

LIDOCHILL(TM) GEL- lidocaine hcl 4%, menthol 1% gel
PureTek Corporation

NDC 59088-240-16
Lidochill™ Gel
(Lidocaine 4% / Menthol 1%)
Topical Anesthetic and Analgesic

Active ingredients (% w/w)

Lidocaine HCL 4%

Menthol 1%

Purposes

Topical Anesthetic (Lidocaine HCL 4%)

Topical Analgesic (Menthol 1%)

Uses

For the temporary relief of pain and itching associated with:

- minor burns ■ sunburn ■ minor cuts ■ scrapes ■ insect bites
- minor skin irritations

Warnings

For external use only

- Do not apply to wounds or damaged skin ■ Do not bandage tightly

When using this product

- Avoid contact with the eyes

Stop use and ask a doctor if

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 physician.

If pregnant or breast-feeding,

ask a health professional before use.

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: do not use, consult a doctor.

Other information

Store at 20° to 25°C (68° to 77°F). [See USP Controlled Room Temperature]

Inactive ingredients

Aqua (Purified Water), Bisabolol, Carbomer, Ethylhexylglycerin, Glycerin, Phenoxyethanol, Sodium Hydroxide.

| | |
|--|--------------------|
| Drug Facts | |
| Active ingredients | Purpose |
| Lidocaine HCl 4%..... | Topical Anesthetic |
| Menthol 1%..... | Topical Analgesic |
| Uses | |
| For the temporary relief of pain and itching associated with: | |
| <ul style="list-style-type: none"> ■ minor burns ■ sunburn ■ minor cuts ■ scrapes ■ insect bites ■ minor skin irritations | |
| Warnings | |
| For external use only | |
| <ul style="list-style-type: none"> ■ Do not apply to wounds or damaged skin ■ Do not bandage tightly | |
| When using this product | |
| <ul style="list-style-type: none"> ■ Avoid contact with the eyes | |
| Stop use and ask a doctor | |
| 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 physician. | |
| If pregnant or breast-feeding, ask a health professional before use. | |
| Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. | |
| Directions | |
| <ul style="list-style-type: none"> ■ 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: do not use, consult a doctor. | |
| Other information | |
| Store at 20° to 25°C (68° to 77°F). [See USP Controlled Room Temperature] | |
| Inactive ingredients | |
| Aqua (Purified Water), Bisabolol, Carbomer, Ethylhexylglycerin, Glycerin, Phenoxyethanol, Sodium Hydroxide. | |

NDC 59088-240-16

DERMACIN[®]

Lidochill[™] Gel

**Lidocaine HCl 4%,
Menthol 1%**

**Topical Anesthetic
and Analgesic**

For external use only

8 fl oz (237 mL)

Manufactured in the USA by:
PureTek Corporation
Panorama City, CA 91402
For questions or information
call toll-free: **877-921-7873**

List No: 240161GA Rev: 39245

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Manufactured in the USA by:
PureTek Corporation
San Fernando, CA 91340
877-921-7873

LIDOCHILL(TM) GEL
lidocaine hcl 4%, menthol 1% gel

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:59088-240 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|--------------------------|-----------------|
| LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987) | LIDOCAINE HYDROCHLORIDE | 40 mg in 1 mL |
| MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) | MENTHOL | 10 mg in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|--|-----------------|
| CARBOMER (UNII: 0A5MM307FC) | |
| BISABOLOL (UNII: 24WE03BX2T) | |
| ETHYLHEXYLGLYCERIN (UNII: 147D247K3P) | |
| SODIUM HYDROXIDE (UNII: 55X04QC32I) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| PHENOXYETHANOL (UNII: HIE492ZZ3T) | |
| WATER (UNII: 059QF0KO0R) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|----------|------------------|---|-----------------------------|---------------------------|
| 1 | NDC:59088-240-16 | 237 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product | 09/08/2025 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------------|---|-----------------------------|---------------------------|
| OTC Monograph Drug | M015 | 09/08/2025 | |

Labeler - PureTek Corporation (785961046)**Establishment**

| Name | Address | ID/FEI | Business Operations |
|---------------------|----------------|---------------|----------------------------|
| PureTek Corporation | | 785961046 | manufacture(59088-240) |