

MAXIMUM STRENGTH PAIN RELIEF LIDOCAINE PATCH- lidocaine patch
Rite Aid Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Maximum Strength Pain Relief Lidocaine Patch

Active Ingredients

Lidocaine 4% Topical Anesthetic

Warnings

For external use only.

Do not Use

- more than one patch on your body at a time
- on cut, irritated or swollen skin
- on puncture wounds
- for more than one week without consulting a doctor
- if you are allergic to any active or inactive ingredients
- if pouch is damaged or opened.

When using this product

- use only as directed
- read and follow all directions and warnings on this carton
- do not allow contact with the eyes
- do not use at the same time as other topical analgesics
- do not bandage tightly or apply local heat (such as heating pads) to the area of use
- do not microwave
- 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 consult a doctor

- condition 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 skin injury, such as pain, swelling or blistering where the product was applied.

Keep out of reach of children and pets.

If swallowed, get medical help or contact a Poison Control Center 800-222-1222 right away.

Dosage and Administration

Directions Adults and children 12 years of age and over :

- Clean and dry affected area
- Carefully remove backing from patch starting at a corner.
- Apply sticky side of patch to affected area.
- use one patch for up to 12 hours.
- Discard patch after single use.
- Children under 12 years of age: consult a physician.

If pregnant or breastfeeding, ask a health professional before use.

Other Safety Information

Store in a clean, dry place outside of direct sunlight. Protect from excessive moisture.

Inactive Ingredients

Aluminum Glycinate, carboxymethylcellulose sodium, Glycerin, iodopropynyl butylcarbamate, Kaolin, petrolatum, phenoxyethanol, polyacrylic acid, Polysorbate 80, Povidone, Propylene Glycol, Sodium Polyacrylate, Tartaric Acid, Titanium Dioxide, Water, 3-(2-ethylhexyloxy)propane-1,2-diol

Uses

Temporarily relieves minor pain.

Uses

Temporarily relieves minor pain.

Drug Facts

Active Ingredient
Lidocaine 4.25% **Purpose**
Topical anesthetic

Uses Temporarily relieves minor pain.

Warnings
For external use only.
Do not use on more than one patch on your body at a time or on cuts, irritated or broken skin or on puncture wounds.
Do not use for more than one week without consulting a doctor. If you are allergic to any active or inactive ingredients, this patch is contraindicated or causes.

Directions
When using this product: Use only as directed. Read and follow all directions and warnings on this carton.
Do not use on irritated skin. Do not use if the nerve line on other labeled analgesics. Do not massage, rub, or apply heat (such as heating pads) to the area of use. Do not microwave in Ziploc® or other plastic in microwave that always keeps product away from children and pets. Lidocaine patches still contain the drug product that can produce serious adverse effects if a child or pet swallows or ingests the patch.
Stop use and consult a doctor if: Condition worsens or Redness is present or Infection develops.
Do not use for more than 7 days or longer use could result within a few days. You experience signs of skin injury, such as pain, swelling or blistering where the product was applied.
If pregnant or breastfeeding, ask a health professional before use.
Keep out of reach of children and pets. If swallowed, get medical help or contact a Poison Control Center 800-222-1222 right away.

Directions
Adults and children 12 years of age and over: Clean and dry affected area. Carefully remove backing from patch starting at a corner. Apply sticky side of patch to affected area. Use one patch for up to 12 hours. Discard patch after single use. Children under 12 years of age: consult a physician.

Other Information
Store in a clean, dry place out of direct sunlight. Protect from excessive moisture.

Inactive Ingredients
sulfonamide plasticizer, methacrylate/cellulose acetate, glycerin, hydroxypropyl butylcarbamate, lauryl, stearic acid, polyacrylate, polyethylene wax, polyethylene glycol, polyethylene glycol, sodium polyacrylate, butadiene, methyl methacrylate, water, 1,2-ethanediol/dipropylene-1,2-diol

Salonpas® is a registered trademark of Heinen Pharmaceuticals Co., Inc. Heinen Pharmaceuticals is not affiliated with Riba Ad® or this product.

1-866-326-1313
Monday - Friday, 8:30 a.m. - 4:30 p.m. CST

MAXIMUM STRENGTH PAIN RELIEF LIDOCAINE PATCH

NDC 11822-6541-0

Compare to the active ingredient of Salonpas® Maximum Strength Lidocaine Patch*

LASTS UP TO 12 HOURS

MAXIMUM STRENGTH PAIN RELIEF LIDOCAINE PATCH

4% LIDOCAINE / TOPICAL ANESTHETIC

Desensitizes aggravated nerves & relieves pain
Medicated for targeted pain relief
Stay-put, flexible patch
No-mess, easy to apply and remove
Odor free

6 PATCHES
3.93 IN x 5.51 IN (10 cm x 14 cm)

MAXIMUM STRENGTH PAIN RELIEF LIDOCAINE PATCH

Child-Resistant Packaging
Model 64-01 1PF
02-0409 01201
V0810048

DISTRIBUTED BY:
SITE RD, 30 HUNTER LANE,
CRAP - HILL, PA 17011
www.zeneca.com
MADE IN CHINA

SATISFACTION GUARANTEE
If you're not satisfied, we'll happily refund your money.

LOT
EXP

MAXIMUM STRENGTH PAIN RELIEF LIDOCAINE PATCH

lidocaine patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-6541
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
POLYACRYLIC ACID (8000 MW) (UNII: 73861X4K5F)	
PETROLATUM (UNII: 4T6H12BN9U)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	

GLYCERIN (UNII: PDC6A3C0OX)	
KAOLIN (UNII: 24H4NWX5CO)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
TARTARIC ACID (UNII: W4888I119H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
WATER (UNII: 059QF0KO0R)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Product Characteristics

Color		Score	
Shape	RECTANGLE	Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-6541-0	6 in 1 CARTON	03/01/2021	
1		1 g in 1 PATCH; Type 0: Not a Combination Product		
2	NDC:11822-6541-1	5 in 1 CARTON	03/01/2021	
2		1 g 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 not final	part348	03/01/2021	

Labeler - Rite Aid Corporation (014578892)

Establishment

Name	Address	ID/FEI	Business Operations
Foshan Aqua Gel Biotech		529128763	manufacture(11822-6541)