

011504 DAPSONE 8.5% / NIACINAMIDE 4% - 011504 dapsone 8.5% / niacinamide 4% 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](#).

011504 DAPSONE 8.5% / NIACINAMIDE 4%

Directions for use



Directions for use

is 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
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.



LS-01-000000-00

Sincerus Florida, LLC adverse reactions



Directions for use

As directed by Physician.

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Sincerus Florida, LLC (8

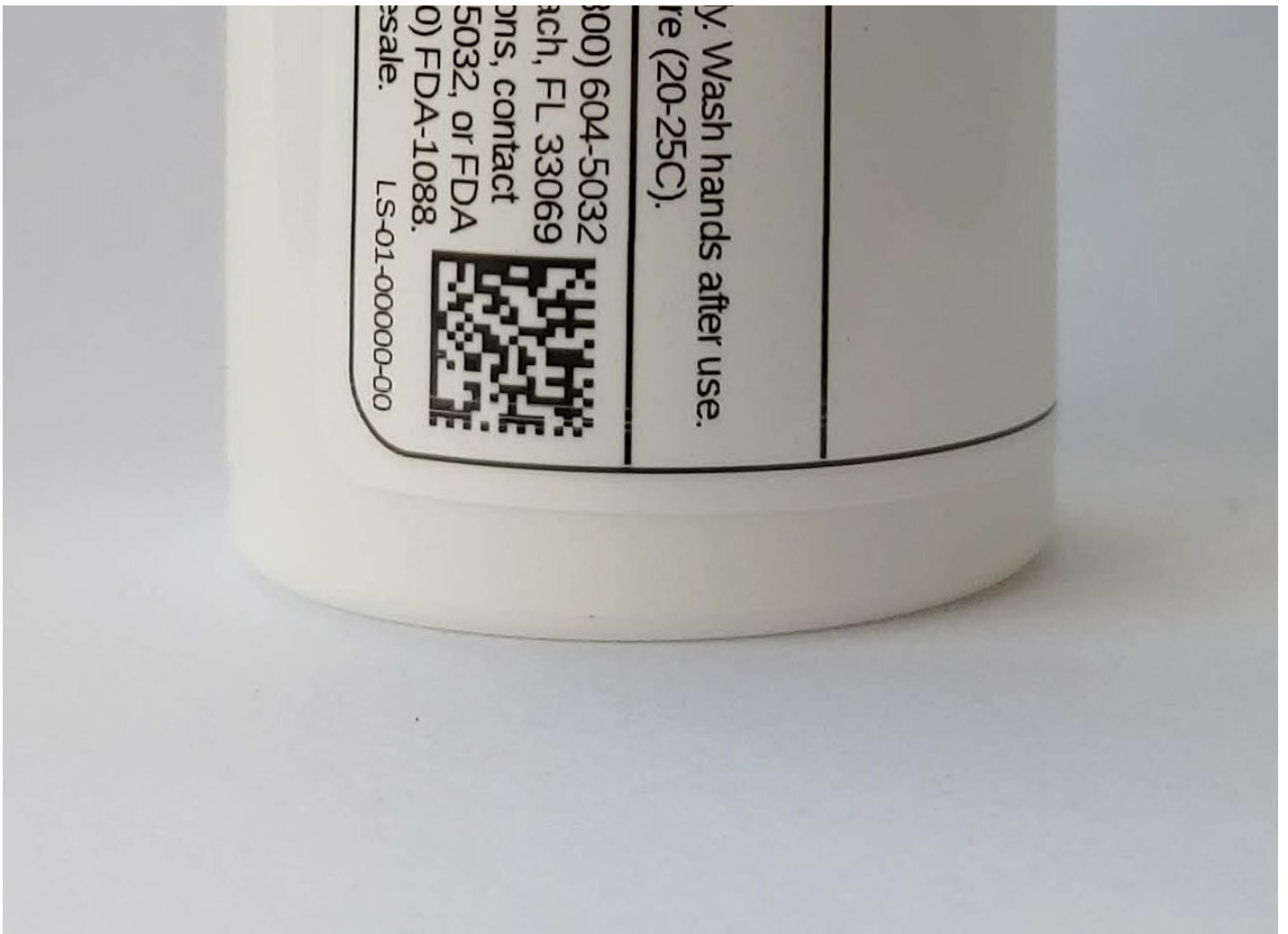
3265 W McNab Rd, Pompano Beach

To report suspected adverse reactions

Sincerus Florida, LLC at (800) 604-

at www.FDA.gov/MedWatch or (800)

Office use only. Not for retail



Active, inactive



011504
DAPSONE
NIACINAMIDE
GEL 30gm

SINCE

Rx only
BUD: XXXXXXXXXX
Lot: 011504XXXXXXXXXXXX@XX
MFG: XXXXXXXXXX

Active ingredients

Dapsone USP 8.5%
Niacinamide USP 4%

Inactive ingredients

Propylene Glycol USP 5%
Suspendisse Gel 82.5%

NDC 72934-1212-2 011504 DAPSONE 8.5% / NIACINAMIDE 4% gel 30 gm

NDC 72934-1212-2

011504

DAPSONE 8.5%

NIACINAMIDE 4%

GEL 30gm

RX only
XXXXXXXXXXXXXXXXXXXX

Lot: 0



011504 DAPSONE 8.5% / NIACINAMIDE 4%

011504 dapsone 8.5% / niacinamide 4% gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-1212
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DAPSONE (UNII: 8W5C518302) (DAPSONE - UNII:8W5C518302)	DAPSONE	8.5 g in 100 g
NIACINAMIDE (UNII: 25X5118RD4) (NIACINAMIDE - UNII:25X5118RD4)	NIACINAMIDE	4 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:72934-1212-	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination	07/02/2020	

2	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(72934-1212)

Revised: 7/2020

Sincerus Florida, LLC