

LACHESIS MUTUS- lachesis mutus 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.

ACTIVE INGREDIENT

LACHESIS MUTUS HPUS 8X and higher

USES

USES: Temporary Relief - Running 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 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

Sucrose

STORAGE

Store in a cool dark place

QUESTIONS OR COMMENTS

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

Rxhomeo, Inc 3200 Commander Dr, Ste 100-W1, Carrollton, TX 75006 USA



rxhomeo.com

LACHESIS MUTUS 8X

Ingredients: Active: As Above, Inactive: Sucrose
USES: Temporary Relief - Running Headache *

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



XXX Pellets

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Distributed by Rxhomeo, Inc 3200 Commander Dr, Ste 100-W1, Carrollton, TX 75006 USA Manufactured by: Rxhomeo Private Limited "Indraathanush", 4-1-424 to 426, Bank Street, Abids, Hyderabad #1 India.

NDC 15631-0261 B.No XXXXXXXXXX MFD XX/XX EXP XX/XX

HOMEPATHIC MEDICINE

<<< MINI LABEL USED
WITH BLISTER PACK FOR
SELECT PACK SIZES



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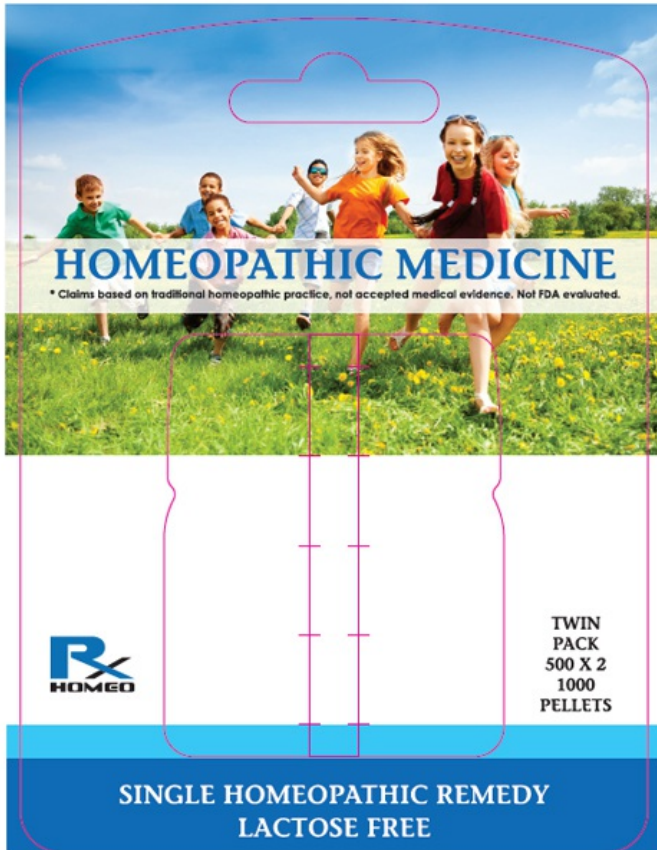
NDC 15631-0261 B.No XXXXXXXXXX MFD XX/XX EXP XX/XX Contents 100 Pellets



HOMEPATHIC MEDICINE



Manufactured according to the Homeopathic Pharmacopoeia of the United States
FDA Est. # 30052969310 | India ML 1/DH/14
info@rxhomeo.com | Rxhomeo.com
1-888-8RYONIA (279-6642)



NOTES RELATED TO PACKING

1. All labels and secondary packing specimens have our India address as the manufacturer, but if the product is manufactured by some other manufacturer, the manufacturers name and address will be changed or removed if those attributes are confidential. However, the manufacturers US FDA Establishment number will be printed at all times.
2. Primary Packing > Depending on the pack size/contents, a bottle/container of appropriate size will be used. Depending on the bottle/container size, a label of appropriate size will be used i.e. regular size or mini size.
3. Secondary Packing > If the bottle/container is small, we will use a secondary packing i.e a Carton box or a Blister pack.

LACHESIS MUTUS

lachesis mutus pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:15631-0261	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
LACHESIS MUTA VENOM (UNII: VSW71SS07I) (LACHESIS MUTA VENOM - UNII:VSW71SS07I)		LACHESIS MUTA VENOM	8 [hp_X]	
Inactive Ingredients				
Ingredient Name		Strength		
SUCROSE (UNII: C151H8M554)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15631-0261-0	100 in 1 VIAL, SINGLE-DOSE; Type 0: Not a Combination Product	01/01/2018	
2	NDC:15631-0261-1	200 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
3	NDC:15631-0261-2	400 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
4	NDC:15631-0261-3	750 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
5	NDC:15631-0261-4	2500 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
6	NDC:15631-0261-5	12500 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
7	NDC:15631-0261-6	500 in 1 PACKAGE; Type 0: Not a Combination Product	08/23/2021	
8	NDC:15631-0261-7	1000 in 1 PACKAGE; Type 0: Not a Combination Product	08/23/2021	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		09/05/2015		

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

Establishment

Name	Address	ID/FEI	Business Operations
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc		650833994	manufacture(15631-0261) , label(15631-0261)

