

**DOLLAR GENERAL COLD-HOT MEDICATED PATCHES- menthol patch**  
**DOLGENCORP, LLC**

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**Dollar General Cold-Hot Medicated Patches**

**ACTIVE INGREDIENT**

| Active Ingredient ..... | Purpose |
|-------------------------|---------|
| Menthol 5%.....         | Topical |
| Analgesic               |         |

**INACTIVE INGREDIENT**

carboxymethylcellulose sodium, Dihydroxyaluminum Aminoacetate, Glycerin, iodopropynyl butylcarbamate, Kaolin, Mineral Oil, Petrolatum, Phenoxyethanol, Polyacrylic Acid, Polysorbate 80, povidone, Propylene Glycol, Sodium Polyacrylate, Tartaric Acid, Titanium Dioxide, Water, 3-(2-ethylhexyloxy)propane-1,2-diol

**KEEP OUT OF REACH OF CHILDREN**

If swallowed, get medical help or contact a Poison Control

**INDICATIONS & USAGE**

Temporarily relieves minor pain associated with: ■ arthritis ■ simple backache ■ bursitis ■ tendonitis ■ muscle strains ■ muscle sprains ■ bruises ■ cramps

**WARNINGS**

**For External Use Only.**

**DOSAGE & ADMINISTRATION**

Adults and children over 12 years: Carefully remove backing from patch. Apply sticky side of patch to affected area.

Wear one patch up to 8 hours. Repeat as necessary, but no more than 4 times daily. Reseal pouch after opening.

Children 12 years or younger: Consult a physician.

**PURPOSE**

Topical Analgesic

## When using this product

Use only as directed ■ Don't bandage tightly or use with heating pad

■ Avoid contact with eyes and mucous membranes ■ Don't apply to wounds or damaged skin.

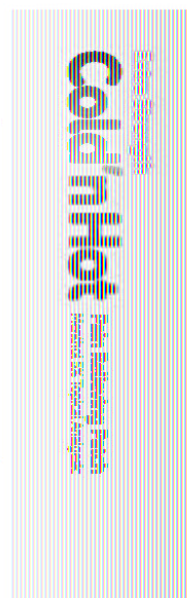
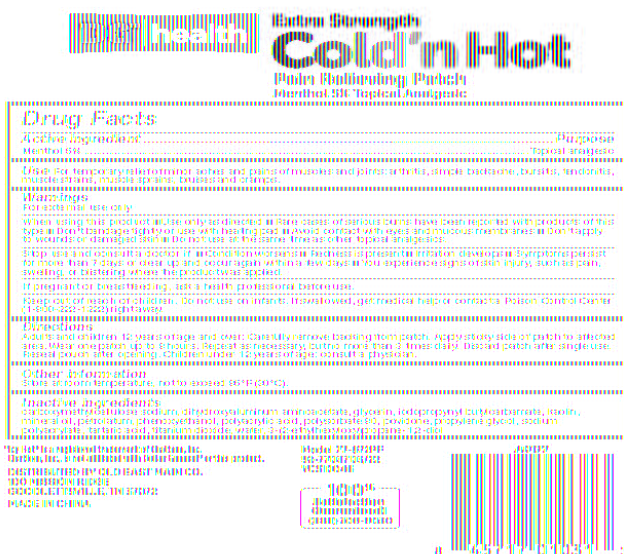
## Stop use and ask a doctor

If condition worsens ■ If redness is present ■ If irritation develops

■ If symptoms persist for more than 7 days or clear up and occur again within a few days.

## If pregnant or breastfeeding

ask a health professional before use.



## DOLLAR GENERAL COLD-HOT MEDICATED PATCHES

menthol patch

### Product Information

| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:55910-972 |
|-------------------------|----------------|--------------------|---------------|
| Route of Administration | TOPICAL        |                    |               |

| Active Ingredient/Active Moiety                               |                   |          |
|---------------------------------------------------------------|-------------------|----------|
| Ingredient Name                                               | Basis of Strength | Strength |
| <b>MENTHOL</b> (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) | MENTHOL           | 5 g      |

| Inactive Ingredients                                     |          |
|----------------------------------------------------------|----------|
| Ingredient Name                                          | Strength |
| <b>PETROLATUM</b> (UNII: 4T6H12BN9U)                     |          |
| <b>MINERAL OIL</b> (UNII: T5L8T28FGP)                    |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                          |          |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                       |          |
| <b>SODIUM POLYACRYLATE (8000 MW)</b> (UNII: 285CYO341L)  |          |
| <b>DIHYDROXYALUMINUM AMINOACETATE</b> (UNII: DO250MG0W6) |          |
| <b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)             |          |
| <b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)                 |          |
| <b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)               |          |
| <b>KAOLIN</b> (UNII: 24H4NWX5CO)                         |          |
| <b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)                 |          |
| <b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)               |          |
| <b>CARBOXYMETHYLCELLULOSE</b> (UNII: 05JZI7B19X)         |          |
| <b>TARTARIC ACID</b> (UNII: W4888I119H)                  |          |
| <b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)          |          |
| <b>POLYACRYLIC ACID (8000 MW)</b> (UNII: 73861X4K5F)     |          |
| <b>IODOPROPYNYL BUTYLCARBAMATE</b> (UNII: 603P14DHEB)    |          |

| Packaging |                  |                                                 |                      |                    |
|-----------|------------------|-------------------------------------------------|----------------------|--------------------|
| #         | Item Code        | Package Description                             | Marketing Start Date | Marketing End Date |
| 1         | NDC:55910-972-06 | 6 in 1 BOX                                      | 10/01/2017           |                    |
| 1         |                  | 1 in 1 POUCH; Type 0: Not a Combination Product |                      |                    |

| Marketing Information |                                          |                      |                    |
|-----------------------|------------------------------------------|----------------------|--------------------|
| Marketing Category    | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC Monograph Drug    | M017                                     | 10/01/2017           |                    |

**Labeler** - DOLGENCORP, LLC (068331990)

| Establishment                      |         |           |                        |
|------------------------------------|---------|-----------|------------------------|
| Name                               | Address | ID/FEI    | Business Operations    |
| Foshan Aqua Gel Biotech Co., Ltd., |         | 529128763 | manufacture(55910-972) |

