LEADER LOPERAMIDE HYDROCHLORIDE- loperamide hcl liquid Cardinal Health

Cardinal Health Loperamide Hydrochloride Oral Suspension 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

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

Ask a doctor or pharmacist before use if you are

taking antibiotics

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
- only use attached measuring cup to dose product

adults and children 12 years and	30 mL (6 tsp) after the first loose stool; 15 mL (3 tsp) after each subsequent loose stool; but no more than 60 mL (12 tsp) in 24 hours
over	
children 9-11 years (60-95 lbs)	15 mL (3 tsp) after the first loose stool; 7.5 mL (1 1/2 tsp) after each subsequent loose stool; but no more than 45 mL (9 tsp) in 24 hours
children 6-8 years (48-59 lbs)	15 mL (3 tsp) after the first loose stool; 7.5 mL (1 1/2 tsp) after each subsequent loose stool; but no more than 30 mL (6 tsp) in 24 hours
children under 6 years (up to 47 lbs)	ask a doctor

Other information

- each 30 mL (6 tsp) contains: sodium 13 mg
- store between 20-25°C (68-77°F)
- do not use if printed plastic neckband is broken or missing
- see side panel for lot number and expiration date

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

Compare to Imodium® A-D active ingredient Loperamide Hydrochloride Oral Suspension Anti-Diarrheal Controls the Symptoms of Diarrhea Mint Flavor 1 mg Loperamide Hydrochloride per 7.5 mL



Drug Facts (continued)	Drug Facts (co	ontinued)	
Ask a doctor before use if you have ■ fever ■ mucus in the stool ■ a history of liver disease	Control Center right away. (1-800-222-1222)		
Ask a doctor or pharmacist before use if you are taking antibiotics	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 ■ only use attached measuring cup to dose product		
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.	adults and children 12 years and over	30 mL (6 tsp) after the first loose stool; 15 mL (3 tsp) after each subsequent loose stool; but no more than 60 mL (12 tsp) in 24 hours	
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	children 9-11 years (60-95 lbs)	15 mL (3 tsp) after the first loose stool; 7.5 mL (1 1/2 tsp) after each subsequent loose stool; but no more than 45 mL (9 tsp) in 24 hours	

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Drug Facts (continued)		Drug Facts (continued)	
children 6-8 years (48-59 lbs)	15 mL (3 tsp) after the first loose stool; 7.5 mL (1 1/2 tsp) after each subsequent loose	■ do not use if printed plastic neckband is broken or missing ■ see side panel for lot number and expiration date	
	stool; but no more than 30 mL (6 tsp) in 24 hours	Inactive ingredients anhydrous citric acid, carboxymethylcellulose sodium, D&C	
children under 6 years (up to 47 lbs)	ask a doctor	yellow no. 10, FD&C blue no. 1, glycerin, microcrystalline cellulose, natural and artificial mint flavor, propylene glycol, purified water, simethicone, sodium benzoate, sucralose,	
Other information ■ each 30 mL (6 tsp) contains: sodium 13 mg ■ store between 20-25°C (68-77°F)		titanium dioxide, xanthan gum (<i>Questions or comments?</i> 1-800-719-9260	
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LEADER LOPERAMIDE HYDROCHLORIDE

lopera	amide hcl liquid							
Pro	duct Informat	tion						
Prod	uct T yp e		HUMAN OTC DRUG	Item Code (Source)		NDC:37205-584		
Rout	e of Administra	tion	ORAL					
Activ	ve Ingredient	t/Active Moi	ety					
		Ingre	dient Name		Basis of Stre	ngth Stre	ngth	
	LOPERAMIDE HYDROCHLORIDE (UNII: 77TI35393C) (LOPERAMIDE -		-	LOPERAMIDE	1 mg			
UNII:6	SX9OC3H4II)				HYDROCHLORIDE	in 7.5	mL	
Inac	tive Ingredie	nts						
mue	uve ingreuie	1105	Ingredient Name			Streng	eth	
ANHY	DROUS CITRIC	ACID (UNII: XF4	-				,	
			DIUM (UNII: K679OBS311)					
D&C	YELLOW NO. 10	0 (UNII: 35SW5U	SQ3G)					
FD&C	C BLUE NO. 1 (U	NII: H3R47K3TBI))					
GLYC	C erin (Unii: PDC	6A3C0OX)						
CELL	ULOSE, MICRO	CRYSTALLINE	(UNII: OP1R32D61U)					
PROF	PYLENE GLYCO	L (UNII: 6DC9Q	167V3)					
	E R (UNII: 059QF0							
	UM BENZOATE		EU)					
	ALOSE (UNII: 96							
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)								
XANTHAN GUM (UNII: TTV12P4NEE)								
Proc	luct Characte	ristics						
Color		GREEN (opac	ue, viscous)		Score			
Shap	e				Size	Size		
Flavo	r	MINT			Imprint Code	Imprint Code		
Conta	Contains							
Pack	kaging							
#	Item Code		Package Description	Μ	arketing Start Date	Marketing End	d Date	
1 ND	C:37205-584-26	120 mL in 1 BOT	TLE; Type 0: Not a Combination	Product 03	/28/2012	06/30/2020		
Marketing Information								
Mar	keting Category	y Applicatio	n Number or Monograph Cit	ation M	arketing Start Date	Marketing End Date		
ANDA	1	ANDA091292		03/2	28/2012	06/30/2020		

Labeler - Cardinal Health (097537435)

Revised: 12/2018

Cardinal Health