

**NEUROMED FA- benzocaine cream**  
**Sambria Pharmaceuticals, Inc.**

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**NeuroMed FA Topical Analgesic**

***Active Ingredients***

Benzocaine 20.0% 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**

**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 or 4 times daily. Children under 2 years of age: consult a physician.

***Inactive Ingredients***

Aqua (Deionized Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Caprylic/Capric Triglyceride, Cetearietj-25, Chondroitin Sulfate, Diethylhexyl Sodium Sulfosuccinate, Emu Oil, Ethoxydiglycol, Ethylhexylglycerin, Glucosamine Sulfate, Glycerin, Isopropyl Palmitate, Laureth-7, Melaleuca alternifolia (Tea Tree) Leaf Oil, Methylfulfonylmenthane (MSM), Phenoxyethanol, Polyacrylamide, Polysorbate-20, Safflower Oil, Stearic Acid, Triethanolamine

***Other Information***

Protect this product from excessive heat or direct sun.

***Questions or Comments?***

FDA Registered: NDC No. 54723-668-03

800-759-6876

**Product label**





## NEUROMED FA

benzocaine cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:54723-722
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZOCAINE</b> (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	20 mg in 100 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ARNICA MONTANA FLOWER</b> (UNII: OZ0E5Y15PZ)	
<b>C13-14 ISOPARAFFIN</b> (UNII: E4F12ROE70)	

<b>MEDIUM-CHAIN TRIGLYCERIDES</b> (UNII: C9H2L21V7U)
<b>CETEARETH-25</b> (UNII: 8FA93U5T67)
<b>CHONDROITIN SULFATE SODIUM (BOVINE)</b> (UNII: 8QTV3DTT8W)
<b>DOCUSATE SODIUM</b> (UNII: F05Q2T2JA0)
<b>EMU OIL</b> (UNII: 344821WD61)
<b>DIETHYLENE GLYCOL MONOETHYL ETHER</b> (UNII: A1A1I8X02B)
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)
<b>GLUCOSAMINE SULFATE</b> (UNII: 1FW7WLR731)
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)
<b>ISOPROPYL PALMITATE</b> (UNII: 8CRQ2TH63M)
<b>LAURETH-7</b> (UNII: Z95S6G8201)
<b>MELALEUCA ALTERNIFOLIA LEAF</b> (UNII: G43C57162K)
<b>DIMETHYL SULFONE</b> (UNII: 9H4PO4Z4FT)
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)
<b>POLYACRYLAMIDE (10000 MW)</b> (UNII: E2KR9C9V2I)
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)
<b>SAFFLOWER OIL</b> (UNII: 65UEH262IS)
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54723-722-05	5000 mg in 1 PACKET; Type 0: Not a Combination Product	08/21/2017	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/21/2017	

**Labeler** - Sambria Pharmaceuticals, Inc. (078676259)

### Establishment

Name	Address	ID/FEI	Business Operations
Southeast Holdings Corp		080504027	manufacture(54723-722)

Revised: 11/2025

Sambria Pharmaceuticals, Inc.