IMODIUM A-D- loperamide hydrochloride tablet, film coated
Johnson & Johnson Consumer Inc.

Imodium ®

A-D

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)
- **do not use if blister unit is broken or torn**

Inactive ingredients

anhydrous lactose, carnauba wax, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, pregelatinized starch

Questions or comments?

1-877-895-3665 (toll-free) or **215-273-8755** (collect)

PRINCIPAL DISPLAY PANEL

NDC 50580-317-06

Imodium ®
A-D

Loperamide Hydrochloride Tablets, 2mg
Anti-Diarrheal

Anti-Diarrheal
Caplets

Controls the symptoms
of diarrhea

Actual Size

*Capsule-Shaped Tablets

12 Caplets*

NDC 50580-317-06

Imodium®

A-D Loperamide Hydrochloride Tablets, 2mg Anti-Diarrheal

Anti-Diarrheal

7N253 MN C3

Drug Facts

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Drug Facts (continued)

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1. Tear off a section
2. Tear at notch. Use scissors if necessary

How can we help?
1-877-895-3665

Active ingredient made in Italy
Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
McNeil Consumer Healthcare Division
Fort Washington, PA 19034 USA
 www.imodium.com
30048667

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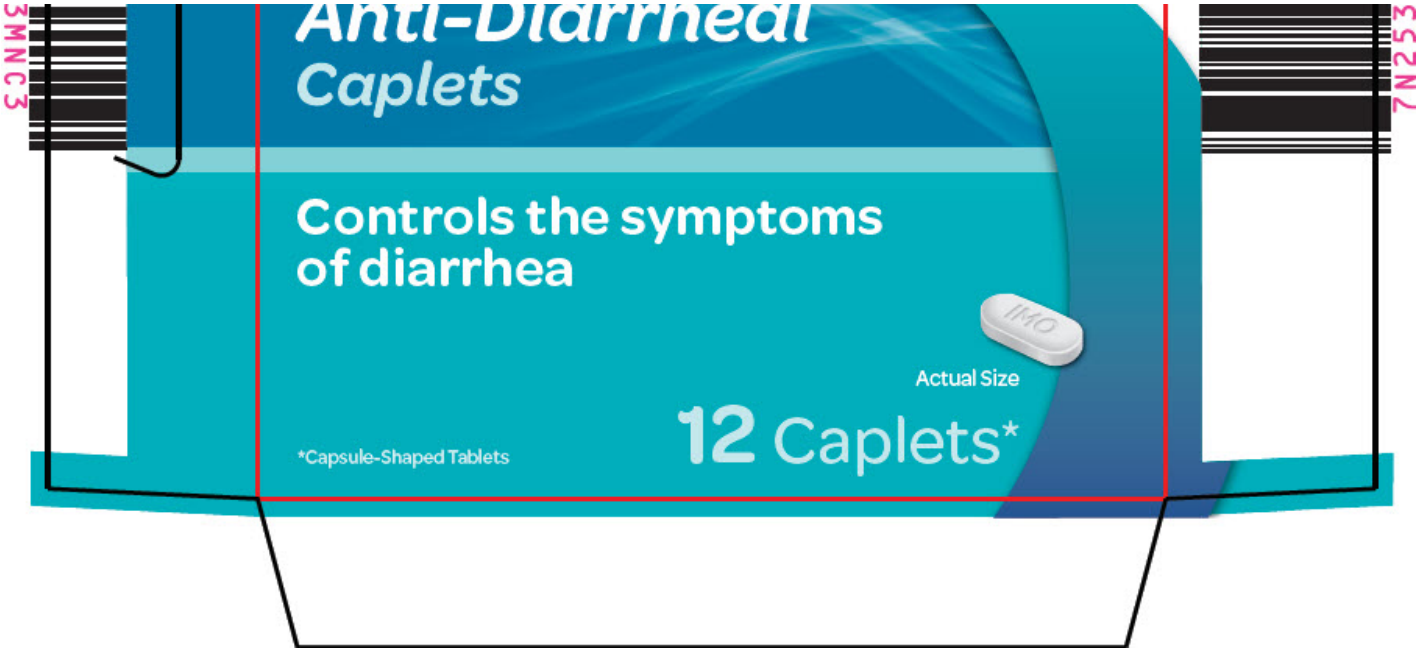

IMODIUM A-D

loperamide hydrochloride tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50580-317
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: 3WQ0SDW1A)	

Product Characteristics			
Color	white	**Score**	2 pieces
Shape	OVAL	**Size**	11mm
Flavor		**Imprint Code**	IMO;2;MG
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-317-01	1 in 1 CARTON	07/13/2015	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:50580-317-03	4 in 1 CARTON	07/13/2015	
2		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:50580-317-04	6 in 1 CARTON	07/13/2015	12/31/2018
3		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:50580-317-05	8 in 1 CARTON	07/13/2015	10/31/2021
4		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:50580-317-06	2 in 1 CARTON	02/10/2020	
5		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:50580-317-12	2 in 1 CARTON	01/29/2024	
6		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	ANDA075232	07/13/2015	

Labeler - Johnson & Johnson Consumer Inc. (878046358)

Revised: 4/2024 Johnson & Johnson Consumer Inc.