# NUTRASTATE SOOTHING RELIEF ROLL-ON GEL- menthol gel NUTRASTATE, LLC

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.

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#### **Active ingredient**

#### Active ingredient Purpose

Menthol 4%......Topical Analgesic

#### Uses

**Uses** Temporary relief from minor aches and pains of muscle aches and pains of muscles and joints associated with arthritis, simple backache, strains and sprains

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#### Warnings

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For external use only.

Flammable: Do not use while smoking or near heat or flame.

## When using this product

When using this product ● avoid contact with the eyes or mucous membranes ● do not apply to wounds or damaged skin ● do not apply to the irritate skin ● do not bandage ● wash hands after use with cool water ● do not use with heating pad or device

## Stop use and ask a doctor if

**Stop use and ask a doctor if** ● condition worsens, or it symptoms persist for more than 7 days, or clear up and occur again within a few days.

## Keep out of reach of children

**Keep out of reach of children.** If accidentally ingested get medical help or contact a Poison Control Center immediately.

Directions ● Adults and children 2 years of age and older:

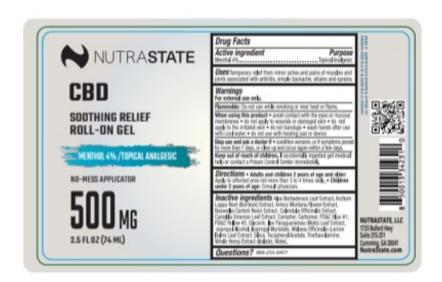
Apply to affected area not more than 3 to 4 times daily. ● Children under 2 years of age: Consult a physician.

#### **Inactive ingredients**

Inactive ingredients Aloe Barbadensis Leaf Extract, Arctium Lappa Root (Burdock) Extract, Arnica Montana Flower Extract, Boswellia Carterii ResinExtract, Calendula Officinalis Extract, Camellia Sinensis Leaf Extract, Camphor Carbomer, FD&C Blue #1, FD&C Yellow #5, Glycerin, Ilex Paraguariensis (Mate) Leaf Extract, Isopropyl Alcohol, Isopropyl Myristate, Melissa Officinalis (Lemon Balm) Leaf Extract, Silica, Tocopheryl Acetate, Triethanolamine, Whole Hemp Extract (Isolate), Water.

#### Questions?

**Questions?** 888-253-0477



#### NUTRASTATE SOOTHING RELIEF ROLL-ON GEL menthol gel **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:81178-001 **Route of Administration TOPICAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) **MENTHOL** 4 g in 100 g **Inactive Ingredients Ingredient Name** Strength FRANKINCENSE (UNII: R9XLF1R1WM)

CALENDULA OFFICINALIS FLOWER (UNII: POM7O4Y7YD)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 903K93S3TK)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)	
WATER (UNII: 059QF0KO0R)	
HEMP (UNII: TD1MUT01Q7)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
<b>CAMPHOR, (-)-</b> (UNII: 213N3S8275)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
MELISSA OFFICINALIS LEAF (UNII: 50D2ZE9219)	

Packaging							
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
		NDC:81178- 001-25	381 g in 1 BOTTLE, WTH APPLICATOR; Type 0: Not a Combination Product	11/23/2020			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part348	11/23/2020				

## Labeler - NUTRASTATE, LLC (114472928)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
InSpec Solutions, LLC		081030372	manufacture(81178-001)				

Revised: 7/2021 NUTRASTATE, LLC