

MENTHOL 5%- menthol 5% patch
Advanced Rx of Tennessee, LLC

Menthol Patch
Menthol 5% | Pain Relieving Patch

ACTIVE INGREDIENTS (IN EACH PATCH)

Menthol 5%

USES

Temporary relief from minor aches and pains of sore muscles and joints

associated with:

- arthritis
- backache
- strains
- sprains

PURPOSE

External Analgesic

WARNINGS

For external use only.

CONSULT WITH A DOCTOR OR PHARMACIST BEFORE CONTINUED USE IF YOU HAVE

Sensitive Skin

WHEN USING THIS PRODUCT

- Use only as directed
- Do not bandage tightly or use with heating pad or device
- Avoid contact with the eyes or mucous membranes
- Do not apply to wounds or damaged skin
- Do not use with other ointments, creams, sprays, or liniments
- Do not apply to irritated skin
- Store in a cool, dry place away from sunlight

STOP USE AND ASK A DOCTOR IF

Burning discomfort or excessive skin irritation develops, condition worsens, or if

symptoms persist for more than 7 days, or clear up and occur again within a few days

IF YOU ARE PREGNANT OR BREAST-FEEDING

IF YOU ARE PREGNANT OR BREAST-FEEDING: Consult with physician before use of this product.

KEEP OUT OF THE REACH OF CHILDREN

If accidentally ingested, get medical help or contact a Poison Control Center immediately

DIRECTIONS:

Adults and Children 12 Years of Age or Over: Clean and dry the affected area, partially peel back the protective film and apply the exposed patch to the site of pain. Carefully remove the remaining film while pressing patch to skin and leave in place for up to 8 hours. Use on affected areas not more than 4 times daily. Wash hands with cool water after use.

Children Under 12 Years of Age: Consult physician.

OTHER INGREDIENTS

Aloe Barbadensis Leaf Extract, Arnica Montana Flower Extract, Boswellia Carterii Resin Extract, Camellia Sinensis Leaf Extract, Carboxymethylcellulose Sodium, Dihydroxyaluminum Aminoacetate, Edetate Disodium, Glycerin, Hydroxyacetophenone, Kaolin, Tartaric Acid, Mineral Oil, Petrolatum, Polyacrylic Acid, Polysorbate 80, Propylene Glycol, Sodium Polyacrylate, Titanium dioxide, Water

QUESTIONS?

Call toll-free: (307) 410-3813 or Email: info@marcellahealth.com



Packed By:
AdvancedRx
Optimizing Provider Care
Nashville, TN 37217



Store between 20° and 25°C
(68° and 77°F)
(See USP Controlled Room Temperature)
Keep Medication out of the reach of children.

MENTHOL 5% PATCH

Compared to: MENTHOL
NDC: 80425-0543-01 Source NDC: 85096-0301-30
Lot Number: XXXXXX Exp: 12/31/2027
MARCELLA HEALTH

#30

Rx Only

MENTHOL 5% PATCH #30
NDC: 80425-0543-01
Source NDC: 85096-0301-30
Lot: XXXXXX Exp: 12/31/2027



S/N: 000000399679
GTIN: 00380425054315

MENTHOL 5%

menthol 5% patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80425-0543(NDC:85096-301)
Route of Administration	TRANSDERMAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
MENTHOL, (+)- (UNII: C6B1OE8P3W) (MENTHOL, (+)- - UNII:C6B1OE8P3W)	MENTHOL, (+)-	500 mg in 10 g

Inactive Ingredients	
Ingredient Name	Strength
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
CAMELLIA SINENSIS LEAF (UNII: W2ZU1RY8B0)	
POVIDONE K90 (UNII: RDH86HJV5Z)	
PETROLATUM (UNII: 4T6H12BN9U)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
KAOLIN (UNII: 24H4NWX5CO)	
TARTARIC ACID (UNII: W4888I119H)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JN12J)	
DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS (UNII: 1K713C615K)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
CELLULOSE GUM (UNII: K679OBS311)	
FRANKINCENSE (UNII: R9XLF1R1WM)	
MINERAL OIL (UNII: T5L8T28FGP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80425-0543-1	30 in 1 BOX	08/08/2025	
1		11 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	08/08/2025	
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Labeler - Advanced Rx of Tennessee, LLC (117023142)

Establishment

Name	Address	ID/FEI	Business Operations
Advanced Rx of Tennessee, LLC		117023142	repack(80425-0543)

Revised: 8/2025

Advanced Rx of Tennessee, LLC