NAUSX- meclizine hydrochloride tablet, chewable Goldman Pharmaceutical Group Inc

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Naus X®

Drug Facts

Active ingredient (in each chewable tablet)	Purpose
Meclizine HCL, USP 25 mg	Antiemetic

Uses

• for the prevention and relief of nausea, vomiting, dizziness and other symptoms associated with motion sickness and/or travelling

Warnings

Do not use with any other product containing meclizine

- *ask a veterinarian before use* if your dog has any chronic illnesses
- ask a veterinarian or pharmacist if your dog is on any other medication before use
- *keep out of reach of children* in case of overdose contact poison control center immediately
- *stop use and contact veterinarian* if symptoms persist for more than 72 hours
- *do not use on dogs* under 26 pounds

Directions

- give one tablet 30 minutes prior to expected travel
- effects last approximately 6 hours
- **DO NOT** exceed two doses in 24 hours
- for dogs 26 75 pounds only

Other information

- store at room temperature 20 25 C (66 78 F)
- protect from moisture
- see end flap for expiration date and lot #

Inactive Ingredients

croscarmellose sodium, dicalcium phosphate dehydrate, magnesium stearate, microcrystalline cellulose, silicon dioxide, sodium sulfate, stearic acid, aspertame, dextrose, fd&c red #40 lake, maltodextrin, natural and artificial flavors, sugar, tricalcium phosphate

Questions or Comments

800.981.7642

petsotc.com

Distributed by

Goldman Pharmaceutical Group Inc.

Holbrook, NY 11741

Tamper-evident: Do not use if carton is open or blister unit is broken or torn

MADE IN AMERICA

PRINCIPAL DISPLAY PANEL - 25 mg Tablet Blister Pack Carton

pet otcTM

over the counter

 $\text{Naus} X^{\text{\circledR}}$

(Meclizine HCl, USP 25 mg) Antiemetic

PREVENTS

AND RELIEVES:

Dizziness

Nausea

Vomiting

TREATS SYMPTOMS

ON THE SPOT!

26 - 75 lbs

10

CHEWABLE

TABLETS

MADE IN THE USA

Recommended by veterinarians for

motion sickness

NDC 72087-012-10



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Goldman Pharmaceutical Group Inc. Holbrook, NY 11741

petsotc.com 2497,186,008

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Questions or Comments

and artificial flavors, sugar, tricalcium phosphate silicon dioxide, sodium sulfate, stearic acid, aspertame, dextrose, fd&c red #40 lake, maltodextrin, natural croscarmellose sodium, dicalcium phosphate dehydrate, magnesium stearate, microcrystalline cellulose,

Inactive Ingredients

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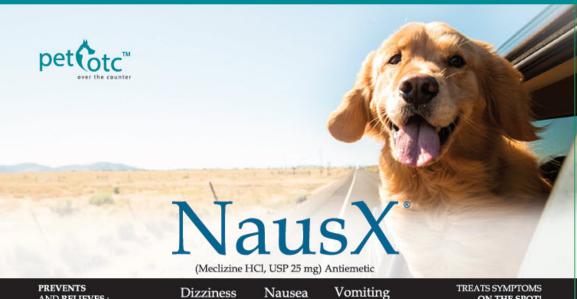
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Meclizine HCL, USP 25 mg Antiemetic Active ingredient (in each chewable tablet)

Drug Facts

NausX°

Vomiting Dizziness Nausea



NausX

Nausea

Vomiting

PREVENTS AND RELIEVES:

TREATS SYMPTOMS ON THE SPOT!



26 - 75 lbs



CHEWABLE TABLETS

MADE IN THE USA

Recommended by veterinarians for motion sickness

NDC 72087-012-10

Vomiting

Nausea

Dizziness



NausX'

NAUSX

meclizine hydrochloride tablet, chewable

Product Information			
Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:72087-012(NDC:0536-1018)
Route of Administration	ORAL		

l	Active Ingredient/Active Moiety			
l	Ingredient Name	Basis of Strength	Strength	
l	MECLIZINE HYDRO CHLO RIDE (UNII: HDP7W44CIO) (MECLIZINE - UNII:3L5TQ84570)	MECLIZINE HYDROCHLORIDE	25 mg	

Inactive Ingredients		
Ingredient Name	Strength	
ASPARTAME (UNII: Z0H242BBR1)		
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)		
DEXTROSE, UNSPECIFIED FORM (UNII: IY9 XDZ35W2)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MALTO DEXTRIN (UNII: 7CVR7L4A2D)		
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
SODIUM SULFATE (UNII: 0 YPR65R21J)		
SUCROSE (UNII: C151H8M554)		
TRICALCIUM PHO SPHATE (UNII: K4C08XP666)		

Product Characteristics			
Color	PINK (pink)	Score	2 pieces
Shape	ROUND (biconvex tablet with bisect)	Size	8 mm
Flavor		Imprint Code	21G
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:72087-012-20	10 in 1 BLISTER PACK		

Marketing Informa	ition		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		0 3/0 1/20 19	

Registrant - Goldman Pharmaceutical Group, Inc. (080389804)

Establishment			
Name	Address	ID/FEI	Business Operations
Goldman Pharmaceutical Group, Inc.		080389804	REPACK

Revised: 2/2019 Goldman Pharmaceutical Group Inc