MEDLINE- menthol patch Medline Industries, LP

972 Medline 5% Menthol Pain Relief Patch

Active ingredient

Menthol 5%

Purpose

Topical analgesic

Uses

for the temporary relief of minor aches and pains of muscles and joints associated with:

- simple backache
- arthritis
- strains
- sprains
- bruises

Warnings

For external use only

Ask doctor before 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
- do not apply to wounds or damaged skin
- do not use with other ointments, creams, sprays, or liniments
- do not apply to irritated skin
- avoid contact with eyes, mucous membranes or rashes

Stop use and ask a doctor if

- burning discomfort or excessive skin irritation develops
- condition worsens
- symptoms persist for more than 7 days
- symptoms 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 right away.

Directions

Adult and children 2 years of age and older:

- 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
- remove patch from the skin after at most 8-hour application

Children under 2 years of age: do not use, consult a doctor

Other information

- protect product from excessive moisture
- store in a cool dry place away from direct sunlight

Inactive 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, I-tartaric acid, mineral oil, petrolatum, polyacrylic acid, polysorbate 80, propylene glycol, pvp, sodium polyacrylate, titanium dioxide, water

Manufacturer information

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Package Label



MEDLINE

menthol patch

menthol patch						
Product Information						
Product Type	HUMAN OTC DRUG	ltem C	ode (Source)	NDC:53329-972		
Route of Administration	TOPICAL					
Active Ingredient/Activ	e Moiety					
Ingredient Name			Basis of Strength	n Strength		
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)			MENTHOL	5 mg in 100 mg		
lue etine lueure die ete						
Inactive Ingredients	In our all and Manage			Charles and the		
				Strength		
EDETATE DISODIUM (UNII: 7FL	ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)					
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)						
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)						
PROPYLENE GLYCOL (UNII: 6DC	•					
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)						
POLYSORBATE 80 (UNII: 60ZP3	9ZG8H)					
PETROLATUM (UNII: 4T6H12BN9U)						
PVP (UNII: FZ 989GH94E)						
WATER (UNII: 059QF0KO0R)						
MINERAL OIL (UNII: T5L8T28FGP)						
CAMELLIA SINENSIS LEAF (UNII: W2ZU1RY8B0)						
SODIUM POLYACRYLATE (800		١				
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6) CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)						
FRANKINCENSE (UNII: R9XLF1R1WM)						
KAOLIN (UNII: 24H4NWX5CO)						
TARTARIC ACID, MESO- (UNII: JQO211TF1A)						
ALOE BARBADENSIS LEAF (UNII: ZY81Z83H0X)						
GLYCERIN (UNII: PDC6A3C0OX)						
Packaging						
# Item Code P	ackage Description		Marketing Start Date	Marketing End Date		
1 NDC:53329-972- 05 5 in 1 BOX		(02/01/2025			
1 1 in 1 POUC	4					
1 500 mg in 1 Product	PATCH; Type 0: Not a Combina	ation				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	505G(a)(3)	02/01/2025			

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 6/2025

Medline Industries, LP