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.

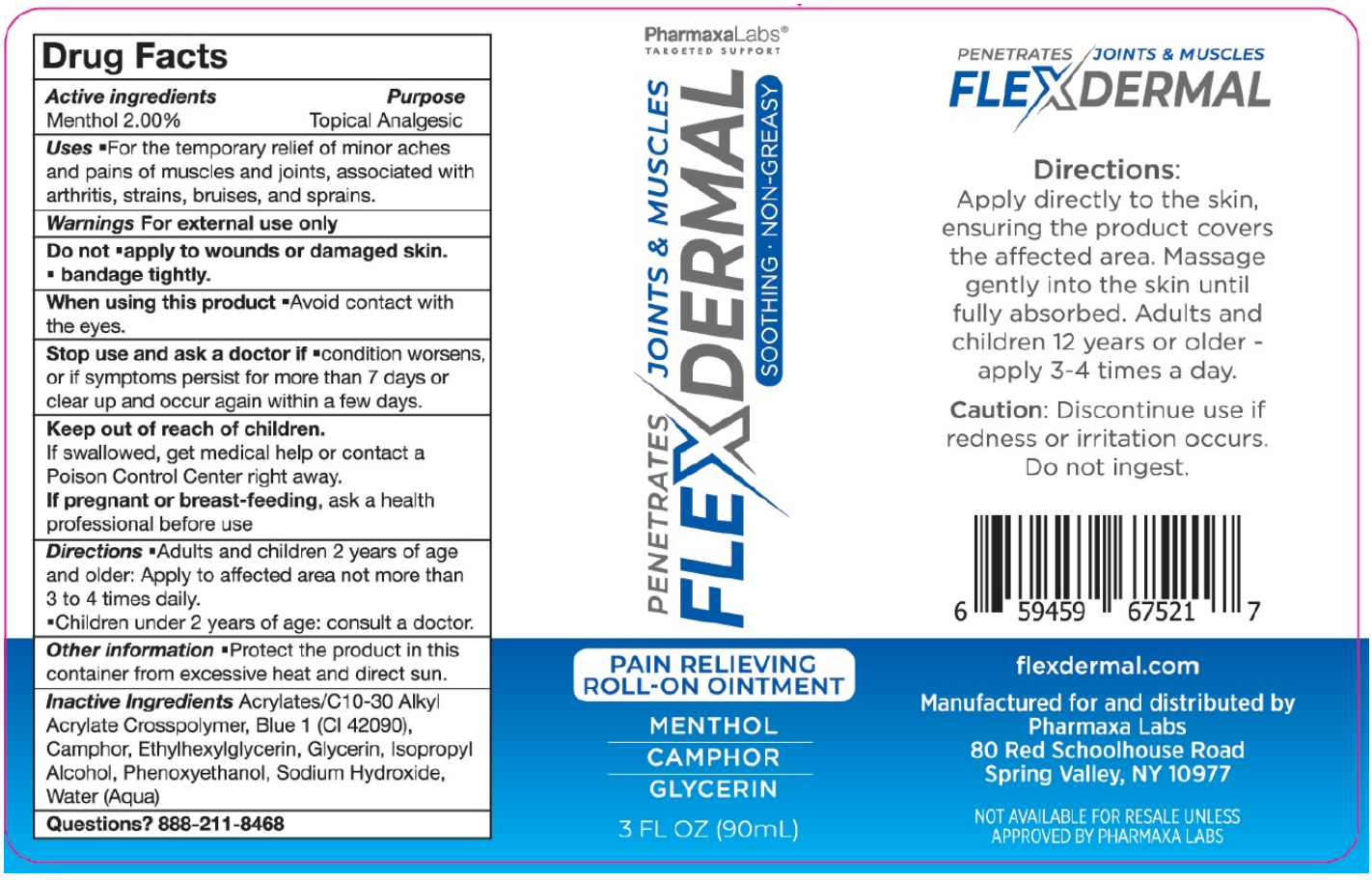
Inactive Ingredients

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:



FLEXDERMAL PAIN RELIEVING ROLL-ON

menthol liquid

Product 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)	

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