011020 NIACINAMIDE 4% / TRETINOIN 0.05%- 011020 niacinamide 4% / 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.

011020 NIACINAMIDE 4% / TRETINOIN 0.05%

Directions for use

Sincerus Florida, LLC adverse reactions

Directions for use

As directed by Physician.

Apply topically. For external use o

Store at controlled room tempera

Sincerus Florida, LLC

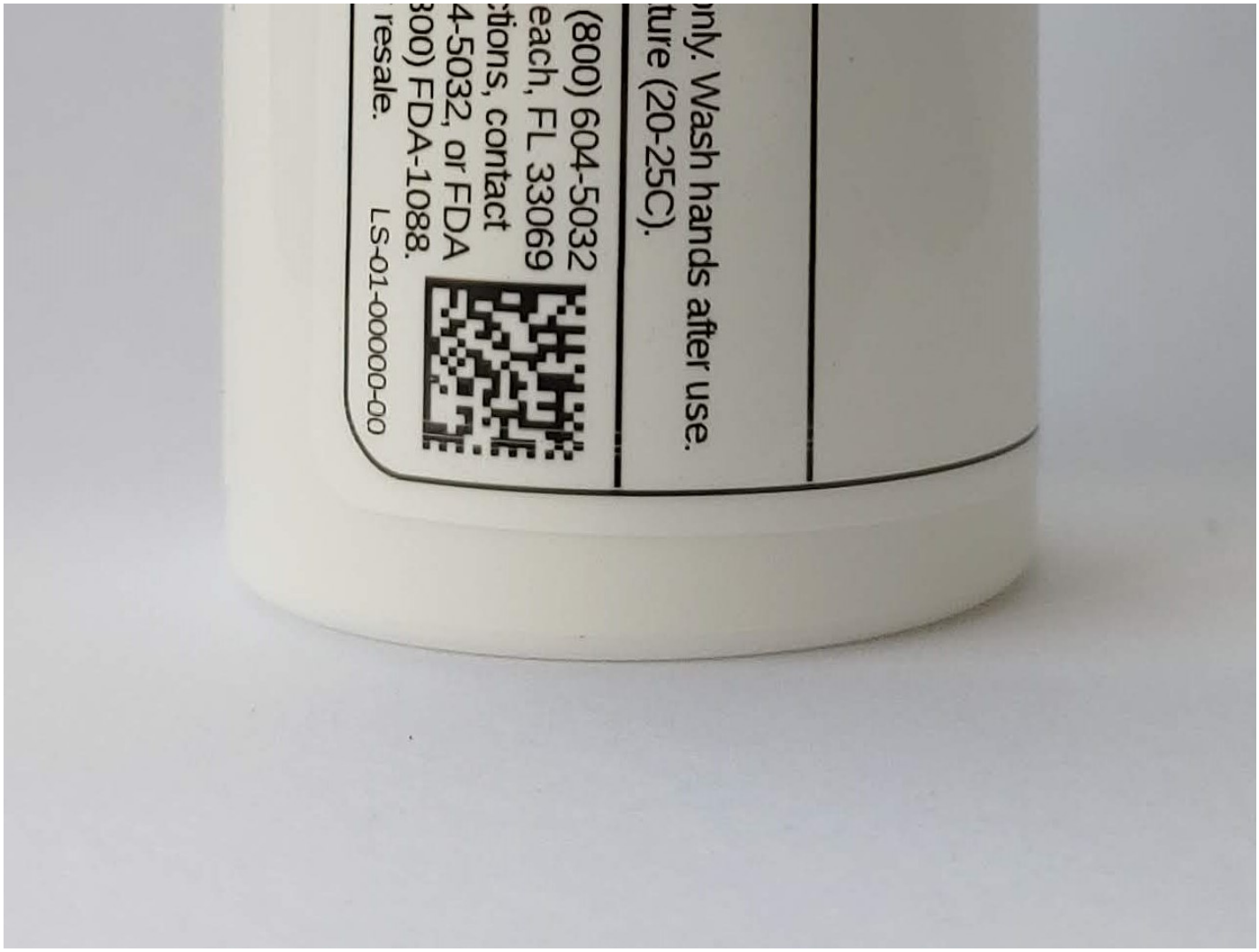
3265 W McNab Rd, Pompano B

To report suspected adverse reac

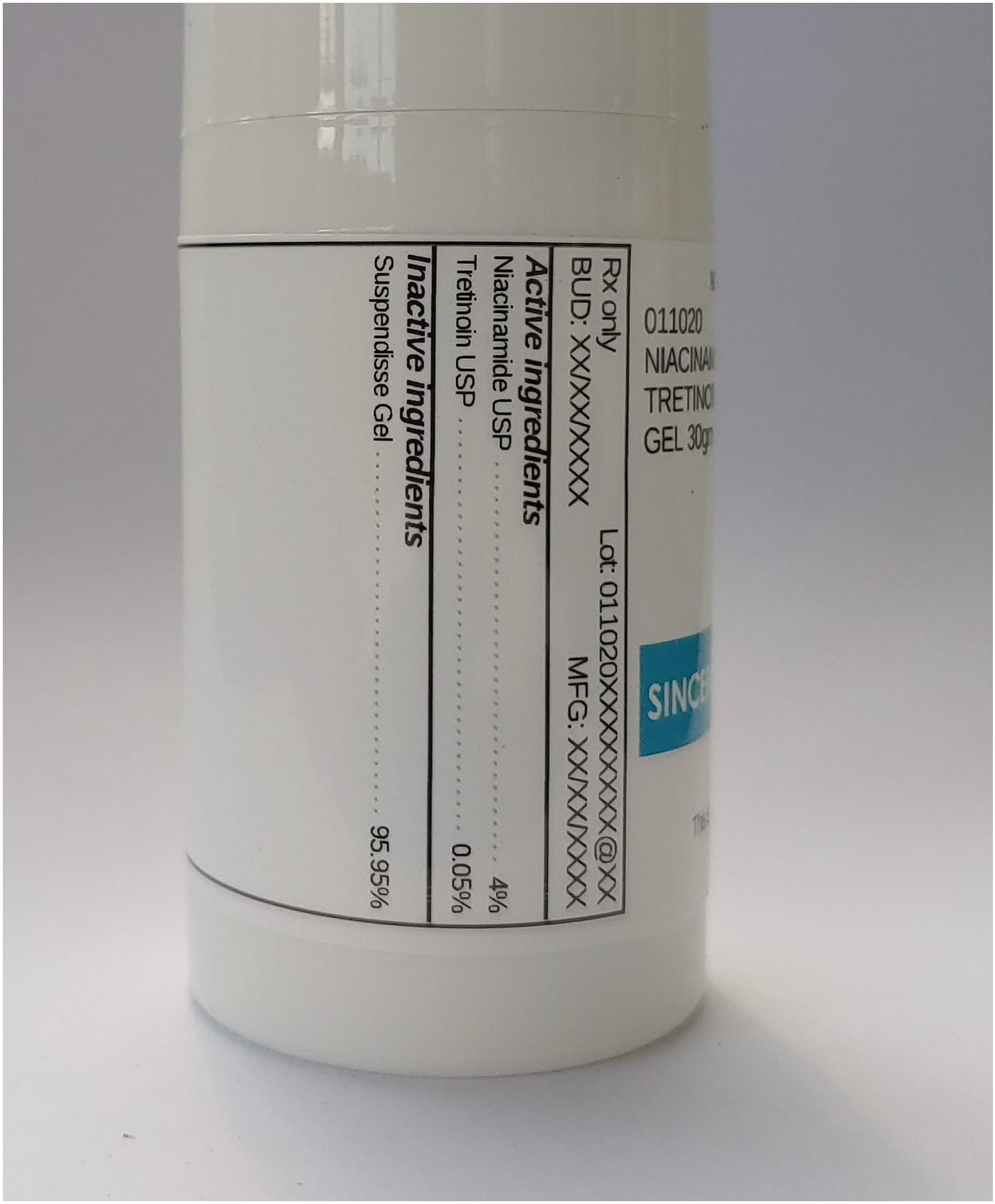
Sincerus Florida, LLC at (800) 60

at www.FDA.gov/MedWatch or (8

Office use only. Not for

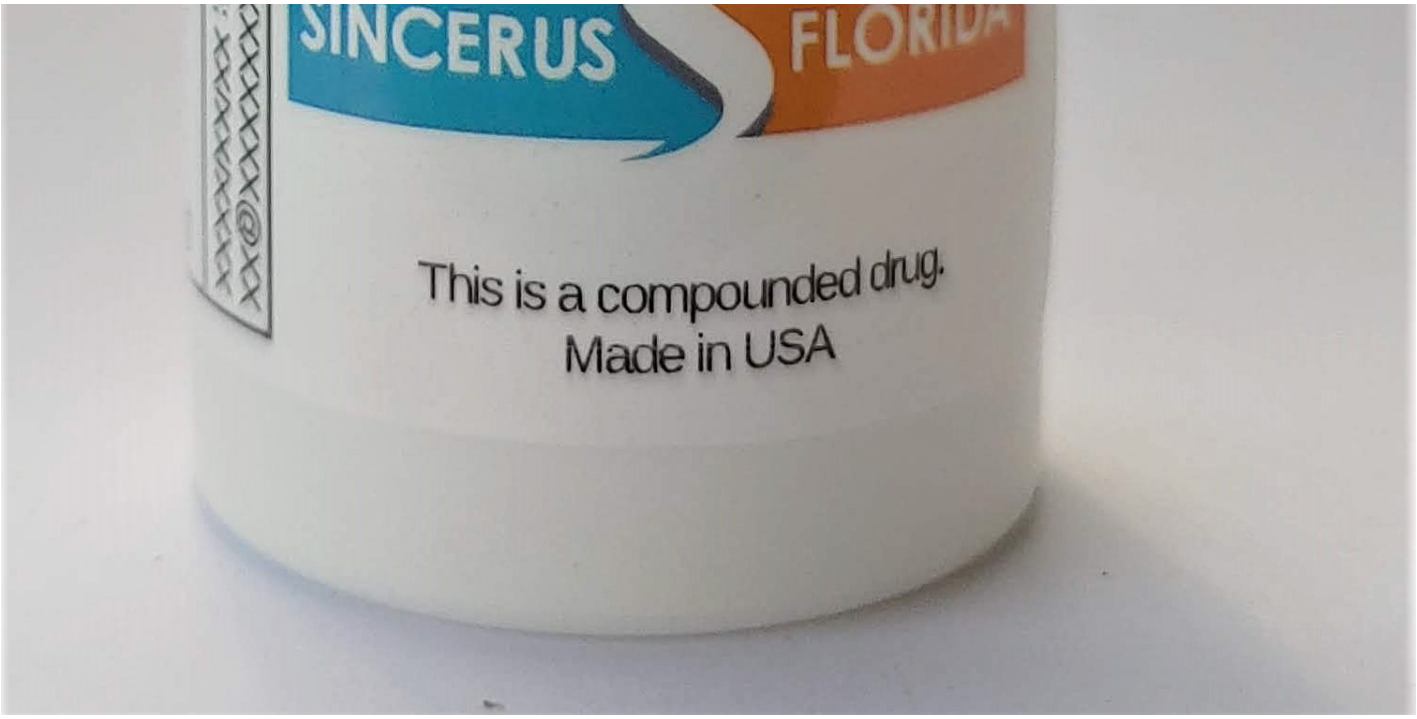


Active, inactive



NDC 72934-1204-2 011020 NIACINAMIDE 4% / TRETINOIN 0.05% gel 30 gm

NDC 72934-1204-2
011020
NIACINAMIDE 4%
TRETINOIN 0.05%
GEL 30gm
Rx only
BUD: XX/XX/XXXX
Lot: 011020XX
MFG



011020 NIACINAMIDE 4% / TRETINOIN 0.05%

011020 niacinamide 4% / tretinoin 0.05% gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-1204
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
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.05 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:72934-1204-2	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(72934-1204)

Revised: 7/2020 Sincerus Florida, LLC