

DESOXIMETASONE 0.05% / NIACINAMIDE 4% - desoximetasone 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](#).

DESOXIMETASONE 0.05% / NIACINAMIDE 4%

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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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

Office use only. Not for resal



Active, inactive



DESOXIMETASONE
NIACINAMIDE
CREAM

SINCE

Rx only
BUD: 01/01/1970

Lot: 351016ABCDEFGH@1
MFG: 01/01/1970

Active ingredients

Desoximetasone USP 0.05%
Niacinamide USP 4%

Inactive ingredients

Suspendisse Cream 95.95%

NDC 72934- 2068-2 DESOXIMETASONE USP 0.05% / NIACINAMIDE USP 4%. Cream 30gm

RX ONLY
NDC 72934-2068-2

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DESOXIMETASONE USP 0.05%
NIACINAMIDE USP 4%
CREAM 30gm



DESOXIMETASONE 0.05% / NIACINAMIDE 4%

desoximetasone 0.05% / niacinamide 4% cream

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2068
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8 RD4) (NIACINAMIDE - UNII:25X51I8 RD4)	NIACINAMIDE	4 g in 100 g
DESOXIMETASONE (UNII: 4E07GXB7AU) (DESOXIMETASONE - UNII:4E07GXB7AU)	DESOXIMETASONE	0.05 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-2068-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/15/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/15/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2068)

Revised: 5/2019

Sincerus Florida, LLC