**PHARMACYS PRESCRIPTION ANALGESIC GEL- menthol gel**
**American Consumer Products Corp**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**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**

**Inactive Ingredients**

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

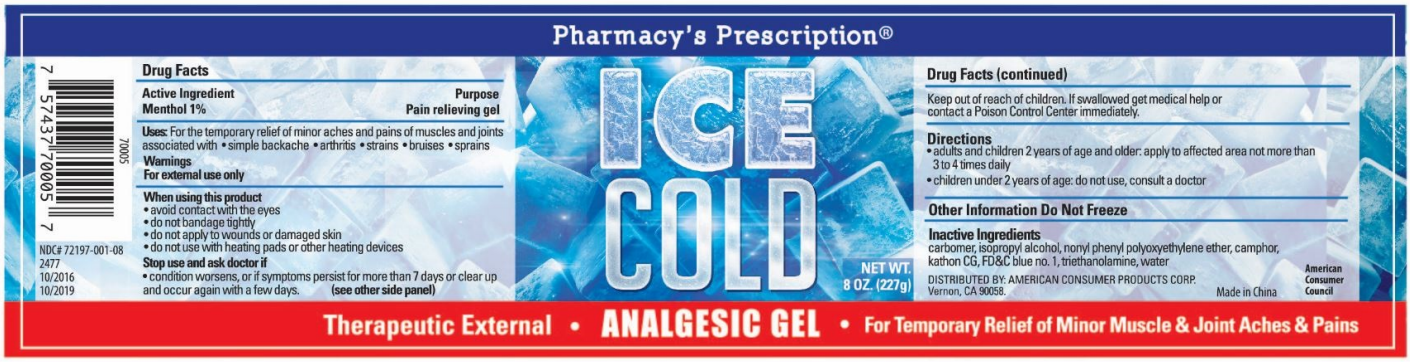
## Dosage & Administration

## 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

## Pharmacy's Prescription 8 OZ Ice Cold Analgesic Gel




# 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 final | part346 | 08/31/2018 | |

## Labeler - American Consumer Products Corp (081101181)

Revised: 5/2023

American Consumer Products Corp