

LIDOCAINE- 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.

73240-004-03

temporarily relieves minor pain associated with:
 arthritis, sprains, muscle strains, cramps, bruises,
 and simple headache.
 Directions: Apply directly to affected area.
 Learn more at
getsunmed.com



LAB REPORT
 LOT 050721BSLR
 EXP 01 / 2024
 Lab Reports at
getsunmed.com



SUNMED
 pain relief roll-on
 with LIDOCAINE

90mL | 3 fl oz



Drug Facts	
Active Ingredients Lidocaine 4%	Purpose Topical Anesthetic
Uses Temporarily relieves minor pain associated with: • arthritis • sprains • simple headache • 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 18 years of age and older. Apply to the affected area not more than 3 to 4 times daily.	
Inactive Ingredients Aqua, Glycerol Stearate, Polyglyceryl-6 Palmitate/ Succinate, Cetearyl Alcohol, Butylene Glycol, Caprylic/ Capric Triglyceride, Hydroxymethoxyphenyl Decanone, Behenyl Alcohol, Cannabis Sativa Seed Oil, Caprylyl Glycol, Hexylene Glycol, Dimethicone, Xanthan Gum, Arnica Montana Flower Extract, Aloe Barbadensis (Aloe Vera) Leaf Juice, Glycerin, Capsicum Annuum Fruit Extract, Phenoxethanol.	
Questions? Concerns? (888) 524-7437 or customersupport@getsunmed.com	

Contains Non-Detectable levels of Delta-9 THC. For adults 18+. Keep out of the reach of children. Not intended for ingestion - Do Not Eat. Do not take if pregnant or nursing. Consult physician before use. This product is not intended to diagnose, treat, cure or prevent any disease. This statement has not been evaluated by the Food and Drug Administration. Learn more at getsunmed.com.

DISTRIBUTED BY: SunFlora, Inc.
 600 8th Ave W Palmetto, FL 34221



LIDOCAINE

lidocaine liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73240-004
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 g

Inactive Ingredients

Ingredient Name	Strength
BUTYLENE GLYCOL (UNII: 3XUS85KORA)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
WATER (UNII: 059QF0KO0R)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
DIMETHICONE 1000 (UNII: MCU2324216)	
XANTHAN GUM (UNII: TTV12P4NEE)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
ARNICA MONTANA FLOWER WATER (UNII: U7L2JP51PR)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73240-004-03	85 g in 1 APPLICATOR; Type 0: Not a Combination Product	01/26/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	01/26/2021	

Labeler - SUNFLORA, INC (067153368)

Revised: 5/2022

SUNFLORA, INC