EXTRA STRENGTH PAIN RELIEF THERAPY - menthol patch United Exchange Corp

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

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

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

If pregnant or breast-feeding, ask a health professional before use.

Directions

- adults and children 12 years of age and older
- peel off protective backing and apply sticky side to affected area
- carefully remove backing from patch
- should be used up to 8 hours should be used no more than 3 times a day
- children under 12 years of age consult a doctor

Other information

• store at room temperature, not to exceed 86°F (30°C)

Inactive ingredients aloe barbadenisis leaf juice, aluminum hydroxide, BHT, castor oil, disodium edetate, gelatin, glycerin, isopropyl myristate, kaolin, magnesium aluminate metasilicate, methyl paraben, peg-40 stearate, polysorbate 80, polyvinyl alcohol, purified water, PVP sodium polyacrylate, tartaric acid, titanium dioxide, tocopherol acetate, Release Liner: polypropylene, Non-Woven Material:

Polyester Non-Woven Fabric.

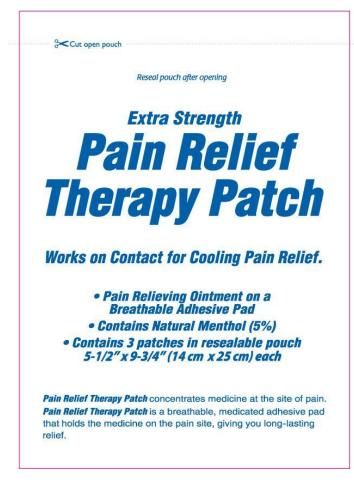
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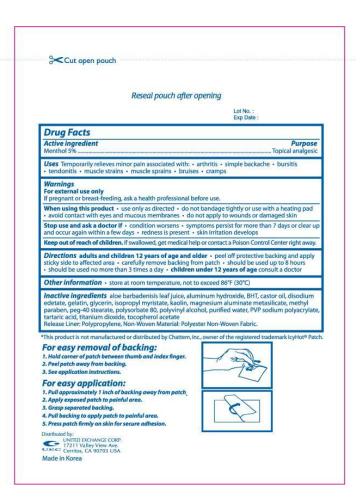
United Exchange Corp

17211 Valley View Ave.

Cerritos, CA 90703 USA

Made in Korea





EXTRA STRENGTH PAIN RELIEF THERAPY

menthol patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-702

Route of Administration TOPICAL

Active ingredient/Active wiole	lient/Active Moiety	e Ingredie	Active
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Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	750 mg

Inactive I	ngredients
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Ingredient Name Strength

ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
ALUMINUM HYDRO XIDE (UNII: 5QB0T2IUN0)	
BUTYLATED HYDRO XYTO LUENE (UNII: 1P9 D0 Z171K)	
CASTOR OIL (UNII: D5340 Y2I9G)	
EDETATE DISO DIUM (UNII: 7FLD91C86K)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)	
KAOLIN (UNII: 24H4NWX5CO)	
METHYLPARABEN (UNII: A2I8 C7HI9 T)	
PEG-40 STEARATE (UNII: ECU18C66Q7)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
WATER (UNII: 059QF0KO0R)	
TARTARIC ACID (UNII: W48881119H)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	
ALPHA-TO COPHEROL ACETATE (UNII: 9E8X80D2L0)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-702-03	3 in 1 POUCH		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	06/30/2013	

Labeler - United Exchange Corp (840130579)

Revised: 6/2013 United Exchange Corp