

**LIDOPRO PAIN RELIEF- capsaicin and lidocaine ointment
Preferred Pharmaceuticals Inc.**

83881-001 LidoPro Transdermal Pain Relief & Applicator

Active ingredient

- Capsicum annum fruit extract (capsaicin)
- Lidocaine HCL
- Menthol
- Methyl Salicylate

Purpose

- counterirritant
- anesthetic
- analgesic

Uses

For the temporary relief of joint pain and muscle pain associated with:

- arthritis
- simple backache
- muscle sprains
- muscle strains

Warnings

For external use only.

Do not use

- on damaged, irritated or infected skin
- with a bandage or heating pad
- if you are allergic to any ingredients in this product

When using this product

avoid contact with the eyes and mucous membranes

Stop use and ask doctor if

- condition worsens
- excessive skin
- irritation develops
- symptoms persist for more than 7 days

- symptoms clear up and occur again within 3 days

If pregnant or breast-feeding

ask a health professional before use.

Keep out of reach of children

If ingested, seek medical help or contact a Poison Control Center immediately.

Flammable

Keep away from excessive heat or open flame.

Directions

Adults and children 12 years of age and older:

- clean and dry the affected area
- apply product directly to skin, up to 4 times daily
- wash hands immediately after use

Children under 12 years of age: Consult physician.

Other information

- Store in a cool, dry place with lid tightly closed
- if the tamper-evident foil seal is not intact, do not use

Inactive ingredient

allantoin, aloe barbadensis leaf juice, ammonium acryloyldimethyltaurate/vp copolymer, cetyl alcohol, chamomilla recutita matricaria flower extract, dimethicone, disodium EDTA, ethylhexylglycerin, glycerin, glyceryl stearate, inulin lauryl carbamate, PEG-100 stearate, phenoxyethanol, stearic acid, triethanolamine, water

Questions?

(800) 224-2048 or info@clinicpharma.com

Relabeled By: Preferred Pharmaceuticals Inc.

NDC 68788-8833-9

Label

LidoPro Topical Pain Relief Oint. & Applicator



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

LidoPro Topical Pain Relief Oint. & Applicator
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Log

Chart

Billing

Patient

Brand Name

Active Ingredients: Lidocaine 4% ...Topical Anesthetic / Menthol 10%...topical analgesic / Capsaicin 0.0325%...counterirritant / Methyl Salicylate 27.5%...counterirritant

Pkg Size: Exp Date: ###/###/####

Ins:
Mfg: Clinic Pharma
Prod#:

Warning

Store in a cool dry place. Keep away from excessive heat or open flame. For external use only. Do not use on damaged, irritated or infected skin; with bandage or heating pad. If condition worsens or symptoms persist for more than 7 days; excessive skin irritation occurs. Avoid contact with eyes, genitals, and other mucous membranes. Keep out of the reach of children. If pregnant or breast-feeding, ask a health professional before use. Package not child-resistant.



Directions English

Apply externally _____ times a day.

GTIN

SN #####
EXP #####



Instrucciones Espanol:

Aplique externamente _____ veces al dia.

LidoPro Topical Pain Relief Oint. & Applicator
Qty: Ins:
Lot: Bat:
Prod# (NDC):

LidoPro Topical Pain Relief Oint. & Applicator
Qty: Ins:
Insurance NDC:
Lot: Bat:

LidoPro Topical Pain Relief Oint. & Applicator
Qty: Ins:
Lot: Bat:
Prod# (NDC):

LIDOPRO PAIN RELIEF

capsaicin and lidocaine ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8833(NDC:83881-001)
Route of Administration	TRANSDERMAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL, (+)- (UNII: C6B1OE8P3W) (MENTHOL, (+)- - UNII:C6B1OE8P3W)	MENTHOL, (+)-	10 mg in 100 mg
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 mg in 100 mg
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	27.5 mg in 100 mg
CAPSAICIN (UNII: S07O44R1ZM) (CAPSAICIN - UNII:S07O44R1ZM)	CAPSAICIN	0.0325 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
MATRICARIA CHAMOMILLA WHOLE (UNII: G0R4UBI2ZZ)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y1O)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PEG-100 MONOSTEARATE (UNII: YD01N1999R)	
ALLANTOIN (UNII: 344S277G0Z)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

GLYCERIN (UNII: PDC6A3C0OX)	
INULIN LAURYL CARBAMATE (UNII: 48RFF58ESG)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8833-9	1 in 1 BOX	02/21/2025	
1		9900 mg in 1 TUBE, WTH APPLICATOR; 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	02/21/2025	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-8833)

Revised: 2/2025

Preferred Pharmaceuticals Inc.