DIMENHYDRINATE- dimenhydrinate tablet Major Pharmaceuticals

1006-Major

Drug Facts

Active ingredient (in each tablet)

Dimenhydrinate 50 mg

Purpose

Antiemetic

Uses

for the prevention and treatment of nausea, vomiting, or dizziness associated with motion sickness

Do not give to children under 2 years of age unless directed by a doctor

Ask a doctor before use if you have

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged 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 professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

■ to prevent motion sickness, the rst dose should be taken one-half to one hour before starting activity

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

Other information

- each tablet contains: calcium 30 mg
- store in a dry place at 15° 30°C (59° 86°F)
- protect from moisture
- see end -ap for expiration date and lot number

■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN

croscarmellose sodium, dicalcium phosphate, magnesium stearate, microcrystalline cellulose

Questions or comments?

1-800-231-4670

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

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

Distributed by: MAJOR® PHARMACEUTICALS Indianapolis, IN 46268 Questions or comments? Call (800) 616-2471 www.majorpharmaceuticals.com

MAJOR ®

NDC 0904-6772-12

Compare to the Active Ingredient in DRAMAMINE ® Original Formula*

Driminate TM

Dimenhydrinate USP, Antiemetic

For Nausea, Dizziness and Vomiting from Motion Sickness

12 Tablets 50mg Each

Use as Directed





Compare to the active ingredient in DRAMAMINE® Original Formula*

R52064

Driminate[™]

For Nausea, Dizziness and Vomiting from Motion Sickness

12 Tablets 50 mg EACH Use as Directed

MAJOR®

NDC 0904-6772-12 Compare to the active ingredient in DRAMAMINE® Original Formula*

Driminate

Dimenhydrinate USP, Antiemetic

For Nausea, Dizziness and Vomiting from Motion Sickness

12 TABLETS 50 mg EACH

Use as Directed

1-800-231-4670 Unestions or comments?

microcrystalline cellulose sodium, dicalcium phosphate, magnesium stearate, Inactive ingredients croscarmellose

Drug Facts (continued)

IS OPENED OR BLISTER IS TORN OR BROKEN ■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE

■ see end flap for expiration date and lot number place at 15° – 30°C (59° – 86°F) ■ protect from moisture ■ each tablet contains: calcium 30 mg ■ store in a dry

Other information

directed by a doctor

6-8 hours; not to exceed 1 1/2 tablets in 24 hours, or as ■ children 2 to under 6 years of age: 1/2 tablet every

directed by a doctor every 6-8 hours; not to exceed 3 tablets in 24 hours, or as

■ children 6 to under 12 years of age: 1/2 to 1 tablet hours, or as directed by a doctor

tablets every 4-6 hours; not to exceed 8 tablets in 24

■ adults and children 12 years of age and over: 1 to 2

■ to prevent or treat motion sickness, see below: taken one-half to one hour before starting activity ■ to prevent motion sickness, the first dose should be

Directions

(1-800-222-1222) right away. medical help or contact a Poison Control Center Keep out of reach of children. In case of overdose, get обыздид шасипету

- muse caution when driving a motor vehicle or increase drowsiness
 - alcohol, sedatives, and tranquilizers may
 - avoid alcoholic drinks
 - marked drowsiness may occur

When using this product

taking sedatives or tranquilizers

Ask a doctor or pharmacist before use if you are

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 - аізпсошя

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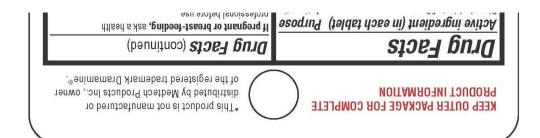
Do not give to children under 2 years of age unless Marnings

assenizzib 🔳 gnitimov 🖿 səsusn 🖿 sesociated with motion sickness: Use for prevention and treatment of these symptoms

M-29

Rev. 09/23

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DIMENHYDRINATE

dimenhydrinate tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-6772
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIMENHYDRINATE (UNII: JB937PER5C) (CHLORTHEOPHYLLINE - 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)			

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904- 6772-12	1 in 1 CARTON	02/14/2019	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information				
Marketing	Application Number or Monograph	Marketing Start	Marketing End	
Category	Citation	Date	Date	

OTC Monograph Drug	M009	02/14/2019	

Labeler - Major Pharmaceuticals (191427277)

Revised: 5/2024 Major Pharmaceuticals