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.

HOMEOPATHIC MEDICINE NDC 60512-1034-1

ACTIVE INGREDIENT HPUS

NATRUM SULPHURICUM 3X & HIGHER (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	80 Pellet	
NATRUM 3X&+		Mfd fo Pellets dispense
SULPHURICUM		for: ser
Sodium Sulfate		3025
NDC 60512-1034-1 BRONCHIAL IRRITATION MADE WORSE BY HUMIDITY *		не L'As
		Sompti
*These claims have not been reviewed by the Food and Drug Administration. They are based on traditional homeopathic practice.		Con, Ma
ACTIVE INGREDIENT HPUS: Listed above. USE: For self-limiting condition listed above or as directed by a health professional.		DLAE ontreal QC H
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relieved or as directed by a health professional. OTHER INFORMATION: Store at room temperature. INACTIVE INGREDIENTS: Lactose, sucrose.	/13	CANAD PROD
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	A	DA

NATRUM SULPHURICUM

natrum sulphuricum pellet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60512-1034
Route of Administration	ORAL		

L

ve Ingredient/Ac	ctive Molety					
Ingredient Name			Basis of Strength	Strength		
UM SULFATE (UNII:	0 YPR65R21J) (SODIUM SULFATE AI	NHYDROUS - UNI	I:36KCS0R750)	SODIUM SULFATE	3 [hp_X]	
tive Ingredients						
	Ingredient Name			Strength		
OSE (UNII: J2B2A4N	98G)					
OSE (UNII: C151H8 M	554)					
kaging						
Item Code	Package Description	Marketing	Start Date	Marketing En	d Date	
C:60512-1034-1	80 in 1 TUBE					
rketing Inform	nation					
rketing Category	Application Number or Monog	raph Citation	Marketing Sta	rt Date Marketing	End Date	
roved homeopathic			10/11/1995			
*						
	uM SULFATE (UNII: tive Ingredients OSE (UNII: J2B2A4N OSE (UNII: C151H8M OSE (UNII: C151H8M tem Code C:60512-1034-1	UM SULFATE (UNII: 0 YPR65R21J) (SODIUM SULFATE AN tive Ingredients Ingredient Name OSE (UNII: J2B2A4N98G) OSE (UNII: C151H8M554) Agging Item Code Package Description C:60512-1034-1 80 in 1 TUBE rketing Information rketing Category Application Number or Monog	Ingredient Name UM SULFATE (UNII: 0 YPR6 5R2 1J) (SO DIUM SULFATE ANHYDRO US - UNI tive Ingredients Ingredient Name COSE (UNII: 12B2A4N98G) OSE (UNII: C151H8M554) Kaging Item Code Package Description Marketing C60512-1034-1 80 in 1 TUBE rketing Information Application Number or Monograph Citation	Ingredient Name Ingredient Name Ingredient Name Ingredient Name Iose (UNII: J2B2A4N98G) OSE (UNII: C151H8M554) GSE (UNII: C151H8M554) Kaging Item Code Package Description Marketing Start Date C:60512-1034-1 80 in 1 TUBE rketing Information Marketing Start Date	Ingredient Name Basis of Strength Marketing Enformation Solution Sulfate Anhydrous - UNIE36KCS0R750) Solution	

Labeler - HOMEOLAB USA INC. (202032533)

Registrant - HOMEOLAB USA INC. (202032533)

Establishment

L.

Name	Address	ID/FEI	Business Operations
HOMEOLAB USA INC.		202032533	manufacture(60512-1034)

Revised: 10/2013

HOMEOLAB USA INC.