LOPERAMIDE HYDROCHLORIDE - loperamide hydrochloride tablet
Amerisource Bergen

Drug Facts

Active ingredient (in each tablet)

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 are

taking 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 (1-800-222-1222) right away.

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

Other information

- store at 20° to 25°C (68° to 77°F).
- **do not use if carton or blister unit is open or torn**
- Meets USP dissolution test 2
- see side panel for lot number and expiration date

Inactive ingredients

colloidal silicon dioxide, D & C yellow No. 10 aluminum lake, FD & C blue No. 1, lactose

monohydrate, magnesium stearate, microcrystalline cellulose, and sodium starch glycolate

Questions or comments?

call **1-855-274-4122**

Distributed By
AmerisourceBergen
1 West First Avenue
Conshohocken, PA 19428
Questions or Concerns?

www.mygnp.com

Made in India

Code : TS/DRUGS/22/2009

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 2 mg Blister Carton 12(2 x 6's Tablets)

GOOD NEIGHBOR PHARMACY®

***Compare to Imodium® A-D active ingredient**
NDC 46122-738-53

Loperamide Hydrochloride Tablets USP 2 mg

Anti-Diarrheal

Controls the symptoms of diarrhea

12 Tablets actual size

Drug Facts

Active ingredient (in each tablet)	*Purpose*
Loperamide hydrochloride USP 2 mg	Anti-diarrheal

Use

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Drug Facts (continued)

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Drug Facts (continued)

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GOOD NEIGHBOR PHARMACY®

*Compare to Imodium® A-D active ingredient

NDC 46122-738-53

Loperamide Hydrochloride Tablets USP 2 mg

Anti-Diarrheal

Controls the symptoms of diarrhea

12 Tablets

actual size

ABC#: 10276227

0 87701 43500 3

To Open The Blister

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1 West First Avenue
Conshohocken, PA 19428
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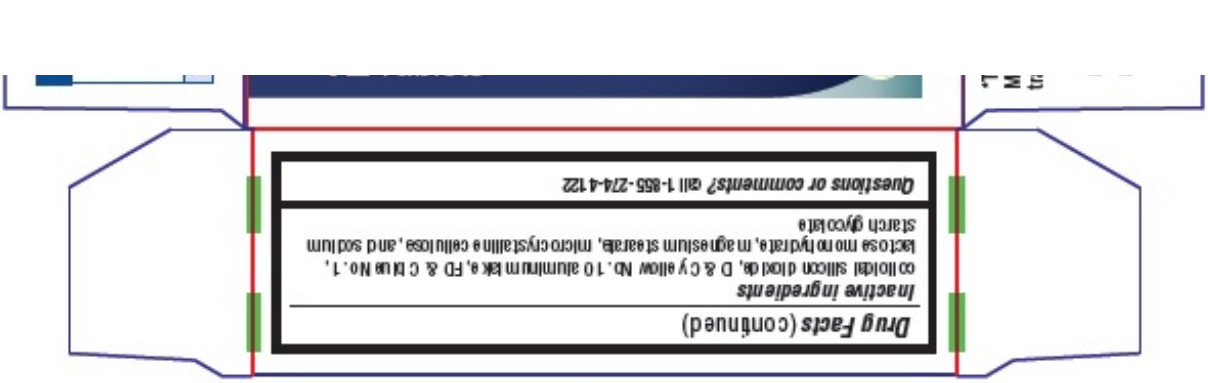
Made in India
Code: TS/DRUGS/22/2009

This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Imodium® A-D.

Lot: P1050530 LM-5068

Exp: NVZ

Batch Coding Area



LOPERAMIDE HYDROCHLORIDE

loperamide hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:46122-738
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LOPERAMIDE HYDROCHLORIDE (UNII: 77TI35393C) (LOPERAMIDE - UNII:6X9OC3H4II)	LOPERAMIDE HYDROCHLORIDE	2 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics			
Color	GREEN (Light Green)	**Score**	2 pieces
Shape	CAPSULE (Biconvex)	**Size**	10mm
Flavor		**Imprint Code**	L;28
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-738-53	2 in 1 CARTON	07/05/2023	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:46122-738-62	4 in 1 CARTON	07/20/2023	
2		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206548	07/05/2023	

Labeler - Amerisource Bergen (007914906)

Registrant - Aurohealth LLC (078728447)

Establishment

Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(46122-738) , MANUFACTURE(46122-738)

Revised: 3/2024 Amerisource Bergen