

ANTI-DIARRHEAL- loperamide hci tablet
Lil' Drug Store Products, Inc.

ANTI-DIARRHEAL loperamide hcl tablet

Drug Facts

Active Ingredient (in each tablet)

Loperamide HCl, USP 2 mg

Purpose

Anti-diarrheal

Use

Controls symptoms of diarrhea, including Travelers' Diarrhea

Warnings

Allergy alert

Allergy alert:Do not use if you have ever had a rash or other allergic reaction to Loperamide HCl.

Heart alert

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 producttiredness, 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 hour
children 9 - 11 years (60 - 95 lbs)	1 caplet after the first loose stool; 1/2 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; 1/2 caplet after each subsequent loose stool; but no more than 2 caplets in 24 hours
children 2 - 5 years (34 - 47 lbs)	ask a doctor
children under 2 (up to 33 lbs)	do not use

Other Information

Other information

- store between 68-77°F (20-25°C)

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 or Comments?

Call **1-800-351-2000 (8:30 AM - 5:00 PM CST)**

*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc, McNeil Consumer Healthcare Division, distributor of Imodium® A-D

Distributed by: Lil' Drug Store Products, Inc., Cedar Rapids, IA 52404

2 mg Loperamide Tablet

Compare to the active ingredient in Imodium® A-D*

Diamode

Anti-Diarrheal Relief

Loperamide HCl

Tablets, USP 2 mg

[tablet image]

actual size

1 CAPLET IN EACH PACKET

READ DRUG FACTS AND INSTRUCTIONS ON PACKET

COMPARE TO THE ACTIVE INGREDIENT IN

Imodium® A-D*

Diamode

Anti-Diarrheal

Loperamide HCl
Tablets, USP 2 mg

Relief



Actual Size

**1 CAPLET
IN EACH PACKET**

READ DRUG FACTS AND INSTRUCTIONS ON PACKET

ANTI-DIARRHEAL

loperamide hci tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:29485-9286
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
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
CROSPROVIDONE (15 MPA.S AT 5%) (UNII: 68401960MK)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYDROGENATED COTTONSEED OIL (UNII: Z82Y2C65EA)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	

Product Characteristics

Color	green	Score	no score
Shape	OVAL	Size	9mm
Flavor		Imprint Code	123
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29485-9286-2	25 in 1 BOX	07/27/2016	
1		1 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:29485-9286-3	30 in 1 BOX	07/27/2016	
2		1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA074091	07/27/2016	

Labeler - Lil' Drug Store Products, Inc. (093103646)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(29485-9286)

Revised: 12/2024

Lil' Drug Store Products, Inc.