

MINTED LEAF COLD THERAPY PAIN RELIEF- menthol gel
MMG Consumer Brands, LLC

Minted Leaf Cold Therapy Pain Relief Gel

Drug Facts

Active Ingredient:

Menthol 4.00%

Purpose

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.

Additional Information:

Store at room temperature.

Other Ingredients:

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arctium Lappa (Burdock) Root Extract, Arnica Montana Flower Extract, Boswellia Serrata Extract, Camellia Sinensis (Green Tea) Extract, Cannabis Sativa (Broad Spectrum Hemp), Copper Sulfate, Ethylhexylglycerin, Ilex Paraguariensis (Yerba Mate') Extract, Isopropyl Alcohol, Isopropyl Myristate, Melissa Officinalis (Lemon Balm) Extract, Methylsulfonylmethane (MSM), Phenoxyethanol, Sodium Hydroxide, Tocopheryl Acetate (Vitamin E), Zemea (Corn) Propanediol.

Package Labeling:

mintedLeaf™

**Cold Therapy
Menthol 4%
Pain Relief**

+ Hemp Extract
150 mg

Cooling
Menthol
Formula

Printing Area

GEL
3 OZ

FAST ACTING

Drug Facts

Active ingredient	Purpose
Menthol 4.00%	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.

Additional Information ■ Store at room temperature.

Other Ingredients: Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arctium Lappa (Burdock) Root Extract, Arnica Montana Flower Extract, Boswellia Serrata Extract, Camellia Sinensis (Green Tea) Extract, Cannabis Sativa (Broad Spectrum Hemp), Copper Sulfate, Ethylhexylglycerin, Ilex Paraguariensis (Yerba Mate) Extract, Isopropyl Alcohol, Isopropyl Myristate, Melissa Officinalis (Lemon Balm) Extract, Methylsulfonylmethane (MSM), Phenoxyethanol, Sodium Hydroxide, Tocopheryl Acetate (Vitamin E), Zemea (Corn) Propanediol.

Item# 10002



Scan for 3rd party lab reports

Distributed by:
MMG Consumer Brands, LLC
Pinecrest, FL.
www.mintedLeafHemp.com



MINTED LEAF COLD THERAPY PAIN RELIEF

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73102-115
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	40 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CANNABIS SATIVA SUBSP. SATIVA FLOWERING TOP (UNII: 8X454SZ22D)	
CUPRIC SULFATE (UNII: LRX7AJ16DT)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
MELISSA OFFICINALIS (UNII: YF70189L0N)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CORN (UNII: 0N8672707O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73102-115-03	88 mL in 1 TUBE; Type 0: Not a Combination Product	05/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/01/2019	

Labeler - MMG Consumer Brands, LLC (117036455)