

NEURACIN TOPICAL GEL- neuracin topical analgesic gel gel

SOLA 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.

Neuracin Topical Analgesic Gel

Active Ingredients

Camphor 4%

Menthol 10%

Methyl Salicylate 30%

Purpose

Topical Analgesic

Uses

Temporarily relieves minor pain associated with:

- Arthritis
- Simple Backache
- Muscle Strains
- Bruises
- Muscle Sprains

Warnings

For External use only

When using this product

- avoid contact with eyes and mucous membranes
- do not apply to wounds or damaged skin
- do not bandage tightly or use with a heating pad
- use only as directed

Stop use and ask a doctor if condition worsens; symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of children. If swallowed, get medical help right away. If pregnant Or Breast-Feeding, Ask A Health Professional Before Use

Directions

- Apply on affected area, not more than 3 to 4 times daily

- Children under 12 years of age; consult a doctor

Other Information

- Store at 20-25°C (68-77°F) Store away from heat

Inactive Ingredients

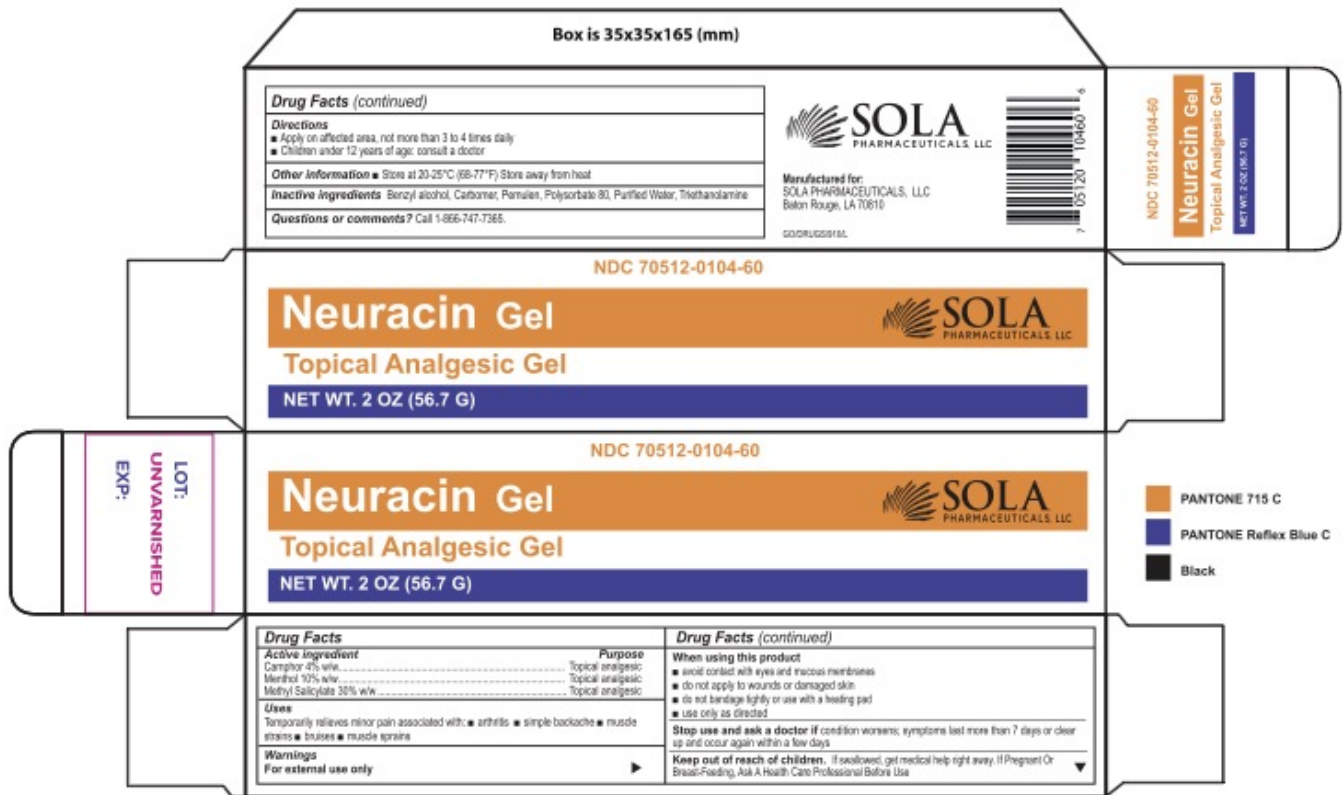
Benzyl alcohol, Carbomer, Pemulen, Polysorbate 80, Purified Water, Triethanolamine

Questions or comments? Call 1-866-747-7365

Neuracin Topical Analgesic Gel

NDC 70512-104-60

Qty 2oz (56.7G)



NEURACIN TOPICAL GEL

neuracin topical analgesic gel gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70512-104
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	300 mg in 1 g
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	40 mg in 1 g
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	100 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70512-104-60	1 in 1 CARTON	05/24/2021	
1		56.7 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	05/24/2021	

Labeler - SOLA Pharmaceuticals (080121345)

Revised: 7/2021

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