

## **MOTION SICKNESS RELIEF- meclizine hcl tablet, chewable**

**Walgreen Company**

*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.*

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**Walgreens 44-404**

### ***Active ingredient (in each tablet)***

Meclizine HCl 25 mg

### ***Purpose***

Antiemetic

### ***Uses***

for prevention and treatment of nausea, vomiting, or dizziness associated with motion sickness

### ***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
- difficulty in urination due to enlargement of the prostate gland
- a breathing problem such as emphysema or chronic bronchitis

#### **Ask a doctor or pharmacist before use if you are**

taking sedatives or tranquilizers

#### **When using this product**

- alcohol, sedatives, and tranquilizers may increase drowsiness
- drowsiness may occur
- 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 (1-800-222-1222) right away.

### ***Directions***

- dosage should be taken one hour before travel starts
- chew or crush tablets completely before swallowing; do not swallow tablets whole
- adults and children 12 years and over: take 1 to 2 chewable tablets once daily or as directed by a

doctor

**Other information**

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- protect from heat and humidity
- see end flap for expiration date and lot number

**Inactive ingredients**

corn starch, FD&C red #40 aluminum lake, flavor, lactose anhydrous, magnesium stearate, saccharin sodium, silicon dioxide

**Questions or comments?**

**1-800-426-9391**

**Principal Display Panel**

NDC 0363-0404-21

Compare to Bonine<sup>®</sup> active ingredient<sup>††</sup>

**Walgreens**

**Motion Sickness Relief**

MECLIZINE HCl TABLETS, 25 mg / CHEWABLE TABLETS / ANTIEMETIC

CHEWABLE LESS DROWSY FORMULA

- Prevents motion sickness
- Chewable, once-a-day protection

Chew or crush tablets completely before swallowing.  
Do not swallow tablets whole.

**16 CHEWABLE  
TABLETS**

ACTUAL SIZE

**RASPBERRY  
FLAVOR**

**TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING**

Walgreens Pharmacist Recommended  
Walgreens Pharmacist Survey

<sup>††</sup>This product is not manufactured or distributed by  
Insight Pharmaceuticals LLC, owner of the registered  
trademark Bonine<sup>®</sup>.

50844 REV1218B40421

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200 WILMOT RD., DEERFIELD, IL 60015

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Walgreens



OR61217-F  
REV0219

Chewable, once-a-day protection  
that prevents motion sickness.

Walgreens

Compare to Bonine®  
active ingredient\*\*

NDC 0363-0404-21

E1042

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RASPBERRY FLAVOR

B-2201-404-21-H  
REV12188-40421

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ITEM 501196 W00000-0000-0



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Do not print Lot and Exp ONLY



Walgreens 44-404

## MOTION SICKNESS RELIEF

meclizine hcl tablet, chewable

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0404
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
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	

### Product Characteristics

Color	PINK	Score	no score
Shape	ROUND	Size	9mm
Flavor	RASPBERRY	Imprint Code	44;404
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0404-21	2 in 1 CARTON	05/29/2002	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part336	05/29/2002	

**Labeler** - Walgreen Company (008965063)

**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867894	MANUFACTURE(0363-0404)

Revised: 3/2020

Walgreen Company