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.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

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

Inactive Ingredients: Water, Salt

Uses: Temporary relief of flu or exhaustion, especially during convalescence or when symptoms first appear.

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.

Questions? Call 866.642.2858
Uriel, East Troy, WI 53120
www.urielpharmacy.com

METEORIC IRON PRUNUS

meteoric iron prunus liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-7052
Route of Administration	ORAL		

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, UNSPECIFIED	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)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:48951-7052-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package		

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(48951-7052)

Revised: 5/2018 Uriel Pharmacy Inc.