KALI MURIATICUM- kali 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:

Kai Muriaticum 6X.

USES:

Helps relieve colds and coughs with white discharges, runny nose, tonsillitis, & skin burns.**

**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 colds and coughs with white discharges, runny nose, tonsillitis, & skin burns.**

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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 71753-8005-1

LACTOSE FREE

Anna's

REMEDIES

Kali Muriaticum

HOMEOPATHIC ORAL SPRAY

1 FL. OZ (30ML)

Active Ingredients:
Kali Muriaticum 6X

Uses:
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LOT: XXXXXX

KALI MURIATICUM

kali muriaticum spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71753-8005
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POTASSIUM CHLORIDE (UNII: 660YQ98I10) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM CATION	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:71753-8005-1	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/01/2020	10/27/2025

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
unapproved homeopathic		10/01/2020	10/27/2025

Labeler - Ratis, LLC (964594324)

Revised: 1/2022 Ratis, LLC