SALINE- nasal spray DISCOUNT DRUG MART, INC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Saline Nasal Spray

Active Ingredients

Sodium Chloride 0.65%

Purpose

Moisturizer

Uses

Naturally provides instant, soothing relief 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 healthcare professional before use. Use of this container by more than one person may spread infection.

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

benzalkonium chloride, benzyl alcohol, purified water, sodium phosphate dibasic, sodium

Questions or comments?

1-866-467-2748

*This product is not manufactured or distributed by Valeant Pharmaceuticals North America LLC, sodium phosphate monobasic.

Saline Nasal Spray

Compare to the active ingredient in Ocean®*

Saline Nasal Spray

Sodium Chloride 0.65%

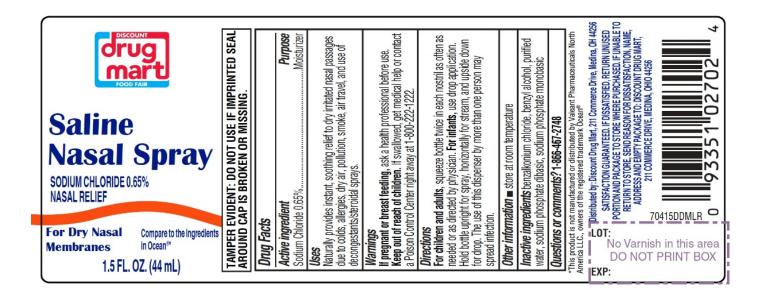
- Instantly relieves dry nasal passages caused by sinus, cold and allergy medications and dry air
- Safe for frequent daily use
- Gentle enough for infants
- Natural, non-medicated relief for stuffy noses

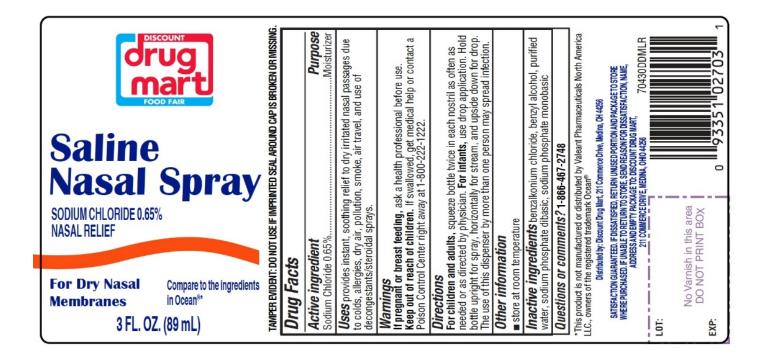
100% SATISFACTION GUARANTEED OR YOUR MONEY BACK

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

Distributed by:

Package label for 44 mL





SALINE

nasal spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53943-704
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		
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)			

Pa	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53943-704- 15	1 in 1 CARTON	09/30/2020	
,		44 mL in 1 BOTTLE; Type 0: Not a Combination		

1		Product		
2	NDC:53943-704- 30	1 in 1 CARTON	09/30/2020	
2		88 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 final	part349	09/30/2020	
o re monegraph mar	parts is	03/30/2020	

Labeler - DISCOUNT DRUG MART, INC. (047741335)

Revised: 1/2022 DISCOUNT DRUG MART, INC.