

**MOTION SICKNESS- dimenhydrinate tablet**  
**Target Corporation**

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**Target 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 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°C-30°C (59°F-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?**

**Call 1-800-910-6874**

## **Principal Display Panel**

Compare to active ingredient in Dramamine® Original Formula\*

## **Motion Sickness Relief**

Dimenhydrinate 50 mg/Antiemetic

## **up&up**

- Helps prevent nausea, dizziness and vomiting from motion sickness
- For adults and children

Actual Size

**24** 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 REV0518F19808

094 01 0726 R00 C-002262-01-034-0000

Distributed by Target Corporation

Minneapolis, MN 55403

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NDC 11673-198-08



Target 44-198

## MOTION SICKNESS

dimenhydrinate tablet

### Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:11673-198 |
| <b>Route of Administration</b> | ORAL           |                           |               |

### Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|-----------------|-------------------|----------|
|-----------------|-------------------|----------|

|  |                      |
|--|----------------------|
| <b>DIMENHYDRINATE</b> (UNII: JB937PER5C) (DIPHENHYDRAMINE - UNII:8GTS82S83M, 8-CHLOROTHEOPHYLLINE - UNII:GE2UA340FM) | DIMENHYDRINATE 50 mg |
|--|----------------------|

### Inactive Ingredients

| Ingredient Name   | Strength |
|---|----------|
| <b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)               |          |
| <b>DIBASIC CALCIUM PHOSPHATE DIHYDRATE</b> (UNII: O7TSZ97GEP) |          |
| <b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)                  |          |
| <b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)          |          |
| <b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)                     |          |
| <b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)                        |          |

### Product Characteristics

|                 |       |                     |          |
|-----------------|-------|---------------------|----------|
| <b>Color</b>    | white | <b>Score</b>        | 2 pieces |
| <b>Shape</b>    | ROUND | <b>Size</b>         | 9mm      |
| <b>Flavor</b>   |       | <b>Imprint Code</b> | 44;198   |
| <b>Contains</b> |       |                     |          |

### Packaging

| # | Item Code        | Package Description                                    | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:11673-198-02 | 2 in 1 CARTON  | 12/01/1992           |                    |
| 1 |                  | 6 in 1 BLISTER PACK; Type 0: Not a Combination Product |                      |                    |
| 2 | NDC:11673-198-08 | 4 in 1 CARTON  | 12/01/1992           |                    |
| 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                                     | 12/01/1992           |                    |

**Labeler** - Target Corporation (006961700)

### Establishment

| Name                    | Address | ID/FEI    | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. |         | 038154464 | pack(11673-198)     |

### Establishment

| Name                    | Address | ID/FEI    | Business Operations                      |
|-------------------------|---------|-----------|--|
| LNK International, Inc. |         | 832867837 | manufacture(11673-198) , pack(11673-198) |

## Establishment

| Name                    | Address | ID/FEI    | Business Operations    |
|-------------------------|---------|-----------|------------------------|
| LNK International, Inc. |         | 117025878 | manufacture(11673-198) |

Revised: 3/2025

Target Corporation