CLINDAMYCIN 1% / NIACINAMIDE 4% / TRETINOIN 0.05%- clindamycin 1% / niacinamide 4% / tretinoin 0.05% solution
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% / TRETINOIN 0.05%

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).

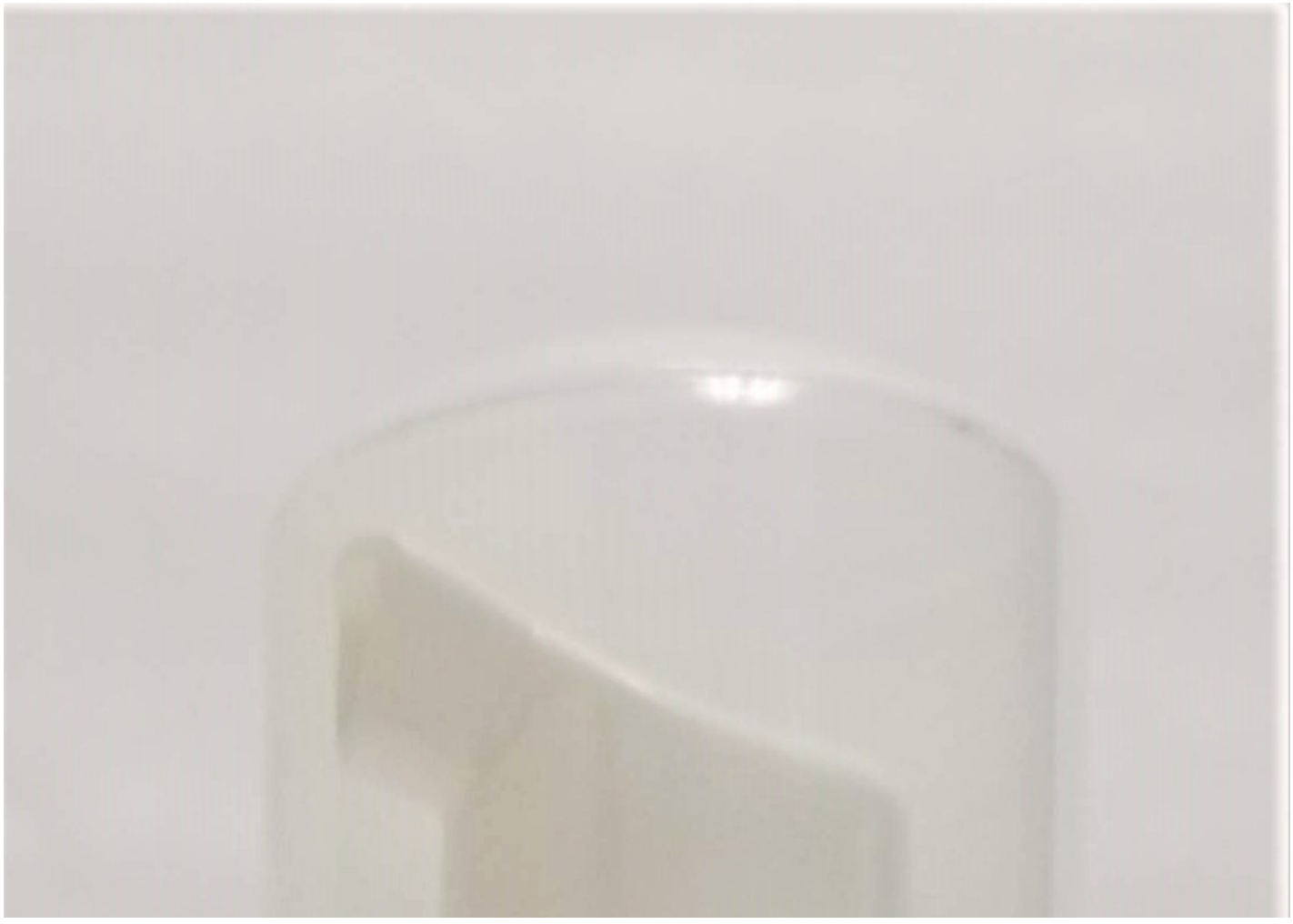
Sincerus Florida, LLC (800) 604-5032

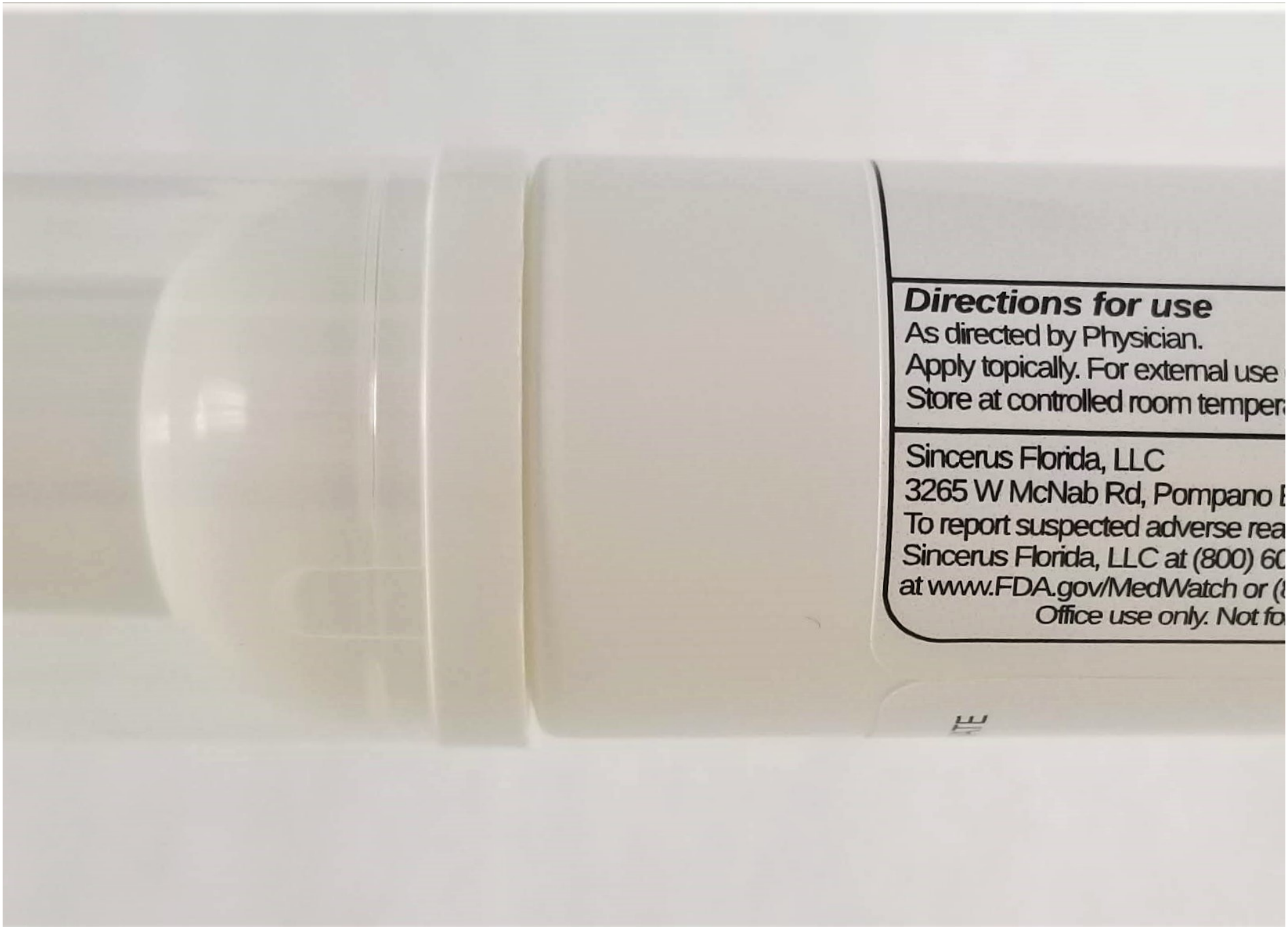
3265 W McNab Rd, Pompano Beach, FL 33069

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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Active, inactive

Rx only
BUD: 01/01/1970

Active ingredien

Clindamycin Phospha

Niacinamide USP ...

Tretinoin USP .....

Inactive ingredie

Butylated Hydroxytolu

Ethyl Alcohol USP ...

Polysorbate 80 NF ...

Preserved Water (para

Propylene Glycol USP

Suspendisse Foam ...

NDC 72934-4054-3 CLINDAMYCIN USP 1% / NIACINAMIDEUSP 4% / TRETINOIN USP 0.05%. Solution 60gm


Rx only
NDC 72934-4054-3
CLINDAMYCIN PHOSPHATE
USP 1%
NIACINAMIDE USP 4%
TRETINOIN USP 0.05%
SOLUTION 60gm

CLINDAMYCIN 1% / NIACINAMIDE 4% / TRETINOIN 0.05%

clindamycin 1% / niacinamide 4% / tretinoin 0.05% solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-4054
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
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	4 g in 100 g
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.05 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:72934-4054-3	60 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/07/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/07/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-4054)

Revised: 5/2019 Sincerus Florida, LLC