

HEALTHWISE MEDICATED COLD HOT 5% PATCH- menthol patch
Veridian Healthcare

HealthWise Medicated Cold Hot 5% Menthol Patch

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

Do not use on infants. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away

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 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. Reseal pouch after opening.

Children under 12 years of age: consult a physician.

PURPOSE

Topical Analgesic

When using this product

When using this product

- 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 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.
- You experience signs injury, such as pain, swelling, or blistering where the product was applied.

If pregnant or breastfeeding

ask a health professional before use.

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 SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)
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:71101-031-06	6 in 1 BOX	04/01/2024	
1		1 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:71101-031-25	25 in 1 BOX	04/01/2024	
2		1 in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:71101-031-12	12 in 1 BOX	10/01/2025	
3		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	04/01/2024	

Labeler - Veridian Healthcare (830437997)

Establishment

Name	Address	ID/FEI	Business Operations
Foshan Aqua Gel Biotech Co., Ltd.,		529128763	manufacture(71101-031)