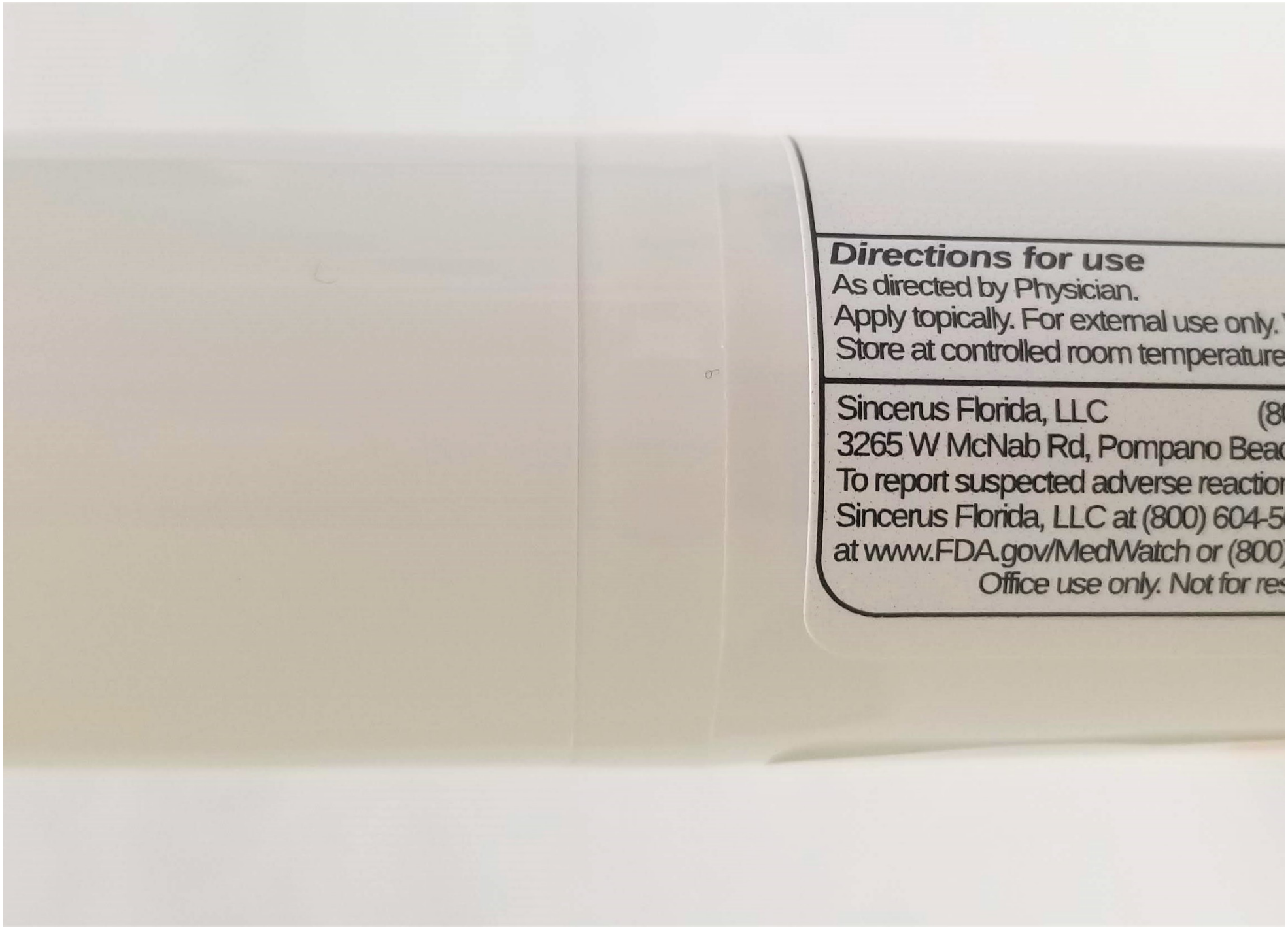
BENZOYL PEROXIDE 2.5% / CLINDAMYCIN 1% / NIACINAMIDE 2% / TRETINOIN 0.025%- benzoyl peroxide 2.5% / clindamycin 1% / niacinamide 2% / tretinoin 0.025% gel
Sincerus Florida, LLC

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.

BENZOYL PEROXIDE 2.5% / CLINDAMYCIN 1% / NIACINAMIDE 2% / TRETINOIN 0.025%

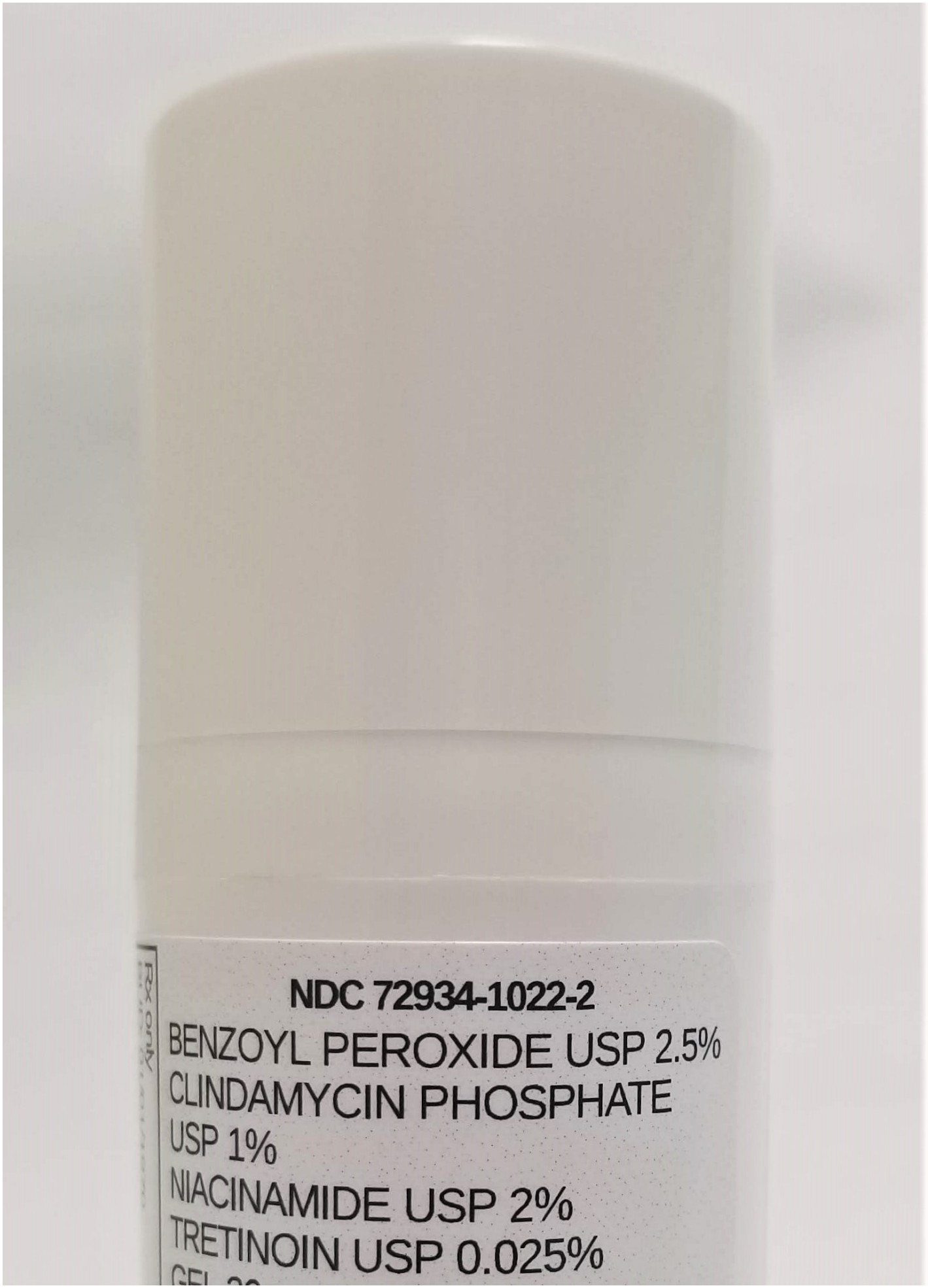
Directions for use

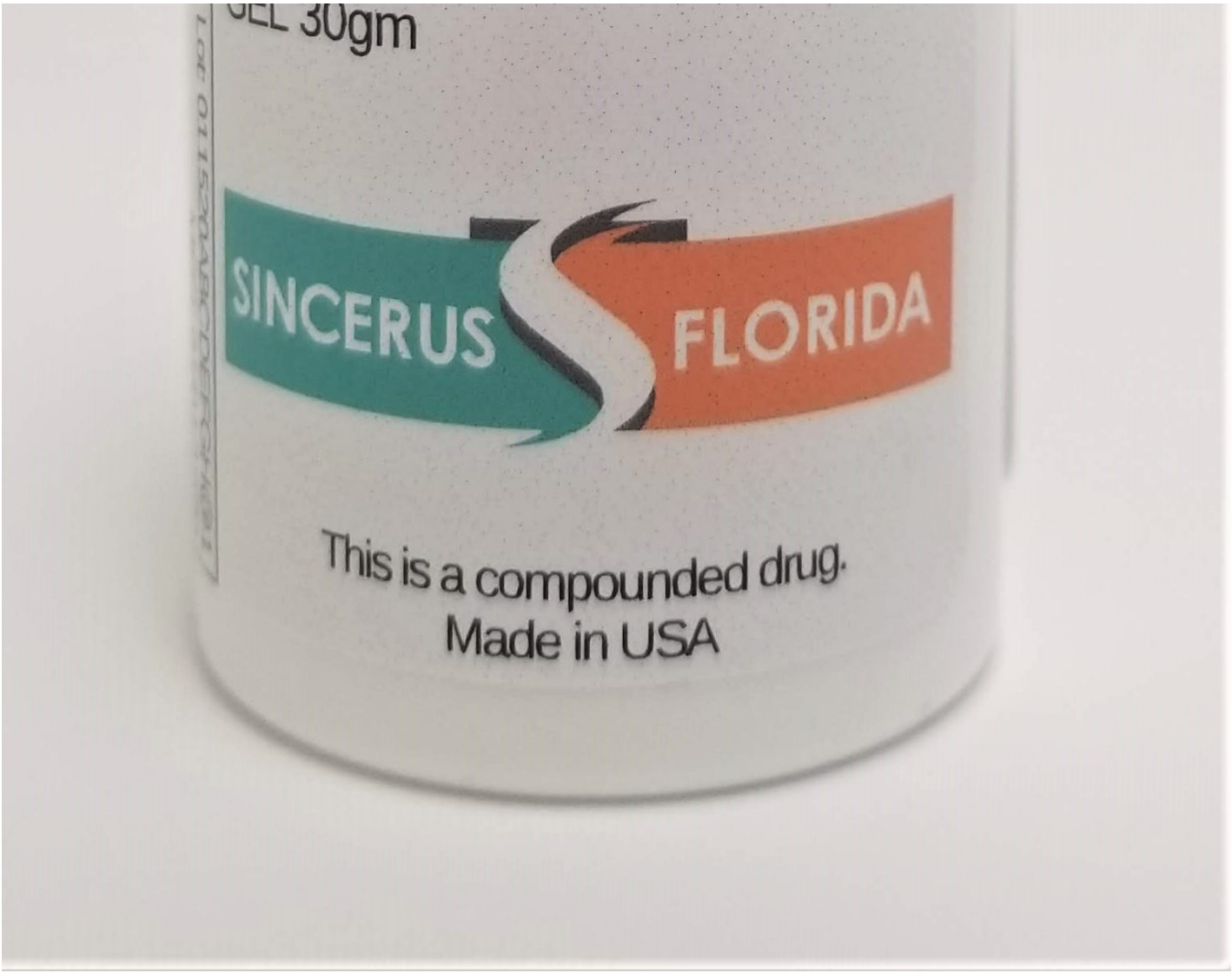
Sincerus Florida, LLC. Adverse reactions


Directions for use
As directed by Physician.
Apply topically. For external use only.
Store at controlled room temperature
Sincerus Florida, LLC
3265 W McNab Rd, Pompano Bea
To report suspected adverse reaction
Sincerus Florida, LLC at (800) 604-5
at www.FDA.gov/MedWatch or (800)
Office use only. Not for re

Active, inactive

NDC 72934- 1022-2 BENZOYL PEROXIDE USP 2.5% / CLINDAMYCIN PHOSPHATE USP 1% / NIACINAMIDE USP 2% / TRETINOIN USP 0.025%. Gel 30 gm


Rx only
NDC 72934-1022-2
BENZOYL PEROXIDE USP 2.5%
CLINDAMYCIN PHOSPHATE
USP 1%
NIACINAMIDE USP 2%
TRETINOIN USP 0.025%



BENZOYL PEROXIDE 2.5% / CLINDAMYCIN 1% / NIACINAMIDE 2% / TRETINOIN 0.025%

benzoyl peroxide 2.5% / clindamycin 1% / niacinamide 2% / tretinoin 0.025% gel

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-1022
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.025 g in 100 g
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	2.5 g in 100 g
CLINDAMYCIN PHOSPHATE (UNII: EH6D7113I8) (CLINDAMYCIN - UNII:3U02EL437C)	CLINDAMYCIN PHOSPHATE	1 g in 100 g
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 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:72934-1022-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/22/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/22/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-1022)

Revised: 5/2019 Sincerus Florida, LLC