BETAMETHASONE DIPROPIONATE 0.05% / NIACINAMIDE 4% - betamethas one dipropionate 0.05% / 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.

BETAMETHASONE DIPROPIONATE 0.05% / NIACINAMIDE 4%

Directions for use. As directed by Physician. Apply topically. For external use only. Wash hands after use. Store at controlled room temperature 20-25 Celsius.





Sincerus Florida, LLC. Adverse reactions

Directions for use

As directed by Physician.

Apply topically. For external use only.

Store at controlled room temperature

Sincerus Florida, LLC (8)
3265 W McNab Rd, Pompano Bear
To report suspected adverse reaction
Sincerus Florida, LLC at (800) 604-5
at www.FDA.gov/MedWatch or (800)
Office use only. Not for res



Active, inactive



BETA DIPA NIAC CRE

Rx only BUD: 01/01/1970 Active ingredients Suspendisse Cream Krisgel 100 Inactive ingredients Betamethasone Dipropionate USP 0.05% Niacinamide USP Lot: 091048ABCDEFGH@1 MFG: 01/01/1970 94.95% 1% 4%

NDC 72934- 2026-2 BETAMETHASONE DIPROPIONATE USP 0.05% / NIACINAMIDE USP 4% Cream 30 gm.

NDC 72934-2026-2 BETAMETHASONE DIPROPIONATE USP 0.05% NIACINAMIDE USP 4% CREAM 30gm



This is a compounded drug. Made in USA

BETAMETHASONE DIPROPIONATE 0.05% / NIACINAMIDE 4%

betamethasone dipropionate 0.05% / niacinamide 4% cream

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2026

Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BETAMETHASONE DIPROPIONATE (UNII: 826 Y60 90 1U) (BETAMETHASONE - UNII: 9842 X06 Q6 M)	BETAMETHASONE	0.05 g in 100 g	
NIACINAMIDE (UNII: 25X5118 RD4) (NIACINAMIDE - UNII:25X5118 RD4)	NIACINAMIDE	4 g in 100 g	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:72934-2026-	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/20/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment				
Name	Address	ID/FEI	Business Operations	
Sincerus Florida, LLC		080105003	manufacture(72934-2026)	

Revised: 5/2019 Sincerus Florida, LLC