

BENZOYL PEROXIDE 5% / CLINDAMYCIN 1% / NIACINAMIDE 2% / TRETINOIN 0.1% - benzoyl peroxide 5% / clindamycin 1% / niacinamide 2% / tretinoin 0.1% 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 5% / CLINDAMYCIN 1% / NIACINAMIDE 2% / TRETINOIN 0.1%

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

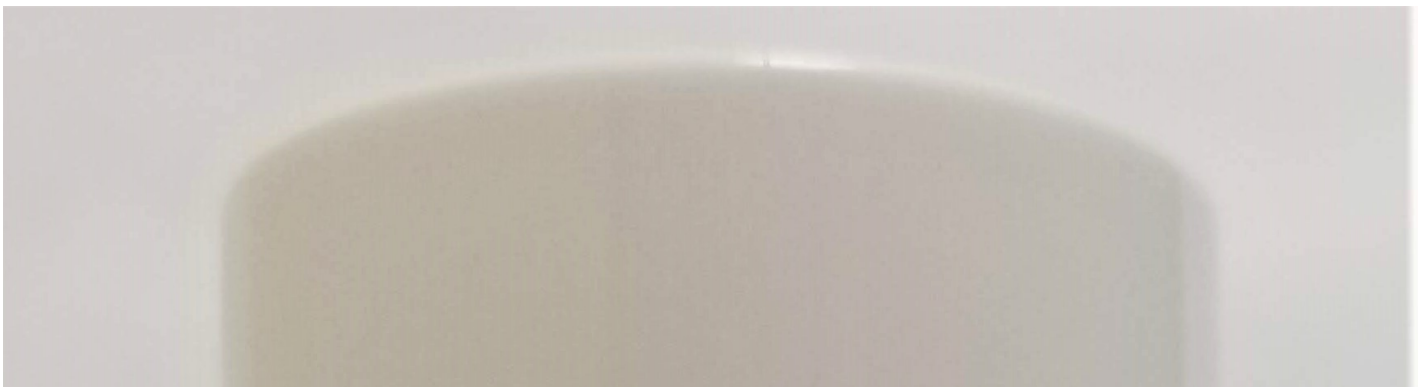
To report suspected adverse reactions, contact

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, Adverse reactions



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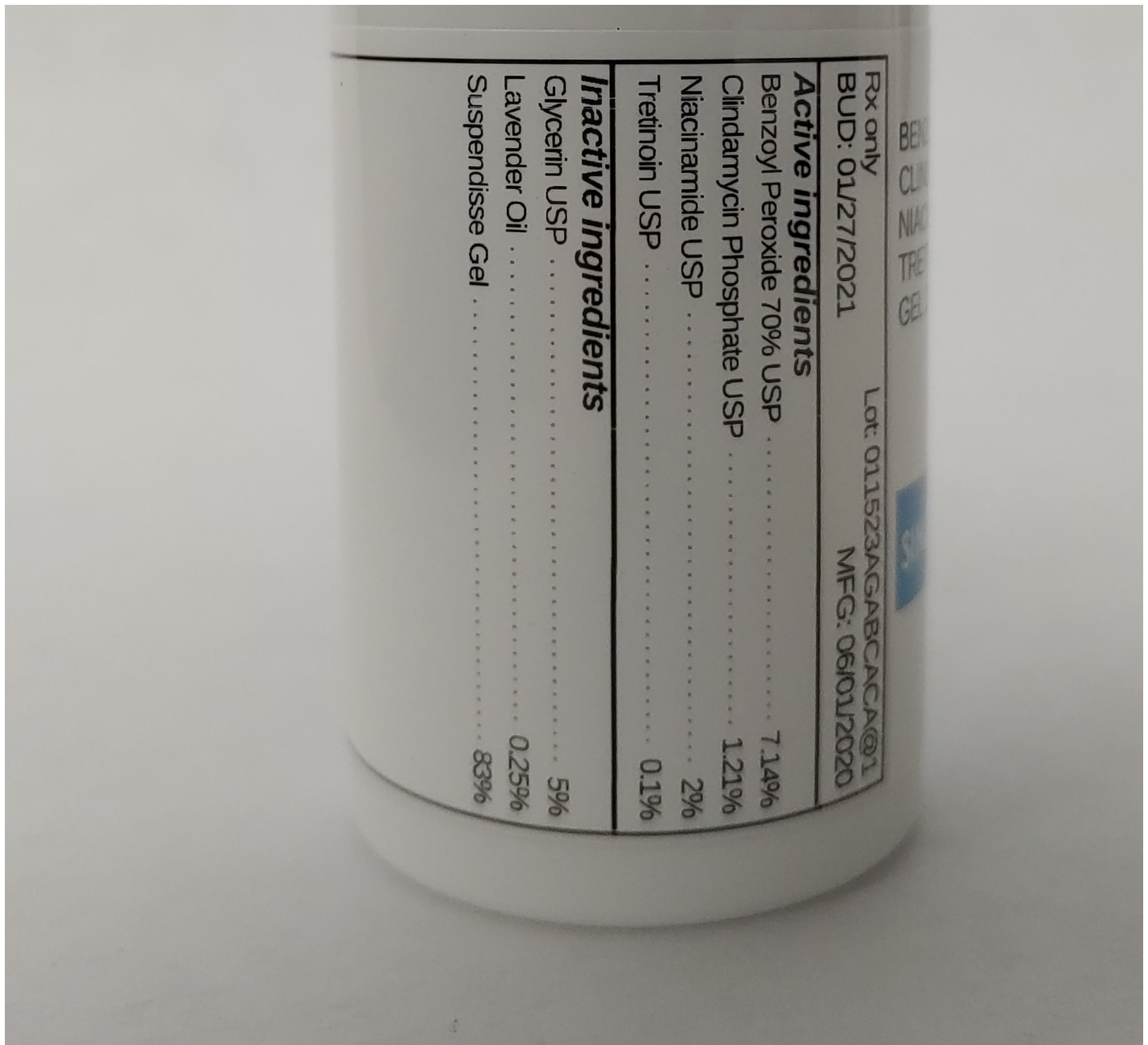
at www.FDA.gov/MedWatch or (800) FDA

Office use only. Not for resale.



Active, inactive





NDC 72934-1018-2 BENZOYL PEROXIDE USP 5% / CLINDAMYCIN USP 1% / NIACINAMIDE 2% / TRETINOIN USP 0.1%. Gel 30gm



This is a compounded drug.
Made in USA

**BENZOYL PEROXIDE 5% / CLINDAMYCIN 1% / NIACINAMIDE 2% /
TRETINOIN 0.1%**

benzoyl peroxide 5% / clindamycin 1% / niacinamide 2% / tretinoin 0.1% gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-1018
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	5 g in 100 g
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 g in 100 g
CLINDAMYCIN PHOSPHATE (UNII: EH6D7113I8) (CLINDAMYCIN - UNII:3U02EL437C)	CLINDAMYCIN PHOSPHATE	1 g in 100 g
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.1 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:72934-1018-2	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(72934-1018)

Revised: 6/2020

Sincerus Florida, LLC