

KATINKO PAIN AND ITCH RELIEVING- camphor (synthetic), menthol, methyl salicylate ointment
Greenstone Pharmaceutical Inc.

Katinko Pain And Itch Relieving Ointment

Active Ingredients:

Camphor 11 percent

Menthol 7.6 percent

Methyl Salicylate 13.0 percent

Purpose

Topical Analgesic

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Uses:

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

Simple Backache Arthritis Rheumatism Muscle Strain Sprains Bruises

For temporary relief of pain and itching associated with:

Insect Bites Minor skin irritation

Warnings

For External Use Only.

When using this product:

Use only as directed.

Do not apply to wounds or damaged skin.

Do not use on the eyes or on the mucous membranes.

Do not use with a heating pad or apply external heat.

Stop use and ask a doctor if:

Skin redness or excessive irritation of the skin occurs

Condition worsens

Symptoms persists for more than 7 days or clears up and occurs again within a few days

Do not bandage tightly

Keep out of reach of children

If swallowed, get medical help or contact a poison control

Directions:

Adults and children 2 yrs and up
times daily

Apply to affected areas not more than 3 to 4

Children under 2 years

Consult a doctor before use

Other Information

This product may provoke allergic reaction in some individuals. Test on small area before use.

Inactive Ingredients

Petroleum Jelly, Paraffin Wax

MANUFACTURED BY:

Greenstone

Pharmaceutical

H.K. Inc.

Anabu Industrial Estate,

Imus, Cavite, Philippines 4103

PRODUCT OF THE PHILIPPINES

MEDICATED

For external use only



DO NOT PURCHASE IF CARTON IS OPENED.

**METHYL SALICYLATE
MENTHOL + CAMPHOR**

KATINKO™
OINTMENT

**TOPICAL
ANALGESIC
OINTMENT**

FOR PAIN
AND ITCH RELIEF

NET WT. 0.36 OZ (10G)

MANUFACTURED BY:
Greenstone Pharmaceutical, Inc.
Anabu Industrial Estate,
Imus, Cavite, Philippines; 4103

UNDER LICENSE OF:
Greenstone
Pharmaceutical
H.K. Ltd.
Hong Kong

MADE IN THE PHILIPPINES

8 52241 01802 9

NDC #: 52241-100-10
WWW.KATINKO.COM

GREENSTONE

Drug Facts
Active ingredients: Purpose
Camphor 11% Topical Analgesic
Menthol 7.6% Topical Analgesic
Methyl Salicylate 13.0% Topical Analgesic

Uses: For temporary relief of minor aches and pains of muscles and joints associated with: ■ Neck and shoulder stiffness ■ Rheumatism ■ Muscle strain ■ Sprain ■ Bruises
For temporary relief of pain and itching associated with: ■ Insect bites ■ Minor skin irritation

Warnings:
■ **For external use only.**
When using this product: ■ Use only as directed. ■ Do not apply to wounds or damaged skin. ■ Do not use on the eyes or on mucous membranes.
■ Do not use with a heating pad or apply external heat. ▶

Drug Facts (continued)
Stop use and ask a doctor if:
■ Skin redness or excessive irritation of the skin occurs ■ Condition worsens
■ Symptoms persist for more than 7 days or clears up and occurs again within a few days.
■ Do not bandage tightly
■ **Keep out of reach of children.**
■ If swallowed, get medical help or contact poison control.

Directions:
Adults and children 2 years and up: Apply to affected area not more than 3 - 4 times daily.
Children under 2 years: Consult a doctor before use.

Other Information: This product may provoke allergic reaction in some individuals. Test on small area before use.

30g Package Label



KATINKO PAIN AND ITCH RELIEVING

camphor (synthetic), menthol, methyl salicylate ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52241-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	11 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	7.6 g in 100 g
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
PARAFFIN (UNII: I9O0E3H2ZE)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52241-100-10	1 in 1 CARTON	05/24/2010	
1		10 g in 1 JAR; Type 0: Not a Combination Product		
2	NDC:52241-100-03	1 in 1 CARTON	05/24/2010	
2		3 g in 1 JAR; Type 0: Not a Combination Product		
3	NDC:52241-100-30	1 in 1 CARTON	05/24/2010	
3		30 g in 1 BOTTLE; 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	05/24/2010	

Labeler - Greenstone Pharmaceutical Inc. (719794307)

Registrant - Greenstone Pharmaceutical Inc. (719794307)

Establishment

Name	Address	ID/FEI	Business Operations
Greenstone Pharmaceutical Inc.		719794307	manufacture(52241-100)

