MOTION SICKNESS ORIGINAL FORMULA- dimenhydrinate tablet Mckesson (Sunmark)

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Dimenhydrinate 50 mg

Drug Facts

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

in 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 drinks
- 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 care 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 $\frac{1}{2}$ to 1 hour before starting activity
- do not exceed recommended dosage. Not for frequent or prolonged use except on the advice of a
 doctor

adults and children 12 years	take 1 to 2 tablets every 4 to 6 hours; not to exceed 8 tablets in 24 hours
of age and over	or as directed by a doctor
children 6 to under 12 years	take 1/2 to 1 tablet every 6 to 8 hours; not to exceed 3 tablets in 24 hours
of age	or as directed by a doctor
children 2 to under 6 years o	f take 1/4 to 1/2 tablet every 6 to 8 hours; not to exceed 1- 1/2 tablets in
age	24 hours or as directed by a doctor.

Other information

- each tablet contains: calcium 30 mg
- store below 86° F (30° C)
- protect from heat and humidity

Inactive ingredients

Croscarmellose Sodium, Dicalcium Phosphate, Magnesium Stearate and Microcrystalline Cellulose.

PRINCIPAL DISPLAY PANEL

Compare to Dramamine® Active Ingredient*

Helps prevent nausea, vomiting or dizziness associated with motion sickness

Motion Sickness

Original formula

For children & Adults

Dimenhydrinate

Fast Acting

THIS PRODUCT IS PACKAGED IN A CHILD RESISTANT AND TAMPER EVIDENT PACKAGE. USE ONLY IF BLISTERS ARE INTACT.

*This product is not manufactured or distributed by McNeil- PPC, Inc., owner of the registered trademark Dramamine®

ANOTHER QUALITY PRODUCT DISTRIBUTED BY McKesson

one post street, San Francisco, CA 94104

Money back gurantee

Please visit us at www.sunmarkbrand.com

Questions or comments? Call toll free: 1-877-753-3935

Product Label

COMPARE TO DRAMAMINE®
ACTIVE INGREDIENT*
NDC 49348-070-02

Helps to prevent nausea, vomiting or dizziness associated with motion sickness

<mark>sun</mark>mark[®]

motion sickness

Original Formula

For children & adults **DIMENHYDRINATE**

FAST ACTING

12 TABLETS 50 mg EACH

THIS PRODUCT IS PACKAGED IN A CHILD RESISTANT AND TAMPER EVIDENT PACKAGE. USE ONLY IF BLISTERS ARE INTACT.

No.: Date:

PLD-A F63SM12

Purpose

Another Quality Product
Distributed By McKesson
One Post Street, San Francisco, CA 94104
Money Back Guarantee
Please visit us at www.sunmarkbrand.com
Questions or comments? Call toll free: 1-877-753-3935

*This product is not manufactured or distributed by McNeil-PPC, Inc., owner of the registered trademark Dramamine@.

Inactive ingredients croscarmellose sodium, dicalcium phosphate, magnesium stearate and

Ochoer information = each tablet contains: calcium 30 mg = store below 86°F (30°C) = protect

take 1/2 to 1 tablet every 6 to 8 hours, not to exceed 3 tablets in 24 hours or as directed by a doctor take 1/4 to 1/2 tablet every 6 to 8 hours, not to exceed 1-1/2 tablets in 24 hours or as directed by a doctor

children 2 to under 6 years of age children 6 to under 12 years of age

forevent motion sickness, the first dose should be taken ½ to 1 hour before starting activity to be done to make a done to exceed recommended dosage, to bot for trepring to 2 habites and this and children 12 years of age and over the start of 24 hours or as directed by a doctor applicable to the start of the st

if **pregnant or breast-feeding**, ask a heelifn care professional before use. **Keep out of reach of children.** In case of overdose, get medical help or contact a Poison Control Center right away

tranquilizers may increase drowsiness 🔳 use caution when driving a motor vehicle or operating machinery When using this product marked drowsiness may occur, mavoid alcoholic drinks malcohol, sedatives, and

Ask a doctor or pharmacist betore use if you are taking sedatives or tranquilizers.

Do not use in children under 2 years of age, unless directed by a doctor.

Ask a doctor before use if you have a a breakhing problem such as emphysems or chronic bronchilis

glaucoms a difficulty in urination due to an enlargement of the prostate gland

Uses for prevention and treatment of these symptoms associated with motion slokness:

nauses

nomitting

Active ingredient (in each tablet) Drug Facts

MOTION SICKNESS ORIGINAL FORMULA

dimenhydrinate tablet

Product Information			
Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:49348-070
Route of Administration	ORAL	DEA Schedule	

Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	DIMENHYDRINATE (DIPHENHYDRAMINE)	DIMENHYDRINATE	50 mg

Inactive Ingredients		
Ingredient Name	Strength	
CROSCARMELLOSE SODIUM		
CALCIUM PHO SPHATE, DIBASIC, ANHYDRO US		
MAGNESIUM STEARATE		
CELLULO SE, MICRO CRYSTALLINE		

Product Characteristics				
Color	WHITE	Score	2 pieces	
Shape	ROUND	Size	9 mm	
Flavor		Imprint Code	1006;1006	
Contains				

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:49348-070-02	12 in 1 CARTON		
1	12 in 1 BLISTER PACK		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part336	06/13/2011	

$\boldsymbol{Labeler} \text{ - } \operatorname{Mckesson}\left(\operatorname{Sunmark}\right) (177667227)$

Revised: 10/2012 Mckesson (Sunmark)