NATRUM MURIATICUM- natrum muriaticum spray
Ratis, LLC

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Drug Facts:

ACTIVE INGREDIENT:

Natrum Muriaticum 6X.

USES:

Helps relieve conditions with excess water or dryness such as runny or stuffy nose, allergies, headaches, migraines, constipation, slow digestion, watery eyes, and dry mouth.**

**These statements are based upon traditional homeopathic practice. They have not been reviewed by the Food and Drug Administration.

WARNINGS:

For oral use only.

If pregnant or breast-feeding, or if symptoms persist or worsen, ask a health care professional.

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

Do not use if tamper evident seal is broken or missing.

KEEP OUT OF REACH OF CHILDREN:

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

DIRECTIONS:

Adults: 2 sprays 3 times a day or as needed.

Children 2-12: 1 spray as above. For children 12 and under, consult a doctor.

INDICATIONS:

Helps relieve conditions with excess water or dryness such as runny or stuffy nose, allergies, headaches, migraines, constipation, slow digestion, watery eyes, and dry

mouth.**

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INACTIVE INGREDIENTS:

Demineralized Water, Organic Ethanol 20%

QUESTIONS:

Comments? Visit

HomeopathyStore.com

or call (888) 405-7551.

Distributed by:

Ratis, LLC,

211 E. Lombard St, STE 303,

Baltimore, MD 21202

PACKAGE LABEL DISPLAY:

NDC 71753-8009-1

LACTOSE FREE

Anna's

REMEDIES

Natrum Muriaticum

HOMEOPATHIC ORAL SPRAY

1 FL. OZ (30ML)

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LOT: XXXXXX

NATRUM MURIATICUM

natrum muriaticum spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71753-8009
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	6 [hp_X] in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71753-8009-1	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/30/2020	10/26/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved			

unapproved homeopathic		09/30/2020	10/26/2025

Labeler - Ratis, LLC (964594324)

Revised: 1/2022 Ratis, LLC