# LIDOCAINE 4% PATCH- lidocaine 4% patch SOLA Pharmaceuticals

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Lidocaine 4% Patch

### **Active Ingredients**

Lidocaine

## **Purpose**

**Topical Anesthetic** 

#### Uses

For the temporary relief of pain

## Warnings

#### For external use only

#### Do not use

- more than 1 patch at a time
- on wounds or damaged skin
- with a heating pad
- if you are allergic to any ingredients of this product

## When using this product

- use only as directed
- avoid contact with the eyes, mucous membranes or rashes
- do not bandage tightly

## Stop use and ask a doctor if

- localized skin reactions occur, such as rash, itching, redness, irritation, pain, swelling and blistering
- conditions worsen
- symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days

**If pregnant or breast-feeding**, ask a health care 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 12 years of age and over:

- Clean and dry the affected area
- Open pouch and remove one patch
- Apply 1 patch at a time to affected area; not more than 3 to 4 times daily
- Reseal pouch containing unused patches after each use
- Remove patch from the skin after at most 8-hour application

Children under 12 years of age: consult a doctor

#### Other Information

- Avoid storing product in direct sunlight
- Protect product from excessive moisture

## Other Ingredients

Dihydroxyaluminum Aminoacetate, Disodium Ethylenediaminetetraacetic Acid, Glycerin, Kaolin, Methylparaben, Polysorbate 80, Propylene Glycol, Polyacrylic Acid, Polyvinylpyrrolidone K90, Propylparaben, Sodium Polyacrylate, Tartaric Acid, Titanium Dioxide, Water.

Ouestions or comments? 866-747-7365

Lidocaine 4% Patch

NDC 70512-812-30

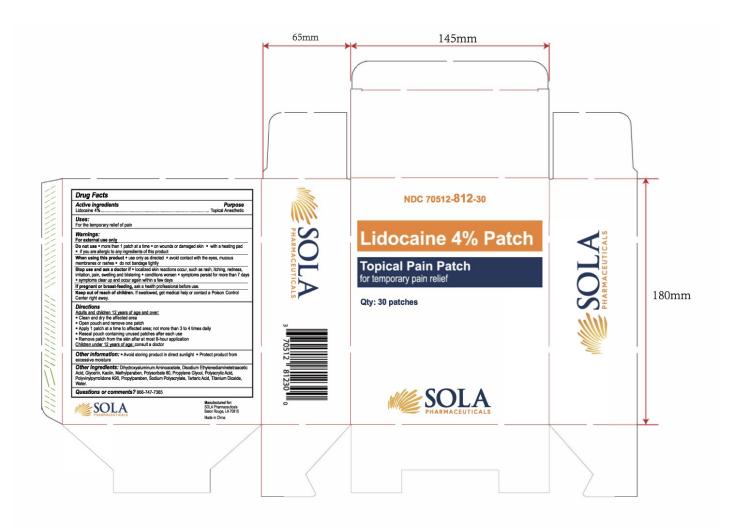
Qty: 30 patches

Manufactured for:

**SOLA Pharmaceuticals** 

Baton Rouge, LA 70810

Made in China



## **LIDOCAINE 4% PATCH**

lidocaine 4% patch

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:70512-812

Route of Administration TOPICAL

## **Active Ingredient/Active Moiety**

Ingredient Name
Basis of Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)
LIDOCAINE (UNII: 98PI200987) 40 mg in 1 g

#### **Inactive Ingredients**

| Ingredient Name                                      | Strength |
|--|----------|
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP)                  |          |
| GLYCERIN (UNII: PDC6A3C0OX)                          |          |
| DISODIUM ETHYLENEDIAMINEDIACETATE (UNII: EQL53S5L0F) |          |
| SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)  |          |
| PROPYLPARABEN (UNII: Z8IX2SC10H)                     |          |
| WATER (UNII: 0590F0KO0R)                             |          |

| KAOLIN (UNII: 24H4NWX5CO)                         |  |
|---|--|
| POLYSORBATE 80 (UNII: 60ZP39ZG8H)                 |  |
| POVIDONE K90 (UNII: RDH86HJV5Z)                   |  |
| DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6) |  |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3)               |  |
| POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)   |  |
| TARTARIC ACID (UNII: W4888I119H)                  |  |

| Product Characteristics |           |              |  |
|-------------------------|-----------|--------------|--|
| Color                   |           | Score        |  |
| Shape                   | RECTANGLE | Size         |  |
| Flavor                  |           | Imprint Code |  |
| Contains                |           |              |  |

| P | Packaging            |   |                         |                       |  |  |
|---|----------------------|---|-------------------------|-----------------------|--|--|
| # | Item Code            | Package Description                               | Marketing Start<br>Date | Marketing End<br>Date |  |  |
| 1 | NDC:70512-812-<br>30 | 30 in 1 BOX                                       | 10/12/2023              |                       |  |  |
| 1 |                      | 1 g in 1 POUCH; Type 0: Not a Combination Product |                         |                       |  |  |

| Marketing Information |   |                         |                       |
|-----------------------|---|-------------------------|-----------------------|
| Marketing<br>Category | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |
| OTC Monograph Drug    | 505G(a)(3)                                  | 10/12/2023              |                       |
|                       |   |                         |                       |

## Labeler - SOLA Pharmaceuticals (080121345)

Revised: 11/2023 SOLA Pharmaceuticals