Loperamide Hydrochloride Capsules, USP Strength: 2 mg Pack Size: HDPE Bottle pack 100's HDPE Bottle pack 500's

Revision No.: 01

EMERGENCY OVERVIEW

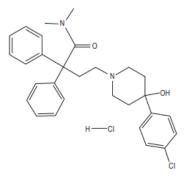
Each Loperamide Hydrochloride Capsules, USP intended for oral administration contains Teriflunomide and excipients generally considered to be non- toxic and non-hazardous in small quantities and under conditions of normal occupational exposure.

Section 1. Identification

Identification of the product

Product Name:Loperamide Hydrochloride Capsules, USPFormula:Loperamide hydrochloride USP, 4-(p-chlorophenyl)-4-hydroxy-
N,N-dimethyl-a,a-diphenyl-1-piperidinebutyramide
monohydrochloride

Chemical Name:



Manufacturer / supplier identification

Company:	Zydus Lifesciences Ltd., N	Aatoda, India
Address:	Zydus Lifescience Limited, Plot No- 1A/1 & 2, Pharmez Special Economic Zone, Sarkhej- Bavla N.H. No. 8A, Near Village Matoda, Tal. Sanand, Dist. Ahmedabad-382 213, India	
Contact for information:	Tel: +91-79-26868100	Fax: +91-79-26868533
Emergency Telephone No.	Tel: +91-79-26868101	
US Customer Service:	1 (877) 993 8779	

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Recommended use /	Loperamide hydrochloride capsules are indicated for the control and
Therapeutic Category	symptomatic relief of acute nonspecific diarrhea in patients 2 years
	of age and older and of chronic diarrhea in adults associated with
	inflammatory bowel disease. Loperamide hydrochloride capsules are
	also indicated for reducing the volume of discharge from
	ileostomies.
Restriction on Use / Contraindications:	 Avoid Loperamide hydrochloride dosages higher than recommended in adult or pediatric patients 2 years of age and older due to the risk of serious cardiac adverse reactions. Loperamide hydrochloride is contraindicated in: Pediatric patients less than 2 years of age due to the risks of respiratory depression and serious cardiac adverse reactions (see WARNINGS). Patients with a known hypersensitivity to loperamide hydrochloride or to any of the excipients. Patients with abdominal pain in the absence of diarrhea. Patients with acute dysentery, which is characterized by blood in stools and high fever. • Patients with bacterial enterocolitis caused by invasive organisms including Salmonella, Shigella and Campylobacter. Patients with pseudomembranous colitis (e.g., Clostridium difficle) associated with the use of broad-spectrum
Section 2. Hazard(s) Identifica	antibiotics.
	• The adverse effects reported during clinical investigations of
	Loperamide hydrochloride are difficult to distinguish from
	symptoms associated with the diarrheal syndrome. Adverse
Adverse Effects	experiences recorded during clinical studies with Loperamide
	hydrochloride were generally of a minor and self-limiting nature.
	They were more commonly observed during the treatment of
	chronic diarrhea.

In Case of Inhalation

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	• The use of higher than recommended loperamide hydrochloride
Over dosage	doses may result in life-threatening cardiac, CNS and respiratory
	adverse reactions.
	Teratology studies have been performed in rats using oral loperamide
	hydrochloride doses of 2.5 mg/kg/day, 10 mg/kg/day and 40
	mg/kg/day and in rabbits using oral doses of 5 mg/ kg/day, 20
	mg/kg/day and 40 mg/kg/day. These studies have revealed no
	evidence of impaired fertility or harm to the fetus at doses up to 10
	mg/kg/day in rats (5 times the human dose based on body surface
Den en en en Commente	area comparison) and 40 mg/kg/day in rabbits (43 times the human
Pregnancy Comments	dose based on body surface area comparison). Treatment of rats with
	oral doses of 40 mg/kg/ day (21 times the human dose based on a
	body surface area comparison) produced marked impairment of
	fertility. The studies produced no evidence of teratogenic activity.
	There are no adequate and well controlled studies in pregnant
	women. Loperamide hydrochloride should be used during pregnancy
	only if the potential benefit justifies the potential risk to the fetus.
Pregnancy Category	Category C

Section 3. Composition / information on ingredients		
Component	Exposure Limit	CAS No.
Principle Component:		
Loperamide Hydrochloride	310 mcg/day	34552-83-5
Inactive ingredients:		
Lactose Monohydrate	Not Found	10039-26-6
Providone K	Not Found	9003-39-8
Sodium starch Glycolate	Not Found	9063-38-1
Microcrystalline Cellulose	Not Found	9004-34-6
Colloidal Silicon Dioxide	Not Found	7631-86-9
Magnesium stearate	Not Found	557-04-0
Section 4. First -aid measures		

Remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give oxygen.

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In Case of Skin Contact	Wash with soap & water for 15 minutes. If irritation persists seek	
III Case of Skill Contact	medical aid.	
In Case of Eye Contact	Flush with copious amounts of water for 15 minutes, separating	
	eyelids with fingers. If irritation persists seek medical aid.	
	Call a physician. Wash out mouth with water. Do not induce	
In Case of Swallowed	vomiting without medical advice	
Section 5. Fire -fighting measu	res	
Fine Fighting Instructions	Wear self-contained breathing apparatus and protective clothing	
Fire Fighting Instructions	to prevent contact with skin and eyes.	
Flammable Properties:	May be combustible at high temperature.	
Hazardous Combustion Products:	Under fire conditions, hazardous fumes will be present.	
Suitable & Unsuitable		
Extinguishing media:	Small fire: dry chemical, CO2 or water spray. Large fire: dry	
	chemical, CO2, alcohol resistant foam or water spray. Do not get	
	water inside containers	
Section 6. Accidental Release M		
Method and material for	On land, sweep or shovel into suitable containers. Minimize	
containment	generation of dust.	
Clean-up Procedures	Do not touch damaged containers or spilled material unless	
	wearing appropriate protective clothing. Wear respirator, chemical	
	safety goggles, rubber boots and heavy rubber gloves. Stop leak if	
	you can do it without risk. Prevent entry into waterways, sewers,	
	basements or confined areas. Shut off all sources of ignition.	
	Evacuate the area. If necessary, employ water fog to disperse the	
	vapors. Absorb the matter with compatible vermiculite or other	
	absorbing material. Place in a suitable container and retain for	
	disposal. Ventilate and clean the affected area. Do not flush into	
	sewerage system or to drains	

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Section 7. Handling and Storage		
	Do not inhale. Avoid contact with eyes, skin and clothing. Avoid	
	prolonged or repeated exposure. Wash thoroughly after handling.	
	Store away from incompatible materials, in a well-ventilated area.	
	Eliminate all sources of ignition. Store in accordance with local	
Precaution and safe handling	regulations. Do not store in unlabelled containers. Containers that	
	have been opened must be carefully resealed and kept upright to	
	prevent leakage. Use appropriate containment to avoid	
	environmental contamination.	
	Store away from incompatible materials, in a well-ventilated area.	
	Eliminate all sources of ignition. Store in accordance with local	
Condition for cofe Storage	regulations. Do not store in unlabelled containers. Containers that	
Condition for safe Storage	have been opened must be carefully resealed and kept upright to	
	prevent leakage. Use appropriate containment to avoid	
	environmental contamination.	
	Store in original container, tightly sealed, protected from direct	
Storage Condition	sunlight and moisture. Preserve in well-closed containers.	
	Protected from light.	

Section 8. Exposure Control / Personal Protection		
Personal Proactive Equipment		
Eyes	Wear appropriate protective eyeglasses or chemical safety goggles.	
	Wear appropriate gloves to prevent skin exposure.	
Skin / Clothing	Wear appropriate protective clothing to minimize contact with	
	skin.	
	Follow WHMIS or OSHA respirator regulations found in 29 CFR	
Respirators	1910.134 or European Standard EN 149. Use a NIOSH/MSHA or	
	European Standard EN 149 approved respirators	

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Thermal Hazards		For products representing a thermal hazard, appropriate Personal		
		Protective Equipment should be used.		
Section 9. Physical and chemical properties				
Physical Sta	ntes			
Solubility	Freely soluble in	Decomposition	Temperature:	225 °C, 437 °F
	methanol and in			
	chloroform;			
	slightly soluble in			
	water and in dilute			
	acids; very			
	slightly soluble in			
	isopropyl alcohol			
Boiling poin	nt	90 °C, 734 °F	Melting Point	220 - 228)°C, (428- 442.4)°F
Evaporation	n rate	No Data	Vapour density	No Data
Reactivity in	n water	No Data	Vapour pressure	No Data
% Volatile I	by volume	No Data	Specific gravity	No Data
Section 10. Stability and Reactivity			1	
Chemical St	tability	Stable under recommended storage conditions		
Condition to		Moisture, sunlight and extreme temperatures		
Incompatib	les	Strong oxidizing agents. Bases. Reducing agents. Acids.		
Hazard Dec	composition	Toxic fumes of carbon monoxide, carbon dioxide, nitrogen oxides		
		and other gases may occur.		
	Toxicological inform		(11141 D50. (
Acute Toxic	eity	Oral: Rat: LD50: (mg/kg): 185 Dermal: Rabbit LD50: (mg/kg): Not		
		available Inhalation: Rat: LC50: (mg/L/4hr): Not available		
	sion/Irritation	Due to lack of data the classification is not possible		
Serious eye	damage / Eye	Due to lack of data the classification is not possible		
Irritation				
Respiratory	Sensitization	Due to lack of data the classification is not possible		

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Skin Sensitization	Based on available data, the classification criteria are not met.	
	Suspected skin sensitiser: The Toolbox profiler Protein binding	
	alerts for skin sensitization by OASIS v1.3 gives an alert for skin	
	sensitisation.	
Section 12. Ecological informat	tion	
Persistence and Degradability	Suspected persistent in the environment: The Danish QSAR	
	database contains information indicating that the substance is	
	predicted as non-readily biodegradable.	
Bio-accumulative Potential	Log Pow: 5.13 (20°C), has potential for bioaccumulation.	
Mobility in Soil	Slightly soluble in water	
	EC50: 48 Hr: Crustacea: (mg/L): Not available	
	LC50: 96 Hr: Fish: (mg/L): Not available	
Torrigity	EC50: 72 or 96 Hr: Algae (or other aqua plants): (mg/L): Not	
Toxicity	available	

Section 13. Disposal Consideration		
	Dispose in accordance with local, state, and federal regulations	
Section 14. Transport Informa	tion	
	The product is not hazardous when shipping via air (IATA), ground	
	(DOT), or sea (IMDG). In accordance with ADR / RID / IMDG /	
	IATA / AND.	
Section 15. Regulatory Informa	ation	
	Under Approval	
Section 16. Other information		
	None	

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Date of issue: 12/09/23

Supersedes edition: New Edition

The information contained herein is based on the state of our knowledge. It characterizes the product with regard to the appropriate safety precautions. It does not represent a guarantee of the properties of the product.