DOLLAR GENERAL COLD-HOT MEDICATED PATCHES- menthol patch
DOLGENCORP, LLC

Dollar General Cold-Hot Medicated Patches

ACTIVE INGREDIENT

Active Ingredient .....Purpose
Menthol 5%.....Topical Analgesic

INACTIVE INGREDIENT

carboxymethylcellulose sodium, Dihydroxyaluminum Aminoacetate, Glycerin, iodopropynyl butylcarbamate, Kaolin, Mineral Oil, Petrolatum, Phenoxyethanol, Polyacrylic Acid, Polysorbate 80, povidone, Propylene Glycol, Sodium Polyacrylate, Tartaric Acid, Titanium Dioxide, Water, 3-(2-ethylhexyloxy)propane-1,2-diol

KEEP OUT OF REACH OF CHILDREN

If swallowed, get medical help or contact a Poison Control

INDICATIONS & USAGE

Temporarily relieves minor pain associated with: ■ arthritis ■ simple backache ■ bursitis ■ tendonitis
■ muscle strains ■ muscle sprains ■ bruises ■ cramps

WARNINGS

For External Use Only.

DOSAGE & ADMINISTRATION

Adults and children over 12 years: Carefully remove backing from patch. Apply sticky side of patch to affected area.

Wear one patch up to 8 hours. Repeat as necessary, but no more than 4 times daily. Reseal pouch after opening.

Children 12 years or younger: Consult a physician.

PURPOSE

Topical Analgesic

When using this product

Use only as directed ■ Don’t bandage tightly or use with heating pad ■ Avoid contact with eyes and mucous membranes ■ Don’t apply to wounds or damaged skin.

Stop use and ask a doctor

If condition worsens ■ If redness is present ■ If irritation develops ■ If symptoms persist for more than 7 days or clear up and occur again within a few days.

If pregnant or breastfeeding

ask a health professional before use.

DOLLAR GENERAL COLD-HOT MEDICATED PATCHES

menthol patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-972
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	5 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
MINERAL OIL (UNII: T5L8T28FGP)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
KAOLIN (UNII: 24H4NWX5CO)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	
TARTARIC ACID (UNII: W4888I119H)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
POLYACRYLIC ACID (8000 MW) (UNII: 73861X4K5F)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-972-06	6 in 1 BOX	10/01/2017	
1		1 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	10/01/2017	

Labeler - DOLGENCORP, LLC (068331990)

Establishment

Name	Address	ID/FEI	Business Operations
Foshan Aqua Gel Biotech Co., Ltd.,		529128763	manufacture(55910-972)