

SAFETY DATA SHEET

Revision date: 01-Mar-2017

Version: 3.0

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1. IDENTIFICATION OF THE SUBSTANCE/MIXTURE AND THE COMPANY/UNDERTAKING Product Identifier

Material Name: Ibutilide Fumarate Injection

Trade Name:Corvert InjectionChemical Family:Mixture

Relevant Identified Uses of the Substance or Mixture and Uses Advised Against Intended Use: Pharmaceutical product used as cardiovascular drug

Details of the Supplier of the Safety Data Sheet Pfizer Inc Pfizer Pharmaceuticals Group 235 East 42nd Street New York, New York 10017 1-800-879-3477

Emergency telephone number: CHEMTREC (24 hours): 1-800-424-9300 Contact E-Mail: pfizer-MSDS@pfizer.com

2. HAZARDS IDENTIFICATION

Classification of the Substance or Mixture		
GHS - Classification	Not classified as hazardous	

Label Elements Signal Word: Hazard Statements:	Not Classified Not classified in accordance with international standards for workplace safety.
Other Hazards	An Occupational Exposure Value has been established for one or more of the ingredients (see Section 8).
Note:	This document has been prepared in accordance with standards for workplace safety, which requires the inclusion of all known hazards of the product or its ingredients regardless of the potential risk. The precautionary statements and warning included may not apply in all cases. Your needs may vary depending upon the potential for exposure in your workplace.

3. COMPOSITION / INFORMATION ON INGREDIENTS

<u>Hazardous</u>

Ingredient	CAS Number	EU	GHS Classification	%
		EINECS/ELINCS		
		List		

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3. COMPOSITION / INFORMATI	ON ON INGRED	IENTS		
Ibutilide Fumarate	122647-32-9	Not Listed	Repr. 1B,H360D; Eye Irrit.	0.01
			2A,H319	

Ingredient	CAS Number	EU EINECS/ELINCS List	GHS Classification	%
Sodium acetate trihydrate	6131-90-4	Not Listed	Not Listed	*
Sodium chloride	7647-14-5	231-598-3	Not Listed	*
Water for Injection	7732-18-5	231-791-2	Not Listed	*

Additional Information:

* Proprietary

Ingredient(s) indicated as hazardous have been assessed under standards for workplace safety. In accordance with 29 CFR 1910.1200, the exact percentage composition of this mixture has been withheld as a trade secret.

For the full text of the CLP/GHS abbreviations mentioned in this Section, see Section 16

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4. FIRST AID MEASURES		

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Description of First Aid Measures

Flush with water while holding eyelids open for at least 15 minutes. Seek medical attention immediately.
Remove contaminated clothing. Flush area with large amounts of water. Use soap. Seek medical attention.
Never give anything by mouth to an unconscious person. Wash out mouth with water. Do not induce vomiting unless directed by medical personnel. Seek medical attention immediately.
Remove to fresh air and keep patient at rest. Seek medical attention immediately.
ects, Both Acute and Delayed
For information on potential signs and symptoms of exposure, See Section 2 - Hazards
Identification and/or Section 11 - Toxicological Information.
None known

Indication of the Immediate Medical Attention and Special Treatment Needed Notes to Physician: None

5. FIRE FIGHTING MEASURES

Extinguishing Media: Extinguish fires with CO2, extinguishing powder, foam, or water.

Special Hazards Arising from the Substance or Mixture

Hazardous CombustionFormation of toxic gases is possible during heating or fire.Products:

Fire / Explosion Hazards: Fine particles (such as dust and mists) may fuel fires/explosions.

Advice for Fire-Fighters

During all fire fighting activities, wear appropriate protective equipment, including self-contained breathing apparatus.

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6. ACCIDENTAL RELEASE MEASURES

Personal Precautions, Protective Equipment and Emergency Procedures

Personnel involved in clean-up should wear appropriate personal protective equipment (see Section 8). Minimize exposure.

Environmental Precautions

Place waste in an appropriately labeled, sealed container for disposal. Care should be taken to avoid environmental release.

Methods and Material for Containme Measures for Cleaning / Collecting:	Int and Cleaning Up Contain the source of spill if it is safe to do so. Collect spill with absorbent material. Clean spill area thoroughly.
Additional Consideration for Large Spills:	Non-essential personnel should be evacuated from affected area. Report emergency situations immediately. Clean up operations should only be undertaken by trained personnel.

7. HANDLING AND STORAGE

Precautions for Safe Handling

Avoid breathing vapor or mist. Avoid contact with eyes, skin and clothing. When handling, use appropriate personal protective equipment (see Section 8). Wash thoroughly after handling. Releases to the environment should be avoided. Review and implement appropriate technical and procedural waste water and waste disposal measures to prevent occupational exposure or environmental releases. Potential points of process emissions of this material to the atmosphere should be controlled with dust collectors, HEPA filtration systems or other equivalent controls.

Conditions for Safe Storage, Including any Incompatibilities

Storage Conditions:	Store as directed by product packaging.
Specific end use(s):	Pharmaceutical drug product

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Control Parameters

Refer to available public information for specific member state Occupational Exposure Limits.

Sodium chloride Latvia OEL - TWA Lithuania OEL - TWA	5 mg/m ³ 5 mg/m ³
Ibutilide Fumarate Pfizer OEL TWA-8 Hr:	5µg/m³
Sodium chloride Pfizer Occupational Exposure Band (OEB):	OEB 1 (control exposure to the range of 1000ug/m ³ to 3000ug/m ³)
Exposure Controls Engineering Controls: Personal Protective Equipment:	Engineering controls should be used as the primary means to control exposures. General room ventilation is adequate unless the process generates dust, mist or fumes. Keep airborne contamination levels below the exposure limits listed above in this section. Refer to applicable national standards and regulations in the selection and use of personal protective equipment (PPE). Contact your safety and health professional or safety equipment supplier for assistance in selecting the correct protective clothing/equipment based on an assessment of the workplace conditions, other chemicals used or present in the workplace and specific operational processes.

Hands:	Impervious disposable gloves (e.g. Nitrile, etc.) (double recommended) if skin contact with drug
	product is possible and for bulk processing operations. (Protective gloves must meet the
	standards in accordance with EN374, ASTM F1001 or international equivalent.)
Eyes:	Wear safety glasses or goggles if eye contact is possible. (Eye protection must meet the
	standards in accordance with EN166, ANSI Z87.1 or international equivalent.)
Skin:	Impervious disposable protective clothing is recommended if skin contact with drug product is
	possible and for bulk processing operations. (Protective clothing must meet the standards in
	accordance with EN13982, ANSI 103 or international equivalent.)
Respiratory protection:	Under normal conditions of use, if the applicable Occupational Exposure Limit (OEL) is
	exceeded, wear an appropriate respirator with a protection factor sufficient to control exposure
	to below the OEL (e.g. particulate respirator with a full mask, P3 filter). (Respirators must mee
	the standards in accordance with EN136, EN143, ASTM F2704-10 or international equivalent.)

9. PHYSICAL AND CHEMICAL PROPERTIES

Physical State: Odor: Molecular Formula:	Liquid No data available. Mixture	Color: Odor Threshold: Molecular Weight:	Colorless No data available. Mixture
Solvent Solubility: Water Solubility: Solubility: pH: Melting/Freezing Point (°C): Boiling Point (°C): Partition Coefficient: (Method, pH, E Sodium acetate trihydrate No data available Sodium chloride No data available Ibutilide Fumarate No data available Decomposition Temperature (°C):	No data available No data available Soluble: Water No data available. No data available No data available. Indpoint, Value)		
Evaporation Rate (Gram/s): Vapor Pressure (kPa): Vapor Density (g/ml): Relative Density: Viscosity: Flammablity: Autoignition Temperature (So Flammability (Solids): Flash Point (Liquid) (°C): Upper Explosive Limits (Liqui Lower Explosive Limits (Liqui	d) (% by Vol.):	No data available No data available No data available No data available No data available	

10. STABILITY AND REACTIVITY

Reactivity: Chemical Stability: Possibility of Hazardous Reactions Oxidizing Properties: Conditions to Avoid:

Stable under normal conditions of use. No data available Fine particles (such as dust and mists) may fuel fires/explosions.

No data available

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10. STABILITY AND REACTIVITY

Incompatible Materials: Hazardous Decomposition Products:

As a precautionary measure, keep away from strong oxidizers No data available

11. TOXICOLOGICAL INFORMATION

Information on Toxicological Effects General Information:	The information included in this section describes the potential hazards of the individual ingredients.
Long Term:	Repeat-dose studies in animals have shown a potential to cause adverse effects on reproductive system, the developing fetus.
Known Clinical Effects:	Convulsions possible at high overdosage. Clinical use of this drug has caused headache, nausea, vomiting, changes in heart rate, impaired heart conduction (atrioventricular block), changes in blood pressure.

Acute Toxicity: (Species, Route, End Point, Dose)

Sodium chloride

Rat Oral LD50 3000 mg/kg Mouse Oral LD50 4000 mg/kg

Ibutilide Fumarate

RatOralLD 50> 500 mg/kgRatPara-periostealLD 50 50mg/kgAcute Toxicity Comments:A g

A greater than symbol (>) indicates that the toxicity endpoint being tested was not achievable at the highest dose used in the test.

Irritation / Sensitization: (Study Type, Species, Severity)

Sodium chloride

Eye Irritation Rabbit Moderate Skin Irritation Rabbit Mild

Ibutilide Fumarate

Eye Irritation Rabbit Severe Skin Irritation Rabbit Non-irritating

Repeated Dose Toxicity: (Duration, Species, Route, Dose, End Point, Target Organ)

Ibutilide Fumarate

14 Day(s) Rat 12.5 mg/kg LOAEL Central nervous system Intravenous 14 Day(s) Dog Intravenous < 2.5 mg/kg/dayNOAEL 90 Day(s) 4 mg/kg/day NOAEL Liver, Gastrointestinal system Rat Oral 90 Day(s) Central Nervous System, Male reproductive system Dog Oral 4 mg/kg/day NOEL

Reproduction & Developmental Toxicity: (Study Type, Species, Route, Dose, End Point, Effect(s))

Ibutilide Fumarate

Reproductive & Fertility	Rat	Oral	5 mg/kg/day	NOAEL	Developmental toxicity
Reproductive & Fertility	Rat	Oral	5 mg/kg/day	NOAEL	Maternal Toxicity

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11. TOXICOLOGICAL INFORMATION

Reproductive & Fertility Rat Oral 5 mg/kg/day NOAEL Paternal toxicity Embryo / Fetal Development Rat Oral 5 mg/kg/day NOAEL Not Teratogenic Oral Embryo / Fetal Development Rabbit 20 mg/kg/day LOAEL Teratogenic

Genetic Toxicity: (Study Type, Cell Type/Organism, Result)

Ibutilide Fumarate

Bacterial Mutagenicity (Ames)SalmonellaNegativeUnscheduled DNA SynthesisNegativeIn Vitro MicronucleusNegativeMammalian Cell MutagenicityMouse LymphomaNegative

Carcinogen Status:

None of the components of this formulation are listed as a carcinogen by IARC, NTP or OSHA.

12. ECOLOGICAL INFORMATION

Environmental Overview:	Environmental properties have not been thoroughly investigated. Releases to the environment should be avoided.
Toxicity:	No data available
Persistence and Degradability:	No data available
Bio-accumulative Potential:	No data available
Mobility in Soil:	No data available

13. DISPOSAL CONSIDERATIONS

Waste Treatment Methods:Dispose of waste in accordance with all applicable laws and regulations. Member State
specific and Community specific provisions must be considered. Considering the relevant
known environmental and human health hazards of the material, review and implement
appropriate technical and procedural waste water and waste disposal measures to prevent
occupational exposure and environmental release. It is recommended that waste minimization
be practiced. The best available technology should be utilized to prevent environmental
releases. This may include destructive techniques for waste and wastewater.

14. TRANSPORT INFORMATION

The following refers to all modes of transportation unless specified below.

Not regulated for transport under USDOT, EUADR, IATA, or IMDG regulations.

Safety, Health and Environmental Regulations/Legislation Specific for the Substance or Mixture

Sodium acetate trihydrate CERCLA/SARA 313 Emission reporting California Proposition 65 Australia (AICS): EU EINECS/ELINCS List	Not Listed Not Listed Present Not Listed
Sodium chloride	
CERCLA/SARA 313 Emission reporting California Proposition 65 Inventory - United States TSCA - Sect. 8(b) Australia (AICS): EU EINECS/ELINCS List	Not Listed Not Listed Present Present 231-598-3
Water for Injection	
CERCLA/SARA 313 Emission reporting California Proposition 65 Inventory - United States TSCA - Sect. 8(b) Australia (AICS): REACH - Annex IV - Exemptions from the obligations of Register: EU EINECS/ELINCS List	Not Listed Not Listed Present Present Present 231-791-2
Ibutilide Fumarate CERCLA/SARA 313 Emission reporting California Proposition 65 EU EINECS/ELINCS List	Not Listed Not Listed Not Listed

16. OTHER INFORMATION

Text of CLP/GHS Classification abbreviations mentioned in Section 3

Reproductive toxicity-Cat.1B; H360D - May damage the unborn child Serious eye damage/eye irritation-Cat.2A; H319 - Causes serious eye irritation

Data Sources:	Pfizer proprietary drug development information. Safety data sheets for individual ingredients. Publicly available toxicity information.
Reasons for Revision:	Updated Section 2 - Hazard Identification. Updated Section 8 - Exposure Controls / Personal Protection.
Revision date:	01-Mar-2017 Product Stewardship Hazard Communication
Prepared by:	Pfizer Global Environment, Health, and Safety Operations

Pfizer Inc believes that the information contained in this Material Safety Data Sheet is accurate, and while it is provided in good faith, it is without warranty of any kind, expressed or implied. If data for a hazard are not included in this document there is no known information at this time.

End of Safety Data Sheet