

DOLOGESIC ROLL-ON- lidocaine hcl liquid
Llorens Pharmaceutical International Division, Inc.

Dologesic Roll-On 513

Active Ingredient

Lidocaine HCL, USP 4%

Purpose

Topical Anesthetic

Use

Temporarily relieves minor pains

Warnings

For external use only

Do not use

- on large areas of the body or on cut, irritated or swollen skin
- on puncture wounds
- for more than one week without consulting a doctor

When using this product

- use only as directed. Read and follow all directions and warnings on this label.
- do not allow contact with eyes
- do not bandage or apply local heat (such as heating pads) to the area of use.

Stop use and ask doctor if

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

If pregnant or breast-feeding, ask a doctor 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 of age
- Apply generously to the affected area as needed every 6-8 hours, not to exceed 3 application in a 24 hour period.
- Not for use in children under 12 years of age.

Inactive ingredients

camphor, glycerin, isopropyl alcohol, menthol, methylparaben, propylparaben, purified water and xantham gum.

Questions

Llorens Pharmaceutical International Division, Inc.

Miami FI 33147

DOLOGESIC
TOPICAL ANESTHESIC

NDC 54859-513-02

PAIN RELIEF

- ▶ Maximum Strength
- ▶ Easy to Apply
- ▶ In a menthol-camphor base

ROLL-ON LIQUID

Contents: 2 fl oz (60 mL)

Drug Facts

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Manufactured by: Code#L-90 Rev.02/18
Llorens Pharmaceutical International Division, Inc.
Miami, FL 33147

For lot number and expiration date, see top of cap. 3  54859 51302 1



DOLOGESIC ROLL-ON

lidocaine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54859-513
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MENTHOL (UNII: L7T10EIP3A)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

WATER (UNII: 059QF0KO0R)

XANTHAN GUM (UNII: TTV12P4NEE)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54859-513-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/01/2018	

Labeler - Llorens Pharmaceutical International Division, Inc. (037342305)

Revised: 12/2025

Llorens Pharmaceutical International Division, Inc.