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	Pharmaceuticals, Inc.	CAUTION: Federal law PROHIBITS transfer of this drug to any person other thean the patient for whom it was prescribed	Meclizine HCL Chewable Tablets Qty: Ins: Lot#: Bat#:	25mg	Log
Tablets Generic for Bonine Each tablet contains: Meclizine HCL, USP2 Smg Pkg Size: Exp Date:	ss.	(S)	Prod# (NDC): Meclizine HCL Chewable Tablets Qty: Ins: Lot#: Bat#: Prod# (NDC):	25mg	Chart
Lot#: Batch#: Ins: Mfg: Unique Pharmaceutical Laboratories Prod#:	Directions English Se drowsiness tablet(s) hours.	ucciones Esp Jsar cia. _ tableta(_ horas.	Meclizine HCL Chewable Tablets Qty: Insurance NDC: Lot#: Bat#:	25mg	Billing
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 urmaining due to an enlarged you are tiking sedatives or tranquilitzers. 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	Dir May cause Takeevery	Instr Puede cau somnolen Toma cada	Meclizine HCL Chewable Tablets Qty: Ins: Lot#: Bat#: Prod# (NDC):	25mg	Patient

MECLIZI	NE HCL
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meclizine hydrochloride chewable tablet, chewable

Product Informat	ion					
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8529	(NDC:16	5571-824)
Route of Administra	tion	ORAL				
Active Ingredient/	Active	Moiety				
	Ingre	dient Name		Basis of Streng	gth	Strength
MECLIZINE HYDROCHLORIDE (UNII: HDP7W44CIO) (MECLIZINE - MECLIZINE UNII:3L5TQ84570) MECLIZINE HYDROCHLORIDE						25 mg
Inactive Ingredier	nts					
		Ingredient Na	me		Strength	
ASPARTAME (UNII: Z0H242BBR1)						
CROSCARMELLOSE SO	DIUM (UN	II: M28OL1HH48)				
DEXTROSE, UNSPECIFI	ED FORM	(UNII: IY9XDZ35W2)				
FD&C RED NO. 40 (UNII	: WZ B912	7XOA)				
MAGNESIUM STEARATE	(UNII: 70	097M6I30)				
MALTODEXTRIN (UNII: 7	CVR7L4A2	D)				
MICROCRYSTALLINE CE	ELLULOSE	(UNII: OP1R32D61U))			
RASPBERRY (UNII: 4N14)	/5R27W)					
SILICON DIOXIDE (UNII:	ETJ7Z6XB	U4)				
SODIUM SULFATE ANH	YDROUS	(UNII: 36KCS0R750)				
SUCROSE (UNII: C151H8	M554)					
TRIBASIC CALCIUM PHO	OSPHATE	(UNII: 91D9GV0Z28)				
Product Characte	ristics					
Color pi	nk (Pink to		Score		2 piec	

Sł	hape	ROUND Size		8mm		
FI	avor	RASPBERRY	Imprint Code	М		
Contains						
D	ackaging					
Packaging						
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date		
1		10 in 1 BOTTLE; Type 0: Not a Combination Product	10/06/2023			
2		30 in 1 BOTTLE; Type 0: Not a Combination Product	10/06/2023			
Marketing Information						
	Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date		
01	FC 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: 10/2023

Preferred Pharmaceuticals Inc.