

BIO-RELIEF- menthol gel
Scientific Solutions Global LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

DRUG FACTS	
Active Ingredient: Menthol 10%	Purpose: Topical Analgesic
Uses: For temporary relief of minor aches and pains of muscles and joints associated with Simple backache, Arthritis, Strains, Sprains.	
Warning: For external use only. Flammable, keep away from fire/flame.	
When using this product : Avoid contact with eyes or mucous membranes. Do not bandage tightly. Do not apply to wound or damaged or irritated skin.	
Stop use and consult a doctor if : Conditions worsens. Symptoms persist for more than 7 days. Clear up and occur again within a few days.	
Keep out of reach of children, if swallowed, get medical help or contact a Poison Control Center right away.	
If pregnant or breast-feeding, ask a health professional before use.	
Directions: Adults & children 2 yrs of age and older: Apply to affected area not more than 4 times daily in one direction, not circular. Children under 2 years of age, consult a physician. Massage not necessary. Shake well before use.	
Other Information: Store in a cool and dry place with lid closed tightly.	
Inactive Ingredients: Aloe Vera Oil, Arnica Flower Tincture, Boswellia Powder, Camphor, Frankincense Oil, Green Tea, Hydroxypropyl Cellulose NF, Isopropyl Alcohol, Lavender Oil, Propylene Glycol, Tea Tree Oil, Thymol, Vitamin E.	
Questions or Comments? E-mail: info@scisolglobal.com	
Manufactured and Distributed by: Scientific Solutions Global LLC, Carle Place, NY, USA	



Menthol 10%

Topical Analgesic

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BIO-RELIEF menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71718-200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	8.9 g in 89 mL

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
ARNICA MONTANA (UNII: O80TY208ZW)	
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	
TEA TREE OIL (UNII: VIF565UC2G)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
THYMOL (UNII: 3J50XA376E)	

Product Characteristics

Color	white (Opaque gel)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71718-200-01	89 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/01/2020	

Labeler - Scientific Solutions Global LLC (097291290)**Registrant** - Scientific Solutions Global LLC (097291290)

Establishment

Name	Address	ID/FEI	Business Operations
Scientific Solutions Global LLC		097291290	manufacture(71718-200)

Revised: 2/2020

Scientific Solutions Global LLC