## 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.

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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

- conditions worsen
- 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 1,3-butylene glycol, aloe vera (powder), metacrylic acid butylacrylate copolymer, dibutylhydroxytoluene, disodium edetate, D-sorbitol solution, gelatin, glycerine, kaolin, light liquid paraffin, magnesium aluminium hydrate, methyl parahydroxybenzoate, polysorbate 80, purified water, sodium metaphosphate, sodium polyacrylate, sorbitan monooleate, tartaric acid, titanium oxide,

tocopherol acetate.

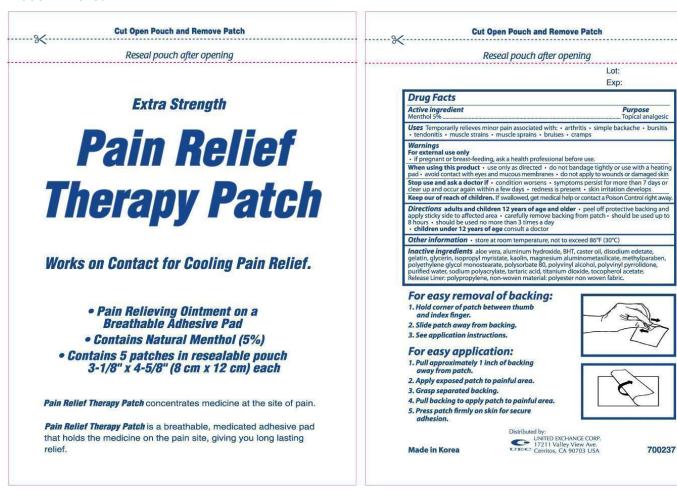
Distributed By:

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:65923-703 Route of Administration TOPICAL

## Active Ingredient/Active Moiety Ingredient Name Basis of Strength MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) MENTHOL 750 mg

### **Inactive Ingredients**

Strength

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:65923-703-05	5 in 1 POUCH; Type 0: Not a Combination Product	06/30/2013	

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: 9/2019 United Exchange Corp