

LESS DROWSY MOTION SICKNESS RELIEF- meclizine hcl tablet
Wal-Mart Stores Inc

Equate 44-403

Active ingredient (in each tablet)

Meclizine HCl 25 mg

Purpose

Antiemetic

Uses

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

Warnings

Do not use

for children under 12 years of age unless directed by a doctor.

Ask a doctor before use if you have

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

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery

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

- **do not take more than directed**
- take first dose ½ 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

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°C-30°C (59°F-86°F)
- protect from heat and humidity
- see end flap for expiration date and lot number

Inactive ingredients

colloidal silicon dioxide, D&C yellow #10 aluminum lake, lactose monohydrate, magnesium stearate, pregelatinized starch

Questions or comments?

1-888-287-1915

Principal Display Panel

NDC 49035-403-19

equate[™]

**Compare to
Dramamine®
Less Drowsy
active
ingredient***

LESS DROWSY FORMULA
**Motion Sickness
Relief**

Meclizine HCl, 25 mg

Antiemetic

• Helps prevent nausea
and dizziness due to
motion sickness

For Ages 12 and Over

Actual Size

**25
mg**

EACH

8 TABLETS

**TAMPER EVIDENT: DO NOT USE IF
PACKAGE IS OPENED OR IF BLISTER
UNIT IS TORN, BROKEN OR SHOWS
ANY SIGNS OF TAMPERING**

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716
PRODUCT OF INDIA

*This product is not manufactured or distributed by Medtech Products Inc.,
owner of the registered trademark Dramamine® Less Drowsy.
50844 ORG042340319

Satisfaction guaranteed –
Or we'll replace it or give you
your money back. For questions
or comments or to report an
undesired reaction or side effect,
please call **1-888-287-1915**.

No Print/No Varnish. Lot & Expiry



Satisfaction guaranteed - Or we'll replace it or give you your money back. For questions or comments or to report an undesired reaction or side effect, please call 1-888-287-1915.

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716
PRODUCT OF: INDIA
owner of the registered trademark Dramamine® Less Drowsy.
50844 0R6042340319

Questions or comments? 1-888-287-1915

Inactive ingredients colloidal silicon dioxide, D&C yellow #10 aluminum lake, lactose monohydrate, magnesium stearate, pregelatinized starch

Other information
■ do not take more than directed
■ 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
■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
■ protect from heat and humidity
■ see end flap for expiration date and lot number

Drug Facts (continued)
Directions
■ do not take more than directed
■ 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

Drug Facts (continued)
Warnings
Do not use for children under 12 years of age unless directed by a doctor.
Ask a doctor before use if you have
■ difficulty in urination due to enlargement of the prostate gland ■ glaucoma
■ a breathing problem such as emphysema or chronic bronchitis
Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers.
When using this product
■ drowsiness may occur
■ avoid alcoholic beverages
■ alcohol, sedatives, and tranquilizers may increase drowsiness
■ use caution when driving a motor vehicle or operating machinery
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.



TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

B-2203-403A-19-R
0R6042340319

Drug Facts
KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION
Active ingredient (in each tablet) Purpose
Meclizine HCl 25 mg Antiemetic
Uses for the prevention and treatment of nausea, vomiting, or dizziness associated with motion sickness.
Drug Facts (continued)

equate™

LESS DROWSY FORMULA

Motion Sickness Relief

Meclizine HCl, 25 mg

Antiemetic

- Helps prevent nausea and dizziness due to motion sickness

For Ages 12 and Over



NDC 49035-403-19



Actual Size

25
mg
EACH

8
TABLETS

Equate 44-403

LESS DROWSY MOTION SICKNESS RELIEF

meclizine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-403
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	yellow	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	44;403
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-403-19	1 in 1 CARTON	06/24/2002	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:49035-403-40	8 in 1 BLISTER PACK; Type 0: Not a Combination Product	06/24/2002	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M009	06/24/2002	

Labeler - Wal-Mart Stores Inc (051957769)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(49035-403)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(49035-403) , pack(49035-403)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(49035-403)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(49035-403)

Revised: 7/2023

Wal-Mart Stores Inc