MECLIZINE HCL 25 MG- meclizine hydrochloride tablet, chewable
Gemini Pharmaceuticals, Inc. dba Plus Pharma

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Meclizine HCl 25 mg Chewable Tablets

DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

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 of age 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 0.28 mg per tablet
- store at room temperature in a dry place

Inactive ingredients Aspartame, croscarmellose sodium, dextrose, FD&C red #40 lake, magnesium stearate, maltodextrin, microcrystalline cellulose, natural and artificial flavors, silicon dioxide, sodium

sulfate, sugar, tricalcium phosphate.

Questions? If you have any questions or comments or to report an adverse event, please contact **(800) 795-9775**.

Distributed by: Plus Pharma, Commack, NY 11725

*Plus Pharma is not affiliated with the owner of the registered trademark Bonine®.

NDC 51645-994-01

Plus Pharma™

MECLIZINE HCl 25 mg

ANTIEMETIC

Treats and Prevents Motion Sickness

*Compare to the Active Ingredient in Bonine®

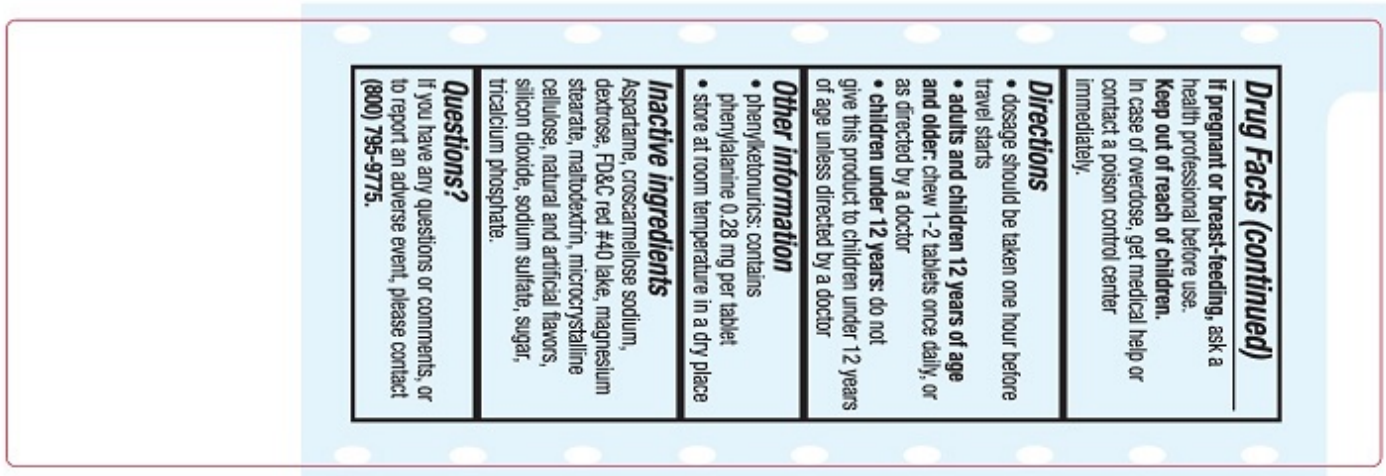
Prevents nausea, dizziness and vomiting

Raspberry

Natural & Artificial Flavor

Contains no ingredient from gluten-containing grain (wheat, barley, or rye).

100 CHEWABLE TABLETS • 25 mg each



THIS PACKAGE FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN

Plus Pharma™

NDC 51645-994-10

MECLIZINE HCl 25 mg

ANTIEMETIC

Treats and Prevents Motion Sickness

Prevents nausea, dizziness and vomiting

*Compare to the Active Ingredient in Bonine®

Raspberry

Natural & Artificial Flavor

Contains no ingredient from a gluten-containing grain (wheat, barley, or rye).

1000 CHEWABLE TABLETS • 25 mg each

MECLIZINE HCL 25 MG

meclizine hydrochloride tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51645-994
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)	
SUCROSE (UNII: C151H8M554)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DEXTROSE (UNII: IY9XDZ35W2)	

FD&C RED NO. 40 (UNII: WZB9127XOA)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	

Product Characteristics

Color	pink (Uncoated)	**Score**	2 pieces
Shape	ROUND (Biconvex)	**Size**	8mm
Flavor	RASPBERRY	**Imprint Code**	21G
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51645-994-01	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/15/2015	
2	NDC:51645-994-10	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/15/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part336	10/15/2015	

Labeler - Gemini Pharmaceuticals, Inc. dba Plus Pharma (055942270)

Establishment

Name	Address	ID/FEI	Business Operations
Gemini Pharmaceuticals, Inc.		055942270	manufacture(51645-994)

Revised: 11/2020 Gemini Pharmaceuticals, Inc. dba Plus Pharma