HEMORRHOID RELIEF- aesculus hippocastanum, arnica montana, calcarea fluorica, carduus marianus, collinsonia canadensis, hamamelis virginiana, muriaticum acidum, ratanhia spray Liddell Laboratories, 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.

DRUG FACTS:

ACTIVE INGREDIENTS:

(in each spray) 12.50 % of Aesculus Hippocastanum 1X, Arnica Montana 3X, Calcarea Fluorica 9X, Carduus Marianus 1X, Collinsonia Canadensis 3X, Hamamelis Virginiana 3X, Muriaticum Acidum 6X, Ratanhia 3X.

INDICATIONS:

May temporarily relieve symptoms associated with hemorrhoids.**

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

WARNINGS:

If symptoms persist for more than 7 days, consult a doctor.

If pregnant or breast feeding, ask a doctor before use.

Keep out of reach of children. In case of overdose, get medical help or call a Poison Control Center right away.

Do not use if TAMPER EVIDENT seal around neck of bottle is missing or broken.

KEEP OUT OF REACH OF CHILDREN:

Keep out of reach of children. In case of overdose, get medical help or call a Poison Control Center right away.

DIRECTIONS:

Adults and children over 12: Spray twice under the tongue 3 times per day.

Children 12 and under: Consult a doctor prior to use.

INDICATIONS:

May temporarily relieve symptoms associated with hemorrhoids.**

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

INACTIVE INGREDIENTS:

Organic alcohol 23% v/v, Purified water.

QUESTIONS:

DIST. BY LIDDELL LABORATORIES WOODBINE, IA 51579 WWW.LIDDELL.NET 1-800-460-7733

PACKAGE LABEL DISPLAY:

ORAL SPRAYS

LIDDELL

LABORATORIES

EST. 1994

²⁷Hem

Hemorrhoid

Relief

HOMEOPATHIC

1.0 FL. OZ. (30 ml)



ACTIVE INGREDIENTS (in each spray) 12.50% of Carduus mar 1X, Collinsonia 3X, Hamamelis 3X, Aesculus hipp 1X, Arnica 3X, Calc fluor 9X, Muriaticum ac 6X, Ratanhia 3X INACTIVE

NGREDIENTS Organic alcohol 23% v/v, Purified

water

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

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LOT: XXXXXX EXP: MM/YY

DIRECTIONS Adults & children over 12: Spray twice under the tongue 3 times per day. Children **NDICATIONS** May temporarily relieve symptoms WARNINGS If symptoms persist for more than **feeding**, ask a doctor before use. **Keep out of reach of children**. In case of overdose, get medical help or call a Poison Control Center 12 and under: Consult a doctor prior to use. days, consult a doctor. If pregnant or breast associated with hemorrhoids.**

HEMORRHOID RELIEF

aesculus hippocastanum, arnica montana, calcarea fluorica, carduus marianus, collinsonia canadensis, hamamelis virginiana, muriaticum acidum, ratanhia spray

Product Information	oduct Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50845-0226		
Route of Administration	ORAL				

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
AESCULUS HIPPOCASTANUM FLOWER (UNII: KK0Z92II8M) (AESCULUS HIPPOCASTANUM FLOWER - UNII: KK0Z92II8M)	AESCULUS HIPPOCASTANUM FLOWER	1 [hp_X] in 1 mL	
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII: O80TY208ZW)	ARNICA MONTANA	3 [hp_X] in 1 mL	
CALCIUM FLUORIDE (UNII: O3B55K4YKI) (FLUORIDE ION - UNII:Q80VPU4080)	CALCIUM FLUORIDE	9 [hp_X] in 1 mL	
MILK THISTLE (UNII: U946SH95EE) (MILK THISTLE - UNII:U946SH95EE)	MILK THISTLE	1 [hp_X] in 1 mL	
COLLINSONIA CANADENSIS ROOT (UNII: O2630F3XDR) (COLLINSONIA CANADENSIS ROOT - UNII: O2630F3XDR)	COLLINSONIA CANADENSIS ROOT	3 [hp_X] in 1 mL	
HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK (UNII: T7S323PKJS) (HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK - UNII:T7S323PKJS)	HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK	3 [hp_X] in 1 mL	
HYDROCHLORIC ACID (UNII: QTT17582CB) (HYDROCHLORIC ACID - UNII:QTT17582CB)	HYDROCHLORIC ACID	6 [hp_X] in 1 mL	
KRAMERIA LAPPACEA ROOT (UNII: P29ZH1A35Z) (KRAMERIA LAPPACEA ROOT - UNII:P29ZH1A35Z)	KRAMERIA LAPPACEA ROOT	3 [hp_X] in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
ALCOHOL (UNII: 3K9958V90M)			

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:50845- 0226-1	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/30/2015	01/29/2025

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		11/30/2015	01/29/2025

Labeler - Liddell Laboratories, Inc. (832264241)

Registrant - Apotheca Company (844330915)

Establishment Name Address ID/FEI Business Operations Apotheca Company 844330915 manufacture(50845-0226) , api manufacture(50845-0226) , label(50845-0226) , pack(50845-0226)

Revised: 4/2022 Liddell Laboratories, Inc.