

ARNICA PAIN RELIEF GEL WITH ALOE- arnica montana gel
HomeoCare Laboratories

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 Pain Relief Gel with Aloe

Active Ingredient:

Arnica Montana 1X HPUS (7%)

Inactive Ingredients:

Aloe vera, Caprylyl Glycol, Hexylene Glycol, Hydroxyethylcellulose, Phenoxyethanol, Potassium Sorbate, Purified water

Warnings:

For external use only. Ask a doctor before use if pregnant or breastfeeding. **Keep out of the reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

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, you are allergic to any of the ingredients.

Warnings:

Keep out of the reach of children.

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).

Questions? Comments?

Call 1-888-466-3622 www.homeocare.com

Manufactured & Distributed by:

Homeocare Laboratories, inc.

Yonkers, NY 10701

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Arnica Pain Relief Gel with Aloe

Bruises

Injuries

Muscle Aches

Pain Relief

Pre/Post Surgery

Arnica Pain Relief Gel with Aloe

Bruises

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Pre/Post Surgery

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NDC 61727-113-02

HOMEOCARE®

ARNICA

Pain Relief Gel



with Aloe

- Bruises
- Injuries
- Muscle Aches
- Pain Relief
- Pre/Post Surgery



Parabens Free

HOMEOPATHIC MEDICINE

NET WT 2 OZ (56.70g)

ARNICA PAIN RELIEF GEL WITH ALOE

arnica montana gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61727-113
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	56.7 g in 1 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
CETYL HYDROXYETHYLCELLULOSE (350000 MW) (UNII: T7SWE4S2TT)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61727-113-02	56.7 [hp_X] in 1 TUBE; Type 0: Not a Combination Product	12/21/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/21/2018	

Labeler - HomeoCare Laboratories (088248828)**Registrant** - HomeoCare Laboratories (088248828)**Establishment**

Name	Address	ID/FEI	Business Operations
HomeoCare Laboratories		088248828	manufacture(61727-113)

Revised: 12/2018

HomeoCare Laboratories