

MECLIZINE HCL- meclizine hydrochloride chewable tablet, chewable Preferred Pharmaceuticals Inc.

Meclizine Hydrochloride Chewable Tablets 25 mg

Drug Facts

Active ingredient (in each chewable tablet)

Meclizine HCl, USP 25 mg

Purpose

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

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers.

When using this product

- 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 the Poison Control Center immediately.

Directions

- Dosage should be taken one hour before travel starts
- **Adults and children 12 years and older:** Chew 1-2 tablets once daily, or as directed by a doctor
- **Children under 12 years:** do not give this product to children under 12 years of age unless directed by a doctor

Other Information

- Phenylketonurics: Contains Phenylalanine 0.0025 mg per tablet
- Store at room temperature in a dry place
- Keep lid tightly closed

Inactive ingredients

aspartame, colloidal silicon dioxide, croscarmellose sodium, dextrose, lake of FD & C Red 40, magnesium stearate, maltodextrin, microcrystalline cellulose, raspberry flavor, sodium sulfate anhydrous, sucrose, tribasic calcium phosphate

Questions or comments?

Call **1-844-474-7464** Monday to Friday 8 AM - 5 PM ET

**TAMPER EVIDENT: DO NOT USE IF FOIL SEAL UNDER CAP, PRINTED WITH "SEALED for YOUR PROTECTION" IS BROKEN OR MISSING.
Rising Pharma Holdings, Inc. is not affiliated with the owner of the registered trademark Bonine®**

**Manufactured by:
Unique Pharmaceutical Laboratories
(A Div. of J.B. Chemicals & Pharmaceuticals Ltd.),
Mumbai 400 030, India**

**Distributed by:
Rising Pharma Holdings, Inc.
East Brunswick, NJ 08816**

**Mfg. Lic. No.: G/1430
Feb 2022**

Repackaged By: Preferred Pharmaceutical Inc.

PRINCIPAL DISPLAY PANEL - 25 mg Chewable Tablet Label

Meclizine Hydrochloride Chewable Tablets

25 mg

68788-8529

Meclizine HCL 25mg Chewable Tablets

Generic for Bonine
Each tablet contains: Meclizine HCL, USP.....2
5mg

Pkg Size: Exp Date:
Lot#:
Batch#:
Ins:

Mfg: Unique Pharmaceutical
Laboratories

Prod#:
Warning

Store at room temperature in a dry place. 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, trouble urinating due to an enlarged prostate gland. Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers. If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. Tablet is round, pink, scored and imprinted with M



CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed

Meclizine HCL 25mg
Chewable Tablets
Qty: Ins:
Lot#: Bat#:

Prod# (NDC):

Meclizine HCL 25mg
Chewable Tablets
Qty: Ins:
Lot#: Bat#:
Prod# (NDC):

Meclizine HCL 25mg
Chewable Tablets
Qty:
Insurance NDC:
Lot#: Bat#:

Meclizine HCL 25mg
Chewable Tablets
Qty: Ins:
Lot#: Bat#:
Prod# (NDC):



Directions English
May cause drowsiness.
Take ___ tablet(s)
every ___ hours.



Instrucciones Espanol:
Puede causar
sornolencia.
Toma ___ tableta(s)
cada ___ horas.

Log
Chart
Billing
Patient

MECLIZINE HCL

meclizine hydrochloride chewable tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8529(NDC:16571-824)
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
ASPARTAME (UNII: Z0H242BBR1)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
RASPBERRY (UNII: 4N14V5R27W)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM SULFATE ANHYDROUS (UNII: 36KCS0R750)	
SUCROSE (UNII: C151H8M554)	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)	

Product Characteristics

Color	pink (Pink to light pink)	Score	2 pieces
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Shape	ROUND	Size	8mm	
Flavor	RASPBERRY	Imprint Code	M	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8529-1	10 in 1 BOTTLE; Type 0: Not a Combination Product	10/06/2023	
2	NDC:68788-8529-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/06/2023	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	336	10/06/2023		

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment			
Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-8529)

Revised: 7/2024

Preferred Pharmaceuticals Inc.