DRAMAMINE LESS DROWSY- meclizine hydrochloride tablet
Medtech Products 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.

Dramamine Less Drowsy

Drug Facts

Active ingredient

(in each tablet)

Meclizine HCl 25 mg

Purpose

Antiemetic

Use

for prevention and treatment of these symptoms associated with motion sickness:

- nausea
- vomiting
- dizziness

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
- 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
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness

- be careful when driving a motor vehicle or operating machinery

If pregnant or breast-feeding,

ask a doctor 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

- take first dose one 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

- store between 20 - 25°C (68 - 77°F)
- do not use if blister is broken or torn

Inactive ingredients

anhydrous lactose, corn starch, colloidal silicon dioxide, D&C yellow # 10 aluminum lake, magnesium stearate, microcrystalline cellulose

Questions or comments?

call 1-800-382-7219

PRINCIPAL DISPLAY PANEL

MECLIZINE HYDROCHLORIDE
TABLETS/ANTIEMETIC
Dramamine®
motion sickness
LESS DROWSY
Dual Action:
Prevents & Relieves Nausea,
Dizziness and Vomiting
8 TABLETS
(25 mg EACH)

DRAMAMINE LESS DROWSY

meclizine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63029-903
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**
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
STARCH, CORN (UNII: O8232NY3SJ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics			
Color	yellow	**Score**	2 pieces
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63029-903-01	1 in 1 BLISTER PACK	09/01/2011	
1		8 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:63029-903-10	2 in 1 CARTON	09/01/2011	
2		5 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	09/01/2011	

Labeler - Medtech Products Inc. (122715688)

Revised: 8/2022 Medtech Products Inc.