

MECLIZINE HYDROCHLORIDE- meclizine hydrochloride tablet, chewable
PD-Rx Pharmaceuticals, Inc.

Drug Facts

Active ingredient (in each chewable tablet)

Meclizine HCl 25 mg

Purpose

Antiemetic

Uses

prevents and treats nausea, vomiting or dizziness due to motion sickness

Do not use in

children under 12 years of age unless directed by a doctor

Ask a doctor before use if you have

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if

you are taking sedatives or tranquilizers

When using this product

- Do not exceed recommended dosage
- may cause drowsiness
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- use caution when driving a motor vehicle or operating machinery

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

- Dosage should be taken one hour before travel starts

adults and children 12 years of age and over
 children under 12 years of age

chew 1 to 2 tablets once daily, or as directed by a doctor
 do not give this product to children under 12 years of age unless directed by a doctor

Other information

- Store at room temperature in a dry place at 15°-30°C (59°-86°F)
- keep lid tightly closed

Croscarmellose Sodium, Crospovidone, FD&C Red #40 Lake, French Vanilla Flavor, Lactose, Magnesium Stearate, Raspberry Flavor, Silica, Sodium Saccharin, Stearic Acid

Questions or comments?

1-800-645-2158

TAMPER EVIDENT: DO NOT USE IF SEAL IS BROKEN OR MISSING FROM BOTTLE.

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Adults and children 12 years of age and over	chew 1 or 2 tablets once daily, or as directed by a doctor
children under 12 years of age	do not give this product to children under 12 years of age unless directed by a doctor

Other information
 • each tablet contains 0.09 mg of Magnesium and 0.82 mg of Sodium
 • store at room temperature in a dry place • keep lid tightly closed

Inactive ingredients Croscarmellose Sodium, Crospovidone, FD&C Red #40 Lake, French Vanilla Flavor, Lactose, Magnesium Stearate, Raspberry Flavor, Silica, Sodium Saccharin, Stearic Acid.

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. (800) 222-1222

Questions or comments?
 1-800-645-2158

Marketed and Packaged by:
 PD-Rx Pharmaceuticals, Inc
 Oklahoma City, OK 73127
 1-405-942-3040 v.8.19.0

GTIN: 00372789250205
 SNO: E22A50000033
 EXP: 01/2024
 LOT: E22A50

20 Tablets
 TAMPER EVIDENT: DO NOT USE IF SEAL IS BROKEN OR MISSING FROM BOTTLE.

MECLIZINE HYDROCHLORIDE

meclizine hydrochloride tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72789-250(NDC:0536-1299)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MECLIZINE HYDROCHLORIDE (UNII: HDP7W44CIO) (MECLIZINE - UNII:3L5TQ84570)	MECLIZINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
CROSPVIDONE (UNII: 2S7830E561)	
VANILLA (UNII: Q74T35078H)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
RASPBERRY (UNII: 4N14V5R27W)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	

Product Characteristics

Color	pink (Rosy)	Score	2 pieces
Shape	ROUND	Size	9mm
Flavor	VANILLA, RASPBERRY	Imprint Code	5172
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72789-250-12	12 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2022	
2	NDC:72789-250-20	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2022	
3	NDC:72789-250-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M009	10/30/2020	

Labeler - PD-Rx Pharmaceuticals, Inc. (156893695)**Registrant** - PD-Rx Pharmaceuticals, Inc. (156893695)

Establishment

Name	Address	ID/FEI	Business Operations
PD-Rx Pharmaceuticals, Inc.		156893695	repack(72789-250)

Revised: 10/2025

PD-Rx Pharmaceuticals, Inc.