

**SALINE NASAL 1.5OZ- sodium chloride 0.65% spray**  
**Velocity Pharma LLC**

*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.*

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**Saline Nasal Spray**

***Active ingredient***

Sodium Chloride, 0.65%

**☐ *Purpose***

Moisturizer

**☐ *Uses***

For dry nasal membranes

**☐ *Warnings***

Do not use if seal is broken or missing.

**Keep out of reach of children.** The use of this dispenser by more than one person may spread infection.

**☐ *Directions***

- Squeeze twice in each nostril as needed
- Upright delivers a spray, horizontally a stream, upside down a drop

**☐ *Inactive ingredients***

Benzalkonium chloride, Disodium phosphate, Phenylcarbinol, Monosodium phosphate, Water

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## SALINE NASAL 1.5OZ

sodium chloride 0.65% spray

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76 168-602
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)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPUU)	
WATER (UNII: 059QF0K00R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76 168-602-	44 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination	11/19/2019	

15	Product	11/19/2019	
<b>Marketing Information</b>			
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph final	part349	11/19/2019	

**Labeler** - Velocity Pharma LLC (962198409)

Revised: 11/2019

Velocity Pharma LLC