

Bortezomib for Injection		Safety Data Sheet (SDS)	
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Section 1 - Identification

(a) Product Identifier: Bortezomib for Injection

(b) Product Code: 70860-225

Common/Trade Name: Bortezomib for Injection

Chemical Name: [(1R)-3-methyl-1-[[(2S)-1-oxo-3-phenyl-2-[(pyrazinylcarbonyl)

amino]propyl]amino]butyl] boronic acid

Chemical Family: Proteasome Inhibitor

(c) **Product Use:** Pharmaceutical, Injectable

Product Type: Regulated Prescription Drug

Container Information: Vial

(d) **Distributor:** Athenex Pharmaceutical Division, 10 N. Martingale Road,

Suite 230, Schaumburg, IL 60173, 847-886-9515

(e) Emergency Telephone: 855-273-0154

Section 2 - Hazards Identification

(a)	Classification:	Reproductive toxicity. Category 1A Specific Target Organ Toxicity – STOT Repeated exposure RE. Category 2 (Nervous System) Reproductive toxicity. Effects on or via lactation.	
(b)	Signal Word:	Danger	
	Hazard Statement	H360 – May damage fertility or the unborn child. H373 – May cause damage to organs through prolonged or repeated exposure. H362 May cause harm to breast-fed children.	
	Symbols		



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P202 - Do not handle until all safety precauunderstood. P260 - Do not breathe dust/fume/gas/mist/P263 - Avoid contact during pregnancy an P280 - Wear protective gloves. Wear eye oprotective clothing. P270 - Do not eat, drink or smoke when us P264 - Wash hands thoroughly after handli Response: P308 + P313 - IF exposed or coattention.		P260 – Do not breathe dust/fume/gas/mist/vapors/spray. P263 – Avoid contact during pregnancy and while nursing. P280 - Wear protective gloves. Wear eye or face protection. Wear protective clothing. P270 - Do not eat, drink or smoke when using this product. P264 - Wash hands thoroughly after handling Response: P308 + P313 - IF exposed or concerned: Get medical
		Storage: P405 - Store locked up. Disposal: P501 - Dispose of contents and container in accordance with all local, regional, national, and international regulations.
(c)	Description of Hazards	Potential Health Effects: This product is intended for therapeutic use only when prescribed by a physician. Potential adverse reactions from prescribed doses and overdoses are described in the package instert. Handling should only be preformed by personnel trained and familiar with handling of potent active pharmaceutical ingredients. May cause irritaiton to eyes, skin and respiratory tract, nausea, comiting, rash, diarrheah. Acute Health Effects: The effects reported ar from use of the drug product clinically or the testing of the drug substance in laboratory animals. In patients given the drug, Bortezomib produces many of the acute offsets typical of drugs that kill rapidly growing cells, including effects on the hematological system, fatigue, nausea, vomiting, diarrheah, and fever. Other organ systems reported to be effected include the respiratory tract (shortness of breath), and nervous system (headache, dizziness, and preipheral neuropathy). Some of the acute effects may be delayed in onset, such as the effects on the hematological sytem, which are characterized by decreased white blood cells, platelets and red blood cells, and may occur serveral days or weeks following the acute exposure. These effects may potentially occur from acute (severe spill) or repeated overexposure in occupational sellings. Chronic Health Effects: based on other antineoplasic agents and data on Bortezomib, the spectrum of effects after chronic exposure would be expected to be similar to those after acute exposure. Initial repeated dose studies in laboratory animals have shown similar effects as acute studies. Target Organs: Eyes, Skin, Respiratory system, Hematological system, peripheral nervous system, liver, kidney, reproductive system.
(d)	Unknown Acute Toxicity	N/A



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Section 3 – Composition / Information on Ingredients

(a) Chemical Name	(b) Common Name / Synonym	% Composition or other measure	(c) CAS No.	(d) Impurities / Stabilizing Additives
Bortezomib	Bortezomib	3.5 mg	390800-88-1	N/A
Mannitol, USP	Mannitol	35 mg	69-65-8	N/A

Section 4 – First Aid Measures

Eye Exposure: Immediately flush eyes with plenty of water for 15 – 20 minutes. Ensure adequate

flushing of the eyes by separating the eyelids with fingers. Get immediate medical

attention.

Skin Exposure: Immediately wash skin with plenty of soap and water for 15-20 minutes, while

removing contaminated clothing and shoes. Get medical attention if irritation

develops or persists.

Ingestion: If swallowed, do NOT induce vomiting. Call a physician or poison control

center immediately. Never give anything by mouth to an unconscious person.

Inhalation: If inhaled, remove to fresh air. If not breathing, give artificial respiration or

give oxygen by trained personnel. Seek immediate medical attention.

Notes to **Physician:**

Material is a cytotoxic anticancer drug

Section 5 – Fire-fighting Measures

(a) Extinguishing Media Use alcohol resistant foam, carbon dioxide, dry chemical or water fog or spray when fighting fires.

Carbon monoxide and nitrogen oxides.

(b) Hazardous Combustion

Byproducts:



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(c) Special Protective Equipment /
Precautions:

As in any fire, wear Self-Contained Breathing Apparatus (SCBA), MSHA/NIOSH (approved or equivalent) and full protective gear.

Evacuate area of unprotected personnel. Use cold water spray to cool fire exposed containers to minimize risk of rupture. Do not enter confined fire space without full protective gear. If possible, contain fire run-off water.

Section 6 - Accidental Release Measures

Spill: Contain spills with an inert, absorbent material such as soil, sand, or oil

dry, then place in a chemical waste container. After removal, flush spill

area with soap and water to remove trace residue.

Release to Air: Avoid personal contact and breathing vapors or mists.

Release to Water: Avoid runoff into storm sewers, ditches, and waterways.

Section 7 - Handling and Storage

General Handling: When handling pharmaceutical products, avoid all contact and

inhalation of vapors, mists and/or fumes. Use with adequate ventilation. Use only in accordance with directions. Handling should only be performed by personnel trained and familiar with

handling of potent active pharmaceutical ingredients.

Storage Conditions: Unopened vials may be stored at controlled room temperature 25°C

(77°F); excursions permitted between 15°C and 30°C (59°F and 86°F).

Retain in original package to protect from light.

Section 8 - Exposure Controls / Personal Protection

(a) Exposure Limits: None

(b) Engineering Controls

Use appropriate engineering control such as process enclosures, local exhaust ventilation, or other engineering controls to control airborne levels below recommended exposure limits. Good general ventilation should be sufficient to control airborne levels. Where such systems are not effective wear suitable personal protective equipment, which performs satisfactorily and meets OSHA or other recognized standards. Consult with local procedures for selection, training, inspection and maintenance of the personal protective equipment.



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(c) Individual Protection Measures

Respiratory Protection:	A NIOSH approved air-purifying respirator with an organic vapor cartridge or canister may be permissible under certain circumstances where airborne concentrations are expected to exceed exposure limits. Protection provided by air purifying respirators is limited. Use a positive pressure air supplied respirator if there is any potential for an uncontrolled release, exposure levels are not known, or any other circumstances where air purifying respirators may not provide adequate protection.
Eye Protection:	Chemical splash goggles. Wear a face shield also when splash hazard exist.
Skin Protection: Protective laboratory coat, apron, or disposable garment recommended. Wear appropriate protective gloves. Consult glov manufacturer's data for permeability data. Nitrile rubber or naturubber gloves are recommended.	
Other Protective Equipment:	Facilities storing or utilizing this material should be equipped with an eyewash and a deluge shower safety station. Decontaminate all protective equipment after use. Exercise extreme care when working with sharps/needles/syringes and potent drugs.

Section 9 - Physical and Chemical Properties

(a)	Appearance	White to off-white cake or powder.
(b)	Odor	Not available
(c)	Odor Threshold	Not available
(d)	рН	Not available
(e)	Melting Point:	140 – 150 °C
(f)	Initial Boiling Point:	Not available
(g)	Flash Point	Not available
(h)	Evaporation Rate:	Not available
(i)	Flammability	Not available
(j)	Upper Lower Flammability or Explosion Limits	Not available
(k)	Vapor Pressure:	Not available
(1)	Vapor Density:	Not available
(m)	Relative Density	Not available



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(n)	Solubility(ies)	Soluble in water (3.3 to 3.8 mg/mL at pH range of 2 to 6.5)
(0)	Partition Coefficient: n-octanol/water	Not available
(p)	Auto-ignition Temperature	Not available
(q)	Decomposition Temperature	Not available
(r)	Viscosity	Not available

Section 10 - Stability and Reactivity

(a)	Reactivity	Stable under normal conditions of use.
(b)	Chemical Stability	Stable at normal temperatures and pressures
(c)	Possibility of Hazardous Reactions	Hazardous polymerization does not occur
(d)	Conditions to Avoid	Protect from light
(e)	Incompatible Materials	Avoid contact with oxidizing agents
(f)	Hazardous Decomposition Products	Carbon oxides. Nitrogen oxides (NOx)

Section 11 - Toxicological Information

(a)	Likely Routes of Exposure	Ingestion, inhalation, skin contact, mucous membranes.
(b)	Symptoms related to the physical, chemical and toxicological characteristics	Eye: Moderate to severe eye irritant Skin: Moderate to severe skin eye irritant Inhalation: Inhalation data not identified. Based on the high acute toxicity reported in non-clinical IV studies, a potential for toxicity via the inhaled route cannot be excluded. Ingestion: Very toxic by ingestion



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(c)	Delayed and immediate effects and chronic effects from short- and long-term exposure	Repeated occupational overexposure may cause fatigue and fever, and effects on the hematological (decreases in hemoglobin/anemia, blood counts and platelets), gastrointestinal (nausea, diarrhea, vomiting, abdominal pain), and nervous systems (headache, peripheral neuropathy). May affect fertility and fetal development based on mechanism and animal toxicity studies. **Mutagenicity**: Bortezomib API showed clastogenic activity in the in vitro chromosomal aberration assay using Chinese hamster ovary cells, but bortezomib was not genotoxic in the in vitro mutagenicity assay (Ames test), or in the in vivo micronucleus assay in mice. **Reproductive Toxicity**: Fertility studies have not been performed, but degenerative effects have been observed in the ovaries and testes of rats in a 6 month toxicity study. Bortezomib API could affect male or female fertility. **Teratogenicity**: Not conducted**
(d)	Other Toxicological Information	Bortezomib API is a cytotoxic antineoplastic compound that exerts its biological effects by inhibiting the activity of the proteasome, a macromolecular complex found in all cells that plays a critical role in protein homeostasis. In local tolerance studies bortezomib showed minimal tissue irritancy; however direct skin contact should be avoided. Bortezomib API is acutely lethal in animals, when delivered intravenously at sufficient doses, and death is due to cardiovascular effects. Systemic, therapeutic exposure to bortezomib in humans is associated with a variety of adverse events, including peripheral neuropathy, hypotension, cardiac disorders, pulmonary disorders, reversible posterior leukoencephalopathy syndrome (RPLS), gastrointestinal disturbances (nausea, diarrhea, constipation, vomiting, decreased platelet count and neutrophil counts in the blood, tumor lysis syndrome, and liver disturbances.

Acute Toxicity: Not available.

Hazardous Chemical Listings

NTP: Not Listed IARC: Not Listed OSHA: Not Listed



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Section 12 - Ecological Information

(a)	Ecotoxicity	No Data Available
(b)	Persistence and degradability	No Data Available
(c)	Bioaccumulative potential	No Data Available
(d)	Mobility in soil	No Data Available
(e)	Other Adverse Effects	No Data Available

Section 13 - Disposal Considerations

Waste Disposal: All wastes containing the material should be properly labeled. Dispose of any waste residues according to prescribed federal, state, and local guidelines, e.g., appropriately permitted chemical waste incinerator. Rinse waters resulting from spill cleanups should be discharged in an environmentally safe manner, e.g., appropriately permitted municipal or on-site wastewater treatment facility.

Section 14 - Transport Information

DOT

DOT Shipping Name:	Not Regulated.
DOT UN Number	Not Regulated.
DOT Hazard Class	Not Regulated.
DOT Packing Group	Not Regulated.

IATA

IATA Shipping Name:	Not Regulated.
IATA UN Number	Not Regulated.
IATA Hazard Class	Not Regulated.
IATA Packing Group	Not Regulated.

Section 15 - Regulatory Information

Below is selected regulatory information chosen primarily for possible Athenex Pharmaceutical Division usage. This section is not a complete analysis or reference to all applicable regulatory information. Please consider all applicable laws and regulations for your country/state.



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U.S. Regulations:

TSCA – Not regulated CERCLA - Not on this list SARA 302 - Not on this list SARA 313 - Not on this list

Section 16 - Other Information

As of the date of effectiveness, we are providing available information relevant to the handling of this material in the workplace. All information contained herein is offered with the good faith belief that it is accurate. THIS SAFETY DATA SHEET SHALL NOT BE DEEMED TO CREATE ANY WARRANTY OF ANY KIND (INCLUDING WARRANTY OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE). In the event of an adverse incident associated with this material, this safety data sheet is not intended to be a substitute for consultation with appropriately trained personnel. Nor is this safety data sheet intended to be a substitute for product literature which may accompany the finished product.

For additional information contact:

Athenex Pharmaceutical Division 10 N. Martingale Road, Suite 230 Schaumburg, IL 60173 847-886-9515

Glossary: This glossary contains definitions of general terms used in SDSs. Not all of these Glossary Terms will apply to this SDS.

ACGIH	American Conference of Governmental Industrial Hygienists
AICS	Australian Inventory of Chemical Substances
AIHA	American Industrial Hygiene Association
ANSI	American National Standards Institute
CAS Number	Chemical Abstract Service Registry Number
CERCLA	Comprehensive Environmental Response Compensation and Liability Act (of
CHAN	Chemical Hazard Alert Notice
CHEMTREC	Chemical Transportation Emergency Center
DOT	Department of Transportation
DSL	Domestic Substances List
ECHA	European Chemicals Agency
EINECS	European Inventory of Existing Commercial Chemical Substances
ELINCS	European List of Notified Chemical Substances
EPA	Environmental Protection Agency
GHS	Globally Harmonized System of Classification and Labelling of Chemicals
HEPA	High Efficiency Particulate Air (Filter)
HMIS	Hazardous Materials Identification System
IARC	International Agency for Research on Cancer
ICAO/IATA	International Civil Aviation Organization/International Air Transport
IMO	International Maritime Organization



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KOWOctanol/Water Partition CoefficientLELLower Explosive LimitMSDSMaterial Safety Data SheetMSHAMine Safety and Health AdministrationNANot Applicable, except in Section 14 where NA = North AmericaNENot EstablishedNADANew Animal Drug ApplicationNAIFNo Applicable Information FoundNCINational Cancer InstituteNDSLNon-Domestic Substances ListNFPANational Fire Protection AssociationNIOSHNational Institute for Occupational Safety and HealthNPDESNational Pollutant Discharge Elimination SystemNOSNot Otherwise Specified
MSDS Material Safety Data Sheet MSHA Mine Safety and Health Administration NA Not Applicable, except in Section 14 where NA = North America NE Not Established NADA New Animal Drug Application NAIF No Applicable Information Found NCI National Cancer Institute NDSL Non-Domestic Substances List NFPA National Fire Protection Association NIOSH National Institute for Occupational Safety and Health NPDES National Pollutant Discharge Elimination System
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NAIF No Applicable Information Found NCI National Cancer Institute NDSL Non-Domestic Substances List NFPA National Fire Protection Association NIOSH National Institute for Occupational Safety and Health NPDES National Pollutant Discharge Elimination System
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NPDES National Pollutant Discharge Elimination System
1 1 1 0 1 0 1 1 1 1 1 1 1 1 1 1 1 1 1 1
NTP National Toxicology Program
OSHA Occupational Safety and Health Administration
OEL Occupational Exposure Limit
PEL Permissible Exposure Limit (OSHA)
RCRA Resource Conservation and Recovery Act
RQ Reportable Quantity
RTECS Registry of Toxic Effects of Chemical Substances
SARA Superfund Amendments and Reauthorization Act
SDS Safety Data Sheet
STEL Short Term Exposure Limit
TLV Threshold Limit Value (ACGIH)
TPQ Threshold Planning Quantity
TSCA Toxic Substances Control Act
TWA Time Weighted Average/8 Hours Unless Otherwise Noted
UEL Upper Explosive Limit
UN United Nations
USP United States Pharmacopeia
WEEL Workplace Environmental Exposure Level (AIHA)
WHMIS Workplace Hazardous Materials Information System

Signature Manifest

Document Number: SDS053 Revision: 02

Title: Bortezomib for Injection **Effective Date:** 04 May 2022

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Change Request

Name/Signature	Title	Date	Meaning/Reason
Tiffany Seger (TSEGER)	Validation Engineer I - Computerized Systems	04 May 2022, 02:54:19 PM	Approved

Dept Approval

Name/Signature	Title	Date	Meaning/Reason
John Andre (EXT-JANDRE)	Regulatory Affairs Consultant	04 May 2022, 03:49:05 PM	Approved

Final QA Approval/Set Effective Date

Name/Signature	Title	Date	Meaning/Reason
Lori Aleshin (LALESHIN)	Director of Quality Assurance	04 May 2022, 04:04:02 PM	Approved