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

Active ingredient Purpose

Menthol 5%.....Topical analgesic

Uses Temporary relieves minor pain associated with:

- arthritis
- simple backache
- bursitis
- tendonitis
- muscle strains
- 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 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 vera, aluminum hydroxide, BHT, caster oil, disodium edetate, gelatin, glycerin, isopropyl myristate, kaolin, magnesium aluminometasilicate, methylparaben, polyethylene glycol monostearate, polysorbate 80, polyvinyl alcohol, polyvinyl pyrrolidone, purified water, sodium polyacrylate, tartaric acid, titanium dioxide, tocopherol acetate. Release Liner: polypropylene, non-

woven material: polyester non woven fabric.

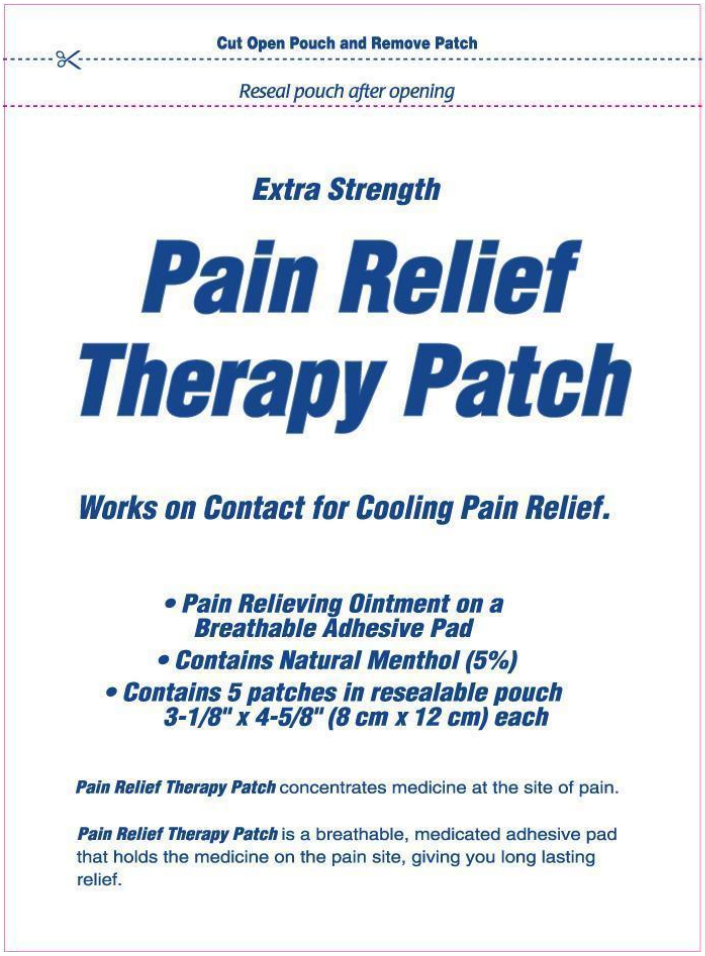
Distributed by:

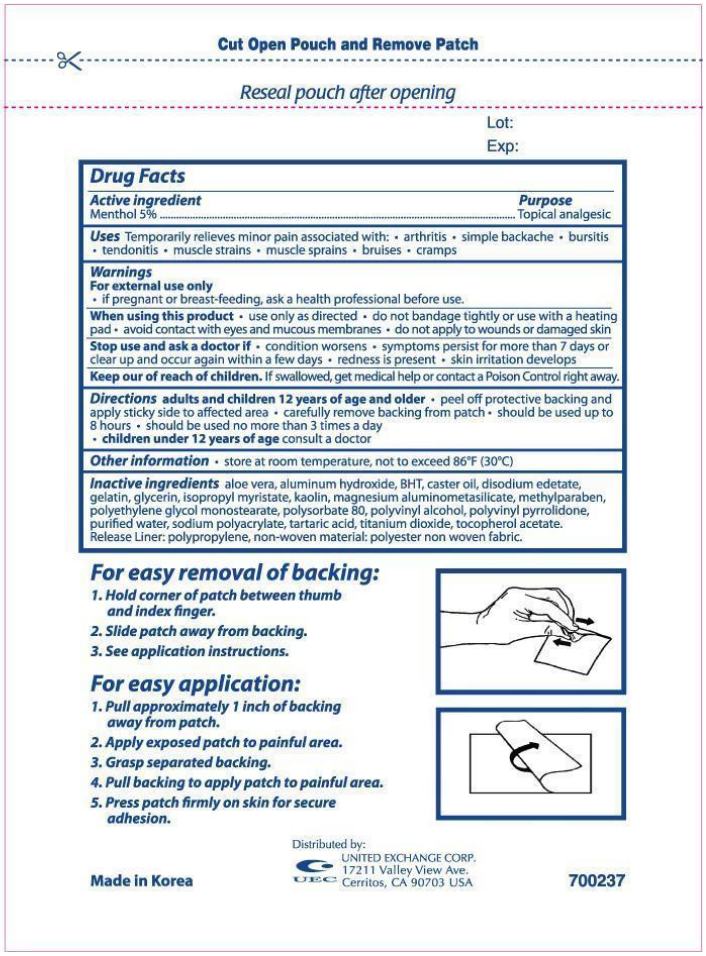
United Exchange Corp

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:65923-701
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**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
KAOLIN (UNII: 24H4NWX5CO)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
WATER (UNII: 059QF0KO0R)	
TARTARIC ACID (UNII: W4888I119H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65923-701-05	5 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