

BASIC CARE LOPERAMIDE HYDROCHLORIDE- loperamide hydrochloride suspension
Amazon.com Services LLC

Amazon Loperamide Hydrochloride Drug Facts

Active ingredient (in each 7.5 mL)

Loperamide HCl 1 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.
- shake well before using
- use only enclosed dosing cup specifically designed for use with this product. Do not use any other dosing device.
- mL = milliliter

adults and children 12 years and over	30 mL after the first loose stool; 15 mL after each subsequent loose stool; but no more than 60 mL in 24 hours
children 9-11 years (60-95 lbs)	15 mL after the first loose stool; 7.5 mL after each subsequent loose stool; but no more than 45 mL in 24 hours
children 6-8 years (48-59 lbs)	15 mL after the first loose stool; 7.5 mL after each subsequent loose stool; but no more than 30 mL 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

- **each 30 mL contains:** sodium 15 mg
- store between 20-25°C (68-77°F)

Inactive ingredients

anhydrous citric acid, carboxymethylcellulose sodium, D&C yellow no. 10, FD&C blue no. 1, glycerin, microcrystalline cellulose, natural and artificial mint flavor, propylene glycol, purified water, simethicone, sodium benzoate, sucralose, titanium dioxide, xanthan gum

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

Anti-Diarrheal Oral Solution

Compare to Imodium® A-D active ingredient

For Ages 6 Years And Up

Loperamide Hydrochloride

Oral Solution, 1 mg per 7.5 mL

Anti-Diarrheal

Controls the symptoms of Diarrhea

Mint Flavor

See New Warning and Directions

4 FL OZ (120 mL)

Drug Facts

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(in each 7.5 mL)
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Peel open to read complete directions and warnings before purchase

amazon

basic care

Anti-Diarrheal Oral Solution

NDC 72288-758-26

For Ages 6 Years And Up

Loperamide Hydrochloride

Oral Solution, 1 mg per 7.5 mL

Anti-Diarrheal

Controls the Symptoms of Diarrhea
Mint Flavor

See New Warning and Directions

4 FL OZ (120 mL)

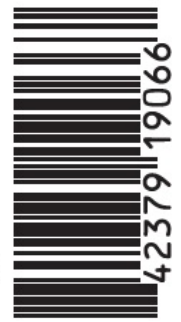
Compare to
Imodium® A-D
active
ingredient*

Do not use if
printed plastic
neckband is
broken or missing

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GLUTEN FREE

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



: 75826 BN FL

Drug Facts (continued)

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artificial mint flavor, propylene glycol, purified water,
simethicone, sodium benzoate, sucralose, titanium
dioxide, xanthan gum

Questions or comments? 1-800-719-9260

BASIC CARE LOPERAMIDE HYDROCHLORIDE

loperamide hydrochloride suspension

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72288-758	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
LOPERAMIDE HYDROCHLORIDE (UNII: 77TI35393C) (LOPERAMIDE - UNII:6X9OC3H4II)		LOPERAMIDE HYDROCHLORIDE	1 mg in 7.5 mL	
Inactive Ingredients				
Ingredient Name			Strength	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)				
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
GLYCERIN (UNII: PDC6A3C0OX)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
DIMETHICONE (UNII: 92RU3N3Y1O)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
XANTHAN GUM (UNII: TTV12P4NEE)				
Product Characteristics				
Color	GREEN (opaque, viscous))	Score		
Shape		Size		
Flavor	MINT	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72288-758-26	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/21/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA091292		12/21/2020	

