MECLIZINE HYDROCHLORIDE- meclizine hydrochloride tablet APNAR PHARMA LP

Meclizine Hydrochloride Tablets

Active Ingredient (in each tablet)

12.5mg tablet: Meclizine HCL.... 12.5 mg

25mg tablet: Meclizine HCL25 mg

Purpose

Antiemetic

Uses

Prevents and treats nausea, vomiting or dizziness associated with motion sickness

Directions

Meclizine HCL 12.5mg tablet :

Dosage should be taken one hour before travel starts -adults & children 12 years and over: take 2 or 4 tablets once daily or as directed by a doctor

Meclizine HCL 25mg tablet:

Dosage should be taken one hour before travel starts -adults & children 12 years and over: take 1 or 2 tablets once daily or as directed by a doctor

Inactive Ingredients

Magnesium stearate, microcrystalline cellulose, silicon dioxide, lactose monohydrate

Other Information:

store at 20° to 25°C (68° to 77°F), excursions permitted between 15° to 30°C (59° to 86°F) [see USP controlled room temperature]

Do not use in children under 12 years of age unless directed by a doctor

Do not take this product, unless directed by a doctor, if you have

- Glaucoma
- A breathing problem such as emphysema or chronic bronchitis
- Trouble urination due to an enlarged prostate gland

Do not take this product if you are taking

Sedatives or tranquilizers, without first consulting your doctor.

When using this product,

- Do not exceed recommended dosage
- may cause drowsiness
- alcohol, sedatives and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- 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 right away (1-800-222-1222)

Questions or Comments?

Call 1-855-642-2594

Rev.: 05/2022

Made in USA

Manufactured by:

APNAR PHARMA LLP

East Windsor, NJ 08520

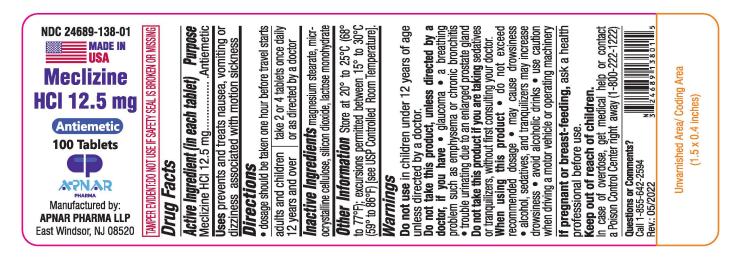
TAMPER EVIDENT: DO NOT USE IF SAFETY SEAL IS BROKEN OR MISSING

NDC: 24689-138-01

Meclizine Hydrochloride Tablets 12.5 mg

Antiemetic

100 Tablets



NDC: 24689-139-01

Meclizine Hydrochloride Tablets 25 mg

Antiemetic

100 Tablets



IPER EVIDENT:DO NOT USE IF SAFETY SEAL IS BROKEN OR MISSIN(

Active Ingredient (in each tablet)

Ises prevents and treats nausea, vomiting or associated with motion sickness Aeclizine HCI 25 ma

dosage should be taken one hour before travel starts take 1 or 2 tablets once daily or as directed by a doctor adults and children 2 years and over

ther Information Store at 20° to 25°C (68° between 15° to 30°C ocrystalline cellulose, silicon dioxide, lactose monohydrate **inactive Ingredients** magnesium stearate, micr-Room Temperature excursions permitted o 77°F); excursions p (59° to 86°F) [see USP

Warnings

Do not use in children under 12 years of age unless directed by a doctor.

Do not take this product, unless directed by a trouble urnating due to an enlarged prostate gland
 Do not take this product if you are taking sedatives

not exceed when driving a motor vehicle or operating machiner sedatives, alcohol,

if pregnant or breast-feeding, ask a healt

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

Questions or Comments? Call 1-855-642-2594 Rev.: 05/2022

Unvarnished Area/ Coding Area (1.5 x 0.4 inches)

MECLIZINE HYDROCHLORIDE

meclizine hydrochloride tablet

Product Information

HUMAN OTC DRUG **Product Type** Item Code (Source) NDC:24689-138

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength MECLIZINE HYDROCHLORIDE (UNII: HDP7W44CIO) (MECLIZINE -**MECLIZ INE** 12.5 mg **HYDROCHLORIDE** UNII:3L5TQ84570)

Inactive Ingredients Ingredient Name Strength MAGNESIUM STEARATE (UNII: 70097M6I30) **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X) MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)

Product Characteristics			
Color	white	Score	2 pieces
Shape	ROUND	Size	7mm
Flavor		Imprint Code	A;C4
Contains			

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24689-138-	100 in 1 BOTTLE; Type 0: Not a Combination	רכחכוחכודח	

Marketing	Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	M009	07/29/2022	

MECLIZINE HYDROCHLORIDE

meclizine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:24689-139
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Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
,, (MECLIZ INE HYDROCHLORIDE	25 mg

Inactive Ingredients		
Ingredient Name	Strength	
SILICON DIOXIDE (UNII: ETJ7Z 6XBU4)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		

Product Characteristics			
Color	white (White to off white)	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	AM7
Contains			

Ш	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:24689-139-	100 in 1 BOTTLE; Type 0: Not a Combination Product	07/29/2022	

Marketing In	formation		
Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M009	07/29/2022	

Labeler - APNAR PHARMA LP (079568229)

Establishment			
Name	Address	ID/FEI	Business Operations
INVAHEALTH INC		116840615	label(24689-138, 24689-139) , pack(24689-138, 24689-139)

Establishment			
Name	Address	,	Business Operations
APNAR PHARMA LLP		118530917	manufacture(24689-138, 24689-139), analysis(24689-138, 24689-139), pack(24689-138, 24689-139)

Revised: 1/2025 APNAR PHARMA LP