FAMILY CARE SALINE- sodium chloride spray
United Exchange Corp.

Active ingredient Purpose

Sodium Chloride 0.65%.....Moisturizer

Use

temporary relief of dry, irritated nasal passages due to colds, flu, allergies, pollution, and the use of decongestants/steroidal sprays

Warnings

For external use only

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.

Directions

- holding bottle upright for spray, horizontally for stream, and upside down for drop applications
- adults and children under 6 years of age: squeeze twice in each nostril as needed
- children under 6 years of age: consult a doctor
- using this dispenser by more than one person may spread infection

Other information

- store at room temperature 20-25°C (68-77°F)
- do not use if printed seal around the cap is broken or missing

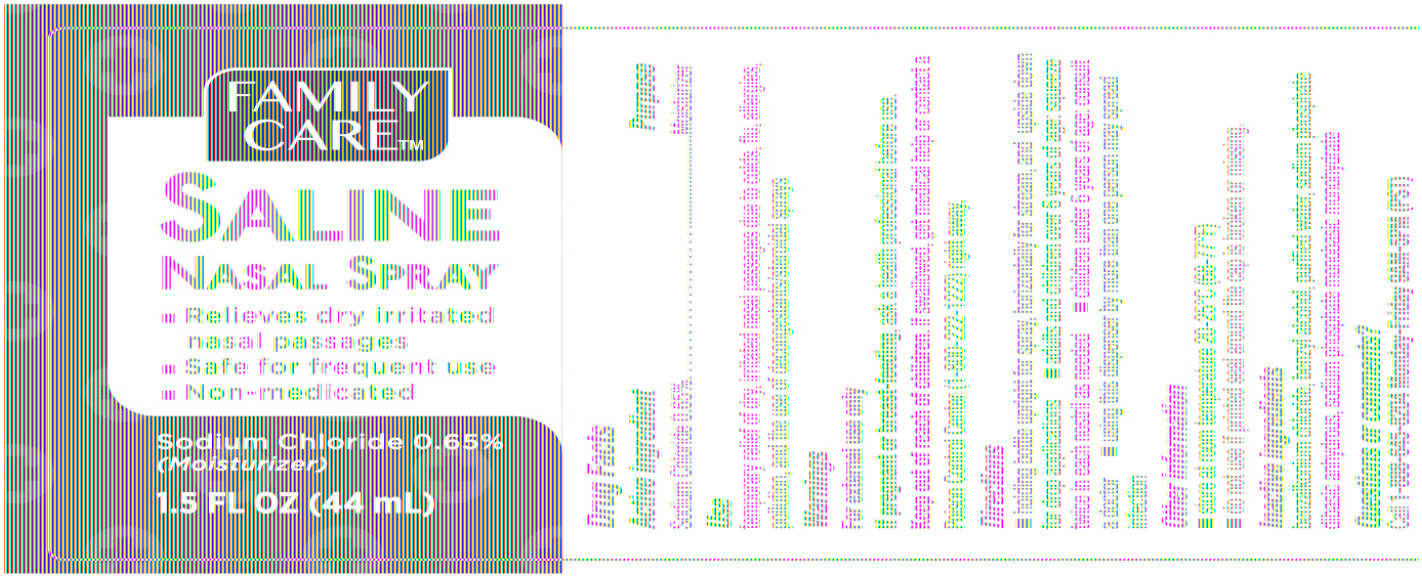
Inactive ingredients

benzalkonium chloride, benzyl alcohol, purified water, sodium phosphate dibasic anhydrous, sodium phosphate monobasic monohydrate

Distributed by: United Exchange Corp.

Cypress, CA 90630 USA

Made in China



FAMILY CARE SALINE

sodium chloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-493
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	6.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM PHOSPHATE DIBASIC DIHYDRATE (UNII: 94255I6E2T)	
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65923-493-44	44 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/28/2023	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M012	09/28/2023	

Labeler - United Exchange Corp. (840130579)

Revised: 2/2024 United Exchange Corp.