

**FLUTICASONE PROPIONATE 0.05% / NIACINAMIDE 4% - fluticasone 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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**FLUTICASONE PROPIONATE 0.05% / NIACINAMIDE 4%**

**Directions for use**



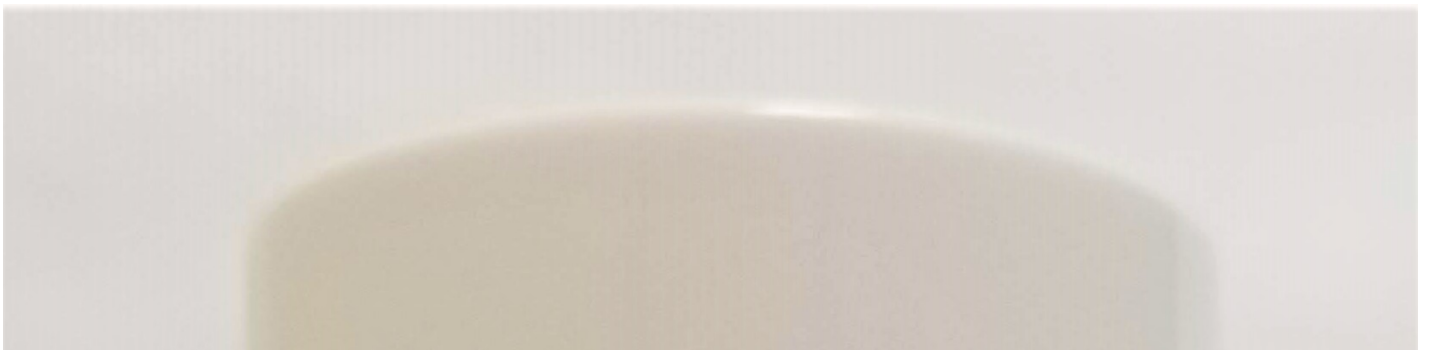
**Directions for use**

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**



**Directions for use**

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Sincerus Florida, LLC (800

3265 W McNab Rd, Pompano Beach

To report suspected adverse reactions

Sincerus Florida, LLC at (800) 604-503

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

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**Active, inactives**





**NDC 72934- 2087-2 FLUTICASONE PROPIONATE USP 0.05% / NIACINAMIDE USP 4%.  
Cream 30gm.**



FLUTICASONONE PROPIONATE  
USP 0.05%  
NIACINAMIDE USP 4%  
CREAM 30gm

SINCERUS

FLORIDA

This is a compounded drug.  
Made in USA

## FLUTICASONONE PROPIONATE 0.05% / NIACINAMIDE 4%

fluticasone propionate 0.05% / niacinamide 4% cream

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2087
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FLUTICASONONE PROPIONATE (UNII: O2GMZ0LF5W) (FLUTICASONONE - UNII:CUT2W21N7U)	FLUTICASONONE PROPIONATE	0.05 g in 100 g

NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)

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:72934-2087-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/14/2019	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/14/2019	

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

**Establishment**

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2087)