EQUATE MENTHOL PAIN RELIEVING- menthol patch
Wal-Mart Stores Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active ingredient

Menthol 5%

Purpose

Menthol - Topical Analgesic

Uses

temporarily relieves minor aches and pains of muscles and joints due to
- simple backache
- arthritis
- strains
- sprains

Warnings

For external use only

When using this product
- use only as directed
- avoid contact with eyes and mucous membranes
- do not apply to wounds or to damaged or very sensitive skin
- do not bandage tightly or use with a heating pad

Stop use and ask a doctor if
- condition worsens
- excessive redness, irritation, burning, or discomfort of the skin develops
- symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 12 years and over: apply to affected area; change patch 1 to 2 times daily
- children under 12 years: ask a doctor
- FOR BEST RESULTS apply to clean, dry skin
- tear open pouch and remove patch; if desired, cut patch to size
- grasp both ends of the patch firmly with thumbs near center
- stretch patch until the backing separates
- remove protective film while applying patch directly to site of pain

Inactive ingredients

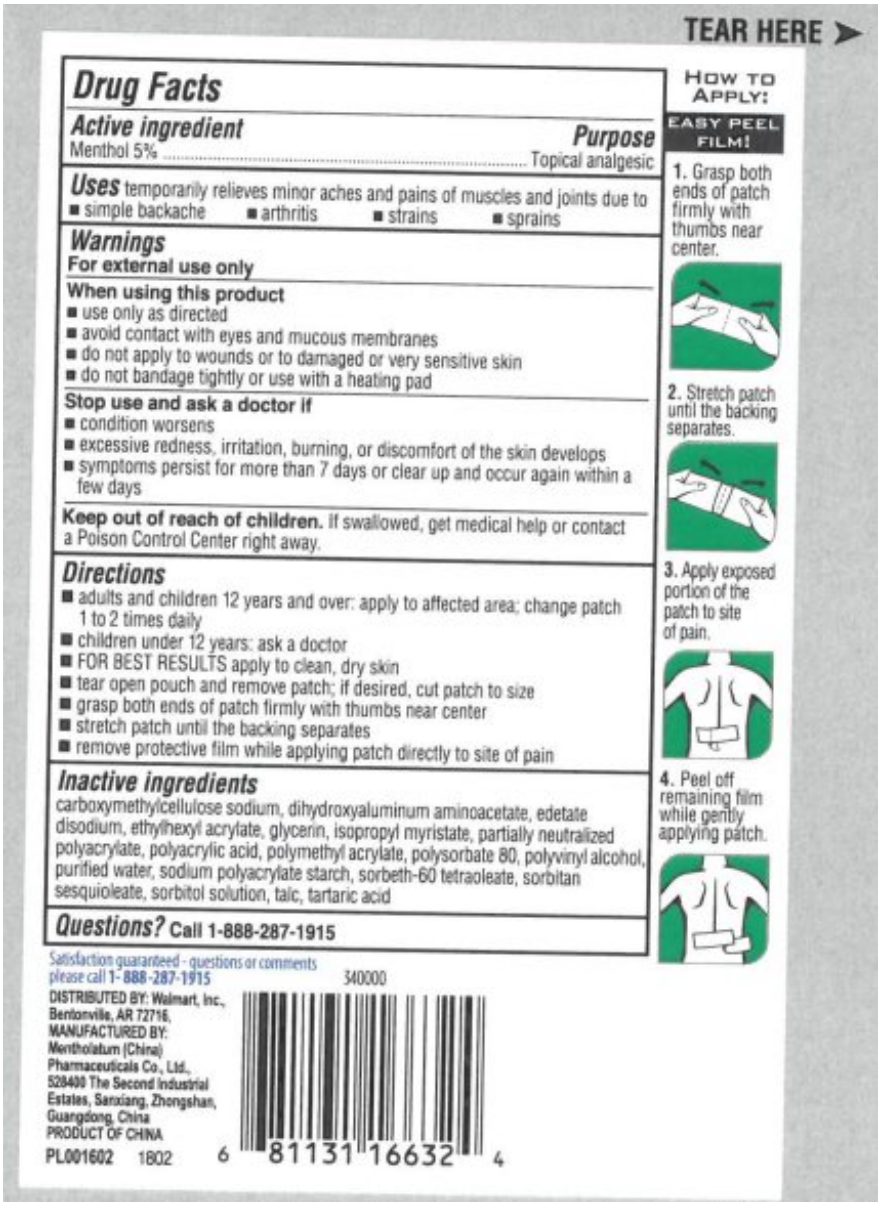
carboxymethylcellulose sodium, dihydroxyaluminum aminoacetate, edetate disodium, ethylhexyl acrylate, glycerin, isopropyl myristate, partially neutralized polyacrylate, polyacrylic acid, polymethyl acrylate, polysorbate 80, polyvinyl alcohol, purified water, sodium polyacrylate starch, sorbeth-60 tetraoleate, sorbitan sesquioleate, sorbitol solution, talc, tartaric acid

Questions?

Call1-888-287-1915

Package/Label Principal Display Panel

Principal Display Panel



EQUATE MENTHOL PAIN RELIEVING

menthol patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-870
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	50 mg

Inactive Ingredients	
Ingredient Name	**Strength**

CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
2-ETHYLHEXYL ACRYLATE (UNII: HR49R9S6XG)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
METHYL ACRYLATE (UNII: WC487PR91H)	
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
WATER (UNII: 059QF0KO0R)	
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
SORBITOL (UNII: 506T60A25R)	
TALC (UNII: 7SEV7J4R1U)	
TARTARIC ACID (UNII: W4888I119H)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-870-01	1 in 1 POUCH; Type 0: Not a Combination Product	01/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	01/01/2018	

Labeler - Wal-Mart Stores Inc (051957769)

Revised: 4/2023 Wal-Mart Stores Inc