

**NIACINAMIDE 4% / TAZAROTENE 0.05% - niacinamide 4% / tazarotene 0.05% 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](#).*

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**NIACINAMIDE 4% / TAZAROTENE 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](http://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  
TAZAROTEN  
CREAM 0.05%

SINCE

Rx only

Lot: 011312ABCD EFGH@1

BUD: 01/01/1970

MFG: 01/01/1970

**Active ingredients**

Niacinamide USP ..... 4%  
Tazarotene ..... 0.05%

**Inactive ingredients**

Suspendisse Cream ..... 95.95%

**NDC 72934-2060-2**  
**NIACINAMIDE 4 / TAZAROTENE 0.05**  
**Cream 30gm**





## NIACINAMIDE 4% / TAZAROTENE 0.05%

niacinamide 4% / tazarotene 0.05% cream

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-2160
<b>Route of Administration</b>	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	0.05 g in 100 g

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-2160-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2019	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/01/2019	

**Labeler** - Sincerus Florida, LLC (080105003)**Establishment**

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2160)

Revised: 4/2019

Sincerus Florida, LLC