

NEUROMED TOPICAL ANALGESIC LA- tetracaine hcl cream
Sambria Pharmaceuticals, Inc.

NeuroMed LA Topical Analgesic

Active Ingredients

Tetracaine HCL 2.0% w/w

Purpose

External Analgesic

Uses

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

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

Inactive Ingredients

Aqua (Deionized Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Chondroitin Sulfate, Emu Oil, Ethoxydiglycol, Ethylhexylglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Melaleuca Alternifolia (Tea Tree) Leaf 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. 54723-667-03

800-759-6876

Product label

NEUROMEDTM LA
2% TETRACAINE LONG-LASTING,
TOPICAL ANESTHETIC

3 ml / .10 fl.oz

TETRACAINE

3

Drug Facts

Active Ingredients	Purpose
Tetracaine HCl, 2.0% w/v	External Anesthetic

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

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 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 a circular motion for 30 to 60 seconds.

Inactive Ingredients
Aqua (Deionized Water), Anise Montana Flower Extract, C15-14 Acipollin, Dodecyl Stearate, Emu Oil, Ethoxyglycol, Ethylhexylglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Methylcobalamin (MSP), Phenylethanol, Polycrylamide, Propylene Glycol, Stearic Acid, Triethanolamine

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

Questions or Comments?
FDA Registered: NDC No. 54723-667-03
800-759-6876

Manufactured for: Samba Pharmaceuticals
1075 Peachtree St. NE Ste. 3450, Atlanta, GA 30339
Made in the USA.



NEUROMED TOPICAL ANALGESIC LA

tetracaine hcl cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54723-310
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TETRACAINE HYDROCHLORIDE (UNII: 5NF5D4OPCI) (TETRACAINE - UNII:0619F35CGV)	TETRACAINE HYDROCHLORIDE	20 mg in 1000 mg

Inactive Ingredients

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

CHONDROITIN SULFATE SODIUM (BOVINE) (UNII: 8QTV3DTT8W)
EMU OIL (UNII: 344821WD61)
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A18X02B)
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)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
STEARIC ACID (UNII: 4ELV7Z65AP)
TROLAMINE (UNII: 9O3K93S3TK)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54723-310-03	3000 mg 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 Drug	M017	02/01/2016	

Labeler - Sambria Pharmaceuticals, Inc. (078676259)

Establishment

Name	Address	ID/FEI	Business Operations
A.I.G. Technologies, Inc.		086365223	manufacture(54723-310)

Revised: 11/2023

Sambria Pharmaceuticals, Inc.