NASAL SOLUTION- nasal solution liquid
Bullet Point LLC

Nasal Solution

Povidone-lodine 1%

Purifed water,xylitol, vitamin D3,polysorbate, gellangum, carrageenan, sodiumhydroxide, aloe vera.

*Gently washes and soothesthe delicate nasal passage.
*Helps clean away viruses,pollen, dust, and otherairborne contaminantsand germs.
Thins and loosens mucussecretion.
*Aids in improving airflow.
*Powerful sinus protection.
*Fast congestion relief.

Uses:
*Congested nose andsinuses.
*lrritated and itchy nose.
*Nasal exposure to bacteria,dust and other airbornecontaminants.

Avoid use if you're allergicto iodine or inactive ingredient(s).Avoid contactin the eyes.
Avoid use onchildren under 3 years old.

Stop use and consult adoctor if symptoms persistor worsen, swelling,infection, rash or feveroccurs.lf pregnant orbreast- feeding, ask a health professional before use.

Keep out of reach ofchildren. lf swallowed, getmedical help or contact the poison control center rightaway.

lf pregnant orbreast- feeding, ask a health professional before use.

Use of container bymore than one person canspread infection.

Directions:
Adults and children over 6years of age: 1 spray in eachnostril not more than 3times in any 24-hour period,Children under é years ofage: Consult a doctor.
For the frst time -Removesafety seal and original capfrom bottle. insert dip tubeinto bottle, and screw on.
Gently blow your nose toclear the nasal passageway.Shake the product well,remove the cap, and thenspray it quickly and frmlyup your nostril.
Do not tilt your headbackward during sprayingAfter use, wipe the spraynozzle clean and securelyreattach the cap.

You can email us at support@noveha.com.com for more help!

NASAL SOLUTION

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84008-101
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
XYLITOL (UNII: VCQ006KQ1E)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
WATER (UNII: 059QF0KO0R)	
GELLAN GUM (LOW ACYL) (UNII: 7593U09I4D)	
CARRAGEENAN (UNII: 5C69YCD2YJ)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84008-101-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/20/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/20/2024	

Labeler - Bullet Point LLC (118190100)

Revised: 7/2024 Bullet Point LLC