

MEDIQUE DIAMODE- loperamide hydrochloride tablet
Proficient Rx LP

MEDIQUE[®] Diamode

Drug Facts

Active ingredient (in each caplet)

Loperamide HCl USP, 2mg

Purpose

Anti-diarrheal

Uses

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

- a 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 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**

Adults and children (12 years and over): Take 2 caplets after the first loose stool; 1 caplet after each subsequent loose stool; but no more than 4 caplets in 24 hours.

Children under 12 years: Do not give to children under 12 years of age.

Other information

- store between 68° - 77°F (20° - 25°C)
- tamper-evident sealed packets
- do not use any opened or torn packet
- see back of packet for lot number and expiration date

Inactive ingredients

anhydrous lactose, croscarmellose sodium, crospovidone, D&C Yellow # 10, FD&C Blue # 1, hydrogenated vegetable oil, magnesium stearate, powdered cellulose, pregelatinized starch

Questions or comments?

1-800-634-7680

Medique Diamode Label

Diamode

Controls the Symptoms of Diarrhea

Pull to Open

See New Warnings Information

Antidiarrheal • Loperamide HCl 2 mg

Relabeled by:

Proficient Rx LP

Thousand Oaks, CA 91320



Scan Here



NDC 71205-316-06

Relabeled By: Proficient Rx LP
Thousand Oaks, CA 91320



Diamode 2mg

#06 Tablets

Each tablet contains: Loperamide HCl USP, 2 mg
Anti-diarrheal

Green, oval shaped caplet, scored in 2 pieces, with imprint code "123"

Product ID: SD031606

Mfr. For: Medique Products Fort Myers, FL 33967

Store between 68°-77°F (20°-25°C)

Keep medication out of the reach of children

Diamode 2mg
#06 Tablets
Lot #:00000 SN# MASTER
NDC 71205-316-06 Exp:00/00/00

Diamode 2mg
#06 Tablets
Lot #:00000 SN# MASTER
NDC 71205-316-06 Exp:00/00/00

Diamode 2mg
#06 Tablets
Lot #:00000 SN# MASTER
NDC 71205-316-06 Exp:00/00/00



GTIN: 00371205316068
SN# MASTER
Exp. 00/00/00
Lot #:00000

MEDIQUE DIAMODE

loperamide hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71205-316(NDC:47682-200)
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)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
CROSPVIDONE (120 .MU.M) (UNII: 68401960MK)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
STARCH, CORN (UNII: O8232NY3SJ)	
HYDROGENATED COTTONSEED OIL (UNII: Z82Y2C65EA)	

Product Characteristics

Color	green (Green)	Score	2 pieces
Shape	OVAL (Caplet)	Size	10mm
Flavor		Imprint Code	123
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71205-316-01	1 in 1 PACKET; Type 0: Not a Combination Product	09/01/2019	
2	NDC:71205-316-06	6 in 1 PACKET; Type 0: Not a Combination Product	10/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA074091	12/30/2008	

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-316) , RELABEL(71205-316)

Revised: 3/2024

Proficient Rx LP