

**GERMA ARNICA RELIEF- arnica montana gel**  
**Caribe Natural LLC**

*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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☐Active Ingredients

Purpose

Arnica Montana 1X HPUS (4%)

Pain, Bruises, Injuries, Muscle Aches

Pain, Bruises, Injuries, Muscle Aches

☐**Keep out of reach of children.**☐ If swallowed, get medical help or contact a Poison Control Center right away.

Germa Arnica Relief Gel

Homeopathic Medicine

Bruising & Swelling / Moreton e Hinchazon

Aches & Pains / Dolores y Molestia Muscular

Joints Pain / Dolores de Articulacion

Back Ache / Dolores de Espalda

Stiffness / Rigidez

Natural Pain Relief

☐**Directions:**

☐**For External Use Only.**☐ Apply generously to affected area, 2 to 3 times daily or as needed. Rub gently until complete absorption. Safe for children over 2 years old (under 2 years ask a Doctor).

☐**Warnings:**

**For External Use Only.** Ask a Doctor before use if pregnant, or breast-feeding. **When using this product:** Do not get into eyes and do not apply on open skin. **Stop use and ask a Doctor if:** Condition worsens, symptoms last more than 7 days, or you are allergic to any of the ingredients.

☐**Inactive Ingredients:** ☐aloe vera, caprylyl glycol, hexylene glycol, hydroxyethylcellulose, phenoxyethanol, potassium sorbate, purified water.



## GERMA ARNICA RELIEF

arnica montana gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59561-100
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	1 [hp_X] in 100 g

### Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
HYDROXYETHYL CELLULOSE (100 MPA.S AT 2%) (UNII: R33S7TK2EP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
WATER (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59561-100-02	57 g in 1 PACKAGE; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		06/23/2015	

**Labeler** - Caribe Natural LLC (624210480)

## Establishment

Name	Address	ID/FEI	Business Operations
Homeocare Laboratories Inc		088248828	manufacture(59561-100)

Revised: 6/2015

Caribe Natural LLC