SOUNDBODY MEDICATED PATCH- camphor, menthol, methyl salicylate patch Kareway Product, Inc.

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.

Soundbody Medicated Pain Relief Patch

Active ingredients

Camphor 3.1%

Menthol 6.0%

Methyl salicylate 10.0%

Purpose

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

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

When using this product

- use only as directed
- avoid contact with the eyes, mucous membranes or rashes
- do not bandage tightly

Stop use and ask a doctor if

- rash, itching or excessive skin irritation develops
- contisions worsen
- symptoms 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-hour 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

hydrogenated poly, pentaerythrityl tetra-di-t-butyl hydroxyhydrocinnamate, petroleum, styrene/isoprene copolymer

Principal Display Panel



SOUNDBODY MEDICATED PATCH

camphor, menthol, methyl salicylate patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67510- 0547
Route of Administration	TOPICAL, PERCUTANEOUS, TRANSDERMAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	31 mg		
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	60 mg		
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:0414PZ4LPZ)	METHYL SALICYLATE	100 mg		

Inactive Ingredients			
Ingredient Name	Strength		
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)			
HYDROGENATED POLY(C6-14 OLEFIN; 2 CST) (UNII: POTX083987)			
LIQUID PETROLEUM (UNII: 6ZAE7X688J)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:67510- 0547-6	1 in 1 BOX	02/19/2020			
1		6 in 1 POUCH; 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	02/19/2002	

Labeler - Kareway Product, Inc. (121840057)

Revised: 2/2023 Kareway Product, Inc.