**MEDIRELIEF ULTRA- menthol patch**
**FARLA MEDICAL HEALTHCARE LTD**

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**MediRelief Ultra Strength Menthol 5% Patch**

***Drug Facts***

**Active Ingredients & Purpose**

| ***Active ingredients (in each patch)*** | ***Purpose*** |
|---|---|
| Menthol 5% | Topical analgesic |

**Uses**

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

**Warnings**

**FOR EXTERNAL USE ONLY. Avoid contact with the eyes.**

**Do not use**

- on wounds or damaged skin
- on a tight bandage
- with a heating pad
- on sensitive skin
- if allergic to any ingredients in this product

**Stop use and ask a doctor if**

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

**If pregnant of 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 & children 12 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 12 years of age: consult a doctor

**Other Information**

- store at room temperature 68-77°F (20-25°C)
- do not store in direct sunlight or expose to excessive heat and moisture

**TAMPER EVIDENT:** Do not use if pouch containing the patch is torn or broken.

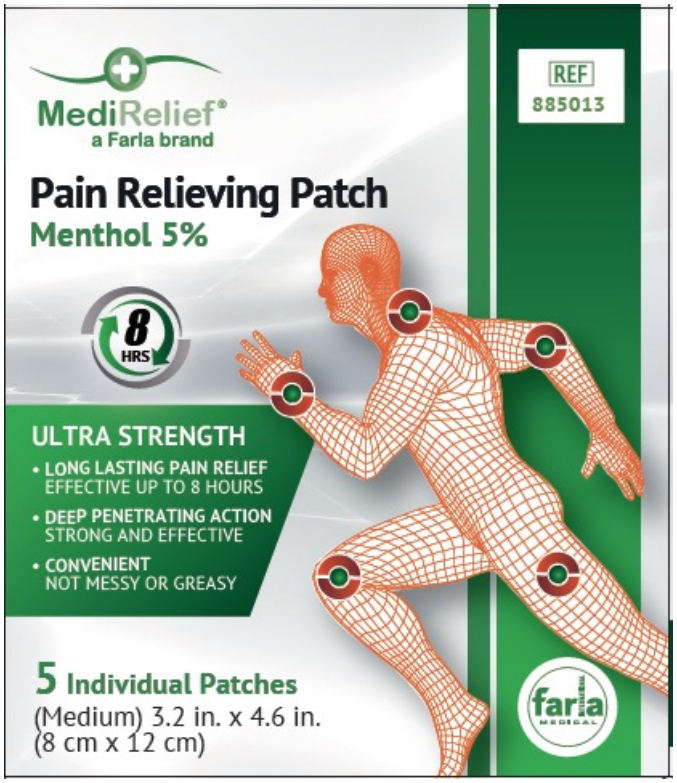
**Inactive Ingredients**

carboxymethylcellulose, dihydroxyaluminum aminoacetate, glycerin, kaolin, methylparaben, mineral oil, petrolatum, polyacrylic acid, polysorbate 80, povidone, propylene glycol, propylparaben, sodium polyacrylate, tartaric acid, titanium dioxide, water.

**Principal Display Panel**

(Medium) Patch
NDC 73486-101-25

(Large) Patch
NDC 73486-101-35




# MEDIRELIEF ULTRA

menthol patch

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:73486-101 |
| **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 | 5 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **SODIUM POLYACRYLATE (8000 MW)** (UNII: 285CYO341L) | |
| **PROPYLPARABEN** (UNII: Z8IX2SC1OH) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **DIHYDROXYALUMINUM AMINOACETATE** (UNII: DO250MG0W6) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **METHYLPARABEN** (UNII: A2I8C7HI9T) | |
| **KAOLIN** (UNII: 24H4NWX5CO) | |
| **POLYSORBATE 80** (UNII: 6OZP39ZG8H) | |
| **POLYACRYLIC ACID (8000 MW)** (UNII: 73861X4K5F) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |
| **TARTARIC ACID** (UNII: W4888I119H) | |
| **PETROLATUM** (UNII: 4T6H12BN9U) | |
| **CARBOXYMETHYLCELLULOSE** (UNII: 05JZI7B19X) | |
| **MINERAL OIL** (UNII: T5L8T28FGP) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| 1 | NDC:73486-101-25 | 5 in 1 CARTON | 05/04/2020 | |
| 1 | | 1 in 1 POUCH; Type 0: Not a Combination Product | | |
| 2 | NDC:73486-101-35 | 5 in 1 CARTON | 05/04/2020 | |
| 2 | | 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 | 05/04/2020 | |

## Labeler - FARLA MEDICAL HEALTHCARE LTD (230061759)

Revised: 2/2024 FARLA MEDICAL HEALTHCARE LTD