MOTION SICKNESS RELIEF- dimenhydrinate tablet Valu Merchandisers Company

Best Choice 44-198

Active ingredient (in each tablet)

Dimenhydrinate 50 mg

Purpose

Antiemetic

Uses

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

- nausea
- vomiting
- dizziness

Warnings

Do not use

for children under 2 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
- difficulty in urination due to enlargement of the prostate gland

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- marked 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-

Directions

 to prevent motion sickness, the first dose should be taken one-half to one hour before starting activity

adults and children 12 years and over	1 to 2 tablets every 4-6 hours; do not exceed 8 tablets in 24 hours, or as directed by a doctor
children 6 to under 12 years	$\frac{1}{2}$ to 1 tablet every 6-8 hours; do not exceed 3 tablets in 24 hours, or as directed by a doctor
Childran / Fo	$\frac{1}{2}$ tablet every 6-8 hours; do not exceed $1\frac{1}{2}$ tablets in 24 hours, or as directed by a doctor

Other information

- each tablet contains: calcium 35 mg
- see end flap for expiration date and lot number
- 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 moisture

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate dihydrate, magnesium stearate, microcrystalline cellulose, silicon dioxide, stearic acid

Questions or comments?

1-800-426-9391

Principal display panel

Best Choice®

Compare to the active ingredient in Dramamine®
Original Formula*

Motion Sickness Relief

Dimenhydrinate 50 mg Antiemetic

Prevents:

- Nausea
- Vomiting
- Dizziness

FOR CHILDREN & ADULTS

Actual Size

12 Tablets

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

*This product is not manufactured or distributed by Medtech Products Inc., owner of the registered trademark Dramamine® Original Formula. 50844 REV0518A19802

PROUDLY DISTRIBUTED BY: ASSOCIATED WHOLESALE GROCERS, INC. KANSAS CITY, KANSAS 66106





Motion Sickness Relief

DIMENHYDRINATE 50 mg ANTIEMETIC



Motion Sickness Relief

DIMENHYDRINATE 50 mg ANTIEMETIC

Compare to the active ingredient in

Original Formula

Dramamine®

Prevents:

- Nausea
- Vomiting
- Dizziness

FOR CHILDREN & ADULTS

12 TABLETS

for more product information. Scan here for more product information. Call 1-844-292-1112



KANSAS CITY, KANSAS 66106 ASSOCIATED WHOLESALE GROCERS, INC. PROUDLY DISTRIBUTED BY:

50844 REV0518A19802 teademark Dramamine® Original Formula. Meditech Products Inc., owner of the registered If his product is not manufactured or distributed by

B-2115-198-02-RR REV0518A19802

28647 1123 LNK BEST CHOICE

No print/No varnish
Lot & Exp date

Gnestions or comments? 1-800-426-9391

microcrystalline cellulose, silicon dioxide, stearic acid calcium phosphate dihydrate, magnesium stearate, INACTIVE INGVEDIENTS croscarmellose sodium, dibasic

■ store at 25°.C (77°F); excursions permitted between 15°-30°C (59°-86°F) ■ protect from moisture OPENED OR BLISTER IS TORN OR BROKEN ■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS

see end flap for expiration date and lot number ■ each tablet contains: calcium 35 mg

Other information

% tablet every 6-8 hours, do not exceed 1% tablets in 24 hours, or as directed by a doctor	children 2 to under 6 years
% to 1 tablet every 6-8 hours, or as exceed 3 tablets in 24 hours, or as directed by a doctor	children 6 to under 12 years
ur before starting activity 1 to 2 tablets every 4-6 hours, or as exceed 8 tablets in 24 hours, or as directed by a doctor	one-half to one ho adults and children 12 years and over

medical help or contact a Poison Control Center right away. Keep out of reach of children. In case of overdose, get before use.

If pregnant or breast-feeding, ask a health professional

шасуливиλ

- use caubon when driving a motor vehicle or operating quom siu ess a snoi q sico pojic penetages
- sicopol, sedatives, and tranquitizers may increase When using this product matked drows ness may occur

sedatives or tranquilizers. Ask a doctor or pharmacist before use if you are taking

- difficulty in unination due to enlargement of the prostate pronchibs
- a preathing problem such as emphysema or chronic Ask a doctor before use if you have 🔳 glaucoma

Do not use for children under 2 years of age unless directed by Warnings

sesociated with motion sickness: a nausea a vomiting USES for prevention and treatment of these symptoms

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

MOTION SICKNESS RELIEF

dimenhydrinate tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63941-198

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIMENHYDRINATE (UNII: JB937PER5C) (DIPHENHYDRAMINE - UNII:8GTS82S83M, 8-CHLOROTHEOPHYLLINE - UNII:GE2UA340FM)	DIMENHYDRINATE	50 mg

Inactive Ingredients	
Ingredient Name	Strength
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	white	Score	2 pieces
Shape	ROUND	Size	9mm
Flavor		Imprint Code	44;198
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:63941-198- 02	2 in 1 CARTON	02/15/2019		
,		6 in 1 BLISTER PACK; Type 0: Not a Combination			

P	roduct		
Marketing In	nformation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M009	02/15/2019	

Labeler - Valu Merchandisers Company (868703513)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(63941-198)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(63941-198) , pack(63941-198)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(63941-198)

Revised: 12/2023 Valu Merchandisers Company