

**NIACINAMIDE 2% / SPIRONOLACTONE 5% / TRETINOIN 0.025% - niacinamide 2% /
spironolactone 5% / tretinoin 0.025% 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 2% / SPIRONOLACTONE 5% / TRETINOIN 0.025%

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



Sincerus Florida, LLC adverse reactions



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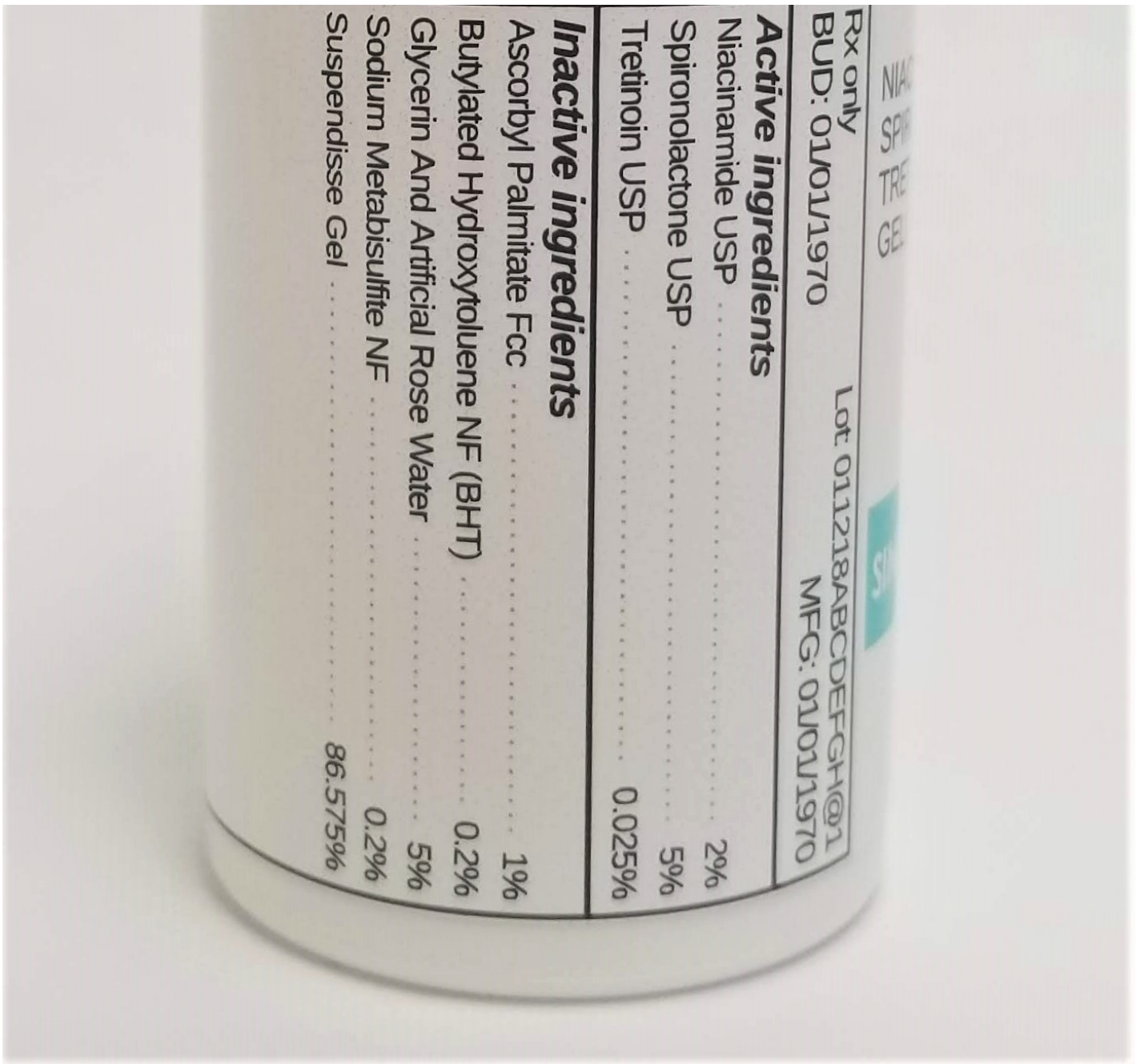
Office use only. Not for resale.





Actives, inactives





NDC 72934-1152-2. NIACINAMIDE USP 2% / SPIRONOLACTONE USP 5% / TRETINOIN USP 0.025%. Gel30gm.



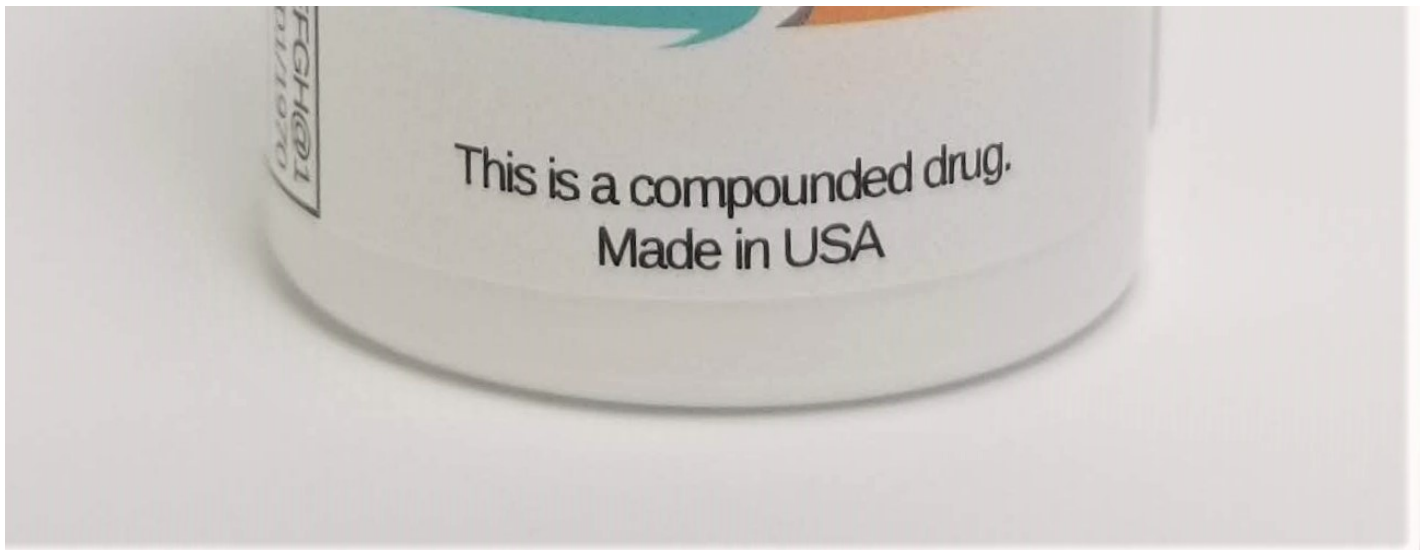
Rx only
BUD: 01/01/1970

Lot: 011218ABCDE
MFG: 01/19

NDC 72934-1152-2

NIACINAMIDE USP 2%
SPIRONOLACTONE USP 5%
TRETINOIN USP 0.025%
GEL 30gm





NIACINAMIDE 2% / SPIRONOLACTONE 5% / TRETINOIN 0.025%

niacinamide 2% / spironolactone 5% / tretinoin 0.025% gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-1152
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 g in 100 g
SPIRONOLACTONE (UNII: 27O7W4T232) (SPIRONOLACTONE - UNII:27O7W4T232)	SPIRONOLACTONE	5 g in 100 g
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.025 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:72934-1152-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/04/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/04/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-1152)

Revised: 5/2019

Sincerus Florida, LLC