DRAMAMINE- meclizine hydrochloride tablet, chewable
Medtech Products Inc.

Dramamine All Day LD Chewable Tablets Raspberry - 63029-913

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 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

■to prevent motion sickness, take first dose ½ to 1 hour before activity

■to treat motion sickness, take at first signs of symptoms

■**adults and children 12 years and over**: 1 to 2 tablets once daily, or as directed by a doctor

Other information

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

Inactive ingredients

corn starch, croscarmellose sodium, crospovidone, flavors, lactose monohydrate, magnesium stearate, maltodextrin, propylene glycol, saccharin sodium, silicon dioxide, stearic acid, sucrose, tricalcium phosphate.

Questions?

1-800-382-7219

Dramamine.com

PRINCIPAL DISPLAY PANEL

All Day Less Drowsy Chewable Formula

Dramamine®

Meclizine Hydrochloride | Antiemetic | 25 mg /Motion Sickness

12 Raspberry Cream Flavored Tablets (25 mg EACH)

DRAMAMINE

meclizine hydrochloride tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63029-913
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MECLIZINE HYDROCHLORIDE (UNII: 008PUV3986) (MECLIZINE - UNII:3L5TQ84570)	MECLIZINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CROSPOVIDONE (UNII: 2S7830E561)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SUCROSE (UNII: C151H8M554)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	

Product Characteristics

Color	white	**Score**	2 pieces
Shape	ROUND	**Size**	9mm
Flavor	RASPBERRY (Raspberry Cream)	**Imprint Code**	258
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63029-913-12	1 in 1 BOX	02/02/2026	
1		12 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	02/02/2026	

Labeler - Medtech Products Inc. (122715688)

Revised: 12/2025 Medtech Products Inc.