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.

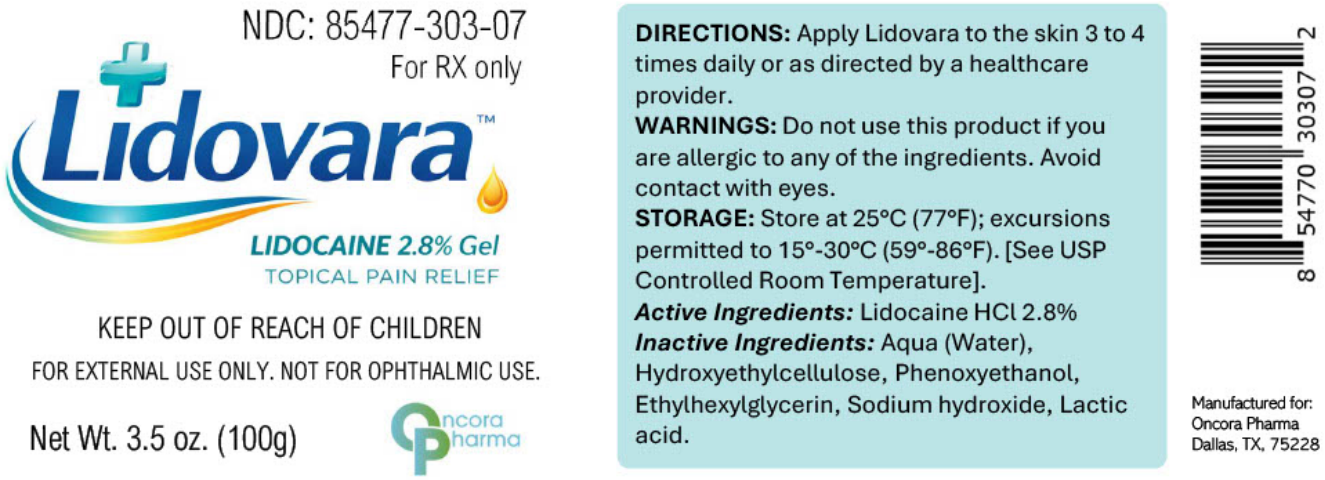
HOW SUPPLIED

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



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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