MEIJER COLD THERAPY PAIN RELIEVING- menthol patch Meijer Inc

Meijer Cold Therapy Pain Relieving Patch, 5 Count

Active Ingredient

Menthol 5%

Purpose

Topical Analgesic

Use

Use For temporary relief of minor aches and pains of muscles and joints: arthritis, simple backache, bursitis, tendonitis, muscle strains, muscle sprains, bruises, and cramps.

Warnings

For External Use Only

When using this product

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- 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

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- 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

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

Directions Adults and children 12 years of age and over: Clean and dry affected area, free of lotions, ointments, and creams. Carefully remove backing from patch. Apply sticky side of patch to affected area. Do not use more than one patch in an 8 hour period. Repeat as necessary. Maximum 3 patches per day. Discard patch after single use. Children under 12 years of age: consult a physician.

Other Information

Store at room temperature, not to exceed 86 F (30C)

Inactive Ingredients

aloe vera extract, arnica montana extract, boswellia carterii resin extract, carboxymethylcellulose sodium, dihydroxyaluminum aminoacetate, ethylhexylglycerin, glycerin, green tea extract, iodopropynyl butylcarbamate, kaolin, mineral oil, petrolatum, phenoxyethanol, polyacrylic acid, polysorbate 80, povidone, propylene glycol, sodium polyacrylate, tartaric acid, titanium dioxide, water



nenthol patch						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC		NDC:7	2:79481-9966	
Route of Administration	TOPICAL					
Active Ingredient/Active	e Moiety					
Ingred	ient Name		Basis of Stren	gth	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)		MENTHOL		5 g in 100 g		
Inactive Ingredients						
Inactive Ingredients	Ingredient Name				Strength	
-					Strength	
DIHYDROXYALUMINUM AMINOA	ACETATE (UNII: DO250MG0				Strength	
DIHYDROXYALUMINUM AMINOA PROPYLENE GLYCOL (UNII: 6DC!	ACETATE (UNII: DO250MG0 9Q167V3)				Strength	
DIHYDROXYALUMINUM AMINOA PROPYLENE GLYCOL (UNII: 6DC BOSWELLIA SERRATA RESIN OI	ACETATE (UNII: DO250MG0 9Q167V3) IL (UNII: 5T1XCE6K8K)				Strength	
DIHYDROXYALUMINUM AMINOA PROPYLENE GLYCOL (UNII: 6DC BOSWELLIA SERRATA RESIN OI POLYACRYLIC ACID (8000 MW)	ACETATE (UNII: DO250MG0 9Q167V3) IL (UNII: 5T1XCE6K8K)				Strength	
Inactive Ingredients DIHYDROXYALUMINUM AMINOA PROPYLENE GLYCOL (UNII: 6DC9 BOSWELLIA SERRATA RESIN OF POLYACRYLIC ACID (8000 MW) WATER (UNII: 059QF0K00R) MINERAL OIL (UNII: T5L8T28FGP)	ACETATE (UNII: DO250MG0 9Q167V3) IL (UNII: 5T1XCE6K8K)) (UNII: 73861X4K5F)				Strength	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
KAOLIN (UNII: 24H4NWX5CO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
PETROLATUM (UNII: 4T6H12BN9U)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)	
TARTARIC ACID (UNII: W4888I119H)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:79481- 9966-5	5 in 1 CARTON	08/01/2024		
1		1 g in 1 PACKAGE; Type 0: Not a Combination Product			
Marketing Information					
	Marketing	Application Number or Monograph	Marketing Start	Marketing End	

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/01/2024	

Labeler - Meijer Inc (006959555)

Establishment

Name	Address	ID/FEI	Business Operations
Foshan Aqua Gel Biotech Co., Ltd.		529128763	manufacture(79481-9966)

Revised: 8/2024

Meijer Inc