

CVS ULTRA STRENGTH PAIN RELIEVING- menthol patch

CVS Pharmacy, 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

Topical Analgesic

□**Uses** □Temporarily relieves minor pain associated with:

- arthritis
- simple backache
- bursitis
- tendonitis
- muscle strains
- muscle sprains
- bruises cramps□

□Warnings

For external use only□

□**When using this product**

- □use only as directed
- do not bandage tightly or use with a heating pad
- avoid contact with eyes and mucous membranes
- do not apply to wounds or damaged skin

□**Stop use and ask a doctor if**

- □condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days
- redness is present
- skin irritation develops

□**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**

- open pouch and remove patch if desired, cut patch to size, peel off protective backing and apply sticky side to affected area
- adults and children 12 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 12 years of age: consult a doctor□

□**Other information**

- □store between 20° and 25° C (68° and 77°F)

□**Inactive ingredients**□ butylated hydroxytoluene, carboxymethylcellulose sodium, castor oil, concentrated glycerin, EDTA-2Na, gelatin, isopropyl myristate, kaolin, magnesium aluminium hydroxide, methylparaben, polyacrylic acid, polysorbate 80, polyvinyl alcohol, purified water, sodium

polyacrylate, sorbitan monooleate, tartaric acid, titanium dioxide

Distributed By:

CVS Pharmacy, Inc.

One CVS Drive.

Woonsocket, RI 02895

Made in Korea

BLEED: _____

NO COPY: _____

GLUE AREA: 



CVS ULTRA STRENGTH PAIN RELIEVING

menthol patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-476
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	750 mg

Inactive Ingredients

Ingredient Name	Strength
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
CASTOR OIL (UNII: D5340Y2I9G)	
GELATIN (UNII: 2G86QN327L)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
KAOLIN (UNII: 24H4NWX5CO)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
WATER (UNII: 059QF0KO0R)	
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)	
TARTARIC ACID (UNII: W4888I119H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-476-04	1 in 1 CARTON		
1		5 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 not final	part348	09/20/2013	

Labeler - CVS Pharmacy, Inc. (062312574)

Revised: 12/2015

CVS Pharmacy, Inc.