

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

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**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**



<b>POLYACRYLIC ACID (8000 MW)</b> (UNII: 73861X4K5F)
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)
<b>SODIUM POLYACRYLATE (8000 MW)</b> (UNII: 285CYO341L)
<b>TARTARIC ACID</b> (UNII: W4888I119H)
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)
<b>WATER</b> (UNII: 059QF0KO0R)
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	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 Drug	M017	04/01/2022	

### Labeler - Safeway (009137209)

### Establishment

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

Revised: 12/2024

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