CANNABICOOL PAIN RELIEVING LIDOCAINE ROLL-ON- lidocaine hcl gel
Market America, 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.

Cannabicool Pain Relieving Lidocaine Roll-On

Active Ingredient

Lidocaine HCl 4%

Purpose

Topical Anesthetic

Uses

For the temporary relief of pain and itching.

For external use only.

Flammable

Keep away from fire or open flame.

Do not use

- In large quantities, particularly over raw surfaces or blistered areas.
- If you are allergic to any of the ingredients in this product.

When using this product

- Avoid contact with eyes.
- Do not apply to wounds or damaged skin.
- Do not bandage tightly.

Stop use and ask a doctor if

- Condition worsens.
- Symptoms persist for more than 7 days or clear up and occur again within a few days.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children

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

Directions

Adults and children over 12 years:

- Apply to affected area not more than 3 to 4 times daily. If medicine comes in contact

with hands, wash with soap and water.
- For children under 12 years of age: ask a doctor.

Other information

- Store tightly closed in a cool, dry place.

Inactive ingredients

Aloe Barbadensis (Aloe Vera) Gel, Beta Caryophyllene, Boswellia Carterii Resin Extract, Calendula Officinalis Flower Extract, Cannabis Sativa Seed Oil, Acrylates/c10-30 Alkyl Acrylate Crosspolymer, Glycerin, Isopropyl Alcohol, Isopropyl Myristate, Melissa Officinalis (Lemon Balm) Leaf Extract, Nigella Sativa (Black Cumin) Seed Oil, Silica, Tocopheryl Acetate, Triethanolamine, Water

CANNABICOOL PAIN RELIEVING LIDOCAINE ROLL-ON

lidocaine hcl gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76209-567
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDAMIDINE HYDROCHLORIDE	40 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARYOPHYLLENE (UNII: BHW853AU9H)	
FRANKINCENSE (UNII: R9XLF1R1WM)	

CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
CARBOMER INTERPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 132584PQMO)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
MELISSA OFFICINALIS LEAF (UNII: 50D2ZE9219)	
NIGELLA SATIVA SEED OIL (UNII: CS4U38E731)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76209-567-01	88 g in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	08/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	08/01/2019	

Labeler - Market America, Inc. (797412236)

Registrant - A.I.G. Technologies, Inc. (086365223)

Establishment

Name	Address	ID/FEI	Business Operations
A.I.G. Technologies, Inc.		086365223	manufacture(76209-567) , label(76209-567)

Revised: 1/2022 Market America, Inc.