ARNICA MONTANA- arnica montana liquid

Newton 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.

Arnica montana

INDICATIONS & USAGE SECTION

Trauma, Injuries, Bruises, Exhaustion, Headache, Sprains, Wounds

DOSAGE & ADMINISTRATION SECTION

Directions: Ages 12 and up, take 6 drops by mouth (ages 0 to 11, give 3 drops) as needed or as directed by a health professional. Sensitive persons begin with 1 drop and gradually increase to full dose.

OTC - ACTIVE INGREDIENT SECTION

Arnica montana 15x, 10x, 200c, 30c.

OTC - PURPOSE SECTION

Trauma, Injuries, Bruises, Exhaustion, Headache, Sprains, Wounds

INACTIVE INGREDIENT SECTION

Inactive Ingredients: USP Purified Water; USP Gluten-free, non-GMO, organic cane alcohol 20%.

QUESTIONS SECTION

www.newtonlabs.net Newton Laboratories, Inc. FDA Est # 1051203 - Conyers, GA 30012 Questions? 1.800.448.7256

WARNINGS SECTION

Warning: Keep out of reach of children. Do not use if tamper-evident seal is broken or missing. If symptoms worsen or persist for more than a few days, consult a doctor. If **pregnant or breast-feeding**, ask a doctor before use.

OTC - PREGNANCY OR BREAST FEEDING SECTION

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

OTC - KEEP OUT OF REACH OF CHILDREN SECTION

Keep out of reach of children.

PACKAGE LABEL



Trauma; Injuries; Bruises; Exhaustion; Headache; Sprains; Wounds

^l fl oz (29.57 ml)

ARNICA MONTANA

arnica montana liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55714-6063
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Route of Administration ORAL

Active Ingredient/Active Moiety

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	Ingredient Name	Basis of Strength	Strength
	ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	15 [hp_X] in 1 mL

Inactive Ingredients

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

Packaging

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ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
ı	1 NDC:55714-6063-1	30 mL in 1 BOTTLE, GLASS			

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved homeopathic		09/01/2011			

Labeler - Newton Laboratories, Inc. (788793610)

Registrant - Newton Laboratories, Inc. (788793610)

Establishment					
Name	Address	ID/FEI	Business Operations		
Newton Laboratories, Inc.		788793610	manufacture(55714-6063)		

Revised: 12/2014 Newton Laboratories, Inc.