

LIDOVARA- lidovara gel
Oncora Pharma, LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Lidovara™ 2.8% Gel

DESCRIPTION

Lidovara™ is a topical gel containing Lidocaine Hydrochloride 28 mg/g (2.8%).

INDICATIONS AND USAGE

Lidovara™ is indicated for topical analgesia as directed by a healthcare provider.

CONTRAINDICATIONS

Lidovara™ is contraindicated in patients with known hypersensitivity to lidocaine or any component of the formulation.

WARNINGS

For external use only. Not for ophthalmic use.

Avoid contact with eyes.

Excessive application may result in increased systemic absorption.

Keep out of reach of children.

OVERDOSAGE

Excessive topical application may result in increased systemic absorption of lidocaine. In the event of overdose, discontinue use and seek medical attention.

DOSAGE AND ADMINISTRATION

For topical use only.

Apply to the affected area 3 to 4 times daily or as directed by a healthcare provider.

HOW SUPPLIED

Lidovara™ (Lidocaine Hydrochloride 28 mg/g) is supplied as a clear topical gel in a 3.5 oz

(100 g) tube.

NDC 85477-303-07

PRINCIPAL DISPLAY PANEL

NDC: 85477-303-07
For RX only



LidovaraTM
LIDOCAINE 2.8% Gel
TOPICAL PAIN RELIEF

KEEP OUT OF REACH OF CHILDREN
FOR EXTERNAL USE ONLY. NOT FOR OPHTHALMIC USE.

Net Wt. 3.5 oz. (100g)



DIRECTIONS: Apply Lidovara to the skin 3 to 4 times daily or as directed by a healthcare provider.

WARNINGS: Do not use this product if you are allergic to any of the ingredients. Avoid contact with eyes.

STORAGE: Store at 25°C (77°F); excursions permitted to 15°-30°C (59°-86°F). [See USP Controlled Room Temperature].

Active Ingredients: Lidocaine HCl 2.8%

Inactive Ingredients: Aqua (Water), Hydroxyethylcellulose, Phenoxyethanol, Ethylhexylglycerin, Sodium hydroxide, Lactic acid.



Manufactured for:
Oncora Pharma
Dallas, TX, 75228

pdp

LIDOVARA

lidovara gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:85477-303
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
-----------------	-------------------	----------

LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)		LIDOCAINE	28 mg in 1 g	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
HYDROXYETHYLCELLULOSE (UNII: T4V6TWG28D)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
LACTIC ACID (UNII: 33X04XA5AT)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85477-303-07	100 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	02/26/2026	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		02/26/2026		

Labeler - Oncora Pharma, LLC (119482542)

Revised: 2/2026

Oncora Pharma, LLC