FERRUM ROSATUM GRAPHITES- ferrum rosatum graphites liquid Uriel Pharmacy 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.

Ferrum rosatum Graphites

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops. Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredients: Rosa cum Ferro (Garden rose, Red iron ore) 3X, Ferrum sidereum (Meteoric iron) 6X, Graphites (Foliated crystalline carbon) 17X

Inactive Ingredients: Distilled water, Organic cane alcohol, Lactose

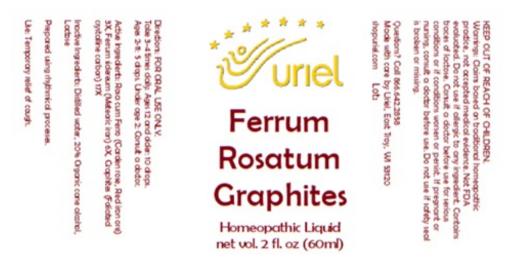
Prepared using rhythmical processes.

Use: Temporary relief of cough.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Contains traces of lactose. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858 Made with care by Uriel, East Troy, WI 53120 shopuriel.com Lot:



FERRUM ROSATUM GRAPHITES

ferrum rosatum graphites liquid

Product Information

Product	Туре
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HUMAN OTC DRUG

Item Code (Source)

Route of Administration ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SEDUM ROSEUM WHOLE (UNII: RH0WP583U3) (SEDUM ROSEUM WHOLE - UNII:RH0WP583U3)	SEDUM ROSEUM WHOLE	3 [hp_X] in 1 mL	
FERRIC OXIDE RED (UNII: 1K09F3G675) (FERRIC OXIDE RED - UNII:1K09F3G675)	FERRIC OXIDE RED	3 [hp_X] in 1 mL	
IRON (UNII: E1UOL152H7) (IRON - UNII:E1UOL152H7)	IRON	6 [hp_X] in 1 mL	
GRAPHITE (UNII: 4QQN74LH4O) (GRAPHITE - UNII:4QQN74LH4O)	GRAPHITE	17 [hp_X] in 1 mL	

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
LACTOSE (UNII: J2B2A4N98G)	
LACTOSE (UNII: J2B2A4N98G)	

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951- 4064-3	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	
Marketing Information				

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment			
Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-4064)

Revised: 6/2025

Uriel Pharmacy Inc.