NATIVEREMEDIES GOUT-GONE- antimon crud, benzoicum ac, ledum, nux vom, quercus, rhododendron, silicea liquid
Silver Star Brands

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.

NativeRemedies® Gout-Gone™

HPUS Active Ingredients

HPUS Active Ingredients: Equal volumes of each ingredient: Antimon crud 6C, Benzoicum ac 6C, Ledum 6C, Nux vom 6C, Quercus 12C, Rhododendron 6C, Silicea 6X.

The letters HPUS indicate that the components in this product are officially monographed in the Homeopathic Pharmacopoeia of the United States.

Uses*:

Uses*: Homeopathic remedy for the relief of acute swelling, inflammation, and burning pain in small joints, especially in the big toe.

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

Warnings

Warnings: If symptoms persist or worsen, consult a healthcare professional. If pregnant or breastfeeding, ask a health professional before use. Keep this and all medication out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

Directions: Mix in 1/4 cup water and sip slowly. Alternatively, drops may be taken directly into the mouth. For all ages: Take 0.50 mL every 20-30 minutes until symptoms improve.

Contains no artificial flavors or colorants. No gluten added.

Tampler resistant for your protection. Use only if safety seal is intact.

Inactive Ingredients

Inactive Ingredients: Purified water, USP grain alcohol (20% of volumn).

Keep this and all medications out of reach of children.

If pregnant or breastfeeding, ask a health professional before use.

Relieves swelling, inflammation, & buring joint pain.

NATIVEREMEDIES GOUT-GONE

antimon crud, benzoicum ac, ledum, nux vom, quercus, rhododendron, silicea liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68703-390
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	6 [hp_X] in 59 mL
ANTIMONY TRISULFIDE (UNII: F79059A38U) (ANTIMONY TRISULFIDE - UNII:F79059A38U)	ANTIMONY TRISULFIDE	6 [hp_C] in 59 mL
BENZOIC ACID (UNII: 8SKN0B0MIM) (BENZOIC ACID - UNII:8SKN0B0MIM)	BENZOIC ACID	6 [hp_C] in 59 mL
QUERCUS ROBUR FLOWER (UNII: ML644HED2V) (QUERCUS ROBUR FLOWER - UNII:ML644HED2V)	QUERCUS ROBUR FLOWER	12 [hp_C] in 59 mL
STRYCHNOS NUX-VOMICA SEED (UNII: 269XH13919) (STRYCHNOS NUX-VOMICA SEED - UNII:269XH13919)	STRYCHNOS NUX-VOMICA SEED	6 [hp_C] in 59 mL
RHODODENDRON AUREUM LEAF (UNII: IV92NQJ73U) (RHODODENDRON AUREUM LEAF - UNII:IV92NQJ73U)	RHODODENDRON AUREUM LEAF	6 [hp_C] in 59 mL
LEDUM PALUSTRE TWIG (UNII: 877L01IZ0P) (LEDUM PALUSTRE TWIG - UNII:877L01IZ0P)	LEDUM PALUSTRE TWIG	6 [hp_C] in 59 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68703-390-02	59 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	04/03/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		04/03/2023	

Labeler - Silver Star Brands (006070379)

Registrant - Silver Star Brands (006070379)

Establishment

Name	Address	ID/FEI	Business Operations
OHM Pharma		030572478	manufacture(68703-390)

Revised: 12/2023 Silver Star Brands