

NUNATURE FIRST AID GEL- lidocaine hci/benzethonium chloride gel
Tec Laboratories Inc

NuNature First Aid Gel

Benzethonium Chloride 0.2%

Lidocaine Hydrochloride 2.5%

Uses

- **First aid to help prevent infection in minor:**
- **cuts**
- **scrapes**
- **burns**
- **For the temporary relief of pain or discomfort in minor:**
- **cuts**
- **scrapes**
- **burns**

Warnings

For external use only

Do not use

- in large quantities particularly over raw surfaces or blister areas

When using this product

avoid contact with eyes

Stop use and ask a doctor if

- symptoms last for more than 7 days or clear up and occur again within a few days
- conditions worsens

KEEP OUT OF REACH OF CHILDREN

if swallowed, get medical help or contact Poison Control center right away

Directions

- **adults and children 2 years of age and older:**clean the affected area whenever possible
- apply to affected area not more than 3 to 4 times daily
- may be covered with a sterile bandage;if bandaged, let dry first
- **children under 2 years of age:**do not use, consult a doctor

Other information

Store at 59-86°F(15-30°C)

allantoin, aminomethyl propanol, carbomer, disodium EDTA, glycerin, polyoxyl 35 castor oil, purified water, sodium carbonate, sodium metabisulfite, tea tree oil, white thyme oil.

Questions?

Call **1-800-482-4464**. Serious side effects associated with this product may be reported to this number.

Topical Analgesic and First Aid Antiseptic

**with Lidocaine
for fast
pain relief**

**Helps prevent
infection**

Lidocaine HCl 2.5%
Benzethonium Chloride 0.2%

**CHILD RESISTANT
PUSH & TURN TO OPEN**

NET WT. 1 OZ. (28.3g)



**FIRST
AID
GEL**

**TOPICAL ANALGESIC
FIRST AID
ANTISEPTIC**

Drug Facts	Drug Facts (continued)
<p>Active ingredients Benzethonium Chloride 0.2% Lidocaine HCl 2.5%</p> <p>Purpose First aid antiseptic Topical analgesic</p> <p>Uses ■ first aid to help prevent infection in minor: ■ cuts ■ scrapes ■ burns ■ For the temporary relief of pain or discomfort in minor: ■ cuts ■ scrapes ■ burns</p> <p>Warnings For external use only</p> <p>Do not use ■ in large quantities particularly over raw surfaces or blister areas</p> <p>When using this product avoid contact with eyes</p> <p>Stop use and ask a doctor if ■ symptoms last for more than 7 days or clear up and occur again within a few days ■ condition worsens</p> <p>KEEP OUT OF REACH OF CHILDREN. If swallowed, get medical help or contact a Poison Control center right away</p>	<p>Directions ■ adults and children 2 years of age and older: clean the affected area whenever possible ■ apply to affected area not more than 3 to 4 times daily ■ may be covered with a sterile bandage; if bandaged, let dry first ■ children under 2 years of age: do not use, consult a doctor</p> <p>Other information Store at 59-86° F (15-30° C)</p> <p>Inactive ingredients allantoin, aminomethyl propanol, carbomer, disodium EDTA, glycerin, polyoxyl 35 castor oil, purified water, sodium carbonate, sodium metabisulfite, tea tree oil, white thyme oil</p> <p>Questions? Call 1-800-482-4464. Serious side effects associated with this product may be reported to this number.</p>

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Mfg. by Tec Laboratories, Inc. Albany, OR 97321 © 2021 Child Resistant Package

NUNATURE FIRST AID GEL

lidocaine hci/benzethonium chloride gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51879-211
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	2 mg in 1 g
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	25 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
TEA TREE OIL (UNII: VIF565UC2G)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
THYME OIL (UNII: 2UK410MY6B)	
ALLANTOIN (UNII: 344S277G0Z)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
POLYOXYL 35 CASTOR OIL (UNII: 6D4M1DAL6O)	
SODIUM CARBONATE (UNII: 45P3261C7T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51879-211-01	28.3 g in 1 TUBE; Type 0: Not a Combination Product	08/11/2022	05/31/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/11/2022	05/31/2026

Labeler - Tec Laboratories Inc (083647792)

Establishment

Name	Address	ID/FEI	Business Operations
Tec Laboratories Inc		083647792	manufacture(51879-211)

Revised: 1/2026

Tec Laboratories Inc