# LIDOCAINE 4% TOPICAL ANESTHETIC CREAM- lidocaine 4% cream cream Mohnark Pharmaceuticals Inc.

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#### **Active Ingredients**

Lidocaine 4% w/w

#### **Purpose**

**Topical Anesthetic** 

Uses temporarily relieves pain and itching due to:

- minor cuts
- minor scrapes
  - sunburn
  - minor skin irritations
  - minor burns
  - insect bites

#### Warnings

For external use only

## When using this product

- do not use in or near the eyes
- do not use in large quantities, particularly over raw surfaces or blistered areas

#### Stop use and ask a doctor if

- allergic reaction occurs
- condition worsens or does not improve within 7 days
- symptoms clear up and return within a few days
- redness, irritation, swelling, pain or other symptoms begin or increase

## Keep out of reach of children

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

#### **Directions**

- Adults and children 2 years and older: apply externally to the affected area up to 3-4 times a day.
- Children under 2 years of age: consult a doctor

#### Other information

• Store at USP controlled room temperature 20-25 °C (68-77 °F).

#### **Inactive ingredients**

benzyl alcohol, cabomer 940, cholesterol, hydrogenated lecithin, polysorbate 80, propylene glycol, tocopheryl acetate, trolamine, water

## **Dosage**

40 milligrams/gram per application

#### Usage

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- Children under 2 years of age: consult a doctor

## Package Label



#### LIDOCAINE 4% TOPICAL ANESTHETIC CREAM

lidocaine 4% cream cream

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73715-001

Route of Administration TOPICAL

## **Active Ingredient/Active Moiety**

Ingredient Name	<b>Basis of Strength</b>	Strength

**LIDOCAINE** (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987) LIDOCAINE 4.27 g in 100 g

#### **Inactive Ingredients**

Ingredient Name	Strength

POLYSORBATE 80 (UNII: 6OZP39ZG8H) 2 g in 100 g

**WATER** (UNII: 059QF0KO0R) 80.48 g in 100 g

#### Packaging

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	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l		NDC:73715-001- 01	30 g in 1 TUBE; Type 0: Not a Combination Product	06/15/2020	

## **Marketing Information**

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	M017	06/15/2020	

## **Labeler - Mohnark Pharmaceuticals Inc.** (117013830)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Mohnark Pharmaceuticals Inc.		117013830	manufacture(73715-001)	

Revised: 1/2024 Mohnark Pharmaceuticals Inc.