

**CLOBETASOL PROPIONATE 0.05% / NIACINAMIDE 4% - clobetasol propionate 0.05% / 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](#).*

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**CLOBETASOL PROPIONATE 0.05% / 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](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**



CLOBETASOL  
USP 0.05%  
NIACINAMIDE  
CREAM



Rx only  
BUD: 01/01/1970

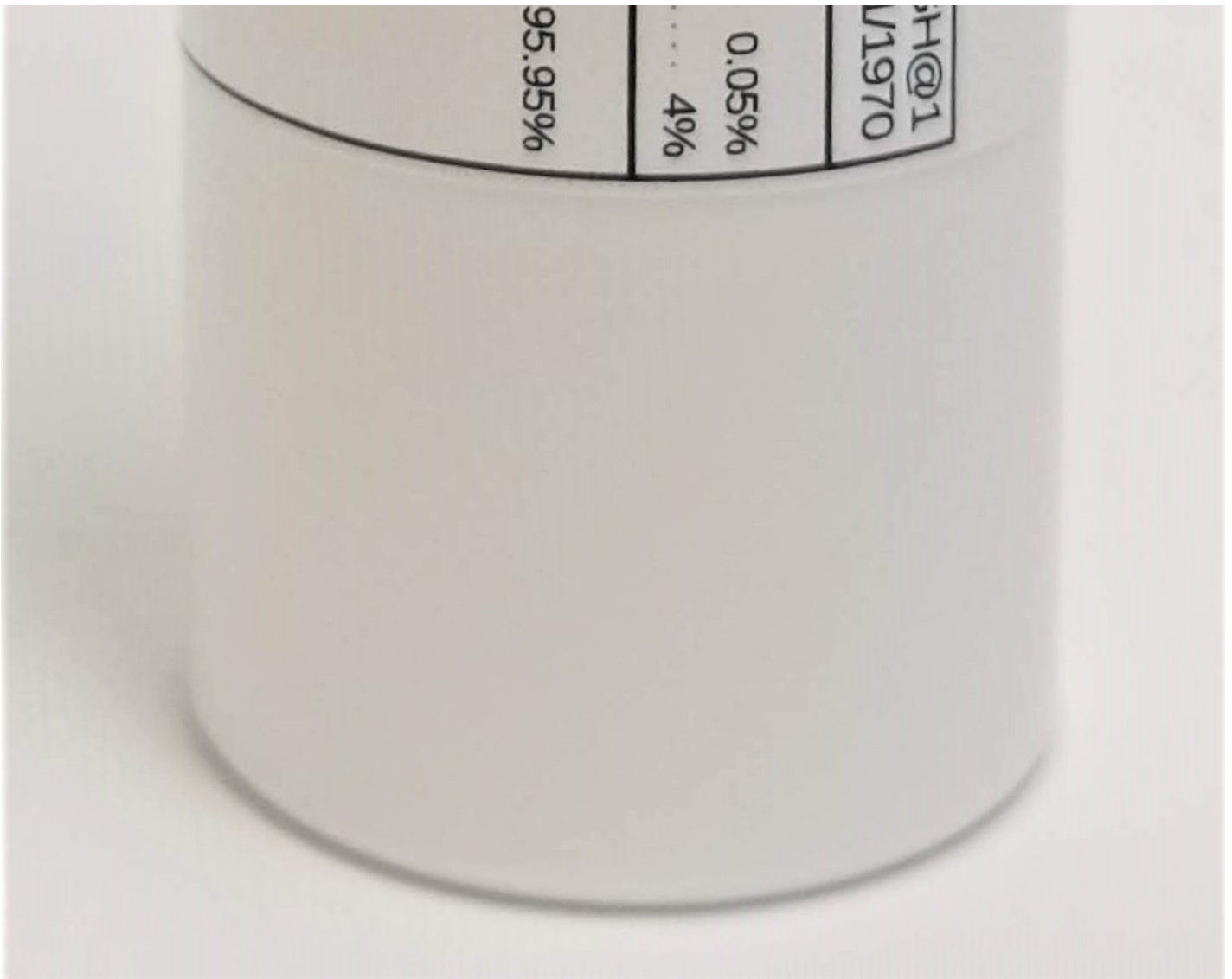
Lot: 091092ABCDEF  
MFG: 01/01

**Active ingredients**

Clobetazol Propionate USP .....  
Niacinamide USP .....

**Inactive ingredients**

Suspendisse Cream .....



**NDC 72934-2059-3 CLOBETASOL PROPIONATE 0.05% / NIACINAMIDE 4%. Cream 60gm**







**NDC 72934-2059-3**

**CLOBETASOL PROPIONATE**

**USP 0.05%**

**NIACINAMIDE USP 4%**

**CREAM 60gm**

**SINCERUS**

**FLORIDA**

This is a compounded drug.  
Made in USA

## CLOBETASOL PROPIONATE 0.05% / NIACINAMIDE 4%

clobetasol propionate 0.05% / niacinamide 4% cream

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-2059
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CLOBETASOL PROPIONATE (UNII: 779619577M) (CLOBETASOL - UNII:ADN79D536H)	CLOBETASOL PROPIONATE	0.05 g in 100 g
NIACINAMIDE (UNII: 25X5118RD4) (NIACINAMIDE - UNII:25X5118RD4)	NIACINAMIDE	4 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:72934-2059-3	60 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(72934-2059)

Revised: 4/2019

Sincerus Florida, LLC