

CLINDAMYCIN 1% / NIACINAMIDE 4% / TRETINOIN 0.025% - clindamycin 1% / 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](#).

CLINDAMYCIN 1% / 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.



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

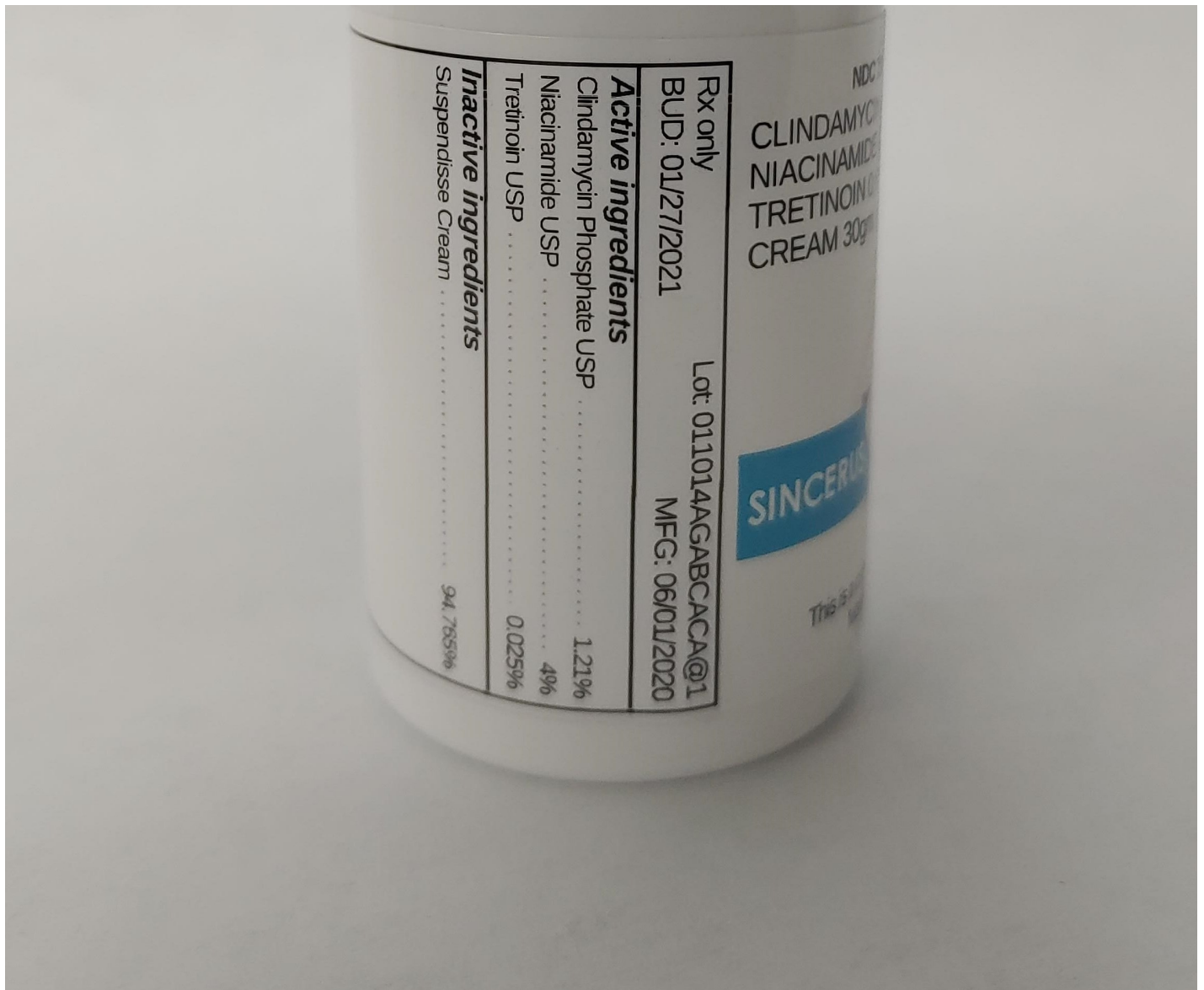
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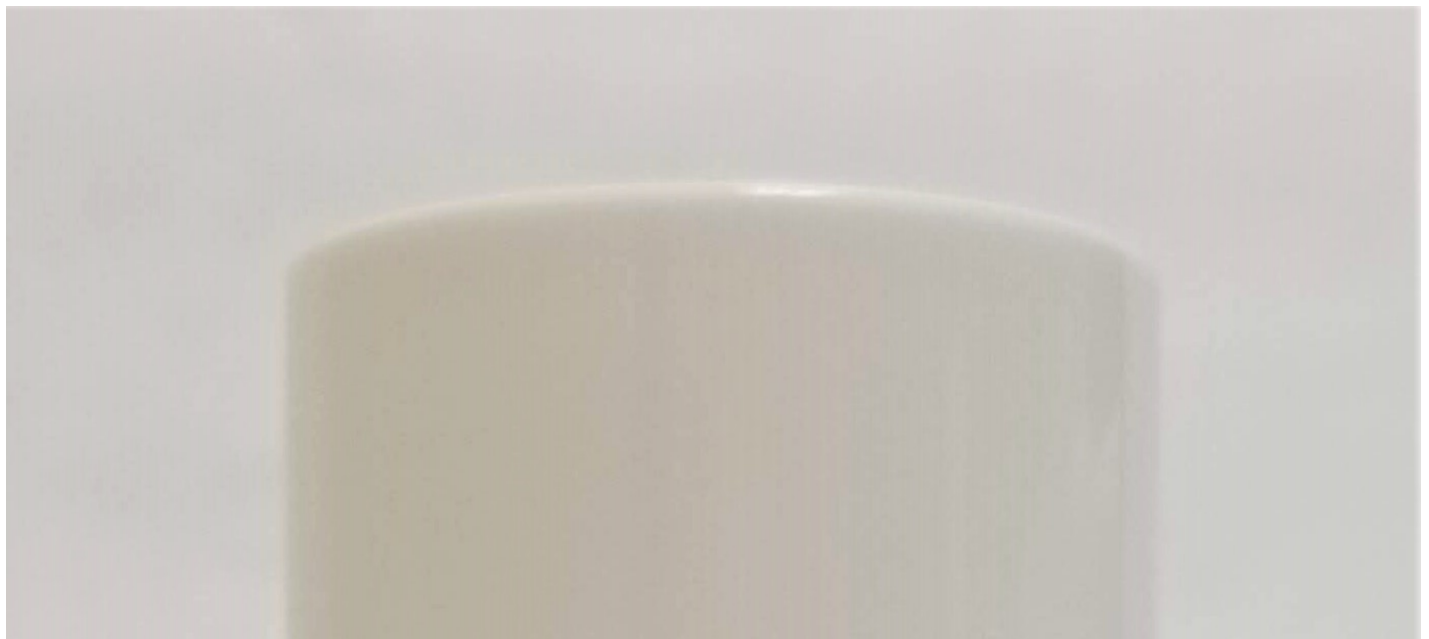


Active/Inactive





NDC 72934-20532-2. CLINDAMYCIN PHOSPHATE USP 1% / NIACINAMIDE USP 4% / TRETINOIN USP 0.025%. Cream 30gm.



NDC 72934-2053-2

CLINDAMYCIN PHOSPHATE
USP 1%

NIACINAMIDE USP 4%
TRETINOIN USP 0.025%
CREAM 30gm

SINCERUS

FLORIDA

This is a compounded drug.
Made in USA

Active Ingredients

Rx only
BUD: 01/01/1970

Lot: 011014ABCDEFGHI@1
MFG: 01/01/1970

CLINDAMYCIN 1% / NIACINAMIDE 4% / TRETINOIN 0.025%

clindamycin 1% / niacinamide 4% / tretinoin 0.025% cream

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2053
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CLINDAMYCIN PHOSPHATE (UNII: EH6D7113I8) (CLINDAMYCIN - UNII:3U02EL437C)	CLINDAMYCIN PHOSPHATE	1 g in 100 g
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.025 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:72934-2053-2	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(72934-2053)

