# ULTRA VEDIC- feminine pain relief patch Pharmaneek, 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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Ultra Vedic<sup>TM</sup>

#### **Feminine Pain Relief Patch**

The unique combination of pain relievers on the transdermal patch relieve pain due to cramps.

## **Active Ingredient**

Menthol 10%

#### **Purpose**

Topical Analgesic

#### Uses

For temporary relief of minor aches and pains in lower back and abdomen associated with cramps.

### Warnings

For External Use Only

Do not use otherwise than as directed.

#### Do not use

- On open wounds, cuts, eyes, and face
- With a heating pad

#### When using this product

- Avoid contact with eyes and mucous membranes
- Do not bandage tightly

#### Ask Your Doctor Before Use if you have

- redness over the affected area
- have sensitive skin
- are pregnant or breast-feeding

#### Stop Use and Ask Your Doctor

- If condition worsens or if symptoms persist for more than 7 days or clear up and occur again within a few days
- If abnormal skin irritation occurs after usage

#### KEEP OUT OF REACH OF CHILDREN

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

#### **Directions**

- For use by Adults and Children 12 year of age and older
- Apply to the affected area no more than 2 times a day
- Dry skin completely at application site area, before applying the patch
- Open pouch and remove patch
- Peel off protective film and apply sticky side to affected area
- If applied on hairy skin, remove gently using mild warm water
- Patch once used cannot be re-pasted or reused
- Dispose properly after use

#### Other Information

• Store at room temperature below 80°F (27°C)

#### **Inactive Ingredients**

Adhesive Plaster, Eucalyptus Oil

#### Questions?

Call 1-866-241-6885 info@pharmaneek.com www.ultravedic.com

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#### PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

Please Reseal After Opening

NDC: 72625-130-05

#### **ULTRA VEDIC**

#### **Feminine Pain Relief Patch**

Pack of 5 Patches Soothe Period Cramps Medicated Patch Fast Acting Apply for 8-12 Hours NDC: 72625-130-05

# **FEMININE PAIN RELIEF PATCH**



# SOOTHE PERIOD CRAMPS



MEDICATED PATCH

FAST ACTING

PLEASE RESEAL AFTER OPENING

#### **FEMININE PAIN RELIEF PATCH**

The unique combination of pain relievers on the transdermal patch relieve

associated with cramps







Peel off one part of protective liner

affected area, and progressively remove rest of the liner

#### DRUG FACTS

| Active Ingredient | Purpose           |  |
|-------------------|-------------------|--|
| Menthol 10%       | Topical Analgesic |  |

For External Use Only

On open wounds, cuts, eyes and face With a heating pad

#### When using this product:

Avoid contact with eyes and mucous membranes
 Do not bandage tightly

#### Ask Your Doctor Before Use if you have:

·redness over the affected area

· have sensitive skin · are pregnant or breast-feeding

#### Stop Use and Ask Your Doctor:

If condition worsens or if symptoms persist for more than 7 days or clear up and occur again within a few days

#### Keep Out of Reach of Children.

If swallowed, seek medical help or contact a Poison Control Center right

#### Directions:

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For use by Adults and Children 12 years of age and older

Apply to the affected area no more than 2 times a day

Dry skin completely at application area, before applying the patch

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#### Other Information:

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#### Inactive Ingredients

#### Questions? Call 1-866-241-6885

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Batch No.: Exp. Date:



#### **ULTRA VEDIC**

feminine pain relief patch

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72625-130

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient NameBasis of StrengthStrengthMENTHOL (UNII: L7T10 EIP3A) (MENTHOL - UNII:L7T10 EIP3A)MENTHOL200 mg

#### **Inactive Ingredients**

Ingredient Name Strength

EUCALYPTUS OIL (UNII: 2R04ONI662)

# Packaging#Item CodePackage DescriptionMarketing Start DateMarketing End Date1NDC:72625-130-051 in 1 PATCH03/01/2019

5 in 1 CARTON; 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                                  | 0 3/0 1/20 19        |                    |
|                         |  |                      |                    |

## Labeler - Pharmaneek, Inc. (063762556)

## **Registrant - Pharmaneek**, Inc. (063762556)

Revised: 12/2019 Pharmaneek, Inc.