DRAMAMINE- dimenhydrinate tablet
Select Corporation

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.

Dramamine®

Drug Facts

Active ingredient (in each tablet)

Dimenhydrinate 50 mg

Purpose

Antiemetic

Use

for prevention and treatment of these symptoms associated with motion sickness:
- nausea
- vomiting
- dizziness

Warnings

Do not give to 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
- 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
- be careful when driving a motor vehicle or operating machinery

If pregnant or breast-feeding, ask a doctor before use.

Keep out of reach of children.

In case of accidental overdose, get medical help or contact a Poison Control Center (1-

800-222-1222) immediately.

Directions

- to prevent motion sickness, the first dose should be taken 1/2 to 1 hour before starting activity
- to prevent or treat motion sickness, see below:

adults and children 12 years and over	• take 1 to 2 chewable tablets every 4-6 hours • do not take more than 8 chewable tablets in 24 hours, or as directed by a doctor
children 6 to under 12 years	• give 1/2 to 1 chewable tablet every 6-8 hours • do not give more than 3 chewable tablets in 24 hours, or as directed by a doctor
children 2 to under 6 years	• give 1/2 chewable tablet every 6-8 hours • do not give more than 1½ chewable tablets in 24 hours, or as directed by a doctor

Other information

- **Phenylketonurics:** contains phenylaline 0.84 mg per tablet
- store at room tempurature 20°- 25°C (68°-77°F)
- **do not use if pouch is opened**
- see bottom of this panel for lot number and expiration date

Inactive ingredients

anhydrous citric acid, aspartame, FD&C yellow #6 aluminum lake, flavors, magnesium stearate, maltodextrin, methacrylic acid copolymer, modified starch, sorbitol

Questions or comments?

call **1-800-382-7219**

Dist By:
Medtech Products, Inc. Tarrytown, NY 10591

Product Repackaged and Distributed
with permission of manufacturer by:
Select Corporation Carrollton, TX 75007

PRINCIPAL DISPLAY PANEL - 50 mg Tablet Packet Carton

#1
Pharmacist
Recommended
BRAND

DIMENHYDRINATE
TABLETS / ANTIEMETIC

Dramamine®

motion sickness

CHEWABLE

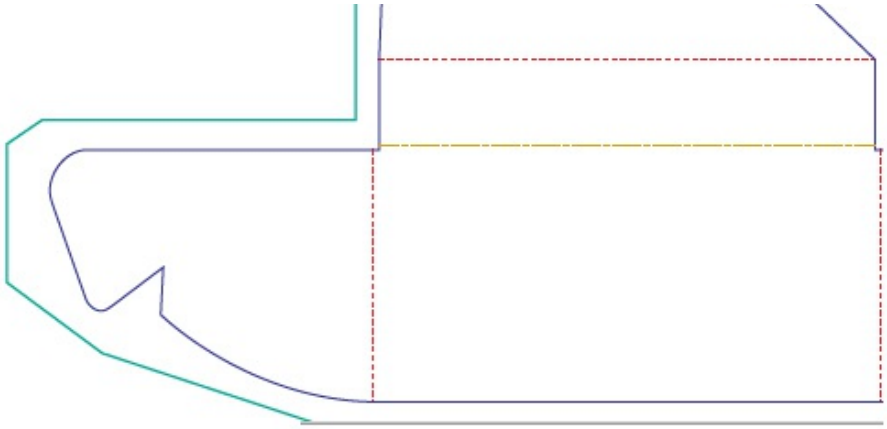
Dual Action:

Prevents & Relieves Nausea,
Dizziness and Vomiting

TO OPEN
PUSH IN TAB AND PULL OUT

25 Packets of 2 Orange Flavor Tablets
(50 mg each)


DIMENHYDRINATE TABLETS / ANTIEMETIC
Dramamine®
motion sickness
CHEWABLE
Dual Action:
Prevents & Relieves Nausea, Dizziness and Vomiting
#1 Pharmacist Recommended BRAND
#1 Pharmacist Recommended BRAND
DIMENHYDRINATE TABLETS / ANTIEMETIC
Dramamine®
motion sickness
CHEWABLE
Dual Action:
Prevents & Relieves Nausea, Dizziness and Vomiting
#1 Pharmacist Recommended BRAND
DIMENHYDRINATE TABLETS / ANTIEMETIC
Dramamine®
motion sickness
CHEWABLE
Dual Action:
Prevents & Relieves Nausea, Dizziness and Vomiting
TO OPEN
PUSH IN TAB AND PULL OUT
25 Packets of 2 Orange Flavor Tablets
(50 mg each)



Drug Facts

Active Ingredient (In each tablet) — **Purpose**

Dimenhydrinate 50 mg ..... Antiemetic

Use

for prevention and treatment of these symptoms associated with motion sickness:

- nausea
- vomiting
- dizziness

Warnings

Do not give to 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
- 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
- be careful when driving a motor vehicle or operating machinery

If pregnant or breast-feeding, ask a doctor before use.

Keep out of reach of children.

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

Directions

- to prevent motion sickness, the first dose should be taken 1/2 to 1 hour before starting activity
- to prevent or treat motion sickness, see below:

adults and children 12 years and over	• take 1 to 2 chewable tablets every 4-6 hours • do not take more than 8 chewable tablets in 24 hours, or as directed by a doctor
children 6 to under 12 years	• give 1/2 to 1 chewable tablet every 6-8 hours • do not give more than 3 chewable tablets in 24 hours, or as directed by a doctor
children 2 to under 6 years	• give 1/2 chewable tablet every 6-8 hours • do not give more than 1½ chewable tablets in 24 hours, or as directed by a doctor

Drug Facts (continued)

Other Information

- **Phenylketonurics:** contains phenylaline 0.84 mg per tablet
- store at room tempurature 20°- 25°C (68°-77°F)
- **do not use if pouch is opened**
- see bottom of this panel for lot number and expiration date

Inactive Ingredients

anhydrous citric acid, aspartame, FD&C yellow #6 aluminum lake, flavors, magnesium stearate, maltodextrin, methacrylic acid copolymer, modified starch, sorbitol

Questions or comments? call 1-800-382-7219

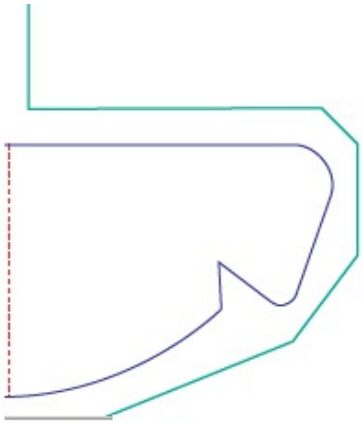
www.dramamine.com

Product Repackaged and Distributed
with permission of manufacturer by:
Select Corporation Carrollton, TX 75007

Select Corporation

6 55708 01962 7

2101962239



DRAMAMINE

dimenhydrinate tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52904-962(NDC:63029-901)
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**
silicon dioxide (UNII: ETJ7Z6XBU4)	
croscarmellose sodium (UNII: M28OL1HH48)	
lactose, unspecified form (UNII: J2B2A4N98G)	
magnesium stearate (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

Product Characteristics			
Color	ORANGE	**Score**	2 pieces
Shape	ROUND	**Size**	8mm
Flavor	ORANGE	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52904-962-04	1 in 1 BLISTER PACK	01/15/2012	
1		2 in 1 POUCH; Type 0: Not a Combination		

1		Product		
2	NDC:52904-962-25	25 in 1 CARTON	01/15/2012	
2		2 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part336	01/15/2012	

Labeler - Select Corporation (053805599)

Revised: 4/2022 Select Corporation