

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

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


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)



Directions English

Apply externally _____tim
es a day.



Instrucciones Espanol:

Aplique
externamente _____vece
s al dia.

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):

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				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8594(NDC:83881-401)
Route of Administration		TOPICAL		
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)			LIDOCAINE	4 mg in 100 mg
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)			MENTHOL	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		

1	8500 mg in 1 PATCH; Type 0: Not a Combination Product		
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/26/2024	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

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