METEORIC IRON PRUNUS- meteoric iron prunus 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.

Meteoric Iron Prunus

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: Prunus spin. (Blackthorn) 2X, Echinacea (Purple coneflower) 3X, Phosphorus (Yellow phosphorus) 6X, Meteoric iron 12X, Quartz (Rock crystal) 12X

Inactive Ingredients: Distilled water, Organic cane alcohol

prepared using rhythmical processes

Uses: Temporary relief of flu symptoms.

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. 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:



METEORIC IRON PRUNUS

meteoric iron prunus liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:48951-7054

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SLOE (UNII: 3MLB4858X7) (SLOE - UNII:3MLB4858X7)	SLOE	2 [hp_X] in 1 mL	
ECHINACEA, UNSPECIFIED (UNII: 4N9P6CC1DX) (ECHINACEA, UNSPECIFIED - UNII:4N9P6CC1DX)	ECHINACEA, UNS PECIFIED	3 [hp_X] in 1 mL	
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	6 [hp_X] in 1 mL	
IRON (UNII: E1UOL152H7) (IRON - UNII:E1UOL152H7)	IRON	12 [hp_X] in 1 mL	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	12 [hp_X] in 1 mL	

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:48951- 7054-3	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(48951-7054)	

Revised: 3/2024

Uriel Pharmacy Inc.