

NATRUM SULPHURICUM- natrum sulphuricum 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 Sulphuricum 6X.

USES:

Helps relieve nausea, vomiting, bitter taste, diarrhea, and influenza. Eliminates toxins and supports liver health.**

**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 nausea, vomiting, bitter taste, diarrhea, and influenza. Eliminates toxins and supports liver health.**

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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 71753-8011-1

LACTOSE FREE

Anna's

REMEDIES

NATRUM SULPHURICUM

HOMEOPATHIC ORAL SPRAY

1 FL. OZ (30ML)

Active Ingredients:
Natrium Sulphuricum 6X

Uses:
Helps relieve nausea, vomiting, bitter taste, diarrhea, and influenza. Eliminates toxins and supports liver health.**

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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.

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LOT: XXXXXX



NATRUM SULPHURICUM

natrium sulphuricum spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71753-8011
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM SULFATE (UNII: 0YPR65R21J) (SODIUM SULFATE ANHYDROUS - UNII:36KCS0R750)	SODIUM SULFATE	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:71753-8011-1	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/30/2020	10/26/2025

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
unapproved homeopathic		09/30/2020	10/26/2025

Labeler - Ratis, LLC (964594324)

Revised: 1/2022

Ratis, LLC