# CAPMENT- menthol, caps aicin patch Original Patch Company Inc

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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## Capment

# Active ingredients:

Capsaicin 0.0375%

Menthol 5.00%

# Purpose

Topical Anesthetic

Topical Analgesic

**Keep out of reach of children.** Consult physician for children under 12.

#### Uses

For the temporary relief of minor aches and muscle pain associated with arthritis, simple backache, strains, muscle soreness and stiffness.

#### **Warnings**

- Only for external use. Use only as directed or by a health professional.
- Do not use: on open wounds, cuts, damaged or infected skin as well as in the eyes, mouth, genitals, or any other mucous membranes.
- Do not cover with bandage.
- **Consult your physician:** If pregnant or pain persists or worsens.

## Directions

Adults and children 12 and over: Apply patch to affected area 1 to 2 times daily or as directed.

Instructions for Use:

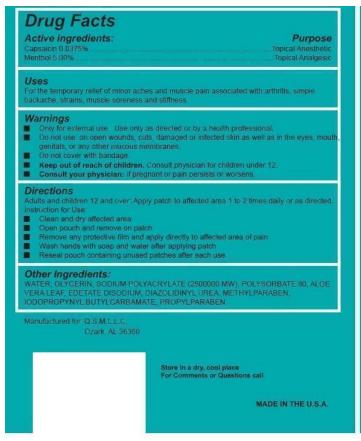
- Clean and dry affected area
- Open pouch and remove on patch
- Remove any protective film and apply directly to affected area of pain
- Wash hands with soap and water after applying patch
- Reseal pouch containing unused patches after each use

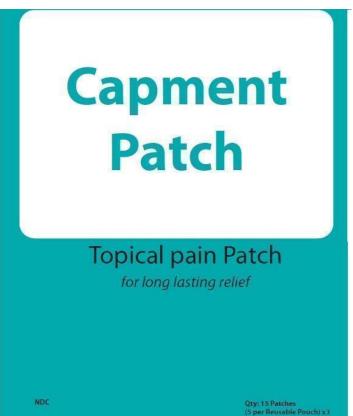
## **Other Ingredients:**

WATER, GLYCERIN, SODIUM POLYACRYLATE (2500000 MW), POLYSORBATE 80, ALOE VERA LEAF, EDETATE DISODIUM, DIAZOLIDINYL UREA, METHYLPARABEN, IODOPROPYNYL BUTYLCARBAMATE, PROPYLPARABEN

Store in a dry, cool place

## **Packaging**





#### **CAPMENT**

menthol, capsaicin patch

## **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69650-001

Route of Administration TOPICAL

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

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	.04 in 100 g
CAPSAICIN (UNII: S07O44R1ZM) (CAPSAICIN - UNII:S07O44R1ZM)	CAPSAICIN	.00375 in 100 g

#### **Inactive Ingredients**

Ingredient Name

SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)

ACRYLIC ACID/SO DIUM ACRYLATE COPOLYMER (1:1; 600 MPA.S AT 0.2%) (UNII: M4PPW69 Y4H)

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#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69650-001-15	15 in 1 BOX				

1 NDC:69650-001-05	1 g in 1 PATCH				
Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved drug other		03/24/2015			

# Labeler - Original Patch Company Inc (079705068)

Revised: 3/2015 Original Patch Company Inc