

NIACINAMIDE 4% / TAZAROTENE 0.1% - niacinamide 4% / tazarotene 0.1% 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](#).

NIACINAMIDE 4% / TAZAROTENE 0.1%

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



NIACINAMIDE
TAZAROTENE
GEL 30g

SING

Rx only

BUD: 01/01/1970

Lot: 011311ABCD EFGH@1

MFG: 01/01/1970

Active ingredients

Niacinamide USP 4%

Tazarotene 0.1%

Inactive ingredients

Suspendisse Gel 95.9%

NDC 72934-1161-2 NIACINAMIDE USP 4% / TAZAROTENE 0.1%. Gel 30gm

Rx only
BUD: 01/01/1970
Lot 011

NDC 72934-1161-2
NIACINAMIDE USP 4%
TAZAROTENE 0.1%
GEL 30gm



NIACINAMIDE 4% / TAZAROTENE 0.1%				
niacinamide 4% / tazarotene 0.1% gel				
Product Information				
Product Type	HUMAN PRESCRIPTION DRUG		Item Code (Source)	NDC:72934-1161
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	
	TAZAROTENE (UNII: 8 1BDR9 Y8 PS) (TAZAROTENE - UNII:8 1BDR9 Y8 PS)	TAZAROTENE	0.1 g in 100 g	
Product Characteristics				
Color	white (clear gel)	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-1161-2	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(72934-1161)

Revised: 5/2019

Sincerus Florida, LLC