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

Maximum Strenght 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.

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MAXIMUM STRENGTH PAIN RELIEF LIDOCAINE PATCH

Pull to separate film













Drug Facts Active ingredient

*Asparcrame® is a registered tradement of Chattern, Inc. Chattern, Inc. is not still leted with Rhis Ald® or this product.

Toll-Free Customer Care Help Line 1-966-326-1313 Monday — Friday, 6:30 s.m. — 4:30 p.m. CST



NDC 11822-6541-1

LASTS UP TO

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



MAXIMUM STRENGTH

PAIN RELIEF

5 PATCHES 183 W x 551 W (10 cm x 14 cm)

Child Resistent Packaging Model 64 911677 82-84011 01/21 VG510048

DISTRIBUTED BY: RITE AD, 30 HUNTER LANE, CAMP HILL, PA 17011 INTRIBUTED BLOOM MADE IN CHINA

SATISFACTION GUARANTEE Pyet're not establed, me'll happly refund year manage

LOT EXP

LIDOCAINE PATCH MAXIMUM STRENGTH



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 Molety				
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: K6790BS311)			
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: W4888119H)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

WATER (UNII: 059QF0KOOR)

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 Drug	M017	03/01/2021	

Labeler - Rite Aid Corporation (014578892)

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

Revised: 12/2024 Rite Aid Corporation