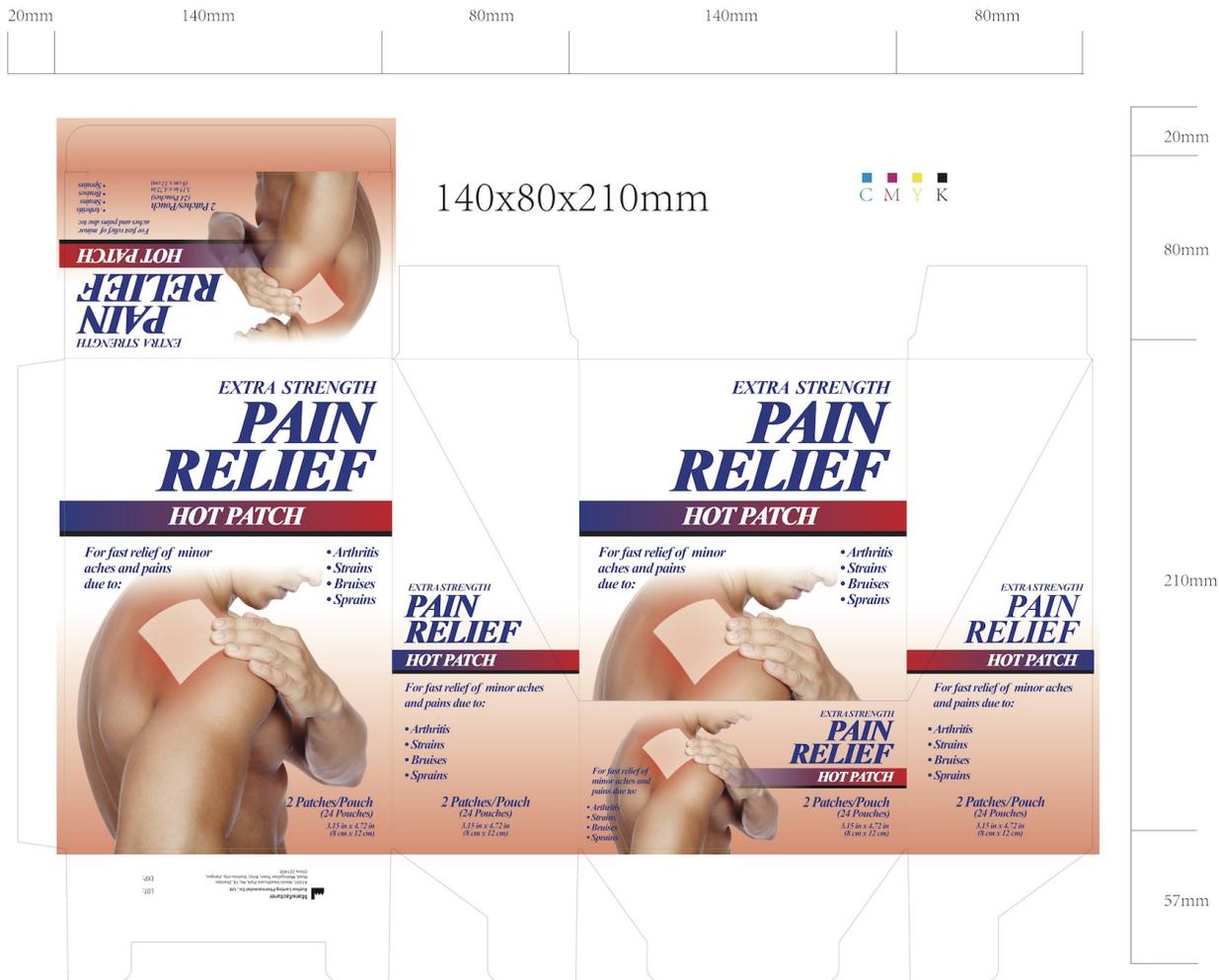


3% MENTHOL PLUS 0.083% CAPSAICIN PAIN RELIEF PATCH- pain relief patch, pain relief strip patch
Xuzhou Lanting Pharmaceutical Co., Ltd

85323-008, 3% Menthol plus 0.083% Capsaicin Pain Relief Patch

3% Menthol plus 0.083% Capsaicin Pain Relief Patch, Package NDC Code: 85323-008-01

3% Menthol plus 0.083% Capsaicin Pain Relief Patch, Package NDC Code: 85323-008-01





黑 专兰 专橘 兰 红 黄 白

白与J24050603共用

Active Ingredient

Active Ingredients (in each gram):

- Menthol 30mg
- Capsaicin 0.83mg

Purpose

Purpose: Topical anesthetic

Uses

Uses

Temporarily relieves minor pain associated with:

- arthritis
- simple backache
- bursitis
- tendonitis
- muscle strains

- bruises
- cramps

Warnings

WARNINGS: EXTERNAL USE ONLY

When using this product

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

Stop use and ask a doctor

- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days
- 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 (1-800-222-1222) immediately.

Other information

Other Information

- store at room temperature 68 to 77°F (20-25°C)

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, Propylene Glycol.

KEEP OUT OF REACH OF CHILDREN.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

DIRECTIONS

Adult and children 12 years of age and older:

- Partially peel back protective film and apply exposed patch to site of pain. Carefully remove remaining film while pressing patch to skin for secure adhesions
- Apply to affected area not more than 3 to 4 times daily

Children under 12 years of age: DO NOT use without consulting a doctor.

Temporarily relieves minor pain associated with:

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

Adult and children 12 years of age and older:

- Partially peel back protective film and apply exposed patch to site of pain. Carefully remove remaining film while pressing patch to skin for secure adhesions
- Apply to affected area not more than 3 to 4 times daily

Children under 12 years of age: DO NOT use without consulting a doctor.

3% MENTHOL PLUS 0.083% CAPSAICIN PAIN RELIEF PATCH

pain relief patch, pain relief strip patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85323-008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	3 g in 100 g
CAPSAICIN (UNII: S07O44R1ZM) (CAPSAICIN - UNII:S07O44R1ZM)	CAPSAICIN	83 mg in 100 g

Inactive Ingredients

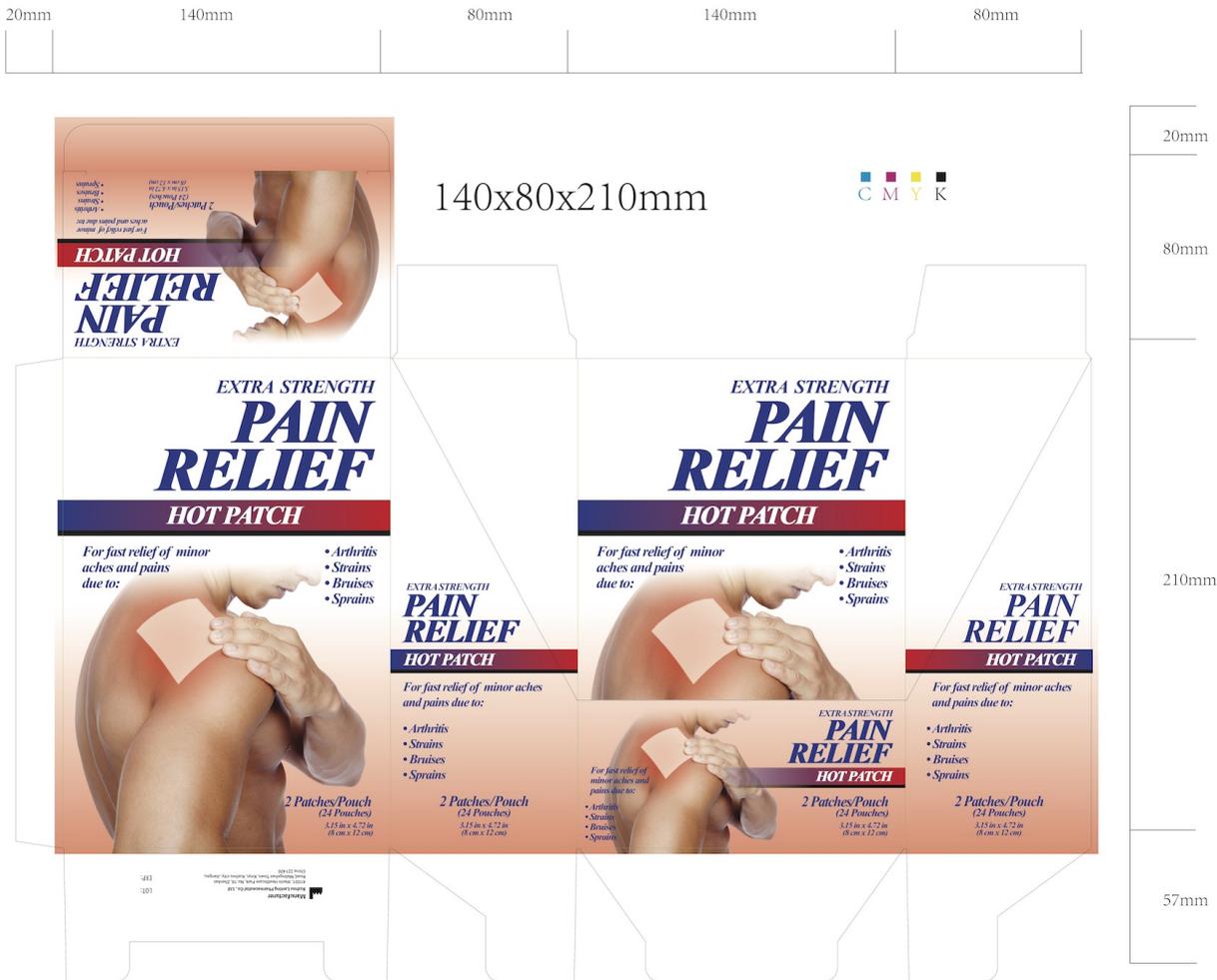
Ingredient Name	Strength
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
ALCOHOL 95% (UNII: 7528N5H79B)	
WATER (UNII: 059QF0KO0R)	
DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS (UNII: 1K713C615K)	
KAOLIN (UNII: 24H4NWX5CO)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CARBOMER 934 (UNII: Z135WT9208)	
TARTARIC ACID (UNII: W4888I119H)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE K90 (UNII: RDH86HJV5Z)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85323-008-01	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	Application Number or Monograph	Marketing Start	Marketing End
-----------	---------------------------------	-----------------	---------------

Category	Citation	Date	Date
OTC Monograph Drug	505G(a)(3)	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(85323-008)

Revised: 6/2025

Xuzhou Lanting Pharmaceutical Co., Ltd