

**LIDOCAINE CREAM, 4%- lidocaine cream**  
**IPG PHARMACEUTICALS, INC.**

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**Active Ingredient**

Lidocaine 4%

**Purposes**

Topical Analgesic

**Uses**

For the temporary relief of pain and itching, associated with minor burns, sun burn, minor cuts, scrapes, insect bites, minor skin irritations, rashes due to poison ivy, poison oak, fever blister, cold sores.

**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.

**Warning**

**Warnings for External Use only**

- Avoid contact with eyes
- Do not use in large quantities, particularly over raw surface or blistered area

**Stop Use and Ask a Doctor if**

- condition worsens
- or if symptoms persist for more than 7 days
- or 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.

**Other Information**

- Store at room temperature, 20 – 25°C (68-77°F).

**Inactive Ingredients**

Allantoin, Aloe Barbadosensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Carbomer, Cetearyl Alcohol, Ethylhexylglycerin, Glycerin, Glyceryl Stearate SE, Helianthus Annuus

(Sunflower) Oil, Phenoxyethanol, Polysorbate-20, Polysorbate-60, Simmondsia Chinensis (Jojoba) Oil, Stearic Acid, Triethanolamine.

**Product label**

**IPG**

NDC 71085-075-19

**Lidocaine Cream, 4%**

Topical Analgesic Cream

Net Wt. 119gm / 4.2oz

**Drug Facts**

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Questions?  
 1-888-711-7116



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Manufactured in USA for IPG Pharmaceuticals, Inc., Tempe, AZ 85281

**LIDOCAINE CREAM, 4%**

lidocaine cream

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:71085-075
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	

<b>CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE</b> (UNII: 0A5MM307FC)
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)
<b>GLYCOL STEARATE</b> (UNII: 0324G66D0E)
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)
<b>SUNFLOWER OIL</b> (UNII: 3W1JG795YI)
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)
<b>POLYSORBATE 60</b> (UNII: CAL22UVI4M)
<b>JOJOBA OIL</b> (UNII: 724GKU717M)
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71085-075-60	60 g in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/13/2025	12/31/2025
2	NDC:71085-075-19	119 g in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/17/2025	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/13/2025	

**Labeler** - IPG PHARMACEUTICALS, INC. (080441238)

Revised: 1/2026

IPG PHARMACEUTICALS, INC.