

CLINDAMYCIN 1% / NIACINAMIDE 4% / TRETINOIN 0.05% - clindamycin 1% / 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](#).

CLINDAMYCIN 1% / NIACINAMIDE 4% / TRETINOIN 0.05%

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



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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

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Active, inactive



Lot: 011022AGABCBACA@1

MF-G: 06/01/2020

1

1.21%

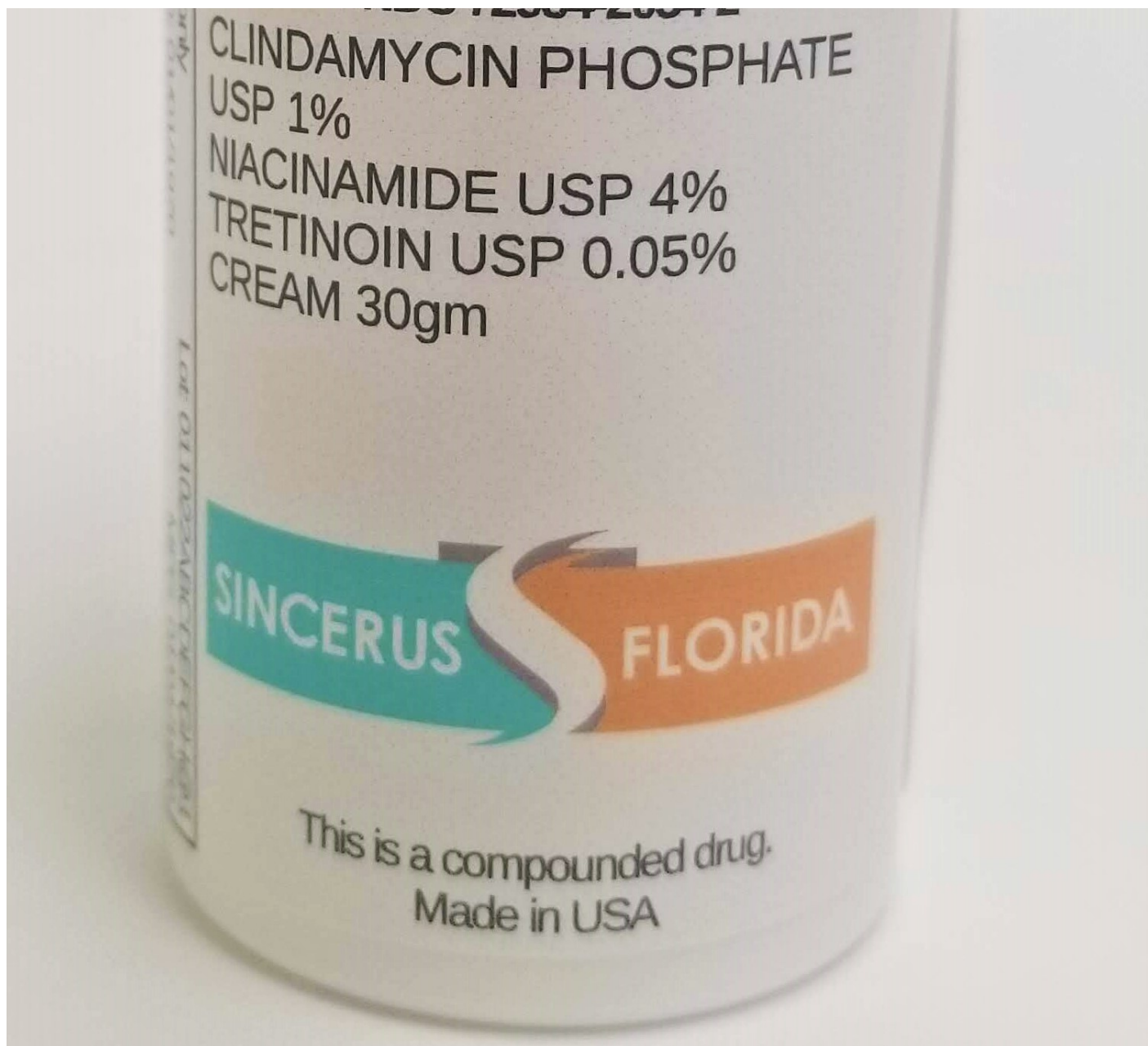
40%

0.05%

94.74%

NDC 72934-2054-2. CLINDAMYCIN 1% / NIACINAMIDE 4% / TRETINOIN 0.05%. Cream 30gm.





CLINDAMYCIN 1% / NIACINAMIDE 4% / TRETINOIN 0.05%

clindamycin 1% / niacinamide 4% / tretinoin 0.05% cream

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2054
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	4 g in 100 g
CLINDAMYCIN PHOSPHATE (UNII: EH6D7113I8) (CLINDAMYCIN - UNII:3U02EL437C)	CLINDAMYCIN PHOSPHATE	1 g in 100 g
ALITRETINOIN (UNII: 1UA8E65KDZ) (ALITRETINOIN - UNII:1UA8E65KDZ)	ALITRETINOIN	0.05 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:72934-2054-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/04/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/04/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment			
Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2054)