LIDOCAINE PATCH- lidocaine patch Foshan Aqua Gel Biotech Co., Ltd.,

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 Lidocaine Patch- bulk label

undefined

Lidocaine 4%

Purpose

Topical Anesthetic

Uses

Temporarily relieves minor pain

Warnings

For external use only

Do Not Use

- more than 1 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

When Using This Product

- use only as directed. Read and follow all directions and warnings on this label.
- do not allow contact with the eyes
- do not bandage tightly or apply local heat (such as heating pads) to the 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 or 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

- 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

If Pregnant or Breast Feeding

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 right away.

Directions

Adults and children over 12 years:

- clean and dry affected area
- remove backing from patch by firmly grasping both ends and gently pulling until backing separates in middle
- carefully remove smaller portion of backing from patch and apply exposed portion of patch to affected area
- once exposed portion of patch is positioned, carefully remove remaining backing to completely apply patch to affected area
- use 1 patch for up to 12 hours

children 12 years or younger: ask a doctor

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Inactive Ingredients

Dihydroxyaluminum aminoacetate, Glycerol, Kaolin, Methylparaben, Polyacrylic acid, Propylene glycol, Propylparaben, PVP, Sodium polyacrylate, Tartaric acid, Titanium dioxide, Tween 80, Water

Outer Label

ODOURLESS LIDOCAINE PAIN RELIEVING PATCH

Active Ingredients:

Batch No. : Mfg. Date : Exp. Date :	Quantity :300 Patches Shipper No. :of Gross Wight : Net Weight :		
STORE IN A COOL DRY PLACE. PROTECT FROM DIRECT SUNLIGHT / MOISTURE / FREEZING.	KEEP OUT OF REACH OF CHILDREN AND PETS. FOR EXTERNAL USE ONLY.		
NDC No.: 69159-003-01	CONFORMANCE WITH THE F.D. & CACT		
Manufactured By:	AND REGULATIONS THEREUNDER		
Foshan Aqua Gel Biotech Co. Ltd	Velocity Labeler Code: 76168		
MADE IN CHINA			
Shipped To: VELOCITY PHARMA LLC 226/B SHERWOOD AVE, FARMINGDALE, NY-11735	0 11822 68614 3		

LIDOCAINE PATCH

lidocaine patch

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Product Information						
Product T ype	HUMAN OTC DRUG	Item Code (Source) NDO		NDC:69	C:69159-003	
Route of Administration	TOPICAL					
Active Ingredient/Active I	Mojety					
Ingredient Name			Basis of Strength		Strength	
LIDO CAINE (UNII: 98 PI200987) (LIDOCAINE - UNII:98 PI200987)		LIDOCAINE		40 mg		
Institute Ingradiants						
Inactive Ingredients					- · ·	
	Ingredient Name				Strength	
SODIUM POLYACRYLATE (800	0 MW) (UNII: 285CYO341L)					
KAOLIN (UNII: 24H4NWX5CO)						
METHYLPARABEN (UNII: A218C7	7HI9T)					
PROPYLPARABEN (UNII: Z8IX2S	С1ОН)					
POVIDONE (UNII: FZ989GH94E)						
DIHYDRO XYALUMINUM AMINO	ACETATE (UNII: DO250 MG0 W6)					

GLYCEROL (1-(12-HY	DROXYSTEARATE)) (UNII: X84XWP4TOC)					
POLYACRYLIC ACID	(UNII: 9G2MAD7J6W) (UNII: 9G2MAD7J6W)					
TARTARIC ACID (UNII	: W4888I119H)					
POLYSORBATE 80 (U	NII: 6OZP39ZG8H)					
WATER (UNII: 059QF0)	KO0R)					
PROPYLENE GLYCOL	. (UNII: 6DC9Q167V3)					
Packaging						
# Item Code	Package Description	N	Iarketing Start Date	Marketing End Date		
1 NDC:69159-003-01	300 in 1 CASE	09	/11/2017			
1	1 in 1 PATCH; Type 0: Not a Combination Product					
Marketing Information						
Marketing Category	y Application Number or Monograph Cita	tion	Marketing Start Date	Marketing End Date		
OTC monograph not fina	al part348		09/11/2017			

Labeler - Foshan Aqua Gel Biotech Co., Ltd., (529128763)

Revised: 10/2017

Foshan Aqua Gel Biotech Co., Ltd.,