

**NIACINAMIDE 2% / SPIRONOLACTONE 5% / TRETINOIN 0.05% - niacinamide 2% /  
spironolactone 5% / tretinoin 0.05% 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](#).*

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**NIACINAMIDE 2% / SPIRONOLACTONE 5% / TRETINOIN 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).

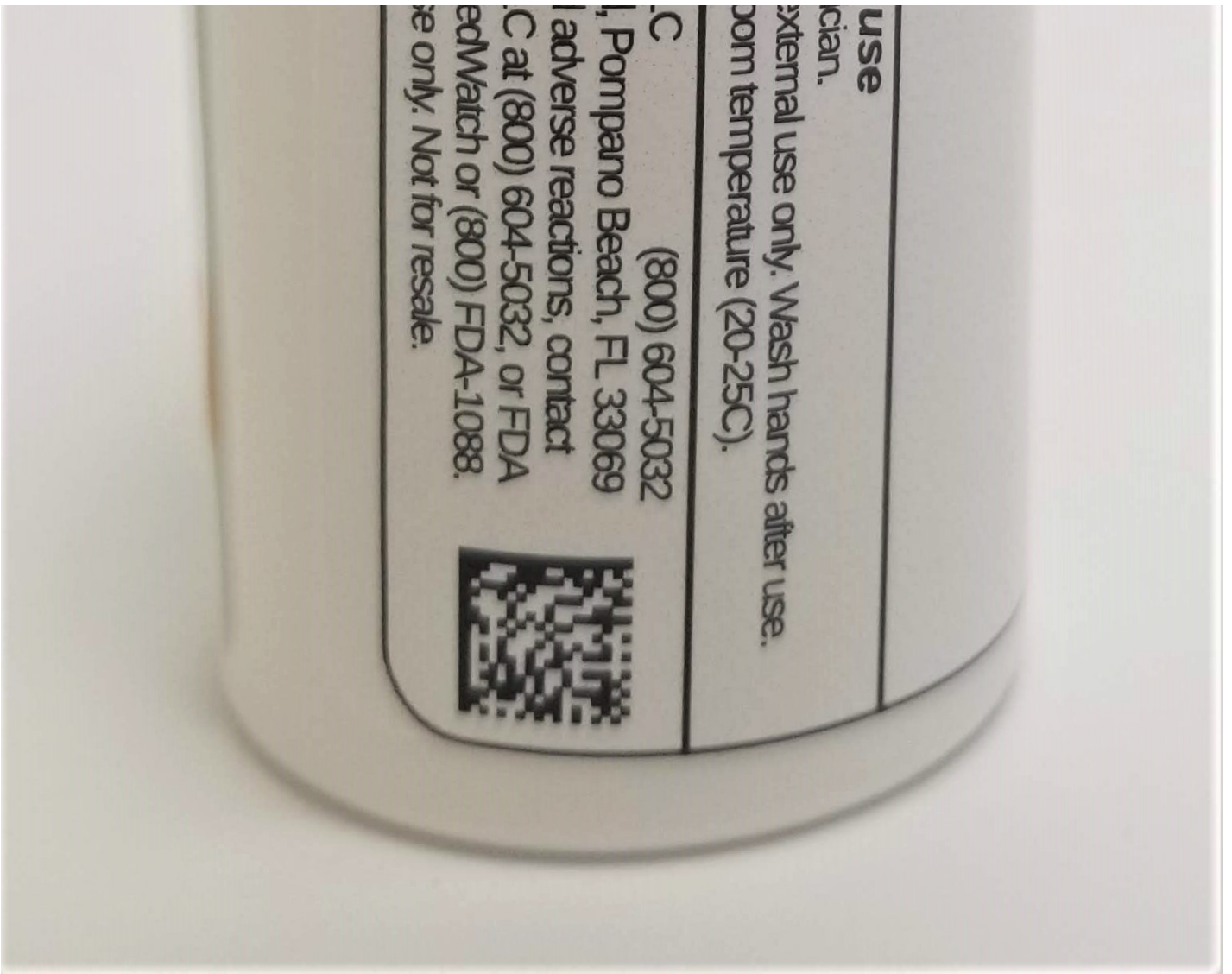
Sincerus Florida, LLC (800) 604-5032  
3265 W McNab Rd, Pompano Beach, FL 33069  
To report suspected adverse reactions, contact  
Sincerus Florida, LLC at (800) 604-5032, or FDA  
at [www.FDA.gov/MedWatch](http://www.FDA.gov/MedWatch) or (800) FDA-1088.  
Office use only. Not for resale.



Sincerus Florida, LLC adverse reactions.

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Office us



**Active, inactive**



25 FEB 83

Rx only

BUD: 01M011970

Lot: 011220ABCDEF GHIJ

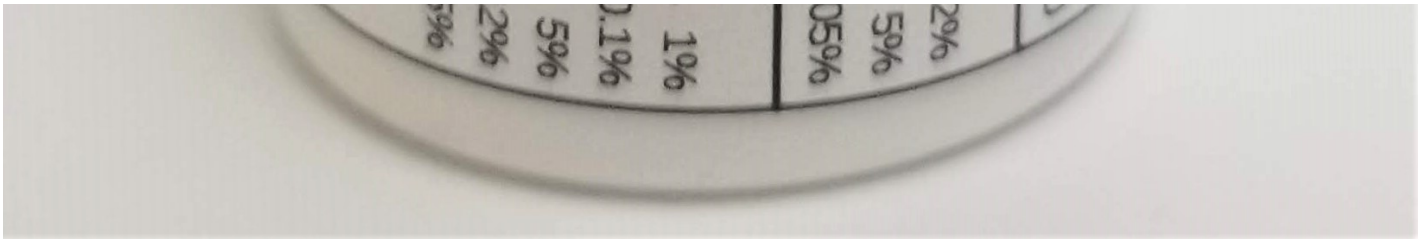
MFG: 01M011970

**Active ingredients**

Niacinamide USP	.....	.....
Spirolactone USP	.....	.....
Tretinoin USP	.....	0.1

**Inactive ingredients**

Ascorbyl Palmirate Fcc	.....	.....
Butylated Hydroxytoluene NF (BHT)	.....	.....
Glycerin And Artificial Rose Water	.....	.....
Sodium Metabisulfite NF	.....	0.
Suspendisse Gel	.....	86.65



**NDC 72934-1153**

**NIACINAMIDE 2% / SPIRONOLACTONE 5% / TRETINOIN 0.05%**

**Gel 30 gm.**





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niacinamide 2% / spironolactone 5% / tretinoin 0.05% gel

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-1153
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 g in 100 g
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.05 g in 100 g
SPIRONOLACTONE (UNII: 27O7W4T232) (SPIRONOLACTONE - UNII:27O7W4T232)	SPIRONOLACTONE	5 g in 100 g

### Product Characteristics

Color	yellow	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-1153-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/03/2019	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/03/2019	

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

### Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida LLC		080105003	manufacture(72934-1153)

Revised: 5/2019

Sincerus Florida LLC