**PULSATILLA TORMENTILLA- pulsatilla tormentilla 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.*

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**Pulsatilla Tormentilla**

Directions: FOR ORAL USE.

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

Active Ingredients: Hepar (Bovine liver) 4X, Magnesium hydroxydatum (Magnesium hydroxide) 4X, Pulsatilla (Pasqueflower) 12X, Tormentilla (Bloodwort) 30X

Inactive Ingredients: Water, Salt

"prepared using rhythmical processes"

Use: Temporary relief of premenstrual syndrome (PMS).

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. If pregnant or nursing, consult a doctor before use.

Questions? Call 866.642.2858 Made by Uriel, East Troy, WI 53120 shopuriel.com Lot:


## PULSATILLA TORMENTILLA

pulsatilla tormentilla liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:48951-8098 |
| **Route of Administration** | ORAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **MAMMAL LIVER** (UNII: D0846624BI) (MAMMAL LIVER - UNII:D0846624BI) | MAMMAL LIVER | 4 [hp_X] in 1 mL |
| **MAGNESIUM HYDROXIDE** (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838) | MAGNESIUM HYDROXIDE | 4 [hp_X] in 1 mL |
| **PULSATILLA VULGARIS** (UNII: I76KB35JEV) (PULSATILLA VULGARIS - UNII:I76KB35JEV) | PULSATILLA VULGARIS | 12 [hp_X] in 1 mL |
| **POTENTILLA ERECTA ROOT** (UNII: BI896CKT6B) (POTENTILLA ERECTA ROOT - UNII:BI896CKT6B) | POTENTILLA ERECTA ROOT | 30 [hp_X] in 1 mL |

## Inactive Ingredients

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

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:48951-8098-1 | 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(48951-8098) |

Revised: 4/2024 Uriel Pharmacy Inc.