PAINGEL- menthol, methyl salicylate spray
NACOR BUSINESS GROUP LLC

85110-301-04

Menthol 10.5%

Methyl Salicylate 30%

External Analgesic

USE

For temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains, bruises and sprains.

- For external use only.
- Do not apply to wounds or damaged skin, with a heating pad or on a child under 12 years of age.

Flammable

Keep away from fire or flame.

- Use only as directed.
- Do not bandage tightly.
- Avoid contact with eyes and mucous membranes.
- If you experience an allergic reaction, discontinue use, and consult a doctor.
- Do not expose the area treated with product to heat or direct sunlight.

Stop use and ask a doctor if

- Condition worsens.
- Redness is present.
- Irritation develops.
- Symptoms persist for more than 7 days or clear up occur again within a few days.
- You experience signs of injury, such as pain, swelling or blistering where the product was applied.

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

Spray two to three pumps or as necessary to cover the affected area not more than

three to four 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).

Alcohol Denat, Water (Aqua), Glycerin, Propylene Glycol, Aloe Barbadensis (Aloe Vera) Leaf Juice.

Questions or comments?

Contact us +1-307-533-7345

PAINGEL SPRAY

PAINGEL

menthol, methyl salicylate spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85110-301
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 g in 100 mL
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID -		28.5 g

METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	28.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85110-301-04	118 mL in 1 BOTTLE, SPRAY; 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