#### ZINCUM METALLICUM- zincum metallicum pellet 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.

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## **ACTIVE INGREDIENT**

ZINCUM METALLICUM HPUS 3X and higher

## USES

USES: Temporary Relief - Headache\*

\* Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

## INDICATIONS

Condition listed above or as directed by the physician

## DOSAGE

Adults- Take 4 or 6 Pellets 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 preganant, 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**

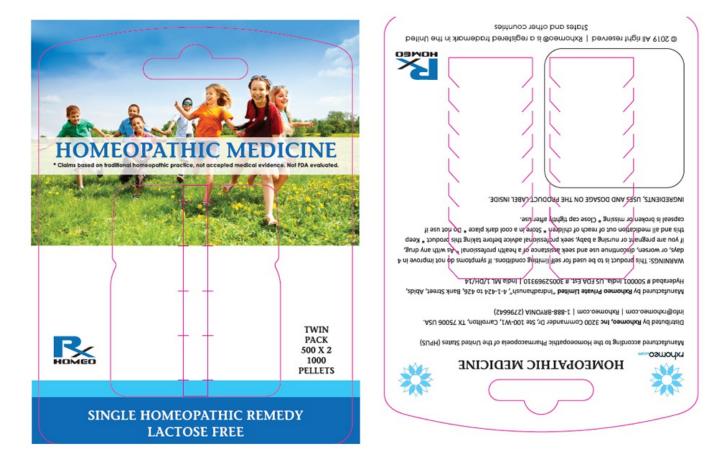
Sucrose

STORAGE

# **QUESTIONS OR COMMENTS**

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# ZINCUM METALLICUM

zincum metallicum pellet

**Product Information** 

Product Type		HUMAN OTC DRUG		ode (Source)		NDC:15631-0469	
R	oute of Admir	nistration	ORAL				
A	ctive Ingred	lient/Active	Moiety				
Ingredient Name					Basis of Strengt		-
ZI	<b>NC</b> (UNII: J41CS)	27QDS) (ZINC - l	JNII:J41CSQ7QDS)		ZINC		3 [hp_X]
lr	nactive Ingr	edients					
		Ing		Strength			
P	ackaging						
#	ltem Code	Pa	ackage Description		Marketin Dat	-	Marketing End Date
1	NDC:15631- 0469-0	100 in 1 VIAL, S Combination Pr	INGLE-DOSE; Type 0: Not oduct	a	01/01/2018		
2	NDC:15631- 0469-1	200 in 1 PACKA Product	GE; Type 0: Not a Combina	ation	01/01/2018		
3	NDC:15631- 0469-2	400 in 1 PACKAGE; Type 0: Not a Combination Product		ation	01/01/2018		
4	NDC:15631- 0469-3	750 in 1 PACKA Product	AGE; Type 0: Not a Combination 01/01/201		01/01/2018		
5	NDC:15631- 0469-4	Product	AGE; Type 0: Not a Combi	01/01/2018			
6	NDC:15631- 0469-5	Product	(AGE; Type 0: Not a Combination		01/01/2018		
7	NDC:15631- 0469-6	Product	GE; Type 0: Not a Combination		08/23/2021		
8	NDC:15631- 0469-7 Product		AGE; Type 0: Not a Combination		08/23/2021		
Μ	larketing	Informat	ion				
	Marketing Category	Applica	tion Number or Mono Citation	ograph	Marketing Date		Marketing End Date
un	approved meopathic				11/01/2015		
٦O	meepaane						

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

Establishment									
Name	Address	ID/FEI	<b>Business Operations</b>						
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc		650833994	manufacture(15631-0469) , label(15631-0469)						