

**NIACINAMIDE 4% / CLOBETASOL PROPIONATE 0.05% - niacinamide 4% / clobetasol propionate 0.05% solution**

**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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**NIACINAMIDE 4% / CLOBETASOL PROPIONATE 0.05%**

**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

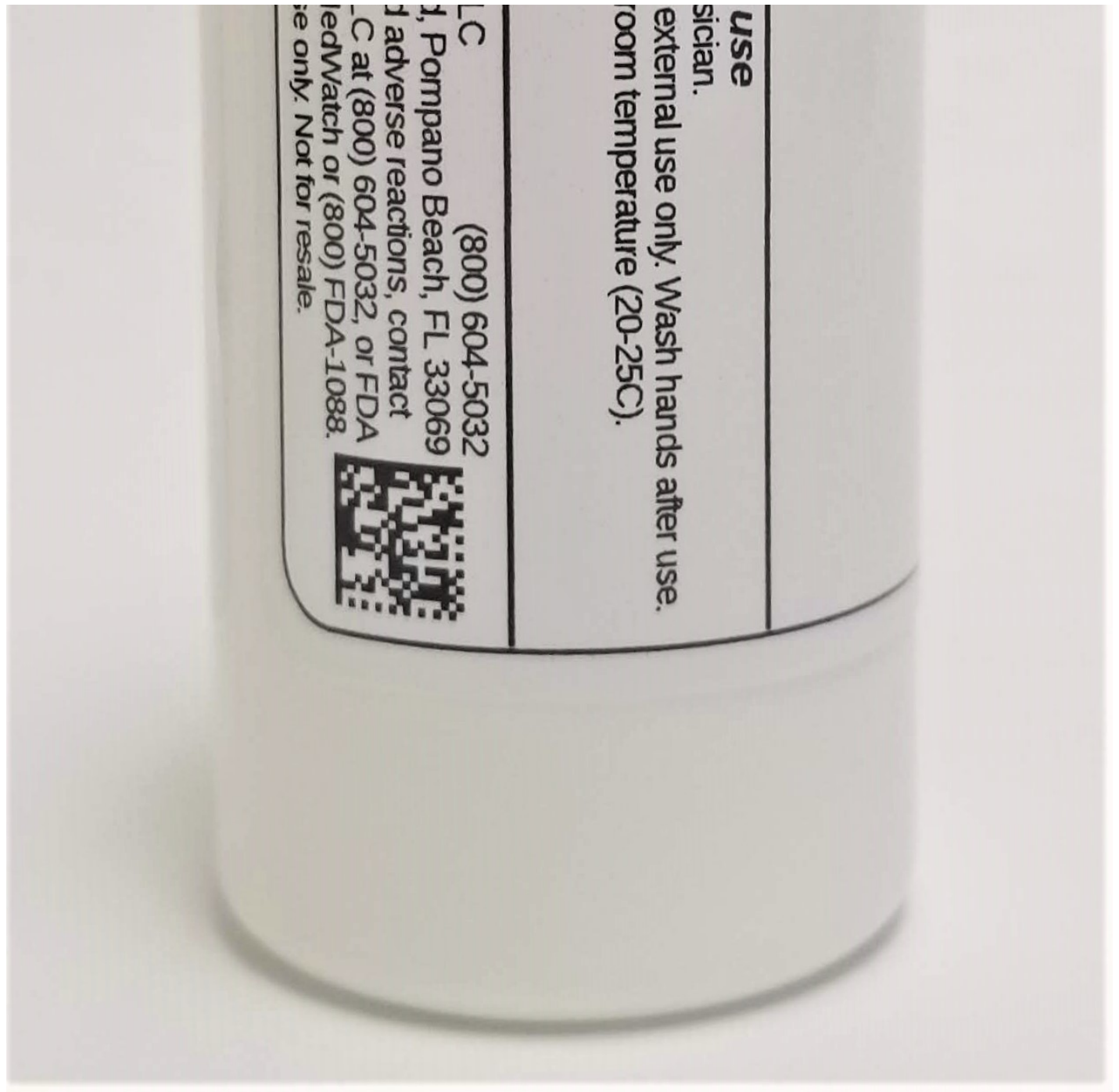
at [www.FDA.gov/MedWatch](http://www.FDA.gov/MedWatch) or (800) FDA-1088.

Office use only. Not for resale.



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Office us



**Active, inactive**





CLOBETASOL  
USP 0.05%  
NIACINAMIDE  
SOLUTION

SINCE

Rx only

BUD: 01/01/1970

Lot: 351033ABC

MFG

**Active ingredients**

Clobetasol Propionate USP .....

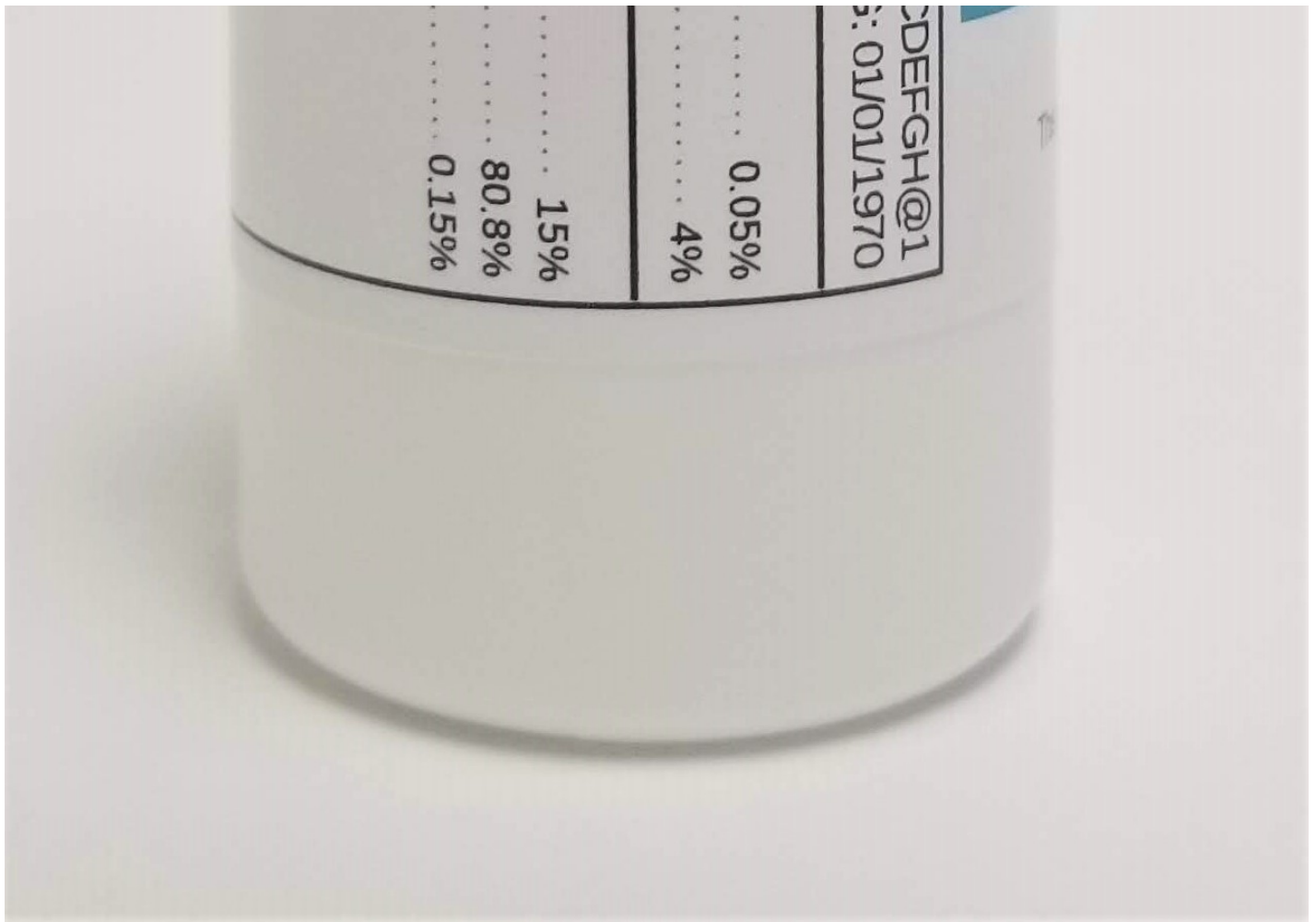
Niacinamide USP .....

**Inactive ingredients**

Ethyl Alcohol USP .....

Purified Water, USP .....

Zinc Pyrithione 50% Dispersion .....



**NDC 72934- 4154-5 NIACINAMIDE USP 4% / CLOBETASOL PROPIONATE USP 0.05%.  
Solution 60 gm.**



Rx only  
BUD: 01/01/1970

Lot 351033AABCDEF@1  
MFG: 01/01/1970

**NDC 72934-4154-5**

**CLOBETASOL PROPIONATE  
USP 0.05%  
NIACINAMIDE USP 4%  
SOLUTION 60gm**



This is a compounded drug.  
Made in USA

## NIACINAMIDE 4% / CLOBETASOL PROPIONATE 0.05%

niacinamide 4% / clobetasol propionate 0.05% solution

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-4154
<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: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	4 g in 100 g

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-4154-5	60 g in 1 CYLINDER; Type 0: Not a Combination Product	05/20/2019	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/20/2019	

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



<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Sincerus Florida, LLC		080105003	manufacture(72934-4154)

Revised: 5/2019

Sincerus Florida, LLC