

METHYL SALICYLATE 30 PERCENT CREAM- methyl salicylate cream
Alexso, Inc

Methyl Salicylate 30 Percent Cream

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Alexso, Inc

Methyl Salicylate 30% Cream

Drug Facts

Active ingredient

Methyl Salicylate 30%

Purpose

Topical analgesic

Uses

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

Warnings

For external use only.

When using this product

- Avoid contact with the eyes
- Do not use in large quantities, particularly over raw surfaces or blistered areas
- Do not apply to wounds or damaged skin
- Do not bandage tightly

Stop use and ask a doctor if

- allergic reaction occurs
- condition worsens or does not improve within 7 days
- symptoms clear up and return within a few days
- redness, irritation, swelling, pain or other symptoms begin or increase

Keep out of reach of children.

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

Directions

adults and children 2 years and older	apply externally to the affected area up to 3 to 4 times a day
children under 2 years	ask a doctor

Other information

- Store at 20-25°C (68-77°F); excursions permitted to 15-30°C (59-86°F). See USP Controlled Room Temperature.

Inactive ingredients

Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, Cetearyl Alcohol, Chondroitin Sulfate, Ethylhexylglycerin, Glucosamine Sulfate, Glycerin, Glyceryl Stearate, C13-14 Isoparaffin, Isostearyl Palmitate, Laureth-7, Methylsulfonylmethane (MSM), PEG-100 Stearate, Phenoxyethanol, Polyacrylamide, Propylene Glycol, Sodium Polyacrylate, Stearic Acid, Triethanolamine

PRINCIPAL DISPLAY PANEL

Methyl Salicylate 30% Cream

NDC: 50488-1030-6

60 grams

PRINCIPAL DISPLAY PANEL

Methyl Salicylate 30% Cream

NDC: 50488-1030-1

120 grams

Manufactured for:
Alexso, Inc
Los Angeles, CA 90064

Drug Facts

Active ingredient **Purpose**
Methyl Salicylate 30%.....Topical Analgesic

Uses ■ For the temporary relief of minor aches and pains of muscles and joints, associated with simple backache, arthritis, strains, bruises, and sprains.

Warnings
For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center.

When using this product ■ Avoid contact with the eyes ■ Do not use in larger quantities, particularly over raw surfaces or blistered areas ■ Do not apply to wounds or damaged skin ■ Do not bandage tightly

Stop use and ask a doctor if
■ allergic reaction occurs ■ condition worsens or does not improve within 7 days ■ symptoms clear up and return within a few days ■ redness, irritation, swelling, pain or other symptoms begin or increase ▶

Methyl Salicylate 30% Cream
2 Fl Oz (60mL)
Topical Analgesic
NDC: 50488-1030-6
Rev 09/25


Drug Facts (continued)

Directions ■ Adults and children 2 years and older apply externally to the affected area up to 3 to 4 times a day ■ Children under 2 years ask a doctor

Other information Store at 0°-25°C (68°-77°F); excursions permitted to 15°-30°C (59°-86°F). See USP Controlled Room Temperature.

Inactive ingredients Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, Cetearyl Alcohol, Chondroitin Sulfate, Ethyl- hexylglycerin, Glucosamine Sulfate, Glycerin, Glyceryl Stearate, C13-14 Isoparaffin, Isostearyl Palmitate, Laureth-7, Methylsulfonylmethane (MSM), PEG-100 Stearate, Phenoxyethanol, Polyacrylamide, Propylene Glycol, Sodium Polyacrylate, Stearic Acid, Triethanolamine.

Manufactured for:
Alexso inc.
2317 Cotner Avenue
Los Angeles, CA 90064
Tel: 888.495.6078



METHYL SALICYLATE 30 PERCENT CREAM

methyl salicylate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50488-1030
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	30 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
BOSWELLIA SERRATA GUM (UNII: 4PW41QCO2M)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
ISOSTEARYL PALMITATE (UNII: 9EHU0R7ER1)	
LAURETH-7 (UNII: Z95S6G8201)	

DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIETHANOLAMINE (UNII: 9O3K93S3TK)	
CHONDROITIN SULFATE (BOVINE) (UNII: 6IC1M3OG5Z)	
POLYACRYLAMIDE (10000 MW) (UNII: E2KR9C9V2I)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50488-1030-1	120 g in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2025	
2	NDC:50488-1030-6	60 g in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/01/2025	

Labeler - Alexso, Inc (963338061)

Revised: 11/2025

Alexso, Inc