

MOTION SICKNESS II- meclizine hcl tablet
GREAT LAKES WHOLESALE, MARKETING, & SALES, INC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Healthcare 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 an 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 (1-800-222-1222) right away.

Directions

- to prevent motion sickness, take the first dose one hour before starting activity
- to prevent or treat motion sickness: 1 to 2 tablets once daily for adults and children 12 years and over, 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°-30°C (59°-86°F)
- protect from heat and humidity
- see end flap for expiration date and lot number

Inactive ingredients

corn starch, D&C yellow #10 aluminum lake, lactose, magnesium stearate, silica gel

Questions or comments?

Call 1-800-426-9391

Principal Display Panel

HEALTHCARE™

NDC 64092-608-08

*Compare to the active ingredient in Dramamine® Less Drowsy Formula

Motion Sickness II

**Meclizine HCl 25 mg Each
Antiemetic**

LESS DROWSY FORMULA

*Helps prevent nausea, vomiting and dizziness
due to motion sickness.*

8 TABLETS

*This product is not manufactured or distributed by Prestige Brands, Inc., owner of the registered trademark Dramamine® Less Drowsy Formula.

50844 REV0715E40319

Distributed by: Great Lakes Wholesale & Marketing L.L.C.
3729 Patterson Ave., S.E., Grand Rapids, MI 49512
www.glwholesale.com

HEALTHCARE GUARANTEE

If you are not completely satisfied with this product, regardless of reason, return your unused portion to Great Lakes Wholesale for a full refund

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

Drug Facts (continued)

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Antiemetic
LESS DROWSY FORMULA
Helps prevent nausea, vomiting and dizziness
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NDC 64092-608-08

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HEALTHCARE

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Drug Facts **KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION**
Active ingredient (in each tablet) Purpose
Meclizine HCl 25 mg Antiemetic

Drug Facts (continued)
Uses for the prevention and treatment of nausea, vomiting, or dizziness associated with motion sickness

meclizine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64092-608
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
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE (UNII: J2B2A4N98G)	

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:64092-608-08	1 in 1 CARTON	06/24/2002	
1		8 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 FINAL	part336	06/24/2002	

Labeler - GREAT LAKES WHOLESAL E, MARKET ING, & SALES, INC. (361925498)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	PACK(64092-608)

Establishment

Name	Address	ID/FEI	Business Operations
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LNK International, Inc.		832867894	MANUFACTURE(64092-608)
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Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	PACK(64092-608)

Revised: 7/2017

GREAT LAKES WHOLESALE, MARKETING, & SALES, INC.