SINUS WASH PRE MIXED READY TO USE RITE AID- sodium chloride, sodium bicarbonate
RITE AID CORPORATION

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.

Rite Aid Sinus Wash Pre-Mixed Ready to Use

Active Ingredients

(in each prefilled bottle)

Sodium Bicarbonate USP 700 mg

Sodium Chloride USP 2300 mg

Active ingredients

(in each packet)

Sodium Bicarbonate USP 700 mg

Sodium Chloride USP 2300 mg

Purpose

Nasal Wash

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Nasal Wash

Uses

Temporarily relieves symptoms associated with

sinusitis, cold, flu or allergies:

-Sneezing

-Runny nose

-Nasal stuffiness

-Post nasal drip

-Removes inhaled irritants (dust, pollen)

-Promotes nasal and sinus drainage

-Helps reduce swelling of nasal membranes

-Moisturizes dry nasal passages

Warnings

Stop use and ask a doctor if washing is uncomfortable or

symptoms are not reduced.

Do not use unfiltered tap water. See instructions inside box

for proper water sources.

Do not use if nasal passages are completely blocked or if you

have an ear infection or blocked ears.

When using this product:

- Use by only one person
- Wash with soap and water after each use
- Do not heat in microwave

Keep out of reach of children

Directions:

Adults and children 4 years and over: use 1 - 2 packets per 8 fl. oz (240 mL) up to every 2 hours as needed

Children under 4 years: Consult a physician

Prior to first use see enclosed instruction sheet for complete directions

Other information

- Inspect saline packets for integrity
- Do not use saline packets if open or torn
- Protect saline packets from excessive heat and moisture
- See saline packets or box for lot no. and expiration date

Inactive ingredients

(in each prefilled bottle)

Purified Water USP, Benzalkonium Chloride USP (Preservative)

Inactive ingredients

(in each packet)

None

Questions?

1-888-547-5492

SINUS WASH PRE MIXED READY TO USE RITE AID

sodium chloride, sodium bicarbonate kit

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11822-0472

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0472-8	1 in 1 KIT; Type 1: Convenience Kit of Co-Package	01/16/2022	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 PACKET	3000 mg
Part 2	1 BOTTLE	240 mg

Part 1 of 2

SINUS WASH PACKET

sodium bicarbonate, sodium chloride powder

Product Information

Route of Administration	NASAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	2300 mg in 3000 mg
SODIUM BICARBONATE (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM BICARBONATE	700 mg in 3000 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		3 in 1 KIT		
1		3000 mg in 1 PACKET; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/16/2019	

Part 2 of 2

SINUS WASH PREFILLED BOTTLE

sodium bicarbonate, sodium chloride solution

Product Information

Route of Administration	NASAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM BICARBONATE	700 mg in 3000 mg
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	2300 mg in 3000 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		1 in 1 KIT		
1		240 mg in 1 BOTTLE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/16/2022	

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/16/2019	

Labeler - RITE AID CORPORATION (014578892)

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