PROFOOT PAIN RELIEF PATCHES- camphor, menthol, methyl salicylate
Profoot, Inc.

Profoot Pain Relief Patches

Active ingredients

Camphor 1.2%

Menthol 5.7%

Methyl salicylate 6.3%

Purpose

Topical Analgesic

Topical Analgesic

Topical Analgesic

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

•arthritis •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 of the 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 • condition worsens •symptoms persist

for more than 7 days • symptoms clear up and occur again within a few days

If pregnant or breastfeeding, 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 3-4 times daily • remove patch from skin after at most 8 hours

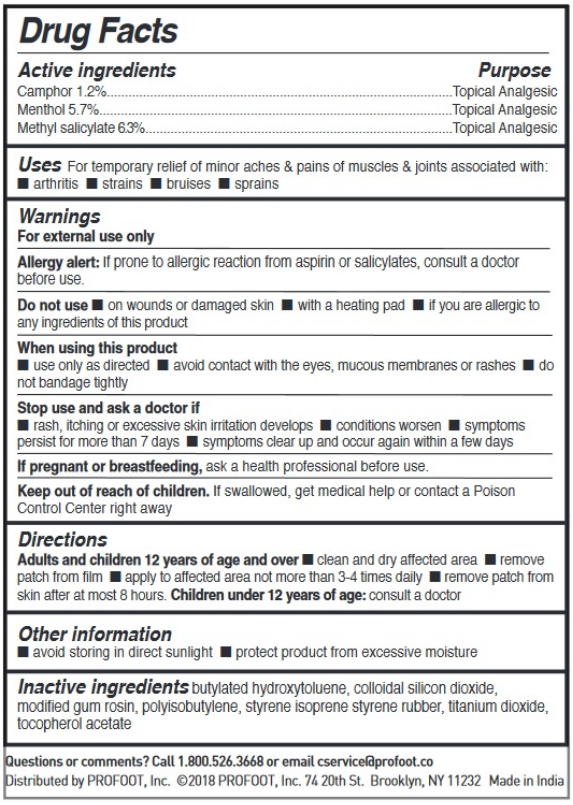
Children under 12 years of age: consult a doctor

Other information

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

Inactive ingredients hydrogenated poly, pentaerythrityl tetra-di-t-butyl Hydroxyhydrocinnamate, white mineral oil, styrene/Isoprene copolymer

Questions or comments? Email cservice@profoot.co



Drug Facts

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 ■ 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 ■ conditions worsen ■ symptoms persist for more than 7 days ■ symptoms clear up and occur again within a few days

If pregnant or breastfeeding, ask a health professional before use.

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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 3-4 times daily ■ remove patch from skin after at most 8 hours. **Children under 12 years of age:** consult a doctor

Other information

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

Inactive ingredients butylated hydroxytoluene, colloidal silicon dioxide, modified gum rosin, polyisobutylene, styrene isoprene styrene rubber, titanium dioxide, tocopherol acetate

Questions or comments? Call 1.800.526.3668 or email cservice@profoot.co

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PROFOOT PAIN RELIEF PATCHES

camphor, menthol, methyl salicylate kit

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:29784-601

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29784-601-01	1 in 1 KIT	10/12/2023	05/31/2027

Quantity of Parts

Part #	Package Quantity	Total Product Quantity

Part 1	1 PATCH	0.36 g
Part 2	1 PATCH	1.3 g

Part 1 of 2

PROFOOT

camphor, menthol, methyl salicylate patch

Product Information

Item Code (Source)	NDC:29784-121
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	63 mg in 1 g
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	12 mg in 1 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	57 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
STYRENE/ISOPRENE/STYRENE BLOCK COPOLYMER (UNII: K7S96QM8DV)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	
MINERAL OIL (UNII: T5L8T28FGP)	
HYDROGENATED C6-20 POLYOLEFIN (100 CST) (UNII: 39EYQ1W9RB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		1 in 1 KIT		
1	NDC:29784-121-36	0.36 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 Drug	M017	10/12/2023	

Part 2 of 2

PROFOOT

camphor, menthol, methyl salicylate patch

Product Information	
Item Code (Source)	NDC:29784-122
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	12 mg in 1 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	57 mg in 1 g
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	63 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
MINERAL OIL (UNII: T5L8T28FGP)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	
HYDROGENATED C6-20 POLYOLEFIN (100 CST) (UNII: 39EYQ1W9RB)	
STYRENE/ISOPRENE/STYRENE BLOCK COPOLYMER (UNII: K7S96QM8DV)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1		2 in 1 KIT		
1	NDC:29784-122-36	1.3 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 Drug	M017	10/12/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	M017	10/12/2023	05/31/2027

Labeler - Profoot, Inc. (107570900)

Revised: 10/2024 Profoot, Inc.