PAINGEL ROLL-ON- menthol gel
NACOR BUSINESS GROUP LLC

85110-003-03

Menthol 3%

Topical Analgesic

Uses

For temporary relief of minor aches and pains in muscles and joints.

- **For external use only.**
- Ask a doctor before use if you have redness over affected area.

- Avoid contact with the eyes or mucous membranes.
- Do not apply to wounds or damaged skin.
- Do not apply to the irritated skin or if excessive irritation develops.
- Do not bandage tightly.
- Do not use with heating pad or device.
- 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.

If pregnant or breast-feeding

Ask a health professional before use.

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

Directions

Adults and children over 12 years of age

- Apply a small amount on the affected area.
- Massage in circular motion, let set for a few seconds.
- Repeat as necessary, but no more than 3 to 4 times daily.

Children under 12 years of age. Do not use, consult a doctor.

Other information

Store tightly closed in a dry place at controlled room temperature between 59°F-86° F(15°C-30° C).

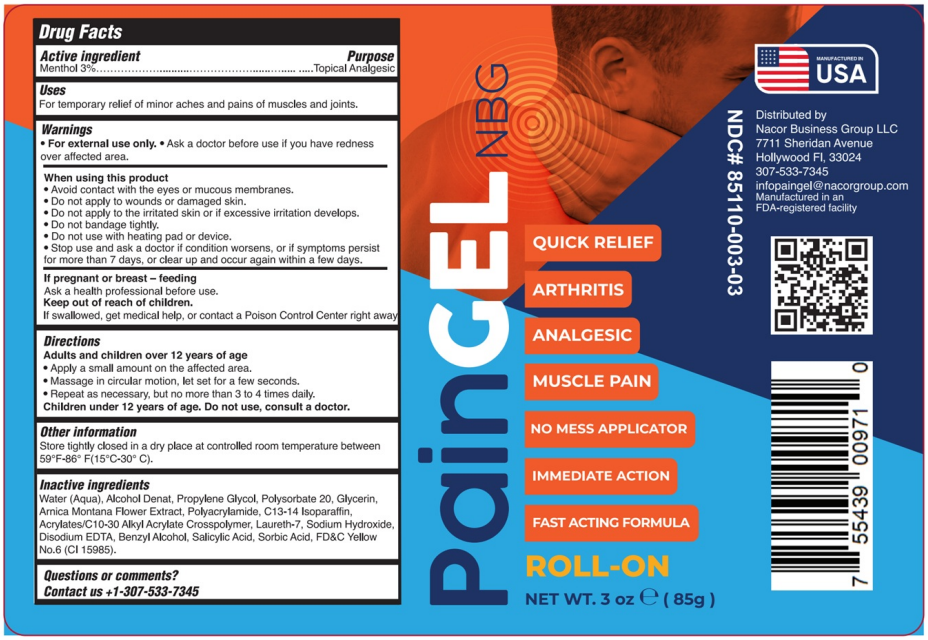
Water (Aqua), Alcohol Denat, Propylene Glycol, Polysorbate 20, Glycerin, Arnica Montana Flower Extract, Polyacrylamide, C13-14 Isoparaffin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Laureth-7, Sodium Hydroxide, Disodium EDTA, Benzyl Alcohol, Salicylic

Acid, Sorbic Acid, FD&C Yellow No.6 (CI 15985).

Questions or comments?

Contact us +1-307-533-7345

PAINGEL ROLL-ON



PAINGEL ROLL-ON

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85110-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	3 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
GLYCERIN (UNII: PDC6A3C0OX)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	

POLYACRYLAMIDE (1500 MW) (UNII: 5D6TC4BRWV)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S AT 1.0%) (UNII: 8Z5ZAL5H3V)	
LAURETH-7 (UNII: Z95S6G8201)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
SORBIC ACID (UNII: X045WJ989B)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85110-003-03	85 g in 1 BOTTLE; Type 0: Not a Combination Product	11/28/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/28/2025	

Labeler - NACOR BUSINESS GROUP LLC (119438840)

Revised: 11/2025 NACOR BUSINESS GROUP LLC