HAMAMELIS AESCULUS- hamamelis aesculus 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.

Hamamelis Aesculus

Directions: FOR ORAL USE.

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

Active Ingredients: Aesculus (Horse chestnut) 3X, Hamamelis (Witch hazel) 3X, Paeonia (Peony) 3X, Pulsatilla (Pasqueflower) 3X, Hirudo (Leech) 3X, Mercurius vivus (Mercury) 17X

Inactive Ingredients: Water, Salt

Uses: Temporary relief of burning and itching due to varicose veins or hemorrhoids.

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 Uriel, East Troy, WI 53120 www.urielpharmacy.com

HAMAMELIS AESCULUS
hamamelis aesculus liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-5031
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HORSE CHESTNUT (UNII: 3C18L6RJAZ) (HORSE CHESTNUT - UNII:3C18L6RJAZ)	HORSE CHESTNUT	3 [hp_X] in 1 mL
HAMAMELIS VIRGINIANA TOP (UNII: UDA30A2JJY) (HAMAMELIS VIRGINIANA TOP - UNII:UDA30A2JJY)	HAMAMELIS VIRGINIANA TOP	3 [hp_X] in 1 mL

PAEONIA OFFICINALIS ROOT (UNII: 8R564U2E1P) (PAEONIA OFFICINALIS ROOT - UNII:8R564U2E1P)	PAEONIA OFFICINALIS ROOT	3 [hp_X] in 1 mL
PULSATILLA VULGARIS (UNII: I76KB35JEV) (PULSATILLA VULGARIS - UNII:I76KB35JEV)	PULSATILLA VULGARIS	3 [hp_X] in 1 mL
HIRUDIN (UNII: MNY7X23SRZ) (HIRUDIN - UNII:MNY7X23SRZ)	HIRUDIN	6 [hp_X] in 1 mL
MERCURY (UNII: FXS1BY2PGL) (MERCURY - UNII:FXS1BY2PGL)	MERCURY	17 [hp_X] in 1 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-5031-1	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(48951-5031)

Revised: 5/2018 Uriel Pharmacy Inc.