

**011054 NIACINAMIDE 4% / SPIRONOLACTONE 5% - 011054 niacinamide 4% /
spironolactone 5% 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](#).

011054 NIACINAMIDE 4% / SPIRONOLACTONE 5%

Directions for use



Directions for use

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.

LS-01-000000-00



Sincerus Florida, LLC adverse reactions

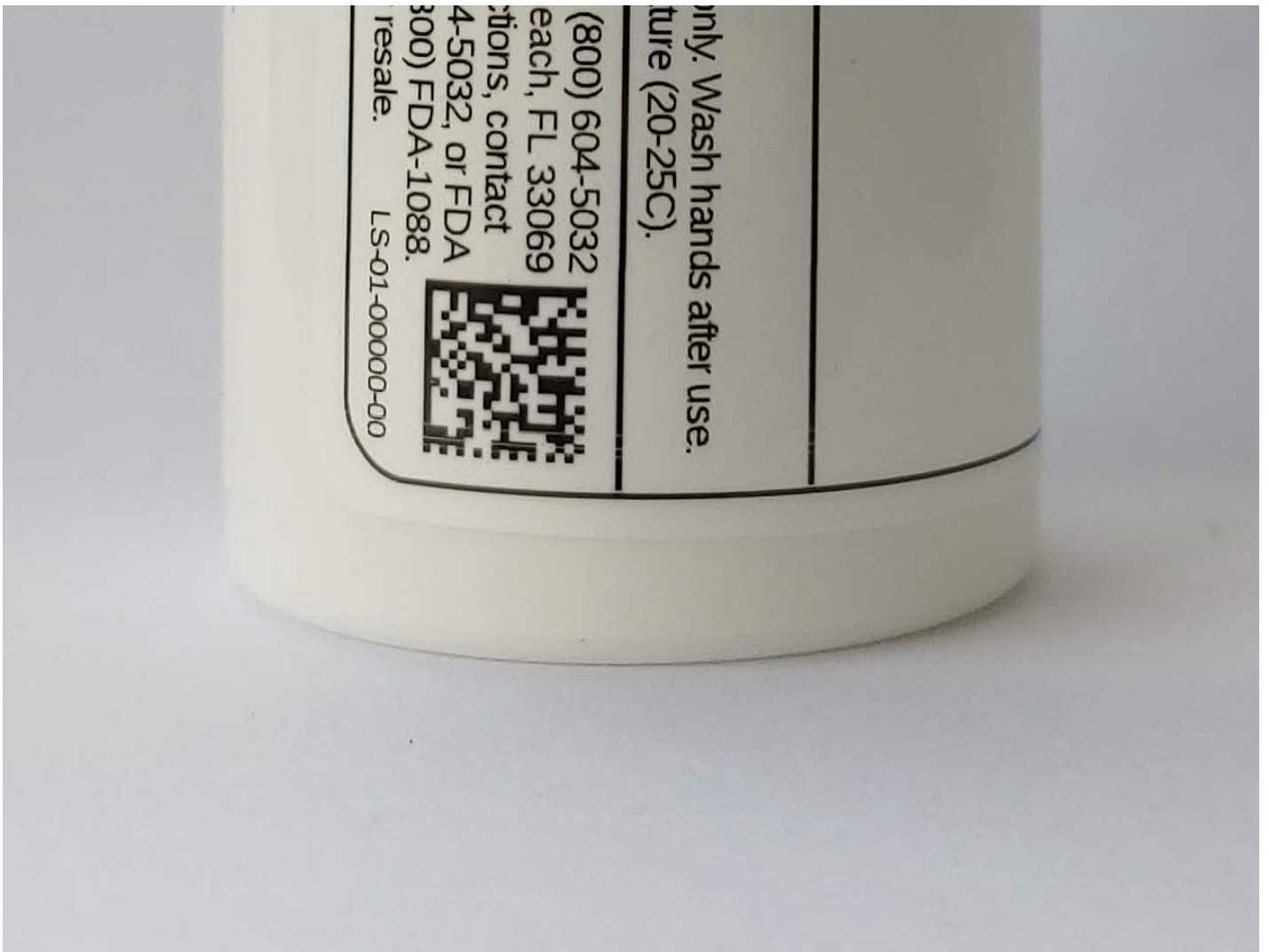
Directions for use

As directed by Physician.

Apply topically. For external use only.
Store at controlled room temperature.

Sincerus Florida, LLC

3265 W McNab Rd, Pompano Beach, FL 33069
To report suspected adverse reactions, contact
Sincerus Florida, LLC at (800) 660-1111 or
at www.FDA.gov/MedWatch or (305) 850-6600.
Office use only. Not for sale.



Active, inactive



011054
NIACINAMIDE
SPIRONOLACTONE
GEL 30g

SINCE

Rx only
BUD: XXXXXXXXX

Lot: 011054XXXXXXXXXX@XX
MFG: XXXXXXXXX

Active ingredients

Niacinamide USP	4%
Spirolactone USP	5%

Inactive ingredients

Glycerin USP	5%
Lavender Oil	0.25%
Suspendisse Gel	84.5%

NDC 72934-1205-2 011054 NIACINAMIDE 4% / SPIRONOLACTONE 5% gel 30 gm



NDC 72934-1205-2

011054

NIACINAMIDE 4%

SPIRONOLACTONE 5%

GEL 30gm

Rx only

Lot: 013



011054 NIACINAMIDE 4% / SPIRONOLACTONE 5%

011054 niacinamide 4% / spironolactone 5% gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-1205
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	4 g in 100 g
SPIRONOLACTONE (UNII: 27O7W4T232) (SPIRONOLACTONE - UNII:27O7W4T232)	SPIRONOLACTONE	5 g in 100 g

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-1205-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/02/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/02/2020	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-1205)

Revised: 7/2020

Sincerus Florida, LLC