5% MENTHOL PAIN RELIEF PATCH- pain relief patch patch
Xuzhou Lanting Pharmaceutical Co., Ltd

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

85323-006, 5% Menthol Pain Relief Patch

5% Menthol Pain Relief Patch, Package NDC Code 85323-006-01

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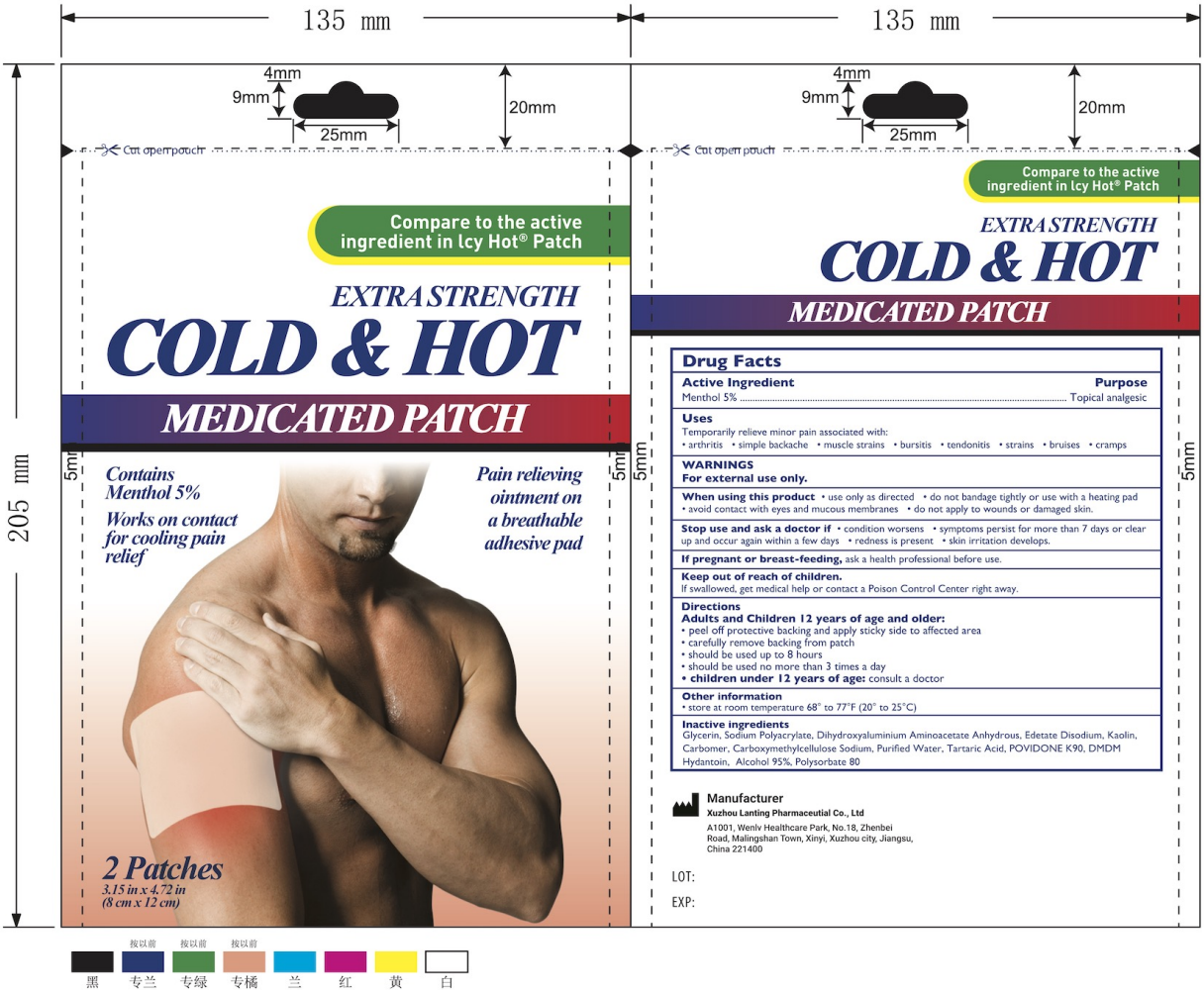

Active Ingredient

Active Ingredient: Menthol 5%

Purpose

Purpose: Topical anesthetic

Warnings

For external use only.

- Do not use
- more than 1 patch at a time
- on wounds or damaged skin
- with a heating pad
- if you are allergic to any ingredients of this product

When using this product

- use only as directed
- do not bandage tightly or use with a heating pad
- avoid contact with the eyes and mucous membranes
- do not apply to wounds or damaged skin.

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few day
- redness is present
- skin irritation develops

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.

Inactive Ingredients

Glycerin, Sodium Polyacrylate, Dihydroxyaluminum Aminoacetate Anhydrous, Edetate Disodium, Kaolin, Carbomer, Carboxymethylcellulose Sodium, Purified Water, Tartaric Acid, Povidone K90, DMDM Hydantoin, Alcohol 95%, Polysorbate 80.

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

Other information

Other Information

- store at roo temperature 68 to 77℉ (20 to 25℃)

Directions

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Adult and children 12 years of age and older:

- peel off protective backing and apply sticky side to affected area
- carefully remove backing from patch
- should be used to 8 hours
- ashould be used no more than 3 times a day
- **Children under 12 years of age:** consult a doctor

Uses

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Temporarily relieves minor pains associated with:

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

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5% MENTHOL PAIN RELIEF PATCH

pain relief patch patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85323-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	5 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
KAOLIN (UNII: 24H4NWX5CO)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
CARBOMER 934 (UNII: Z135WT9208)	
GLYCERIN (UNII: PDC6A3C0OX)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS (UNII: 1K713C615K)	
POVIDONE K90 (UNII: RDH86HJV5Z)	
TARTARIC ACID (UNII: W4888I119H)	
ALCOHOL 95% (UNII: 7528N5H79B)	

Product Characteristics			
Color		**Score**	

Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85323-006-01	24 in 1 BOX	06/10/2025	
1		2 in 1 POUCH		
1		8.5 g in 1 PATCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

unapproved drug other		06/10/2025	

Labeler - Xuzhou Lanting Pharmaceutical Co., Ltd (457641059)

Registrant - Xuzhou Lanting Pharmaceutical Co., Ltd (457641059)

Establishment

Name	Address	ID/FEI	Business Operations
Xuzhou Lanting Pharmaceutical Co., Ltd		457641059	manufacture(85323-006)

Revised: 6/2025 Xuzhou Lanting Pharmaceutical Co., Ltd