MYDERM PRO FREEZE COOLING MENTHOL PAIN RELIEF ROLL-ON-
menthol gel
Inspec Solutions

MyDerm Pro Freeze Cooling Menthol Pain Relief Roll-On (IS0252)

Active Ingredient

Menthol 6%

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

Menthol 6%----------Topical Analgesic

For external use only.

Stop use and ask doctor if condition worsens, or if symtoms persist for more than 7 days, or clean up and reoccur again within a few days.

keep out of reach of children if swallowed,get medical help or contact a poson Control Center Immediately.

Inactive Ingredient

Aloe Barbadensis Leaf Juice

Arctium Lappa Root Extract

Arnica Montana Flower Extract

Boswellia Carterii resin Extract

Calendula Officinalis Extract

Camellia Sinensis Leaf Extract

Camphor

Carbomer

FD & C Blue #1

FD & C Yellow #5

Glycerin

Isopropyl Alcohol

Isopropyl Myristate

Ilex Paraguariensis Leaf Extract

Melissa Officinalis Leaf Extract

Phenoxyethanol

Silicon dioxide

Tocopheryl Acetate

Triethanolamine

Water

Direction: Apply to affectered area not more that 3 to 4 times daily.

150252

myDerm™ PRO

FREEZE

cooling menthol

PAIN RELIEF ROLL-ON

50% MORE MENTHOL!

3 FL OZ (89 mL)

Drug Facts

Active ingredient	Purpose
Menthol 6%	Topical Analgesic

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

Warnings

For external use only.

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

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

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

Keep out of reach of children. If swallowed, 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 physician.

Inactive ingredients Aloe Barbadensis Leaf Extract, Arctium Lappa Root (Burdock) Extract, Arnica Montana Flower Extract, Boswellia Carterii Resin Extract, 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, Water.

Questions? 1-855-MYDERM1

Distributed by: myDerm
Daytona Beach, FL 32117
1-855-MYDERM1
www.myderm.com

MYDERM PRO FREEZE COOLING MENTHOL PAIN RELIEF ROLL-ON

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72667-073
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	6 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CAMPHOR, (-)- (UNII: 213N3S8275)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
FRANKINCENSE (UNII: R9XLF1R1WM)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
MELISSA OFFICINALIS LEAF (UNII: 50D2ZE9219)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72667-073-01	89 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/06/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/06/2024	

Labeler - Inspec Solutions (081030372)

Establishment

Name	Address	ID/FEI	Business Operations
Inspec Solutions		081030372	manufacture(72667-073)

Revised: 6/2024

Inspec Solutions