LOPERAMIDE HYDROCHLORIDE- loperamide hydrochloride tablet Chain Drug Consortium, LLC

Loperamide Hydrochloride

Drug Facts

Active ingredient (in each caplet)

Loperamide Hydrochloride USP, 2 mg

Purpose

Anti-diarrheal

Use

controls symptoms of diarrhea, including Travelers' Diarrhea

Warnings

Allergy alert

Do not use if you have ever had a rash or other allergic reaction to loperamide hydrochloride

Heart alert

Taking more than directed can cause serious heart problems or death

Do not use if you have bloody or black stool

Ask a doctor before use if you have

- fever
- mucus in the stool
- a history of liver disease
- a history of abnormal heart rhythm

Ask a doctor or pharmacist before use if you aretaking a prescription drug. Loperamide may interact with certain prescription drugs.

When using this product tiredness, drowsiness or dizziness may occur. Be careful when driving or operating machinery.

Stop use and ask a doctor if

- symptoms get worse
- diarrhea lasts for more than 2 days
- you get abdominal swelling or bulging. These may be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

- drink plenty of clear fluids to help prevent dehydration caused by diarrhea
- find right dose on chart. If possible, use weight to dose; otherwise, use age.

adults and children 12 years and over	2 caplets after the first loose stool; 1 caplet after each subsequent loose stool; but no more than 4 caplets in 24 hours	
children 9-11 years (60-95 lbs)	1 caplet after the first loose stool; ½ caplet after each subsequent loose stool; but no more than 3 caplets in 24 hours	
children 6-8 years (48-59 lbs)	1 caplet after the first loose stool; ½ caplet after each subsequent loose stool; but no more than 2 caplets in 24 hours	
children 2-5 years (34 to 47 lbs)	ask a doctor	
children under 2 years (up to 33 lbs)	do not use	

Other information

- store between 20° 25°C (68° 77°F)
- see side panel for lot number and expiration date
- TAMPER EVIDENT: THIS PRODUCT IS PROTECTED WITH SEALED BLISTER UNITS. DO NOT USE IF ANY ARE TORN OR BROKEN.

Inactive ingredients

anhydrous lactose, croscarmellose sodium, crospovidone, D&C yellow no. 10 aluminum lake, FD&C blue no. 1 aluminum lake, hydrogenated vegetable oil, magnesium stearate, powdered cellulose, pregelatinized starch

Questions?

Call toll-free Monday-Friday 8:30 am to 5 pm EST at **1800-406-7984**.

Distributed By: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087

PRINCIPAL DISPLAY PANEL - 2 mg Caplet Blister Pack Carton

COMPARE TO THE ACTIVE INGREDIENT OF IMODIUM [®]A-D [†]

See New Warnings and Directions

Premier Value ®

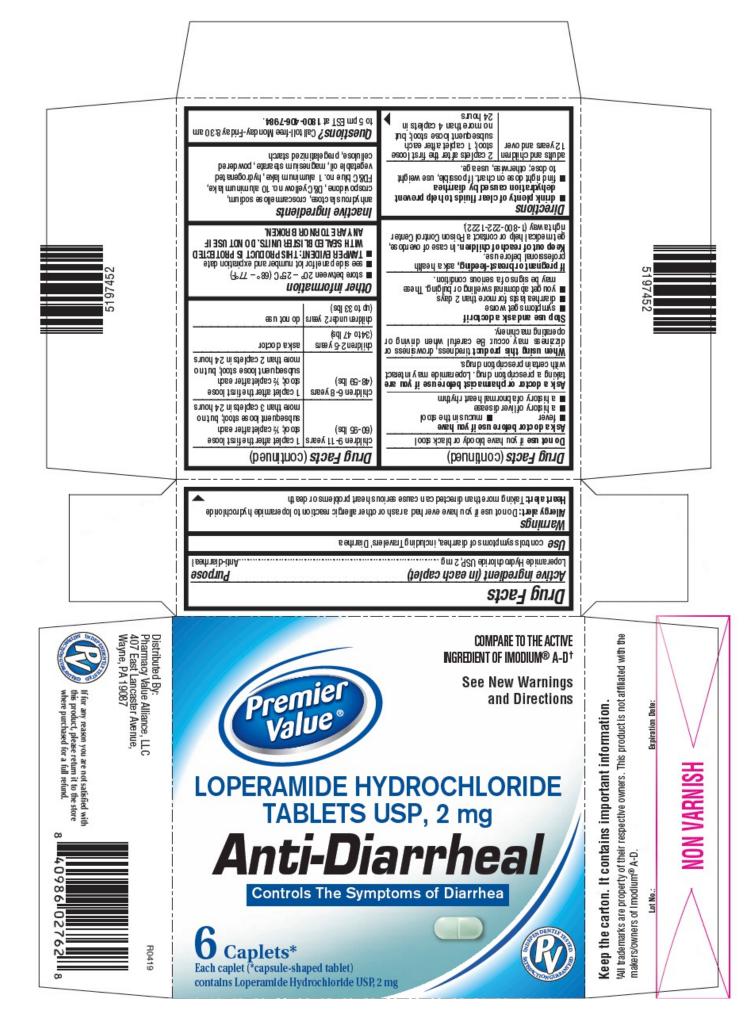
LOPERAMIDE HYDROCHLORIDE TABLETS USP, 2 mg

Anti-Diarrheal

Controls The Symptoms of Diarrhea

6 Caplets* Each caplet (*capsule-shaped tablet) contains Loperamide Hydrochloride USP, 2 mg

INDEPENDENTLY TESTED PV SATISFACTION GUARANTEED





LOPERAMID		OCHLORIDE						
loperamide hydro			-					
Product Infor	mation							
Product Type		HUMAN OTC DRUG	G It	em Co	ode (S	ource)	NDC:68016-123	
Route of Admin	istration	ORAL						
Active Ingred	ient/Active	Moiety						
	Ingre	edient Name				Basis of St	rength	Strength
LOPERAMIDE HYD UNII:6X9OC3H4II)	ROCHLORIDE	(UNII: 77TI35393C)	(LOPERAMII	DE -		LOPERAMIDE HYDROCHLORIDE		2 mg
less stilles la sure	alla untra							
Inactive Ingre	alents						-	
			lame				5	trength
ANHYDROUS LACT								
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48) CROSPOVIDONE (15 MPA.S AT 5%) (UNII: 68401960MK)								
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)								
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)								
HYDROGENATED	OTTONSEED	OIL (UNII: Z82Y2C6	65EA)					
MAGNESIUM STEA	RATE (UNII: 70	097M6I30)						
POWDERED CELL	ULOSE (UNII: S	MD1X3XO9M)						
STARCH, CORN (UNII: 08232NY3SJ)								
Product Chara	acteristics							
Color	gree	en	Score				2 pieces	
Shape		SULE	Size			9mm		
Flavor			Imprint Code			123		
Contains								
Packaging								
# Item Code	Pa	ackage Descrip	tion		Mark	eting Start Date		eting End Date
1 NDC:68016- 123-06	6 in 1 BLISTER Product	R PACK; Type 0: Not	a Combina	tion	02/01/1	993		

M	larketing	Information		
4	NDC:68016- 123-97 96 in 1 BOTTLE; Type 0: Not a Combination Product 02/01/1993			
3	NDC:68016- 123-24	24 in 1 BLISTER PACK; Type 0: Not a Combination Product	02/01/1993	
2	NDC:68016- 123-18	18 in 1 BLISTER PACK; Type 0: Not a Combination Product	02/01/1993	

Labeler - Chain Drug Consortium, LLC (101668460)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment							
Name	Address	ID/FEI	Business Operations				
Ohm Laboratories Inc.		051565745	manufacture(68016-123)				

Revised: 1/2024

Chain Drug Consortium, LLC