

ADAPALENE 0.1% / BENZOYL PEROXIDE 2.5% / CLINDAMYCIN 1% - adapalene 0.1% / benzoyl peroxide 2.5% / clindamycin 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](#).

ADAPALENE 0.1% / BENZOYL PEROXIDE 2.5% / CLINDAMYCIN 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, 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

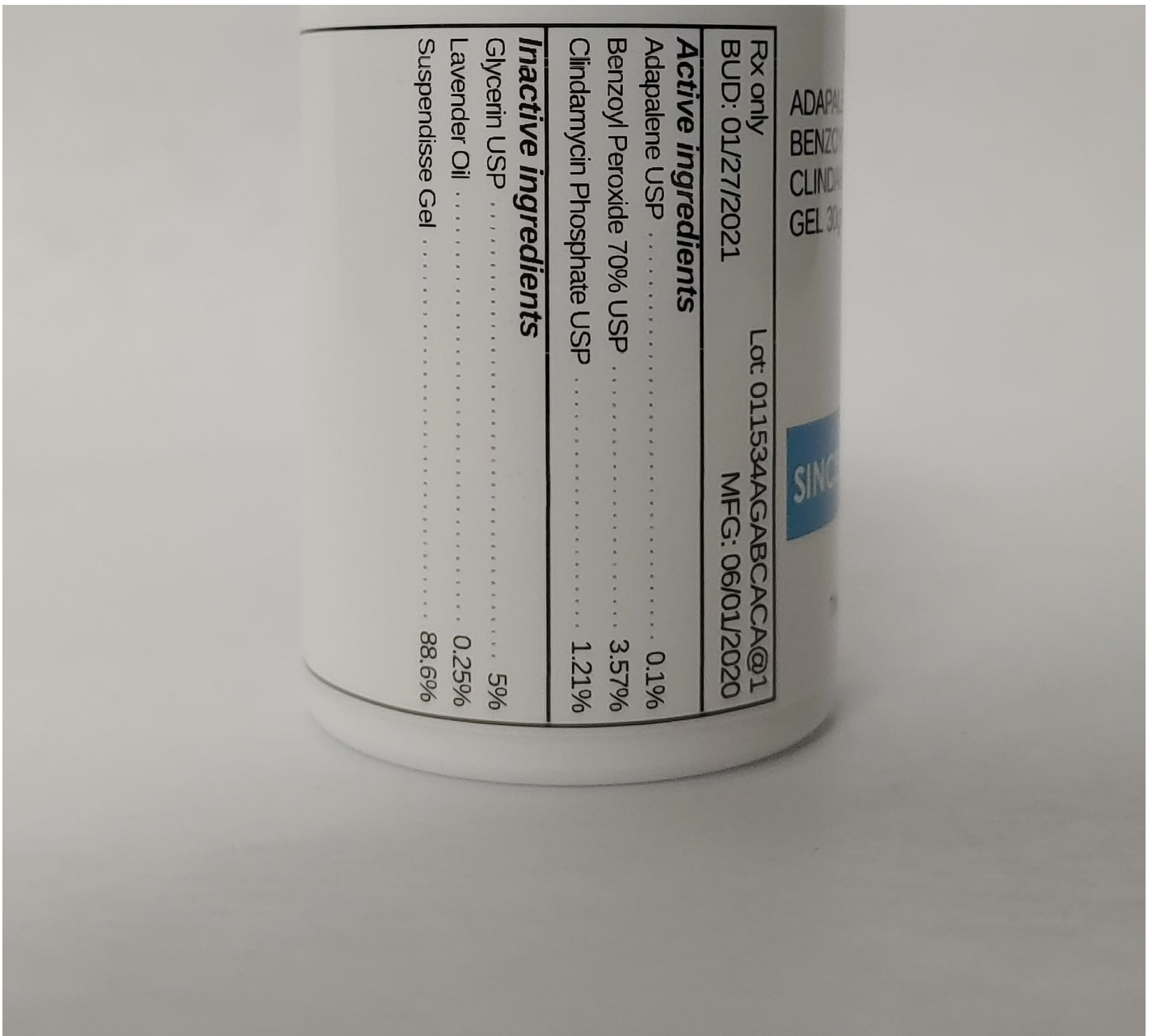
at www.FDA.gov/MedWatch or (800) F

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Active and Inactive





NDC 72934-1002-2 ADAPALENE USP 0.1% / BENZOYL PEROXIDE USP 2.5% / CLINDAMYCIN USP 1%. Gel 30gm



NDC 72934-1002-2

**ADAPALENE USP 0.1%
BENZOYL PEROXIDE USP
2.5%**

**CLINDAMYCIN PHOSPHATE
USP 1%
GEL 30gm**

Rx only
BUD: 01/01/1970

Lot 0111



ADAPALENE 0.1% / BENZOYL PEROXIDE 2.5% / CLINDAMYCIN 1%

adapalene 0.1% / benzoyl peroxide 2.5% / clindamycin 1% gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-1002
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
ADAPALENE (UNII: 1L4806J2QF) (ADAPALENE - UNII:1L4806J2QF)	ADAPALENE	0.1 g in 100 g
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	2.5 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:72934-1002-2	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(72934-1002)

Revised: 6/2020

Sincerus Florida, LLC