2.5% MENTHOL PAIN RELIEF- menthol gel
Universal Distribution Center LLC

2.5% Menthol PAIN RELIEF GEL

Drug facts

Active ingredient (in each gram)

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 immediately.

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

Cinnamomum camphora(camphor) bark oil, Carbomer, Ethylhexylglycerin, Glycerin, Isopropyl Alcohol, Methylparaben, Polysorbate 20, Propylparaben, Purified Water, Sodium Hydroxide.

COOLING MENTHOL FORMULA

✓ Relives back, neck & joint pain

✓ Fast-absorbing & non-greasy

Soothes sore muscles

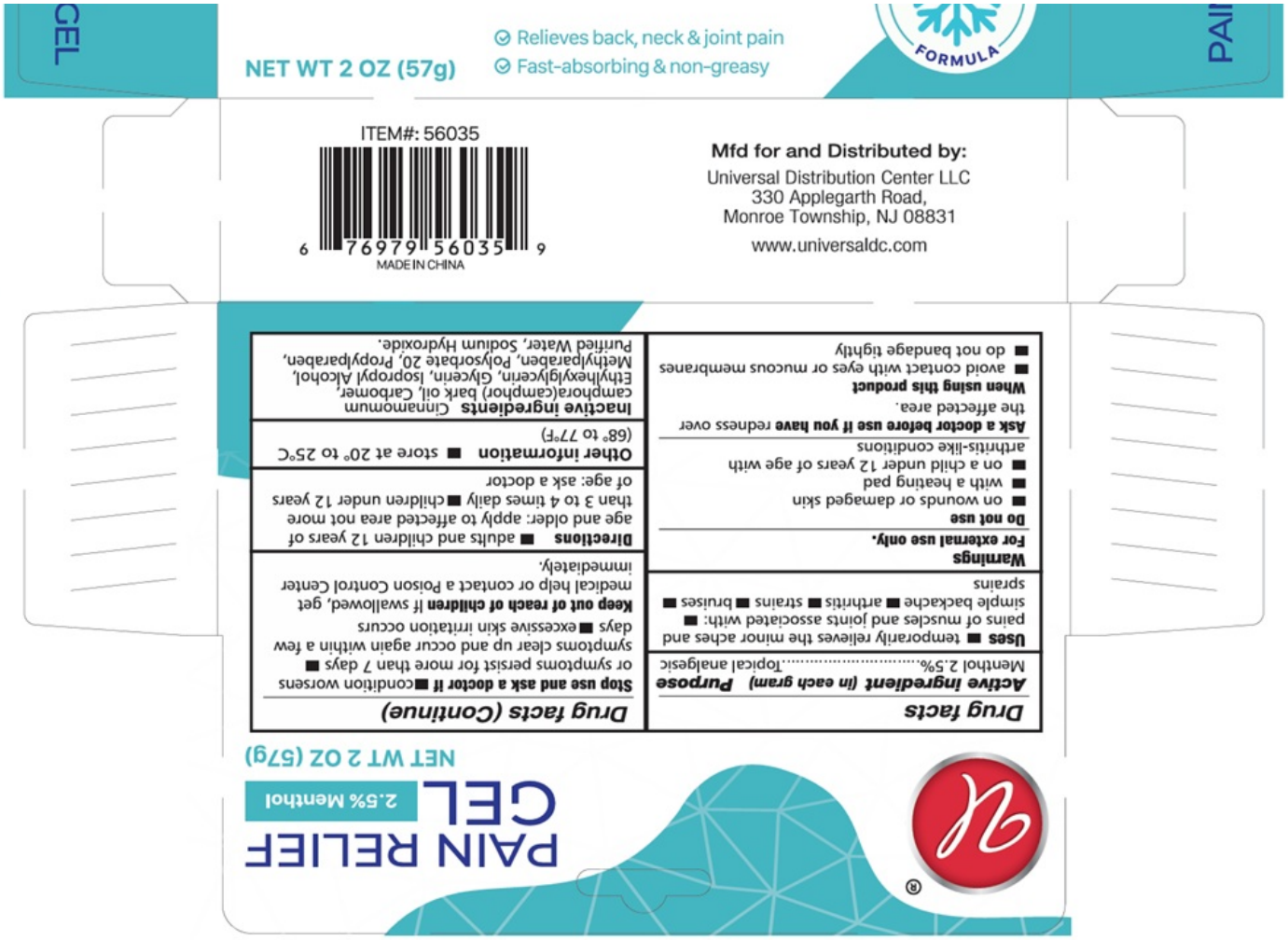
Mfd for and Distributed by:
Universal Distribution Center LLC
330 Applegarth Road,
Monroe Township, NJ 08831

www.universaldc.com

MADE IN CHINA

Packaging

Outer Package label



Inner Package Label

2.5% MENTHOL PAIN RELIEF

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-440
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	25 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
CAMPHOR OIL (UNII: 75IZZ8Y727)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52000-440-02	1 in 1 BOX	01/19/2026	
1		57 g in 1 TUBE; Type 0: Not a Combination		

1		Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/19/2026	

Labeler - Universal Distribution Center LLC (019180459)

Revised: 1/2026 Universal Distribution Center LLC