## NATRUM SULPHURICUM- natrum sulphuricum pellet HOMEOLAB USA 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.

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#### **HOMEOPATHIC MEDICINE NDC 60512-6350-1**

#### **ACTIVE INGREDIENT HPUS**

NATRUM SULPHURICUM 1X

(Sodium Sulfate)

BRONCHIAL IRRITATION MADE WORSE BY HUMIDITY

#### **USE**

For self-limiting condition listed above or as directed by a health professional.

#### **WARNINGS**

**Do not use** if pellet-dispenser seal is broken.

**Stop use and ask a doctor if** symptoms persist more than 3 days or worsen.

**If pregnant or breast-feeding**, ask a health professional before use.

Keep out of reach of children.

#### **DIRECTIONS**

**Adults:** Allow 3 or 4 pellets to dissolve in the mouth 3 times a day until symptoms are relieved or as directed by a health professional.

#### OTHER INFORMATION

Store at room temperature.

#### **INACTIVE INGREDIENTS**

Lactose, sucrose.

#### **QUESTIONS?**

#### 1-800-404-4666

The letters 'HPUS' indicate that the component in this product is officially monographed in the Homeopathic Pharmacopoeia of the United States.

These claims have not been reviewed by the Food and Drug Administration. They are based on traditional homeopathic practice.

80 Pellets

Pellet dispenser

Mfd for: HOMEOLAB USA INC., 3025 De L'Assomption, Montreal, QC, H1N 2H2, CANADA

Product of Canada

LABEL

### HOMEOPATHIC MEDICINE

# NATRUM SULPHURICUM

**1**X

Sodium Sulfate

NDC 60512-6350-1

BRONCHIAL IRRITATION MADE WORSE BY HUMIDITY \*

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#### NATRUM SULPHURICUM

natrum sulphuricum pellet

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:60512-6350

Route of Administration ORAL

Active Ingredient/Active Moiety					
Ingredient Name	<b>Basis of Strength</b>	Strength			
SODIUM SULFATE (UNII: 0 YPR65R21J) (SODIUM SULFATE ANHYDROUS - UNII:36KCS0R750)	SODIUM SULFATE	1 [hp_X]			

Inactive Ingredients				
Ingredient Name	Strength			
LACTOSE (UNII: J2B2A4N98G)				
SUCROSE (UNII: C151H8 M554)				

]	Packaging					
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:60512-6350-1	80 in 1 TUBE				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		10/11/1995		

## Labeler - HOMEOLAB USA INC. (202032533)

## Registrant - HOMEOLAB USA INC. (202032533)

Establishment				
Name	Address	ID/FEI	Business Operations	
HOMEOLAB USA INC.		202032533	manufacture(60512-6350)	

Revised: 10/2013 HOMEOLAB USA INC.