

**DRAMAMINE- dimenhydrinate tablet, chewable**  
**Lil' Drug Store Products, Inc.**

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**Dramamine**

**DRUG FACTS**

***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 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
- 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) right away.

## **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:

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Adults and children 12 years and over	<ul style="list-style-type: none"><li>▪ Take 1 to 2 chewable tablets every 4-6 hours</li><li>▪ Do not take more than 8 chewable tablets in 24 hours, or as directed by a doctor</li></ul>
Children 6 to under 12 years	<ul style="list-style-type: none"><li>▪ Give 1/2 to 1 chewable tablet every 6-8 hours</li><li>▪ Do not give more than 3 chewable tablets in 24 hours, or as directed by a doctor</li></ul>
Children 2 to under 6 years	Give 1/2 chewable tablet every 6-8 hours Do not give more than 1 1/2 chewable tablets in 24 hours, or as directed by a doctor

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## **Other Information**

- **Phenylketonurics:** contains phenylalanine 0.84 mg per tablet.
- store at room temperature 68°-77°F (20°-25°C)
- **do not use if packet is torn**
- see bottom 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?**

1-800-382-7219 Dramamine.com

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## **PRINCIPAL DISPLAY PANEL - 50mg Tablet Pouch Blister**

#1  
Pharmacist  
Recommended  
Brand  
  
Dimenhydrinate  
Tablets/Antiemetic

Dramamine®

motion sickness

CHEWABLE

Dual Action:

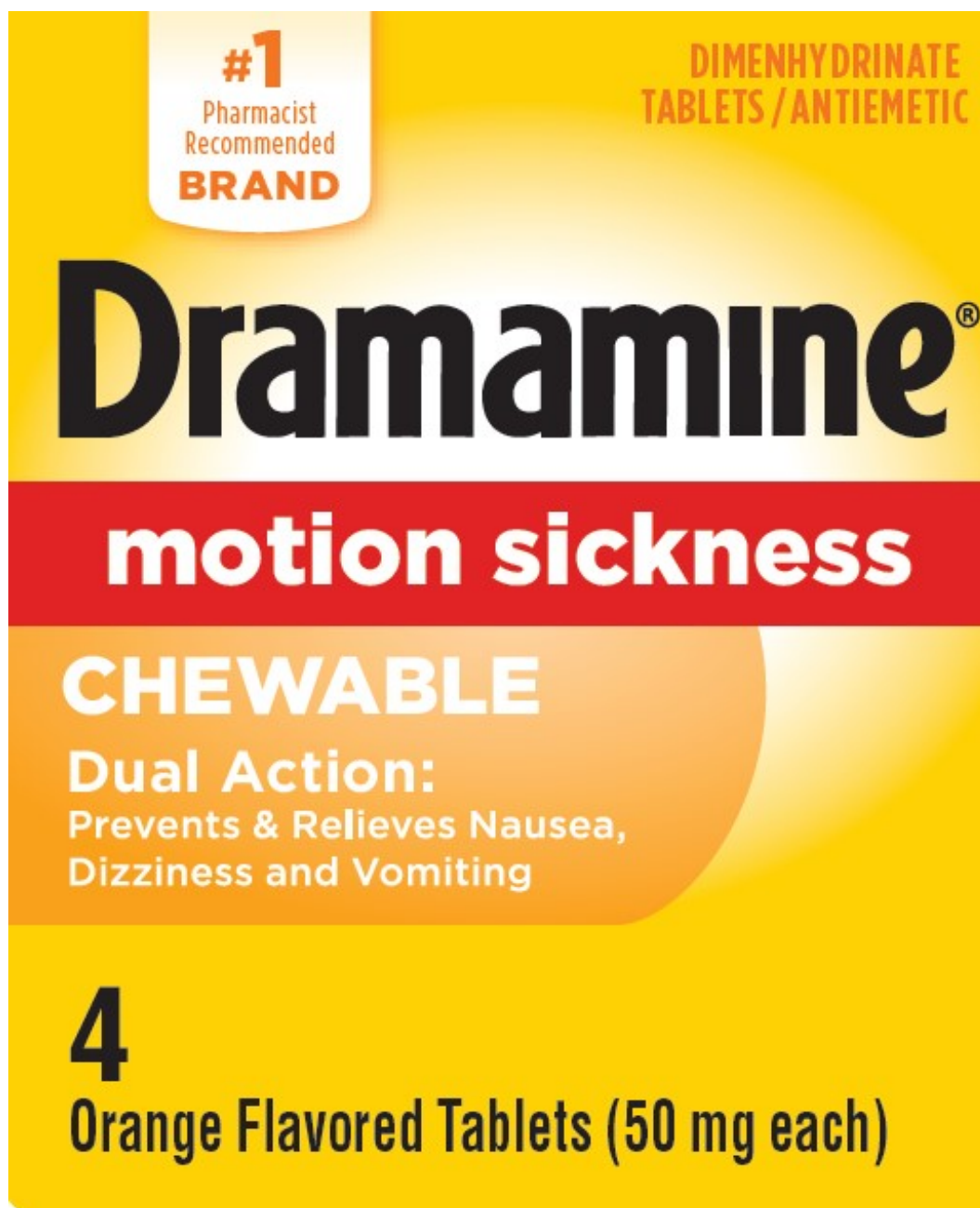
Prevents & Relieves Nausea,

Dizziness and Vomiting

4

Orange Flavored

Tablets (50 mg each)



The image shows the packaging for Dramamine motion sickness chewable tablets. The packaging is primarily yellow with a red horizontal band. At the top left, there is a white badge with a red border that says "#1 Pharmacist Recommended BRAND". To the right of this badge, the text "DIMENHYDRINATE TABLETS / ANTIEMETIC" is printed in red. The brand name "Dramamine®" is prominently displayed in large, bold, black letters. Below this, the words "motion sickness" are written in white on the red band. Underneath the red band, the word "CHEWABLE" is written in large, bold, white letters. Below "CHEWABLE", the text "Dual Action: Prevents & Relieves Nausea, Dizziness and Vomiting" is written in white. At the bottom left, the number "4" is written in large, bold, black letters, and below it, the text "Orange Flavored Tablets (50 mg each)" is written in bold, black letters.

**#1**  
Pharmacist  
Recommended  
**BRAND**

**DIMENHYDRINATE  
TABLETS / ANTIEMETIC**

**Dramamine®**

**motion sickness**

**CHEWABLE**

**Dual Action:**  
Prevents & Relieves Nausea,  
Dizziness and Vomiting

**4**  
**Orange Flavored Tablets (50 mg each)**

## PRINCIPAL DISPLAY PANEL - 50mg Tablet Pouch Blister

**#1**

Pharmacist

Recommended

**Brand**

Dimenhydrinate

Tablets/Antiemetic

**Dramamine** ®

motion sickness

**CHEWABLE**

Dual Action:

Prevents & Relieves Nausea,

Dizziness and Vomiting

**6**

**Orange Flavored Tablets (50 mg each)**



## PRINCIPAL DISPLAY PANEL - 50mg Tablet Pouch Box

#1  
 Pharmacist  
 Recommended  
 Brand  
 Dimenhydrinate  
 Tablets/Antiemetic  
 Dramamine ®

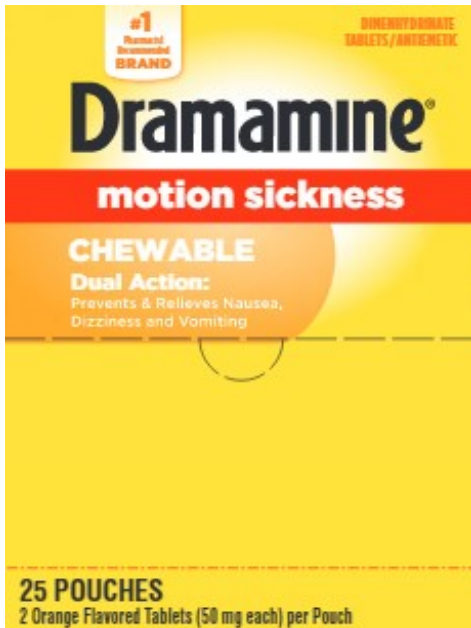
motion sickness

CHEWABLE  
 Dual Action:

Prevents & Relieves Nausea,  
 Dizziness and Vomiting

25 POUCHES

2 Orange Flavored Tablets (50 mg each) per Pouch



**DRAMAMINE**

dimenhydrinate tablet, chewable

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:29485-1216
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
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**DIMENHYDRINATE** (UNII: JB937PER5C) (CHLORTHEOPHYLLINE - UNII:GE2UA340FM) DIMENHYDRINATE 50 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1)</b> (UNII: 74G4R6TH13)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>ASPARTAME</b> (UNII: Z0H242BBR1)	
<b>STARCH, CORN</b> (UNII: O8232NY3S)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	

### Product Characteristics

<b>Color</b>	orange	<b>Score</b>	2 pieces
<b>Shape</b>	ROUND	<b>Size</b>	12mm
<b>Flavor</b>	ORANGE	<b>Imprint Code</b>	DRA
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29485-1216-2	2 in 1 BLISTER PACK	05/10/2010	12/31/2024
1		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 Drug	M009	01/01/1994	12/31/2024

## DRAMAMINE

dimenhydrinate tablet, chewable

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:29485-6925
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
DIMENHYDRINATE (UNII: JB937PER5C) (CHLORTHEOPHYLLINE - UNII:GE2UA340FM)		DIMENHYDRINATE	50 mg	
<b>Inactive Ingredients</b>				
Ingredient Name			Strength	
ALUMINUM OXIDE (UNII: LMI26O6933)				
MALTODEXTRIN (UNII: 7CVR7L4A2D)				
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)				
SORBITOL (UNII: 506T60A25R)				
ASPARTAME (UNII: Z0H242BBR1)				
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
STARCH, CORN (UNII: O8232NY3SJ)				
<b>Product Characteristics</b>				
Color	orange	Score	2 pieces	
Shape	ROUND	Size	12mm	
Flavor	ORANGE	Imprint Code	DRA	
Contains				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29485-6925-3	3 in 1 BLISTER PACK	02/06/2017	12/31/2024
1		2 in 1 POUCH; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M009	02/06/2017	12/31/2024	

<b>DRAMAMINE</b>			
dimenhydrinate tablet, chewable			
<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:29485-8048
Route of Administration	ORAL		
<b>Active Ingredient/Active Moiety</b>			

Ingredient Name		Basis of Strength	Strength
DIMENHYDRINATE (UNII: JB937PER5C) (CHLORTHEOPHYLLINE - UNII:GE2UA340FM)		DIMENHYDRINATE	50 mg

Inactive Ingredients		
Ingredient Name	Strength	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		
ALUMINUM OXIDE (UNII: LMI26O6933)		
MALTODEXTRIN (UNII: 7CVR7L4A2D)		
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)		
SORBITOL (UNII: 506T60A25R)		
ASPARTAME (UNII: Z0H242BBR1)		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
STARCH, CORN (UNII: O8232NY3S)		

Product Characteristics			
Color	orange	Score	2 pieces
Shape	ROUND	Size	12mm
Flavor	ORANGE	Imprint Code	DRA
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29485-8048-2	25 in 1 BOX	07/10/2020	12/31/2025
1		1 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 Drug	M009	01/01/1994	12/31/2025

**Labeler** - Lil' Drug Store Products, Inc. (093103646)

Revised: 12/2023

Lil' Drug Store Products, Inc.