

**LACTIC ACID 30% / SALICYLIC ACID 30% - lactic acid 30% / salicylic acid 30% 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](#).*

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**LACTIC ACID 30% / SALICYLIC ACID 30%**

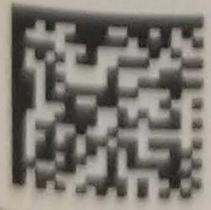
**Directions for use**



**Directions for use**

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](http://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  
SALIC  
CREAM

SINO

Rx only  
BUD: 01/01/1970

Lot: 221032ABCD EFGH@1  
MFG: 01/01/1970

**Active ingredients**

Lactic Acid USP ..... 30%  
Salicylic Acid USP ..... 30%

**Inactive ingredients**

Krisgel 100 ..... 1%  
Salt Stable Ls Cream ..... 39%

**NDC 72934- 2136-2 LACTIC ACID USP 30% / SALICYLIC ACID USP 30%. Cream 30gm**



Rx only  
EXP: 01/01/1970

**NDC 72934-2136-2**

**LACTIC ACID USP 30%**

**SALICYLIC ACID USP 30%**

**CREAM 30gm**



## LACTIC ACID 30% / SALICYLIC ACID 30%

lactic acid 30% / salicylic acid 30% cream

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-2136
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	30 g in 100 g
LACTIC ACID (UNII: 33X04XA5AT) (LACTIC ACID - UNII:33X04XA5AT)	LACTIC ACID	30 g in 100 g

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-2136-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/17/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/17/2019	

**Labeler** - Sincerus Florida, LLC (080105003)

### Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2136)

Revised: 5/2019

Sincerus Florida, LLC