## SINUCLEANSE- sodium bicarbonate, sodium chloride powder, for solution 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.

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## SinuCleanse

**Active Ingredients** (in each packet) Sodium Bicarbonate USP 700 mg Sodium Chloride USP 2300 mg Purpose 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 relieved. 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

-Top rack of dishwasher safe

-Do not heat in microwave

Keep out of reach of children

Directions

Adults and children 4 years and over: use 1/2 - 1 packet per 4 fl oz (120 mL)up to every two hours as needed

Children under 4 years: Consult a physician

See enclosed instruction sheet for complete directions and proper use

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 # and expiration date

Inactive Ingredients

None

Questions?

1-888-547-5492



sodium bicarbonate, sodium chloride powder, for solution

Product Information									
Product Type	HUMAN OTC DRUG	Item Code (Source)	1	NDC:42	829-408				
Route of Administration	NASAL								
Active Ingredient/Active Moiety									
Active myreulent/Active molecy									
Ingredient Name			Basis of Strength		Strength				
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4M0NH37, BICARBONATE ION - UNII:HN1ZRA3Q20)			SODIUM BICARBOI		22.2 mg in 100 mg				
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37)			SODIUM CHLORIDE	E	77.8 mg in 100 mg				

In	active Ingre	edients					
Ingredient Name			Strength				
WATER (UNII: 059QF0KO0R)							
Packaging							
#	ltem Code	Package Description	Marketing Start Date		Marketing End Date		
1	NDC:42829- 408-60	60 in 1 CARTON	02/01/2015				
1		3000 mg in 1 PACKET; Type 0: Not a Combination Product					
Marketing Information							
	Marketing Category	Application Number or Monograph Citation	Marketi Da	ng Start Ite	Marketing End Date		
	approved drug ner		02/01/2015				

Labeler - ASCENT CONSUMER PRODUCTS, INC. (078396381)

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ASCENT CONSUMER PRODUCTS, INC.