GOOD SENSE ANTI DIARRHEAL- loperamide hydrochloride tablet, film coated Proficient Rx LP

Perrigo 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

- 1. fever
- 2. mucus in the stool
- 3. a history of liver disease

Ask a doctor or pharmacist before use if you are

taking antibiotics

When using this product

1. tiredness, drowsiness or dizziness may occur. Be careful when driving or operating machinery.

Stop use and ask a doctor if

- 1. symptoms get worse
- 2. diarrhea lasts for more than 2 days
- 3. 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	1 caplet after the first loose stool; 1/2 caplet after each subsequent loose
(60-95 lbs)	stool; but no more than 3 caplets in 24 hours
children 6-8 years	1 caplet after the first loose stool; 1/2 caplet after each subsequent loose
(48-59 lbs)	stool; but no more than 2 caplets in 24 hours
children 2-5 years	ask a doctor
(34 to 47 lbs)	
children under 2 years	do not use
(up to 33 lbs)	

Other information

- 1. store at 20°-25°C (68°-77°F)
- 2. 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

Anti-Diarrheal

See New Warnings and Directions

Loperamide Hydrochloride Tablets, 2 mg

Anti-Diarrheal

Controls the Symptoms of Diarrhea

Actual Size

Compare to active ingredient of Imodium® A-D

100% SATISFACTION GUARANTEED

12 Caplets*

*Capsule-Shaped Tablets

Relabeled by:

Proficient Rx LP





NDC 71205-079-12

Lot #:00000 Exp. 00/00/00 SN# MASTER

Loperamide HCI 2mg

#12 Tablets Lot #:00000 NDC 71205-079-12

SN# MASTER Exp:00/00/00

Loperamide HCI 2mg

#12 Tablets
Lot #:00000
NDC 71205-079-12

SN# MASTER Exp:00/00/00

Loperamide HCI 2mg

NDC 71205-079-12

SN# MASTER Exp:00/00/00

Tablets Lot #:00000

> Relabeled By: Proficient Rx LP Thousand Oaks, CA 91320

Loperamide HCI 2mg

#12 **Tablets**

Each tablet contains: Loperamide HCI, 2mg Anti diarrheal

See Box

Product ID: SL007912 Dist. By: Perrigo Allegan, MI 49010 Store at 20°-25°C (68°-77°F)

Keep medication out of the reach of children

GOOD SENSE ANTI DIARRHEAL

loperamide hydrochloride tablet, film coated

Product Information

Product Type HUMAN OTC DRUG NDC:71205-079(NDC:0113-0224) Item Code (Source)

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) MICRO CRYSTALLINE CELLULO SE (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	10 mm
Flavor		Imprint Code	L2
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:71205-079-12	12 in 1 CARTON	08/01/2018	
1	$1\ \text{in}\ 1\ \text{BLISTER}$ PACK; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075232	02/25/2003	

Labeler - Proficient Rx LP (079196022)

Establishment				
Name	Address	ID/FEI	Business Operations	
Proficient Rx LP		079196022	RELABEL(71205-079)	

Revised: 10/2019 Proficient Rx LP