ICY HOT MEDICATED- menthol patch ICY HOT MEDICATED, XL- menthol patch Lead Chemical Co., Ltd.

Icy Hot® Medicated Patch

Active Ingredient

<u>Icy Hot Arm, Neck and Leg Patches; and Icy Hot XL Patches</u>
Menthol 5%

Purpose

Topical analgesic

Uses

<u>Icy Hot Arm, Neck and Leg Patches; and Icy Hot XL Patches</u>

temporarily relieves minor pain associated with:

- arthritis
- simple backache
- bursitis
- tendonitis
- muscle strains
- sprains
- bruises
- cramps

Warnings

For external use only

When using this product

Icy Hot Arm, Neck and Leg Patches; and Icy Hot XL Patches

- use only as directed. Read and follow all directions and warnings on this label.
- avoid contact with eyes and mucous membranes
- rare cases of serious burns have been reported with products of this type
- do not apply to wounds or damaged, broken or irritated skin
- do not bandage tightly or apply local heat (such as heating pads) to the area of use
- a transient burning sensation may occur upon application but generally disappears in several days

Stop use and ask a doctor if

<u>Icy Hot Arm, Neck and Leg Patches</u>; and Icy Hot XL Patches

- 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.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

<u>Icy Hot Arm, Neck and Leg Patches</u>; and Icy Hot XL Patches

adults and children over 12 years:

- 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
- wear one Icy Hot patch for up to 8 hours
- repeat as necessary, but no more than 3 times daily

children 12 years or younger: ask a doctor

Inactive ingredients

Icy Hot Arm, Neck and Leg Patches; and Icy Hot XL Patches

aluminum hydroxide, cellulose gum, glycerin, isopropyl myristate, methyl acrylate/2-ethylhexyl acrylate copolymer, nonoxynol-30, polyacrylic acid, polysorbate 80, sodium polyacrylate, sorbitan sesquioleate, starch/acrylic acid graft copolymer sodium salt, talc, tartaric acid, titanium dioxide, water

Principal Display Panel - Arm, Neck & Leg Pouch

MENTHOL 5%

ICY HOT® ORIGINAL Pain Relief Patch

ARM, NECK & LEG PATCH

FEEL IT WORKING INSTANTLY

- Powerful Targeted Relief
- Icy to Dull, Hot to Relax
- Comfortable Fabric
- Stays in Place

Wear For Up To 8 Hours

Contains 5 Patches in 1 Resealable Pouch 3-1/8" x 4-5/8" (8 cm x 12 cm) each

CHATTEM®

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Pouch Label - Arm, Neck, & Leg

Principal Display Panel - XL Back

MENTHOL 5%

ICY HOT®
ORIGINAL
Pain Relief Patch

XL PATCH

FEEL IT WORKING INSTANTLY

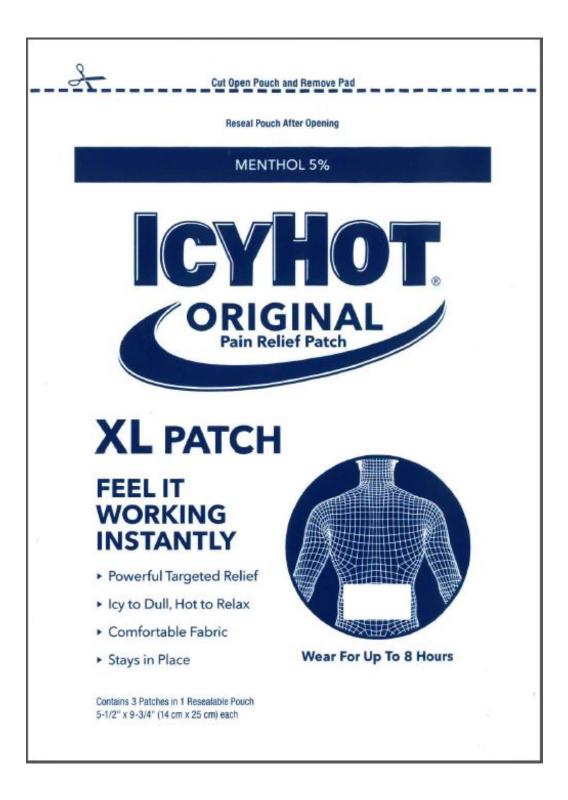
- Powerful Targeted Relief
- Icy to Dull, Hot to Relax
- Comfortable Fabric
- Stays in Place

Wear For Up To 8 Hours

Contains 3 Patches in 1 Resealable Pouch 5-1/2" x 9-3/4" (14 cm x 25 cm) each

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Pouch Label - XL Back

ICY HOT MEDICATED

menthol patch					
Product Information	Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62168-0084		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength			
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	205.5 mg		

Inactive Ingredients			
Ingredient Name	Strength		
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)			
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)			
GLYCERIN (UNII: PDC6A3C0OX)			
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)			
NONOXYNOL-30 (UNII: JJX07DG188)			
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)			
POLYSORBATE 80 (UNII: 6OZP39ZG8H)			
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)			
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)			
TALC (UNII: 7SEV7J4R1U)			
TARTARIC ACID (UNII: W48881119H)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
WATER (UNII: 059QF0KO0R)			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:62168- 0084-3	270 in 1 BOX	06/09/2021	06/30/2026	
1		5 in 1 POUCH; Type 0: Not a Combination Product			
2	NDC:62168- 0084-4	160 in 1 CARTON	12/08/2023		
2		5 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)	11/01/2000		

ICY HOT MEDICATED, XL

menthol patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62168-0847
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	750 mg		

Inactive Ingredients			
Ingredient Name	Strength		
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)			
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)			
GLYCERIN (UNII: PDC6A3C0OX)			
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)			
NONOXYNOL-30 (UNII: JJX07DG188)			
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)			
POLYSORBATE 80 (UNII: 6OZP39ZG8H)			
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05115JN12J)			
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)			
TALC (UNII: 7SEV7J4R1U)			
TARTARIC ACID (UNII: W4888I119H)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
WATER (UNII: 059QF0KO0R)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:62168- 0847-5	120 in 1 CARTON	05/17/2021	07/31/2026		
1		3 in 1 POUCH; Type 0: Not a Combination Product				
2	NDC:62168- 0847-7	100 in 1 CARTON	09/19/2023			
2		3 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)	12/01/2007		

Labeler - Lead Chemical Co., Ltd. (693727091)

Revised: 4/2025 Lead Chemical Co., Ltd.