

CLINDAMYCIN 1% / NIACINAMIDE 4%- clindamycin 1% / niacinamide 4% gel
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%

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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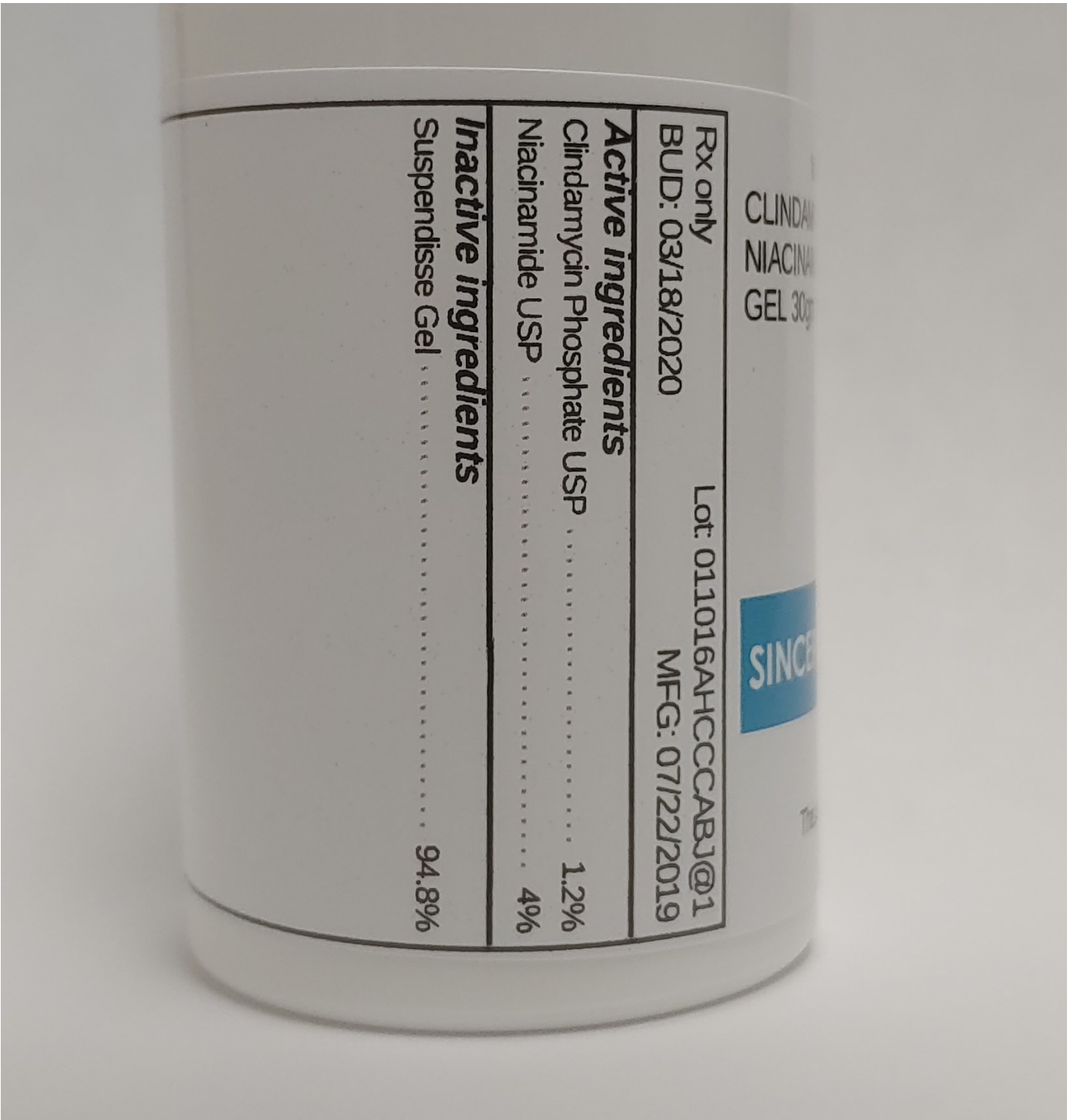
at www.FDA.gov/MedWatch or (800) FD

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Active, inactive





NDC 72934- 1051-2 CLINDAMYCIN PHOSPHATE USP 1% / NIACINAMIDE USP 4%. Gel 30gm



RX ONLY
NDC 72934-1051-2

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CLINDAMYCIN PHOSPHATE

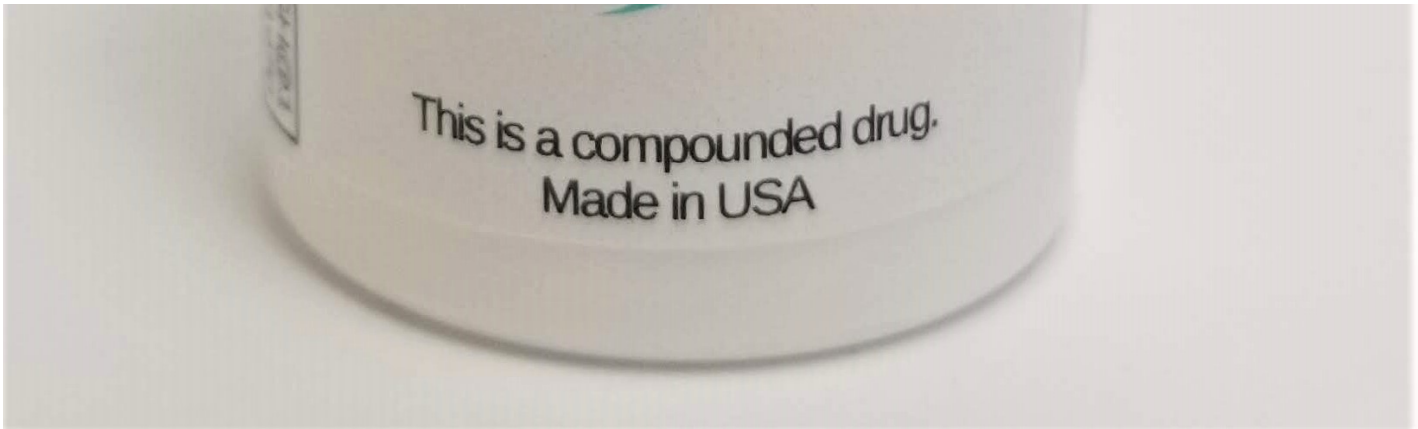
USP 1%

NIACINAMIDE USP 4%

GEL 30gm

SINCERUS

FLORIDA



CLINDAMYCIN 1% / NIACINAMIDE 4%
 clindamycin 1% / niacinamide 4% gel

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-1051
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
CLINDAMYCIN PHOSPHATE (UNII: EH6D7113I8) (CLINDAMYCIN - UNII:3U02EL437C)	CLINDAMYCIN PHOSPHATE	1 g in 100 g

Product Characteristics			
Color	white (Clear gel)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-1051-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/10/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/10/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment			
Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-1051)

Revised: 8/2019

Sincerus Florida, LLC