

CALCIPOTRIENE 0.005% / NIACINAMIDE 4% - calcipotriene 0.005% / 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](#).

CALCIPOTRIENE 0.005% / 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 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



CALCIPOTRIENE
USP 0.005%
NIACINAMIDE
CREAM

SINCE

Rx only
BUD: 01/01/1970

Lot: 161021ABCD EFGH@1
MFG: 01/01/1970

Active ingredients

Calcipotriene Anhydrous USP 0.005%
Niacinamide USP 4%

Inactive ingredients

Suspendisse Cream 95.995%

NDC 72934- 2033-2 CALCIPOTRIENE 0.005% / NIACINAMIDE 4%. Cream 30gm

RX ONLY
1 (813) 812-1111

NDC 72934-2033-2

**CALCIPOTRIENE ANHYDROUS
USP 0.005%
NIACINAMIDE USP 4%
CREAM 30gm**



This is a compounded drug.

Made in USA

CALCIPOTRIENE 0.005% / NIACINAMIDE 4%

calcipotriene 0.005% / niacinamide 4% cream

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2033
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8 RD4) (NIACINAMIDE - UNII:25X51I8 RD4)	NIACINAMIDE	4 g in 100 g
CALCIPO TRIENE (UNII: 143NQ3779B) (CALCIPOTRIENE - UNII:143NQ3779B)	CALCIPOTRIENE	0.005 g in 100 g

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-2033-2	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(72934-2033)

