NAT SULPH- sodium sulfate anhydrous 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.

NAT. SULPH.

DIRECTIONS

Adults & Children ages 7 - 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 flu,nausea and vomiting.

FORMULA

Natrum Sulphuricum 6X HPUS

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

“HPUS” indicates that the active ingredients are in the official Homeopathic Pharmacopœia of the United States.

Warnings

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

If symptoms persist for more than 7 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 all medications out of the reach of children

QUESTIONS?

(800) 624-9659

PRINCIPAL DISPLAY PANEL - 500 Tablet Bottle Label

HOMEOPATHIC

NDC 54973-6440-1

***Hyland's*®**

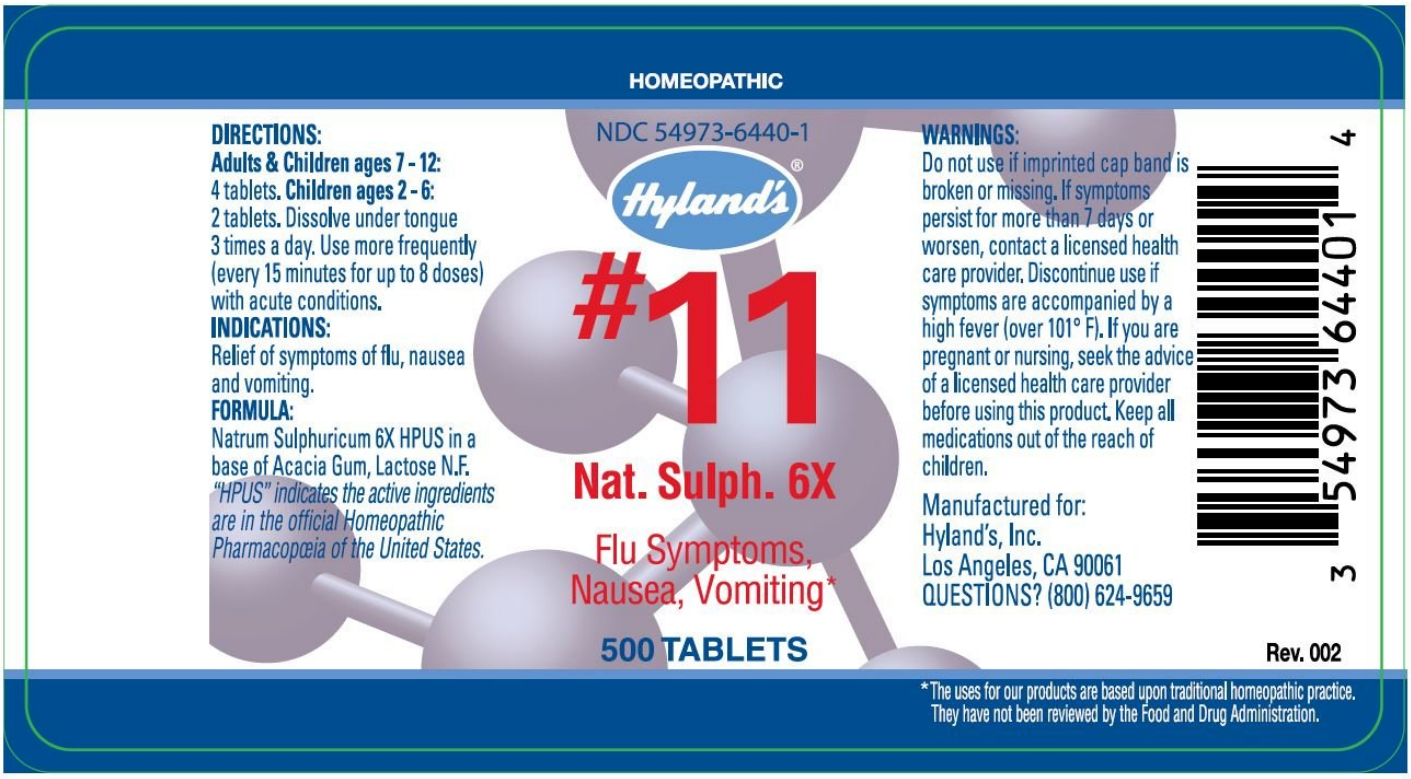
#11

Nat. Sulph. 6X

Flu Symptoms,

Nausea,Vomiting*

500 TABLETS



NAT SULPH

sodium sulfate anhydrous tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54973-6440
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM SULFATE ANHYDROUS (UNII: 36KCS0R750) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM SULFATE ANHYDROUS	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:54973-6440-1	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1940	
2	NDC:54973-6440-2	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1940	

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(54973-6440) , pack(54973-6440) , label(54973-6440)

Revised: 12/2022 Hyland's Inc.