

NASAL SPA NATURAL SEA SALT- sodium chloride spray
NACUR HEALTHCARE LTD

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.

Drug Facts

Active Ingredient:

Natural Sea Salt 9.5 mg/mL

Purpose

Nasal Moisturizer

Uses

- For cleansing and moisturizing the nose
- Helps dry nasal congestion caused by allergens, irritants, colds, occupational and environmental pollutants

Warnings

- Do not share between users -Do not use after the expiry date
- Use within 6 months of opening

Keep out of reach of children

Directions

- Spray 2 - 3 times into each nostril and inhale through your nose
- For best results, use as often as needed

Other Information

- Store below 25C (77F)
- Suitable for children and adults

Inactive Ingredient

Purified water

For expiry date see top lid.

Use Nasal Spa Nasal Spray

to help:

- nasal congestion
- dry nose
- runny clear mucus
- rinse out allergens and pollutants
- remove excess mucus
- dripping sensation in the back of the throat

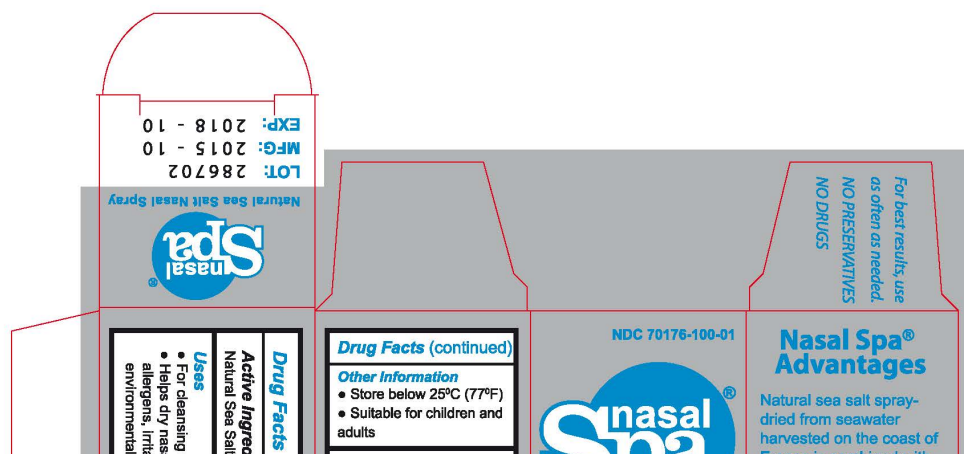
The patented applicator of

the Nasal Spa Nasal Spray contains an highly effective valve and an **oligo-dynamically active silver coil** which protects the solution and the dosage applied against microbial congestion.

The sea salt is manufactured in a highly technical process (spray-drying) harvesting seawater taken from an archipelago off the coast of France in Europe.

Compared to standard sodium chloride solutions, the gentle, isotonic natural sea salt solution (0.95%) of the Nasal Spa Spray is rich in minerals and trace elements.

MADE IN GERMANY





NASAL SPA NATURAL SEA SALT

sodium chloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70176-100
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8 X) (SODIUM CATION - UNII:L YR4M0NH37)	SODIUM CHLORIDE	9.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70 176-100-01	1 in 1 CARTON	10/29/2015	
1		20 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/29/2015	

Labeler - NACUR HEALTHCARE LTD (346989317)

Revised: 4/2016

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