EQUATE PREMIUM SALINE- sodium chloride 0.65% spray
Walmart Stores, 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.

Equate Premium Nasal Saline Spray Sodium Chloride 0.65%

Drug Facts

Active ingredient

Sodium Chloride 0.65%

Purpose

Nasal Moisturizer

Uses

Provides instant, soothing relief to dry irritated nasal passages due to colds, allergies, dry air, pollution, smoke, air travel, and use of decongestants/steroidal sprays.

Warnings

If pregnant or breast feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away. The use of this dispenser by more than one person may spread infection.

Directions

Squeeze bottle twice in each nostril as needed or as directed by physician. **For children 3 years old and younger**, consult a physician before use. **For infants**, use drop applicaiton. Hold bottle upright for spray, horizontally for stream, and upside down for drop.

Other information

store at room temperature

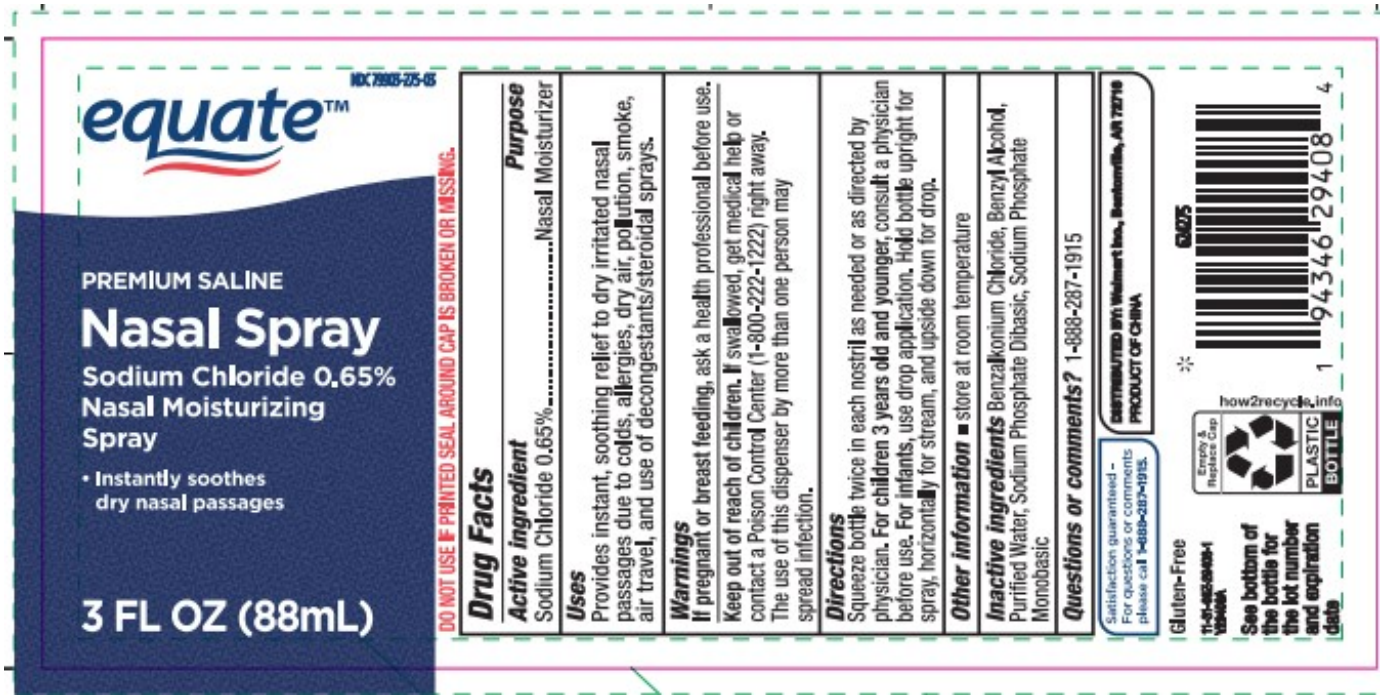
Inactive ingredients

Benzalkonium Chloride, Benzyl Alcohol, Purified Water, Sodium Phosphate Dibasic, Sodium Phosphate Monobasic

Questions or comments?

1-888-287-1915

Principal display panel



EQUATE PREMIUM SALINE

sodium chloride 0.65% spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79903-275
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	0.65 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	

BENZYL ALCOHOL (UNII: LKG8494WBH)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPUU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-275-03	88 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/30/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/30/2024	

Labeler - Walmart Stores, Inc. (051957769)

Revised: 9/2024 Walmart Stores, Inc.