ZENTRIP MOTION SICKNESS- meclizine hydrochloride tablet Sato Pharmaceutical Co., LTD

ZenTrip Motion Sickness (12 Tablets - 25mg each)

Active ingredient (in each tablet)

Meclizine hydrochloride 25mg

Purpose

Antiemetic

Uses

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

Warnings

Do not use in children under 12 years of age unless directed by a physician

Ask a doctor before use if you have

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

Ask a physician or pharmacist before use if you are

• taking sedatives or tranquilizers

When using this product

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

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

- to prevent motion sickness, take it at least one hour before traveling
- adults and children 12 years of age and over: take 1 to 2 tablets (25 to 50mg) once

daily, or as directed by a physician.

Other information

store at 20-30°C (68-86°F)

Inactive ingredients

acesulfame potassium, erythritol, hydroxypropyl cellulose, mannitol, menthol, silicon dioxide colloidal, sodium stearyl fumarate, and yellow ferric oxide.

ZenTrip Motion Sickness - 12Tablets





富山スガキ株式会社

ZENTRIP MOTION SICKNESS

meclizine hydrochloride tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49873-805	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MECLIZINE HYDROCHLORIDE (UNII: HDP7W44CIO) (MECLIZINE - UNII:3L5TQ84570)	MECLIZ INE HYDROCHLORIDE	25 mg	

Inactive Ingredients				
Ingredient Name	Strength			
ACESULFAME POTASSIUM (UNII: 230V73Q5G9)				
MANNITOL (UNII: 3OWL53L36A)				
MENTHOL (UNII: L7T10EIP3A)				
FERRIC OXIDE YELLOW (UNII: EX43802MRT)				
ERYTHRITOL (UNII: RA96B954X6)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)				
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)				

Product Characteristics				
Color	yellow	Score	score with uneven pieces	

Shape	ROUND	Size	13mm
Flavor		Imprint Code	SJ
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:49873-805- 01	2 in 1 BOX	05/01/2021	
	1		6 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 Drug	M009	05/01/2021	

Labeler - Sato Pharmaceutical Co., LTD (690575642)

Establishment				
Name	Address	ID/FEI	Business Operations	
Sato Pharmaceutical Co., LTD		715699133	manufacture(49873-805), pack(49873-805), label(49873-805)	

Revised: 11/2023 Sato Pharmaceutical Co., LTD