

NIACINAMIDE 4% / TRETINOIN 0.1% - niacinamide 4% / tretinoin 0.1% 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](#).

NIACINAMIDE 4% / TRETINOIN 0.1%

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



Sincerus Florida, LLC adverse reactions.



Directions for use

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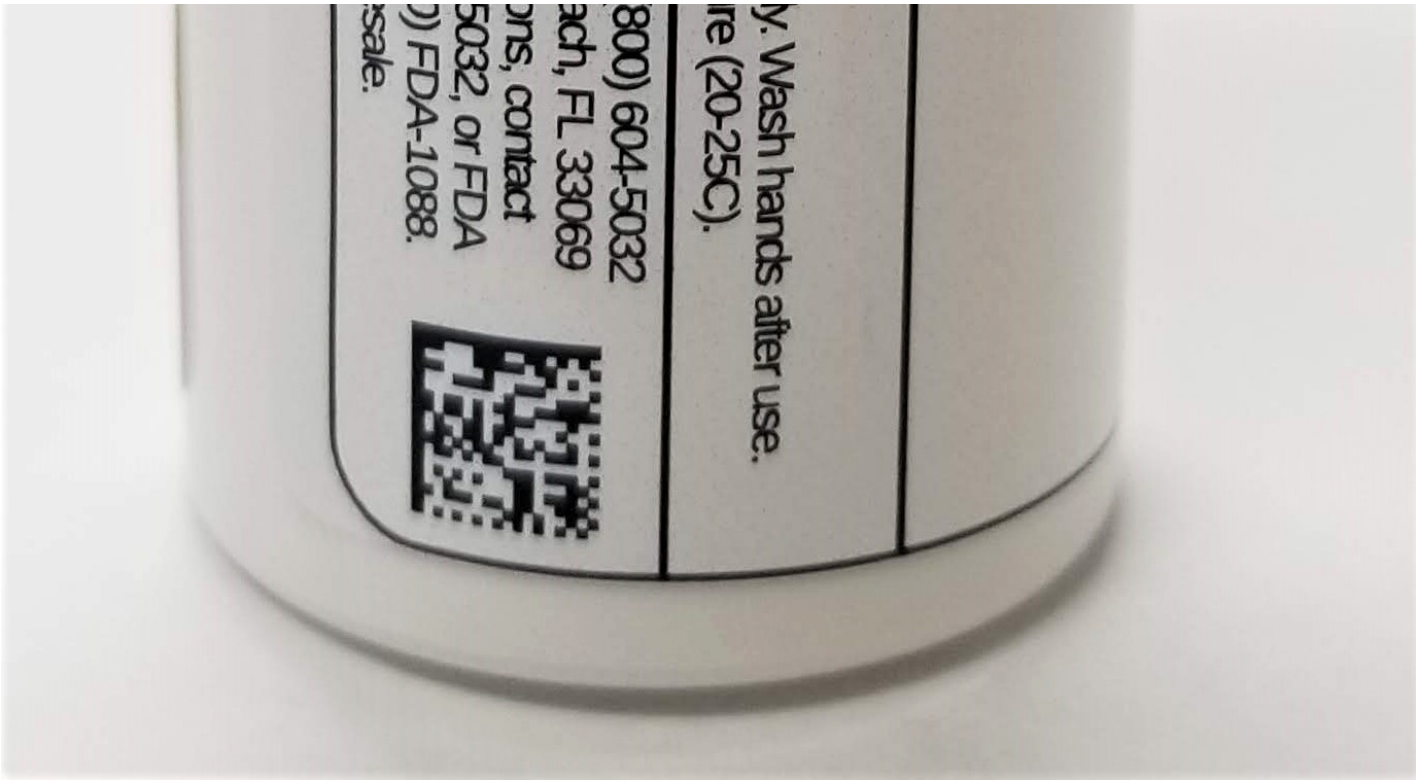
Sincerus Florida, LLC

3265 W McNab Rd, Pompano Beach, FL 33069

To report suspected adverse reactions, contact Sincerus Florida, LLC at (800) 604-4444.

For more information, visit www.FDA.gov/MedWatch or (800) 332-1088.

Office use only. Not for retail sale.



Active, inactive



NDC 72934-2168-2
NIACINAMIDE 4 / TRETINOIN 0.01
Cream 30gm

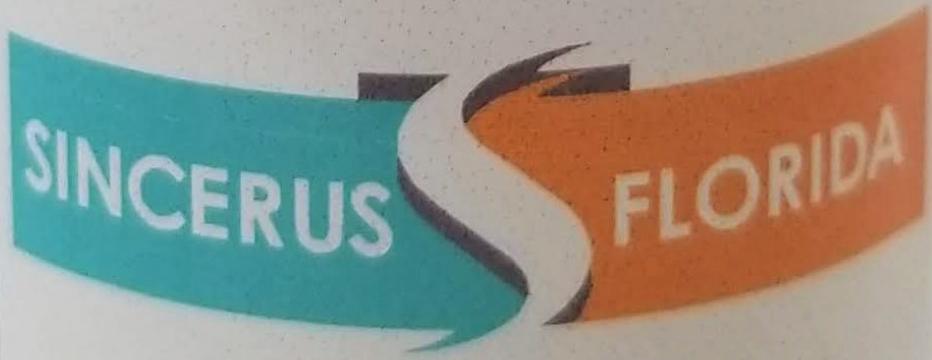


Rx only
BUD: 01/01/1970

Lot: 011031ABCD EFGHIJ
MFG: 01/01/1970

NDC 72934-2168-2

**NIACINAMIDE USP 4%
TRETINOIN USP 0.1%
CREAM 30gm**



This is a compounded drug.
Made in USA

NIACINAMIDE 4% / TRETINOIN 0.1%

niacinamide 4% / tretinoin 0.1% cream

Product Information

Product Type

HUMAN PRESCRIPTION DRUG

Item Code (Source)

NDC:72934-2168

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
TRETINO IN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.1 g in 100 g

Product Characteristics

Color	yellow	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-2168-2	30 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(72934-2168)

Revised: 1/2020

Sincerus Florida, LLC