

AURUM MURIATICUM NATRONATUM- aurum muriaticum natronatum tablet
Rxhomeo Private Limited d.b.a. Rxhomeo, 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.

ACTIVE INGREDIENT

AURUM MURIATICUM NATRONATUM HPUS 3X and Higher

USES

Headache & Hemorrhoids

INDICATIONS

Condition listed above or as directed by the physician

DOSAGE

Adults- Take 4 or 6 Tablets by mouth, three times daily or as suggested by physician. Children 2 years and older- take 1/2 the adult dose.

WARNINGS

This product is to be used for self-limiting conditions

If symptoms do not improve in 4 days, or worsen, discontinue use and seek assistance of health professional

As with any drug, if you are pregnant, or nursing a baby, seek professional advice before taking this product

Keep this and all medication out of reach of children

Do not use if capseal is broken or missing.

Close the cap tightly after use.

INACTIVE INGREDIENTS

Lactose

STORAGE

Store in a cool dark place

QUESTIONS OR COMMENTS

www.Rxhomeo.com | 1.888.2796642 | info@rxhomeo.com

Rxhomeo, Inc 9415 Burnet Road, Suite 312, Austin, TX 78758

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



AURUM MURIATICUM NATRONATUM 3X

Ingredients: Active: As Above, Inactive: Lactose
USES: Headache & Hemorrhoids



Manufactured according to the Homeopathic
Pharmacopoeia of the United States
Est. # 30052969310
info@rxhomeo.com/RxHomeo.com
1-888-8RYONIA (279-6442)



Distributed in the US by Rxhomeo, Inc 9415 Burnet Road, Suite 312, Austin, TX 78758 Manufactured by: Rxhomeo Private Limited "Indradhanush", 4-1-424 to 426, Bank Street, Abids, Hyderabad #500001 India.

NDC 15631-0529-3 B.No XXXXXXXXXX MFD XX/XX EXP XX/XX Contents 100 Tablets

AURUM MURIATICUM NATRONATUM

aurum muriaticum natronatum tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:15631-0529
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM TETRACHLORO AURATE (UNII: 7FT6QUT299) (TETRACHLORO AURATE ION - UNII:ZNL6IP5PJX)	SODIUM TETRACHLORO AURATE	3 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	3mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15631-0529-0	1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:15631-0529-1	4 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:15631-0529-2	50 in 1 CONTAINER; Type 0: Not a Combination Product		

4	NDC:15631-0529-3	100 in 1 CONTAINER; Type 0: Not a Combination Product		
5	NDC:15631-0529-4	250 in 1 CONTAINER; Type 0: Not a Combination Product		
6	NDC:15631-0529-5	500 in 1 CONTAINER; Type 0: Not a Combination Product		
7	NDC:15631-0529-6	1000 in 1 CONTAINER; Type 0: Not a Combination Product		
8	NDC:15631-0529-7	10000 in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/30/2015	

Labeler - Rxhomeo Private Limited d.b.a. Rxhomeo, Inc (650833994)

Establishment

Name	Address	ID/FEI	Business Operations
Rxhomeo, Inc		832534981	wholesale drug distributor(15631-0529)

Establishment

Name	Address	ID/FEI	Business Operations
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc		650833994	manufacture(15631-0529) , label(15631-0529)

Revised: 12/2015

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