NEURAXCIN NEUROMETHACIN- menthol roll-on liquid
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.

Neuraxcin Neuromethacin, Pain Relief Roll-on

Menthol

Topical Analgesic

Indications

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

Warnings

-For external use only.

-Avoid contact with eyes.

-If symptoms persist for more than seven days, discontinue use and consult physician.

Keep out of reach of children

-If swallowed, consult physician.

-Do not apply to wounds or damaged skin.

-Do not bandage tightly.

If pregnant or breast feeding

Contact physician prior to use.

Directions

-Adults and children two-years of age or older. Apply to affected area not more than three to four times daily.

-Children under two-years of age, consult a physician.

Storage

Store at room temperature.

Other Ingredients

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montant Flower Extract, Boswellia Serrata Extract, Camellia Sinensis (Green Tea) Extract, Cannabis Sativa (Full Spectrum Hemp) Oil, Caprylyl Glycol, Cetearyl Olivate, Cetyl Alcohol, Citric Acid, Glucosamine Sulfate, Glycerin, Glyceryl Stearate, Helianthus Annus (Sunflower) Oil, Isopropyl Alcohol, Isopropyl Myristate, Methylsufonylmethane (MSM), Phenoxyethanol, Sorbitan Olivate, Stearic Acid, Xanthan Gum.

NEURAXCIN NEUROMETHACIN

menthol roll-on liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70512-105
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)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
WATER (UNII: 059QF0KO0R)	

INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70512-105-90	90 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	12/10/2020	

Labeler - Sola Pharmaceuticals (080121345)

Revised: 12/2020 Sola Pharmaceuticals