ALBERTSON LIDOCAINE, MENTHOL PAIN RELIEF MEDICATED PATCH- lidocaine and menthol patch Safeway

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.

Albertson Signature Care Lidocaine + Menthol Pain Relief Medicated Patch

Active ingredients

Lidocaine 4%

Menthol 1%

Purposes

Topical anesthetic

Topical analgesic

Use

for the temporary relief minor pain

Warnings

For external use only

Do not use

- more than 1 patch on your body at a time
- on a 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 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 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 800-222-1222 right away.

Directions

adults and children 12 years of age and older:

- clean and dry affected area
- Carefully remove backing from patch starting at a corner
- Apply sticky side of patch to affected area
- use 1 patch for up to 12 hours
- Discard patch after single use

children under 12 years of age: consult a physician

Inactive ingredients

carboxymethylcellulose sodium, dihydroxyaluminum aminoacetate, glycerin, iodopropynyl butylcarbamate, kaolin, petrolatum, phenoxyethanol, polyacrylic acid, polysorbate 80, povidone, propylene glycol, sodium polyacrylate, tartaric acid, titanium dioxide, water, 3-(2ethylhexyloxy)propane-1,2-diol.

PRINCIPAL DISPLAY PANEL



LOT EXP

ALBERTSON LIDOCAINE, MENTHOL PAIN RELIEF MEDICATED PATCH

lidocaine and menthol patch

PALATE RIM SING OF PALA OF OR SING OF OR S

Drug Facts

DISTRIBUTED BY BETTER LINNING BRANDS LLC, P.O. BICK 80, PLEYSANTON, CA 94596-0008 1-456-775-9079 VANN BA

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-961
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	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g	

Inactive Ingredients	
Ingredient Name	Strength
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	
KAOLIN (UNII: 24H4NWX5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
POLYACRYLIC ACID (8000 MW) (UNII: 73861X4K5F)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PETROLATUM (UNII: 4T6H12BN9U)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
TARTARIC ACID (UNII: W4888I119H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
WATER (UNII: 059QF0KO0R)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:21130-961- 05	5 g in 1 CARTON; Type 0: Not a Combination Product	04/01/2022	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	04/01/2022		

Labeler - Safeway (009137209)

Establishment				
Name	Address	ID/FEI	Business Operations	
Foshan Aqua Gel Biotech Co., Ltd.,		529128763	manufacture(21130-961)	

Revised: 4/2022 Safeway