CORALITE PAIN RELIEF- dl-camphor, l-menthol, methylsalicylate 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.

Coralite Pain Relief Patch 20 Ct

Active ingredients Purpose

DL-Camphor 1.2%..... Topical Analgesic

L-Menthol 5.7%..... Topical Analgesic

Methyl Salicylate 6.3%..... Topical Analgesic

Uses

For temporary relief of minor aches and pains of muscles and joints associated with:

- arthritis
- simple backache
- strains
- bruises
- sprains

Warnings

For external use only

Do not use

- on wounds or damaged skin
- if you are allergic to aspirin or salicylates
- with a heating pad
- with, or at the same time as, other external analgesic products

Ask a doctor before use if you allergic to any ingredients of this product

When using this product

- do not use other than directed
- avoid contact with the eyes, mucous membranes or rashes

Stop use and ask a doctor if

- rash, itching or excessive skin irritation develops
- conditions worsen symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days

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

Caution: This product contains natural rubber latex which may cause allergic reactions

Other information

- avoid storing product in direct sunlight
- protect from excessive moisture

Inactive ingredients BHT, calcium carbonate, C4-6 olefin/styrene copolymer, glyceryl abiertate, isopropyl myristate, polybutene, polyisobutylene, propylene carbonaterubber latex, sorbitan stearate, ys

resin, zinc oxide

DISTRIBUTED BY:

UNITED EXCHANGE CORP.

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CERRITOS, CA 90703 USA

CORALITE PAIN RELIEF

dl-camphor, l-menthol, methylsalicylate patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-156
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	1.2 g in 100 g
LEVOMENTHOL (UNII: BZ1R15MTK7) (LEVOMENTHOL - UNII:BZ1R15MTK7)	LEVOMENTHOL	5.7 g in 100 g
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	6.3 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
STYRENE (UNII: 44LJ2U959V)	
GLYCERYL ABIETATE (UNII: 2F22LY70Q1)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
POLYISOBUTYLENE (1000 MW) (UNII: 5XB3A63Y52)	
POLYBUTENE (1400 MW) (UNII: 1NA5AO9GH7)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
ZINC OXIDE (UNII: SOI2LOH54Z)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65923-156-20	20 in 1 PACKAGE	04/18/2016	
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	03/26/2014	

Labeler - United Exchange Corp. (840130579)

Revised: 4/2016 United Exchange Corp.