# EQUATE PAIN RELIEVING ULTRA STRENGTH- camphor, menthol, methyl salicylate cream Wal-Mart Stores Inc

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 ingredients**

Camphor 4%

Menthol 10%

Methyl salicylate 30%

## **Purpose**

Camphor - Topical analgesic

Menthol - Topical analgesic

Methyl salicylate - Topical analgesic

#### Uses

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

- simple backache
- arthritis
- strains
- bruises
- sprains

## **Warnings**

# For external use only

# When using this product

- use only as directed
- do not get into eyes or on mucous membranes
- do not apply to wounds or damaged skin
- do not bandage tightly
- do not use with heating pad or apply external heat

# Stop use and ask a doctor if

- skin redness or excessive irritation of the skin occurs
- condition worsens

 symptoms persist for more than 7 days or 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 to avoid accidental ingestion.

If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

- adults and children 12 years and over: apply to affected area not more than 3 to 4 times daily
- children under 12 years: ask a doctor

## **Inactive ingredients**

aloe barbadensis leaf extract, carbomer 940, cocos nucifera (coconut) oil, edetate disodium, glyceryl stearate, isopropyl myristate, lanolin, mineral oil, polysorbate 60, purified water, sodium lauryl sulfate, stearic acid, trolamine, vitamin E

## **Questions?**

Call: 1-888-287-1915

Package/Label Principal Display Panel



# **EQUATE PAIN RELIEVING ULTRA STRENGTH**

camphor, menthol, methyl salicylate cream

|   |   |   |   | -  |   | - |    |   |   | _  |  |
|---|---|---|---|----|---|---|----|---|---|----|--|
| m | - | m | м | 73 | n | - | ct | П | 9 | 96 |  |
|   |   |   |   |    |   |   |    |   |   |    |  |

Product Type HUMAN OTC DRUG Item Code (Source) NDC:49035-700

| Active Ingredient/Active Moiety  |                              |                  |  |  |  |  |
|--|------------------------------|------------------|--|--|--|--|
| Ingredient Name  | <b>Basis of Strength</b>     | Strength         |  |  |  |  |
| CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)             | CAMPHOR (SYNTHETIC)          | 40 mg<br>in 1 g  |  |  |  |  |
| MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A) | MENTHOL,<br>UNSPECIFIED FORM | 100 mg<br>in 1 g |  |  |  |  |
| METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:0414PZ4LPZ)                    | METHYL SALICYLATE            | 300 mg<br>in 1 g |  |  |  |  |

| Inactive Ingredients   |          |  |  |  |  |
|--|----------|--|--|--|--|
| Ingredient Name  | Strength |  |  |  |  |
| ALOE VERA LEAF (UNII: ZY81Z83H0X)  |          |  |  |  |  |
| CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E) |          |  |  |  |  |
| COCONUT OIL (UNII: Q9L0O73W7L)   |          |  |  |  |  |
| EDETATE DISODIUM (UNII: 7FLD91C86K)  |          |  |  |  |  |
| GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)   |          |  |  |  |  |
| ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)   |          |  |  |  |  |
| LANOLIN (UNII: 7EV65EAW6H)   |          |  |  |  |  |
| MINERAL OIL (UNII: T5L8T28FGP)   |          |  |  |  |  |
| POLYSORBATE 60 (UNII: CAL22UVI4M)  |          |  |  |  |  |
| WATER (UNII: 059QF0KO0R)   |          |  |  |  |  |
| SODIUM LAURYL SULFATE (UNII: 368GB5141J)   |          |  |  |  |  |
| STEARIC ACID (UNII: 4ELV7Z65AP)  |          |  |  |  |  |
| TROLAMINE (UNII: 903K93S3TK)   |          |  |  |  |  |
| .ALPHATOCOPHEROL (UNII: H4N855PNZ1)  |          |  |  |  |  |

| Packaging        |  |                         |                       |  |  |  |  |
|------------------|--|-------------------------|-----------------------|--|--|--|--|
| # Item Code      | Package Description                                  | Marketing Start<br>Date | Marketing End<br>Date |  |  |  |  |
| 1 NDC:49035-700- | 1 in 1 CARTON  | 06/13/2011              |                       |  |  |  |  |
| 1                | 113.3 g in 1 TUBE; Type 0: Not a Combination Product |                         |                       |  |  |  |  |

| Marketing Information   |   |                         |                       |  |  |  |
|-------------------------|---|-------------------------|-----------------------|--|--|--|
| Marketing<br>Category   | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |  |  |  |
| OTC monograph not final | part348                                     | 06/13/2011              |                       |  |  |  |
|                         |   |                         |                       |  |  |  |

Revised: 2/2023 Wal-Mart Stores Inc