

ANTI-DIARRHEAL- loperamide hcl tablet
Chain Drug Consortium

Premier Value 44-375

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

- a 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 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 (34-47 lbs)	ask a doctor
children under 2 years (up to 33 lbs)	do not use

Other information

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store between 20°-25°C (68°-77°F)
- see end flap for expiration date and lot number

Inactive ingredients

corn starch, D&C yellow #10 aluminum lake, dibasic calcium phosphate dihydrate, FD&C blue #1 aluminum lake, magnesium stearate, microcrystalline cellulose, silicon dioxide

Questions or comments?

Call 1-800-426-9391 8:30 AM-4:00 PM ET, Monday-Friday

Principal Display Panel

*COMPARE TO THE ACTIVE INGREDIENT IN IMODIUM® A-D

Premier Value®

Anti-diarrheal

Loperamide HCl Tablets, 2 mg

ANTI-DIARRHEAL

Controls the symptoms of diarrhea

6 Caplets

actual size

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Imodium® A-D.

50844 ORG061937545

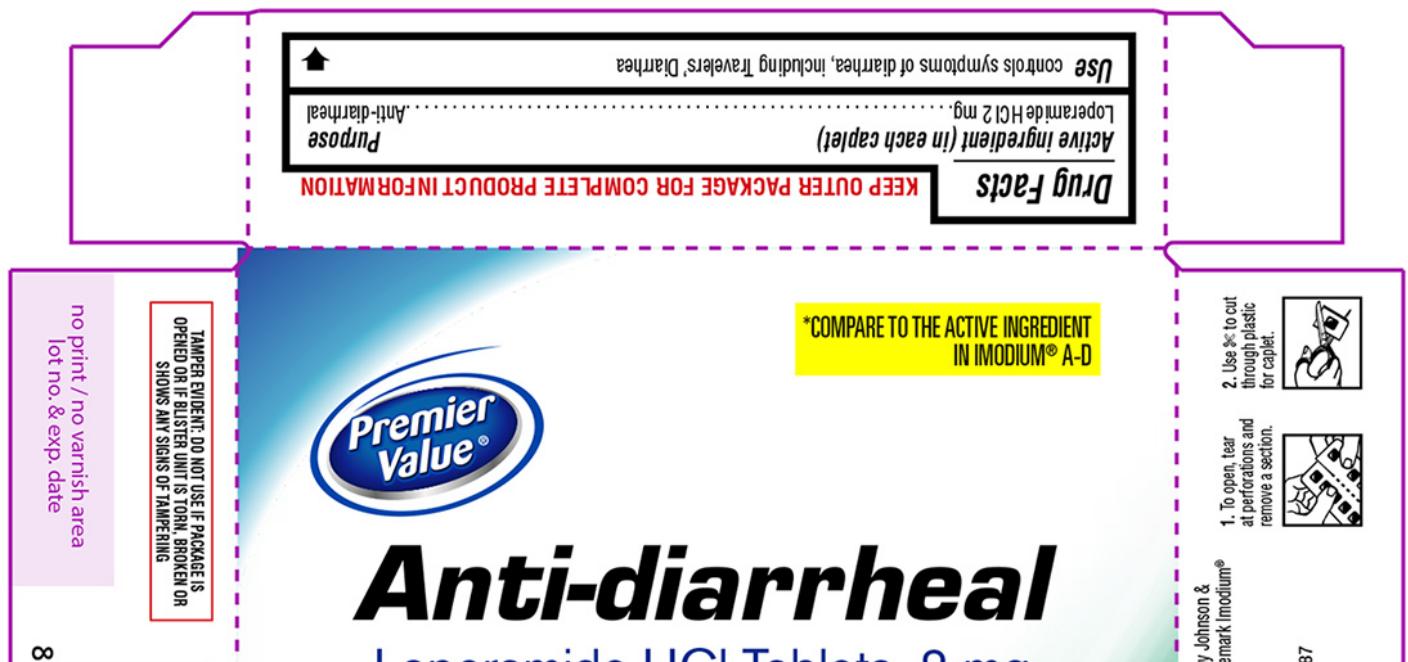
**Distributed by:
Pharmacy Value Alliance, LLC
407 East Lancaster Avenue, Wayne, PA 19087**

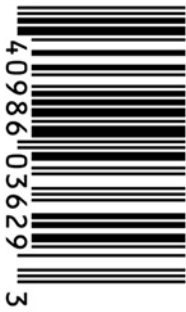
INDEPENDENTLY TESTED

PV

SATISFACTION GUARANTEED

If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.





Loperamide HCl Tablets, 2 mg

ANTI-DIARRHEAL

Controls the symptoms of diarrhea

6 Caplets

actual size



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Distributed By: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 190



If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.

B-1580-375-45
ORG061937545

Drug Facts (continued)

Other Information ■ store between 20°-25°C (68°-77°F) ■ see end flap for expiration date and lot number ■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN

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Drug Facts (continued)

Warnings 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 ■ a fever ■ mucus in the stool ■ a history of liver disease ■ 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.

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Premier Value 44-375

ANTI-DIARRHEAL

loperamide hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-301
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LOPERAMIDE HYDROCHLORIDE (UNII: 77TI35393C) (LOPERAMIDE -	LOPERAMIDE	2 mg

UNII:6X9OC3H4II)

HYDROCHLORIDE

2 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	green	Score	2 pieces
Shape	OVAL	Size	10mm
Flavor		Imprint Code	44;375
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-301-06	1 in 1 CARTON	12/06/2019	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:68016-301-18	3 in 1 CARTON	12/06/2019	
2		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:68016-301-24	4 in 1 CARTON	12/06/2019	
3		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	ANDA076497	12/06/2019	

Labeler - Chain Drug Consortium (101668460)**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(68016-301) , pack(68016-301)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(68016-301)

Revised: 12/2023

Chain Drug Consortium