

LIDOZEN PATCH- lidocaine, menthol patch
Village Pharma LLC

Lidozen Patch

DRUG FACTS:

ACTIVE INGREDIENTS:

Lidocaine 4.00%

Menthol 1.00%

Topical Anesthetic

External Analgesic

USES:

For temporary relief of pain

WARNINGS:

- For external use only.
- Avoid contact with eyes.
- If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a physician.

Do not use

- in large quantities, particularly over raw surfaces or blistered areas.

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.

DIRECTIONS (Adults and Children Over 12 Years):

Clean and dry affected area.

Remove patch from backing and apply to affected area.

Use only one patch at a time, and maximum of four patches / day.

Leave patch on affected area for up to 8-hours.

Do not use patches for longer than five consecutive days.

Children under 12 should consult physician prior to use.

INACTIVE INGREDIENTS

Water, Glycerol, Sodium Polyacrylate, Propylene Glycol, Polysorbate 80, Tartaric Acid, Dihydroxyaluminium Aminoacetate, Methylparaben

Store below 25 degrees Celsius. Avoid direct sunlight.

Package Labeling:

LidozenPatch[®]

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Manufactured For:
Village Pharma, LLC
Agoura Hills, CA 91301
For Questions or Comments please email: info@villagepharma.com

Made in China. Patent Pending



LidozenPatch[®]

(Lidocaine 4% / Menthol 1%)

VillagePharma

NDC 71574-905-05
Patch Size 10 x 14cm
Qty 5 Patches
TOPICAL ANALGESIC PATCH

LidozenPatch®

LIDOZEN PATCH

lidocaine, menthol patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71574-905
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	40 mg in 1 g
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
TARTARIC ACID (UNII: W4888I119H)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	
METHYLPARABEN (UNII: A2I8C7HI9T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71574-905-05	5 in 1 POUCH	09/01/2022	
1		1 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	M017	09/01/2022	

Labeler - Village Pharma LLC (080749749)

Registrant - Village Pharma LLC (080749749)

Establishment

Name	Address	ID/FEI	Business Operations
Beijing HKKY Medical Tech. Co., Ltd.		544434817	MANUFACTURE(71574-905)

Revised: 1/2026

Village Pharma LLC