

**LANACANE MAXIMUM STRENGTH ANTI-ITCH- benzethonium chloride and benzocaine cream**  
**RB Health (US) LLC**

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**Lanacane ® Maximum Strength Anti-Itch Cream**

***Drug Facts***

<b><i>Active ingredients</i></b>	<b><i>Purpose</i></b>
Benzethonium chloride 0.2%	First-aid antiseptic
Benzocaine 20%	Pain relief cream

**Uses**

first aid for the temporary relief of pain and itching and to help prevent infection in minor cuts, scrapes and burns

**Warnings**

**For external use only**

**Do not use**

- in the eyes
- over large areas of the body

**Ask a doctor before use if you have**

- deep or puncture wounds
- animal bites
- serious burns

**Stop use and ask a doctor if**

- condition worsens
- needed for longer than 1 week

**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: clean the affected area and apply a small amount to the affected area not more than 1 to 3 times daily
- children under 2 years of age: ask a doctor
- may be covered with a sterile bandage

## **Other information**

- store at 20-25°C (68-77°F)
- retain carton for future reference

## **Inactive ingredients**

acetylated lanolin alcohol, aloe, cetyl acetate, cetyl alcohol, cholecalciferol, corn oil, dimethicone, dl-alpha tocopherol acetate, fragrance, glycerin, glyceryl monostearate, isopropyl myristate, methylparaben, mineral oil, PEG-100 stearate, polyvinylpyrrolidone/eicosene copolymer, propylparaben, pyrithione zinc, retinyl palmitate, sorbitan monostearate, stearamidopropyl PG-dimonium chloride phosphate, water

## **Questions?**

**1-866-252-5327.**

You may also report side effects to this phone number.

Distributed by: RB Health (US)  
Parsippany, NJ 07054-0224

## **PRINCIPAL DISPLAY PANEL - 28 g Tube Carton**

NDC 63824-810-01

LANACANE ®

Maximum Strength

ANTI-ITCH  
CREAM

Benzethonium chloride 0.2%  
(First-Aid Antiseptic)  
Benzocaine 20%  
(Pain Relieving Cream)

2-in-1

FAST Acting  
Itch Relief

+

Kills Germs\*

\*unlike Hydrocortisones

Temporary itch relief from:

- Insect bites
- Rashes
- Dry, Itchy skin

NET WT. 1 OZ (28 g)



LOT:  
EXP:

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Maximum Strength Anti-Itch Cream

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Benzocaine 20%.....	Pain relieving cream

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**Warnings**

For external use only

Do not use

- in the eyes
- over large areas of the body

Consult a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and consult 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 2 years of age and older: clean the affected area and apply a small amount of this product on the area not more than 1 to 3 times daily
- children under 2 years of age: consult a doctor
- may be covered with a sterile bandage

**Other information**

- store at 20-25°C (68-77°F)
- retain carton for future reference

**Inactive ingredients**

acetylated lanolin alcohol, aloe, cetyl acetate, cetyl alcohol, cholecalciferol, corn oil, dimethicone, di-alpha tocopherol acetate, fragrance, glycerin, glyceryl monostearate, isopropyl myristate, methylparaben, mineral oil, PEG-100 stearate, polyvinylpyrrolidone/ethylene copolymer, propylparaben, pyriithione zinc, retinyl palmitate, sorbitan monostearate, stearamidopropyl PG-dimonium chloride phosphate, water

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**LANACANE MAXIMUM STRENGTH ANTI-ITCH**

benzethonium chloride and benzocaine cream

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63824-810
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZETHONIUM CHLORIDE</b> (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	2 mg in 1 g
<b>BENZOCAINE</b> (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	200 mg in 1 g

### Inactive Ingredients

Ingredient Name	Strength
<b>ACETYLATED LANOLIN ALCOHOLS</b> (UNII: SNN716810P)	
<b>ALOE</b> (UNII: V5VD430YW9)	
<b>CETYL ACETATE</b> (UNII: 4Q43814HXS)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>CHOLECALCIFEROL</b> (UNII: 1C6V77QF41)	
<b>CORN OIL</b> (UNII: 8470G57WFM)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>.ALPHA.-TOCOPHEROL ACETATE, DL-</b> (UNII: WR1WPI7EW8)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>ISOPROPYL MYRISTATE</b> (UNII: ORE8K4LNJS)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>PYRITHIONE ZINC</b> (UNII: R953O2RHZ5)	
<b>SORBITAN MONOSTEARATE</b> (UNII: NVZ4I0H58X)	
<b>STEARAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE</b> (UNII: W6000VEI5Y)	
<b>VITAMIN A PALMITATE</b> (UNII: 1D1K0N0VVC)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63824-810-01	1 in 1 CARTON	09/28/2012	08/25/2025
1		28 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 Drug	M003	09/28/2012	08/25/2025

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**Labeler** - RB Health (US) LLC (081049410)

Revised: 11/2023

RB Health (US) LLC