

THIOSINAMINUM- allylthiourea pellet
Washington Homeopathic Products

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.

Thiosinaminum Kit Refill 30c

ACTIVE INGREDIENTS

THIOSINAMINUM 30C

USES

Indicated for Scarring.

KEEP OUT OF REACH OF CHILDREN

As with all medications, keep out of reach of children.

INDICATIONS

THIOSINAMINUM Scarring

STOP USE AND ASK DOCTOR

If symptoms persist or recur, discontinue use. If pregnant or nursing, consult a licensed practitioner before using this product.

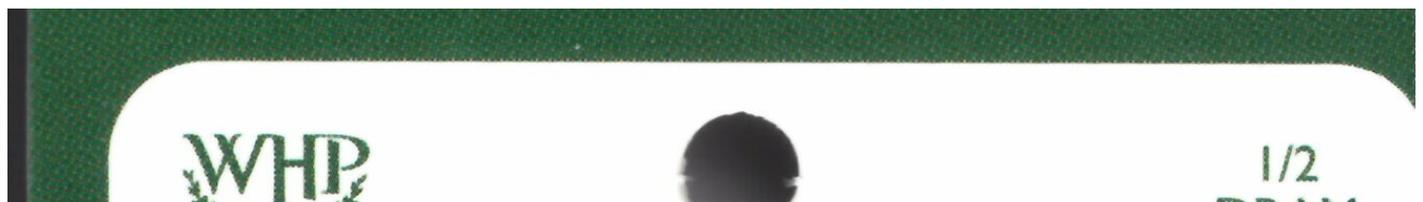
DIRECTIONS

(Adults or Children): Dissolve 3 to 5 pellets under tongue 3 times a day or as directed by a licensed practitioner. Take at greater intervals as condition subsides.

INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL



CONTENTS: **HOMEOPATHIC MEDICINE (HPUS*)**

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Take at greater intervals as condition subsides.

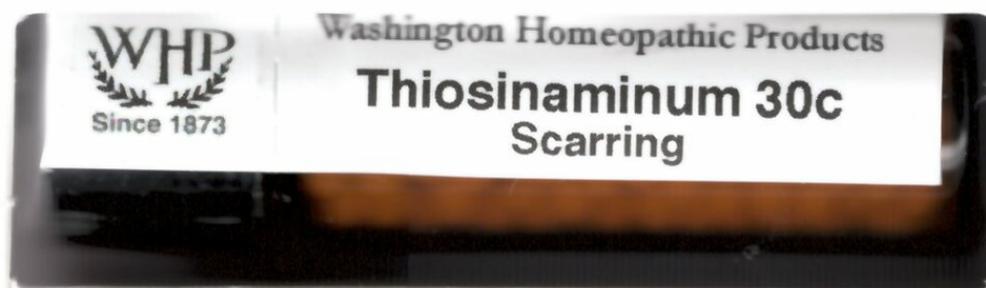
WARNING: If symptoms worsen or persist for more than 7 days, or if pregnant or nursing, consult your practitioner.
Keep out of reach of children.

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

Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

DO NOT USE IF SEAL IS BROKEN.

SUCROSE/LACTOSE PELLETS



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EST. # 3006489197

THIOSINAMINUM

allylthiourea pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71919-819
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLYLTHIOUREA (UNII: 706IDJ14B7) (ALLYLTHIOUREA - UNII:706IDJ14B7)	ALLYLTHIOUREA	30 [hp_C]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
LACTOSE (UNII: J2B2A4N98G)	

Product Characteristics

Color	white (white)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71919-819-01	350 in 1 VIAL, GLASS; Type 0: Not a Combination Product	11/08/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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unapproved homeopathic		11/08/2023	
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Labeler - Washington Homeopathic Products (084929389)

Establishment

Name	Address	ID/FEI	Business Operations
Washington Homeopathic Products		084929389	manufacture(71919-819)

Revised: 1/2025

Washington Homeopathic Products