

RALLI RAPID PAIN RELIEF- menthol gel
Troy Manufacturing, Inc.

Ralli Rapid Pain Relief Gel

Drug Facts

Active Ingredient

Menthol 10.0%

Purpose

Topical Analgesic

Uses

For the temporary relief of minor aches and pains of muscles and joints associated with: • simple backache • arthritis • strains • bruises • sprains

Warnings

For external use only. Avoid contact with eyes.

Flammable: Keep away from fire or flame.

When using this product

• use only as directed • do not bandage tightly or use a heating pad • do not apply to wounds or damaged skin

Stop use and ask a doctor if

• condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days • redness is present • excessive irritation of the skin develops

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults and children 2 years of age and older: • apply directly onto affected area without the need to bandage • repeat if necessary, but do not apply more than 3 to 4 times daily.

Children under 2 Years of age: do not use, consult a doctor.

Other Information

- Store at room temperature

Inactive Ingredients

acrylic acid/vinyl ester copolymer, dimethylsulfone (MSM), eucalyptus oil, glucosamine sulfate, hydroxypropylcellulose, PEG-8 dimethicone, pentylene glycol, peppermint oil, SD alcohol 39C, triethanolamine, water (USP).

Package Labeling:

rallipainrelief.com

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Manufactured for:
Wellness Formulations, LLC
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Hazleton, PA 18202
1-866-323-4285



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ralli™

Bounce Back from Pain

RAPID PAIN RELIEF

10%
ACTIVE MENTHOL

GEL

3 FL OZ (88ML)

RALLI RAPID PAIN RELIEF

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63936-9752
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ACRYLIC ACID (UNII: J94PBK7X8S)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
PEG-8 DIMETHICONE (600 CST) (UNII: GIA7T764OD)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KOOR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63936-9752-1	88 mL in 1 TUBE; Type 0: Not a Combination Product	02/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/01/2024	

Labeler - Troy Manufacturing, Inc. (160075248)

Revised: 2/2024

Troy Manufacturing, Inc.