HEALTH MART PAIN RELIEF- camphor, menthol, and methyl salicylate patch
McKesson

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 ingredients	**Purpose**
Camphor 1.2%	Topical analgesic
Menthol 5.7%	Topical analgesic
Methyl salicylate 6.3%	Topical analgesic

Uses

for temporary relief of minor aches & pains of muscles & joints associated with:

- arthritis
- simple backache
- strains
- bruises
- sprains

Warnings

For external use only

Allergy alert: If prone to allergic reaction from aspirin or salicylates, consult a doctor before use.

Do not use

- on wounds or damaged skin
- with a heating pad
- if you are allergic to any ingredients of this product

Stop use and ask a doctor if

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

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

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

Directions

adults and children 12 years of age and over:

- clean and dry affected area
- remove patch from film
- apply to affected area not more than 3 to 4 times daily
- remove patch from the skin after at most 8 hours' application

children under 12 years of age:

- consult a doctor

Other information

- avoid storing product in direct sunlight

- protect product from excessive moisture

Inactive ingredients

glyceryl hydrogenated rosinate, hydrated silica, mineral oil, polyethylene glycol 400, polyisobutylene (1300 MW), styrene/isoprene copolymer, YS resin, zinc oxide

Distributed by:

McKesson

One Post Street

San Franciso, CA 94104

www.healthmart.com/healthmartbrand

HEALTH MART PAIN RELIEF

camphor, menthol, and methyl salicylate patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62011-0328
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	7.1 mg
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	33 mg
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	36 mg

Inactive Ingredients

Ingredient Name	Strength
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYISOBUTYLENE (1300 MW) (UNII: 241BN7J12Y)	
STYRENE (UNII: 44LJ2U959V)	
ISOPRENE (UNII: 0A62964IBU)	
ZINC OXIDE (UNII: SOI2LOH54Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62011-0328-1	40 in 1 BOX; Type 0: Not a Combination Product	01/23/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	01/23/2017	

Labeler - McKesson (177667227)

Registrant - United Exchange Corp. (840130579)

Establishment

Name	Address	ID/FEI	Business Operations
Foshan Aqua Gel Biotech Co., Ltd.		529128763	manufacture(62011-0328)

Revised: 1/2017 McKesson