SULPHUR- sulphur 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 60512-1047-1

ACTIVE INGREDIENT HPUS

SULPHUR 1X & UP

Sulphur

SKIN PROBLEMS*

USE

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

WARNINGS

Enter section text here

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.

The letters 'HPUS' indicate that the component in this product is officially monographed in the Homoeopathic 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

3025 De L'Assomption, Montreal, QC, H1N 2H2, CANADA

LABEL

HOMEOPATHIC MEDICINE

SULPHUR

1 X &+

Sulphur NDC 60512-1047-1 SKIN PROBLEMS *

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ACTIVE INGREDIENT HPUS: Listed above. 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.

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Rev. 10/13

SULPHUR

sulphur pellet

Product Informati	on
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:60512-1047

Route of Administration ORAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
SULFUR (UNII: 70 FD1KFU70) (SULFUR - UNII:70 FD1KFU70)	SULFUR	1 [hp_X]

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

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:60512-1047-1	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)

Establishment			
Name	Address	ID/FEI	Business Operations
HOMEOLAB USA INC		202032533	manufacture(60512-1047)

Revised: 5/2014 HOMEOLAB USA INC