

BENZOYL PEROXIDE 5% / CLINDAMYCIN 1% / NIACINAMIDE 4% - benzoyl peroxide 5% / 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](#).

BENZOYL PEROXIDE 5% / CLINDAMYCIN 1% / NIACINAMIDE 4%

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).

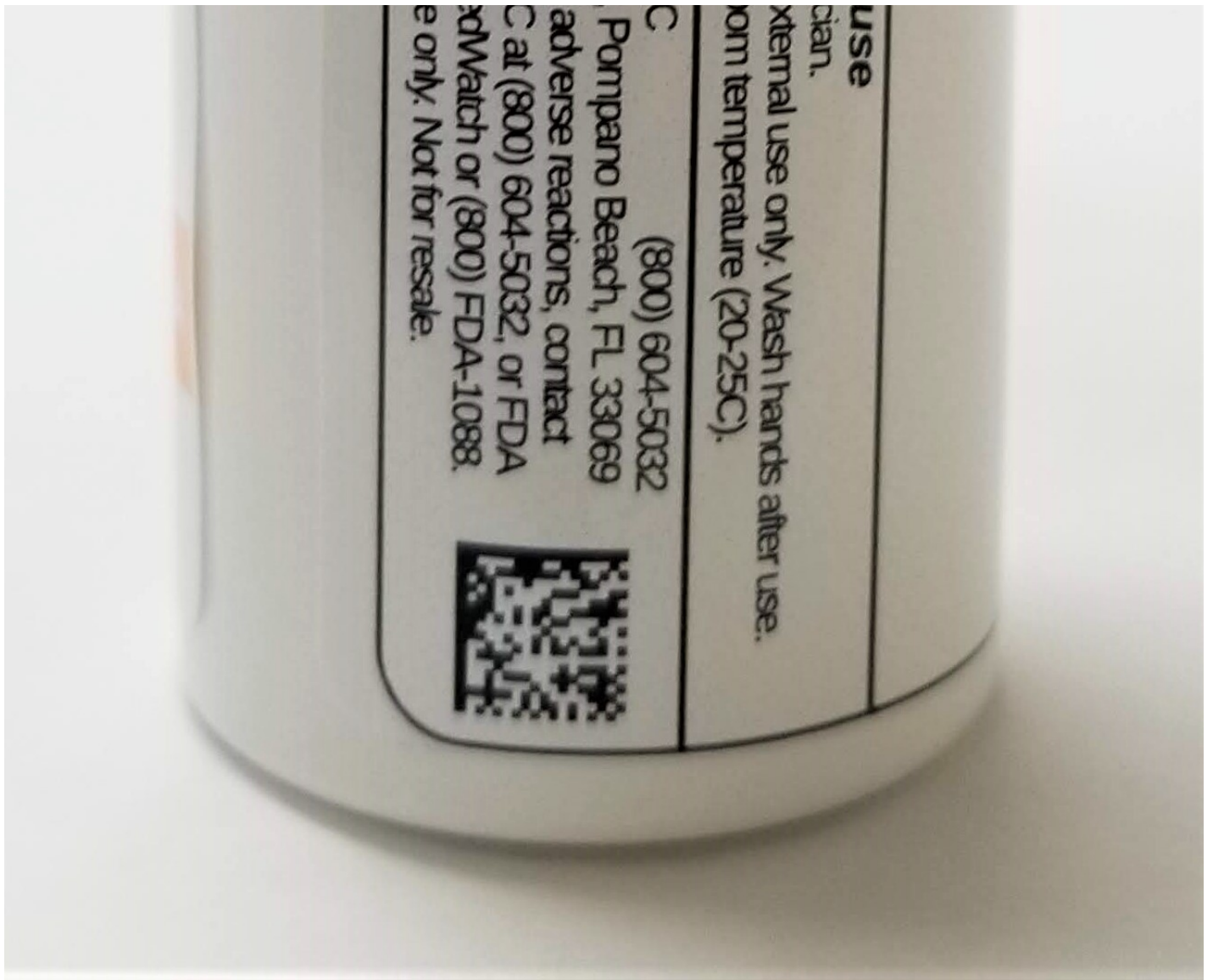
Sincerus Florida, LLC (800) 604-5032
3265 W McNab Rd, Pompano Beach, FL 33069
To report suspected adverse reactions, contact
Sincerus Florida, LLC at (800) 604-5032, or FDA
at www.FDA.gov/MedWatch or (800) FDA-1088.
Office use only. Not for resale.



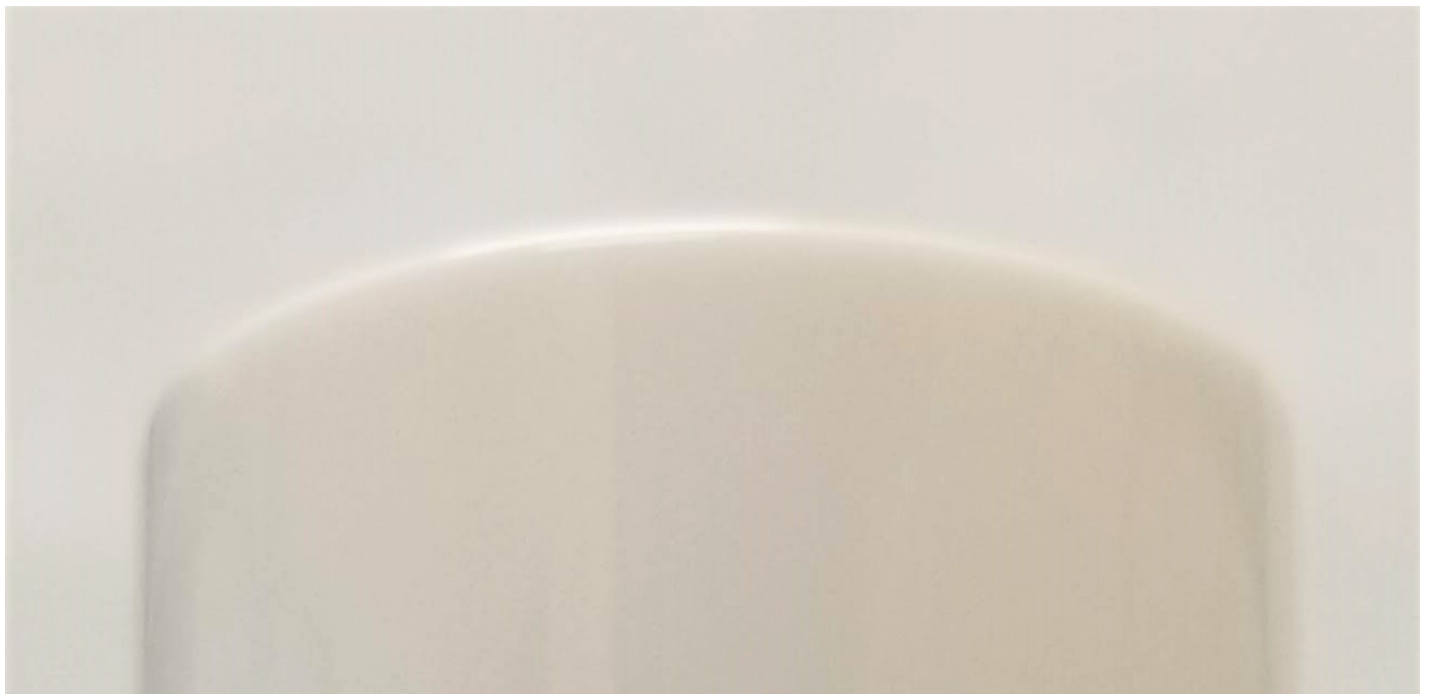
Sincerus Florida, LLC adverse reactions.

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Active, inactive



BENZ
CLIN
USP
NIA
GEL

Rx only

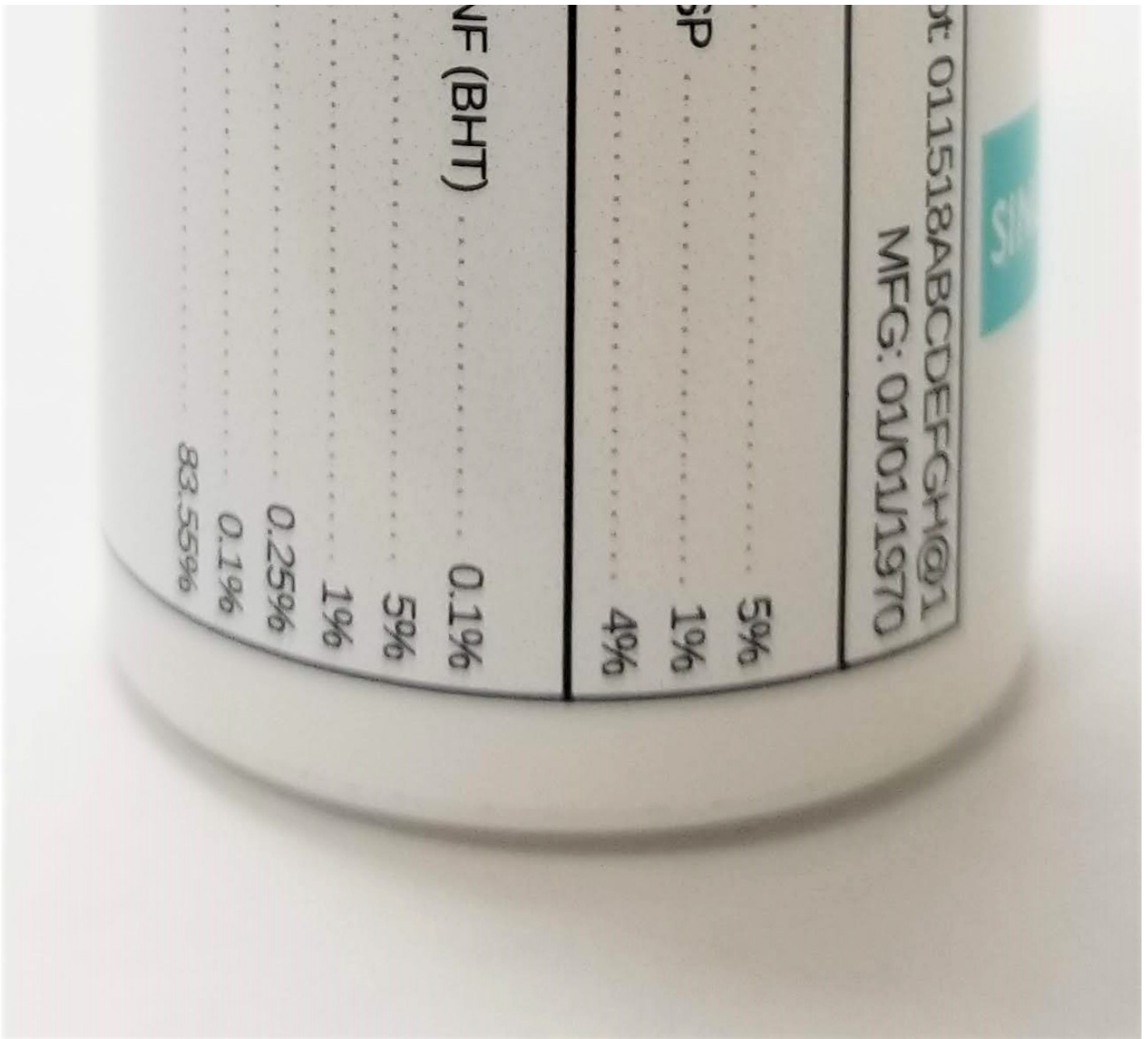
BUD: 01/01/1970

Active ingredients

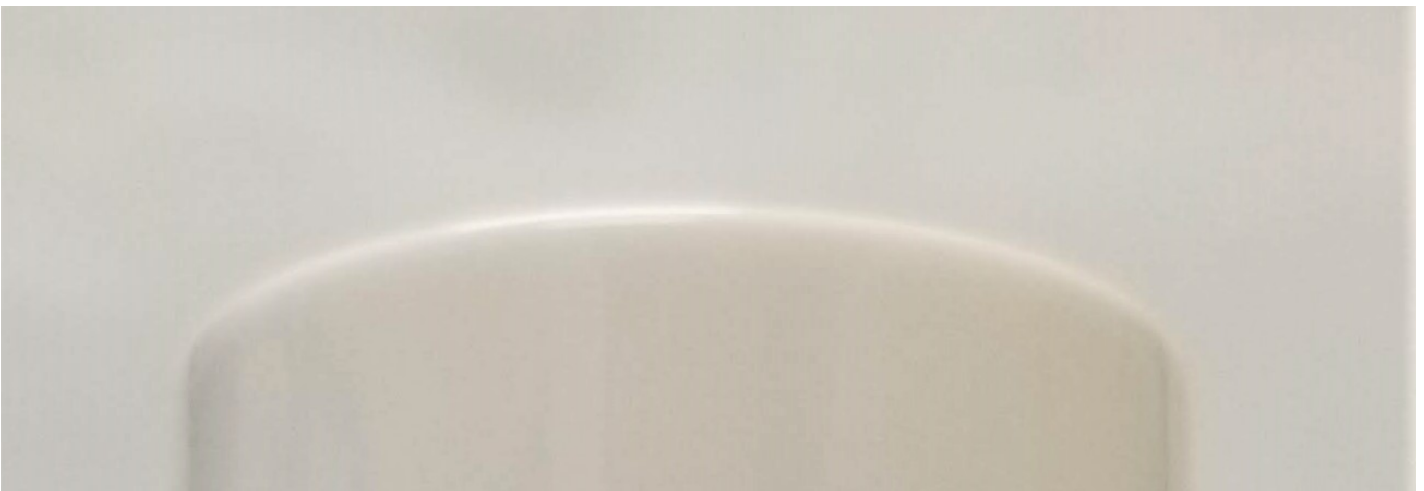
Benzoyl Peroxide USP
Clindamycin Phosphate US
Niacinamide USP

Inactive ingredients

Butylated Hydroxytoluene
Glycerin USP
Krisgel 100
Lavender Oil
Sodium Metabisulfite NF
Suspendisse Gel



NDC 72934-1019-2
ENZOYL PEROXIDE 5 / CLINDAMYCIN PHOSPHATE 1/ NIACINAMIDE 4
gel 30gm

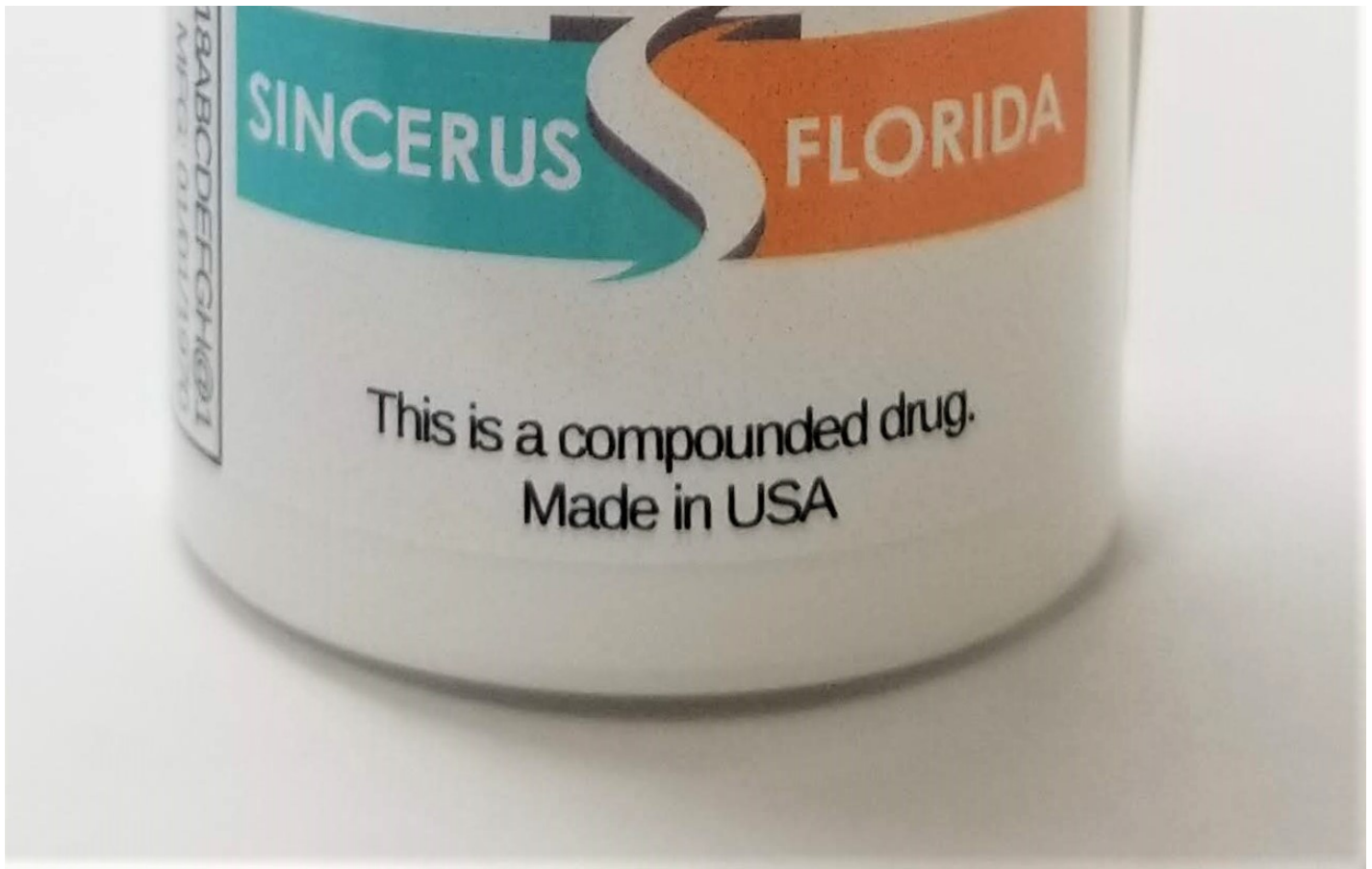


NDC 72934-1019-2

BENZOYL PEROXIDE USP 5%
CLINDAMYCIN PHOSPHATE
USP 1%
NIACINAMIDE USP 4%
GEL 30gm

Rx only
BUD: 01/01/1970

Lot 0115



BENZOYL PEROXIDE 5% / CLINDAMYCIN 1% / NIACINAMIDE 4%

benzoyl peroxide 5% / clindamycin 1% / niacinamide 4% gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-1019
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
CLINDAMYCIN PHOSPHATE (UNII: EH6D7113I8) (CLINDAMYCIN - UNII:3U02EL437C)	CLINDAMYCIN PHOSPHATE	1 g in 100 g
NIACINAMIDE (UNII: 25X5118RD4) (NIACINAMIDE - UNII:25X5118RD4)	NIACINAMIDE	4 g in 100 g

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-1019-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/01/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment			
Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-1019)

Revised: 4/2019

Sincerus Florida, LLC