

**LIDOPRO PATCH- lidocaine, menthol, and methyl salicylate patch**  
**Preferred Pharmaceuticals Inc.**

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**LidoPro Patch**

**Active Ingredient**

Lidocaine 4%

**Purpose**

Topical Anesthetic

**Active Ingredient**

Menthol .5%

**Purpose**

Topical Analgesic

**Active Ingredient**

Methyl Salicylate .1%

**Purpose**

Topical Counterirritant

**Uses**

Temporarily relieves minor pain

**Warnings**

**For external use only**

**Do not use**

- more than one patch on your body at a time or on cuts, irritated, or swollen skin
- on puncture wounds
- for more than one week without consulting a doctor.

**When using this product**

- Use only as directed. Read and follow directions and warning on this packaging.
- Do not apply to wounds or damaged, broken, or irritated skin
- Avoid contact with the eyes or mucous membranes
- Do not bandage tightly or apply local heat (such as heating pads) to area of use
- Do not use at the same time as other topical analgesics
- Dispose of used patch in manner that always keeps product away from children and pets. Used patches still contain the drug product that can produce serious adverse effects if a child or pet chews or ingests this patch.

**Stop use and ask a doctor if**

- conditions worsens
- redness is present
- irritation develops
- symptoms persist for more than 7 days or clear up and occur again within a few days
- you experience signs of injury, such as pain, swelling, or blistering where the product was applied

**If pregnant or breast feeding,**

ask health professional before use.

**Keep out of reach of children and pets.**

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

**DIRECTIONS**

**adults and children over 12 years:**

- clean and dry the affected area
- remove patch from plastic liner and place on the affected area
- use 1 patch for up to 12 hours
- place used patch on the liner when not in use
- re-use the patch up to 2 times.

**children 12 years or younger: ask a doctor**

**OTHER INGREDIENTS:**

Vegan Glycerol, Polyacrylate, Aqua. Polysorbate 80

**Principal Display Panel**

LidoPro patch

**Relabeled By: Preferred Pharmaceuticals Inc.**

**LidoPro® Patch**  
Brand Name

In each patch: Lidocaine 4% ...Topical Anesthetic / Methyl Salicylate .1%...Topical Counterirritant / Menthol .5%... Topical Analgesic

**Pkg Size:** Exp Date: #####  
Lot#: Batch#:  
Ins:  
Mfg: Clinic Pharma  
Prod#:

**Warning**  
Use: Temporarily relieves minor pain. For external use only. Do not use more than one patch on your body at a time or on cut, irritated, or swollen skin, on puncture wounds. For more than one week without consulting a doctor. Avoid contact with the eyes, or mucous membranes, with a heating pad. Avoid storing in direct sunlight, protect product from excessive moisture. Store at 67°-77°F (19°-25°C)



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

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



**Directions English**  
Apply externally \_\_\_\_\_ times a day.



GTIN  
#####  
SN #####  
EXP #####

**Instrucciones Espanol:**  
Aplice externamente \_\_\_\_\_ veces al dia.

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

LidoPro® Patch  
Qty:  
Insurance NDC:  
Lot: Bat:

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

Log  
Chart  
Billing  
Patient

NDC 68788-8594-3

LIDOPRO PATCH				
lidocaine, menthol, and methyl salicylate patch				
Product Information				
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68788-8594(NDC:83881-401)	
<b>Route of Administration</b>	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)		LIDOCAINE	4 mg in 100 mg	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)		MENTHOL, UNSPECIFIED FORM	0.5 mg in 100 mg	
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)		METHYL SALICYLATE	0.1 mg in 100 mg	
Inactive Ingredients				
Ingredient Name			Strength	
GLYCERIN (UNII: PDC6A3C0OX)				
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)				
POLYSORBATE 80 (UNII: 6OZP39ZG8H)				
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8594-3	15 in 1 BOX	02/26/2024	
1		2 in 1 POUCH		
		9500 mg in 1 PATCH; Type 0; Not a Combination		

<b>1</b>	8500 mg in 1 PATCH; Type O: Not a Combination Product		
<b>Marketing Information</b>			
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M017	02/26/2024	

**Labeler** - Preferred Pharmaceuticals Inc. (791119022)

**Registrant** - Preferred Pharmaceuticals Inc. (791119022)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-8594)

Revised: 5/2026

Preferred Pharmaceuticals Inc.