MOTION SICKNESS- dimenhydrinate tablet
Geiss, Destin & Dunn Inc.

GoodSense 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°-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

GoodSense®

NDC 50804-198-02

Original Formula

Motion Sickness
Dimenhydrinate 50 mg
Antiemetic

Prevents Nausea, Vomiting & Dizziness
for Children & Adults

12 Tablets

actual size

Compare to the active ingredient of
Dramamine® Original Formula

100%
SATISFACTION
GUARANTEED

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 REV0518B19802

Distributed by: Perrigo Direct, Inc., Peachtree City, GA 30269
www.PerrigoDirect.com (1-800-426-9391)
GoodSense® is a registered trademark of
L. Perrigo Company.

GOODSENSE®

NDC 50804-198-02

Motion Sickness

Dimenhydrinate 50 mg
Antiemetic

12 Tablets

GOODSENSE®

Motion Sickness

GOODSENSE®

NDC 50804-198-02

Original Formula

Motion Sickness

Dimenhydrinate 50 mg
Antiemetic

Prevents Nausea, Vomiting & Dizziness for Children & Adults

actual size

12 Tablets

***Compare to the active ingredient of Dramamine® Original Formula**

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50844 REV0518819802

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L. Perrigo Company.

8 46036 00089 4

Z-Fold
LEBG30210⁴

8-0560-198-02-R
REV0518819802

no print / no varnish area
lot no. & exp. date

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

GOODSENSE®

Motion Sickness

Dimenhydrinate 50 mg
Antiemetic

■ nausea ■ vomiting ■ dizziness
associated with motion sickness.

Warnings

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■ 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.

children 2 to under 6 years	¼ tablet every 6-8 hours; do not exceed 1½ 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
12 years and over	1 to 2 tablets every 6-8 hours, or as directed by a doctor; do not exceed 6 tablets in 24 hours

Other information

■ **each tablet contains:** calcium 35 mg
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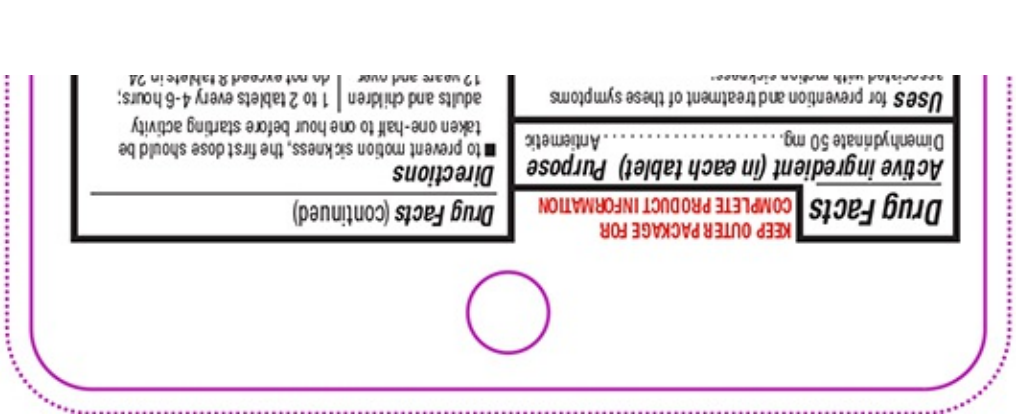

Good Sense 44-198

MOTION SICKNESS

dimenhydrinate tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-198
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:50804-198-02	2 in 1 CARTON	04/07/2020	

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	04/07/2020	

Labeler - Geiss, Destin & Dunn Inc. (076059836)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(50804-198)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(50804-198) , pack(50804-198)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(50804-198)

Revised: 11/2023 Geiss, Destin & Dunn Inc.