

CLINDAMYCIN 1% / HYDROCORTISONE 2.5% / NIACINAMIDE 4% - clindamycin 1% / hydrocortisone 2.5% / niacinamide 4% cream

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% / HYDROCORTISONE 2.5% / 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

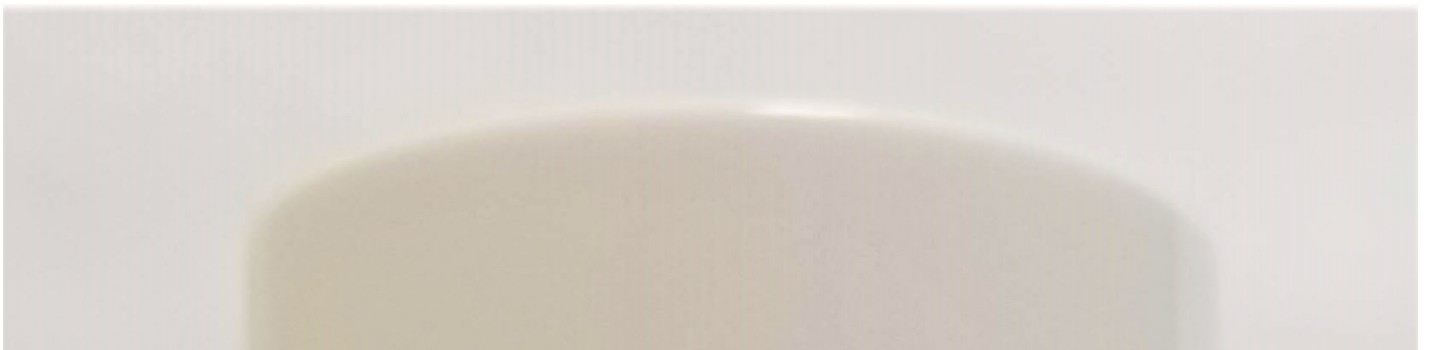
To report suspected adverse reactions, contact

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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Sincerus Florida, LLC (800

3265 W McNab Rd, Pompano Beach

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Sincerus Florida, LLC at (800) 604-503

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

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Active, inactive



CLINDAMYCIN
USP
HYDROCORTISONE
NIACINAMIDE
CREAM



Rx only

BUD: 01/01/1970

Lot: 011026ABCDEF@1

MFG: 01/01/1970

Active ingredients

Clindamycin Phosphate USP	1%
Hydrocortisone USP	2.5%
Niacinamide USP	4%

Inactive ingredients

Glycerin USP	5%
Suspendisse Cream	87.5%

**NDC 72934-2048-2 CLINDAMYCIN PHOSPHATE USP 1% / HYDROCORTISONE USP 2.5%
/ NIACINAMIDE USP 4%. Cream 30gm**





CLINDAMYCIN 1% / HYDROCORTISONE 2.5% / NIACINAMIDE 4%

clindamycin 1% / hydrocortisone 2.5% / niacinamide 4% cream

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2048
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)		HYDROCORTISONE	2.5 g in 100 g
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)		NIACINAMIDE	4 g in 100 g
CLINDAMYCIN PHOSPHATE (UNII: EH6D7113I8) (CLINDAMYCIN - UNII:3U02EL437C)		CLINDAMYCIN PHOSPHATE	1 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:72934-2048-2	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(72934-2048)

Revised: 5/2019

Sincerus Florida, LLC