LIDOEASE 2%- lidocaine hcl 2% gel
PureTek Corporation

NDC 59088-221 Lidoease™ 2% Gel

Active Ingredient

Lidocaine HCl 2%

Purpose

Topical Analgesic

Uses

For the temporarily relief of pain and itching associated with:
■ minor burns ■ sunburn ■ minor cuts ■ scrapes ■ insect bites ■ minor skin irritations

Warnings

For external use only.

When using this product:

■ Avoid contact with the eyes. ■ Do not use in large quantities, particularly over raw surfaces or blistered areas.

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

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 or older: Apply externally to the affected area up to 3-4 times daily.
■ Children under 2 years: do not use, consult a doctor.

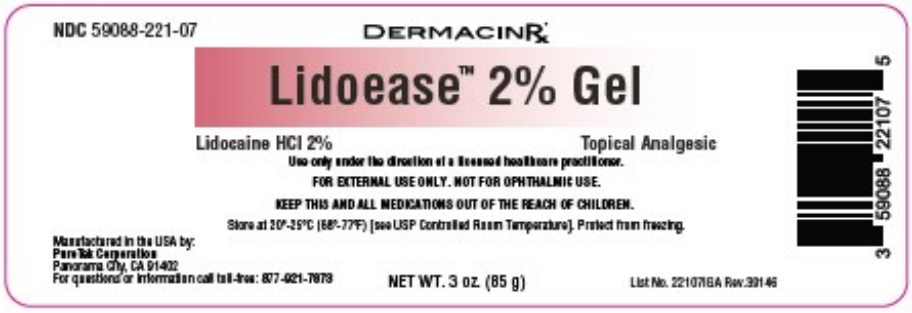
Other information

■ Store at USP controlled room temperature 20° to 25°C (68° to 77°F)

Inactive ingredients:

Aloe Barbadensis (Aloe Vera) Leaf Juice, Aqua (Purified Water), Caprylyl Glycol, Carbomer, Chlorphenesin, Cucumis Sativus (Cucumber) Fruit Extract, Dimethicone, Glycerin, Phenoxyethanol, Propanediol, Propylene Glycol, Sodium Hydroxide.

Lidoease™ 2% Gel



LIDOEASE 2%

lidocaine hcl 2% gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59088-221
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
DIMETHICONE (UNII: 92RU3N3Y1O)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER (UNII: 0A5MM307FC)	
CUCUMBER (UNII: YY7C30VXJT)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPANEDIOL (UNII: 5965N8W85T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59088-221-07	85 g in 1 TUBE; Type 0: Not a Combination Product	08/06/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/06/2025	

Labeler - PureTek Corporation (785961046)

Revised: 8/2025 PureTek Corporation