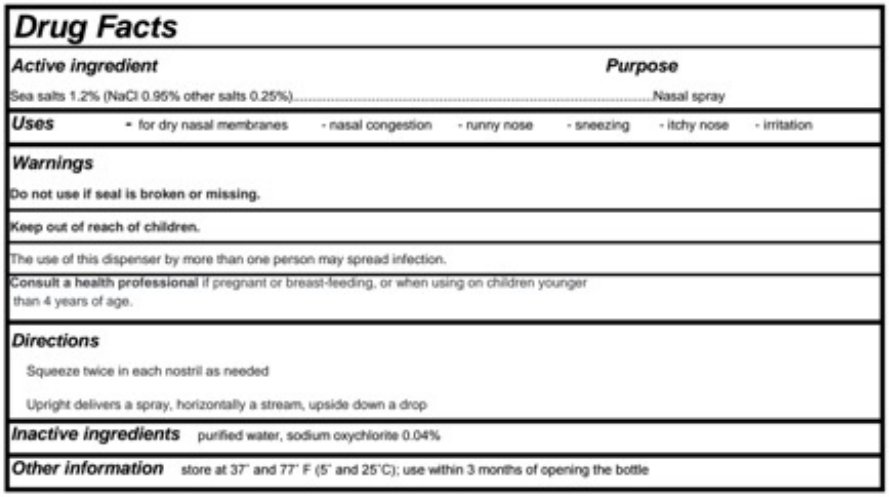
NASOXINSPRAY- sodium chloride spray spray
FRIANA LLC

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.

Nasoxin Nasal Spray



Drug Facts

Active ingredient — **Purpose**

Sea salts 1.2% (NaCl 0.95% other salts 0.25%) ..... Nasal spray

Uses • for dry nasal membranes • nasal congestion • runny nose • sneezing • itchy nose • irritation

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.

Consult a health professional if pregnant or breast-feeding, or when using on children younger than 4 years of age.

Directions

Squeeze twice in each nostril as needed

Upright delivers a spray, horizontally a stream, upside down a drop

Inactive ingredients purified water, sodium oxychlorite 0.04%

Other information store at 37° and 77° F (5° and 25°C); use within 3 months of opening the bottle

Sodium Chloride (NaCl 0.95%, Sea salts 1.2%, other salts 0.25%)

Keep out of reach of children.

Moisturizes nasal passages

for dry nasal membranes
nasal congestion
runny nose
sneezing
itchy nose
irritation

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The use of this dispenser by more than one person may spread infection.
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Purified water
Sodium Oxychlorite 0.04%

NASOXINSPRAY

sodium chloride spray spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70508-1601
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM CHLORIDE	1.2 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM HYPOCHLORITE (UNII: DY38VHM5OD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70508-1601-1	15 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/04/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	04/04/2016	

Labeler - FRIANA LLC (079770345)

Registrant - FRIANA LLC (079770345)

Establishment

Name	Address	ID/FEI	Business Operations
FRIANA LLC		079770345	manufacture(70508-1601)

Revised: 4/2016 FRIANA LLC