DERMFREE PAIN RELIEF- arnica montana hpus 7% pain relief cream
Jiangxi Hemei Pharmaceutical Co., Ltd

84010-016

Active Ingredient

Arnica montana HPUS7%

Purpose

Relieves muscle pain &stiffness swelling from injuries, discoloration from bruises

Use

temporarily relieves muscle pain and stiffness due to:minor injuriesmoverexertionmfalls
reduces symptoms of bruisingsuch as:mpainmswellingmdiscoloration

Warnings

For external use only

Do not use

Do not use if you are allergic to dermfreecream or to any of this product's inactive ingredients

When Using

When usingthis product avoid contact with eyes, mucous membranes, woundsdamaged or irritated skin.use only as directed dryness or irritationmay occur
do not tightly wrap or bandage the treated areado not apply heat or ice to treated areaimmediately beforeor after use

Stop Use

ocondition persists for more than 3 days or worsensosymptoms clear up and occur again within a few days.

Ask Doctor

ocondition persists for more than 3 days or worsensosymptoms clear up and occur again within a few days.

Keep Oot Of Reach Of Children

lfswallowed, get medical helpor contact a Poison Control Center right away

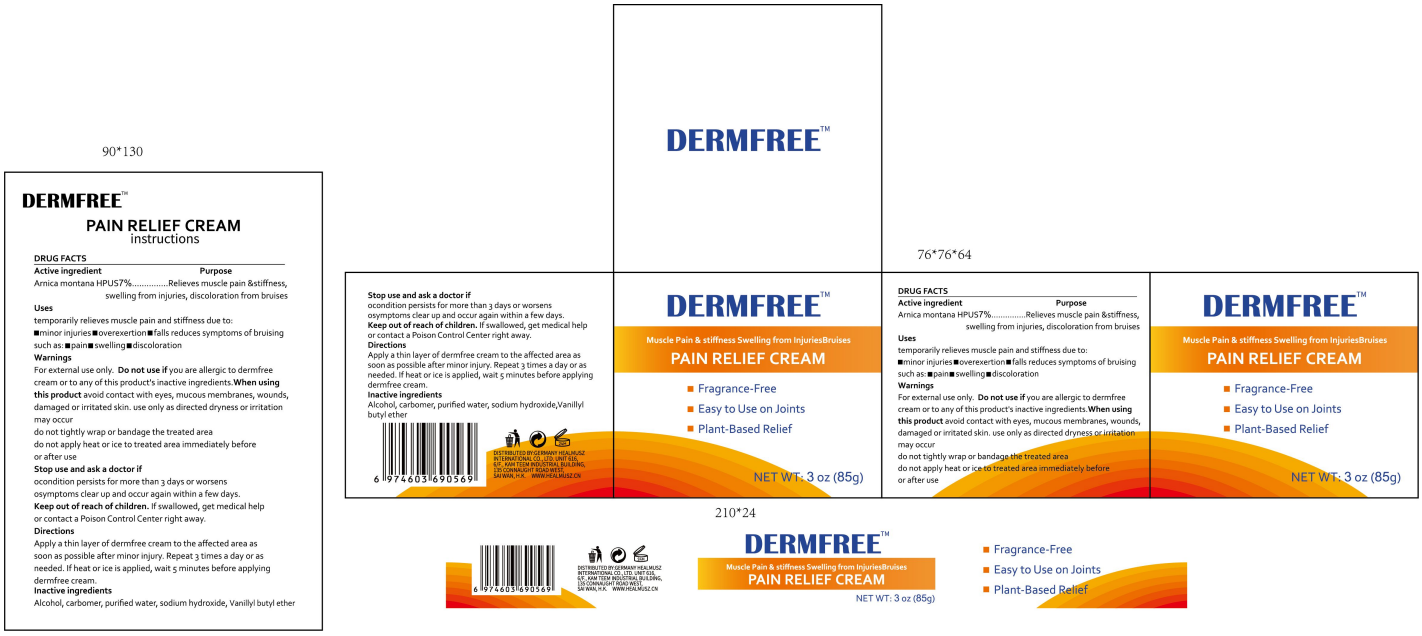
Directions

Apply a thin layer of dermfree cream to the affected area assoon as possible after minor injury. Repeat 3 times a day or asneeded.lf heat orice is applied, wait 5 minutes before applyingdermfree cream.

Inactive ingredients

Alcohol, carbomer, purifed water, sodium hydroxide, Vanillyl butyl ether

PRINCIPAL DISPLAY PANEL



DERMFREE PAIN RELIEF

arnica montana hpus 7% pain relief cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84010-016
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	7 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALCOHOL (UNII: 3K9958V90M)	
VANILLYL BUTYL ETHER (UNII: S2ULN37C9R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84010-016-01	85 g in 1 JAR; Type 0: Not a Combination Product	05/07/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/07/2024	

Labeler - Jiangxi Hemei Pharmaceutical Co., Ltd (724892056)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-016)

Revised: 5/2024 Jiangxi Hemei Pharmaceutical Co., Ltd