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.

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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 hemmorrhage 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.

premier lumicain ™ Topical Hemostatic Solution 60cc

For Rapid Control of Minor Hemorrhage

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

In an aqueous base.

Enter section text here

LUMICAIN

aluminium chloride hexahydrate solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:10733-412
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
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.