

AMINO RIP MUSCLE PAIN RELIEF- lidocaine hcl liquid
LLORENS PHARMACEUTICALS INTERNATIONAL DIVISION

Amino Rip Muscle 415

Active Ingredients

Lidocaine HCl

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 the reach of children
- If swallowed, get medical help or contact a Poison Control Center right away.

Keep out of reach of children.

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 applications 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, xanthan gum

NDC 54859-415-02

AMINORIP[®]**MUSCLE
PAIN
RELIEF**

• TOPICAL ANESTHETIC •

**UPPER & LOWER BACK
NECK | THIGHS | CALVES****ROLL-ON**

2 FL OZ (60 mL)

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Manufactured byLlorens Pharmaceutical International Division, Inc
Miami FL, 33147

For Lot Number and Expiration Date, see top of cap.

LLORENS
REPORTS DIVISION
WWW.AMINORIP.COM

Code: L-22 Rev.: 05/19

AMINO RIP MUSCLE PAIN RELIEF

lidocaine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54859-415
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MENTHOL (UNII: L7T10EIP3A)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54859-415-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/01/2019	

Labeler - LLORENS PHARMACEUTICALS INTERNATIONAL DIVISION (037342305)

Revised: 1/2026

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