# COLD AND HOT MEDICATED PAIN RELIEF LARGE- menthol patch Valu Merchandisers Company, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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#### **Best Choice Cold & Hot Large Medicated Patch**

## **Active Ingredient**

Menthol 5%

#### **Purpose**

Topical analgesic

#### Uses

Temporarily relieves minor pain associated with:

- arthritis
- muscle strains
- simple backache
- bursitis
- cramps
- tendonitis
- muscle sprains
- bruises

#### Warnings

# For external use only

#### When using this product

- use only as directed
- do not bandage tightly or use with heating pad
- avoid contact with eyes and mucous membranes
- do not apply to wounds or damaged skin
- do not use if you are allergic to any ingredients of this product

#### Stop use and ask a doctor if

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

#### 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 immediately.

#### **Directions**

## Adults and children over 12 years:

- carefully remove backing from patch
- apply one patch to affected area
- repeat as necessary, but no more than 4 times daily

# Children under 12 years of age: Ask a doctor

## **Inactive ingredients**

dihydroxyaluminum aminoacetate, glycerin, kaolin, methylparaben, polyacrylic acid, polyethylene glycol 400, polysorbate 80, povidone, propylparaben, purified water, sodium polyacrylate, tartaric acid, titanium dioxide

#### **Questions or Comments?**

call 1-800-883-0085

Reseal pouch after opening

## package label

Best Choice Cold and Hot Large Medicated Patch



# COLD AND HOT MEDICATED PAIN RELIEF LARGE

menthol patch

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63941-949		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	875 mg	

Inactive Ingredients	
Ingredient Name	Strength
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
PO VIDO NE (UNII: FZ989 GH94E)	
DIHYDRO XYALUMINUM AMINO ACETATE (UNII: DO 250 MG0 W6)	
GLYCERIN (UNII: PDC6A3C0OX)	
KAOLIN (UNII: 24H4NWX5CO)	
METHYLPARABEN (UNII: A2I8 C7HI9 T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
WATER (UNII: 059QF0KO0R)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYACRYLIC ACID (8000 MW) (UNII: 73861X4K5F)	
TARTARIC ACID (UNII: W48881119 H)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	

Packaging				
# Item Code	Package Description	Marketing Start Date	<b>Marketing End Date</b>	
1 NDC:63941-949-05	1 in 1 CARTON	12/29/2018		
1	5 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 not final	part348	09/12/2011			

Labeler - Valu Merchandisers Company, Inc. (868703513)

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