)benzenamine **Proper Shipping Name** 2-TRIFLUOROMETHYLANILINE **Chemical Formula**  $C_7H_6F_3N$ Other means of identification Not available **CAS Number** 88-17-5

#### Relevant identified uses of the substance or mixture and uses advised against:

7	
Relevant identified uses	Intermediate.

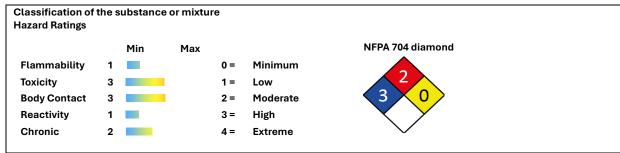
#### Details of the manufacturer or supplier of the safety data sheet:

Registered company name	Cohizon Life Sciences Limited	
Address	Plot No. 6102/3, 6117-19, 5809-10, GIDC, Ankleshwar Gujarat 393002 India	
Telephone	Not Available	
Fax	Not Available	
Website	Not Available	
Email	Not Available	

#### **Emergency telephone number:**

Ī	Association / Organisation	Cohizon Life Sciences Limited
	Emergency telephone number(s)	(+91) 7046611150/51
	Other emergency telephone number(s)	(+91) 6357684904

#### **SECTION 2 Hazards identification**



Note: The hazard category numbers found in GHS classification in section 2 of this SDSs are NOT to be used to fill in the NFPA 704 diamond. Blue = Health Red = Fire Yellow = Reactivity White = Special (Oxidizer or water reactive substances

	Flammable Liquids Category 4, Acute Toxicity (Oral, Dermal and Inhalation) Category 3,
	Skin Corrosion/Irritation Category 2, Serious Eye Damage/Eye Irritation Category 1,
Classification Specific Target Organ Toxicity - Single Exposure (Respiratory Tract Irritation	
	Specific Target Organ Toxicity - Repeated Exposure Category 2, Hazardous to the
	Aquatic Environment Long-Term Hazard Category 2

## Label elements

Hazard pictogram(s)		
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Signal word	Danger



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## Hazard statement(s)

H227	Combustible liquid.	
H301+H311+H331	Toxic if swallowed, in contact with skin or if inhaled.	
H315	Causes skin irritation.	
H318	Causes serious eye damage.	
H335	May cause respiratory irritation.	
H373	May cause damage to organs through prolonged or repeated exposure.	
H411	Toxic to aquatic life with long lasting effects.	

#### Precautionary statement(s) Prevention

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P210	Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No	
	smoking.	
P260	Do not breathe mist/vapours/spray.	
P264	Wash all exposed external body areas thoroughly after handling.	
P270	Do not eat, drink or smoke when using this product.	
P271	Use only outdoors or in a well-ventilated area.	
P280	Wear protective gloves, protective clothing, eye protection and face protection.	
P273	Avoid release to the environment.	

#### Precautionary statement(s) Response

	(o)opoo
P301+P310	IF SWALLOWED: Immediately call a POISON CENTER/doctor/physician/first aider.
P302+P352	IF ON SKIN: Wash with plenty of water.
P304+P340	IF INHALED: Remove person to fresh air and keep comfortable for breathing.
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if
	present and easy to do. Continue rinsing.
P330	Rinse mouth.
P361+P364	Take off immediately all contaminated clothing and wash it before reuse.
P370+P378	In case of fire: Use alcohol resistant foam or fine spray/water fog to extinguish.
P391	Collect spillage.
P332+P313	If skin irritation occurs: Get medical advice/attention.

#### Precautionary statement(s) Storage

P405	Store locked up.
P403+P233	Store in a well-ventilated place. Keep container tightly closed.

#### Precautionary statement(s) Disposal

P501	Dispose of contents/container to authorised hazardous or special waste collection
	point in accordance with any local regulation

### SECTION 3 Composition / information on ingredients

#### Substances

Cas No.	%[weight]	Name
88-17-5	>98	<u>2-Aminobenzotrifluride</u>

#### Mixtures

See Section above for composition of substances

## **SECTION 4 First aid measures**

## Description of first aid measures

	If this product comes in contact with the eyes:
	<ul> <li>Immediately hold eyelids apart and flush the eye continuously with running water.</li> </ul>
Eye Contact	<ul> <li>Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids.</li> </ul>
	<ul> <li>Continue flushing until advised to stop by the Poisons         Information Centre or a doctor, or for at least 15 minutes.     </li> <li>Transport to hospital or doctor without delay.</li> </ul>



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	Removal of contact lenses after an eye injury should only be undertaken by
	skilled personnel
	If skin or hair contact occurs:
	<ul> <li>Quickly but gently, wipe material off skin with a dry, clean cloth.</li> </ul>
Skin Contact	Immediately remove all contaminated clothing, including footwear.
	<ul> <li>Wash skin and hair with running water. Continue flushing with water until</li> </ul>
	advised to stop by the Poisons Information Centre.
	Transport to hospital, or doctor.
	If fumes or combustion products are inhaled remove from contaminated
	area.
	<ul> <li>Lay patient down. Keep warm and rested.</li> <li>Prostheses such as false teeth, which may block airway, should be removed,</li> </ul>
Inhalation	where possible, prior to initiating first aid procedures.
	<ul> <li>Apply artificial respiration if not breathing, preferably with a demand valve</li> </ul>
	resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR
	if necessary.
	Transport to hospital, or doctor, without delay
	<ul> <li>IF SWALLOWED, REFER FOR MEDICAL ATTENTION, WHERE POSSIBLE, WITHOUT DELAY.</li> </ul>
	<ul> <li>For advice, contact a Poisons Information Centre or a doctor</li> </ul>
	<ul> <li>Urgent hospital treatment is likely to be needed</li> </ul>
	<ul> <li>In the meantime, qualified first-aid personnel should treat the patient</li> </ul>
	following observation and employing supportive measures as indicated by the patient's condition.
	<ul> <li>If the services of a medical officer or medical doctor are readily available,</li> </ul>
	the patient should be placed in his/her care and a copy of the SDS should
Ingestion	be provided. Further action will be the responsibility of the medical
	specialist.
	<ul> <li>If medical attention is not available on the worksite or surroundings send the patient to a hospital together with a copy of the SDS.</li> </ul>
	Where medical attention is not immediately available or where the patient is
	more than 15 minutes from a hospital or unless instructed otherwise:
	INDUCE vomiting with fingers down the back of the throat, ONLY IF
	CONSCIOUS. Lean patient forward or place on left side (head-down
	position, if possible) to maintain open airway and prevent aspiration.
	NOTE: Wear a protective glove when inducing vomiting by mechanical means

### Indication of any immediate medical attention and special treatment needed

The material may induce methaemoglobinaemia following exposure.

- Initial attention should be directed at oxygen delivery and assisted ventilation if necessary. Hyperbaric oxygen has not demonstrated substantial benefits.
- Hypotension should respond to Trendelenburg's position and intravenous fluids; otherwise dopamine may be needed.
- Symptomatic patients with methaemoglobin levels over 30% should receive methylene blue. (Cyanosis, alone, is not an indication for treatment). The usual dose is 1-2 mg/kg of a 1% solution (10 mg/ml) IV over 50 minutes; repeat, using the same dose, if symptoms of hypoxia fail to subside within 1 hour.
- Thorough cleansing of the entire contaminated area of the body, including the scalp and nails, is of utmost importance. BIOLOGICAL EXPOSURE INDEX - BEI
- These represent the determinants observed in specimens collected from a healthy worker exposed at the Exposure Standard (ES or TLV):

Determinant Index Sampling Time Comment
1. Methemoglobin in blood 1.5% of hemoglobin During or end of shift B, NS, SQ

B: Background levels occur in specimens collected from subjects  ${\bf NOT}$  exposed

NS: Non-specific determinant; also observed after exposure to other materials

SQ: Semi-quantitative determinant - Interpretation may be ambiguous; should be used as a screening test or confirmatory test



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## SECTION 5 Firefighting measures

#### Extinguishing media

- Foam.
- Dry chemical powder
- BCF (where regulations permit).
- Carbon dioxide.
- Water spray or fog Large fires only

#### Special hazards arising from the substrate or mixture

Fire Incompatibility	Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine	
	bleaches, pool chlorine etc. as ignition may result	

Advice for firefighters			
	<ul> <li>Alert Fire Brigade and tell them location and nature of hazard.</li> </ul>		
	<ul> <li>Wear breathing apparatus plus protective gloves</li> </ul>		
	<ul> <li>Prevent, by any means available, spillage from entering drains or water courses</li> </ul>		
Fire Fighting	<ul> <li>Use water delivered as a fine spray to control fire and cool adjacent area</li> </ul>		
	DO NOT approach containers suspected to be hot		
	<ul> <li>Cool fire exposed containers with water spray from a protected location</li> </ul>		
	<ul> <li>If safe to do so, remove containers from path of fire</li> </ul>		
	<ul> <li>Equipment should be thoroughly decontaminated after use</li> </ul>		
	Combustible.		
	<ul> <li>Slight fire hazard when exposed to heat or flame.</li> </ul>		
	Heating may cause expansion or		
	decomposition leading to violent rupture of		
	containers.		
	On combustion, may emit toxic fumes of		
	carbon monoxide (CO).		
Fire/Explosion Hazard	May emit acrid smoke.		
	Mists containing combustible materials may be explosive		
	Combustion products include.		
	carbon monoxide (CO)		
	carbon dioxide (CO2)		
	<ul> <li>nitrogen oxides (NOx)</li> </ul>		
	<ul> <li>other pyrolysis products typical of burning organic material</li> </ul>		
	May emit poisonous fumes		
	May emit corrosive fumes		

### SECTION 6 Accidental release measures

### Personal precautions, protective equipment and emergency procedures

See section 8

#### **Environmental precautions**

See section 12

#### Methods and material for containment and cleaning up

Minor Spills	Remove all ignition sources.
	Clean up all spills immediately.
	Avoid contact with skin and eyes.
	<ul> <li>Control personal contact with the substance, by using protective equipment.</li> </ul>
	<ul> <li>Use dry clean up procedures and avoid generating dust.</li> </ul>
	<ul> <li>Place in a suitable, labelled container for waste disposal.</li> </ul>
	Clear area of personnel and move upwind.
Major Spills	<ul> <li>Alert fire Brigade and tell them location and nature of hazard.</li> </ul>
	<ul> <li>Control personal contact by wearing protective clothing.</li> </ul>



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	Prevent, by any means available, spillage from entering drains or water
	courses.
	Stop leak if safe to do so.
	Contain spill with sand, earth or vermiculite.
	Collect recoverable product into labelled containers for recycling.
	Neutralise/decontaminate residue (see Section 13 for specific agent).
	Collect solid residues and seal in labelled drums for disposal.
•	Wash area and prevent runoff into drains.
	After clean up operations, decontaminate and launder all protective clothing
	and equipment before storing and re-using.
	If contamination of drains or waterways occurs, advise emergency services

Personal Protective Equipment advice is contained in Section 8 of the SDS

## SECTION 7 Handling and storage

Precautions for safe handl	ing		
	Avoid all personal contact, including inhalation.		
	Wear protective clothing when risk of exposure occurs		
	Use in a well-ventilated area		
	<ul> <li>Prevent concentration in hollows and sumps.</li> </ul>		
	<ul> <li>DO NOT enter confined spaces until the atmosphere has been checked.</li> </ul>		
Safe Handling	<ul> <li>DO NOT allow material to contact humans, exposed food or food utensils</li> </ul>		
	Avoid contact with incompatible materials		
	When handling, DO NOT eat, drink or smoke.		
	Keep containers securely sealed when not in use		
	Avoid physical damage to containers		
	<ul> <li>Always wash hands with soap and water after handling</li> </ul>		
	Work clothes should be laundered separately. Launder contaminated clothing		
	before re-use		
	Use good occupational work practice		
	<ul> <li>Observe manufacturer's storage and handling recommendations contained within this SDS.</li> </ul>		
	Atmosphere should be regularly checked against established exposure		
	standards to ensure safe working conditions are maintained.		
	DO NOT allow clothing wet with material to stay in contact with skin		
	Store in original containers.		
	Keep containers securely sealed.		
	Store in a cool, dry, well-ventilated area.		
Other information	Store away from incompatible materials and foodstuff containers.		
	<ul> <li>Protect containers against physical damage and check regularly for leaks.</li> </ul>		
	Observe manufacturer's storage and handling recommendations contained within this SDS		

### Conditions for safe storage, including any incompatibilities

Contactions for date of orange,	, including any incompatibilities
	Lined metal can, lined metal pail/ can.
	Plastic pail.
	Polyliner drum.
	Packing as recommended by manufacturer.
	<ul> <li>Check all containers are clearly labelled and free from leaks. For low viscosity materials</li> </ul>
	<ul> <li>Drums and jerricans must be of the non-removable head type.</li> </ul>
Suitable container	<ul> <li>Where a can is to be used as an inner package, the can must have a screwed enclosure.</li> </ul>
	<ul> <li>For materials with a viscosity of at least 2680 cSt. (23 deg. C) and solids (between 15 C deg. and 40 deg C.)</li> </ul>
	Removable head packaging;
	<ul> <li>Cans with friction closures and low pressure tubes and cartridges may be used.</li> </ul>
	Where combination packages are used, and the inner packages are of glass,



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	there must be sufficient inert cushioning material in contact with inner and outer packages.	
	<ul> <li>In addition, where inner packaging's are glass and contain liquids of packing group I and II there must be sufficient inert absorbent to absorb any spillage *.</li> </ul>	
	<ul> <li>unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic.</li> </ul>	
	<ul> <li>Many arylamines (aromatic amines such as aniline, N-ethylaniline, o-toluidine, xylidine etc. and their mixtures) are hypergolic (ignite spontaneously) with red fuming nitric acid. When the amines are dissolved in triethylamine, ignition occurs at -60 deg. C or less.</li> </ul>	
Storage incompatibility	<ul> <li>Various metal oxides and their salts may promote ignition of amine-red fuming nitric acid systems. Soluble materials such as copper(I) oxide, ammonium metavanadate are effective; insoluble materials such as copper(II) oxide, iron(II) oxide, potassium dichromate are also effective.</li> </ul>	
	<ul> <li>Avoid oxidising agents, acids, acid chlorides, acid anhydrides, chloroformates.</li> </ul>	

#### **SECTION 8 Exposure controls / personal protection**

#### **Control Parameters**

Occupational Exposure Limits (OEL)

#### INGREDIENT DATA

Not Available

Ingredient	Original IDLH	Revised IDLH
2-aminobenzotrifluoride	Not Available	Not Available

#### **Exposure controls**

Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection.

The basic types of engineering controls are:

Process controls which involve changing the way a job activity or process is done to reduce the risk.

Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation system must match the particular process and chemical or contaminant in use. Employers may need to use multiple types of controls to prevent employee overexposure.

# Appropriate engineering controls

- Local exhaust ventilation usually required. If risk of overexposure exists, wear approved respirator. Correct fit is essential to obtain adequate protection. Supplied-air type respirator may be required in special circumstances. Correct fit is essential to ensure adequate protection.
- An approved self contained breathing apparatus (SCBA) may be required in some situations.
- Provide adequate ventilation in warehouse or closed storage area.
   Air contaminants generated in the workplace possess varying
   "escape" velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contaminant:

Type of Contaminant	Air Speed
solvent, vapours, degreasing etc., evaporating from tank (in still air).	.25-0.5 m/s (50- 100 f/min.)
aerosols, fumes from pouring operations, intermittent container filling, low speed conveyer transfers, welding, spray drift, plating acid fumes, pickling (released at low velocity into zone of active generation)	0.5-1 m/s (100- 200 f/min.)



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	conveyer loading, crusher dusts, gas discharge (active generation into zone of rapid air motion)		1-2.5 m/s (200-500 ft/min)
			,
	grinding, abrasive blasting, tumbling, high spee	d wheel	2.5-10 m/s (500-
	generated dusts (released at high initial velocit	y into zone of	2000 ft/min)
	very high rapid air motion		
	Which each range the appropriate value depe		
	Lower end of the range	Upper end of th	
	Room air currents minimal or favourable to capture	1. Disturbing ro	oom air currents
	Contaminants of low toxicity or of     Contaminants of high toxicity		nts of high toxicity
	nuisance value only 3. Intermittent, low production	3: High produc	tion, heavy use
	4. Large hood or large air mass in motion		local control only
	Simple theory shows that air velocity falls rapidly with distance away from the		
	opening of a simple extraction pipe. Velocity gedistance from the extraction point (in simple calextraction point should be adjusted, according the contaminating source. The air velocity at the beal minimum of 4-10 m/s (800-2000 ft/min) for generated 2 metres distant from the extraction considerations, producing performance deficits	ases). Therefore, the air speed at the gly, after reference to distance from ne extraction fan, for example, should or extraction of crusher dusts n point. Other mechanical	
	make it essential that theoretical air velocities are multiplied by factors of 10 or more		
Individual protection	when extraction systems are installed or used		
measures, such as personal			
protective			
equipment			
equipment			
Eye and face protection	<ul> <li>Safety glasses with side shields.</li> <li>Chemical goggles. [AS/NZS 1337.1, EN166 or national equivalent]</li> <li>Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants. A written policy document, describing the wearing of lenses or restrictions on use, should be created for each workplace or task. This should include a review of lens absorption and adsorption for the class of chemicals in use and an account of injury experience.</li> <li>Medical and first-aid personnel should be trained in their removal, and suitable equipment should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove contact lens as soon as practicable. Lens should be removed at the first signs of eye redness or irritation - lens should be removed in a clean environment only after workers</li> <li>have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59</li> </ul>		
Skin protection	See Hand protection below		
	NOTE:  • Wear chemical protective gloves, e.g. PVC. • Wear safety footwear or safety gumboots, e.g. Rubber  The selection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer.  Where the chemical is a preparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application.  The exact break through time for substances has to be obtained from the manufacturer of the protective gloves and has to be observed when making a final choice.  Personal hygiene is a key element of effective hand care. Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturizer is recommended.  Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include:  • frequency and duration of contact,		
Hands/feet protection	Where the chemical is a preparation of several sthe glove material can not be calculated in advance checked prior to the application.  The exact break through time for substances ha manufacturer of the protective gloves and has tochoice.  Personal hygiene is a key element of effective hon clean hands. After using gloves, hands shou Application of a non-perfumed moisturizer is resultability and durability of glove type is dependent the selection of gloves include:	substances, the ance and has the as to be obtained to be observed wand care. Gloveld be washed ar commended.	e resistance of erefore to be d from the when making a final s must only be worn and dried thoroughly.



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•	glove thickness and
•	dexterity

Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739, AS/NZS 2161.1 or national equivalent).

- When prolonged or frequently repeated contact may occur, a glove with a protection class of 5 or higher (breakthrough time greater than 240 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended.
- When only brief contact is expected, a glove with a protection class of 3 or higher (breakthrough time greater than 60 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended.
- Some glove polymer types are less affected by movement and this should be taken into account when considering gloves for long-term use.
- Contaminated gloves should be replaced.

As defined in ASTM F-739-96 in any application, gloves are rated as:

- Excellent when breakthrough time > 480 min
- Good when breakthrough time > 20 min
- Fair when breakthrough time < 20 min
- · Poor when glove material degrades

For general applications, gloves with thickness typically greater than 0.35 mm, are recommended.

It should be emphasized that glove thickness is not necessarily a good predictor of glove resistance to a specific chemical, as the permeation efficiency of the glove will be dependent on the exact composition of the glove material. Therefore, glove selection should also be based on consideration of the task requirements and knowledge of breakthrough times.

Glove thickness may also vary depending on the glove manufacturer, the glove type and the glove model. Therefore, the manufacturers' technical data should always be taken into account to ensure selection of the most appropriate glove for the task. Note: Depending on the activity being conducted, gloves of varying thickness may be required for specific tasks. For example:

- Thinner gloves (down to 0.1 mm or less) may be required where a high
  degree of manual dexterity is needed. However, these gloves are only
  likely to give short duration protection and would normally be just for
  single use applications, then disposed of.
- Thicker gloves (up to 3 mm or more) may be required where there is a mechanical (as well as a chemical) risk i.e. where there is abrasion or puncture potential

Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturiser is recommended.

Neoprene gloves

Body protection	See Other protection below
	Overalls
Other protection	Eye wash unit
Other protection	Barrier Cream
	Skin Cleansing cream

#### Respiratory protection

Type -P Filter of sufficient capacity. (AS/NZS 1716 & 1715, EN 143:2000 & 149:2001, ANSI Z88 or national equivalent Selection of the Class and Type of respirator will depend upon the level of breathing zone contaminant and the chemical nature of the contaminant. Protection Factors (defined as the ratio of contaminant outside and inside the mask) may also be important.

Required minimum protection factor	Maximum gas/vapour concentration present in air p.p.m. (by volume)	Half-face Respirator	Full-Face Respirator	
up to 10	1000	A-AUS / Class1	-	



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up to 50	1000	-	A-AUS / Class 1
up to 50	5000	Airline *	-
up to 100	5000	-	A-2
up to 100	10000	-	A-3
100+			Airline**

Continuous Flow \*\* - Continuous-flow or positive pressure demand

A(All classes) = Organic vapours, B AUS or B1 = Acid gasses, B2 = Acid gas or hydrogen cyanide(HCN), B3 = Acid gas or hydrogen cyanide(HCN), E = Sulfur dioxide(SO2), G = Agricultural chemicals, K = Ammonia(NH3), Hg = Mercury, NO = Oxides of nitrogen, MB = Methyl bromide, AX = Low boiling point organic compounds(below 65 degC)

- Cartridge respirators should never be used for emergency ingress or in areas of unknown vapor concentrations or oxygen content.
- The wearer must be warned to leave the contaminated area immediately on detecting any odors through the respirator. The odor may indicate that the mask is not functioning properly, that the vapor concentration is too high, or that the mask is not properly fitted. Because of these limitations, only restricted use of cartridge respirators is considered appropriate.
- Cartridge performance is affected by humidity. Cartridges should be changed after 2 hr of continuous use unless it is determined that the humidity is less than 75%, in which case, cartridges can be used for 4 hr. Used cartridges should be discarded daily, regardless of the length of time used

#### SECTION 9 Physical and chemical properties

Information on basic physical and chemical properties

Appearance	Off-white crystalline	powder; does not mix well with wat	ter.
Physical state	Liquid	Relative density (Water = 1)	1.296
Odour	Not Available	Partition coefficient noctanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Applicable	Decomposition temperature (°C)	Not Available
Melting point / freezing point (°C)	-3028°C	Viscosity (cSt)	Not Applicable
Initial boiling point and boiling range (°C)	170-173℃	Molecular weight (g/mol)	161.13
Flash point (°C)	67°C	Taste	Not Available
Evaporation rate	Not Applicable	<b>Explosive properties</b>	Not Available
Flammability	Not Applicable	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Applicable
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Negligible	Gas group	Not Available
Solubility in water	Partly miscible	pH as a solution (1%)	Not Applicable
Vapour density (Air = 1)	>1	VOC g/L	Not Applicable
Heat of Combustion (kJ/g)	Not Available	Ignition Distance (cm)	Not Available
Flame Height (cm)	Not Available	Flame Duration (s)	Not Available
Enclosed Space Ignition Time		Enclosed Space	
Equivalent (s/m3)	Not Available	Ignition Deflagration	Not Available
		Density (g/m3)	



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## SECTION 10 Stability and reactivity

Reactivity	See section 7
Chemical stability	<ul><li>Unstable in the presence of incompatible materials</li><li>Product is considered stable</li></ul>
	Hazardous polymerisation will not occur
Possibility of hazardous reactions	See section 7
Conditions to avoid	See section 7
Incompatible materials	See section 7
Hazardous decomposition products	See section 5

## SECTION 11 Toxicological information

### Information on toxicological effects

miormation on toxicotogical on	
a) Acute Toxicity	There is sufficient evidence to classify this material as acutely toxic.
b) Skin Irritation/Corrosion	There is sufficient evidence to classify this material as skin corrosive or irritating.
c) Serious Eye Damage/Irritation	There is sufficient evidence to classify this material as eye damaging or irritating
d) Respiratory or Skin sensitisation	Based on available data, the classification criteria are not met.
e) Mutagenicity	Based on available data, the classification criteria are not met.
f) Carcinogenicity	Based on available data, the classification criteria are not met.
g) Reproductivity	Based on available data, the classification criteria are not met.
h) STOT - Single Exposure	There is sufficient evidence to classify this material as toxic to specific organs through single exposure
i) STOT - Repeated Exposure	There is sufficient evidence to classify this material as toxic to specific organs through repeated exposure
j) Aspiration Hazard	Based on available data, the classification criteria are not met.

Inhaled	<ul> <li>Inhalation of vapours, aerosols (mists, fumes) or dusts, generated by the material during the course of normal handling, may be harmful</li> <li>The material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage.</li> <li>Inhalation of vapours may cause drowsiness and dizziness. This may be accompanied by sleepiness, reduced alertness, loss of reflexes, lack of co-ordination, and vertigo.</li> <li>Inhalation hazard is increased at higher temperatures.</li> <li>Inhalation of high concentrations of gas/vapour causes lung irritation with coughing and nausea, central nervous depression with headache and dizziness, slowing of reflexes, fatigue and inco-ordination.</li> </ul>
	<ul> <li>Toxic effects may result from the accidental ingestion of the material; animal experiments indicate that ingestion of less than 40 gram may be fatal or may produce serious damage to the health of the individual.</li> <li>The substance and/or its metabolites may bind to haemoglobin inhibiting normal uptake of oxygen. This condition, known as "methaemoglobinemia", is a form of oxygen starvation (anoxia).</li> </ul>
Ingestion	<ul> <li>Symptoms include cyanosis (a bluish discolouration skin and mucous membranes) and breathing difficulties. Symptoms may not be evident until several hours after exposure.</li> </ul>
	<ul> <li>At about 15% concentration of blood methaemoglobin there is observable cyanosis of the lips, nose and earlobes. Symptoms may be absent although euphoria, flushed face and headache are commonly experienced. At 25-40%, cyanosis is marked but little disability occurs other than that produced on physical exertion. At 40-60%, symptoms include weakness, dizziness, lightheadedness, increasingly severe</li> </ul>



## 2-Aminobenzotrifluoride

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	headache, ataxia, rapid shallow respiration, drowsiness, nausea,
	vomiting, confusion, lethargy and stupor. Above 60% symptoms include
	dyspnea, respiratory depression, tachycardia or bradycardia, and
	convulsions. Levels exceeding 70% may be fatal.
	<ul> <li>Aromatic nitriles, unlike aliphatic nitriles, do not appear to liberate</li> </ul>
	cyanide within the body
	Skin contact with the material may produce toxic effects; systemic effects may
	result following absorption
	This material can cause inflammation of the skin on contact in some persons
	The material may accentuate any pre-existing dermatitis condition
Skin Contact	Open cuts, abraded or irritated skin should not be exposed to this material
	Entry into the blood-stream, through, for example, cuts, abrasions or
	lesions, may produce systemic injury with harmful effects
	Examine the skin prior to the use of the material and ensure that any
	external damage is suitably protected
Eye	If applied to the eyes, this material causes severe eye damage
	<ul> <li>Chronic exposure of the related product, 3-aminobenzotrifluoride produced dystrophic changes in the brain cells, liver, kidney and lungs of rats.</li> </ul>
	<ul> <li>Repeated or long-term occupational exposure is likely to produce cumulative health effects involving organs or biochemical systems.</li> </ul>
Chronic	<ul> <li>Long-term exposure to respiratory irritants may result in airways disease,</li> </ul>
	involving difficulty breathing and related whole-body problems.
	Most arylamines are very toxic to the blood cell-forming system, and they
	produce methaemoglobinaemia in humans. High doses
	<ul> <li>congest the spleen and then cause formation of sarcomas (a type of malignant tumour)</li> </ul>

2-AMINOBENZOTRIFLUORIDE	TOXICITY	IRRITATION
	Not Available	Not Available
Legend:	1. Value obtained from Eur	ope ECHA Registered Substances - Acute toxicity
	2. Value obtained from ma	nufacturer's SDS. Unless otherwise specified data
	extracted from RTECS - Re	gister of Toxic Effect of chemical Substances

2-AMINOBENZOTRIFLUORIDE	Asthma-like symptoms may continue for months or even years after exposure to the material ends. This may be due to a non- allergic condition known as reactive airways dysfunction syndrome (RADS) which can occur after exposure to high levels of highly irritating compound. Main criteria for diagnosing RADS include the absence of previous airways disease in a non-atopic individual, with sudden onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. Other criteria for diagnosis of RADS include a reversible airflow pattern on lung function tests, moderate to severe bronchial hyperreactivity on methacholine challenge testing, and the lack of minimal lymphocytic inflammation, without eosinophilia. RADS (or asthma) following an irritating inhelation is an infraguent disorder with rates related to the concentration of and
	inflammation, without eosinophilia. RADS (or asthma) following an irritating inhalation is an infrequent disorder with rates related to the concentration of and duration of exposure to the irritating substance. On the other hand, industrial
	bronchitis is a disorder that occurs as a result of exposure due to high
	concentrations of irritating substance (often particles) and is completely reversible
	after exposure ceases. The disorder is characterized by difficulty breathing, cough
	and mucus production.
	No significant acute toxicological data identified in literature search



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Acute Toxicity	~	Carcinogenicity	×
Skin Irritation/Corrosion	•	Reproductivity	×
Serious Eye Damage/Irritation	•	STOT - Single Exposure	•
Respiratory or Skin sensitisation	×	STOT - Repeated Exposure	•
Mutagenicity	×	Aspiration Hazard	×

## SECTION 12 Ecological information

#### Toxicity

A AMINORENZOTRIELHORIDE	End Point	Test Duration (hr.)	Species	Value	Source
2-AMINOBENZOTRIFLUORIDE	EC50(ECx)	24h	Crustacea	10mg/l	2
	LC50	96h	Fish	>25<35mg/l	2
Legand:	Ecotoxicologic 4. US EPA, Eco Assessment Da	Extracted from 1. IUCLID Toxicity Data 2. Europe ECHA Registered Substances - Ecotoxicological Information - Aquatic Toxicity 4. US EPA, Ecotox database - Aquatic Toxicity Data 5. ECETOC Aquatic Hazard Assessment Data 6. NITE (Japan) - Bioconcentration Data 7. METI (Japan) - Bioconcentration Data 8. Vendor Data			

log P 2.41 Henry's Law Constant: 1.65E-005 atm-m3/mole, 25 C Atmospheric OH Rate Constant: 2.164E-011 cm3/molecule-sec Toxic to aquatic organisms.

Do NOT allow product to come in contact with surface waters or to intertidal areas below the mean high water mark. Do not contaminate water when cleaning equipment or disposing of equipment wash-waters

Wastes resulting from use of the product must be disposed of on site or at approved waste sites. May cause long-term adverse effects in the aquatic environment.

For Arylamines (Aromatic Amines):

Aquatic Fate - Arylamines, particularly aromatic amines, irreversibly bind with humic substances present in most natural waters. The estimated half-life of aromatic amines in water is approximetly 100 days.

Ecotoxicity: Anilines, benzidines and toluidines are of environmental concern. Anilines and benzidines are both acutely toxic and toxic depending on the specific aquatic species (except algae). Toluidines represent a similar concern, It has been speculated that aqueous solutions of aromatic amines can be oxidized by organic radicals. The estimated half-life of aromatic amines in water is approximately 100 days.

**DO NOT** discharge into sewer or waterways

### Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
2-Aminobenzotrifluoride	High	High

## Bio accumulative potential

Ingredient	Bioaccumulation
2-Aminobenzotrifluoride	Low (Log KOW = 2.41)

## Mobility in soil

Ingredient	Mobility
2-Aminobenzotrifluoride	LOW (Log KOC = 326.2)



#### 2-Aminobenzotrifluoride

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### SECTION 13 Disposal considerations

#### Waste treatment methods

#### **Product / Packaging** Containers may still present a chemical hazard/ danger when empty disposal Return to supplier for reuse/recycling if possible. Otherwise: If container cannot be cleaned sufficiently well to ensure that residuals do not remain or if the container cannot be used to store the same product, then puncture containers, to prevent re-use, and bury at an authorised Where possible retain label warnings and SDS and observe all notices pertaining to the product Legislation addressing waste disposal requirements may differ by country, state and/ or territory. Each user must refer to laws operating in their area. In some areas, certain wastes must be tracked. A Hierarchy of Controls seems to be common - the user should investigate: Reduction Reuse Recycling Disposal (if all else fails) This material may be recycled if unused, or if it has not been contaminated so as to make it unsuitable for its intended use. Shelf life considerations should also be applied in making decisions of this type. Note that properties of a material may change in use, and recycling or reuse may not always be appropriate. In most instances the supplier of the material should be consulted. DO NOT allow wash water from cleaning or process equipment to enter drains It may be necessary to collect all wash water for treatment before disposal In all cases disposal to sewer may be subject to local laws and regulations and these should be considered first. Where in doubt contact the responsible authority Recycle wherever possible or Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified. Bury or incinerate residue at an approved site.

### **SECTION 14 Transport information**

Labels Required	
	6
Marine Pollutant	

Recycle containers if possible, or dispose of in an authorised landfill.

#### Land transport (UN)

14.1. UN number or ID number	2942	
14.2. UN proper shipping name	2-TRIFLUOROMETHYLANILINE	
14.3. Transport hazard	Class	6.1
class(es)	Subsidiary Hazard	Not Applicable
14.4. Packing group	· ·	
14.5. Environmental hazard	Environmentally hazardous	
	Special Provisions	Not Applicable



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14.6. Special precautions for	Limited Ouality	51
user	Limited Quality	36

#### Air transport (ICAO-IATA / DGR)

14.1. UN number	2942		
14.2. UN proper shipping name	2-Trifluoromethylaniline		
14.3. Transport hazard	ICAO/IATA Class	6.1	
class(es)	ICAO / IATA Subsidiary Hazard	Not Applicable	
	ERG Code	6L	
14.4. Packing group	III		
14.5. Environmental hazard	Environmentally hazardous		
14.6. Special precautions for	Special provisions Not Applicable		Not Applicable
user	Cargo Only Packing Instructions 663		663
	Cargo Only Maximum Qty / Pack 220 L		220 L
	Passenger and Cargo Packing Instructions 655		655
	Passenger and Cargo Maximum Qty / Pack 60 L		60 L
	Passenger and Cargo Limited Quantity Packing Y642		Y642
	Instructions		
	Passenger and Cargo Lim	Passenger and Cargo Limited Maximum Qty / Pack 2 L	

#### Sea transport (IMDG-Code/ GGVSee)

oca danisport (ii ibo code) covoce)		
14.1. UN number	2942	
14.2. UN proper shipping name	2-TRIFLUOROMETHYLANILINE	
14.3. Transport hazard	IMDG Class 6.1	
class(es)	IMDG Subsidiary	Not Applicable
	Hazard	
14.4. Packing group	III	
14.5 Environmental hazard	Marine Pollutant	
14.6. Special precautions for	EMS Number	F-A,S-A
user	Special provisions	Not Applicable
	Limited Quantities	5 L

### 14.7 Maritime transport in bulk according to IMO instruments

#### 14.7.1 Transport in bulk in accordance to Annex II of MARPOL and the IBC Code

Product Name	Group
2-Aminobenzotrifluoride	Not Available

### 14.7.2 Transport in bulk in accordance with MARPOL Annex V and the IMSBC Code

Product Name	Ship Type
2-Aminobenzotrifluoride	Not Available

#### 14.7.3 Transport in bulk in accordance with the IGC Code

Product Name	Ship Type
2-Aminobenzotrifluoride	Not Available

## SECTION 15 Regulatory information

Safety, health and environmental regulations / legislation specific for the substance or mixture

#### 2-Aminobenzotrifluoride is found on the following lists

Not Applicable

### Additional Regulatory Information

Not Applicable

#### National Inventory Status

National Inventory	Status
Australia - AIIC / Australia Non-	No (2-Aminobenzotrifluoride)
Industrial Use	
Canada - DSL	No (2-Aminobenzotrifluoride)



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Canada - NDSL	Yes	
China - IECSC	Yes	
Europe - EINEC / ELINCS / NLP	Yes	
Japan - ENCS	Yes	
Korea - KECI	No (2-Aminobenzotrifluoride)	
New Zealand - NZIoC	Yes	
Philippines - PICCS	Yes	
USA - TSCA	All chemical substances in this product have been designated as TSCA Inventory	
	'Active'	
Taiwan - TCSI	Yes	
Mexico - INSQ	Yes	
Vietnam - NCI	Yes	
Russia - FBEPH	No (2-Aminobenzotrifluoride)	
	Yes = All CAS declared ingredients are on the inventory	
Legend:	No = One or more of the CAS listed ingredients are not on the inventory. These	
2050	ingredients may be exempt or will require registration.	

#### **SECTION 16 Other information**

Revision Date	20/06/2022
Initial Date	12/05/2005

#### **SDS Revision Summary**

Revision	Date of Update	Section Update
5.1	20/06/2022	Expiration. Review and Update

#### Other information

Classification of the preparation and its individual components has drawn on official and authoritative sources.

The SDS is a Hazard Communication tool and should be used to assist in the Risk Assessment. Many factors determine whether the reported Hazards are Risks in the workplace or other settings. Risks may be determined by reference to Exposures Scenarios. Scale of use, frequency of use and current or available engineering controls must be considered.

#### **Definitions and abbreviations**

PC - TWA: Permissible Concentration-Time Weighted Average

PC - STEL: Permissible Concentration-Short Term Exposure Limit

IARC: International Agency for Research on Cancer

ACGIH: American Conference of Governmental Industrial Hygienists

STEL: Short Term Exposure Limit

TEEL: Temporary Emergency Exposure Limit

 ${\tt IDLH: Immediately \, Dangerous \, to \, Life \, or \, Health \, Concentrations.}$ 

ES: Exposure Standard OSF: Odour Safety Factor

NOAEL: No Observed Adverse Effect Level LOAEL: Lowest Observed Adverse Effect Level

TLV: Threshold Limit Value LOD: Limit Of Detection OTV: Odour Threshold Value BCF: Bio Concentration Factors BEI: Biological Exposure Index DNEL: Derived No-Effect Level

PNEC: Predicted no-effect concentration

MARPOL: International Convention for the Prevention of Pollution from Ships

IMSBC: International Maritime Solid Bulk Cargoes Code

IGC: International Gas Carrier Code IBC: International Bulk Chemical Code

AIIC: Australian Inventory of Industrial Chemicals

DSL: Domestic Substances List NDSL: Non-Domestic Substances List

IECSC: Inventory of Existing Chemical Substance in China

EINECS: European INventory of Existing Commercial chemical Substances

ELINCS: European List of Notified Chemical Substances

NLP: No-Longer Polymers



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ENCS: Existing and New Chemical Substances Inventory

KECI: Korea Existing Chemicals Inventory NZIoC: New Zealand Inventory of Chemicals

PICCS: Philippine Inventory of Chemicals and Chemical Substances

TSCA: Toxic Substances Control Act

TCSI: Taiwan Chemical Substance Inventory INSQ: Inventario Nacional de Sustancias Químicas

NCI: National Chemical Inventory

FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances

This Material Safety Data Sheet (MSDS) has been prepared in accordance with our company standards and is intended solely for the use of trained personnel. The information provided herein is believed to be accurate as of the date of issue, but no warranty, express or implied, is made regarding its accuracy, completeness, or suitability for any particular purpose. The user is responsible for ensuring safe conditions and compliance with applicable laws and regulations.