

QUALITY CHOICE SALINE- plus soothing aloe spray
QUALITY CHOICE (CHAIN DRUG MARKETING ASSOCIATION)

Quality Choice Saline Nasal Spray

Drug Facts

Active Ingredients

Sodium Chloride 0.65%

Purpose

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 right away at 1-800-222-1222

Directions

For children and adults, squeeze bottle twice in each nostril as often as needed or as directed by physician. **For infants**, use drop application. Hold bottle upright for spray, horizontally for stream, and upside down for drop. The use of this dispenser by more than one person may spread infection.

Other Information

- store at room temperature

Inactive Ingredients

Aloe barbadensis leaf juice (aloe vera gel), benzalkonium chloride, benzyl alcohol, purified water, sodium phosphate dibasic, sodium phosphate monobasic

Questions or comments?

1-866-467-2748

Distributed by: CDMA, Inc.

Novi, MI, 48375

www.qualitychoice.com

Questions: 800-935-2362

Saline Nasal Spray

NDC: 83324-341-30

Saline Nasal Spray

+Soothing Aloe

Sodium Chloride 0.65%

Relief for Stuffy Noses

Gentle Enough for Infants

3 FL OZ (89 mL)

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL AROUND CAP IS BROKEN OR MISSING.

2.375" Height

5" Width

NDC 83324-341-30

QC
QUALITY
CHOICE

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6 35515 99670 0

LOT:

EXP:

**SATISFACTION
100%
GUARANTEE**

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No Varnish in this area
DO NOT PRINT BOX

QUALITY CHOICE SALINE

plus soothing aloe spray

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	
Route of Administration		NASAL	NDC:83324-341	
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
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
BENZYL ALCOHOL (UNII: LKG8494WBH)				
WATER (UNII: 059QF0KO0R)				
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)				
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-341-30	1 in 1 CARTON	07/11/2025	
1		89 mL in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M022	07/11/2025	

Labeler - QUALITY CHOICE (CHAIN DRUG MARKETING ASSOCIATION) (011920774)