# MOTION SICKNESS RELIEF- dimenhydrinate tablet Topco Associates, LLC

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

# 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**

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

adults and	1 to 2 tablets every 4-6
children 12	hours; do not exceed 8
years and	tablets in 24 hours, or as
over	directed by a doctor
children 6	½ to 1 tablet every 6-8
to	hours; do not exceed 3
under 12	tablets in 24 hours, or as
years	directed by a doctor
children 2	½ tablet every 6-8 hours; do not
to	exceed 1½ tablets in 24
under 6 years	hours, or as directed by a doctor
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#### Other information

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

## Inactive ingredients

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

### Questions or comments?

1-888-423-0139

# Principal display panel

**TopCare**<sub>®</sub> health

NDC 36800-930-12

COMPARE TO DRAMAMINE® ORIGINAL FORMULA ACTIVE INGREDIENT\*

ORIGINAL FORMULA

Motion Sickness Relief

DIMENHYDRINATE, 50 mg - ANTIEMETIC

### **Prevents Nausea, Dizziness & Vomiting**

#### 12 TABLETS

For Children & Adults

actual size

\*This product is not manufactured or distributed by Medtech Products Inc., owner of the registered trademark Dramamine® Original Formula. 50844 REV0518C19802

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





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#### QUALITY GUARANTEED

B-1910-198-02-RR REV0518C19802

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Scan here for more information or call 1-888-423-0139

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children 2 years under 6 years	% tablet every 6-8 hours, or as directed 1% tablets in 24 hours, or as directed by a doctor
children 6 to under 12 years	1/s to 1 tablet every 6-8 hours, do not exceed 3 tablets in 24 hours, or as directed by a doctor
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KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

Drug Facts



**TopCare 44-198** 

### **MOTION SICKNESS RELIEF**

dimenhydrinate tablet

<b>Product Inf</b>	formation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:36800-930

Route of Administration ORAL

### **Active Ingredient/Active Moiety**

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

DIMENHYDRINATE 50 mg

### **Inactive Ingredients**

**Ingredient Name** 

Strength

CROSCARMELLOSE SODIUM (UNII: M280L1HH48)

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:36800-930- 12	2 in 1 CARTON	12/01/1992			
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product				

Marketing Information				
Marketing Application Number or Monograph Marketing Start Marketing Category Citation Date Date				
OTC Monograph Drug	M009	12/01/1992		

# Labeler - Topco Associates, LLC (006935977)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		038154464	pack(36800-930)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(36800-930) , pack(36800-930)

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		117025878	manufacture(36800-930)	

Revised: 5/2024 Topco Associates, LLC