

**NEUROMED TOPICAL ANESTHETIC 7- lidocaine hydrochloride cream**  
**Sambria Pharmaceuticals**

*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.*

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**NeuroMed 7 Topical Anesthetic**

***Active Ingredient***

Lidocaine HCL 4.00% w/w

***Purpose***

External Analgesic

***Uses***

For temporary relief of pain and itching due to minor skin irritation.

***Warnings***

☐ **For external use only**

☐ **Avoid contact with eyes**

☐ **Do not use** ☐ in large quantities, particularly over raw surfaces or blistered areas

☐ **Stop use and ask a doctor if** ☐

- Condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days. Discontinue use.

☐ **Keep out of reach of children** ☐

- If product is swallowed, get medical help or contact a Poison Control Center right away.

***Directions***

For adults and children two-years or older: Apply to affected area not more than 3 or 4 times daily.  
Children under 2 years of age: consult a physician. Apply in a circular motion for 50 to 60 seconds.

***Inactive Ingredients***

Aqua (Deionized Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Chondroitin Sulfate, Emu Oil, Ethoxydiglycol, Ethylhexyglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Melaleuca Alternifolia (Tea Tree) Oil, Methylsulfonylmethane (MSM), Phenoxyethanol, Polyacrylamide, Propylene Glycol, Stearic Acid, Triethanolamine.

☐ ***Other Information***

Protect this product from excessive heat and direct sun.

☐ ***Questions or Comments?***

FDA Registered: NDC No. 54723-667-04

**NEUROMED 7**  
7-FLUOROCARNE TOPICAL ANALGESIC

4 ml / .14 fl.oz

LIDOCAINE  
  


**Drug Facts**

Active Ingredients	Purpose
Lidocaine HCl, 4.0% w/w	External Analgesic

**Uses**  
 For temporary relief of pain and itching due to minor skin irritation.

**Warnings**  
**For external use only.**  
**Avoid contact with eyes.**  
 Do not use in large quantities, particularly over raw surfaces or broken areas.  
**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. Discontinue use.  
**Keep out of reach of children**  
 • If product is swallowed, get medical help or contact a Poison Control Center right away.

**Directions**  
 For adults and children two-years or older: Apply to affected area not more than 3 to 4 times daily. Children under 2 years of age, consult a physician. Apply in a circular motion for 30 to 90 seconds.

**Inactive Ingredients**  
 Aqua (Deionized Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Uroindolein Sulfate, Iru Oil, Ethoxydiglycol, Ethylhexylglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Methylsilylphenylmethane (MSM), Phenoxymethanol, Polyoxyethylene Propylene Glycol, Stearic Acid, Triphenylamine.

**Other Information**  
 Protect this product from excessive heat and direct sun.

**Questions or Comments?**  
 FDA Registered: NDC No. 54721-900-04  
 800-749-6801

Manufactured for Samba Pharmaceuticals  
 1075 Peachtree St. NE Ste 5650, Atlanta, GA 30359  
 Made in the USA



## NEUROMED TOPICAL ANESTHETIC 7

lidocaine hydrochloride cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:54723-300
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1 g

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)
SODIUM CHONDROITIN SULFATE (PORCINE; 5500 MW) (UNII: H5BJH23Z9A)
EMU OIL (UNII: 344821WD61)
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A118 X02B)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)
LAURETH-7 (UNII: Z95S6G8201)
MELALEUCA ALTERNIFOLIA LEAF (UNII: G43C57162K)
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
POLYACRYLAMIDE (10000 MW) (UNII: E2KR9C9V2I)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
STEARIC ACID (UNII: 4ELV7Z65AP)
TROLAMINE (UNII: 9O3K93S3TK)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54723-300-04	4 g in 1 PACKET; Type 0: Not a Combination Product	02/01/2016	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	08/20/2015	

**Labeler** - Sambria Pharmaceuticals (078676259)

Establishment			
Name	Address	ID/FEI	Business Operations
A.I.G. Technologies, Inc.		171837367	manufacture(54723-300)

Establishment			
Name	Address	ID/FEI	Business Operations
JP Packaging LLC		151369456	repack(54723-300)