

FERRUM ARSENICOSUM 6- ferrum arsenicosum 6 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 arsenicosum 6

Directions: FOR ORAL USE.

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

Active Ingredient: Ferrum arsenicosum (iron arsenite solution) 6X

Inactive Ingredients: Water, Salt, Lactose

Use: Temporary relief of digestive upset.

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.

Questions? Call 866.642.2858

Uriel, East Troy, WI 53120

www.urielpharmacy.com

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

Active Ingredient: Ferrum arsenicosum (iron arsenite solution) 6X

Inactive Ingredients: Water, Salt, Lactose

Use: Temporary relief of headache.

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.

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Uriel, East Troy, WI 53120
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Ferrum
arsenicum 6X

Homeopathic Ampules
net vol. 0.3 fl. oz (10 x 1 ml)

Ferrum arsenicosum 6X

FERRUM ARSENICOSUM 6

ferrum arsenicosum 6 liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-4057
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
FERROUS ARSENATE (UNII: 129CO35H12) (FERROUS ARSENATE - UNII:129CO35H12)		FERROUS ARSENATE	6 [hp_X] in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
WATER (UNII: 059QF0K00R)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
LACTOSE (UNII: J2B2A4N98G)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-4057-1	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(48951-4057)

Revised: 4/2018

Uriel Pharmacy Inc.