LIBERDOL TOPICAL ANALGESIC- methyl salicylate, menthol, camphor (synthetic) spray ViaDerma Distribution, Inc

Liberdol Topical Analgesic Spray

Drug Facts

Active Ingredient

Methyl Salicylate 300mg Menthol 100mg

Camphor 40mg

Purpose

Analgesic

Uses

Soothing on-the-go tempoary relief from minor aches and pains of sore muscles and joint associated with:

- arthritis
- backache
- strains
- sprains

Warnings

For external use only. : Keep away form excessive heat or open flame Flammable

Ask a doctor before use if you have:

Sensitive skin, are pregnant or are bleeding

When using this product:

- Avoid contact with eyes or mucus membranes
- Do not apply to open wounds or damaged skin
- Do not use with other ointments, creams, sprays or liniments
- Do not apply to irritated skin, or if excessive irritation develops
- Do not bandage
- Do not use with heating pad or device

Stop use and ask a doctor if:

Condition worsens or if symptoms persist more than 7 days, or clear up or reoccur

Keep out of reach of children.

If swallowed, get medical help or contact a Poison control Center right away

Directions

- Apply to affected areas not more than 4 times daily Adults and children 2 years of age an older:
- Consult a physician Children under 2 years of age:

Other Information

• Store in a cool dry place out of direct sunlight

Inactive Ingredients

acetic acid, arnica montana flower extract, ascorbic acid, chlorhexidine gluconate, cholecalciferol, dimethyl sulfoxide, dipropylene glycol, glucono delta lactone, glycerin, histidine, hydroxethyl cellulose, magnesium stearate, sodium hydroxide, sorbic acid, stearic acid, water

Package Labeling:





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KEEP THIS AND ALL DRUGS AWAY FROM CHILDREN. IN CASE OF ACCIDENTAL INGESTION, SEEK MEDICAL ASSISTANCE OR CONTACT A POSION CONTROL IMMEDIATELY.



BERDOL

Topical Analgesic

Analgesic Spray

30% Methyl Salicylate 10% Menthol 4% Camphor

(active ingredients)







L/BERDOL

Topical Analgesic

Analgesic Spray

30% Methyl Salicylate 10% Menthol 4% Camphor

(active ingredients)



1.85 FL OZ (55ml)

Licensed to ViaDerma Distribution by: ViaDerma, Inc. Distributed by: ViaDerma Distribution Report any side effects to: ViaDerma Distribution 2235 E. Flamingo Rd, Suite 152 Las Vegas, NV 89119 Tel: 1-800-585-8685 | viadermainc.com



NDC# 71262-002-011



L/BERDOL

Topical Analgesic

Analgesic Spray

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ViaDerma, Inc. Distributed by: **ViaDern** icensed to **ViaDerma Distribution** by



BERDO

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Analgesic Spray

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MEDICAL ASSISTANCE OR CONTACT **KEEP THIS AND ALL DRUGS AWAY** POISON CONTROL IMMEDIATEL FROM CHILDREN. IN CASE OF

LIBERDOL TOPICAL ANALGESIC

methyl salicylate, menthol, camphor (synthetic) spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71262-008

Route of Administration TOPICAL

Active Ingredient/Active Moiety Basis of Ingredient Name Strength Strength 300 mg METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ) METHYL SALICYLATE in 1 mL 100 mg MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) **MENTHOL** in 1 mL CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) -**CAMPHOR** 40 mg UNII:5TJD82A1ET) (SYNTHETIC) in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
ACETIC ACID (UNII: Q40Q9N063P)		
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)		
ASCORBIC ACID (UNII: PQ6CK8PD0R)		
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)		
CHOLECALCIFEROL (UNII: 1C6V77QF41)		
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)		
DIPROPYLENE GLYCOL (UNII: E107L85C40)		
GLUCONOLACTONE (UNII: WQ29KQ9POT)		
GLYCERIN (UNII: PDC6A3C0OX)		

HISTIDINE (UNII: 4QD397987E)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SORBIC ACID (UNII: X045WJ989B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
WATER (UNII: 059QF0KO0R)	

ı	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:71262-008- 55	1 in 1 CARTON	06/01/2021		
	L	55 mL in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	06/01/2021		

Labeler - ViaDerma Distribution, Inc (081113521)

Revised: 12/2023 ViaDerma Distribution, Inc