

LIDOSOLVE PAIN RELIEF- lidocaine hydrochloride liquid
HENNIKA HEALTH, LLC

LIDOSOLVE PAIN RELIEF SPRAY

Drug Facts

Active ingredients

Lidocaine HCL 4.00%

Purpose

Topical Analgesic

Uses:

- For the temporary relief of pain.

Warnings:

not intended for ingestion. **For external use only**

Do not use

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

When using this product

- Avoid contact with the eyes.

Stop use and ask a doctor if

- condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children.

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

If pregnant or breast-feeding,

ask a health professional before use.

Directions:

- Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- Children under 2 years of age: consult a doctor.

Other information:

- Store at 20-25°C (68-77°F) and protect from moisture.

Inactive ingredients:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Calendula Officinalis Extract, Chamomilla Matricaria (Chamomile) Extract, Cyanocobalamin (Vitamin B 12), Ethylhexylglycerin, Ethyl Vanillin, Glycerin, Phenoxyethanol, Polysorbate-20, Pyridoxine HCL (Vitamin B6), SD-Alcohol 40B, Sodium Metabisulfite, Tetrasodium EDTA, Zemea (Corn) Propanediol

Questions?

1-800-272-1875 / www.hennikahealth.com

Package Labeling:

lidosolve

PAIN RELIEF SPRAY

4% LIDOCAINE HCL

- ✦ Aloe Vera
- ✦ Vitamins B6 & B12

Natural Healing with a Fresh Vanilla Scent



NET WT 6 OZ (170G)



Distributed by:
Hennika Health, LLC
Avondale, AZ 85323

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LIDOSOLVE PAIN RELIEF

lidocaine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83646-420
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII: 98PI200987)	LIDOCAINE	40 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
CHAMOMILE (UNII: FGL3685T2X)	
CYANOCOBALAMIN (UNII: P6YC3EG204)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ETHYL VANILLIN (UNII: YC9ST449YJ)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CORN (UNII: 0N8672707O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83646-420-00	170 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/22/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/22/2023	

Labeler - HENNIKA HEALTH, LLC (118847402)

Revised: 11/2023

HENNIKA HEALTH, LLC