# LIDOPRO PATCH (LIDOCAINE 4%, MENTHOL 1%)- lidocaine and menthol patch patch

Preferred Pharmaceuticals Inc.

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83881-402 LidoPro Patch (Lidocaine 4%, Menthol 1%)

#### **Active ingredient**

Lidocaine 4%

Menthol 1%

#### **Purpose**

External Anesthetic

External Analgesic

#### Uses

For the temporary relief of pain.

#### Warnings

For external use only.

#### Do not use

- on the face or rashes, on wounds or damaged skin
- in the eyes, mouth, or other mucous membranes
- on genitals
- with a heating pad
- right before or after heart surgery
- any patch from a pouch that has been opened for 7 or more days
- in large quantities, particularly over raw surfaces or blistered areas
- if tamper-evident seal is torn, broken, or missing
- more than 2 patches per day unless directed by a doctor
- children under 18 years of age

## Ask a doctor before use if you have

- allergies to topical products
- high blood pressure, heart disease, or kidney disease

## When using this product

- Avoid contact with eyes. If eye contact occurs, rinse thoroughly with water.
- The risk of heart attack or stroke may increase if you use more than directed or for

longer than directed.

#### Stop use and ask doctor if

condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

#### If pregnant of breast-feeding, ask a health professional before use

Do not use during last 3 months of pregnancy because it may cause problems in the unborn child or complications during delivery.

#### Keep out of reach of children.

If put in mouth, seek medical help or contact a Poison Control Center right away. Dispose if the used patches by folding sticky ends together.

#### **Directions**

Adults 18 years and older:

- clean and dry affected area
- open pouch and remove one patch
- remove protective film from patch
- apply one patch to the affected area of pain and leave in place for 8 to 12 hours

#### Other information

- Some individuals may not experience pain relief until several minutes or hours after applying the patch
- avoid storing product in direct sunlight
- protect product from excessive moisture
- store at 67-77°F (19-25°C)

### Inactive ingredients

Carboxymethylcellose sodium, Dihydroxyaluminum aminoacetate, Edetate sodium, Glycerin, Kaolin, Methyl parahydroxybenzoate, Partially neutralized Sodium polyacrylate, Polyacrylic acid, Polysorbate 80, Propyl parahydroxybenzoate, Propylene glycol, Sodium Polyacrylate, Tartaric acid, Titanium dioxide, water

### **Questions?**

Call toll-free (800) 224-2048

# LidoPro® Patch ( Pain Reliever)

Brand Name

In each patch: Lidocaine 4% ...Topical Anesthetic / Menthol 1% ... Topical Analgesic

Pkg Size: Exp Date: ##/##/### Lot#: Batch#:

Ins: Mfg: Clinic Pharma Prod#:

Prod#:

Warning
For external use only. Do not use on the face or realways on wounds or damaged ske not use on the face or realways mouth, or other mucous membranes, on genitals, with a heating pad, allergue to any NSAIDS, rapid before or after head surely of the padding the product contains an NSAID, which may cause stomach bleeding. Ask a doctor before use if you are allergie to you, you are taking a duretie, you have high bloodless to you, you are taking a duretie, you have high bloodless to you, you from the product of the





CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed.



Instrucciones Espanol:
Aplique
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LidoPro® Patch (Pain Reliever) Qty: Ins: Lot: Bat: Prod# (NDC):

Log

Chart

Billing

Patient

LidoPro® Patch (Pain Reliever) Qty: Ins: Lot: Bat: Prod# (NDC):

LidoPro® Patch (Pain Reliever) Qty: Insurance NDC: Lot: Bat:

LidoPro® Patch (Pain Reliever)
Qty: Ins:
Lot: Bat:
Prod# (NDC):

#### **LIDOPRO PATCH (LIDOCAINE 4%, MENTHOL 1%)**

lidocaine and menthol patch patch

**Product Information** 

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68788-8879(NDC:83881-402)

Route of Administration TRANS DERMAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	84 mg in 8400 mg		
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	336 mg in 8400 mg		

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
METHYLPARABEN (UNII: A218C7H19T)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
TARTARIC ACID (UNII: W4888I119H)	
WATER (UNII: 059QF0KO0R)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	
POLYACRYLIC ACID (8000 MW) (UNII: 73861X4K5F)	
PROPYLPARABEN (UNII: Z8IX2SC10H)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
KAOLIN (UNII: 24H4NWX5CO)	
EDETATE SODIUM (UNII: MP1J8420LU)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
PROPANEDIOL (UNII: 5965N8W85T)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)	

Packaging				
# Item C	Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:6878	38- 15 mg in 3 Product	1 POUCH; Type 0: Not a Combination	05/20/2025	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	05/20/2025		

# Labeler - Preferred Pharmaceuticals Inc. (791119022)

# Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-8879)	

Revised: 5/2025 Preferred Pharmaceuticals Inc.