SOUND BODY EXTRA STRENGTH COLD AND HOT MEDICATED- 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.

Active ingredeint Purpose

Menthol 5%..... Topical analgesic

Uses

Temporarily relieves minor pain associated with:

- arthritis
- simple backache
- muscle strains
- bursitis
- tendonitis
- 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
- redness is present
- skin irritation develops

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. In case of accidental ingestion, get medical help or contact a Poison Control Center right away

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 12 years of younger: ask a doctor

Other information

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

Inactive ingredients

aluminum hydroxide, camphor, eucalyptus oil, glycerine, methylparaben, polyacrylic acid, propaneidol, propylparaben, purified water

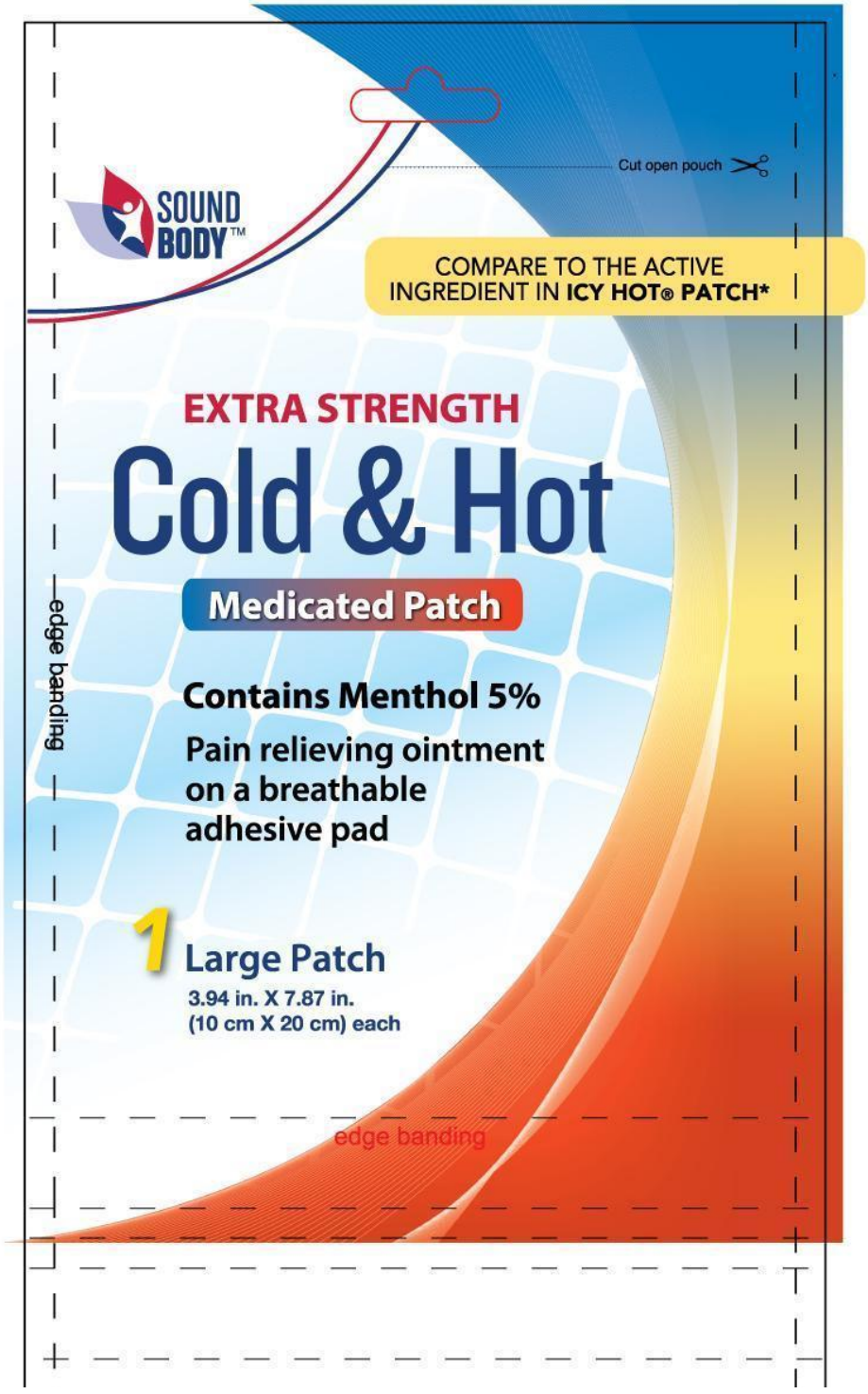
DISTRIBUTED BY:

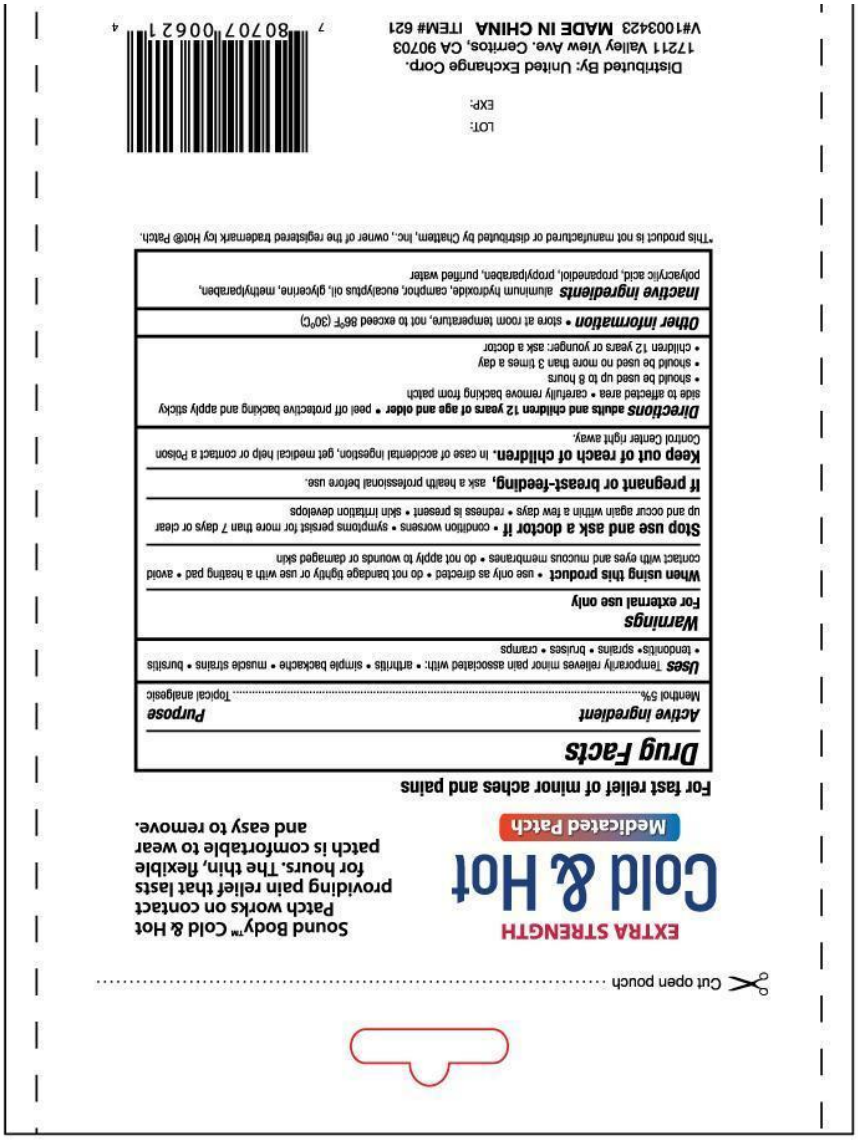
UNITED EXCHANGE CORP.

17211 VALLEY VIEW AVE.

CERRITOS, CA 90703

MADE IN CHINA





SOUND BODY EXTRA STRENGTH COLD AND HOT MEDICATED

menthol patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-621
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	50 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**

ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-621-01	1 in 1 POUCH		
1		1 g in 1 PATCH; 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	02/25/2015	

Labeler - United Exchange Corp. (840130579)

Revised: 8/2015 United Exchange Corp.