NERVE ACTIVE PAIN RELIEF CREAM- menthol cream TOP DOG DIRECT, LLC

NERVE ACTIVE Pain Relief Cream

Drug Facts

Active Ingredients

Menthol 3.00%

Purpose

Topical Analgesic

Uses:

• For the temporary relief of minor aches and pains of muscles and joints, associated with simple backache, arthritis, strains, bruises, and sprains.

Warnings:

For external use only

Do not use

• on damaged or broken skin.

When using this product

- Avoid contact with the eyes.
- Do not bandage tightly.

Stop use and ask a doctor if

- rash or irritation develops and lasts.
- condition worsens, or if symptoms persist for more than 7 days or clear up and occur agani within a few days.

Keep out of reach of children.

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

If pregnant or breast-feeding,

ask a health professional before use

Directions:

 Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily. • Children under 2 years of age: consult a doctor.

Other information:

• Protect the product in this container from excessive head and direct sun.

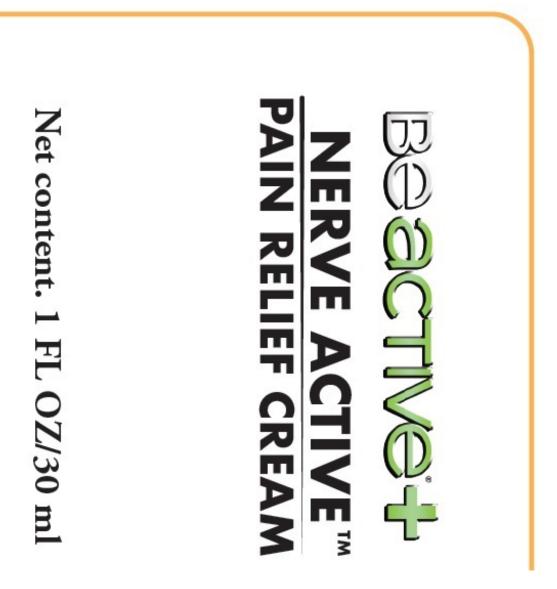
Inactive ingredients:

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, Cetyl Alcohol, Ethylhexylglycerin, Glyceryl Stearate, Glycyrrhiza Glabra (Licorice) Extract, Ilex Paraguariensis (Yerba Mate') Extract, Isopropyl Myristate, Magnesium Sulfate, Methyl Salicylate, PEG-8, Phenoxyethanol, Polysorbate-20, Triethanolamine.

Questions?

800-340-3418

Package Labeling:



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NERVE ACTIVE PAIN RELIEF CREAM

menthol cream

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:83690-422

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	30 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
TROLAMINE (UNII: 903K93S3TK)	

Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83690-422- 00	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/18/2023	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	09/18/2023		

Labeler - TOP DOG DIRECT, LLC (011971091)

Revised: 11/2023 TOP DOG DIRECT, LLC