MYDERM COOLING MENTHOL PAIN RELIEF WITH 50 PERCENT MORE MENTHOL- menthol spray Inspec Solutions 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.

myDerm Cooling Menthol Pain Relief Spray with 50% More Menthol Active ingredient

Menthol 10.5%

Menthol 10.4% Topical Analgesic

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

For external use only

When using this product, avoid contact with eyes and mucous membranes., do not apply to wound or damage skin, do not bandage, do not use with heating pad or device.

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

Keep out of reach of children, if swallowed, get medical help or contact a Poison Control Center right away.

Adults and children 2 years of age and older: Apply to affected area no more than 3 to 4 times daily. Wash hands with soap. Children under 2 years of age: consult a doctor.

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

Full Spectrum Industrial Hemp Extract

Glycerin

Ilex Paraguariensis (Mate) Leaf Extract

Isopropyl Alcohol

Isopropyl Myristate

Melissa Officinalis (Lemon Balm) Leaf Extract

Silica

Tocopheryl Acetate

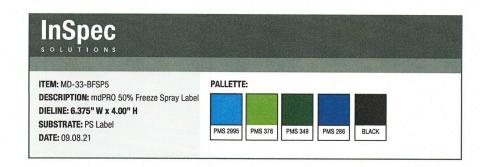
Triethanolamine

Water



4.00"

Trim Line
Text Safe Line
Bleed Line



MENTHOL

menthol spray

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Prod	uct	Intori	mation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72667-020

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) MENTHOL 15.75 g in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
FRANKINCENSE (UNII: R9XLF1R1WM)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
GLYCERIN (UNII: PDC6A3C0OX)	
MELISSA OFFICINALIS LEAF (UNII: 50D2ZE9219)	
TROLAMINE (UNII: 903K93S3TK)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CAMPHOR (NATURAL) (UNII: N20HL7Q941)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

l	Packaging				
	#	Item Code Package Description		Marketing Start Date	Marketing End Date
	1	NDC:72667- 020-01	89 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/27/2021	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	12/27/2021		

Labeler - Inspec Solutions LLC (081030372)

Establishment				
Name	Address	ID/FEI	Business Operations	
Inspec Solutions LLC.		081030372	manufacture(72667-020)	

Revised: 1/2023 Inspec Solutions LLC