

# 0.025% CAPSAICIN PAIN RELIEF PATCH- capsaicin pain relief patch, capsaicin pain relief strip patch

Xuzhou Lanting Pharmaceutical Co., Ltd

0.025% Capsaicin Pain relief Patch, Package NDC Code: 85323-001-00

0.025% Capsaicin Pain relief Patch, Package NDC Code: 85323-001-00



## Active Ingredient

Active Ingredient: 0.025% Capsaicin

## Purpose

Purpose: Topical anesthetic

## Warnings

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
- avoid contact with the eyes, mucous membranes or rashes
- do not bandage tightly

Stop use and ask a doctor if

- localized skin reactions occur, such as rash, itching, redness, irritation, pain, swelling and blistering
- condition worsen
- 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. If swallowed, get medical help or contact a Poison Control Center right away.

## **Uses**

Uses For temporary relief of minor aches & pains of muscles & joints associated with:

- simple backache
- arthritis
- strains
- bruises
- sprains

## **Other information**

- Avoid storing product in direct sunlight
- Protect product from excessive moisture

## **Inactive Ingredients**

Glycerin, dihydroxyaluminum aminoacetate anhydrous, edetate disodium, tartaric acid, povidone K90, dmdm hydantoin, polysorbate 80, propylene glycol, carboxymethylcellulose sodium, kaolin, carbomer, sodium polyacrylate, water.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

## **Directions**

Uses For temporary relief of minor aches & pains of muscles & joints associated with:

- simple backache
- arthritis
- strains
- bruises
- sprains

Adult 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  
 remove patch from the skin after at most 8-hour application  
 Children under 12 years of age: consult a doctor

Adult 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  
 remove patch from the skin after at most 8-hour application

Children under 12 years of age: consult a doctor

## 0.025% CAPSAICIN PAIN RELIEF PATCH

capsaicin pain relief patch, capsaicin pain relief strip patch

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:85323-001
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CAPSAICIN</b> (UNII: S07O44R1ZM) (CAPSAICIN - UNII:S07O44R1ZM)	CAPSAICIN	0.025 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>KAOLIN</b> (UNII: 24H4NWX5CO)	
<b>CARBOMER</b> (UNII: 0A5MM307FC)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>SODIUM POLYACRYLATE (2500000 MW)</b> (UNII: 05I15JN12J)	
<b>GLYCERIN</b> (UNII: PDC6A3C00X)	
<b>DMDM HYDANTOIN</b> (UNII: BYR0546TOW)	
<b>DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS</b> (UNII: 1K713C615K)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>TARTARIC ACID</b> (UNII: W4888I119H)	
<b>POVIDONE K90</b> (UNII: RDH86HJV5Z)	
<b>CARBOXYMETHYLCELLULOSE SODIUM</b> (UNII: K679OBS311)	

### Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	

<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85323-001-00	5 in 1 BOX	03/01/2025	
1		1 in 1 POUCH		
1		11 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
OTC Monograph Drug	505G(a)(3)	03/01/2025	

**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(85323-001)

Revised: 3/2025

Xuzhou Lanting Pharmaceutical Co., Ltd