

**LUMICAIN- aluminium chloride hexahydrate solution**  
**Medical Products Laboratories, Inc.**

*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.*

**lumicain Topical Hemostatic Solution**

FOR TOPICAL APPLICATION ONLY

For Granulation Tissue Growth: Allow packing saturated in Lumicain to remain in nail groove for 48 hours; repeat if necessary.

If hemorrhage is profuse, dress wound with gauze saturated with Lumicain and allow to remain for 24 hours or longer.

CAUTION: Federal Law restricts sale and use to physician or licensed practitioner.

NDC 10733-412-60		Directions: For Granulation Tissue Growth: Allow packing saturated with Lumicain to remain in nail groove for 48 hours; repeat if necessary.
Manufactured for: Premier® Medical Products 1710 Romano Drive Plymouth Meeting, PA 19462 U.S.A.	<b>Lumicain®</b> Topical Hemostatic Solution	If hemorrhage is profuse, dress wound with gauze saturated with Lumicain and allow to remain for 24 hours or longer.
<b>Rx ONLY</b> Store at 68°F-77°F (20°C-25°C)	For Rapid Control of Minor Hemorrhage	Each Gram Contains: 250mg of Aluminum Chloride-6-Hydrate in an aqueous base.
To obtain an SDS visit www.premusa.com or call Premier at 610-239-6000.	60cc  9045010	Made in U.S.A. 0818067 Rev3 MPL 1800068
Manufactured by: Medical Products Laboratories, Inc., Philadelphia, PA 19115-1083, U.S.A.		

premier lumicain™ Topical Hemostatic Solution 60cc

For Rapid Control of Minor Hemorrhage

Each Gram Contains: Aluminium Chloride.....250 mg.

In an aqueous base.

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<b>LUMICAIN</b>			
aluminium chloride hexahydrate solution			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:10733-412
<b>Route of Administration</b>	TOPICAL		
<b>Active Ingredient/Active Moiety</b>			
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
	ALUMINUM CHLORIDE (UNII: 3CYT62D3GA) (ALUMINUM CATION - UNII:3XHB1D032B)	ALUMINUM CHLORIDE	250 mg in 1 g

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10733-412-60	67 g in 1 BOTTLE; Type 0: Not a Combination Product	09/29/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/29/2010	

**Labeler** - Medical Products Laboratories, Inc. (002290302)**Registrant** - Medical Products Laboratories, Inc. (002290302)**Establishment**

Name	Address	ID/FEI	Business Operations
Medical Products Laboratories, Inc.		002290302	analysis(10733-412) , manufacture(10733-412) , label(10733-412) , pack(10733-412)

Revised: 12/2020

Medical Products Laboratories, Inc.