GOOD SENSE ANTI DIARRHEAL- loperamide hydrochloride tablet, film coated Northwind Pharmaceuticals

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
- 4. 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

- 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 (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

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

Inactive ingredients

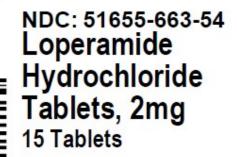
anhydrous lactose, carnauba wax, D&C yellow no. 10 aluminum lake, FD&C blue no. 1 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, pregelatinized starch

Questions or comments?

1-800-719-9260

Principal Display Panel

NDC: 51655-663-54



Dosage: See package insert Store at 20° - 25°C (68° - 77°F) (See USP Controlled Room Temperature)

Keep out of the reach of children. Store in original container.

Do not use if you have ever had a rash or other allergic reaction to loperamide HCl. Taking more than directed can cause serious heart problems or death.

LCN#: 00 Rev. A 03/21

Ask a doctor before use if you have fever, mucus in the stool, a history of liver disease, or a history of abnormal heart rhythm. Active Ingredient (in each caplet) Loperamide HCI 2mg Repackaged From: 0113-0224-62 Perrigo, Lot 0000000000

you have bloody or black stool

Do not use if

Repackaged By: Northwind Pharmaceuticals Indianapolis, IN 46203 GTIN: 00351655663547 S/N: 00000000000000000000

GOOD SENSE ANTI DIARRHEAL

loperamide hydrochloride tablet, film coated

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:51655-663(NDC:0113-0224)

Route of Administration ORAL

Active Ingredient/Active Moiety

Basis of Strength Strength **Ingredient Name** LOPERAMIDE HYDROCHLORIDE (UNII: 77TI35393C) (LOPERAMIDE -**LOPERAMIDE** 2 mg UNII:6X9OC3H4II) **HYDROCHLORIDE**

Inactive Ingredients Ingredient Name Strenath ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK) CARNAUBA WAX (UNII: R12CBM0EIZ) HYPROMELLOSES (UNII: 3NXW29V3WO) MAGNESIUM STEARATE (UNII: 70097M6I30) CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U) POLYETHYLENE GLYCOL (UNII: 3M/Q0SDW1A) 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	g		
# Item C	ode Package Descript	ion Marketing Star Date	t Marketing End Date
1 NDC:51655	5-663- 15 in 1 BAG; Type 0: Not a Comb Product	oination 04/02/2021	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA075232	04/02/2021		

Labeler - Northwind Pharmaceuticals (036986393)

Registrant - Northwind Pharmaceuticals (036986393)

Establishment				
Name	Address	ID/FEI	Business Operations	
Northwind Pharmaceuticals		036986393	relabel(51655-663)	

Revised: 1/2023 Northwind Pharmaceuticals