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 as k 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

When using this product:

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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 Injuries Muscle Aches Pain Relief

Pre/Post Surgery

Arnica Pain Relief Gel with Aloe

Active Ingredient: Arnica montana 1X HPUS(7%)

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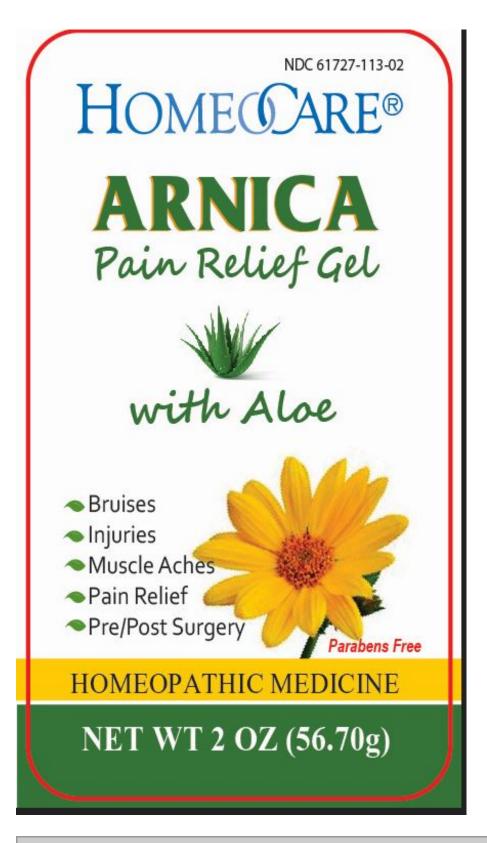
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ARNICA PAIN RELIEF GEL WITH ALOE

arnica montana gel

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

Active Ingredier	nt/Active Moiety				
	Ingredient Name	Basis of Streng	gth Strength		
ARNICA MONTANA	(UNII: 080TY208ZW) (ARNICA MONTANA - UNII:080TY20	8ZW) ARNICA MONTAI	NA 56.7 g in 1 [hp_X]		
Inactive Ingredi	ents				
	Ingredient Name		Strength		
CETYL HYDROXYETHYLCELLULOSE (350000 MW) (UNII: T7SWE4S2TT)					
PHENOXYETHANOL (UNII: HIE492ZZ3T)					
WATER (UNII: 059QF	0KO0R)				
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)					
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)					
HEXYLENE GLYCOL (UNII: KEH0A3F75J)					
CAPRYLYL GLYCOL (UNII: 00 YIU5438U)					
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 In	formation				
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved homeopa	hic	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