

**PAINLESS TATTOO NUMBING- lidocaine hydrochloride liquid**  
**Sambria Pharmaceuticals, LLC**

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***Drug Facts***

***Active ingredient***

Lidocaine HCL 4.0%

***Purpose***

External Analgesic

***Uses***

For temporary relief of pain and itching.

***Warnings***

**For external use only.**

**Flammable:** Do not use near heat, flame, or while smoking.

**Do not use** on wounds or damaged skin, in large quantities, or if you are allergic to any ingredients of this product.

**When using this product** use only as directed. Keep away from face to avoid breathing it. Avoid contact with eyes. Do not store at temperatures above 120°F. Do not puncture or incinerate.

**Stop use and ask doctor 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** and pets If swallowed get medical help or contact a Poison Control Center right away.

***Directions***

Shake well. Do not spray on face. Adults and children 12 years of age and over: Clean and dry affected area, apply to affected area not more than 3 to 4 times daily. Children 12 years of age or younger: ask a doctor.

***Other Information***

Protect this product from excessive heat and direct sun.

## Inactive Ingredients

Aloe Barbadensis Leaf Extract, Arnica Montana Flower Extract, Aqua (Deionized Water), Dimethyl Sulfone, Disodium EDTA, Ethoxydiglycol, Ethylhexylglycerin, Glycerin, Phenoxyethanol, Propylene Glycol, SD Alcohol 40-B

## Product label

**PT PAINLESS TATTOO**

**NUMBING SPRAY**  
TOPICAL ANESTHETIC

**ALL GAIN NO PAIN**

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## PAINLESS TATTOO NUMBING

lidocaine hydrochloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:54723-019
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	4 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
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<b>WATER</b> (UNII: 059QF0KO0R)
<b>ALCOHOL</b> (UNII: 3K9958V90M)
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)
<b>DIMETHYL SULFONE</b> (UNII: 9H4PO4Z4FT)
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)
<b>DIETHYLENE GLYCOL MONOETHYL ETHER</b> (UNII: A1A1I8X02B)
<b>ARNICA MONTANA FLOWER</b> (UNII: OZ0E5Y15PZ)
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54723-019-01	41 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2024	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/09/2024	

**Labeler** - Sambria Pharmaceuticals, LLC (078676259)

### Establishment

Name	Address	ID/FEI	Business Operations
Southeast Holdings Corp		080504027	manufacture(54723-019)

Revised: 11/2025

Sambria Pharmaceuticals, LLC