**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 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 | ½ to 1 tablet every 6-8 hours; do not exceed 3 tablets in 24 hours, or as directed by a doctor |
| children 2 to under 6 years | ½ tablet every 6-8 hours; do not exceed 1½ tablets in 24 hours, or as directed by a doctor |

***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®**
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**

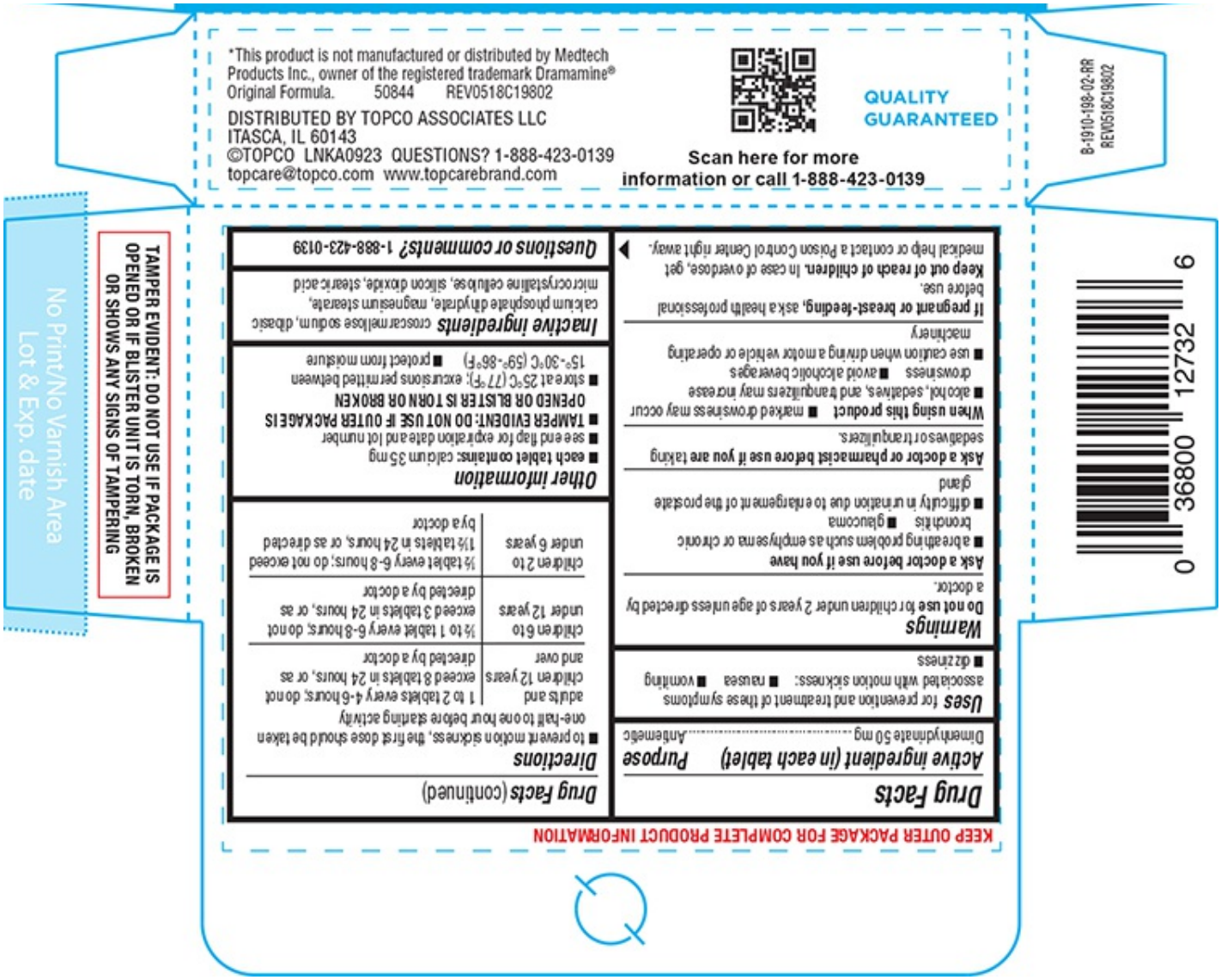
For Children & Adults

actual size

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

DISTRIBUTED BY TOPCO ASSOCIATES LLC
ITASCA, IL 60143
©TOPCO LNKA0923 QUESTIONS? 1-888-423-0139
topcare@topco.com www.topcarebrand.com

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




**TopCare 44-198**

# MOTION SICKNESS RELIEF

dimenhydrinate tablet

| Product Information | | | |
|---|---|---|---|
| **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** |
| **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** | | | |

## 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 Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M009 | 12/01/1992 | |

## Labeler - Topco Associates, LLC (006935977)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| LNK International, Inc. | | 038154464 | pack(36800-930) |

## Establishment

| 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