

MOTION SICKNESS RELIEF- dimenhydrinate tablet
Rite Aid Corporation

Rite Aid 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

- alcohol, sedatives, and tranquilizers may increase drowsiness
- marked drowsiness may occur
- 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°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?

1-800-426-9391

Principal Display Panel

NDC 11822-0198-7

Compare to the active ingredient in
Dramamine® Original Formula*

MOTION SICKNESS

RELIEF

DIMENHYDRINATE 50 mg

ANTIEMETIC**PREVENTS MOTION SICKNESS**

Prevents nausea, vomiting & dizziness

For children & adults

ACTUAL SIZE

36 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 REV0518G19807

DISTRIBUTED BY: RITE AID,
200 NEWBERRY COMMONS
ETTERS, PA 17319
www.riteaid.com

**SATISFACTION
GUARANTEE**

If you're not satisfied, we'll
happily refund your money.

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:11822-0198-2	2 in 1 CARTON	12/01/1992	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11822-0198-7	6 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 - Rite Aid Corporation (014578892)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(11822-0198)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(11822-0198) , pack(11822-0198)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(11822-0198)