

ANTI DIARRHEAL- loperamide hydrochloride tablet, film coated
Meijer Distribution Inc

Meijer Distribution, Inc. Anti-Diarrheal Drug Facts

Active ingredient (in each caplet)

Loperamide HCl 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 HCl

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 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; 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 to 47 lbs)	ask a doctor
children under 2 years (up to 33 lbs)	do not use

Other information

- store at 20°-25°C (68°-77°F)
- see end panel for lot number and expiration date

Inactive ingredients

anhydrous lactose, carnauba wax, D&C yellow no. 10 aluminum lake, FD&C blue no. 1 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, pregelatinized starch

Questions or comments?

1-800-719-9260

Principal Display Panel

meijer®

Compare to Imodium® A-D active ingredient

anti-diarrheal

Loperamide Hydrochloride Tablets, 2 mg Anti-diarrheal

SEE NEW WARNINGS

Controls the Symptoms of Diarrhea

12 Caplets*

*capsule-shaped tablets

actual size

PID 467573

DO NOT USE IF BLISTER UNIT IS BROKEN OR TORN



meijer

NDC 41250-224-53

Compare to Imodium® A-D active ingredient**

anti-diarrheal

Loperamide Hydrochloride Tablets, 2 mg
Anti-diarrheal

SEE NEW WARNINGS

Controls the Symptoms of Diarrhea

12 Caplets*

*capsule-shaped tablets



actual size

22453 6E C1 1

OPEN OTHER ENDS

Drug Facts

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- a history of an abnormal heart rhythm

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Drug Facts (continued)

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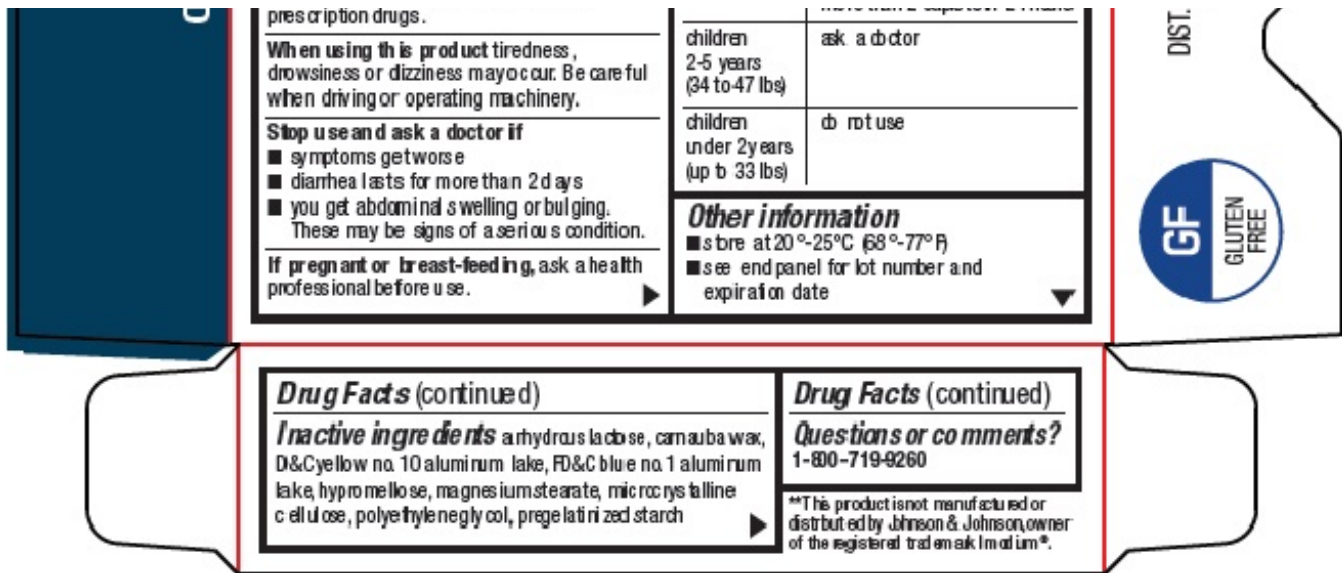
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BY MEIJER DISTRIBUTION, INC.
GRAND RAPIDS, MI 49544
www.meijer.com



ANTI DIARRHEAL

loperamide hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-224
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
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics

Color	GREEN	Score	2 pieces
Shape	OVAL	Size	10mm
Flavor		Imprint Code	L2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-224-53	12 in 1 CARTON	03/24/2003	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:41250-224-67	1 in 1 CARTON	02/08/2004	08/31/2021
2		48 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:41250-224-82	1 in 1 CARTON	04/01/2015	09/30/2021
3		200 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:41250-224-91	6 in 1 CARTON	03/16/2016	
4		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:41250-224-80	1 in 1 CARTON	04/30/2003	12/19/2016
5		96 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:41250-224-01	2 in 1 CARTON	04/30/2003	12/19/2016
6		96 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075232	03/05/2003	

Labeler - Meijer Distribution Inc (006959555)

Revised: 10/2022

Meijer Distribution Inc