

**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.*

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**Arnica 9028L**

**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 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**

newtonlabs.net - Questions? 800.448.7256

Newton Laboratories, Inc. FDA Est # 1051203 - Conyers, GA 30013

**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



## ARNICA MONTANA

arnica montana liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDc:55714-9028
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ARNICA MONTANA</b> (UNII: 080TY208ZW) (ARNICA MONTANA - UNII:080TY208ZW)	ARNICA MONTANA	30 [hp_C] in 1 mL

## Inactive Ingredients

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

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55714-9028-1	30 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	01/28/2026	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/28/2026	

**Labeler** - Newton Laboratories, Inc. (788793610)

**Registrant** - Newton Laboratories, Inc. (788793610)

## Establishment

Name	Address	ID/FEI	Business Operations
Newton Laboratories, Inc.		788793610	manufacture(55714-9028)

Revised: 1/2026

Newton Laboratories, Inc.