KINKAN COOL- diphenhydramine hydrochloride, menthol liniment
KINKANDO CO., LTD .

Drug Facts

Active ingredients

Diphenhydramine Hydrochloride 2%

Menthol 1%

Purpose

Antihistamine

External Analgesic

Uses

Temporary relief of itching associate with insect bite

Warnings

For external use only

Flammable keep away from fire or flame

Do not use on

- wounds
- damaged skin

When using this product

- avoid contact with the eyes and mouth

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days

Keep out of reach of children.

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

Directions

Adults and children 6 years of age and older: Apply to affected area not more than 3 to 4 times daily. Children under 6 years of age: Do not use, consult a doctor.

Other Information

- Store at room temperature under 86°F (30°C)
- Flammable. keep away from fire or flame

Inactive ingredients

Glycerin
Geraniol-denatured Alcohol

Questions and comments

Call 1-310-661-7260 Mon.-Fri. (2p.m. - 3p.m. PST)

Principal Display Panel - 50 mL Bottle Label

NDC 51027-0330-2

KINKAN
COOL
Liniment

EXTERNAL
ANTIPRURITIC
LINIMENT

MANUFACTURED BY
KINKANDO CO., LTD.350-2 KODAMACHOKYOEI,
HONJO-SHI, SAITAMA 367-0206,
JAPAN

DISTRIBUTED BY
PMAI21112 Figueroa Street Unit B

Carson, CA 90745

1.7 FL OZ (50mL)

KINKAN COOL

diphenhydramine hydrochloride, menthol liniment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51027-0330

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	1 g in 50 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.5 g in 50 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51027-0330-2	96 in 1 CARTON	04/26/2000	
1		1 in 1 BOX		
1		50 mL 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	04/26/2000	

Labeler - KINKANDO CO., LTD . (694329470)

Establishment

Name	Address	ID/FEI	Business Operations
KINKANDO CO., LTD .		715451980	manufacture(51027-0330)

Revised: 6/2025 KINKANDO CO., LTD .