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.

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

carboxymethylcellulose, dihydroxyaluminum aminoacetate, glycerin, kaolin, methylparaben, mineral oil, petrolatum, polyacrylic acid, polysorbate 80, propylene glycol, povidone, propylparaben, sodium polyacrylate, tartaric acid, titanium dioxide, water

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:63941-948
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	550 mg

Inactive Ingredients	
Ingredient Name	**Strength**
PETROLATUM (UNII: 4T6H12BN9U)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
POVIDONE (UNII: FZ989GH94E)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	
GLYCERIN (UNII: PDC6A3C0OX)	

KAOLIN (UNII: 24H4NWX5CO)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
WATER (UNII: 059QF0KO0R)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYACRYLIC ACID (8000 MW) (UNII: 73861X4K5F)	
TARTARIC ACID (UNII: W4888I119H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
MINERAL OIL (UNII: T5L8T28FGP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63941-948-05	1 in 1 CARTON	07/06/2022	
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	07/06/2022	

Labeler - Valu Merchandisers Company, Inc. (868703513)

Revised: 7/2022 Valu Merchandisers Company, Inc.