

MECLIZINE- meclizine hcl 25mg tablet, chewable
NuCare Pharmaceuticals, Inc.

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.

Drug Facts

Active ingredient (in each tablet)

Meclizine HCl 25mg

Purpose

Antiemetic

Uses

prevents and treats nausea, vomiting or dizziness associated with motion sickness

Warnings

Do not use for children under 12 years of age unless directed by a doctor.

Do not take unless directed by a doctor if you have

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

Do not take if you are taking sedatives or tranquilizers, without first consulting your doctor.

When using this product

- do not exceed recommended dosage
- drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- be careful 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.

Directions

- doage should be taken 1 hour before travel starts

Adults and children 12 years and over	take 1 or 2 tablets once daily or as directed by doctor
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Other information

- **Tamper Evident:** do not use if safety seal under cap is broken or missing
- store at room temperature 20°-25°C (68°-77°F)

Inactive ingredients

Croscarmellose sodium, dextrose, FD&C Red#40, flavor, magnesium stearate, microcrystalline cellulose, silicon dioxide, sodium saccharine, stearic acid

Questions?

Adverse drug event call (800) 687-0176 (M - F, 8AM - 4PM EST).

NuCare Pharmaceuticals, Inc.



Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

Caution Instructions:

Chew _____ every _____ hours _____ times a day.

GTIN 00388071498660
Serial# 00000000002
Exp. Date 00-00
LOT#: 000000

Rev 01/01/19

NDC: 68071-4986-6
Meclizine HCl 25mg
#6 Chewtabs

Round Pink Scored Tablet Debossed: "PH 051" on the un-scored side Each tablet contains: Meclizine HCl 25mgAntiemetic

Warnings: Do not use for children under 12 years of age unless directed by a doctor. Do not take unless directed by a doctor if you have glaucoma, trouble urinating due to an enlarged prostate gland, a breathing problem such as emphysema or chronic bronchitis.

Do not take if you are taking sedatives or tranquilizers, without first consulting your doctor. When using this product do not exceed recommended dosage, drowsiness may occur, alcohol, sedatives, and tranquilizers may increase drowsiness, avoid alcohol drinks, be careful when driving a motor vehicle or operating machinery. If pregnant or breast-feeding, ask a health professional before use.

Product #: P1781006ER



6 8 0 7 1 4 9 8 6 6 0

Meclizine HCl 25mg
#6 Chewtabs Serial# 0000000002
Lot: 000000 NDC: 68071-4986-06
Exp.: 00-00 MFR NDC: 66424-0387-01

WARNING: KEEP OUT OF REACH OF CHILDREN STORE AT CONTROLLED TEMPERATURE 68-77°F.

MECLIZINE

meclizine hcl 25mg tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-4986(NDC:66424-387)
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)				
DEXTROSE (UNII: IY9XDZ35W2)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
Product Characteristics				
Color	pink (LIGHT PINK COLOR)		Score	2 pieces
Shape	ROUND (ROUND TABLET)		Size	8mm
Flavor			Imprint Code	PH051
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-4986-6	6 in 1 BOTTLE; Type 0: Not a Combination Product	07/24/2019	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part336	02/01/2018		

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	repack(68071-4986)

Revised: 2/2021

NuCare Pharmaceuticals,Inc.