ICY HOT- lidocaine, menthol patch
Lead Chemical Co. Ltd.

Icy Hot Max Pain Relief Patch

Active ingredients

Lidocaine 4%
Menthol 1%

Purposes

Topical anesthetic
Topical analgesic

Use

for the temporary relief of pain

Warnings

For external use only

Do not use

- on puncture wounds, cuts, irritated, damaged or swollen skin
- more than 1 patch on your body at a time or with other topical analgesics at the same time
- with a heating pad or apply local heat to the area of use

When using this product

- use only as directed
- do not bandage tightly
- avoid contact with the eyes and mucous membranes
- rare cases of serious burns have been reported with products of this type
- a transient burning sensation may occur upon application but generally disappears in several days
- 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 or symptoms persist for more than 7 days

- symptoms clear up and occur again within a few days
- severe burning sensation, redness, or irritation develops
- 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 12 years of age and older:

- 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 at a time and not more than 3 to 4 times daily

children under 12 years of age: consult a doctor

Inactive ingredients

aluminum glycinate, aluminum hydroxide, cellulose gum, glycerin, methyl acrylate/2-ethylhexyl acrylate copolymer, methylparaben, nonoxynol-30, polyacrylic acid, polysorbate 80, propylene glycol, silica, sodium polyacrylate, tartaric acid, titanium dioxide, urea, water

Package/Label Principal Display Panel

LIDOCAINE 4% + MENTHOL 1%

ICY HOT®
MAX
Pain Relief Patch

LIDOCAINE
PATCH

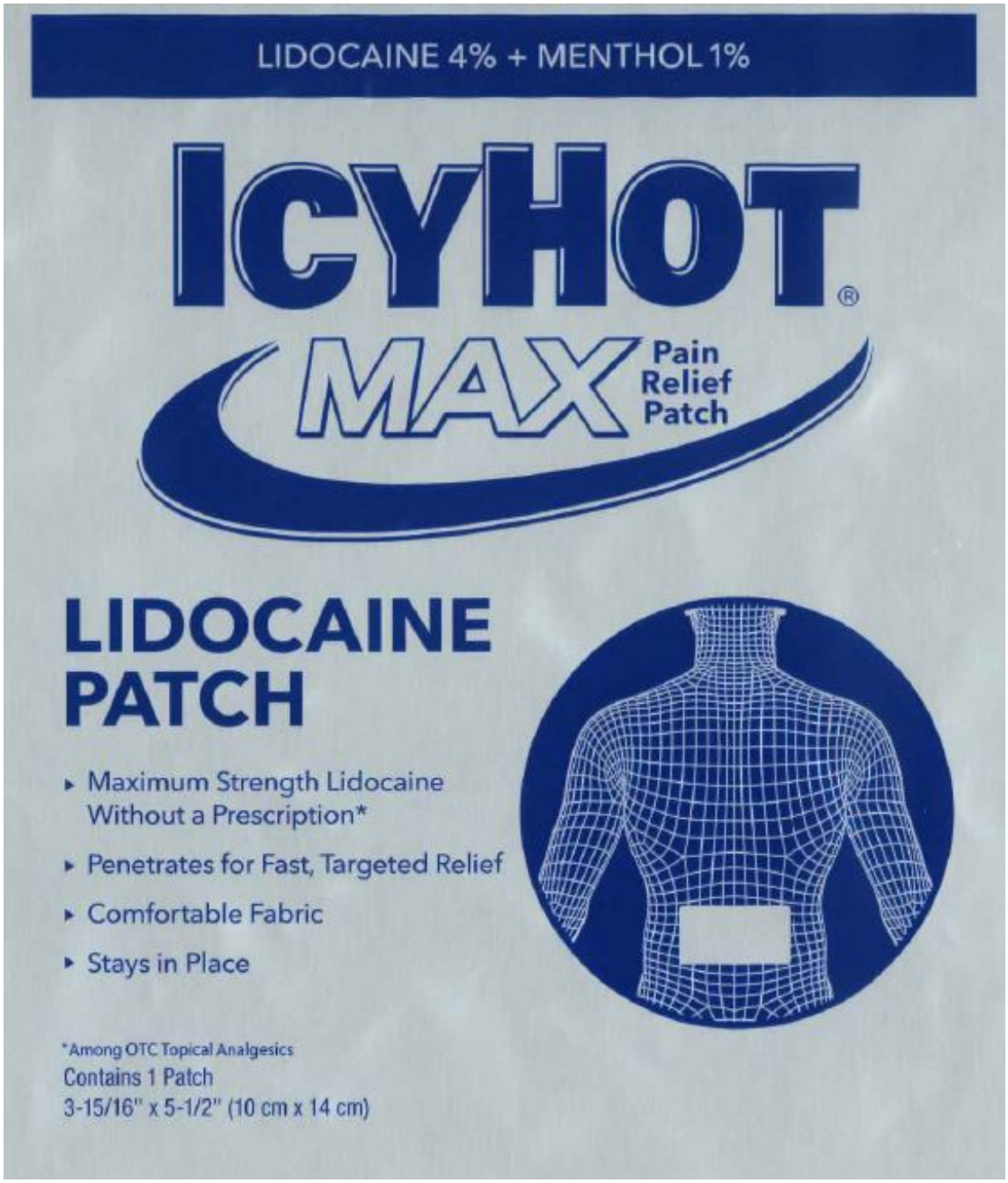
- Maximum Strength Lidocaine
  Without a Prescription*
- Penetrates for Fast, Targeted Relief

- Comfortable Fabric
- Stays in Place

*Among OTC Topical Analgesics
Contains 1 Patch
3-15/16” x 5-1/2” (10 cm x 14 cm)

CHATTEM®
A SANOFI COMPANY
Dist. by Chattem, Inc., a Sanofi Company
P.O. Box 2219
Chattanooga, TN 37409-0219 USA

www.icyhot.com Made in Japan.



Label

ICY HOT

lidocaine, menthol patch

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62168-1720
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	240 mg
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	60 mg

Inactive Ingredients

Ingredient Name	Strength
DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS (UNII: 1K713C615K)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
NONOXYNOL-30 (UNII: JJX07DG188)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TARTARIC ACID (UNII: W4888I119H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
UREA (UNII: 8W8T17847W)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62168-1720-2	400 in 1 CARTON	05/10/2021	
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	505G(a)(3)	01/15/2017	

Labeler - Lead Chemical Co. Ltd. (693727091)

Revised: 1/2024 Lead Chemical Co. Ltd.