NASAL WASH SALINE- sodium chloride liquid Shenzhen Jun&Liang Media Tech Limited

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.

Active ingredient(in each dosage unit): Sodium chloride

Purpose: Clean the nasal cavity

Uses: Work with Gailyus's nasal washer

Warnings: Do not use with illegal drugs, alcohol, or non-prescribed medications

Ask a doctor before use:

If you have an ear infection or if your ears are blocked.

If your nasal passages are completely blocked.

Ask a doctor or pharmacist before use: if you recently had ear or nose surgery.

When using this product:

Do not use regular tap water. It recommends working with distilled, sterile or previously boiled water.

Stop use and ask a doctor if:

Your symptoms persist or worsen; or your nasal illness doesn't improve after rinsing.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Water

Directions:

Place a new, unused saline capsule in the Gailyus nasal washer's crushing chamber, and keep the foil-side down.

The optimal water temperature is around 35°C-40°C (95°F-104°F) for rinsing nasal.

Questions?You can email us at Gailyus@outlook.com for more help!

Place a new, unused saline capsule in the Gailyus nasal washer's crushing chamber, and keep the foil-side down.

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Drug Facts

Manufacturer: Guangdong XIN RUN TAO Technology Co., Ltd.

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label

NASAL WASH SALINE

sodium chloride liquid

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83660-001
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Route of Administration NASAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32Z N48698)	SODIUM CHLORIDE	245 mg in 9.5 mL

nts

Ingredient Name	Strength			
WATER (LINII) OFOCEOVOOR)	7 1725 ml in 0.5 ml			

WATER (UNII: 059QF0KO0R) 7.1725 mL in 9.5 mL

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:83660- 001-12	30 in 1 BOX	07/28/2023		
1	NDC:83660- 001-11	9.5 mL in 1 CAPSULE; Type 0: Not a Combination Product			
2	NDC:83660- 001-22	60 in 1 BOX	07/28/2023		
2	NDC:83660- 001-21	9.5 mL in 1 CAPSULE; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		07/28/2023		

Labeler - Shenzhen Jun&Liang Media Tech Limited (444563661)

Registrant - Guangdong XINRUNTAO Technology Co., Ltd. (403789850)

Establishment				
Name	Address	ID/FEI	Business Operations	
Guangdong XINRUNTAO Technology Co., Ltd.		403789850	manufacture(83660-001) , label(83660-001)	

Revised: 8/2023 Shenzhen Jun&Liang Media Tech Limited