

SALINE AMPOULES- sodium chloride 0.9% spray
Trifecta Pharmaceuticals USA, LLC.

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.

Enjoy Good Health
Pure and Sterile
0.9% Sodium Chloride Isotonic
5ml Single Use Saline Ampoules

Active Ingredient

Sodium Chloride 0.9%

Purpose

Moisturizer

This product may be used for conditions in the respiratory passages, nasal cavity or as directed by your healthcare provider.

Warnings

For External Use Only

Not for parenteral administration

When using this product

When using this product a mild and temporary irritation may be experienced at the beginning of use.

Keep out of the reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

If pregnant or breast feeding

Ask a doctor before use.

Do Not Use

- Same applicator on more than one individual
- If sensitive to any of the components

Directions

For Adults and children 4 years and older. Use as often as needed.

Children under 4 years of age consult a doctor.

Hold vial gently and twist top off

Squeeze vial to dispense

Discard after use

Other information

- Store at controlled room temperature 20°-25°C (68°-77°F) and protect from sunlight and sources of heat.
- See bottom of carton for Lot No. and Expiration data
- No Bacteriostatic or other preservatives added

Inactive Ingredients

Purified Water

Questions or Comments

Call 1-888-296-9067

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Enjoy Good Health

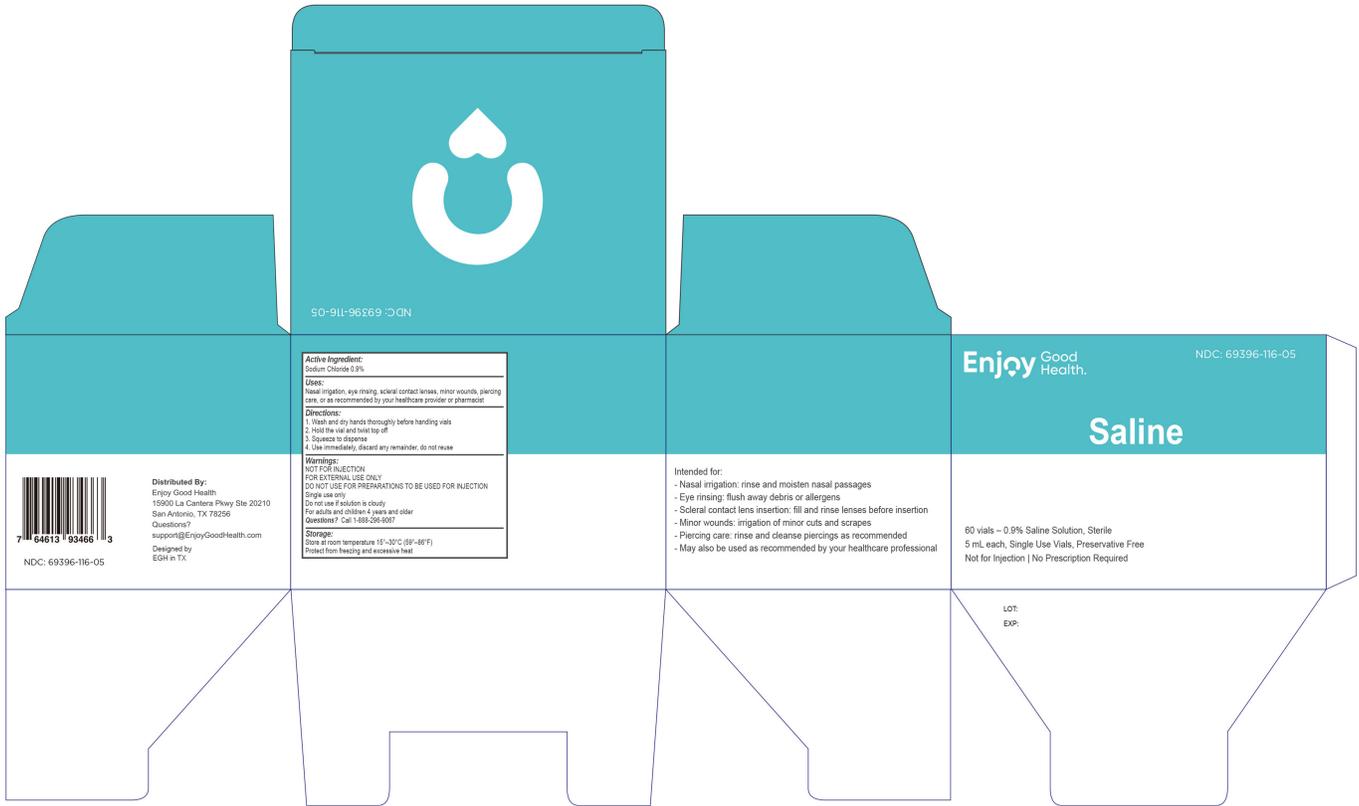
15900 La Cantera Pkwy Ste 20210

San Antonio, TX 78256

Questions: support@enjoygoodhealth.com

Label

BOX



LABEL



SALINE AMPOULES

sodium chloride 0.9% spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69396-116
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	9 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:69396-116-05	60 in 1 CARTON	11/22/2022	
1		5 mL in 1 VIAL; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved drug other			07/28/2022	

Labeler - Trifecta Pharmaceuticals USA, LLC. (079424163)

Revised: 10/2025

Trifecta Pharmaceuticals USA, LLC.