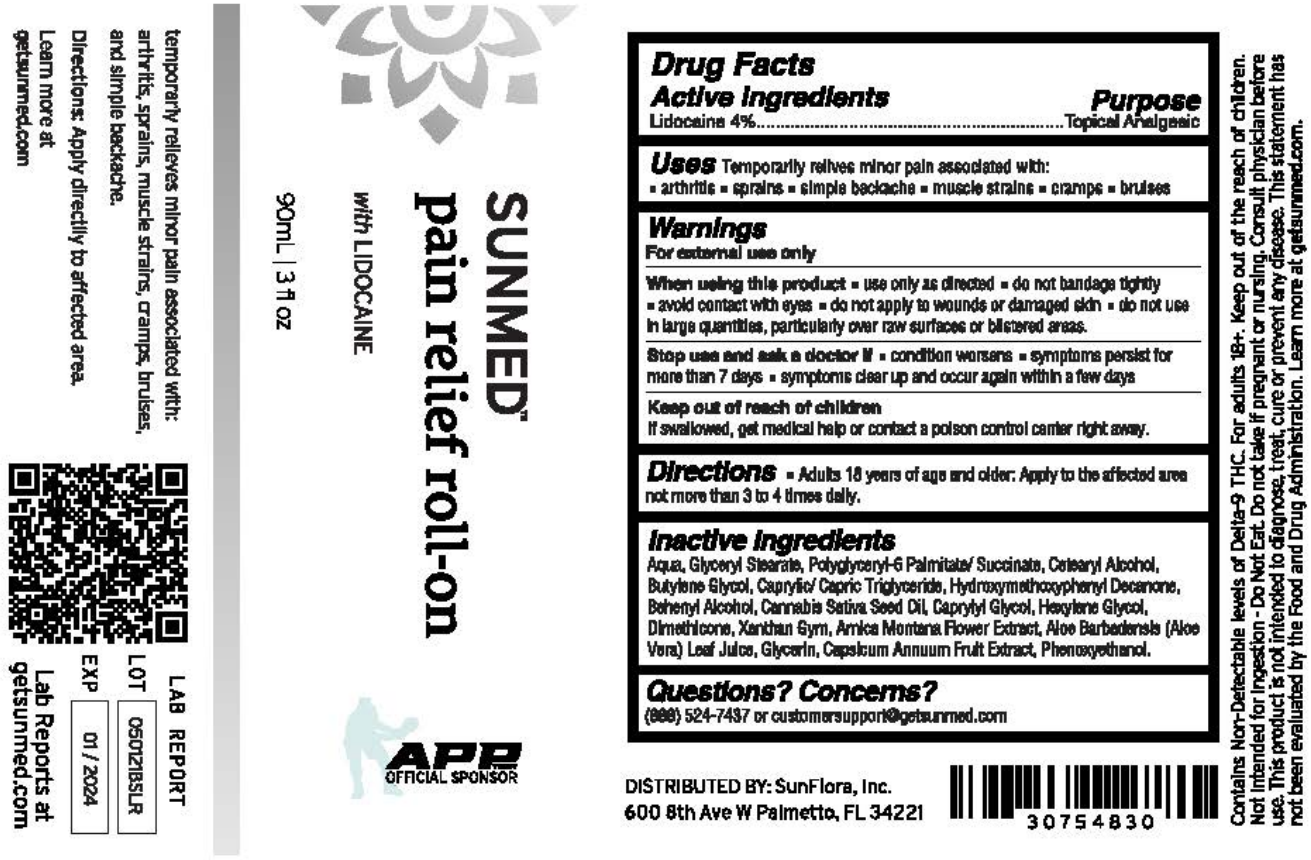
LIDOCAINE- lidocaine liquid
SUNFLORA, INC

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



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: 3XUS85K0RA)	
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