

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

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***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.

Meclizine 25 mg

Antiemetic

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Active ingredient (in each chewable tablet)	Purpose
Meclizine HCl 25mg	Antiemetic

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

**Warnings**  
 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

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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

**Directions**  
 • dosage should be taken one hour before travel starts

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

<b>MECLIZINE HYDROCHLORIDE</b>			
meclizine hydrochloride tablet, chewable			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:72789-250(NDC:0536-1299)
<b>Route of Administration</b>	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: 4/2024

PD-Rx Pharmaceuticals, Inc.