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.

Pulsatilla Tormentilla

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops. Ages 2-11: 5 drops. Under age 2: Consult a doctor.

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

Inactive Ingredients: Distilled water, Propolis

Use: Temporarily relieves symptoms 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. Do not use if safety seal is broken or missing.

REFRIGERATE AFTER OPENING.
BEST WHEN USED WITHIN 30 DAYS OF OPENING.

Questions? Call 866.642.2858 Made with care by Uriel, East Troy, WI 53120 www.urielpharmacy.com

PULSATILLA TORMENTILLA

pulsatilla tormentilla liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-8099
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)	
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8099-3	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	

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(48951-8099)

Revised: 4/2018 Uriel Pharmacy Inc.