UP AND UP LIDOCAINE PAIN RELIEF- lidocaine patch
Target 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.

Target Corporation Lidocaine Pain-Relief Patches Drug Facts

Active ingredient

Lidocaine 4%

Purpose

Topical anesthetic

Uses

For temporary relief of pain

Warnings

For external use only

Do not use

- more than 1 patch at a time
- on wounds or damaged skin
- with a heating pad
- if you have ever had an allergic reaction to this product or any of its ingredients

When using this product

- use only as directed
- avoid contact with the eyes, mucous membranes or rashes
- do not bandage tightly

Stop use and ask a doctor if

- localized skin reactions occur, such as rash, itching, redness, irritation, pain, swelling and blistering
- conditions worsen
- symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

Adults and children 12 years of age and over:

- clean and dry affected area
- remove film from patch and apply to the skin (see illustration)
- apply 1 patch at a time to affected area, not more than 3 to 4 times daily
- remove patch from the skin after at most 8 hours of application

Children under 12 years of age: consult a doctor

Other information

- avoid storing product in direct sunlight
- protect product from excessive moisture
- store at 20-25°C (68-77°F)

Inactive ingredients

carboxymethylcellulose sodium, dihydroxyaluminum aminoacetate, edetate disodium, glycerin, kaolin, methylparaben, polyacrylic acid, polyvinyl alcohol, propylene glycol, propylparaben, purified water, sodium polyacrylate, sodium polyacrylate starch, sorbitol solution, tartaric acid, urea

Questions?

Call 1-888-547-7400

Principal Display Panel

Compare to active ingredient in Salonpas® Lidocaine Patch

up to 8 hours

maximum strength

lidocaine pain-relief patches

4% lidocaine/topical anesthetic

desensitizes aggravated nerves in back, neck, shoulders, knees and elbows

for temporary relief of pain

numbing

unscented

6 PATCHES 3 15⁄16 IN x 5 ½ IN (10 cm x 14 cm) EACH

Do not use if printed pouch is torn or punctured.

Compare to active ingredient in
Salonpas® Lidocaine Patch*

NDC 11673-744-91

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4% lidocaine/topical anesthetic

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up&up™

6 PATCHES
3 15/16 IN x 5 ½ IN (10 cm x 14 cm) EACH

up&up™

7U291 UW C1

HOW TO APPLY

1. 

2. 

3. 

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100% satisfaction guaranteed or your money back.

Distributed by Target Corporation
Minneapolis, MN 55403
Made in Taiwan

*This product is not manufactured or distributed by Hisamitsu America, Inc., distributor of Salonpas® Lidocaine Patch.

PAPER BOX | MULTI-LAYER POUCH how2recycle.info

up&up™

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UP AND UP LIDOCAINE PAIN RELIEF

lidocaine patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-744
Route of Administration	TOPICAL, PERCUTANEOUS, TRANSDERMAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	560 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
KAOLIN (UNII: 24H4NWX5CO)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
TARTARIC ACID (UNII: W4888I119H)	
UREA (UNII: 8W8T17847W)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-744-91	6 in 1 CARTON	07/15/2019	
1		1 in 1 POUCH; 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	07/15/2019	

Labeler - Target Corporation (006961700)

Revised: 7/2019 Target Corporation