FLEXDERMAL PAIN RELIEVING ROLL-ON- menthol liquid SOLVADERM LLC

FlexDermal Pain Relieving Roll-On Ointment

Drug Facts

Active ingredients

Menthol 2.00%

Purpose

Topical Analgesic

Uses

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

Warnings

For external use only

Do not

- apply to wounds or damaged skin.
- bandage tightly.

When using this product

Avoid contact with the eyes.

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.

Keep out of reach of children.

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

If pregnant or breast-feeding,

ask a health professional before use

Directions

- Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- Children under 2 years of age: consult a doctor.

Other information

• Protect the product in this container from excessive heat and direct sun.

Inactive Ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Blue 1 (CI 42090), Camphor, Ethylhexylglycerin, Glycerin, Isopropyl Alcohol, Phenoxyethanol, Sodium Hydroxide, Water(Aqua)

Questions?

888-211-8468

Package Labeling:

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Apply directly to the skin, ensuring the product covers the affected area. Massage gently into the skin until fully absorbed. Adults and children 12 years or older apply 3-4 times a day.

Caution: Discontinue use if redness or irritation occurs.

Do not ingest.



PAIN RELIEVING ROLL-ON OINTMENT

MENTHOL CAMPHOR GLYCERIN

3 FL OZ (90mL)

flexdermal.com

Manufactured for and distributed by Pharmaxa Labs 80 Red Schoolhouse Road Spring Valley, NY 10977

> NOT AVAILABLE FOR RESALE UNLESS APPROVED BY PHARMAXA LABS

FLEXDERMAL PAIN RELIEVING ROLL-ON

menthol liquid

Product Information	roduct Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84197-867	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	20 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
GLYCERIN (UNII: PDC6A3C0OX)		
ISOPROPYL ALCOHOL (UNII: ND2M416302)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
WATER (UNII: 059QF0KO0R)		

ı	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:84197-867- 00	90 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/24/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/24/2025	

Labeler - SOLVADERM LLC (093262409)

Revised: 2/2025 SOLVADERM LLC