

LACTIC ACID 10% / NIACINAMIDE 4% - lactic acid 10% / niacinamide 4% cream
Sincerus Florida, 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](#).

LACTIC ACID 10% / NIACINAMIDE 4%

Directions for use



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As directed by Physician.

Apply topically. For external use only. Wash hands after use.
Store at controlled room temperature (20-25C).

Sincerus Florida, LLC (800) 604-5032

3265 W McNab Rd, Pompano Beach, FL 33069

To report suspected adverse reactions, contact

Sincerus Florida, LLC at (800) 604-5032, or FDA
at www.FDA.gov/MedWatch or (800) FDA-1088.

Office use only. Not for resale.



Sincerus Florida, LLC adverse reactions.



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Active, inactive



LACTIC
NIACINAMIDE
CREAM

SINCERUS

Rx only

BUD: 01/01/1970

Lot: 1A1068ABCD EFGH @1

MFG: 01/01/1970

Active ingredients

Lactic Acid USP 10%

Niacinamide USP 4%

Inactive ingredients

Ascorbyl Palmitate FCC 1%

Butylated Hydroxytoluene NF (BHT) 0.1%

Kojic Acid 4%

Potassium Azeloyl Diglycinate 10%

Suspendisse Cream 70.9%

NDC 72934-2134-2 LACTIC ACID USP 10% / NIACINAMIDE USP 4% Cream 30 gm Sincerus

NDC 72934-2134-2

**LACTIC ACID USP 10%
NIACINAMIDE USP 4%
CREAM 30gm**

Rx only
BUD: 01/01/1970
Active ingredients



LACTIC ACID 10% / NIACINAMIDE 4%

lactic acid 10% / niacinamide 4% cream

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2134
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)		NIACINAMIDE	4 g in 100 g	
LACTIC ACID (UNII: 33X04XA5AT) (LACTIC ACID - UNII:33X04XA5AT)		LACTIC ACID	10 g in 100 g	
Product Characteristics				
Color	yellow	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-2134-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2019	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		05/01/2019		

Labeler - Sincerus Florida, LLC (080105003)

Establishment			
Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2134)