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 application. 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	R MSSING.	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.	Marnings f 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 application. Hold bottle upright for spray, horizontally for stream, and upside down for drop.	e	Inactive ingredients Benzalkonium Chloride, Benzy Alcohol, Purified Water, Sodium Phosphate Dibasic, Sodium Phosphate Monobasic		, Bentomille, AN 72716	s	
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Nasal Spray Sodium Chloride 0.65%	ID CAP		elief t rgies, gesta	askah	f swa (1-80 ore tha	stril as old an p appli nd ups	at roon	alkonit te Dibë	1-886	UTED BY W	*	
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	Drug F	Active in Sodium Ch	Uses Provides ir passages air travel,	Warnings f pregnant (Keep out of reac contact a Poison The use of this di spread infection.	Directions squeeze bott ohysician For pefore use. Fo spray, horizon	ther infe	Inactive i Purified Wat Monobasic	lestions	Satisfaction guaranteed - For questions or commen please cal 1-889-287-1915	Gluten-Free	bottom of bottom of bottine for ot numbe
3 FL OZ (88mL)		A S	Pro Pro air	3±	3 S H S	e seas	9	2 28	9	For		S a a a

EQUATE PREMIUM S sodium chloride 0.65% spray					
Product Information					
Product Type	HUMAN OTC DRUG	ltem Code (Source)	NDC:	79903-275
Route of Administration	NASAL				
Active Ingredient/Active	Moiety				
Ingred	lient Name		Basis of Streng	yth	Strength
SODIUM CHLORIDE (UNII: 451W47 UNII:Q32ZN48698)	/IQ8X) (CHLORIDE ION -		SODIUM CHLORIDE	0	.65 g in 100 mL
Inactive Ingredients					
	Ingredient Name				Strength
SODIUM PHOSPHATE, DIBASIC,	ANHYDROUS (UNII: 22ADO	53M6F)			

R (UNII: 0590 M PHOSPH aging m Code :79903-	ATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPU Package Description 88 mL in 1 BOTTLE, SPRAY; Type 0: Not a	Marketing Start Date	Marketing End Date				
M PHOSPH aging m Code :79903-	ATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPU Package Description 88 mL in 1 BOTTLE, SPRAY; Type 0: Not a	Marketing Start Date					
aging m Code :79903-	Package Description 88 mL in 1 BOTTLE, SPRAY; Type 0: Not a	Marketing Start Date					
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:79903-	88 mL in 1 BOTTLE, SPRAY; Type 0: Not a	Date					
-03	Combination Product	09/30/2024					
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Marketing Information							
arketing ategory	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
oved drug		09/30/2024					
i	arketing ategory	ategory Citation	Arketing ategoryApplication Number or Monograph CitationMarketing Start DateDate				

Labeler - Walmart Stores, Inc. (051957769)

Revised: 9/2024

Walmart Stores, Inc.