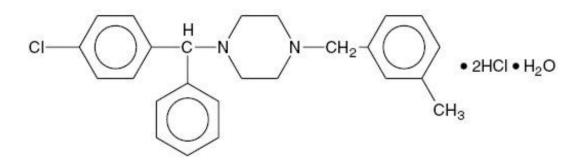
#### MECLIZINE HYDROCHLORIDE - meclizine hydrochloride tablet Life Line Home Care Services, Inc.

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# MECLIZINE HYDROCHLORIDE TABLETS, USP Rx only

#### DESCRIPTION

Chemically, Meclizine HCl is  $1-(p-chloro-\alpha-phenylbenzyl)-4-(m-methylbenzyl)$  piperazine dihydrochloride monohydrate.



Meclizine HCI Tablets, USP are available in two different strengths: 12.5 mg and 25 mg. In addition each tablet contains the following inactive ingredients: Colloidal Silicon Dioxide, Croscarmellose Sodium, Lactose Monohydrate, Magnesium Stearate, Microcrystalline Cellulose. Also, Meclizine HCI Tablets USP, 12.5 mg contains FD&C Blue #1 Aluminum Lake (11-13%) and Meclizine HCI Tablets USP, 25 mg contains D&C Yellow #10 Aluminum Lake (15-20%).

# **CLINICAL PHARMACOLOGY**

Meclizine Hydrochloride is an antihistamine which shows marked protective activity against nebulized histamine and lethal doses of intravenously injected histamine in guinea pigs. It has a marked effect in blocking the vasodepressor response to histamine, but only a slight blocking action against acetylcholine. Its activity is relatively weak in inhibiting the spasmogenic action of histamine on isolated guinea pig ileum.

#### INDICATIONS AND USAGE

For the management of nausea and vomiting, and dizziness associated with motion sickness.

# CONTRAINDICATIONS

Meclizine Hydrochloride is contraindicated in individuals who have shown a previous hypersensitivity to it.

#### WARNINGS

Since drowsiness may, on occasion, occur with use of this drug, patients should be warned of this possibility and cautioned against driving a car or operating dangerous machinery.

Patients should avoid alcoholic beverages while taking this drug.

Due to its potential anticholinergic action, this drug should be used with caution in patients with asthma, glaucoma or enlargement of the prostate gland.

# PRECAUTIONS

#### PREGNANCY, Teratogenic Effects

Pregnancy Category B. Reproduction studies in rats have shown cleft palates at 25-50 times the human dose. Epidemiological studies in pregnant women, however, do not indicate that medicine increases the risk of abnormalities when administered during pregnancy. Despite the animal findings, it would appear that the possibility of fetal harm is remote. Nevertheless, meclizine, or any other medication, should be used during pregnancy only if clearly necessary.

#### Pediatric Use

Clinical studies establishing safety and effectiveness in children have not been done; therefore, usage is not recommended in children under 12 years of age.

## **ADVERSE REACTIONS**

Drowsiness, dry mouth and, on rare occasions, blurred vision have been reported.

## DOSAGE AND ADMINISTRATION

#### **Motion Sickness**

The initial dose of 25 to 50 mg of Meclizine HCI should be taken one hour prior to travel for protection against motion sickness. Thereafter, the dose may be repeated every 24 hours for the duration of the journey.

#### HOW SUPPLIED

Meclizine HCI Tablets, USP are available in the following strengths and package sizes:

12.5 mg (Blue, oval-shaped, scored, debossed with TL122)

Bottles of 100	NDC 59746-122-06
Bottles of 1000	NDC 59746-122-10

25 mg (Yellow, oval-shaped, scored, debossed with TL121)

Bottles of 100	NDC 59746-121-06
Bottles of 1000	NDC 59746-121-10

Store at 20-25°C (68-77°F) (See USP Controlled Room Temperature].

Manufactured By:

Jubilant Cadista Pharmaceuticals Inc. Salisbury, MD 21801, USA.

Revised 03/11

#### PRINCIPAL DISPLAY PANEL

MECLIZINE HCL 25 MG QTY:30 TABLETS MFG: CADISTA MECLIZINE HCL MFG LOT: 1P584 MFG NDC: 597 4601 2106 PROD: NOP 02/20/12 10:13 AM	LIFE LINE 161 0 Ma dison Av e. Tif ton, GA 31794 NDC: 75921012130 EXP: 09/30/13
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MECLIZINE HYDROCHLORIDE						
meclizine hydrochloride ta	blet					
<b>Product Information</b>						
Product T ype	HUMAN PRESC	RIPTION DRUG	Item Code (Source)	NDC:75921-	121(NDC:	59746-121)
Route of Administration	ORAL					
Active Ingredient/Acti	ve Moiety					
Ingredient Name					rength	Strength
MECLIZINE HYDRO CHLOR	IDE (UNII: HDP7W44CIO)	(MECLIZINE - UN	III:3L5TQ84570)	MECLIZINE		25 mg
Inactive Ingredients						
Ingredient Name					St	rength
SILICON DIOXIDE (UNII: ET	J7Z6XBU4)					
CROSCARMELLOSE SODIU	J <b>M</b> (UNII: M28OL1HH48)					
LACTOSE MONOHYDRATE	E (UNII: EWQ57Q8I5X)					
MAGNESIUM STEARATE (U						
CELLULOSE, MICROCRYS		6 1U)				
D&C YELLOW NO. 10 (UNII						
ALUMINUM OXIDE (UNII: LN	MI26O6933)					
Product Characteristic		6		) nia		
Color	yellow OVAL			2 pieces		
Shape	OVAL			13mm		
Flavor		Imprint Code		TL121		
Contains						
Packaging						
# Item Code	Package Description	on Ma	rketing Start Date	Mark	eting Er	nd Date
	30 in 1 BOTTLE		Incling Start Date	IVIAT K	eting El	
1 1100.70521-121-50						

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA040659	06/04/2010		

Labeler - Life Line Home Care Services, Inc. (844985374)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
Life Line Home Care Services, Inc.		844985374	repack, relabel

Revised: 2/2012

Life Line Home Care Services, Inc.