

PAINGEL- menthol, methyl salicylate spray
NACOR BUSINESS GROUP LLC

85110-207-04

Menthol 7%

Methyl Salicylate 20%

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

Pain GEL NBG

Pain GEL SPRAY

QUICK RELIEF

ARTHRITIS

ANALGESIC

MUSCLE PAIN

4 fl oz e (118 mL)

NDC# 85110-207-04

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USA

Distributed by
Nacor Business Group LLC
7711 Sheridan Avenue
Hollywood Fl, 33024
307-533-7345
infoaingel@nacorgroup.com
Manufactured in an
FDA-registered facility

Drug Facts

Active ingredient	Purpose
Menthol 7 %	External Analgesic
Methyl Salicylate 20%	External Analgesic

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

Warnings
• 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.

When using this product
• Use only as directed.
• Do not bandage tightly.
• Avoid contact with eyes and mucous membranes.
• Do not apply to wounds or damaged, broken or irritated skin.
• 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.

Keep out of reach of children.
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, 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).

Inactive ingredients
Alcohol Denat, Water (Aqua), Glycerin, Propylene Glycol, Aloe Barbadensis (Aloe Vera) Leaf Extract.

Questions or comments?
Contact us +1-307-533-7345

PAINGEL

menthol, methyl salicylate spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85110-207	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)		MENTHOL	7 g in 100 mL	
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)		METHYL SALICYLATE	20 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:85110-207-04	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: 2/2026

NACOR BUSINESS GROUP LLC