ZIMS MAX FREEZE- lidocaine hydrochloride and menthol, patch
KOBAYASHI Healthcare International, Inc.

ZIM'S™ MAX FREEZE

Drug Facts

Active ingredients	***Purpose***
Lidocaine HCl 4%	Topical anesthetic
Menthol 1%	Topical analgesic

Uses

For the temporary relief of pain

Warnings

For external use only

Do not use

- more than 1 patch on your body at a time
- if you are allergic to any ingredients in this product
- on open wounds, damaged or irritated skin
- with a heating pad or TENS device or other topical analgesics
- after expiration date

When using this product

- do not bandage tightly
- do not use otherwise than as directed
- avoid contact with eyes or mucous membranes
- may cause a burning sensation wherever it is applied, but is usually mild and gradually lessens over time
- if severe burning sensation occurs, discontinue use immediately
- may leave a blue stain on skin, which is safe and can be removed by washing with soap and water
- avoid contact with silk
- dispose of used patch in manner that always keeps product away from children and pets. Used patches still contain the active ingredients that can produce serious adverse effects if a child or pet chews or ingests this patch

Stop use and ask a doctor if

- burning sensation is painful and persistent
- condition worsens
- rash, itching, redness, swelling, blistering or excessive irritation of the skin develops
- 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 and pets. If swallowed, get medical help or contact a Poison Control Center right away.

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

Children under 12 years of age:

- consult a doctor

Other information

- store in a cool place away from direct sunlight

Inactive ingredients

FD&C Blue No. 1, Glycerin, Methylparaben, Polyacrylic Acid, Polysorbate 80, Polyvinyl Alcohol, Povidone, Propylene Glycol, Propylparaben, Sodium Polyacrylate, Sodium Polyacrylate Starch, Sorbitan Oleate, Sorbitol, Tartaric Acid, Vanillyl Butyl Ether, Water

Distributed by:
KOBAYASHI Consumer Products, LLC
P.O. Box 1191, Dalton, GA 30722

PRINCIPAL DISPLAY PANEL - 3 Patch Pouch Box

ZIM'S™
MAX-
FREEZE

LIDOCAINE 4%
PAIN RELIEF PATCH

FREEZING SENSATION

NUMBS
AWAY PAIN

Fast Acting & Long Lasting
Hydrated Gel Patch

3 PATCHES

3 15/16" X 5 1/2"
(10 cm X 14 cm)
Individually Packaged

LIDOCAINE 4% & MENTHOL 1%

ZIM'S™ MAX FREEZE

ZIM'S™ MAX FREEZE

LIDOCAINE 4%
PAIN RELIEF PATCH

- FREEZING SENSATION -

DESENSITIZES AGGRAVATED NERVES

On Back

On Neck

On Shoulder

placement only
7 81485 25132 2

ZIM'S MAX FREEZE

LIDOCAINE 4%
PAIN RELIEF PATCH
FREEZING SENSATION

3 PATCHES
3 15/16" X 5 1/2" (10 cm X 14 cm)
Individually Packaged

Features

1. Our unique patch formula infuses 4% Lidocaine into a hydro gel to relieve pain quickly.
2. Provides targeted pain relief that desensitizes aggravated nerves.
3. Easy to use patch with a No Mess application.

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KOBAYASHI

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P.O. Box 1191, Dalton, GA 30722

Consumer Comments Call: 1-800-432-8629
Share your feedback at: **feedback.ZimsUSA.com**
for a chance to win an assortment of Kobayashi products.
Made in Taiwan **www.ZimsUSA.com**

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P251320

EASY-TO-USE

K / O for
Lot Printing

ZIMS MAX FREEZE

lidocaine hydrochloride and menthol, patch

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54273-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	760 mg
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	190 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM POLYACRYLATE STARCH (400 MICROMETER PARTICLE) (UNII: Y58AEZ2NBY)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYACRYLIC ACID (300000 MW) (UNII: A8371R0U5J)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)	
SORBITOL (UNII: 506T60A25R)	
TARTARIC ACID (UNII: W4888I119H)	
VANILLYL BUTYL ETHER (UNII: S2ULN37C9R)	
WATER (UNII: 059QF0KO0R)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54273-012-01	3 in 1 BOX	05/10/2023	
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 drug	M017	05/10/2023	

Labeler - KOBAYASHI Healthcare International, Inc. (156391729)

Revised: 11/2023 KOBAYASHI Healthcare International, Inc.