

**MECLIZINE HCL- meclizine hcl 25mg chewable tablets tablet, chewable**  
**Advance Pharmaceutical 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.*

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**Motion Sickness**

**Active ingredient**

Meclizine Hydrochloride 25 mg

**Purpose**

Antiemetic

**Uses**

For the prevention and treatment of nausea, vomiting or dizziness associated with motion sickness. For other uses consult your doctor

**Warnings**

**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**

- you may get drowsy
- avoid alcoholic drinks
- alcohol, sedatives and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

**Directions**

**take dose one hour before travel starts**  
**tablets can be chewed or swallowed whole with water**

**adults & children 12 years and over:**

- take 1-2 tablets once daily

**children under 12 years:**

- ask a doctor

**Keep out of reach of children.**

In case of overdose, get medical help or contact a Poison Control Center right away.

**OTHER INFORMATION**

Phenylketonurics:  
 each tablet contains:  
 phenylalanine 0.28 mg  
 store at room temperature 15 - 30 °C

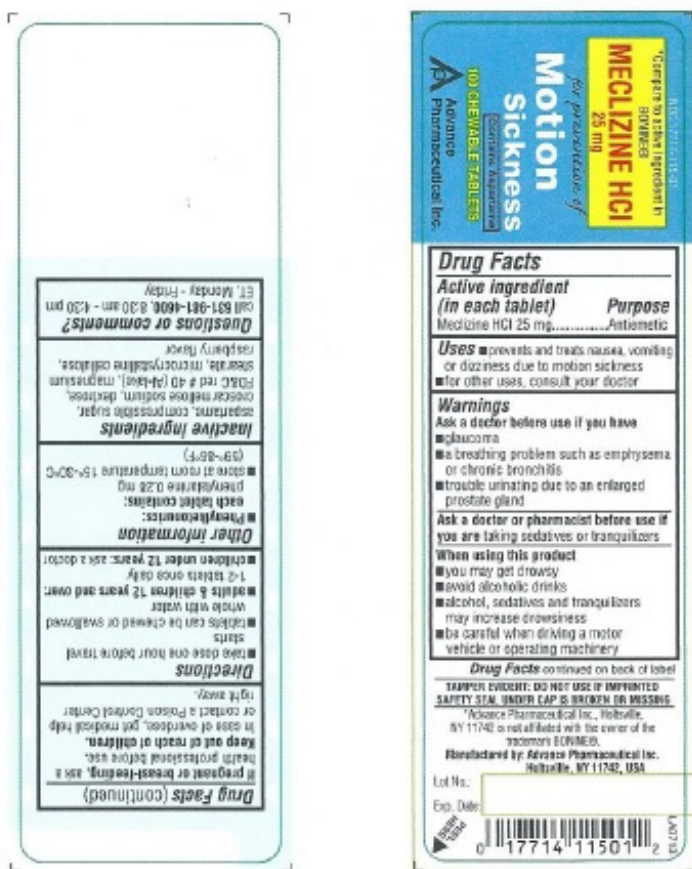
### Questions or Comments

call 631-981-4600, 8.30 am-4.30 pm ET, Monday - Friday  
 TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

### Inactive ingredients

Aspartame, compressible sugar, croscarmellose sodium, dextrose, FD&C red 40 (al-lake), magnesium stearate, microcrystalline cellulose, raspberry flavor

### Package/Label Principal Display Panel



## MECLIZINE HCL

meclizine hcl 25mg chewable tablets tablet, chewable

### Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:17714-115

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)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DEXTROSE (UNII: IY9XDZ35W2)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
RASPBERRY (UNII: 4N14V5R27W)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SUCROSE (UNII: C151H8M554)	

### Product Characteristics

Color	red	Score	2 pieces
Shape	ROUND	Size	8mm
Flavor	RASPBERRY	Imprint Code	AP;115
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17714-115-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2013	
2	NDC:17714-115-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2013	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part336	07/01/2013	

**Labeler** - Advance Pharmaceutical Inc. (078301063)

**Registrant** - Advance Pharmaceutical Inc. (078301063)

### Establishment

Name	Address	ID/FEI	Business Operations
Advance Pharmaceutical Inc.		078301063	manufacture(17714-115)

