

NIACINAMIDE 4% / TRETINOIN 0.05% - niacinamide 4% / tretinoin 0.05% 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 0.05%

Directions for use



Directions for use

is 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
265 W McNab Rd, Pompano Beach, FL 33069
To report suspected adverse reactions, contact
Sincerus Florida, LLC at (800) 604-5032, or FDA
www.FDA.gov/MedWatch or (800) FDA-1088.
Office use only. Not for resale.



Sincerus Florida, LLC adverse reactions



Directions for use

As directed by Physician.

Apply topically. For external use only. Wash hands

Store at controlled room temperature (20-25C).

Sincerus Florida, LLC (800) 604-5032

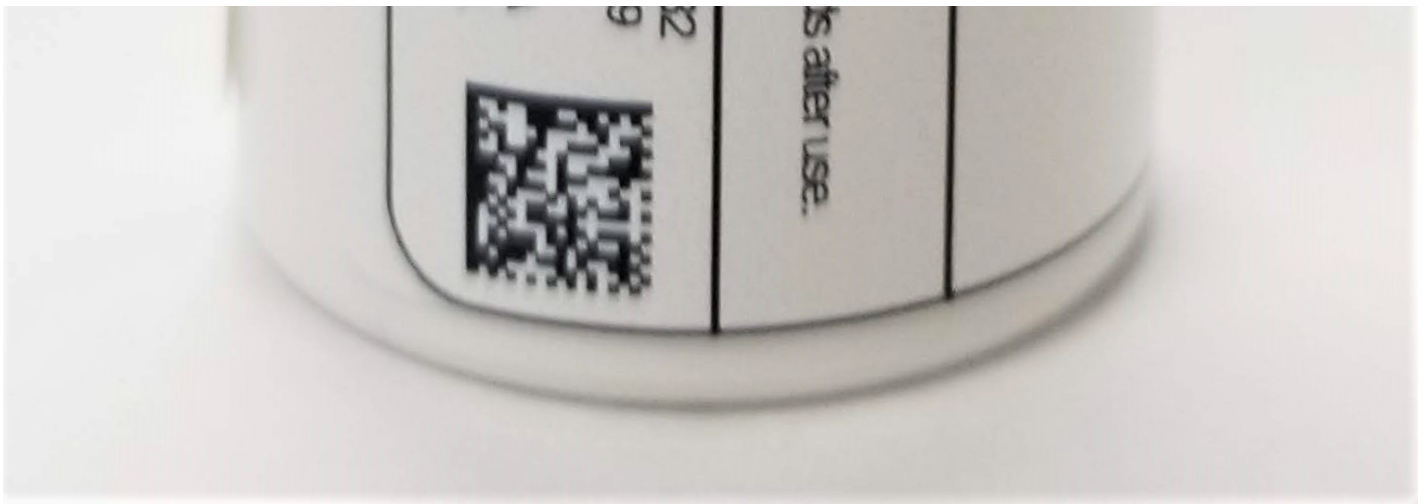
3265 W McNab Rd, Pompano Beach, FL 33068

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.



Active, inactive



NIACINAMIDE
TRETINOIN
CREAM

SINCE

Rx only
BUD: 01/01/1970

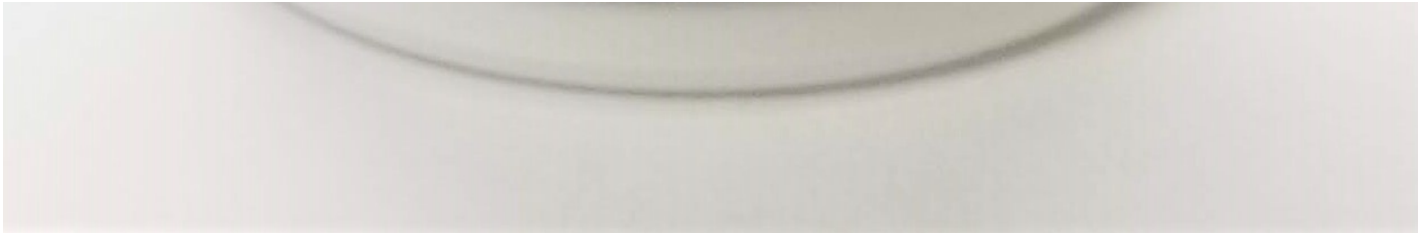
Lot: 0111021ABCDEFGHI@1
MFG: 01/01/1970

Active ingredients

Niacinamide USP 4%
Tretinoin USP 0.05%

Inactive ingredients

Suspendisse Cream 95.95%



NDC 72934-2163-2
NIACINAMIDE 4% / TRETINOIN 0.05%
Cream 30gm





NIACINAMIDE 4% / TRETINOIN 0.05%

niacinamide 4% / tretinoin 0.05% cream

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2163
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
-----------------	-------------------	----------

TRETINO IN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.05 g in 100 g
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	4 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:72934-2163-2	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(72934-2163)

Revised: 4/2019

Sincerus Florida, LLC