CBD MIST TOPICAL ANESTHETIC- lidocaine hydrochloride liquid
Meneks, Michael dba Doctor Energy, 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.

CBD MIST Topical Anesthetic

Drug Facts

Active Ingredient

Lidocaine HCl 4.00%

Purpose

Topical Analgesic

Uses:

For temporary relief of pain.

Warnings:

• For external use only.

• Avoid contact with eyes.
• If symptoms persist for more than seven days, discontinue use and consult physician.

Keep out of reach of children.
- If swallowed, consult physician.

Do not use
- in large quantities, particularly over raw surfaces or blistered areas.

If pregnant or breast feeding,

contact physician prior to use.

Directions:

• Adults and children two-years of age or older: Apply to affected area not more than three to four times daily.
• Children under two-years of age: consult a physician.

Additional Information:

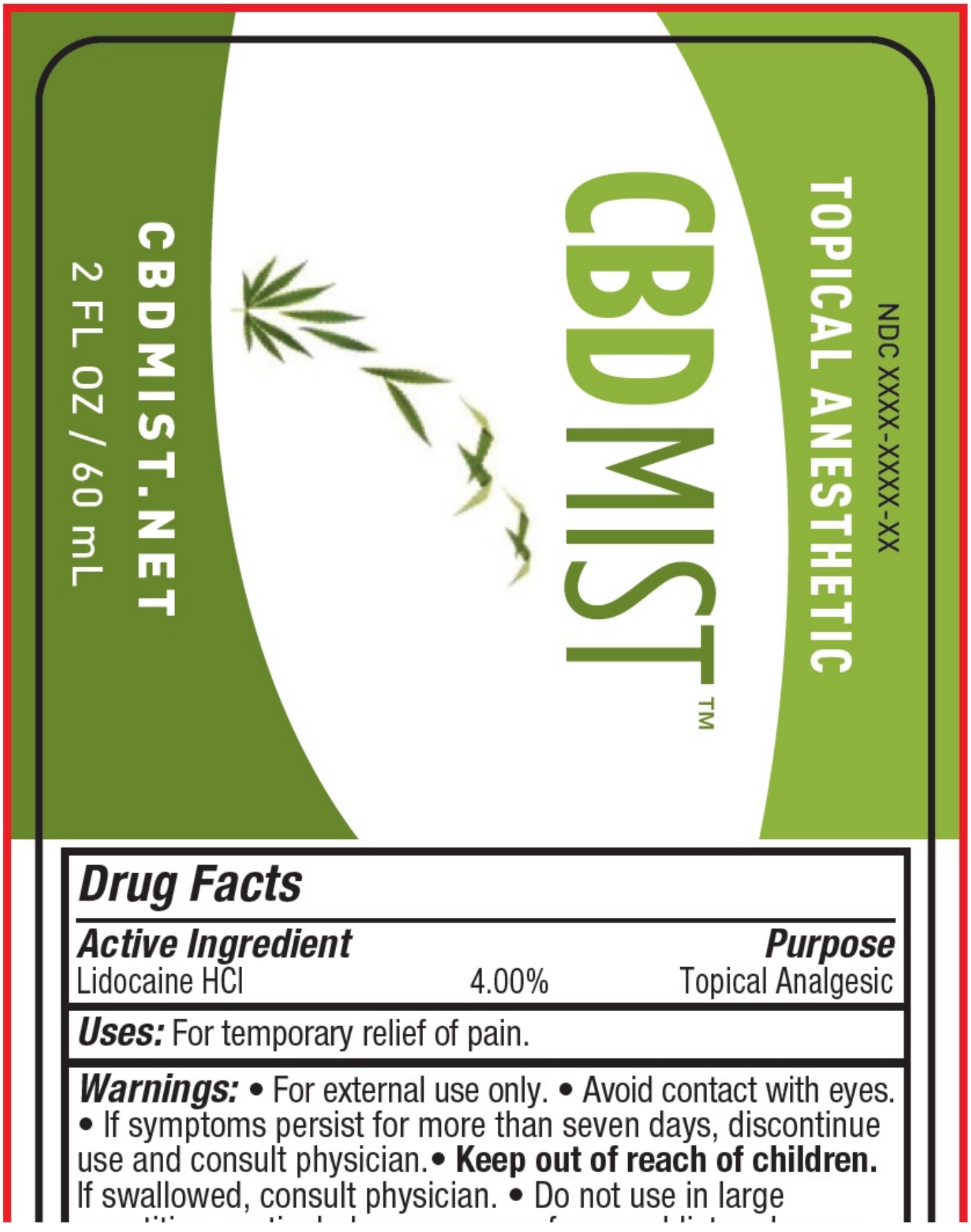
Store at room temperature.

Other Ingredients:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, Cannabis Sativa (Full Spectrum Hemp) Oil, Ethylhexylglycerin,

Isopropyl Alcohol, Methylsulfonylmethane (MSM), Phenoxyethanol, Polysorbate 20, Rosmarinus Officinalis (Rosemary) Oil, Simmondsia Chinensis (Jojoba) Oil.

Package Labeling:



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Distributed by: Doctor Energy, Inc.
North Miami Beach FL 33160 **www.doctorenergyinc.com**

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CBD MIST TOPICAL ANESTHETIC

lidocaine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73524-095
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
ROSEMARY (UNII: IJ67X351P9)	
JOJOBA OIL (UNII: 724GKU717M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73524-095-02	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	12/01/2019	

Labeler - Meneks, Michael dba Doctor Energy, Inc. (009963305)

Revised: 12/2019 Meneks, Michael dba Doctor Energy, Inc.