

**FLUOCINOLONE ACETONIDE 0.01/ NIACINAMIDE 4- fluocinolone acetoneide 0.01/
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](#).

FLUOCINOLONE ACETONIDE 0.025% / NIACINAMIDE 4%

Diections 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 or (800) FDA-1088.
Office use only. Not for resale.



Sincerus Florida, LLC adverse reactions.



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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 or (800) F

Office use only. Not for resal



Active, inactive



FLUOCINOLONE
USP 0.025%
NIACINAMIDE
CREAM 30g

SINCE

Rx only
BUD: 01/01/1970

Lot: 091025ABCD EFGH@1
MFG: 01/01/1970

Active ingredients

Fluocinolone Acetonide USP 0.025%
Niacinamide USP 4%

Inactive ingredients

Suspendisse Cream 95.975%

NDC 72934-2084-2
FLUOCINOLONE ACETONIDE 0.025% / NIACINAMIDE 4%
Cream 30gm.

Rx only
BUD: 01/01/197

NDC 72934-2084-2

**FLUOCINOLONE ACETONIDE
USP 0.025%
NIACINAMIDE USP 4%
CREAM**



FLUOCINOLONE ACETONIDE 0.01/ NIACINAMIDE 4

fluocinolone acetonide 0.01/ niacinamide 4 cream

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2084
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FLUOCINOLONE ACETONIDE (UNII: 0 CD5FD6S2M) (FLUOCINOLONE ACETONIDE - UNII:0 CD5FD6S2M)	FLUOCINOLONE ACETONIDE	0.025 g in 100 g
NIACINAMIDE (UNII: 25X51I8 RD4) (NIACINAMIDE - UNII:25X51I8 RD4)	NIACINAMIDE	4 g in 100 g

Product Characteristics

Color	white	Score	
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Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-2084-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/13/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/13/2019	

Labeler - Sincerus Florida LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida LLC		080105003	manufacture(72934-2084)

Revised: 5/2019

Sincerus Florida LLC