

EXTERNAL ANALGESIC- camphor (synthetic), capsicum, menthol patch
Zhengzhou Daohe Medical Technology Co.,Ltd

EXTERNAL ANALGESIC PATCH

Drug Facts

Active ingredients

Camphor 9.6%

Capsicum containing 0.05% Capsaicin

Menthol 7.8%

Purpose

External Analgesic

Uses

For the temporary relief of minor aches and pains of muscles and joints associated with

- simple backache
- arthritis
- strains
- bruises
- sprains

Warnings

For external use only

When using this product

- avoid contact with the eyes
- do not apply to wounds or damaged skin
- do not bandage tightly

Stop use and ask a doctor if

- condition worsens
- if 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.

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

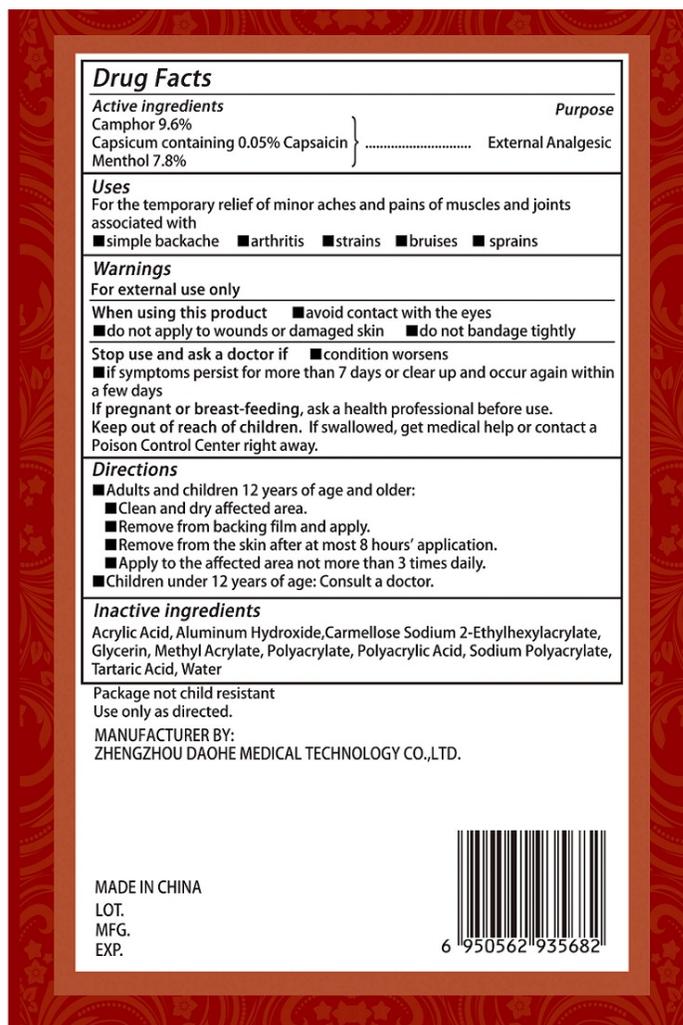
Directions

- Adults and children 12 years of age and older:
- Clean and dry affected area.
- Remove from backing film and apply.
- Remove from the skin after at most 8 hour's application
- Apply to the affected area not more than 3 times daily.
- Children under 12 years of age: Consult a doctor.

Inactive ingredients

Acrylic Acid, Aluminum Hydroxide, Carmellose Sodium 2-Ethylhexylacrylate, Glycerin, Methyl Acrylate, Polyacrylate, Polyacrylic Acid, Sodium Polyacrylate, Tartaric Acid, Water

Package Labeling:



EXTERNAL ANALGESIC

camphor (synthetic), capsiicum, menthol patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73567-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	96 mg in 1 mL
CAPSICUM (UNII: 00UK7646FG) (CAPSICUM - UNII:00UK7646FG)	CAPSICUM	0.5 mg in 1 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	78 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ACRYLIC ACID (UNII: J94PBK7X8S)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYL ACRYLATE (UNII: WC487PR91H)	
TARTARIC ACID (UNII: W4888I119H)	
WATER (UNII: 059QF0KOOR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73567-001-00	1 in 1 POUCH	04/01/2020	
1		8 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/01/2020	

Labeler - Zhengzhou Daohe Medical Technology Co.,Ltd (406230407)

Establishment

Name	Address	ID/FEI	Business Operations
Zhengzhou Daohe Medical Technology Co.,Ltd		406230407	manufacture(73567-001)