

NEUROMED FA- benzocaine cream
Sambria Pharmaceuticals, Inc.

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
2% BENZOCAINE TOPICAL ANESTHETIC
3 ml / .10 fl.oz

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DRUG FACTS

Active Ingredients	Purpose
Benzocaine 20.0% w/w	External Anesthetic

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 5 to 4 times daily. Children under 2 years of age: consult a physician. Apply in circular motion for 30 to 60 seconds.

Inactive Ingredients
Aqua (Distilled Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Lactic Acid, Triethyl Citrate, Citronellol, Citronellal, Citral, Ethylhexylglycerin, Sodium Lauryl Sulfate, Ethyl Ol. Ethylhexylglycol, Ethylhexylglycerin, Glycerin, Isopropyl Alcohol, Isobutyl Alcohol, Laureth-7, Methylacrylate Copolymer (Res. Free) Oil, Methylparaben, Methylphenylisothiourea (MSM), Phenoxyethanol, Polysorbate-20, Polysorbate-80, Stearic Acid, Tetrahydrofuran.

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

Questions or Comments?
FDA Registered: NDC No. 54723-668-03
800-759-6876

Manufactured for: Serrbia Pharmaceuticals
1075 Peachtree St. NE, Ste. 3650, Atlanta, GA 30339
Made in the USA



CLOSE

NEUROMED FA

benzocaine cream

Product Information

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

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)		BENZOCAINE	20 mg in 100 mg	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)				
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)				
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)				
CETEARETH-25 (UNII: 8FA93U5T67)				
CHONDROITIN SULFATE SODIUM (BOVINE) (UNII: 8QTV3DTT8W)				
DOCUSATE SODIUM (UNII: F05Q2T2JA0)				
EMU OIL (UNII: 344821WD61)				
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A18X02B)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)				
GLYCERIN (UNII: PDC6A3C0OX)				
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)				
POLYSORBATE 20 (UNII: 7T1F30V5YH)				
SAFFLOWER OIL (UNII: 65UEH262IS)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
TROLAMINE (UNII: 9O3K93S3TK)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54723-722-03	3000 mg in 1 PACKET; Type 0: Not a Combination Product	02/01/2016	
2	NDC:54723-722-04	1500 mg in 1 PACKET; Type 0: Not a Combination Product	07/29/2024	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	02/01/2016		

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.