

**MECLIZINE HCL- meclizine hcl tablet**  
**Aidarex Pharmaceuticals LLC**

*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.*

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**Travel Sickness**

**Active Ingredient**  
**(in each tablet)**

**Meclizine HCl 25 mg**

**Purpose**

**Antiemetic**

**Uses**

prevents and treats nausea, vomiting, or dizziness due to motion sickness

**WARNINGS**

**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**

- you may get drowsy
- avoid alcoholic drinks
- alcohol, sedatives & tranquilizers may increase drowsiness
- be careful 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**

- take dose one hour before travel starts
- tablets can be chewed or swallowed whole with water
- **adults & children 12 years and over:** 1-2 tablets once daily
- **children under 12 years:** ask a doctor

**Other Information**

- **phenylketonurics: each tablet contains:** phenylalanine 0.28mg
- store at room temperature 15°-30°C (59°-86°F)

**Inactive Ingredients**

aspartame, compressible sugar, croscarmellose sodium, dextrose, FD&C red # 40(Al-lake), magnesium stearate, microcrystalline cellulose, raspberry flavor

### Questions or Comments

Call 1-800-645-2158, 9 am – 5 pm ET, Monday - Friday

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

**Rugby® Duluth, Georgia 30097**

Repackaged By :  
Aidarex Pharmaceuticals LLC,  
Corona, CA 92880

### PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

Meclizine HCl

COMPARE TO ACTIVE INGREDIENT IN BONINE

Meclizine HCl 25 mg (ANTIEMETIC)

NDC 33261-0413-30

30 CHEWABLE TABLETS

Contains Aspartame

Rugby

Duluth, Georgia 30097

Repackaged By :  
Aidarex Pharmaceuticals LLC,  
Corona, CA 92880

CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT. KEEP OUT OF REACH OF CHILDREN. STORE AT CONTROLLED ROOM TEMP 15-30C (59-86F)

Packaged and Distributed by: **AIDAREX PHARMACEUTICALS LLC.**

**MECLIZINE CHEW.**

**25mg**

**30 TABS**

EACH TABLET CONTAINS THE FOLLOWING ACTIVE INGREDIENTS:  
MECLIZINE.....25 mg  
HYDROCHLORIDE  
CHEWABLE

PINK ROUND TABLET W/ LOGO P 115 ON ONE SIDE & SCORE ON REVERSE

GENERIC FOR : ANTIVERT  
NDC: 33261-0413-30

TAKE \_\_\_\_\_ EVERY \_\_\_\_\_ HOURS \_\_\_\_\_ TIMES A DAY  
TOME \_\_\_\_\_ CADA \_\_\_\_\_ HORAS \_\_\_\_\_ VECES AL DIA

MFG: FOR: RUGBY DULUTH, GEORGIA 30097

MECLIZINE CHEW. 25mg 30  
NDC: 33261-0413-30  
RXQLS0000

MECLIZINE CHEW. 25mg 30  
NDC: 33261-0413-30  
RXQLS0000

MECLIZINE CHEW. 25mg 30  
NDC: 33261-0413-30  
RXQLS0000

PEEL HERE  
PATIENT LOG CHART  
PEEL HERE

## MECLIZINE HCL

meclizine hcl tablet

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:33261-413(NDC:0536-3990)
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
MECLIZINE HYDROCHLORIDE (UNII: HDP7W44CIO) (MECLIZINE - UNII:3L5TQ84570)	MECLIZINE HYDROCHLORIDE	25 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
ASPARTAME (UNII: Z0H242BBR1)	
MAGNESIUM STEARATE (UNII: 70097M6B30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SUCROSE (UNII: C151H8M554)	
DEXTROSE (UNII: IY9XDZ35W2)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
RASPBERRY (UNII: 4N14V5R27W)	

**Product Characteristics**

<b>Color</b>	RED	<b>Score</b>	2 pieces
<b>Shape</b>	ROUND	<b>Size</b>	8mm
<b>Flavor</b>	RASPBERRY	<b>Imprint Code</b>	AP;115
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:33261-413-30	30 in 1 BOTTLE, PLASTIC		
2	NDC:33261-413-60	60 in 1 BOTTLE, PLASTIC		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC MONOGRAPH FINAL	part336	07/22/2008	

**Labeler** - Aidarex Pharmaceuticals LLC (801503249)