

**MAXIMUM STRENGTH 4% LIDOCAINE PATCH GOOD NEIGHBOR PHARMACY-
lidocaine patch
InterMed Laboratories Private Limited**

Maximum Strength 4% Lidocaine Patch

Active Ingredient

Lidocaine 4%

Purpose

External Analgesic

Uses

For temporary relief of pain

Warnings

For external use only

Do Not Use

- more than 1 patch on your body at a time or on cut, irritated or swollen skin
- for more than one week without consulting a doctor
- over raw surfaces or blistered areas
- if pouch is damaged or opened

When Using This Product

- use only as directed. Read and follow all directions and warnings on this label.
- do not allow contact with the eyes

Stop Use and Ask a Doctor if

- condition worsens
- skin irritation develops
- symptoms persist for more than 7 days or 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 and Pets

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

Directions

Adults and children over 2 years:

- clean and dry affected area free of lotions, ointments and creams.
- carefully remove backing from patch starting at a corner. Apply sticky side of patch to affected area.
- Do not use more than one patch in a 12 hour period. Maximum 2 patches per day. Discard patch after single use.

children 2 years or younger:ask a physician.

Inactive Ingredients

dihydroxy aluminium aminoacetate, disodium EDTA, glycerin, methyl paraben, polyvinyl alcohol, propyl paraben, propylene glycol, sodium carboxy methyl cellulose, sodium polyacrylate, tartaric acid, titanium dioxide, water

Bulk Package Label



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lidocaine patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71811-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	400 mg

Inactive Ingredients

Ingredient Name	Strength
TARTARIC ACID (UNII: W4888I19H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	

GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JN12J)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71811-010-05	5 in 1 CARTON	11/06/2025	
1		1 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	M017	11/06/2025	

Labeler - InterMed Laboratories Private Limited (676244169)

Revised: 2/2026

InterMed Laboratories Private Limited