MOTION SICKNESS RELIEF- meclizine hcl tablet, film coated Shield Pharmaceuticals Corp

Motion Sickness Relief

Drug Facts

Active ingredient (in each liquid-filled capsule)

Meclizine HCl 25 mg

Purpose

Antiemetic

∏Uses

for the 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

- drowsiness may occur n avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- 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

· do not take more than directed

- take first dose ½ to 1 hour before starting activity
- adults and children 12 years and over: 1 to 2 tablets once daily, or as directed by a doctor

Other information

- TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR SAFETY SEAL UNDER CAP IS BROKEN OR MISSING
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F) protect from heat and humidity

Inactive ingredients

colloidal silicon dioxide, D&C yellow #10 aluminum lake, lactose monohydrate, magnesium stearate, pregelatinized starch.

Questions or comments?

1-800-373-6981 (toll-free)

Shield Pharmaceuticals Corp. Carle Place, NY 11514

PRINCIPAL DISPLAY PANEL

Compare to Dramamine® Less Drowsy active ingredient

LESS DROWSY FORMULA

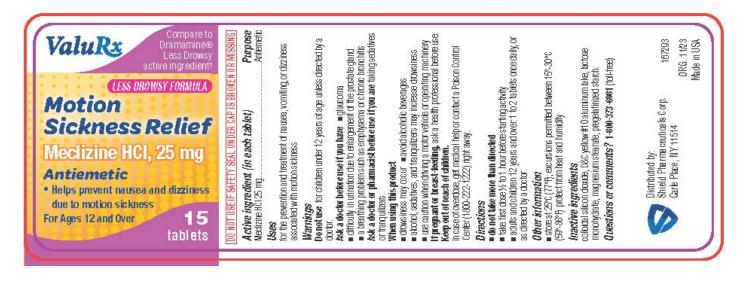
Motion Sickness Relief

Meclizine HCl, 25 mg

Antiemetic

Helps prevent nausea and dizziness due to motion sickness

For Ages 12 and Over





MOTION SICKNESS RELIEF

meclizine hcl tablet, film coated

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:83059-0007 Route of Administration ORAL

	Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength		
	MECLIZ INE IYDROCHLORIDE	25 mg		

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics				
Color	yellow	Score	no score	
Shape	ROUND	Size	8mm	
Flavor		Imprint Code	44;403	
Contains				

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:83059- 0007-1	1 in 1 PACKAGE	12/06/2023				
1		15 in 1 BOTTLE; Type 0: Not a Combination Product					

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M002	12/06/2023		

Labeler - Shield Pharmaceuticals Corp (118724924)

Revised: 12/2023 Shield Pharmaceuticals Corp