PHARMACYS PRESCRIPTION ANALGESIC GEL- menthol gel
American Consumer Products Corp

Pharmacy's Prescription 8OZ Ice Cold Analgesic Gel

Active Ingredient

Active Ingredient: Menthol 1%

Purpose

Purpose: Pain relieving gel

Warnings

Warnings: For external use only

Stop Use

Stop use and ask doctor if

- condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children

Keep our of reach of children. If swallowed get medical help or contact a Poison Control Center immediately.

Inactive Ingredients

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carbomer, isopropyl alcohol, nonyl phenyl polyoxyethylene ether, camphor, kathon CG, FD&C blue no. 1, triethanolamine, water

Indications & Usage Section

Uses: For the temporary relief of minor aches and pains of muscles and joints associated with - simple backache - arthritis - strains - bruses - sprains

When using this product

- avoid contact with eyes

- do not bandage tightly

- do not apply to wounds or damaged skin

- do not use with heating pads or other heating devices

Dosage & Administration

Directions

- adults and childrent 2 years of age and older: apply to affected area not more than 3 to 4 times daily

- children under 2 years of age: do not use, consult a doctor

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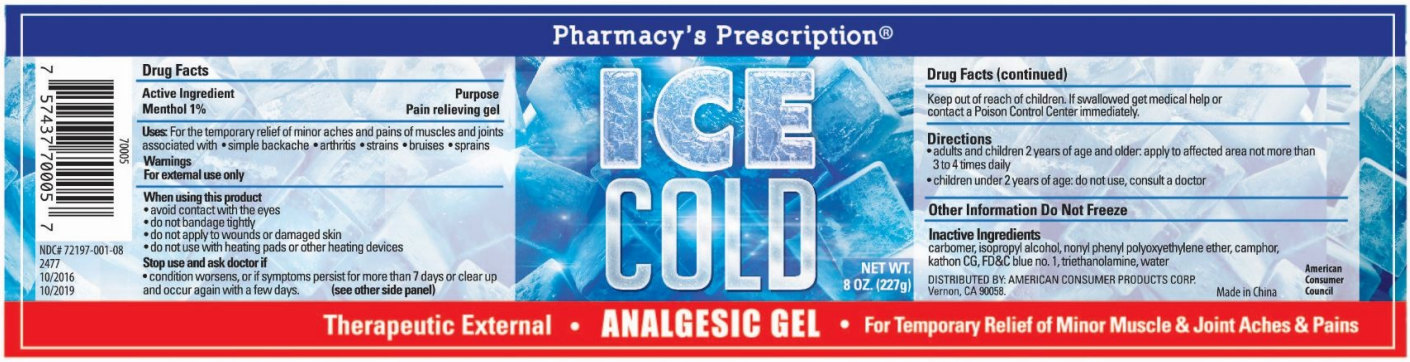

PHARMACYS PRESCRIPTION ANALGESIC GEL

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72197-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
CAMPHOR, (-)- (UNII: 213N3S8275)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
METHYLCHLOROISOTHIAZOLINONE/METHYLISOTHIAZOLINONE MIXTURE (UNII: 15O9QS218W)	
2,2',3,3',4,4',5,5',6-NONACHLORODIPHENYL ETHER (UNII: 4S0765P9W8)	
TRIETHANOLAMINE LAURYL SULFATE (UNII: E8458C1KAA)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72197-001-08	227 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/31/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	08/31/2018	

Labeler - American Consumer Products Corp (081101181)

Revised: 1/2025

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