LIDOZEN- lidocaine hydrochloride, menthol gel
Village Pharma, LLC

Lidozen Gel

DRUG FACTS:

ACTIVE INGREDIENTS:

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

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):

Apply directly to affected area. Do not use more than four times per day.

INACTIVE INGREDIENTS:

Aloe Barbadensis Leaf (Aloe Vera Juice) Gel, Aqua (Deionized Water), Arnica Montana Extract, Boswellia Serrata Extract, Camellia Sinensis Leaf (Green Tea) Extract, Carbomer, Ethylhexylglycerin, Glycerin, Isopropyl Myristate, PEG-8, Phenoxyethanol, Polysorbte-80, Sodium Lauryl Sulfate, Triethanolamine, FD&C Blue #1, FD&C Yellow #5

Package Labeling:

NDC 71574-300-72

LidozenGel®

(Lidocaine 4% / Menthol 1%)

VillagePharma

120mL (4 fl oz)
TOPICAL ANALGESIC GEL

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Menthol	1.00%	External Analgesic

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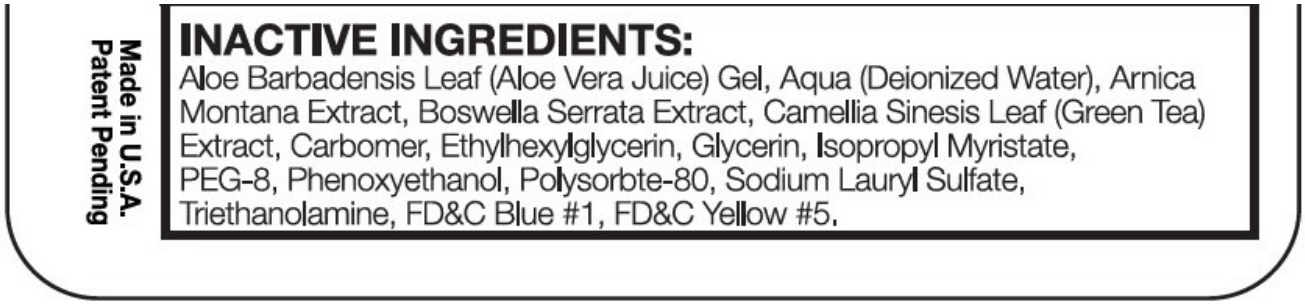
Manufactured For:
Village Pharma, LLC
Agoura Hills, CA 91301

For Questions or Comments
Please E-mail:
info@villagepharma.com



LIDOZEN

lidocaine hydrochloride, menthol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71574-300
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	40 mg in 1 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ARNICA MONTANA (UNII: O80TY208ZW)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71574-300-72	1 in 1 BOX	07/22/2017	
1		120 mL in 1 BOTTLE; 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	07/22/2017	

Labeler - Village Pharma, LLC (080749749)

Revised: 11/2023 Village Pharma, LLC