LACHESIS BELLADONNA- lachesis belladonna 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.

Lachesis Belladonna

Directions: FOR ORAL USE.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredients: Atropa belladonna (Nightshade) 4X, Mercurialis (Dog's mercury) 6X,

Hepar sulfuris (Liver of lime) 7X, Lachesis (Bushmaster venom) 12X

Inactive Ingredients: Water, Salt

Use: Promotes healing of minor wounds.

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. Consult a doctor before use for serious conditions or if conditions worsen or persist. Contains traces of lactose. If pregnant or nursing, consult a doctor before use.

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



Lachesis Belladonna

LACHESIS BELLADONNA

lachesis belladonna liquid

| Product Information | | | |
|-------------------------|----------------|--------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:48951-6012 |
| Route of Administration | ORAL | | |
| | | | |
| | | | |

| Active Ingredient/Active Moiety | | | |
|--|-------------------------|----------------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII: WQZ3G9PF0H) | ATROPA BELLADONNA | 4 [hp_X] in 1 mL | |
| MERCURIALIS PERENNIS (UNII: Q35465A1MA) (MERCURIALIS PERENNIS - UNII:Q35465A1MA) | MERCURIALIS PERENNIS | 6 [hp_X] in 1 mL | |
| CALCIUM SULFIDE (UNII: 1MBW07J51Q) (CALCIUM SULFIDE - UNII:1MBW07J51Q) | CALCIUM SULFIDE | 7 [hp_X] in 1 mL | |
| LACHESIS MUTA VENOM (UNII: VSW71SS07I) (LACHESIS MUTA VENOM - UNII: VSW71SS07I) | LACHESIS MUTA VENOM | 12 [hp_X] in 1 mL | |

| Inactive Ingredients | | |
|------------------------------------|----------|--|
| Ingredient Name | Strength | |
| WATER (UNII: 059QF0KO0R) | | |
| SODIUM CHLORIDE (UNII: 451W47IQ8X) | | |

| P | Packaging | | | |
|---|---------------------------------|---|-------------------------|-----------------------|
| # | # Item Code Package Description | | Marketing Start Date | Marketing End Date |
| 1 | NDC:48951- 6012-1 | 10 in 1 BOX | 09/01/2009 | |
| 1 | | 1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package | | |

| 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 | Business Operations | | |
| Uriel Pharmacy Inc. | | 043471163 | manufacture(48951-6012) | | |

Revised: 11/2023 Uriel Pharmacy Inc.