ICEQUAKE COLDN HOT MEDICATED PATCH- menthol cloth Southern Sales & Service, Inc.

Icequake Cold'n Hot medicated patch

Drug Facts

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
- do not use a 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 conctact a Poison Control Center right away

Directions

Adults and children over 12 years:

store at room temperature Children under 12 years of age:

- Carefully remove backing film from patch
- Apply one patch to affected area
- Use one patch at a time, 1 or 2 times a day

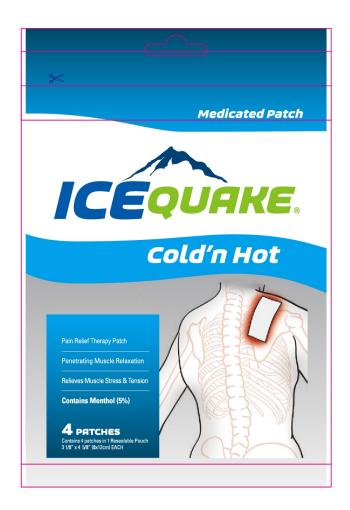
Other Information

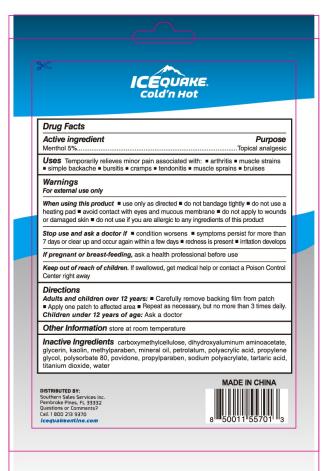
store at room temperature

Inactive Ingredients

CARBOXYMETHYLCELLULOSE, DIHYDROXYALUMINUM AMINOACETATE, GLYCERIN, KAOLIN, METHYLPARABEN, MINERAL OIL, PETROLATUM, POLYACRYLIC ACID, PROPYLENE GLYCOL, POLYSORBATE 80, POVIDONE, PROPYLPARABEN, SODIUM POLYACRYLATE, TARTARIC ACID, TITANIUM DIOXIDE, WATER

Package Labeling:





ICEQUAKE COLDN HOT MEDICATED PATCH

menthol cloth

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69822-014

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)

MENTHOL

M

Inactive Ingredients

Ingredient Name		Strength
TARTARIC ACID (UNII: W48881119H)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

POLYSORBATE 80 (UNII: 60ZP39ZG8H)

CARBOXYMETHYLCELLULOSE (UNII: 05JZ17B19X)

DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)

GLYCERIN (UNII: PDC6A3C0OX)

KAOLIN (UNII: 24H4NWX5CO)	
POVIDONE (UNII: FZ 989GH94E)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
PROPYLPARABEN (UNII: Z8IX2SC10H)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69822-014- 04	4 in 1 POUCH	11/02/2022			
1		9.5 g in 1 PATCH; 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	11/02/2022			

Labeler - Southern Sales & Service, Inc. (013114906)

Revised: 11/2023 Southern Sales & Service, Inc.