

JET-AVERT MOTION SICKNESS AID- meclizine hydrochloride tablet
Bell 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.

Jet-Avert Motion Sickness Aid

Drug Facts

Active ingredient (in each tablet)

Meclizine HCL 25mg

Purpose

Antiemetic

Uses

- prevent 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

Ask a doctor before use if you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- difficulty in urination due to enlargement prostate gland

Ask a doctor or pharmacist before use if you are taking

- Sedatives • tranquilizers

When using this product

• drowsiness may occur • alcohol, sedatives, and tranquilizers may increase drowsiness • avoid alcoholic drinks • be careful when driving 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

- take first dose one hour before starting activity

adults and children 12 years and over
children under 12 years

take 1 to 2 tablets once daily, or as directed by a doctor
do not use unless directed by doctor

Other information

- store at room temperature between 59°-86°F (15°-30°C)
- protect from excessive moisture
- don not use if tamper-evident seal is broken or missing

Inactive ingredients

calcium stearate, FD&C red#40 aluminum lake, lactose, microcrystalline cellulose, silicon dioxide, sodium starch glycolate

Questions or comments?

1-800-328-5890 Weekdays 8:30-5:00 CST

Package Labeling:

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REV 7/16

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Manufactured in the USA
For Bell Pharmaceuticals, Inc.
200 W. Beaver St.
Belle Plaine, MN 56011
Visit www.jet-avert.com

JET-AVERT MOTION SICKNESS AID

meclizine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:15579-837
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MECLIZINE HYDRO CHLORIDE (UNII: HDP7W44CIO) (MECLIZINE - UNII:3L5TQ84570)	MECLIZINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
CALCIUM STEARATE (UNII: 776XM7047L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
LACTOSE (UNII: J2B2A4N98G)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	4mm
Flavor		Imprint Code	25
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15579-837-23	3 in 1 BOX	10/07/2017	
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	10/07/2017	

Labeler - Bell Pharmaceuticals, Inc. (140653770)

Revised: 10/2017

Bell Pharmaceuticals, Inc.