KALI SULPH- potassium sulfate tablet
Hyland's Inc.

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.

#7 KALI SULPH. 6X 500 Tablets

DIRECTIONS

Adults & Children ages 6 - 12: 4 tablets. **Children ages 2 - 6:** 2 tablets. Dissolve under tongue 3 times a day. Use more frequently (every 15 minutes for up to 8 doses) with acute conditions.

INDICATIONS

Relief of symptoms of the common cold. Relief of symptoms of pimples and blemishes associated with common acne.

FORMULA

Kali Sulphuricum 6X HPUS

In a base of Acacia Gum, Lactose N.F.

Warnings

Do not use if imprinted cap band is broken or missing.

If symptoms persist for more than seven days or worsen, contact a licensed health care provider.

Discontinue use if symptoms are accompanied by a high fever (over 101° F).

If you are pregnant or nursing, seek the advice of a licensed health care provider before using this product.

Keep this and all medications out of the reach of children.

QUESTIONS?

800.624.9659

PRINCIPAL DISPLAY PANEL - 500 Tablet Bottle Label

HOMEOPATHIC

NDC 54973-5230-01

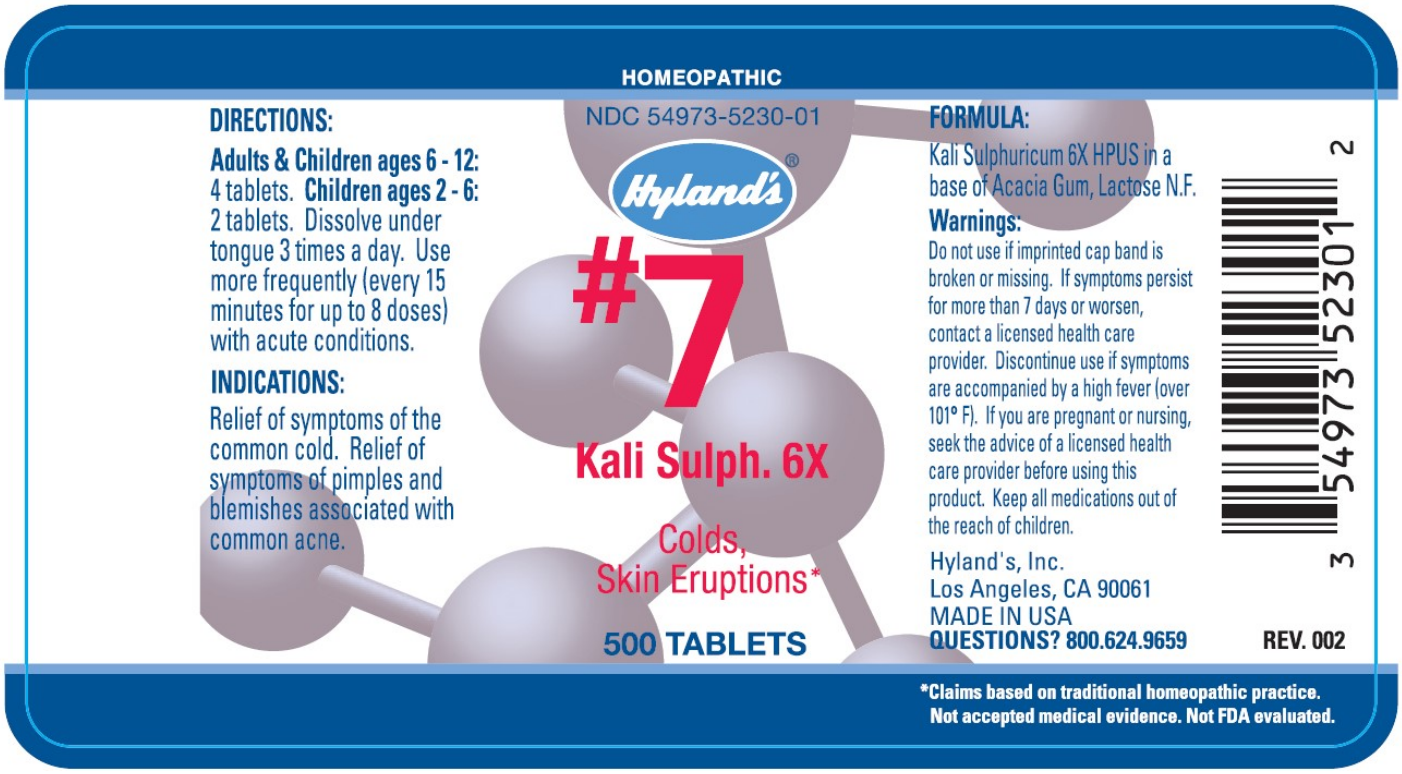
Hyland's®

#7

Kali Sulph. 6X

Colds,
Skin Eruptions

500 TABLETS



KALI SULPH

potassium sulfate tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54973-5230
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POTASSIUM SULFATE (UNII: 1K573LC5TV) (SULFATE ION - UNII:7IS9N8KPMG)	POTASSIUM SULFATE	6 [hp_X]

Inactive Ingredients	
Ingredient Name	**Strength**
ACACIA (UNII: 5C5403N26O)	
LACTOSE (UNII: J2B2A4N98G)	

Product Characteristics			
Color	white	**Score**	no score
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54973-5230-1	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1940	
2	NDC:54973-5230-2	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1940	12/28/2021

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/01/1940	

Labeler - Hyland's Inc. (008316655)

Establishment

Name	Address	ID/FEI	Business Operations
Hyland's Inc.		008316655	manufacture(54973-5230) , pack(54973-5230) , label(54973-5230)

Revised: 12/2022 Hyland's Inc.