

011013 NIACINAMIDE 4% / TRETINOIN 0.025% - 011013 niacinamide 4% / tretinoin 0.025% 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](#).

011013 NIACINAMIDE 4% / TRETINOIN 0.025%

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.

LS-01-000000-00



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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



011013
NIACINAMIDE
TRETINOIN
CREAM 30

SINCE

Rx only
BUD: XXXXXXXXXXXX

Lot: 011013XXXXXXXXXX@XX
MFG: XXXXXXXXXXXX

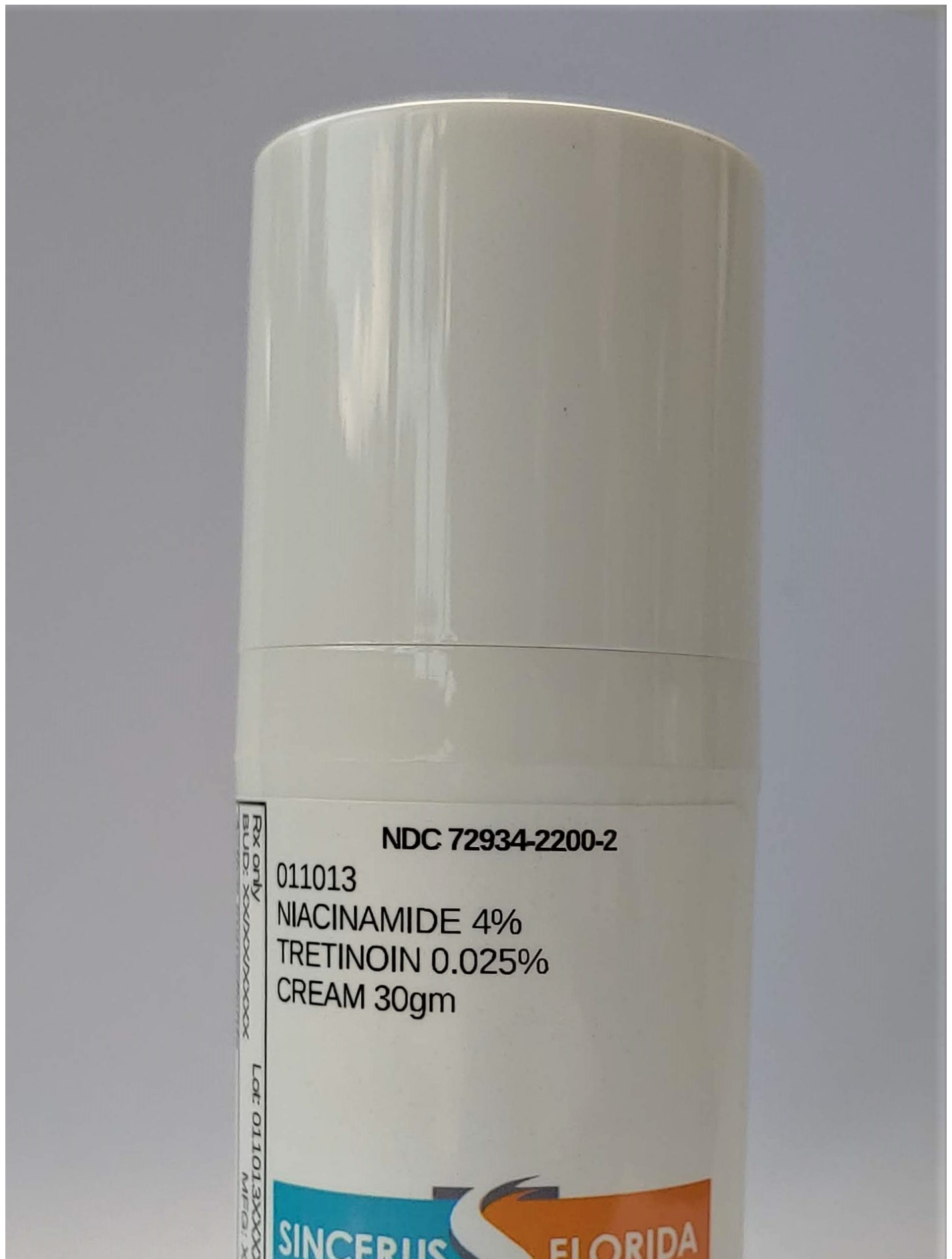
Active ingredients

Niacinamide USP 4%
Tretinoin USP 0.025%

Inactive ingredients

Suspendisse Cream 95.975%

NDC 72934-2200-2 011013 NIACINAMIDE 4% / TRETINOIN 0.025% cream 30 gm

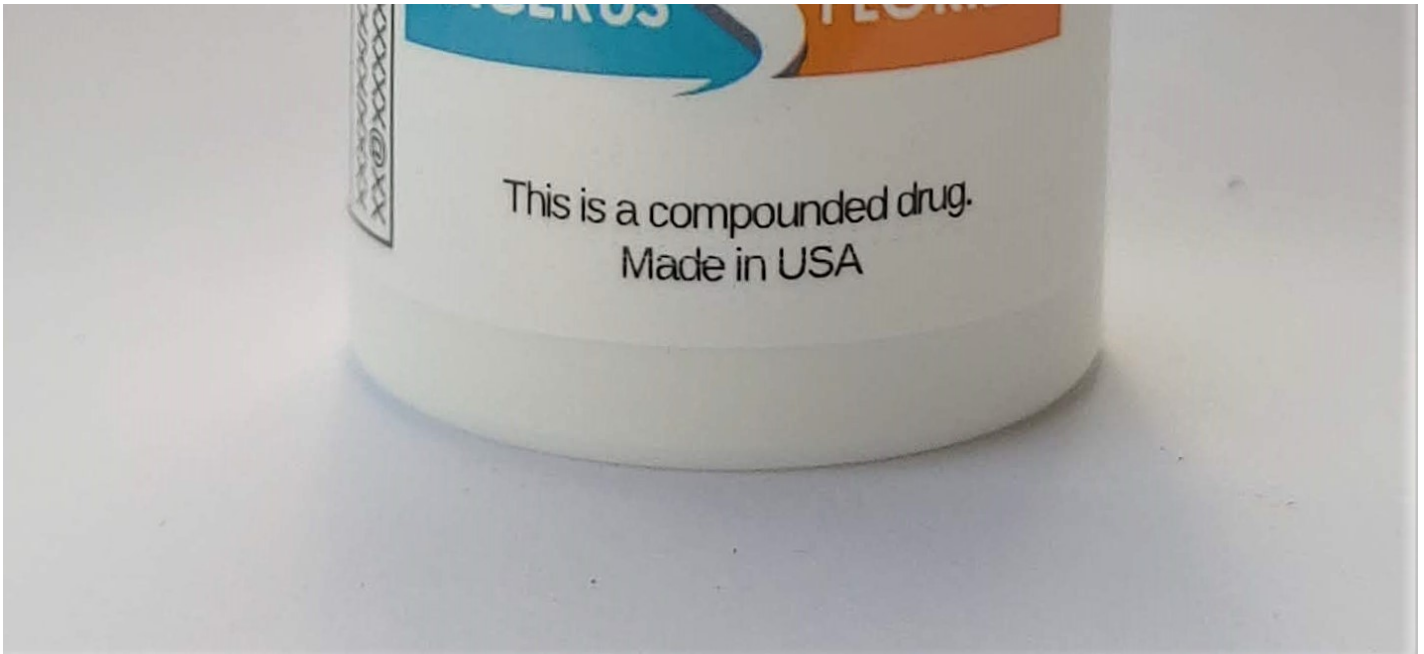


Rx only
NDC: XXXXXXXXX
Lot: 011013XXX
MFG: X

NDC 72934-2200-2

011013
NIACINAMIDE 4%
TRETINOIN 0.025%
CREAM 30gm

SINCERUS FLORIDA



011013 NIACINAMIDE 4% / TRETINOIN 0.025%
 011013 niacinamide 4% / tretinoin 0.025% cream

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2200
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
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.025 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:72934-2200-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/02/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

unapproved drug other		07/02/2020	
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Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2200)

Revised: 7/2020

Sincerus Florida, LLC