UP AND UP ANTI DIARRHEAL- loperamide hcl suspension Target Corporation

Target Corporation Anti-Diarrheal 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 under 6 years (up 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

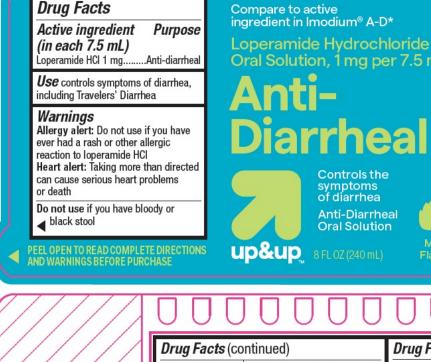
Questions or comments?

1-888-547-7400

Package/Label Principal Display Panel

Compare to active ingredient in Imodium[®] A-D Loperamide Hydrochloride Oral Solution, 1 mg per 7.5 mL Anti-Diarrheal Controls the symptoms of diarrhea Anti-Diarrheal Oral Solution up&up[™] 8 FL OZ (240 mL) Mint Flavor

Drug Facts



Oral Solution, 1 mg per 7.5 mL Consumer Inc., Diarrheal distributor of Imodium® A-D. Controls the symptoms of diarrhea Anti-Diarrheal **Oral Solution** up&up 8 FL 0Z (240 mL) Flavor

DO NOT USE IF **PRINTED PLASTIC**

*This product is not manufactured or distributed by Johnson & Johnson



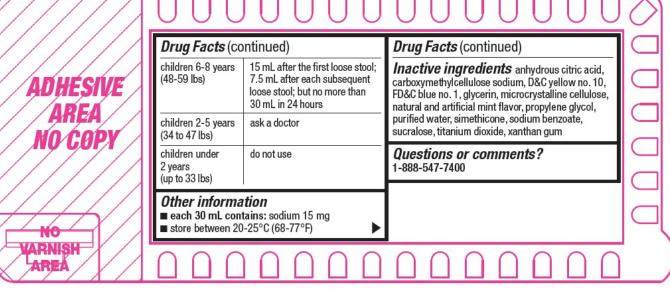
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ſ	Drug Facts (continued)	Drug Facts (cor	ntinued)	
	Ask a doctor before use if you have ■ fever ■ mucus in the stool ■ a history of liver disease ■ a history of abnormal heart rhythm	Directions ■ drink plenty of clo dehydration caus	ear fluids to help prevent ed by diarrhea	
	Ask a doctor or pharmacist before use if you are taking a prescription drug. Loperamide may interact with certain prescription drugs.	dose; otherwise us ■ shake well before	using	
	When using this product tiredness, drowsiness or dizziness may occur. Be careful when driving or operating machinery.		dosing cup specifically designed roduct. Do not use any other	AREA
	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.	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	NO CO
	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)	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	

NDC:11673-900- 34	240 mL in 1 BC Product	OTTLE; Type 0: Not a Comb	ination 02/2	26/2016			
titem Code	Pa	ckage Description	м	arketing Start Date	Marl	keting I Date	End
Packaging							
ontanis							
lavor Contains				Imprint Code			
hape lavor	MINT			Size			
olor	GREEN (OP	aque, viscous)		Score			
				500-00			
Product Chara	acteristics						
	(UNII: ETJ/26XB	U4)					
ANTHAN GUM (UN							
ITANIUM DIOXIDE		-					
UCRALOSE (UNII:							
ODIUM BENZOAT	-	E5EU)					
VATER (UNII: 059Q							
ROPYLENE GLYC		Q167V3)					
		(UNII: OP1R32D61U)					
IYCERIN (UNII: PD	DC6A3C0OX)						
D&C BLUE NO. 1	(UNII: H3R47K3	TBD)					
&C YELLOW NO.	10 (UNII: 355W	/5USQ3G)					
ARBOXYMETHYL	CELLULOSE S	DDIUM, UNSPECIFIED (UN	III: K679OBS	311)			
NHYDROUS CITR	IC ACID (UNII:)	(F417D3PSL)					
		Ingredient Name				Stren	gth
nactive Ingre	dients						
NII:6X9OC3H4II)				HYDROCHLORIDE		in 7.5 r	ΠĽ
	ROCHLORIDE	(UNII: 77TI35393C) (LOPERA	MIDE -			1 mg	
	Ingred	lient Name		Basis of Str	ength	Stren	ngth
Active Ingredi	ent/Active	Moiety					
Route of Admini	stration	ORAL					
Product Type		HUMAN OTC DRUG	Item Code	e (Source)	NDC:1	1673-900	
Product Infor	mation						

Category	Citation	Date	Date
ANDA	ANDA091292	02/26/2016	

Labeler - Target Corporation (006961700)

Revised: 4/2025

Target Corporation