

MUSCLE RUB PAIN RELIEVER GEL- menthol, unspecified form gel
Universal Distribution Centre LLC

Pain Reliever Gel

Drug Facts

Active ingredient

Menthol 2.5%

Purpose

Topical analgesic

Uses

temporarily relieves the minor aches and pains of muscles and joints associated with:

- simple backache
- arthritis
- strains
- bruises
- sprains

Warnings

For external use only.

Do not use

- on wounds or damaged skin
- with a heating pad
- on a child under 12 years of age with arthritis-like conditions

Ask a doctor before use if you have redness over the affected area

When using this product

- avoid contact with eyes or mucous membranes
- do not bandage tightly

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days
- excessive skin irritation occurs

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

Directions

- adults and children 12 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 12 years of age: ask a doctor

Other information

store at 20° to 25°C (68° to 77°F)

Inactive ingredients

camphor, carbomer, DMDM hydantoin, isoceteth-20, isopropyl alcohol, PEG-40 hydrogenated castor oil, sodium hydroxide, water

Questions?

call **1-800-223-0182**(toll-free) or **215-273-8755**(collect)

Distributed by:

**JOHNSON & JOHNSON
CONSUMER INC.**

Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 57 g Tube Carton

86071 1.25OZ MUSCLE RUB - 24



Ingredient Name		Basis of Strength	Strength	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)		MENTHOL, UNSPECIFIED FORM	25 mg in 1 g	
Inactive Ingredients				
Ingredient Name			Strength	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)				
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)				
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)				
ISOCETETH-20 (UNII: O020065R7Z)				
ISOPROPYL ALCOHOL (UNII: ND2M416302)				
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-072-04	1 in 1 CARTON	01/24/2021	
1	NDC:52000-072-03	35.4 g in 1 TUBE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	01/24/2021		

Labeler - Universal Distribution Centre LLC (019180459)

Registrant - Sawy Care & Cosmetics Pvt. Ltd. (915039748)

Establishment

Name	Address	ID/FEI	Business Operations
Sawy Care & Cosmetics Pvt. Ltd.		915039748	manufacture(52000-072)

Revised: 5/2024

Universal Distribution Centre LLC