

DRAMAMINE LESS DROWSY- meclizine hydrochloride tablet
Navajo Manufacturing Company Inc.

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 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 first dose 1/2 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

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

Inactive ingredients

anhydrous lactose, corn starch, colloidal silicon dioxide, D&C yellow no. 10 aluminum lake, magnesium stearate, microcrystalline cellulose

Questions?

1-800-382-7219

Dramamine.com

PRINCIPAL DISPLAY PANEL



DRAMAMINE LESS DROWSY

meclizine hydrochloride tablet

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:67751-242(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)	
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:67751-242-02	1 in 1 CARTON	02/18/2026	
1		2 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:67751-242-04	2 in 1 CARTON	02/18/2026	
2		2 in 1 POUCH; 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	02/18/2026	

Labeler - Navajo Manufacturing Company Inc. (091917799)

Registrant - Navajo Manufacturing Company Inc. (136941411)

Establishment

Name	Address	ID/FEI	Business Operations
Navajo Manufacturing Company Inc.		136941411	repack(67751-242)

Revised: 2/2026

Navajo Manufacturing Company Inc.