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

**Product Identifier** 

Material Name: Buprenorphine Hydrochloride Injection (Hospira Inc.)

Trade Name: Buprenorphine Hydrochloride Injection

Chemical Family: Not determined

Relevant Identified Uses of the Substance or Mixture and Uses Advised Against

Intended Use: Pharmaceutical product used as analgesic

Details of the Supplier of the Safety Data Sheet

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

**Emergency telephone number:** 

1-800-879-3477

Hospira UK Limited

Horizon Honey Lane Hurley

Maidenhead, SL6 6RJ United Kingdom

**Emergency telephone number:** 

International CHEMTREC (24 hours): +1-703-527-3887

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: Not Classified

Hazard Statements: 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

**Hazardous** 

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Ingredient	CAS Number	EU EINECS/ELINCS	GHS Classification	%
		List		
HYDROCHLORIC ACID	7647-01-0	231-595-7	Skin Corr.1B (H314)	**
			STOT SE 3 (H335)	
Buprenorphine Hydrochloride	53152-21-9	Not Listed	Acute Tox 4 (H302)	0.0324
			STOT SE 3 (H336)	

Ingredient	CAS Number	EU EINECS/ELINCS	GHS Classification	%
		List		
Water for Injection	7732-18-5	231-791-2	Not Listed	*
Dextrose	14431-43-7	Not Listed	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 thoroughly with plenty of water, also under the eyelids. 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 Effects, Both Acute and Delayed

Symptoms and Effects of For information on potential signs and symptoms of exposure, See Section 2 - Hazards

**Exposure:** Identification and/or Section 11 - Toxicological Information.

Medical Conditions None known

Aggravated by Exposure:

Indication of the Immediate Medical Attention and Special Treatment Needed

Notes to Physician: None

## 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:** 

Fire / Explosion Hazards: Not applicable

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Advice for Fire-Fighters

During all fire fighting 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 / Contain the source of spill if it is safe to do so. Collect spill with absorbent material. Clean spill

**Collecting:** area thoroughly.

Additional Consideration for Non-essential personnel should be evacuated from affected area. Report emergency

Large Spills: situations immediately. Clean up operations should only be undertaken by trained personnel.

## 7. HANDLING AND STORAGE

### **Precautions for Safe Handling**

Restrict access to work area. 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.

#### HYDROCHLORIC ACID

 ACGIH Ceiling Threshold Limit:
 2 ppm

 Australia PEAK
 5 ppm

 7.5 mg/m³

 Austria OEL - MAKs
 5 ppm

 8 mg/m³

**Belgium OEL - TWA** 5 ppm 8 mg/m<sup>3</sup>

 Bulgaria OEL - TWA
 5 ppm

 8.0 mg/m³
 5 ppm

 Cyprus OEL - TWA
 5 ppm

8 mg/m<sup>3</sup>

 Czech Republic OEL - TWA
 8 mg/m³

 Estonia OEL - TWA
 5 ppm

 8 mg/m³
 8 mg/m³

**Germany - TRGS 900 - TWAs** 2 ppm 3 mg/m<sup>3</sup>

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8. EXPOSURE CONTROLS / PERSONAL PROTECTION

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> 5 ppm **Italy OEL - TWA** 8 mg/m<sup>3</sup> 2 ppm Japan - OELs - Ceilings 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> 5 ppm **Luxembourg OEL - TWA** 8 mg/m<sup>3</sup> Malta OEL - TWA 5 ppm 8 mg/m<sup>3</sup> **Netherlands OEL - TWA**  $8 \text{ mg/m}^3$ **Poland OEL - TWA** 5 mg/m<sup>3</sup> 5 ppm Portugal 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 mg/m<sup>3</sup> 5 ppm Slovenia OEL - TWA 8 mg/m<sup>3</sup> Spain OEL - TWA 5 ppm 7.6 mg/m<sup>3</sup> **Switzerland OEL -TWAs** 2 ppm 3.0 mg/m<sup>3</sup> Vietnam OEL - TWAs 5 mg/m<sup>3</sup>

The purpose of the Occupational Exposure Band (OEB) classification system is to separate substances into different Hazard categories when the available data are sufficient to do so, but inadequate to establish an Occupational Exposure Limit (OEL). The OEB given is based upon an analysis of all currently available data; as such, this value may be subject to revision when new information becomes available.

**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. It is recommended

that all operations be fully enclosed and no air recirculated.

**Personal Protective** 

Equipment:

Refer to applicable national standards and regulations in the selection and use of personal

protective equipment (PPE).

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.)

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## 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

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 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: No data available

Water Solubility: Soluble pH: Soluble 3.5-5.5

Melting/Freezing Point (°C):

Boiling Point (°C):

No data available.

No data available.

Partition Coefficient: (Method, pH, Endpoint, Value)

HYDROCHLORIC ACID
No data available
Dextrose

No data available

**Buprenorphine Hydrochloride** 

No data available Water for Injection No data available

**Decomposition Temperature (°C):** No data available.

Evaporation Rate (Gram/s):

Vapor Pressure (kPa):

Vapor Density (g/ml):

Relative Density:

No data available

Flammablity:

Autoignition Temperature (Solid) (°C):

Flammability (Solids):

Flash Point (Liquid) (°C):

Upper Explosive Limits (Liquid) (% by Vol.):

Lower Explosive Limits (Liquid) (% by Vol.):

No data available
No data available
No data available

## 10. STABILITY AND REACTIVITY

Reactivity: No data available

Chemical Stability: 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:** 

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# 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:** May cause central nervous system effects.

**Known Clinical Effects:** The most common adverse effects seen during clinical use of this drug include drowsiness,

nausea, vomiting, decrease in blood pressure (hypotension), increase in blood pressure (hypertension), respiratory depression, decreased heart rate (bradycardia), increased heart

rate (tachycardia), dizziness, sweating, headache, dry mouth, hallucinations.

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

HYDROCHLORIC ACID

Rat Oral LD 50 238-277 mg/kg

**Buprenorphine Hydrochloride** 

Rat Oral LD50 > 1000 mg/kg Mouse Oral LD50 800mg/kg Rat IV LD50 62mg/kg Mouse IV LD50 72mg/kg

Acute Toxicity Comments: A greater than symbol (>) indicates that the toxicity endpoint being tested was not achievable

at the highest dose used in the test.

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

**Buprenorphine Hydrochloride** 

Reproductive & Fertility Rat Oral80 mg/kg/day NOAEL Negative, Fertility

Embryo / Fetal Development Rat Intramuscular Subcutaneous 5 mg/kg/day NOAEL Not Teratogenic Embryo / Fetal Development Rabbit Intramuscular Subcutaneous 5 mg/kg/day NOAEL Not Teratogenic

Embryo / Fetal Development Rat Oral 160 mg/kg/day NOAEL Not Teratogenic

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

HYDROCHLORIC ACID

Bacterial Mutagenicity (Ames) Salmonella Negative

In Vivo Micronucleus Rat Negative

**Buprenorphine Hydrochloride** 

Bacterial Mutagenicity (Ames) Equivocal

Mammalian Cell Mutagenicity Hamster Bone Marrow Negative Mammalian Cell Mutagenicity Mouse Lymphoma Negative

Carcinogenicity: (Duration, Species, Route, Dose, End Point, Effect(s))

**Buprenorphine Hydrochloride** 

27 Month(s) Rat Oral, in feed 56 mg/kg/day NOAEL Not carcinogenic 86 Week(s) Mouse Oral, in feed 100 mg/kg/day NOAEL Not carcinogenic

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

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# 11. TOXICOLOGICAL INFORMATION

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

HYDROCHLORIC ACID

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5000 lb

# 15. REGULATORY INFORMATION

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** 

Substances EPCRA RQs

California Proposition 65
Inventory - United States TSCA - Sect. 8(b)
Australia (AICS):
Standard for the Uniform Scheduling
for Drugs and Poisons:
Schedule 6
EU EINECS/ELINCS List
Not Listed
Present
Schedule 5
Schedule 6
231-595-7

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

Present

Present

obligations of Register:

EU EINECS/ELINCS List 231-791-2

**Buprenorphine Hydrochloride** 

CERCLA/SARA 313 Emission reporting

California Proposition 65

EU EINECS/ELINCS List

Not Listed

Not Listed

Dextrose

CERCLA/SARA 313 Emission reporting

California Proposition 65

Australia (AICS):

Present

EU EINECS/ELINCS List

Not Listed

Not Listed

# 16. OTHER INFORMATION

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

Acute toxicity, oral-Cat.4; H302 - Harmful if swallowed

Skin corrosion/irritation-Cat.1B; H314 - Causes severe skin burns and eye damage

Specific target organ toxicity, single exposure; Respiratory tract irritation-Cat.3; H335 - May cause respiratory irritation Specific target organ toxicity, single exposure; Narcotic effects-Cat.3; H336 - May cause drowsiness and dizziness

**Data Sources:** Pfizer proprietary drug development information. Publicly available toxicity information.

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Product Stewardship Hazard Communication
Pfizer Global Environment, Health, and Safety Operations

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Prepared by:

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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** 

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