SALISPRAY DAILY SALINE- nasal mist aerosol, spray PONTIKA AEROTECH LIMITED

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.

Active ingredient

Sodium Chloride 0.9%

Purpose

Nasal Moisturizer

Uses

- Restores vital moisture to provide prompt relief for dry, crusted, and inflamed nasal membranes due to chronic sinusitis, colds, low humidity, overuse of nasal decongestant drops and sprays, allergies, minor nose bleeds, and other minor nasal irritations.
- Helps remove inhaled irritants such as dust, dirt, and pollen.

Warnings

- This product may cause temporary discomfort such as burning, stinging, sneezing, or an increase in nasal discharge.
- The use of this dispenser by more than one person may spread infection.
- Avoid spraying in eyes.
- Content under pressure.
- Do not puncture or incinerate.
- Store between 59º and 86º F (15º and 30º C)

Keep out of reach of children

Directions

Read HOW TO USE instructions before use

- Adults and children 4 years of age an older: Use as often as needed or as
 directed by a physician
- Children under 4 years: Consult a physician.

Other information

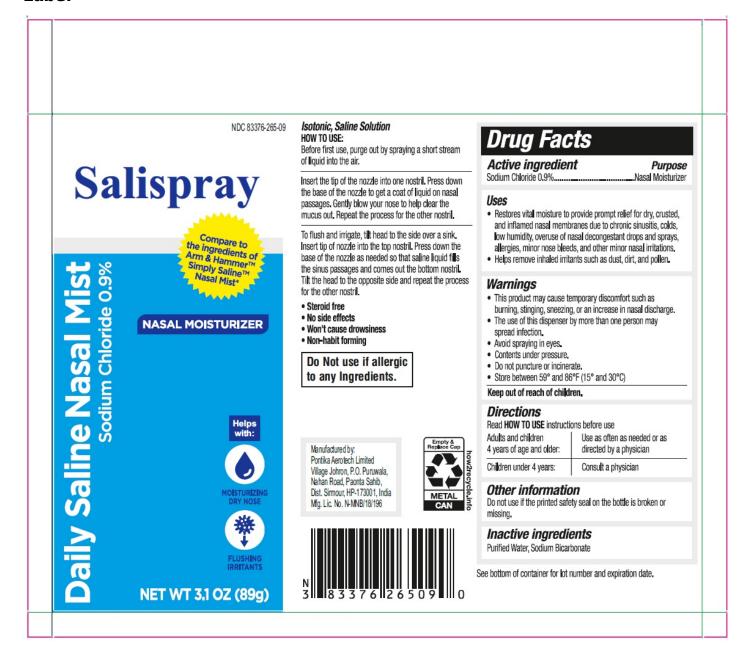
Do not use if the printed safety seal on the bottle is broken or missing.

Inactive ingredients

Purified Water, Sodium Bicarbonate

PRINCIPAL DISPLAY PANEL

Daily Saline Nasal Mist Sodium Chloride 0.9% - NDC 83376-265-09 - Bottle Label



SALISPRAY DAILY SALINE

nasal mist aerosol, spray

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83376-265	
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

WATER (UNII: 059QF0KO0R)

SODIUM BICARBONATE (UNII: 8MDF5V39QO)

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# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:83376-265-09	89 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/05/2024	

Marketing Information

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		09/05/2024		

Labeler - PONTIKA AEROTECH LIMITED (861299010)

Establishment					
Name	Address	ID/FEI	Business Operations		
PONTIKA AFROTECH LIMITED		861200010	manufacture(83376-265) analysis (83376-265)		

Revised: 9/2024 PONTIKA AEROTECH LIMITED