SALINE NASAL 1 FL OZ- sodium chloride 2.65% spray
Velocity Pharma 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.

Allergy and Sinus: Hypertonic Saline Mist: CVS Health

Active ingredient

Sodium Chloride, 2.65%

Purpose

Nasal Wash to flush allergens and dust from Nasal Passages

Uses

Nasal congestion
Removes inhaled irritants (dust, dirt, pollen)
Moisturizes dry nasal passages

Warnings

Do not use if seal is broken or missing.

Warnings

Use of this product by more than one person may spread infection
Contents under pressure
Do not puncture or incinerate
Avoid spraying in eyes

Keep out of reach of children. The use of this dispenser by more than one person may spread infection.

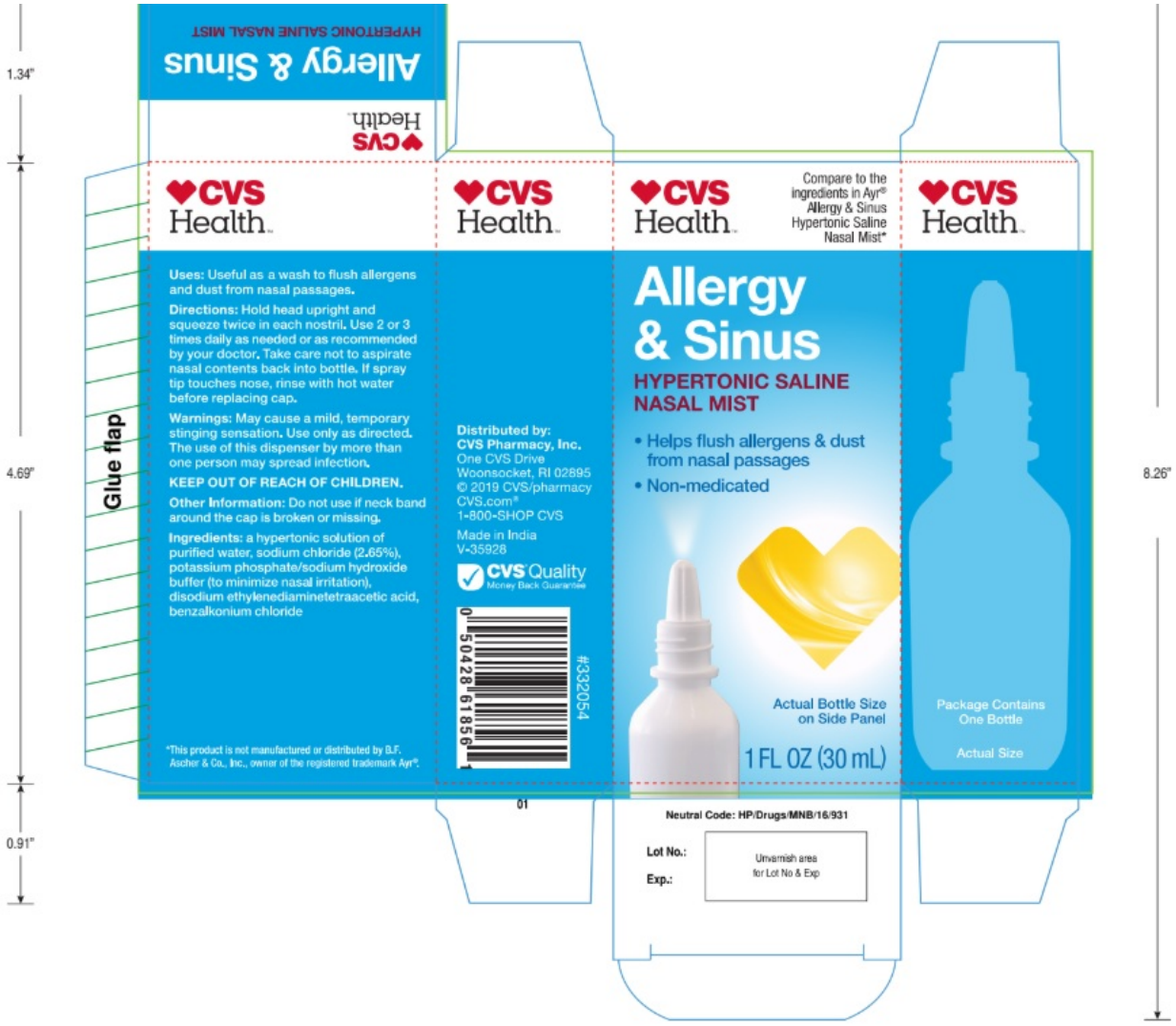
Directions

- Squeeze twice in each nostril as needed
- Upright delivers a spray, horizontally a stream, upside down a drop
- Take care not to aspirate nasal contents back into bottle
- If spray tip touches nose, rinse with hot water before replacing cap

Inactive ingredients

A hypertonic solution of purified water, potassium phosphate/Sodium hydroxide buffer, disodium ethylenediaminetetraacetic acid, benzalkonium chloride

Label



SALINE NASAL 1 FL OZ

sodium chloride 2.65% spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76168-701
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	2.65 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76168-701-30	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/19/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/19/2019	

Labeler - Velocity Pharma LLC (962198409)

Revised: 4/2021

Velocity Pharma LLC