YETI POT- SCHNOZ WASH- nasal cleaning salt granule, for solution HEALTHLAND LLC

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.

Warnings

Stop use and ask doctor if washing is uncomfortable or symptoms are not relieved.

When using this product

Use by only one person

Wash with soap and water after each use

Top rack of dishwasher safe

Do not heat in microwave

Keep out of reach of children

Other Information

- -Inspect saline solution packets for integrity
- -Do not use if open or torn
- -Protect saline solution packets from excessive heat andmoisture

Uses

- -Temporarily relieves systoms associated with rhinorrhea, cold, flu or allergies:
- -Sneezing Runny nose Nasal stuffiness Post Nasal drip
- -Removes inhaled irritants(dust, pollen)
- -Promotesnasal and sinus drainage
- -Help reduce swelling of nasal membranes

Drug Facts

Active Ingredients Purpose

(in each Packet)

Sodium Bicarbonate USP 515 mg Nasal Wash

Sodium Chloride USP 1685 mg

Directions

Adults and children 6 years and over: Use 1-2 packets every 2 hours as needed

Children under 6 years : Consult a physician

EASY TO USE INSTRUCTIONS

STEP ONE: Mix one packet of Schnoz Wash with distilled water up to the 240 ml mark in the Yeti Pot.

Replace the lid and shake well so that the solution mixes witht he water.

STEP TWO: Tilt your head and insert the Yeti Pot spout gently into the raised nostril, creating a seal between the pot and nostril. Raise the pot slowly to develop a steady flow through the upper nostril and out the lower nostril.

STEP THREE: During the process breathe through your mouth. Upon completion, exhale firmly several times to clear the nasal passages. Reverse the tilt of your head and repeat the other process on the other side.

Inactive Ingredient

None

Yeti Pot -Schnoz wash label



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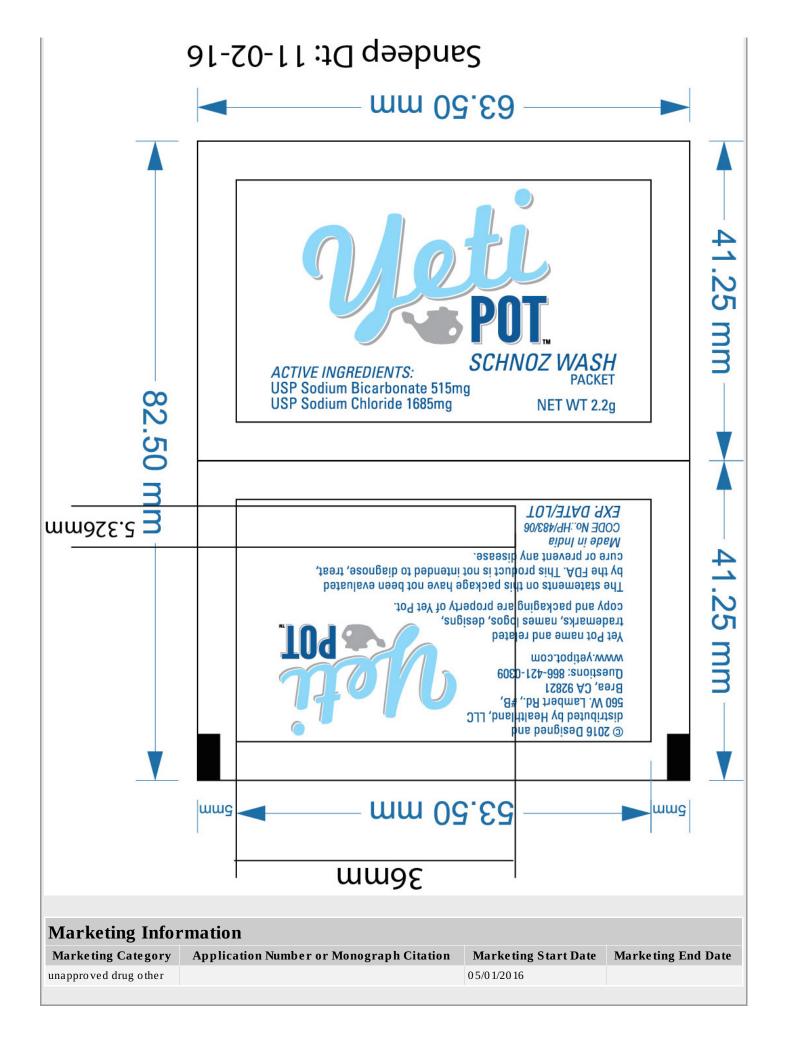
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70708-786	
Route of Administration	NASAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
SODIUM BICARBONATE (UNII: 8MDF5V39QO) (BICARBONATE ION - UNII:HN1ZRA3Q20)	SODIUM BICARBONATE	515 mg in 1 mg		
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM CHLORIDE	1685 mg in 1 mg		

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	240 mg in 1 mg

Product Characteristics				
Color	white (white crystalline granule powder)	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:70708-786-00	2200 mg in 1 PACKET; Type 0: Not a Combination Product	05/01/2016		



Labeler - HEALTHLAND LLC (079671169)

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