

COLD SPOT POINT RELIEF- menthol, methyl salicylate gel Fabrication Enterprises

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.

Cold Spot Point Relief Pain relieving Gel - 32 oz.

Active Ingredients: Menthol, methyl salicylate

Inactive Ingredients: deionized water, arnica, chondroitin sulfate, citric acid, euclayptus oil, glucosamine sulfate, ilex paraguariesis leaf, isopropyl alcohol, peppermint oil, dimethyl sulfone, polysorbate-20, SD alcohol 40B.

Keep out of reach of children. If swallowed consult physician

Warnings Section: For external use only, avoid contact with eyes, do not apply to open wounds or damaged skin, if symptoms persist for more than seven days discontinue use and consult physician, keep out of reach of children and if swallowed consult physician, do not bandage tightly.

pain relieving gel

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

Apply directly to effected area. Do not use more than four times per day.

ColdSpot Point Relief Pain Relieving spray, all natural ingredients.

Drug Facts	
Active Ingredients: menthol - USP 1.2% methyl salicylate 4%	Purpose: external analgesic external analgesic
Uses: For temporary relief of minor aches and pains of the muscles and joints associated with simple backache, arthritis, bruises, strains and/or sprains.	
Warnings: <ul style="list-style-type: none">• For external use only• Avoid contact with eyes• Do not apply to open wounds or damaged skin• 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 bandage tightly	
Directions: Apply directly to affected area. Do not use more than four times per day.	
Other Ingredients: aqua (deionized water), arnica montana flower (arnica) extract, boswellia serrata extract, bromelain, carbomer, diazolidinyl urea, eucalyptus globulus oil, glycyrrhiza glabra (licorice) extract, ilex paraguariensis leaf (yerba mate) extract, mentha piperita (peppermint) oil, methyl paraben, MSM (dimethyl sulfone), polysorbate-80, propyl paraben, propylene glycol, SD-alcohol 40B, triethanolamine	

Manufactured For:
Fabrication Enterprises Inc.
Post Office Box 1500
White Plains, New York 10602 USA
tel: 914-345-9300 fax: 914-345-9800
www.FabricationEnterprises.com

soothing menthol relief

ColdSpot™

POINT RELIEF™

pain relieving gel

all natural ingredients

really hits the spot

net wt. 32fl. oz. / 960mL.

Authorized CE representative: RMS UK Ltd, 28 Trinity Road, Nailsea, Somerset BS48 4NU (UK) ©FEI 2010, all rights reserved. ColdSpot™ and Point Relief™ are trademarks of FEI

11-0711-1 ColdSpot™ 32oz. gel

7 14905 02507 4

NDC 61462-001-32

FEI FABRICATION ENTERPRISES INC. made in USA

COLD SPOT POINT RELIEF
menthol, methyl salicylate gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51452-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EP3A) (MENTHOL - UNII:L7T10EP3A)	MENTHOL	120 mg in 1 mL
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ARNICA CORDIFOLIA FLOWER (UNII: JCG1OSZ7A8)	
CARBOMER 1342 (UNII: 809Y72KV36)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51452-001-32	960 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/24/2010	02/01/2021

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	08/24/2010	02/01/2021

Labeler - Fabrication Enterprises (070577218)**Registrant** - Fabrication Enterprises (070577218)**Establishment**

Name	Address	ID/FEI	Business Operations
Pure Source, LLC		080354456	manufacture(51452-001)

