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

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.

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 quantitiies 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 bandange;if bandaged, let dry first
- **children under 2 years of age:** do not use, consult a doctor

Other information

Store at 59-86ºF(15-30ºC)

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

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	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	08/11/2022	

Labeler - Tec Laboratories Inc (083647792)

Establishment

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

Revised: 10/2023 Tec Laboratories Inc