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

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.

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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
PMAI
1700 W. WALNUT PARKWAY,
COMPTON, CA 90220

1.7 FL OZ (50mL)
KINKAN COOL

diphenhydramine hydrochloride, menthol liniment
# Product Information

<table>
<thead>
<tr>
<th><strong>Product Type</strong></th>
<th>HUMAN OTC DRUG</th>
<th><strong>Item Code (Source)</strong></th>
<th>NDC:51027-0330</th>
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</thead>
<tbody>
<tr>
<td><strong>Route of Administration</strong></td>
<td>TOPICAL</td>
<td></td>
<td></td>
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</table>

## Active Ingredient/Active Moiety

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>diphenhydramine hydrochloride (UNII: TC2D6JAD40) (diphenhydramine - UNII:8GTS82S83M)</td>
<td>diphenhydramine hydrochloride</td>
<td>1.00 g in 50 mL</td>
</tr>
<tr>
<td>Menthol (UNII: L7T10EIP3A) (Menthol - UNII:L7T10EIP3A)</td>
<td>Menthol</td>
<td>0.50 g in 50 mL</td>
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</tbody>
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## Inactive Ingredients

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
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</thead>
<tbody>
<tr>
<td>glycerin (UNII: PDC6A3C0OX)</td>
<td></td>
</tr>
<tr>
<td>alcohol (UNII: 3K9958V90M)</td>
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## Packaging

<table>
<thead>
<tr>
<th>#</th>
<th>Item Code</th>
<th>Package Description</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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<tbody>
<tr>
<td>1</td>
<td>NDC:51027-0330-2</td>
<td>96 in 1 CARTON</td>
<td>04/26/2000</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td></td>
<td>1 in 1 BOX</td>
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<tr>
<td>1</td>
<td></td>
<td>50 mL in 1 BOTTLE; Type 0: Not a Combination Product</td>
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## Marketing Information

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<th>Marketing Category</th>
<th>Application Number or Monograph Citation</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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<tr>
<td>OTC monograph not final</td>
<td>part348</td>
<td>04/26/2000</td>
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## Labeler - KINKANDO CO., LTD. (694329470)

### Establishment

<table>
<thead>
<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
<th>Business Operations</th>
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<tr>
<td>KINKANDO CO., LTD.</td>
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<td>715451980</td>
<td>MANUFACTURE(51027-0330)</td>
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Revised: 10/2019