

PAIN RELIEVING PATCH- lidocaine patch
FlexiGo Inc.



13.5*11*16cm



13.5*11*16cm



Drug Facts

Active ingredient Lidocaine 4% Purpose Topical Analgesic

Uses

Temporarily relieves muscle soreness and minor joint pains in the wrist, knees, back, hips, shoulders, elbows.

Warnings

For external use only

Do not use

On cuts or damaged skin
With a heating pad or device
With other ointments, creams, sprays, or oils.

If you are allergic to any ingredients of this product

When using this product

Avoid contact with the eyes, mucous membranes, or rashes
Do not swallow.

Stop use and ask a doctor if

Skin reactions such as redness, swelling, blistering or other discomfort occur.

Symptoms persist for more than 7 days.

If pregnant or breast-feeding, ask a health professional before use.

KEEP OUT OF REACH OF CHILDREN
If swallowed accidentally, get medical help or contact Poison Control Center immediately.

Directions

Clean and dry the affected area
Tear off the protective film and apply the exposed part of the patch to the affected area.

Carefully remove remaining film while pressing the patch firmly on the skin.
Remove the patch from the skin after at most 8-hour application.

Children under age of 12: Consult a doctor.

Inactive ingredients

ALUMINUM HYDROXIDE, SODIUM POLYACRYLATE (2000000 MW), AQUA, TITANIUM DIOXIDE, POLYISOBUTYLENE, POLYSORBATE 80, GLYCERIN, DMDM HYDANTION, CAPRYLIC OIL, C12-15 CARBOXYMETHYL CELLULOSE SODIUM (1:3 CARBOXYMETHYL SUBSTITUTION PER DAU), SODIUM 1,2-HEXYL SALT 2%, SILICA, TITANIUM DIOXIDE, ALUMINUM CYANATE

Distributed by:
FlexGo Inc
9815 Taylor St
Englewood Colorado 80110

Design by USA, Made in P.R.C

Usage



MFG:11/25/2025

EXP:11/24/2028

BATCH NUMBER:20251125-01

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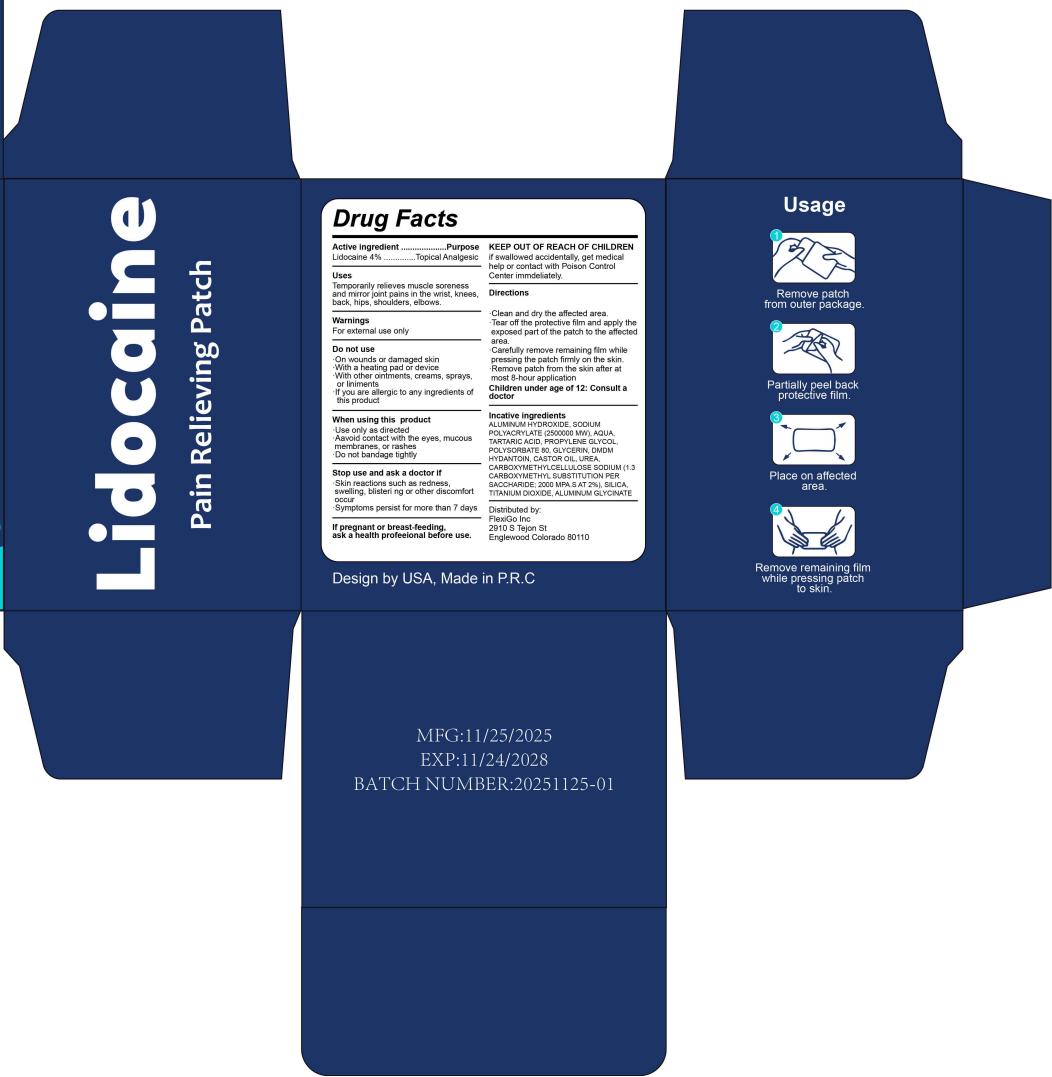
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PAIN RELIEVING PATCH

lidocaine patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDc:85524-2601
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	0.04 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CY0341L)	
AQUA (UNII: 059QF0KO0R)	
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	
TARTARIC ACID (UNII: W4888I119H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
GLYCERIN (UNII: PDC6A3C00X)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
CASTOR OIL (UNII: D5340Y2I9G)	
UREA (UNII: 8W8T17847W)	
CARBOXYMETHYLCELLULOSE SODIUM (1.3 CARBOXYMETHYL SUBSTITUTION PER SACCHARIDE; 2000 MPAS AT 2%) (UNII: 8UX21M67IJ)	
SILICA (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ALUMINUM GLYCINATE (UNII: 1K713C615K)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85524-2601-1	30 g in 1 BOX; Type 0: Not a Combination Product	01/18/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/18/2026	

Labeler - FlexiGo Inc. (118951527)

Establishment

Name	Address	ID/FEI	Business Operations
FlexiGo Inc		118951527	label(85524-2601) , manufacture(85524-2601)