HOT/COLD MEDICATED PATCH- hot cold patch patch
Dynarex

1451 Cold/Hot Medictaed Patches Arm/Neck 67777-201-10
1452 Cold/Hot Medicated Patches Back 67777-201-20

Active Ingredient

Menthol 5%

Purpose

Topical analgesic

Use

Temporarily relieves minor aches and pains of muscles and joints due to
- Arthritis
- Simple backache
- Strains
- Sprains
- Bruises

Warnings

For external use only

Do not use
- On wounds or damaged skin
- With a heating pad
- If you are allergic to any ingredients of this product

When using this product
- Use only as directed
- Avoid contact with the eyes, mucous membranes or rashes

Stop use and ask a doctor if
- Excessive redness or irritation is present
- Conditions worsen
- Symptoms persist for more than 7 days or clear up and occur again within a few days

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 (1-800-222-1222) right away.

Directions

Adults and children 12 years of age and over

- Clean and dry affected area
- Remove film from patch and apply to the skin
- Apply 1 patch at a time to affected area,not more than 3 to 4 times daily
- Wear each patch up to 8 hours maximum

Children under 12 years of age, consult a doctor

Other information

- Avoid storing product in direct sunlight
- Protect product from excessive moisture

Inactive ingredients

Aluminum Hydroxide, Castor Oil, Disodium EDTA, Glycerin, Isopropyl Myristate, Kaolin, Polysorbate 80, Polyvinyl Alcohol, Sodium Polyacrylate, Tartaric Acid, Titanium Dioxide, Water

Questions and Comments?

1-888-396-2739 Monday - Friday 9AM - 5PM EST

Label

1451 Hot Cold Medicated Patch

Label

1452 Hot Cold Medicated Patch

HOT/COLD MEDICATED PATCH

hot cold patch patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-201
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	285 mg in 5700 mg

Inactive Ingredients	
Ingredient Name	**Strength**
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
CASTOR OIL (UNII: D5340Y2I9G)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
TARTARIC ACID (UNII: W4888I119H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
KAOLIN (UNII: 24H4NWX5CO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-201-10	180 in 1 CASE	12/13/2019	
1	NDC:67777-201-11	5 in 1 BOX		
1		285 mg in 1 PATCH; Type 0: Not a Combination Product		
2	NDC:67777-201-20	180 in 1 CASE	12/13/2019	
2	NDC:67777-201-21	5 in 1 BOX		
2		285 mg 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	12/13/2019	

Labeler - Dynarex (008124539)

Registrant - Dynarex (008124539)

Revised: 12/2023 Dynarex