

Revision date: 08-Apr-2019

Version: 2.0

Page 1 of 9

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

Material Name: Dobutamine Injection, USP (Hospira Inc.)

Trade Name: Chemical Family: Dobutamine Injection, USP Not determined

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 Hospira, A Pfizer Company 275 North Field Drive Lake Forest, Illinois 60045 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

Serious Eye Damage/Eye Irritation: Category 2A

### Label Elements

Signal Word:	Warning
Hazard Statements:	H319 - Causes serious eye irritation
Precautionary Statements:	P264 - Wash hands thoroughly after handling P280 - Wear protective gloves/protective clothing/eye protection/face protection P305 + P351 + P338 - IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing P337 + P313 - If eye irritation persists: Get medical advice/attention



PZ03082

Hospira UK Limited Horizon Honey Lane Hurley Maidenhead, SL6 6RJ United Kingdom Emergency telephone number: International Chemtrec (24 hours): +1-703-527-3887

Material Name: Dobutamine Injection, USP (Hospira Inc.) Revision date: 08-Apr-2019

**Other Hazards** An Occupational Exposure Value has been established for one or more of the ingredients (see Section 8). This document has been prepared in accordance with standards for workplace safety, which Note: 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**

### Hazardous

Ingredient	CAS Number	EU	GHS Classification	%
		EINECS/ELINCS		
		List		
Dobutamine Hydrochloride	49745-95-1	256-464-1	Eye Dam 1 (H318)	1.25
SODIUM HYDROXIDE	1310-73-2	215-185-5	Skin Corr. 1A (H314)	**
HYDROCHLORIC ACID	7647-01-0	231-595-7	Skin Corr.1B (H314)	**
			STOT SE 3 (H335)	

Ingredient	CAS Number	EU EINECS/ELINCS List	GHS Classification	%
sodium metabisulphite	8681-57-4	Not Listed	Not Listed	*
Disodium EDTA (dihydrate)	6381-92-6	Not Listed	Not Listed	*
Water for Injection	7732-18-5	231-791-2	Not Listed	*

### Additional Information:

\* Proprietary

\*\* to adjust pH

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

### 4. FIRST AID MEASURES

Description of First Aid Measures	
Eye Contact:	Rinse immediately with plenty of water, also under the eyelids, for at least 15 minutes. If irritation occurs or persists, get medical attention.
Skin Contact:	Wash off immediately with soap and plenty of water. If skin irritation persists, call a physician.
Ingestion:	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.
Inhalation:	Move to fresh air If discomfort occurs, get medical attention.
Most Important Symptoms and Effect Symptoms and Effects of Exposure: Medical Conditions Aggravated by Exposure:	cts, 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 Notes to Physician:	Attention and Special Treatment Needed None

Notes to Physician:

Material Name: Dobutamine Injection, USP (Hospira Inc.) Revision date: 08-Apr-2019

### 5. FIRE FIGHTING MEASURES

**Extinguishing Media:** As for primary cause of fire.

 Special Hazards Arising from the Substance or Mixture

 Hazardous Combustion
 Formation of toxic gases is possible during heating or fire.

 Products:
 Products:

Fire / Explosion Hazards: Not applicable

### **Advice for Fire-Fighters**

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

### 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 Containment and Cleaning Up

Measures for Cleaning / Collecting:	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. Cleanup 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.
Incompatible Materials:	None known
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.

Dobutamine Hydrochloride Pfizer OEL TWA-8 Hr:

Pfizer OEL TWA-8 Hr:	300µg/m <sup>3</sup> , Severe Eye Irritant
SODIUM HYDROXIDE	
ACGIH Ceiling Threshold Limit:	2 mg/m <sup>3</sup>
Australia PEAK	2 mg/m <sup>3</sup>
Austria OEL - MAKs	2 mg/m <sup>3</sup>
Bulgaria OEL - TWA	2.0 mg/m <sup>3</sup>

Page 4 of 9 Version: 2.0

3. EXPOSURE CONTROLS / PERSONAL F	PROTECTION	
Czech Republic OEL - TWA	1 mg/m <sup>3</sup>	
Estonia OEL - TWA	1 mg/m <sup>3</sup>	
France OEL - TWA	2 mg/m <sup>3</sup>	
Greece OEL - TWA	2 mg/m <sup>3</sup>	
Hungary OEL - TWA	2 mg/m <sup>3</sup>	
Japan - OELs - Ceilings	$2 \text{ mg/m}^3$	
Latvia OEL - TWA	0.5 mg/m <sup>3</sup>	
OSHA - Final PELS - TWAs:	$2 \text{ mg/m}^3$	
Poland OEL - TWA	0.5 mg/m <sup>3</sup>	
Slovakia OEL - TWA	$2 \text{ mg/m}^3$	
Slovenia OEL - TWA	2 mg/m <sup>3</sup>	
Sweden OEL - TWAs	1 mg/m <sup>3</sup>	
Switzerland OEL -TWAs	$2 \text{ mg/m}^3$	
IYDROCHLORIC ACID		
ACGIH Ceiling Threshold Limit:	2 ppm	
Australia PEAK	5 ppm 7.5 mg/m³	
Austria OEL - MAKs	5 ppm 8 mg/m <sup>3</sup>	
Belgium OEL - TWA	5 ppm 8 mg/m <sup>3</sup>	
Bulgaria OEL - TWA	5 ppm 8.0 mg/m³	
Cyprus OEL - TWA	5 ppm 8 mg/m <sup>3</sup>	
Czech Republic OEL - TWA	8 mg/m <sup>3</sup>	
Estonia OEL - TWA	5 ppm 8 mg/m <sup>3</sup>	
Germany - TRGS 900 - TWAs	2 ppm 3 mg/m <sup>3</sup>	
Germany (DFG) - MAK	2 ppm 3.0 mg/m <sup>3</sup>	
Greece OEL - TWA	5 ppm 7 mg/m <sup>3</sup>	
Hungary OEL - TWA	8 mg/m <sup>3</sup>	
Ireland OEL - TWAs	5 ppm 8 mg/m <sup>3</sup>	
Italy OEL - TWA	5 ppm 8 mg/m <sup>3</sup>	
Japan - OELs - Ceilings	2 ppm 3.0 mg/m <sup>3</sup>	
Latvia OEL - TWA	5 ppm 8 mg/m <sup>3</sup>	
Lithuania OEL - TWA	5 ppm 8 mg/m <sup>3</sup>	
Luxembourg OEL - TWA	5 ppm 8 mg/m <sup>3</sup>	
Malta OEL - TWA	5 ppm 8 mg/m <sup>3</sup>	
Netherlands OEL - TWA	8 mg/m <sup>3</sup>	
Poland OEL - TWA	5 mg/m <sup>3</sup>	

Page 5 of 9 Version: 2.0

8. EXPOSURE CONTROLS		
Portugal OEL - TWA	5 ppm	
Fortugal OEL - TWA	8 mg/m <sup>3</sup>	
Romania OEL - TWA	5 ppm	
	8 mg/m <sup>3</sup>	
Slovakia OEL - TWA	5 ppm	
	$8.0 \text{ mg/m}^3$	
Slovenia OEL - TWA	5 ppm	
	8 mg/m <sup>3</sup>	
Spain OEL - TWA	5 ppm	
	$7.6 \text{ mg/m}^3$	
Switzerland OEL -TWAs	2 ppm	
Switzenand OLE -TWAS	$3.0 \text{ mg/m}^3$	
Vietnam OEL - TWAs	5 mg/m <sup>3</sup>	
	3 mg/m	
Exposure Controls		
Engineering Controls:	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.	
Personal Protective	Refer to applicable national standards and regulations in the selection and use of personal	
Equipment:	protective equipment (PPE).	
Hands:	Impervious gloves (e.g. Nitrile, etc.) are 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 goggles if eye contact is possible (face shield recommended if splashing is possible). (Eye protection must meet the standards in accordance with EN166, ANSI Z87.1 or international equivalent.)	
Skin:	Wear impervious protective clothing to prevent skin contact. (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 exposures to below the OEL (e.g. particulate respirator with a full mask, P3 filter). (Respirators must meet the standards in accordance with EN136, EN143, ASTM F2704-10 or international equivalent.)	

# 9. PHYSICAL AND CHEMICAL PROPERTIES

Physical State:	Liquid	Color:	Colorless
Odor:	No data available.	Odor Threshold:	No data available.
Molecular Formula:	Mixture	Molecular Weight:	Mixture
Solvent Solubility: Water Solubility: pH: Melting/Freezing Point (°C): Boiling Point (°C): Partition Coefficient: (Method, pH, E Water for Injection No data available HYDROCHLORIC ACID No data available SODIUM HYDROXIDE No data available Disodium EDTA (dihydrate)	No data available Soluble 2.5-5.5 No data available No data available. Endpoint, Value)		

Page 6 of 9 Version: 2.0

### 9. PHYSICAL AND CHEMICAL PROPERTIES

No data available		
sodium metabisulphite		
No data available		
Dobutamine Hydrochloride		
No data available		
Decomposition Temperature (°C):	No data available.	
Evaporation Rate (Gram/s):	No data available	
Vapor Pressure (kPa):	No data available	
Vapor Density (g/ml):	No data available	
Relative Density:	No data available	
Viscosity:	No data available	
Flammablity:		
Autoignition Temperature (Solid) (°C):		No data available
Flammability (Solids):		No data available
Flash Point (Liquid) (°C):		No data available
		No data available
Lower Explosive Limits (Liquid) (% by Vol.):		No data available

### **10. STABILITY AND REACTIVITY**

Reactivity: Chemical Stability:	No data available Stable under normal conditions of use.
Possibility of Hazardous Reactions	
Oxidizing Properties:	None
Conditions to Avoid:	None known
Incompatible Materials:	None known
Hazardous Decomposition	None known
Products:	

# **11. TOXICOLOGICAL INFORMATION**

Information on Toxicological Effects General Information:	The information included in this section describes the potential hazards of the individual ingredients.
Short Term: Known Clinical Effects:	May cause eye irritation (based on components). The most common adverse effects seen during clinical use of this drug include headache, nausea, shortness of breath (dyspnea), palpitations, chest pain, increased heart rate (tachycardia), increase in blood pressure (hypertension).

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

### HYDROCHLORIC ACID Rat Oral LD 50 238-277 mg/kg

### **Dobutamine Hydrochloride**

Rat Oral LD50 2296 mg/kg Mouse Oral LD50 1324mg/kg Rat Intravenous LD50 59.6mg/kg Mouse Intravenous LD50 34.3mg/kg

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

Material Name: Dobutamine Injection, USP (Hospira Inc.) Revision date: 08-Apr-2019

### 11. TOXICOLOGICAL INFORMATION

#### **Dobutamine Hydrochloride**

Skin Irritation Rabbit Non-irritating Eye Irritation Rabbit Corrosive

### Reproduction & Development Toxicity: (Duration, Species, Route, Dose, End Point, Effect(s))

<b>Dobutamine Hydrochloride</b> Embryo / Fetal Development Embryo / Fetal Development	Rat No route specified14.4 mg/kg/day NOAEL Not teratogenic Rabbit No route specified 28.8 mg/kg/day NOAEL Not Teratogenic
HYDROCHLORIC ACID Bacterial Mutagenicity (Ames) In Vivo Micronucleus Rat	Salmonella Negative Negative
Carcinogen Status:	None of the components of this formulation are listed as a carcinogen by IARC, NTP or OSHA.
HYDROCHLORIC ACID IARC:	Group 3 (Not Classifiable)

### 12. ECOLOGICAL INFORMATION

Environmental Overview:	Environmental properties have not been 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.

# **15. REGULATORY INFORMATION**

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

Dobutamine Hydrochloride	Netlisted
CERCLA/SARA 313 Emission reporting	Not Listed Not Listed
California Proposition 65 EU EINECS/ELINCS List	256-464-1
EU EINECS/ELINCS LISt	200-404-1
sodium metabisulphite	
CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
EU EINECS/ELINCS List	Not Listed
SODIUM HYDROXIDE	
CERCLA/SARA 313 Emission reporting	Not Listed
CERCLA/SARA Hazardous Substances	1000 lb
and their Reportable Quantities:	454 kg
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
Standard for the Uniform Scheduling	Schedule 5
for Drugs and Poisons:	Schedule 6
EU EINECS/ELINCS List	215-185-5
Disodium EDTA (dihydrate)	
CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Australia (AICS):	Present
EU EINECS/ELINCS List	Not Listed
HYDROCHLORIC ACID	
CERCLA/SARA 313 Emission reporting	1.0 %
CERCLA/SARA Hazardous Substances	5000 lb
and their Reportable Quantities:	2270 kg
CERCLA/SARA - Section 302 Extremely Hazardous	500 lb
TPQs	
CERCLA/SARA - Section 302 Extremely Hazardous	5000 lb
Substances EPCRA RQs	<b>NI</b> 211 2 2
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present

Material Name: Dobutamine Injection, USP (Hospira Inc.) Revision date: 08-Apr-2019

15. REGULATORY INFORMATION	
Standard for the Uniform Scheduling	Schedule 5
for Drugs and Poisons:	Schedule 6
EU EINECS/ELINCS List	231-595-7
Water for Injection	
CERCLA/SARA 313 Emission reporting	Not Listed
California Proposition 65	Not Listed
Inventory - United States TSCA - Sect. 8(b)	Present
Australia (AICS):	Present
REACH - Annex IV - Exemptions from the obligations of Register:	Present
EU EINECS/ELINCS List	231-791-2

### **16. OTHER INFORMATION**

### Text of CLP/GHS Classification abbreviations mentioned in Section 3

Skin corrosion/irritation-Cat.1A; Skin corrosion/irritation-Cat.1B; H314 - Causes severe skin burns and eye damage Serious eye damage/eye irritation-Cat.1; H318 - Causes serious eye damage Specific target organ toxicity, single exposure; Respiratory tract irritation-Cat.3; H335 - May cause respiratory irritation

Data Sources:	Pfizer proprietary drug development information. Publicly available toxicity information.
Reasons for Revision:	Updated Section 8 - Exposure Controls / Personal Protection.
Revision date:	08-Apr-2019 Product Stewardship Hazard Communication
Prepared by:	Pfizer Global Environment, Health, and Safety Operations

Pfizer Inc believes that the information contained in this 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