SALINE NASAL MOISTURIZING WALGREENS- sodium chloride gel
WALGREENS COMPANY

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.

Walgreens Saline Nasal Gel

Active Ingredient

Sodium Chloride 0.9%

Sterile, Hypertonic Saline Solution,

Sodium Bicarbonate to adjust pH

Purpose

Nasal moisturizer

Uses

provides moisture to soothe and hydrate dry nasal passages caused by:

- indoor heat
- dry climate
- air travel
- high altitude
- oxygen use
- CPAP machine use

Directions

Adults and children 6 years and over: Use 1-2 sprays in each nostril up to every 2 hours as needed

Children under 6 years- Consult a physician

Warnings

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Stop use and ask a doctor if use is uncomfortable or dryness persists

Do not use if allergic to any of the ingredients

Do not use if safety seal is missing or broken

For nasal use only, do not use for dry mouth

Keep out of reach of chlidren.

Other Information

Store in cool dry place and protect from freezing

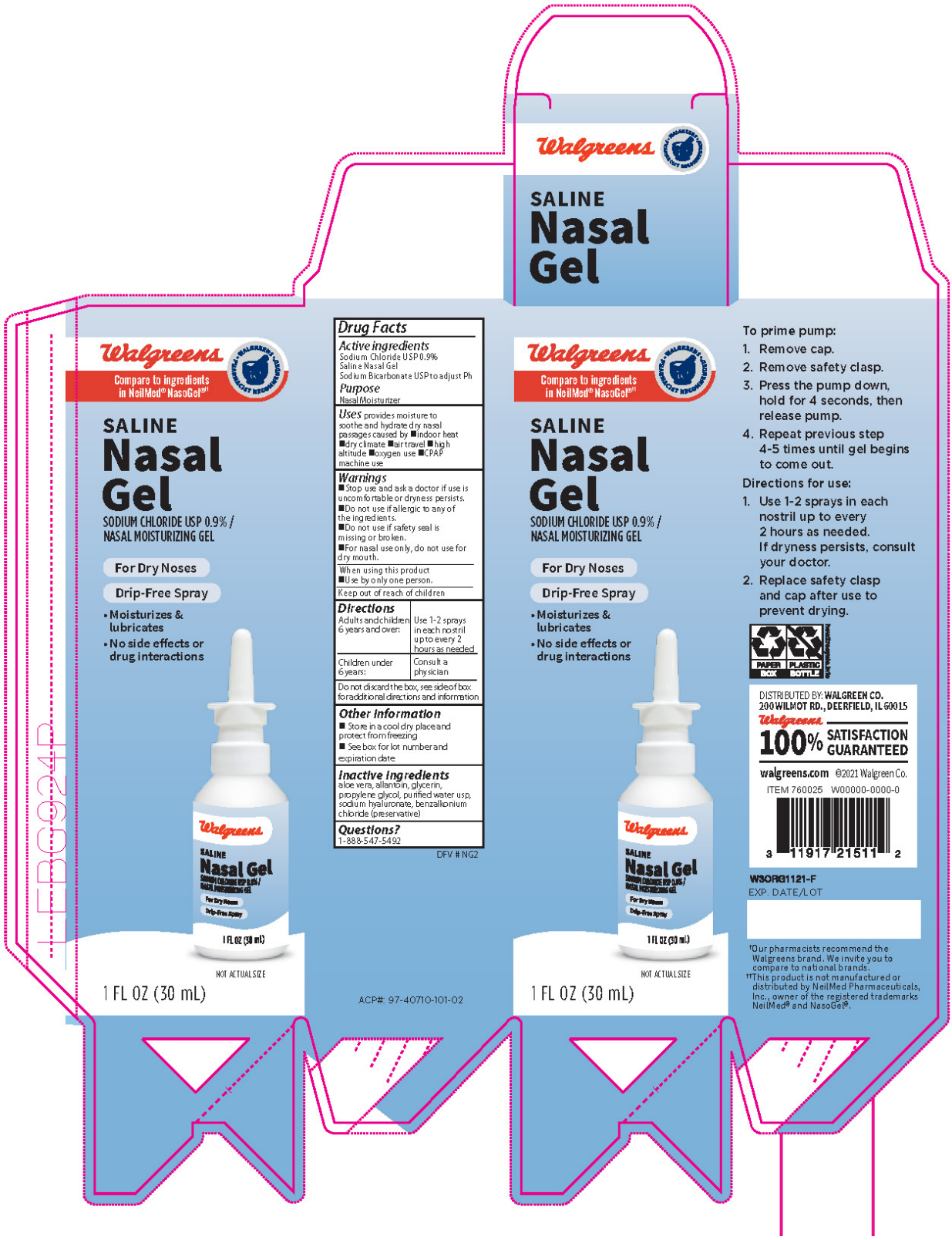
Inactive ingredients

Purfied Water, allantoin, glycerin, propylene glycol, sodium hyaluronate, benzalkonium chloride as preservative

QUESTIONS ?

1-888-547-5492

Saline Nasal Spray product label



SALINE NASAL MOISTURIZING WALGREENS

sodium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-5557
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	0.9 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
WATER (UNII: 059QF0KO0R)	
ALLANTOIN (UNII: 344S277G0Z)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0363-5557-30	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/13/2022	

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
unapproved drug other		01/13/2022	

Labeler - WALGREENS COMPANY (008965063)

Revised: 1/2022 WALGREENS COMPANY