SUNMED MOTION- lidocaine liquid
Sunflora Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

SUNMED MOTION

Drug Facts

Active Ingredients

Lidocaine 4%

Purpose

Topical Analgesic

Uses

Temporarily relieves minor pain associated with:

• arthritis • sprains • simple backache • muscle strains • cramps • bruises

Warnings

For external use only

When using this product

• use only as directed • do not bandage tightly • avoid contact with eyes • do not apply to wounds or damaged skin • do not use in large quantities, particularly over raw surfaces or blistered areas.

Stop use and ask a doctor if

• condition worsens • symptoms persist for more than 7 days • symptoms 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.

Directions

• Adults and children 2 years of age and older: Apply to the affected area not more than 3 to 4 times daily.

• Children under 2 years of age: Consult a doctor.

Inactive Ingredients

Allantoin, Caprylic/Capric Triglyceride, Capsicum Annuum Fruit Powder, Calcium Disodium EDTA, Cannabis Sativa (Aerial) Extract Oil, Dimethicone, dl-alpha Tocopheryl Acetate, Ethyl Alcohol, Ethylhexylglycerin, Ethylhexyl Stearate, Fragrance, Glycerin, Phenoxyethanol, Polysorbate 80, Propanediol, Propylene Glycol, Purified Water, Simethicone, Sodium Polyacrylate, Trideceth-6, Vitis

Vinifera (Grape) Seed Oil, Xanthan Gum.

Visit us at SUNFLORA.org

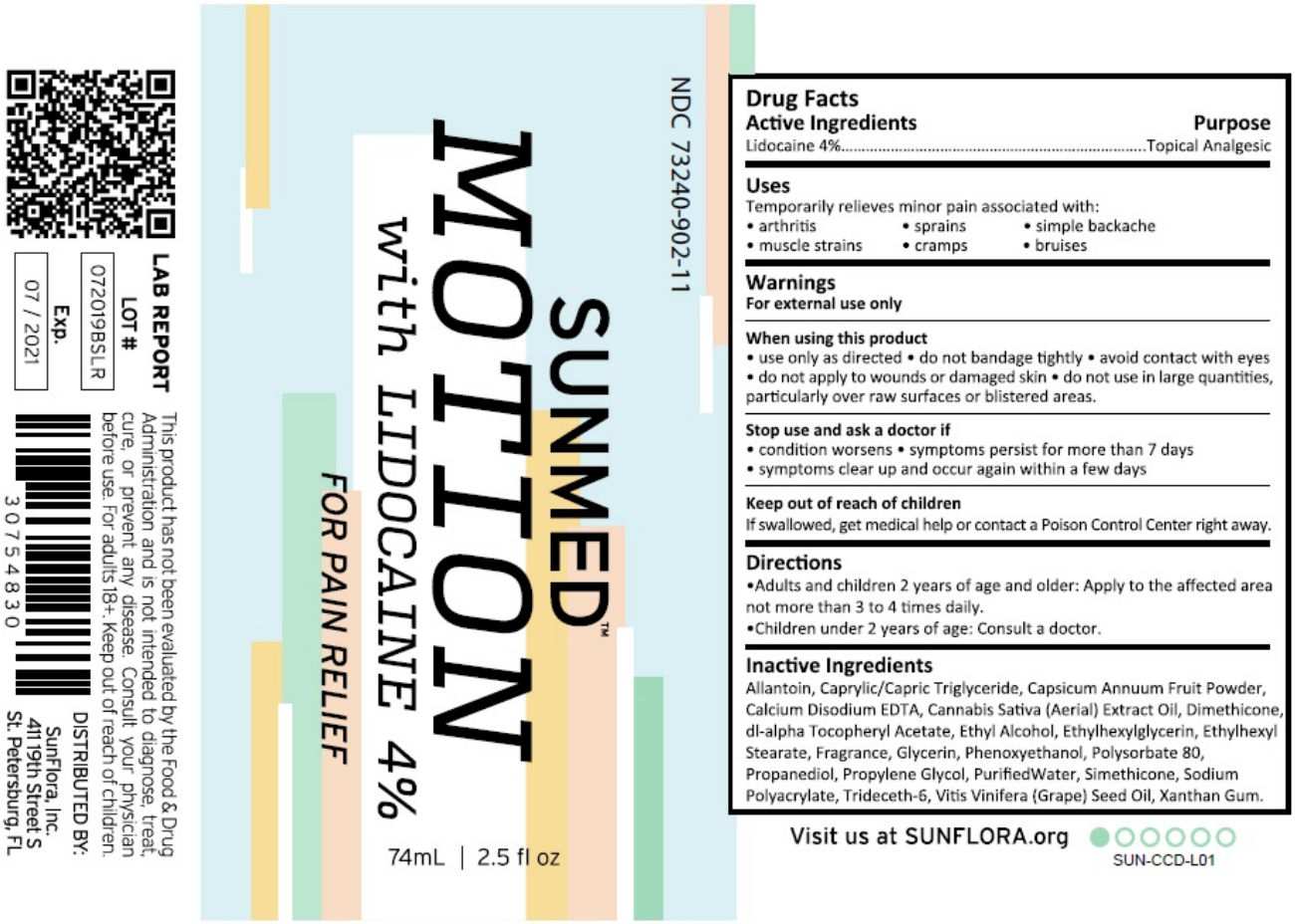
This product has not been evaluated by the Food & Drug Administration and is not intended to diagnose, treat, cure, or prevent any disease. Consult your physician before use. For adults 18+.

DISTRIBUTED BY:

SunFlora, Inc.
411 19th street S

St. Petersberg, FL

Packaging



FRONT PANEL

SUNMED MOTION
lidocaine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73240-902
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALLANTOIN (UNII: 344S277G0Z)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
PAPRIKA (UNII: X72Z47861V)	
EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955)	
CANNABIS SATIVA SUBSP. SATIVA FLOWERING TOP (UNII: 8X454SZ22D)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
ALCOHOL (UNII: 3K9958V90M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPANEDIOL (UNII: 5965N8W85T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
GRAPE SEED OIL (UNII: 930MLC8XGG)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73240-902-11	1 in 1 BOX	08/22/2019	
1		74 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 not final	part348	08/22/2019	

Labeler - Sunflora Inc (067153368)