FEMININE PAIN RELIEF- menthol, unspecified form patch
Unexo Life Sciences Private Limited

FEMININE PAIN RELIEF PATCH - 50 cm2

DRUG FACTS

Uses

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

Active Ingredient

Menthol 10%

Purpose

Topical Analgesic

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 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
- 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

Manufactured By:
Unexo Life Sciences Pvt. Ltd.
B-16, Sector 4, Bawana Industrial Area,
Delhi -110039 (INDIA)

PRINCIPAL DISPLAY PANEL - 4 Patch Pouch Label

NDC: 71391-131-04

UNEXO LIFE SCIENCES

FEMININE PAIN RELIEF PATCH
50 cm2

MENTHOL

SOOTHE PERIOD CRAMPS
FAST ACTING
APPLY FOR 8-12 HOURS

4 PATCHES

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FEMININE PAIN RELIEF PATCH

Drug Facts

Active ingredient ..... **Purpose**
Menthol 10% ..... Topical analgesic

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

Warnings
For external use only.

When using this product ■ Use only as directed ■ Rare cases of serious burns have been reported with products of this type ■ Don't bandage tightly or use with heating pad ■ Avoid contact with eyes and mucous membranes ■ Don't apply to wounds or damaged skin ■ Do not use at the same time as other topical analgesics.

Stop use and consult a doctor if ■ Condition worsens ■ Redness is present ■ Irritation develops ■ Symptoms persist for more than 7 days or clear up and occur again within a few days ■ You experience signs of skin injury, such as pain, swelling, or blistering where the product was applied.

If pregnant or breastfeeding, ask a health professional before use.

Keep out of reach of children. Do not use on infants. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions
Adults and children 12 years of age and over: Carefully remove backing from patch.
Apply sticky side of patch to affected area. Wear one patch up to 8 hours. Repeat as necessary.
Maximum 3 patches per day. Properly discard patch after single use.
Children under 12 years of age: consult a physician.

Other information
Store at room temperature, not to exceed 80°F (27°C).

Inactive ingredients
adhesive plaster, eucalyptus oil

Manufactured By:
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NC/DL-378-A&U

1 PATCH

LOT
EXP

FEMININE PAIN RELIEF
menthol, unspecified form patch

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71391-131
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	100 mg

Inactive Ingredients

Ingredient Name	Strength
Eucalyptus oil (UNII: 2R04ONI662)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71391-131-24	24 in 1 BOX	03/06/2024	
1	NDC:71391-131-04	4 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	03/06/2024	

Labeler - Unexo Life Sciences Private Limited (872260479)

Establishment

Name	Address	ID/FEI	Business Operations
Unexo Life Sciences Private Limited		872260479	MANUFACTURE(71391-131)

Revised: 3/2024 Unexo Life Sciences Private Limited