DRAMAMINE- meclizine hydrochloride powder
Medtech Products Inc.

Dramamine Less Drowsy Powder Orange

Drug Facts

Active ingredient (in each pack)

Meclizine HCl 50 mg

Purpose

Antiemetic

Use

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

■nausea ■vomiting ■dizziness

Warnings

Do not give to

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 dose ½ to 1 hour before starting activity

■**adults and children 12 years and over:** place 1 packet on tongue once daily, or as directed by a doctor

Other information

■PHENYLKETONURICS: contains phenylalanine 1.68 mg per packet.

■store at room temperature 20°–25°C (68°-77°F)

Inactive ingredients

anhydrous citric acid, ascorbic acid, aspartame, butylated hydroxytoluene, corn starch, guar gum, maltodextrin, mannitol, microcrystalline cellulose, natural orange flavor, natural pink lemonade flavor, silicon dioxide, sorbitol.

Questions?

1-800-382-7219

Dramamine.com

PRINCIPAL DISPLAY PANEL

Dramamine®

Meclizine Hydrochloride | Antiemetic | 50 mg

Motion Sickness

Less Drowsy Powder

4 Orange Flavored Powder Packs (50 MG EACH)

DRAMAMINE

meclizine hydrochloride powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63029-912
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MECLIZINE DIHYDROCHLORIDE (UNII: HDP7W44CIO) (MECLIZINE - UNII:3L5TQ84570)	MECLIZINE DIHYDROCHLORIDE	50 mg

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
ASPARTAME (UNII: Z0H242BBR1)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
STARCH, CORN (UNII: O8232NY3SJ)	
GUAR GUM (UNII: E89I1637KE)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MANNITOL (UNII: 3OWL53L36A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor	ORANGE	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63029-912-04	4 in 1 BOX; Type 0: Not a Combination Product	02/02/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M009	02/02/2026	

Labeler - Medtech Products Inc. (122715688)

Revised: 12/2025 Medtech Products Inc.