

**INSTA RELIEF DM- lidocaine hydrochloride, menthol cream
DM LAB CORP**

INSTA-RELIEF DM

DRUG FACTS

ACTIVE INGREDIENTS

Lidocaine HCL 4.00%

Menthol 1.00%

Topical Anesthetic

INDICATIONS:

For the temporary relief of pain and itching associated with minor burns sunburn, minor cuts, scrapes, insect bites or minor skin irritations.

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 apply

- to wounds or damaged skin.
- Do not bandage tightly.

If pregnant or breast feeding,

contact physician

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.

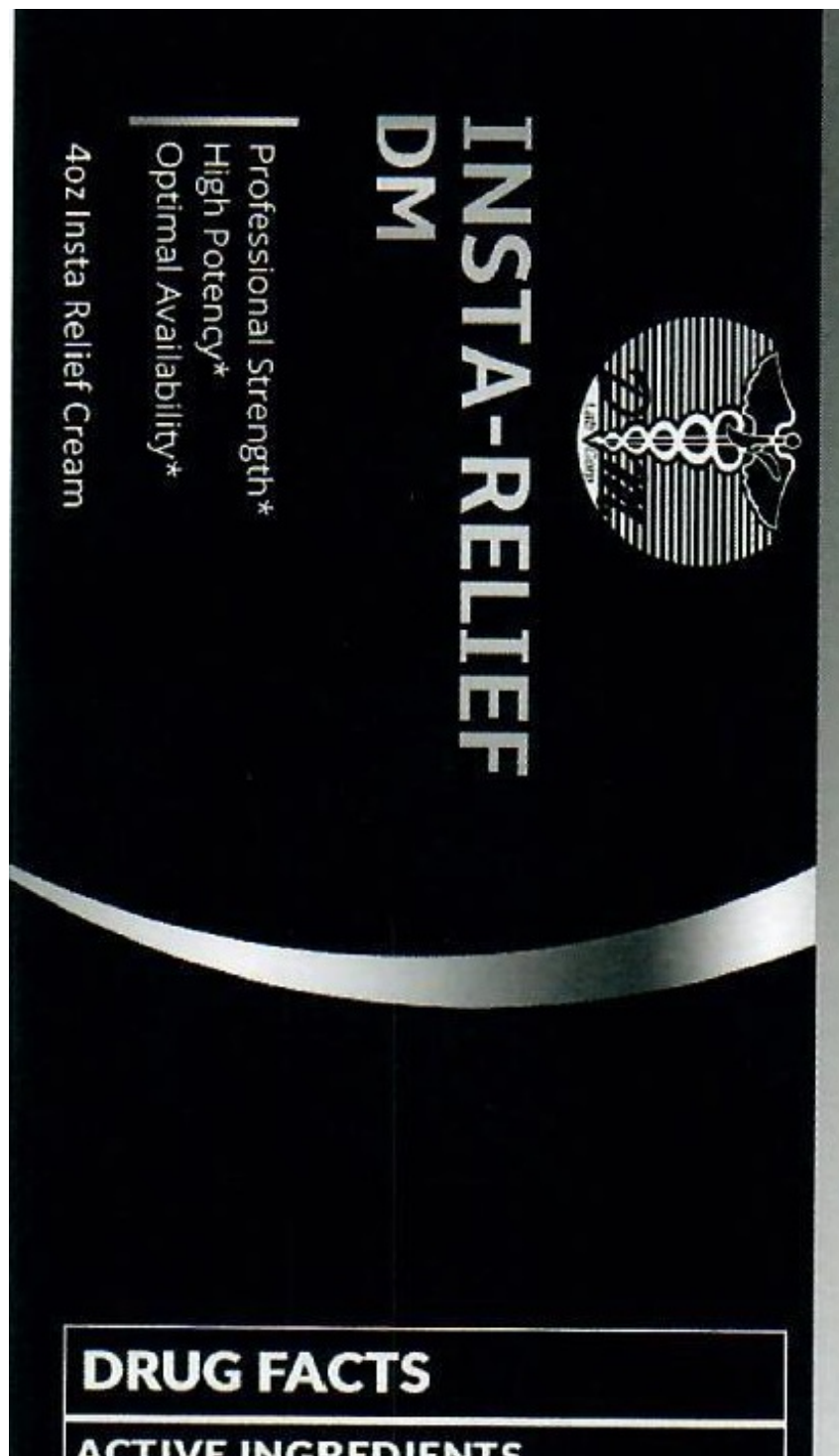
Additional Information:

Store at room temperature.

OTHER INGREDIENTS:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Benzyl Alcohol, Caprylic/Capric Triglyceride, Carbomer, Cetearyl Alcohol, Cetereth-20, Cetyl Alcohol, Citrus Aurantifolia (Orange) Oil, Daucus Carota Sativa (Carrot) Extract, Ethylhexylglycerin, Eucalyptus Globulus Oil, Glycerin, 1-Arginine, Lavandula Angustifolia (Lavender) Oil, Lecithin, Methylsulfonylmethane (MSM), Phenoxyethanol, Potassium Hydroxide

Package Labeling:



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Lidocaine HCL	4.00%	Topical Anesthetic
Menthol	1.00%	Topical Analgesic

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Manufactured for:

DM Lab Corp
425 Broadhollow Rd.
Suite 25
Melville, NY 11747
631 662-9200
web: dmlabcorp.com
email: info@dmlabcorp.com



INSTA RELIEF DM

lidocaine hydrochloride, menthol cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72465-600
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
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
LIME OIL (UNII: UZH29XGA8G)	
CARROT (UNII: L56Z1JK48B)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:72465-600-00	118 mL in 1 JAR; Type 0: Not a Combination Product	07/05/2018	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017		07/05/2018	

Labeler - DM LAB CORP (099396086)

Revised: 11/2023

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