CUPRITE 3 SPECIAL ORDER- cuprite 3 special order powder 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.

Cuprite 3 Special Order

Directions: FOR ORAL USE ONLY.

Ages 12 and older: 1/8 teaspoon. Ages 2-11: 1/16 teaspoon. Under age 2: Consult a doctor.

Active Ingredient: Cuprite 3X Inactive Ingredient: 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 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 mssing.

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

Chrestions FUNC CHALL DIS CHALL'S Ages 25 older 18 teappoon. Ages 2-th: 176 teappoon. Under age 2. Consult a dector.

Active Ingredient: Cuprite 3X inactive Ingredient: Lactose. Use: Temporary relief of headache.

Questions? Call 866.642.2888

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www.uriels/harmary.com. Lob.

Cuprite
3X
Special Order
Homeopathic Powder
net wt. 1.7 oz (50g)

Withing Claim based on traditional homeopethic practice, and ecosphed medical homeopethic practice, and ecosphed medical evidence. Net FDA controlled, Co not use if a largis to any ingredient. Contains factors. Combin a decide before use for serious conditions or conditions, uneman or partial. If programs or number, consult a decide before use, Do net use if safety seed is broken or methog.

CUPRITE 3 SPECIAL ORDER

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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-3137
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CUPRIC OXIDE (UNII: V1XJO704R4) (CUPRIC CATION - UNII:8CBV67279L)	CUPRIC CATION	3 [hp X] in 1 g

Inactive Ingredients	
Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	

	eting End Date
1 NDC:48951-3137- 4 60 g in 1 BOTTLE, GLASS; Type 0: Not a Combination Product 09/01/2009	

Marketing Inform	mation		
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 (48951-3137)

Revised: 4/2018 Uriel Pharmacy Inc.