

LIBERDOL ANALGESIC- menthol, unspecified form, methyl salicylate liquid
ViaDerma Distribution, Inc.

Liberdol Analgesic Spray

Drug Facts

Active ingredients

Camphor 4% Menthol 10%

Methyl Salicylate 30%

Purpose

Topical Analgesic

Use

- For the temporary relief of minor aches and pains of muscles and joints

Warnings

For external use only

When using this product

- avoid contact with eyes or mucous membranes
- do not apply to wounds or damaged skin
- do not bandage tightly
- do not use otherwise than as directed

Stop use and ask a doctor if

- condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days
- excessive irritation of the skin develops

Keep out of reach of children to avoid accidental poisoning.

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

Directions

- Adults and children 3 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- Children under 3 years of age: Consult a doctor.

Other information

- Store in a cool dry place out of direct sunlight

Inactive ingredients

acetic acid, arnica 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

Questions or Comments?

1-800-585-8685

Package Labeling:

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Drug Facts (continued)

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Questions or Comments?

1-800-585-8685

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

Analgesic Spray
Topical Analgesic
LIBERDOL™
VIADERMA®
Distribution

VIADERMA®
Distribution
LIBERDOL™
Topical Analgesic
Analgesic Spray
30% Methyl Salicylate
10% Menthol
4% Camphor
(active ingredients)

4 fl oz (118 mL)

VIADERMA®
Distribution

LIBERDOL™
Topical Analgesic
Analgesic Spray

30% Methyl Salicylate
10% Menthol
4% Camphor
(active ingredients)



4 fl oz (118 mL)

LICENSED TO VIADERMA DISTRIBUTION BY: VIADERMA, INC.
DISTRIBUTED BY: VIADERMA DISTRIBUTION
2235 E. FLAMINGO RD, SUITE 152, LAS VEGAS, NV 89119
TO REPORT A SERIOUS ADVERSE EVENT, CONTACT 1-800-585-8685
WWW.VIADERMA.NET



NDC# 71262-002-011

LICENSED TO VIADERMA DISTRIBUTION BY: VIADERMA, INC.
 DISTRIBUTED BY: VIADERMA DISTRIBUTION
 2235 E. FLAMINGO RD, SUITE 152, LAS VEGAS, NV 89119
 TO REPORT A SERIOUS ADVERSE EVENT, CONTACT 1-800-585-8685
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VIADERMA[®]
Distribution

LIBERDOL[™]
 Topical Analgesic

Analgesic Spray

**30% Methyl Salicylate,
 10% Menthol, 4% Camphor**

4 fl oz (118 mL)

THIS UNIT NOT LABELED FOR RETAIL SALE

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 CHILDREN. IN CASE OF ACCIDENTAL INGESTION,
 SEEK MEDICAL ASSISTANCE OR CONTACT
 A POISON CONTROL IMMEDIATELY.

LIBERDOL ANALGESIC

menthol, unspecified form, methyl salicylate liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71262-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
GLYCERIN (UNII: PDC6A3C0OX)	
HISTIDINE (UNII: 4QD397987E)	
MAGNESIUM STEARATE (UNII: 70097M6130)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

SORBIC ACID (UNII: X045WJ989B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71262-004-18	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/01/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/01/2019	

Labeler - ViaDerma Distribution, Inc. (081113521)

Revised: 12/2023

ViaDerma Distribution, Inc.