

MECLIZINE HYDROCHLORIDE- meclizine hydrochloride tablet, chewable
NuCare Pharmaceuticals, Inc.

5172C- Rubgy

Drug Facts

Active ingredient (in each chewable tablet)

Meclizine HCl 25 mg

Purpose

Antiemetic

Uses

prevents and treats nausea, vomiting or dizziness due to motion sickness

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

- ☐ Do not exceed recommended dosage
- ☐ 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 a Poison Control Center right away (1-800-222-1222).

Directions

□ Dosage should be taken one hour before travel starts

adults and children 12 years of age and over	chew 1 to 2 tablets once daily, or as directed by a doctor
children under 12 years of age	do not give this product to children under 12 years of age unless directed by a doctor

Other information

- Each tablet contains 0.09 mg of Magnesium and 0.82 mg of Sodium
- store at room temperature in a dry place
- keep lid tightly closed

Croscarmellose Sodium, Crospovidone, FD&C Red #40 Lake, French Vanilla Flavor, Lactose, Magnesium Stearate, Raspberry Flavor, Silica, Sodium Saccharin, Stearic Acid

Questions or comments?


1-800-645-2158

THIS PACKAGE FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN

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

*This product is not manufactured or distributed by Prestige Brands, Inc owner of the registered trademark Bonine®.

Distributed by:
RUGBY® LABORATORIES
17177 N Laurel Park Dr., Suite 233
Livonia, MI 48152
www.rugbylaboratories.com

 NuCare Pharmaceuticals, Inc.

NDC: 68071-2574-2

Meclizine HCl 25mg


#20 Chewtabs

Each chewable tablet contains: Meclizine HCl 25mg..... Antiemetic

Warnings: Do not use for 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, drowsiness may occur, 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. In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222) Round pink score chewtab imprint 5172 on the score side

Meclizine HCl 25mg
Lot: 00000 NDC: 68071-2574-02
MFR NDC: 0536-1299-01 Exp.: 00-00
Serial# 0000000002

Meclizine HCl 25mg
Lot: 00000 NDC: 68071-2574-02
MFR NDC: 0536-1299-01 Exp.: 00-00
Serial# 0000000002



GTIN 00368071257427
Serial# 0000000002
Exp. Date 00-00
LOT#: 00000

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

Product #: P1763020

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 59-86°F.

Distributed by:
Rugby Laboratories Indianapolis, IN 46268
Packaged By:
NuCare Pharmaceuticals, Inc.
Orange, CA 92867

Chew _____ times a day, every _____ hours

Rev 01/01/19

MECLIZINE HYDROCHLORIDE

meclizine hydrochloride tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-2574(NDC:0536-1299)
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
CROSPVIDONE (UNII: 2S7830E561)	
VANILLA (UNII: Q74T35078H)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
RASPBERRY (UNII: 4N14V5R27W)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	pink (Rosy)	Score	2 pieces
Shape	ROUND	Size	9mm
Flavor	VANILLA, RASPBERRY	Imprint Code	5172
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-2574-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2021	
2	NDC:68071-2574-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2021	
3	NDC:68071-2574-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2021	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
-----------	---------------------------------	-----------------	---------------

Category	Citation	Date	Date
OTC Monograph Drug	M009	10/30/2020	

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	repack(68071-2574)

Revised: 1/2026

NuCare Pharmaceuticals,Inc.