## THYREOIDEA FERRUM- thyreoidea ferrum 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.

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## Thyreoidea Ferrum

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: Fucus ves. (Bladderweed) 3X, Urtica dioica (Stinging nettle) 3X, Glandula thyreoidea (Bovine thyroid gland) 4X, Ferrum arsenicosum (Iron arsenite solution) 6X, Ferrum siderum (Meteoric iron) 6X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol, Lactose

Prepared using rhythmical processes.

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. Contains traces of lactose. 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:



## THYREOIDEA FERRUM

thyreoidea ferrum liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-9390	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
FUCUS VESICULOSUS (UNII: 535G2ABX9M) (FUCUS VESICULOSUS - UNII:535G2ABX9M)	FUCUS VESICULOSUS	3 [hp_X] in 1 mL	
IRON (UNII: E1UOL152H7) (IRON - UNII:E1UOL152H7)	IRON	6 [hp_X] in 1 mL	
URTICA DIOICA (UNII: 710FLW4U46) (URTICA DIOICA - UNII:710FLW4U46)	URTICA DIOICA	3 [hp_X] in 1 mL	
THYROID, UNSPECIFIED (UNII: 0B4FDL9I6P) (THYROID, UNSPECIFIED - UNII: 0B4FDL9I6P)	THYROID, UNSPECIFIED	4 [hp_X] in 1 mL	
FERROUS ARSENATE (UNII: 129CO35H12) (FERROUS ARSENATE - UNII:129CO35H12)	FERROUS ARSENATE	6 [hp_X] in 1 mL	

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

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:48951- 9390-3	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	<b>Business Operations</b>		
Uriel Pharmacy, Inc		043471163	manufacture(48951-9390)		

Revised: 3/2025 Uriel Pharmacy, Inc