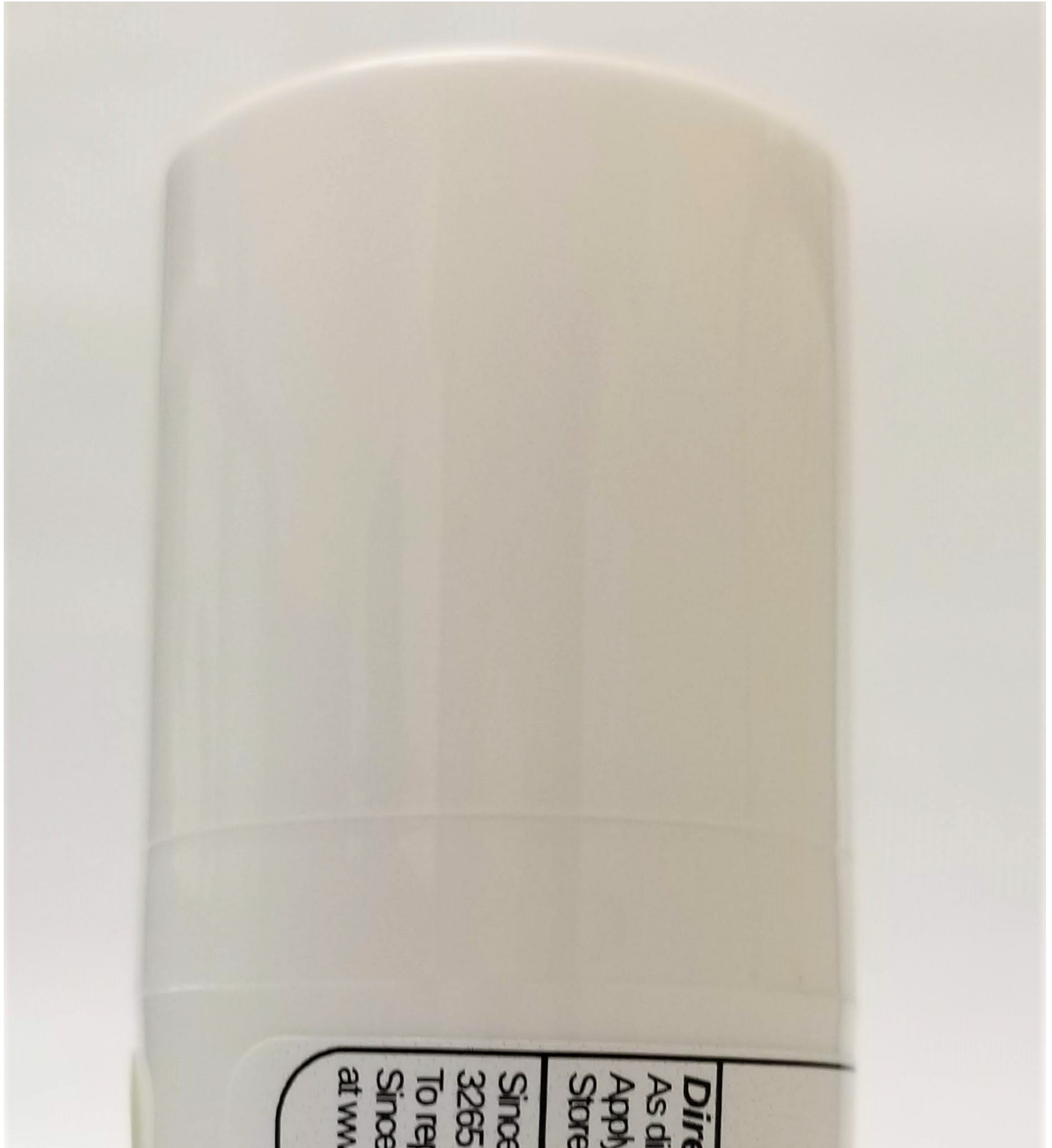


DAPSONE 8.5% / NIACINAMIDE 4% - dapsone 8.5% / niacinamide 4% gel
Sincerus Florida, LLC

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](#).

DAPSONE 8.5% / NIACINAMIDE 4%

Directions for use



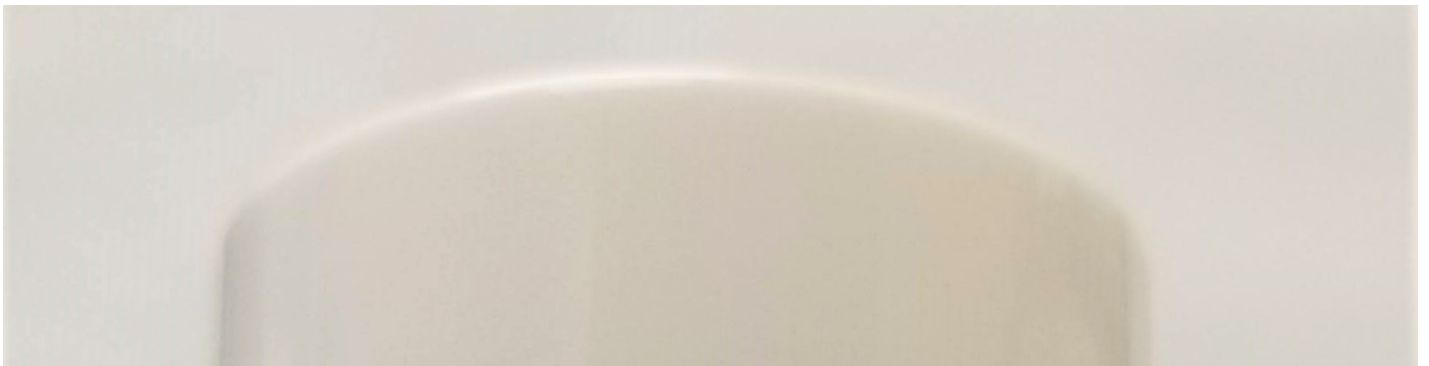
actions for use

ected by Physician.
/topically. For external use only. Wash hands after use.
at controlled room temperature (20-25C).

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



Sincerus Florida, LLC adverse reactions.



Directions for use

As directed by Physician.

Apply topically. For external use only. Wash

Store at controlled room temperature (20-25

Sincerus Florida, LLC (800) 604

3265 W McNab Rd, Pompano Beach, FL 3

To report suspected adverse reactions, cont

Sincerus Florida, LLC at (800) 604-5032, or

at www.FDA.gov/MedWatch or (800) FDA-1

Office use only. Not for resale.



Active, inactive



DAPSONE
NIACINAMIDE
GEL 300

SING

Rx only

BUD: 01/01/1970

Lot: 011504ABCDEF@1

MFG: 01/01/1970

Active ingredients

Dapsone USP 8.5%
Niacinamide USP 4%

Inactive ingredients

Propylene Glycol USP 5%
Suspendisse Gel 82.5%

NDC 72934-1066-2
DAPSONE 8.5 / NIACINAMIDE 4
Gel 30gm



Rx only
NDC 72934-1066-2

NDC 72934-1066-2

DAPSONE USP 8.5%

NIACINAMIDE USP 4%

GEL 30gm



DAPSONE 8.5% / NIACINAMIDE 4%

dapsone 8.5% / niacinamide 4% gel

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-1066
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: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	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:72934-1066-2	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(72934-1066)

Revised: 4/2019

Sincerus Florida, LLC