

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

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**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:

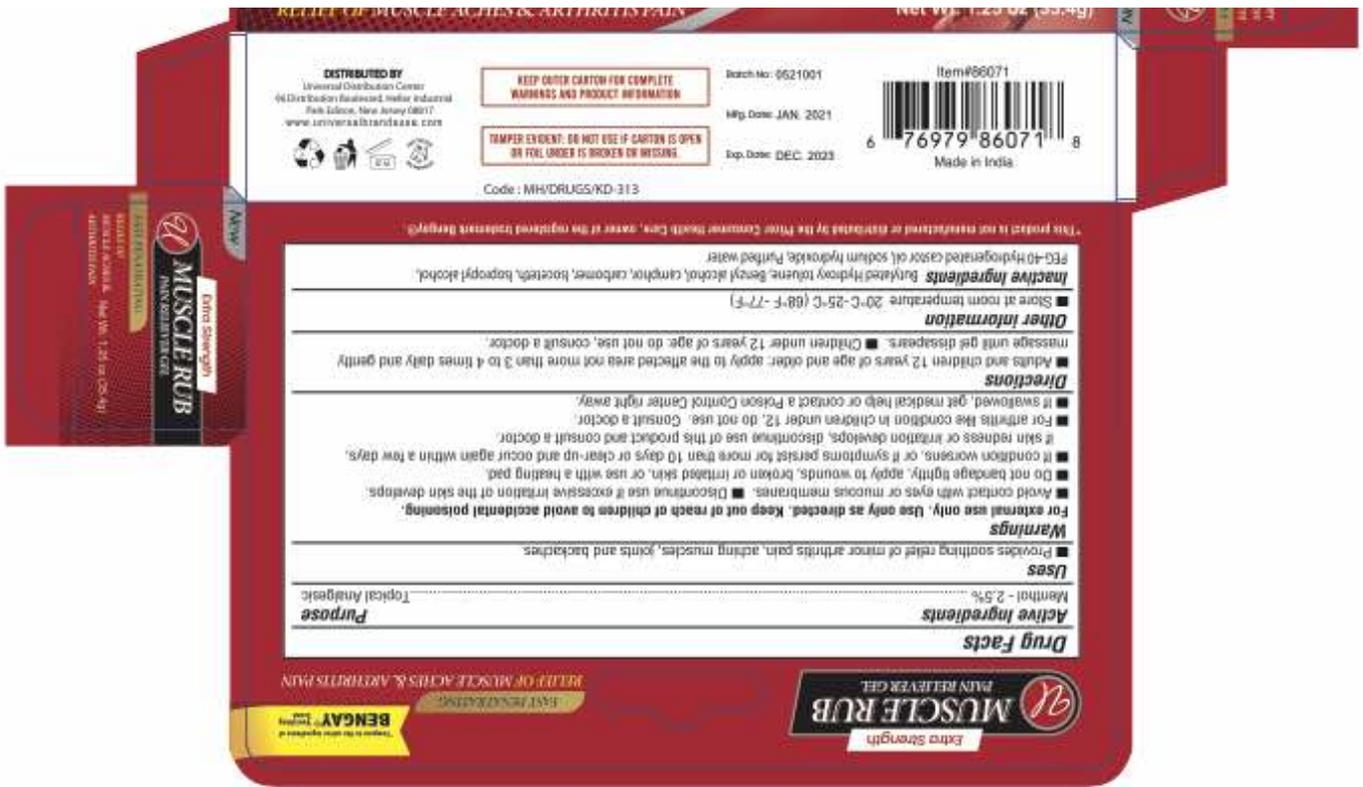
**JOHNSON & JOHNSON  
CONSUMER INC.**

Skillman, NJ 08558

## PRINCIPAL DISPLAY PANEL - 57 g Tube Carton

86071 1.25OZ MUSCLE RUB - 24





**CYAN MAGENTA YELLOW BLACK**

**Size : 60 X 30 X 40 X 100 X 153 MM**

Extra Strength Muscle Rub

Pain Reliever Gel

Fast Penetrating

**MENTHOL 2.5% TOPICAL ANALGESIC GEL**

**NET WT 1 OZ (35.4 g)**

## MUSCLE RUB PAIN RELIEVER GEL

menthol, unspecified form gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:52000-072
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
<b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)		MENTHOL, UNSPECIFIED FORM	25 mg in 1 g	
<b>Inactive Ingredients</b>				
Ingredient Name			Strength	
<b>BUTYLATED HYDROXYTOLUENE</b> (UNII: 1P9D0Z171K)				
<b>CAMPHOR (SYNTHETIC)</b> (UNII: 5TJD82A1ET)				
<b>CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE</b> (UNII: 0A5MM307FC)				
<b>ISOCETETH-20</b> (UNII: O020065R7Z)				
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302)				
<b>POLYOXYL 40 HYDROGENATED CASTOR OIL</b> (UNII: 7YC686GQ8F)				
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)				
<b>WATER</b> (UNII: 059QF0KO0R)				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-072-02	1 in 1 CARTON	01/24/2021	
1	NDC:52000-072-01	35.4 g in 1 TUBE; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
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)

Revised: 12/2024

Universal Distribution Centre LLC