

SUSTAFIX- camphor cream
Hakimian Global LLC

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.

Sustafix (Hakimian Global LLC | 82225-002-10)

Active Ingredient

Camphor 1.00%

Purpose

External Analgesic

Uses

For temporary relief of pain

Warnings

For external use only. Contains bee venom.

Do Not use

In large quantities, particularly over raw surfaces or blistered areas. • If condition worsens or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a doctor.

When Using This Product

Avoid contact with the eyes

Stop using and Ask a Doctor

if irritation or rash occurs.

Keep Out Of Reach of Children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily. Children under 2 years of age: consult a doctor.

Other Information

Keep dry. Store at 15°C and 30°C. Avoid direct sunlight.

Inactive Ingredients

Acetylated glycol stearate, Aesculus hippocastanum extract, Amica montana flower extract, Beeswax, Benzyl alcohol, Butyrospermum parkii butter, Ceteth-251 Cet:yl alcohol, Cyclopentasiloxane, Dimethicone, Ethylhexylglycerin, Ethylhexyl stearate, Eucalyptus globulus leaf oil, Harpagophytum procumbens root extract, Limonene, Linalool, PEG-9 stearate, Phenoxyethanol, Propolis extract, Propylene glycol, Rosmarinus officinalis leaf oil, Sesamum indicum seed oil, Sodium polyacrylate, Trideceth-61 Water, Bee venom, BHT1 Methyl nicotinate

Principal Display

Made in the EU

Manufactured for Likasso, LTD

Grafenberger Alle 115

4023 Dusseldorf, Germany

NDC: 82225-002-10

- How to use:**
1. Take a small amount of the cream
 2. Massage it onto the affected area
 3. Let the product be fully absorbed
 4. Repeat daily
- * Use once per day

Drug Facts

Active Ingredient	Purpose
Camphor 1.00% (wt/wt)	External Analgesic
Uses For temporary relief of pain	
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Made in The EU:
Manufactured for Likasso LTD
Grafenberger Allee 115,
4023 Düsseldorf,
Germany

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sales@sustafix.com
www.instagram.com/sustafix



SustaFix

Pain Relieving Cream



- For your body
- Improves mobility

Net Contents 3.4 fl. oz./100 mL e

● For your body
● Improves mobility

SustaFix
Pain Relieving Cream



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Net Contents 3.4 fl. oz./100 mL e

● For your body
● Improves mobility

SustaFix
Pain Relieving Cream



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* Use once per day



SCAN ME



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SUSTAFIX

camphor cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82225-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
AESCULUS HIPPOCASTANUM FLOWER (UNII: KK0Z92II8M)	
CETETH-25 (UNII: 5KLY4IOG20)	

CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)
EUCALYPTUS GLOBULUS LEAF (UNII: S546YWLW6E6)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
ROSMARINUS OFFICINALIS FLOWER (UNII: NR1A27F29O)
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)
BENZYL ALCOHOL (UNII: LKG8494WBH)
LINALOOL, (+)- (UNII: F4VNO44C09)
SODIUM BENZOATE (UNII: OJ245FE5EU)
WATER (UNII: 059QF0KO0R)
LIMONENE, (+)- (UNII: GFD7C86Q1W)
SYNTHETIC BEESWAX (UNII: 08MNR5YE2R)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)
APIS MELLIFERA VENOM (UNII: 76013O881M)
GLYCOL STEARATE (UNII: 0324G66D0E)
CETYL ALCOHOL (UNII: 936JST6JCN)
DIMETHICONE (UNII: 92RU3N3Y1O)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)
METHYL NICOTINATE (UNII: 7B1AVU9DJN)
PEG-9 STEARATE (UNII: WZU67V0H4I)
HARPAGOPHYTUM PROCUMBENS ROOT (UNII: 1OYM338E89)
PROPOLIS WAX (UNII: 6Y8XYV2NOF)
SESAMUM INDICUM WHOLE (UNII: JD6YPE8XLT)
TRIDECETH-6 (UNII: 3T5PCR2H0C)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82225-002-10	1 in 1 BOX	10/27/2021	
1		100 g in 1 TUBE; 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	10/27/2021	

Labeler - Hakimian Global LLC (117010019)