

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

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
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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.

NDC: 68071-4584-2

Meclizine HCl 25mg #20 Chewtabs

Each chewtab contains:
Meclizine HCl 25mg.....Antiemetic

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. Round Pink Scored Chewtab Debossed: "PH 051" on the un-scored side

Product #: P1763020

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 68-77°F.

Meclizine HCl 25mg
Lot: 000000 NDC: 68071-4584-02
MFR NDC: 66424-0387-01 Exp.: 00-00

Meclizine HCl 25mg
Lot: 000000 NDC: 68071-4584-02
MFR NDC: 66424-0387-01 Exp.: 00-00

GTIN 00368071458428
Serial# 00000000002
Exp. Date 00-00
LOT#: 000000

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

Distributed by: SDA Laboratories, Inc., Greenwich, CT 06830
Packaged By: NuCare Pharmaceuticals, Inc. Orange, CA 92867

Patent Instructions: 88071458402*20*000000*000000

Chew _____ times a day, every _____ hours

Rev 01/01/19

MECLIZINE

meclizine hcl 25mg tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-4584(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
CROSCARMELOSE 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:68071-4584-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	04/04/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	02/01/2018	

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NUCARE PHARMACEUTICALS INC		010632300	repack(68071-4584)

Revised: 6/2024

NuCare Pharmaceuticals, Inc.