

DAPSONE 6% / NIACINAMIDE 4% - dapsone 6% / 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](#).

DAPSONE 6% / NIACINAMIDE 4%

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).

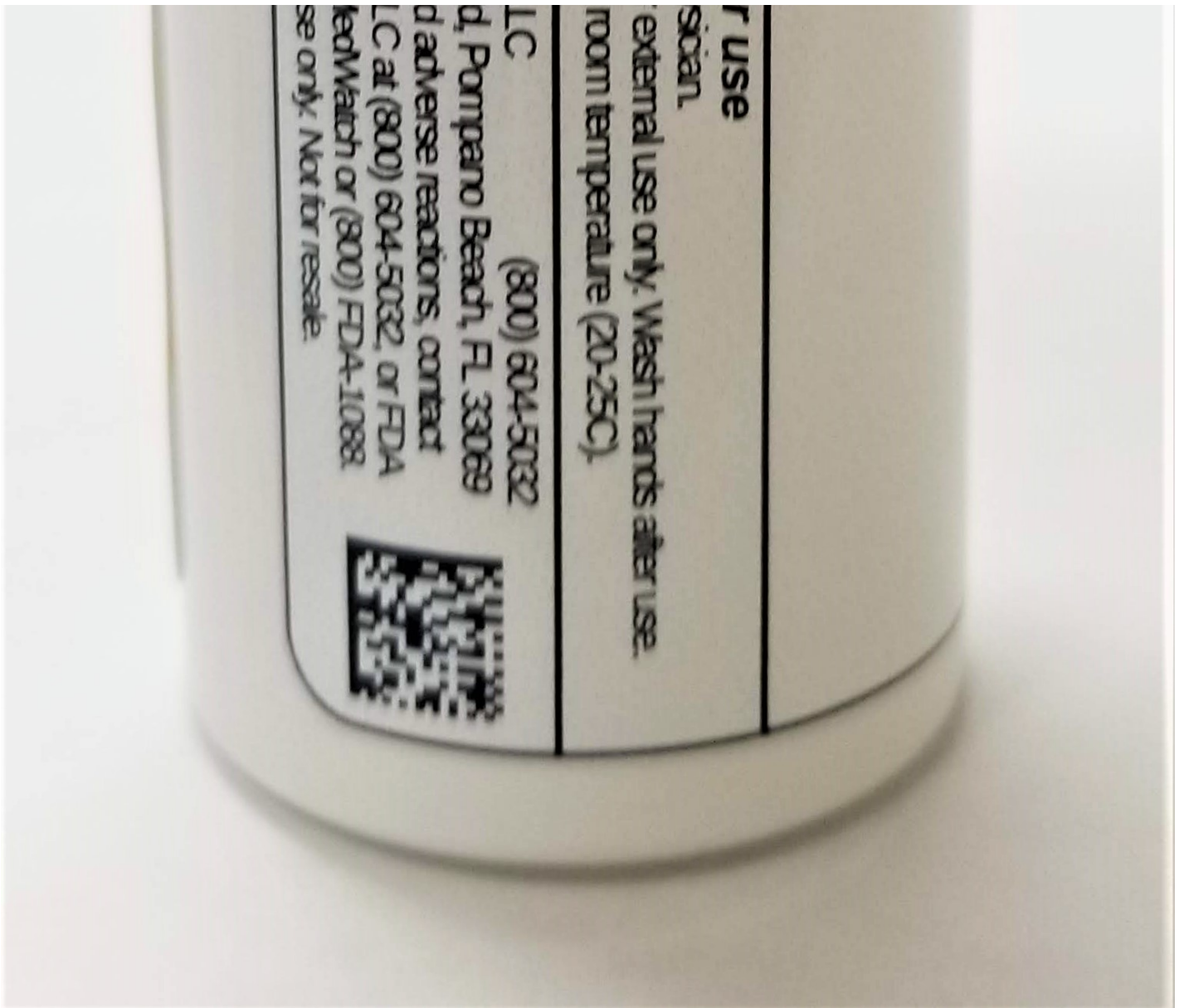
Sincerus Florida, LLC (800) 604-5032
3265 W McNab Rd, Pompano Beach, FL 33069
To report suspected adverse reactions, contact
Sincerus Florida, LLC at (800) 604-5032, or FDA
at www.FDA.gov/medwatch or (800) FDA-1088.
Office use only. Not for resale.



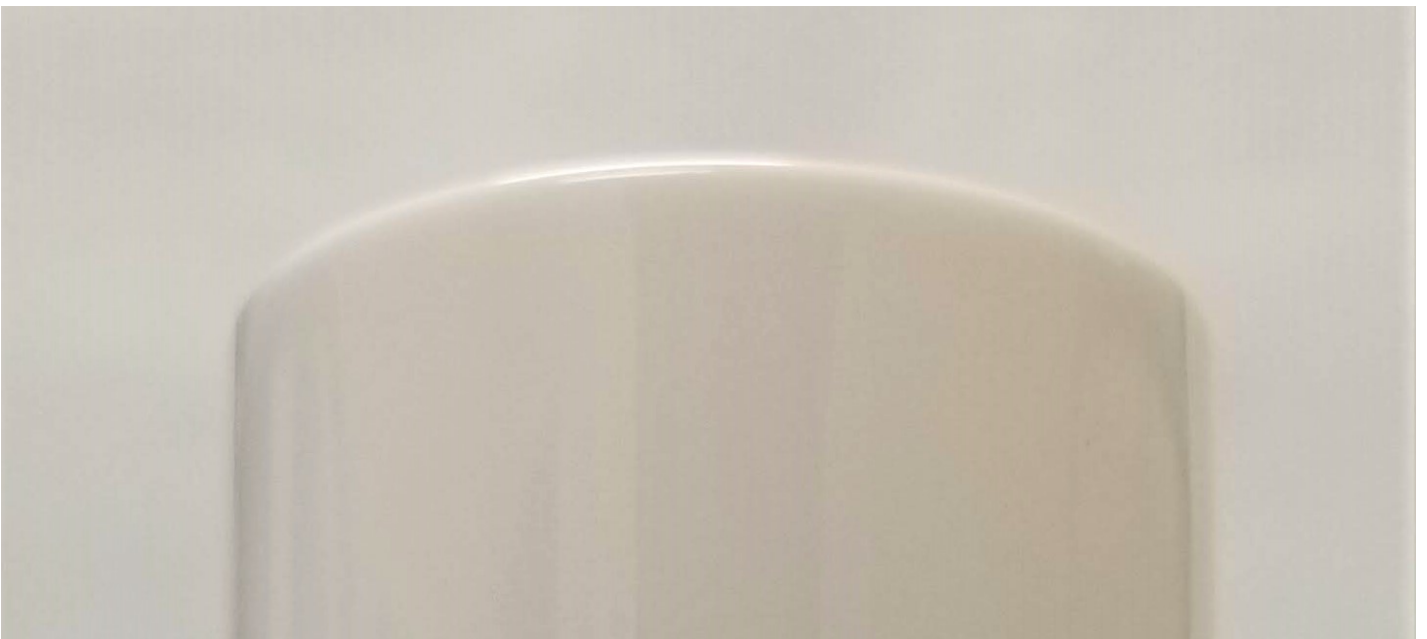
Sincerus Florida, LLC adverse reactions.

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3265 W McNab R
To report suspecte
Sinoerus Florida, L
at www.FDA.gov/M
Office US



Active, inactive



DAPS
NIAC
GEL

SIN

Rx only
BUD: 01/01/1970

Lot: 011500ABCD
MFG: C

Active ingredients

Dapsone USP
Niacinamide USP

Inactive ingredients

Butylated Hydroxytoluene NF (BHT)
Propylene Glycol USP
Suspendisse Gel



NDC 72934-1063-2
DAPSONE 6 / NIACINAMIDE 4
Gel 30gm



NDC 72934-1063-2

**DAPSONE USP 6%
NIACINAMIDE USP 4%
GEL 30gm**

Rx only
BUD: 01/01/1970

Lot: 011500ABCDEF@1
MFG: 01/01/1970

SINCERUS

FLORIDA

This is a compounded drug.
Made in USA

DAPSONE 6% / NIACINAMIDE 4%

dapsone 6% / niacinamide 4% gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-1063
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DAPSONE (UNII: 8W5C518302) (DAPSONE - UNII:8W5C518302)	DAPSONE	6 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:72934-1063-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/01/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-1063)