SINUCLEANSE NASAL DRIP- sodium chloride gel
ASCENT CONSUMER PRODUCTS, INC

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.

SinuCleanse Breathe Moisturizing Nasal Gel - Drip Free Spray

Active Ingredient

Sodium Chloride 0.9%

Saline Nasal Gel,

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 machineuse

Directions

Adults and children 4 years and over: Use as often as needed
Children under 4 years: Consult a physician
See directions on can for complere instructions

For nasal use only.

Before use, expel a shoet stream of mist into the air. Insert tip of nozzle into one nostril and press down on the textured area at the base of the nozzle so that a gentle mist coats nasal passages. Blow your nose very gently to clear the mucus out. Repeat for the other nostriil.

To flush and irrigate, tilt head to the side over sink. Insert top of nozzle into the top nostril, pressing down on the textured area at the base of the nozzle so that a gentle mist fills sinus passages and flows out the opposite nostril. Repeat in other nostril.

Wipe nozzle after each use.

Warnings

Warnings

-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 the 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 freezing

-See box for lot number and expiration date

Inactive ingredients

aloe vera, allantoin, glycerin, propylene glycol, purified water USP, sodium hyaluronate, benzalkonium chloride

QUESTIONS ?

1-888-547-5492

Product label

SINUCLEANSE NASAL DRIP

sodium chloride gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42829-407
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 mg in 100 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42829-407-10	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/08/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/08/2022	

Labeler - ASCENT CONSUMER PRODUCTS, INC (078396381)

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