

MOTION SICKNESS RELIEF- dimenhydrinate tablet
Chain Drug Marketing Association, Inc.

Quality 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
- 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 | 1/2 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 | 1/2 tablet every 6-8 hours; do not exceed 1&1/2 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?

1-800-426-9391

Principal Display Panel

NDC 83324-032-12

**QC®
QUALITY
CHOICE**

***Compare to the
Active Ingredient in
Dramamine® Original Formula**

**Motion Sickness Relief
Original Formula
Dimenhydrinate 50 mg | Antiemetic**

Prevents: Nausea, Vomiting & Dizziness
for Children & Adults

12 Tablets

**actual
size**

**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

**SATISFACTION
GUARANTEED
100% QC**

Distributed by CDMA, Inc.
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362



Motion Sickness Relief

Original Formula

Dimenhydrinate 50 mg | Antiemetic

Prevents: Nausea, Vomiting & Dizziness
for Children & Adults

NDC 83324-032-12



*Compare to the
Active Ingredient in
Dramamine® Original Formula

Motion Sickness Relief

Original Formula

Dimenhydrinate 50 mg | Antiemetic

Prevents: Nausea, Vomiting & Dizziness
for Children & Adults

actual
size



12 Tablets

E7732RZ

Distributed by CDMA, Inc.
Novi, MI 48375
Questions: 800-935-2362
www.qualitychoice.com



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

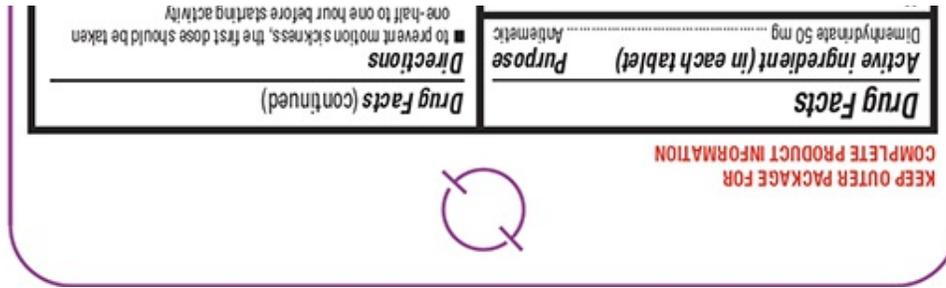
REV0518A19802
B-0220-188-02-R

No print/No varnish
Lot & Exp date

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

| | |
|--|--|
| <p>Uses for prevention and treatment of these symptoms associated with motion sickness: ■ nausea ■ vomiting ■ dizziness</p> | |
| <p>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 ■ sedatives or tranquilizers. Ask a doctor or pharmacist before use if you are taking ■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ marked drowsiness may occur ■ use caution when driving a motor vehicle or operating machinery ■ avoid alcoholic beverages</p> | |
| <p>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.</p> | |
| <p>Other information ■ each tablet contains: calcium 3 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</p> | |
| <p>Inactive ingredients croscarmellose sodium, dibasic calcium phosphate dihydrate, magnesium stearate, microcrystalline cellulose, silicon dioxide, stearic acid</p> | |
| <p>Questions or comments? 1-800-426-9391</p> | |





Quality_Chocie_44-198

MOTION SICKNESS RELIEF

dimenhydrinate tablet

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:83324-032 |
| 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 |
|---|----------|
| CROSCARMELOSE 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:83324-032-12 | 2 in 1 CARTON | 04/04/2024 | |
| 1 | | 6 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |

| | | | | |
|---|------------------|--|------------|--|
| 2 | NDC:83324-032-24 | 4 in 1 CARTON | 04/04/2024 | |
| 2 | | 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 | 04/04/2024 | |

Labeler - Chain Drug Marketing Association, Inc. (011920774)

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 038154464 | pack(83324-032) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|--|
| LNK International, Inc. | | 832867837 | manufacture(83324-032) , pack(83324-032) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|------------------------|
| LNK International, Inc. | | 117025878 | manufacture(83324-032) |

Revised: 5/2025

Chain Drug Marketing Association, Inc.