MQFORU PAIN PATCH- pain patch patch Zhengzhou Miaoqi Medical Technology Co., Ltd.

83781-010

Active Ingredient

Menthol 2.0%

Borneol 1.2%

Camphor 1.5%

Purpose

Topical Analgesic

Use

for sub healthy people with discomfort caused by neck, shoulder, waist and leg pain.

Warnings

For external use only.

Do not use

When using this product

- use only as directed
- do not bandage tighty or use with a heating pad
- avoid contact with eyes and mucous membranes
- do not apply to wounds or damaged, broken or imitated skin
- do not use at the same time as other topical analgesics

When Using

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Stop Use

Stop use and ask doctor if:

- rash, itching or excessive skin irritation develops
- condition worsens
- symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days If pregnant or breastfeeding, 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

Ask Doctor

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Keep Out Of Reach Of Children

■ Keep out of reach of children

Directions

adults and children over 12 years:

- ■clean and dry the affected area
- ■remove backing from patch and apply patch to affected area
- ■use 1 patch per knee for 8 to 12 hours, once per day children 12 years or younger: ask a doctor

Other information

store at room temperature, not to exceed 86°F (30°C)

Inactive ingredients

Radix Angelicae Pubescentis, Drynaria Rhizome, Myrrh, Frankincense, Elderberry, Safflower, Angelicae, and Trachlospermi Caulis Et Folium

Questions

Questions or comments?

Email: mqforu86@gmail.com

PRINCIPAL DISPLAY PANEL





MQFORU

PAIN PATCH







MQFORU®PAIN PATCH

- Lasts Up to 24 Hours
- Comfortable Flexible Fabric
- Easy to Use & Remove

PATCHES 2.8"×3.9" 7cm×10cm

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MQFORU

PAIN PATCH

cles, backaches, and joint pain





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MQFORU PAIN PATCH

pain patch patch

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:83781-010

Route of Administration TOPICAL

| Active Ingredient/Active Moiety | | | |
|---------------------------------------------------------------------------------------|----------------------|-----------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) | MENTHOL | 2 g in 100 | |
| CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET) | CAMPHOR (SYNTHETIC) | 1.5 g in 100 | |
| BORNEOL (UNII: M89NIB437X) (BORNEOL - UNII:M89NIB437X) | BORNEOL | 1.2 g in 100 | |
| | | | |

| Inactive Ingredients | |
|-----------------------------------------------------|----------|
| Ingredient Name | Strength |
| ANGELICA BISERRATA WHOLE (UNII: IAA753UT7B) | |
| MYRRH (UNII: JC71GJ1F3L) | |
| ANGELICIN (UNII: CZZ080D7BD) | |
| TRACHELOSPERMUM JASMINOIDES STEM (UNII: 7YF691N2XM) | |
| EUROPEAN ELDERBERRY (UNII: BQY1UBX046) | |
| DRYNARIA FORTUNEI ROOT (UNII: 731W842X8Q) | |
| SAFFLOWER (UNII: 4VBL71TY4Y) | |
| FRANKINCENSE (UNII: R9XLF1R1WM) | |

| P | Packaging | | | |
|---|----------------------|----------------------------------------------------|-------------------------|-----------------------|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:83781-010- 01 | 10 in 1 BOX; Type 0: Not a Combination Product | 08/13/2024 | |
| 2 | NDC:83781-010- 02 | 20 in 1 BOX; Type 0: Not a Combination Product | 08/13/2024 | |
| 3 | NDC:83781-010- 03 | 30 in 1 BOX; Type 0: Not a Combination Product | 08/13/2024 | |
| 4 | NDC:83781-010- 04 | 40 in 1 BOX; Type 1: Convenience Kit of Co-Package | 08/13/2024 | |
| 5 | NDC:83781-010- 05 | 50 in 1 BOX; Type 0: Not a Combination Product | 08/13/2024 | |
| 6 | NDC:83781-010- 06 | 60 in 1 BOX; Type 0: Not a Combination Product | 08/13/2024 | |
| 7 | NDC:83781-010- 07 | 100 in 1 BOX; Type 0: Not a Combination Product | 08/13/2024 | |

| Marketing Information | | | | |
|-----------------------|---------------------------------------------|-------------------------|-----------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC Monograph Drug | M017 | 08/13/2024 | | |
| | | | | |

Labeler - Zhengzhou Miaoqi Medical Technology Co., Ltd. (701762807)

| Establishment | | | |
|-----------------------------------------------|---------|-----------|-------------------------------------------|
| Name | Address | ID/FEI | Business Operations |
| Zhengzhou Miaoqi Medical Technology Co., Ltd. | | 701762807 | manufacture(83781-010) , label(83781-010) |

Revised: 8/2024 Zhengzhou Miaoqi Medical Technology Co., Ltd.