**RITE AID COLD AND HOT MEDICATED- menthol patch**
**Rite Aid Corporation**

*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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**Drug Facts**

**Active ingredient**

Menthol 5%

**Purpose**

Menthol - Topical Analgesic

**Uses**

temporarily relieves minor aches and pains of muscles and joints due to

- arthritis
- bursitis
- simple backache
- tendonitis
- strains
- bruises
- sprains
- cramps

**Warnings**

**For external use only**

**When using this product**

- use only as directed
- avoid contact with eyes or on mucous membranes
- do not apply to wounds or to damaged or very sensitive skin
- do not bandage tightly or use a heating pad

**Stop use and ask a doctor if**

- excessive redness or irritation is present
- condition worsens
- pain persists for more than 7 days
- symptoms clear up and occur again within a few days

**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 12 years and over: apply patch to affected area as needed but no more than 4 times daily
- children under 12 years: ask a doctor
- for easy application: grasp both ends of pad firmly, pull at both ends. Stretch pad until backing separates. Remove protective film while applying pad directly to site of pain.

**Inactive ingredients**

carbomer homopolymer, carboxymethylcellulose sodium, castor oil, dihydroxyaluminum aminoacetate, edetate disodium, glycerin, hydroxypropyl cellulose, kaolin, partially neutralized polyacrylate, polyvinyl alcohol, purified water, sorbitol solution, tartaric acid

**Package/Label Principal Display Panel**


| **RITE AID COLD AND HOT MEDICATED** |
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| menthol patch |
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## Product Information

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|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:11822-3729 |
| **Route of Administration** | TOPICAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **MENTHOL, UNSPECIFIED FORM** (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A) | MENTHOL, UNSPECIFIED FORM | 50 g |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED)** (UNII: HHT01ZNK31) | |
| **CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM** (UNII: K679OBS311) | |
| **CASTOR OIL** (UNII: D5340Y2I9G) | |
| **DIHYDROXYALUMINUM AMINOACETATE** (UNII: DO250MG0W6) | |
| **EDETATE DISODIUM** (UNII: 7FLD91C86K) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **HYDROXYPROPYL CELLULOSE, UNSPECIFIED** (UNII: 9XZ8H6N6OH) | |
| **KAOLIN** (UNII: 24H4NWX5CO) | |
| **POLYACRYLIC ACID (250000 MW)** (UNII: 9G2MAD7J6W) | |
| **POLYVINYL ALCOHOL, UNSPECIFIED** (UNII: 532B59J990) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SORBITOL** (UNII: 506T60A25R) | |
| **TARTARIC ACID** (UNII: W4888I119H) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:11822-3729-7 | 1 in 1 CARTON | 06/29/2010 | |
| **1** | | 5 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 not final | part348 | 06/29/2010 | |

## Labeler - Rite Aid Corporation (014578892)

Revised: 2/2023 Rite Aid Corporation