

**BENZOYL PEROXIDE 8% / NIACINAMIDE 4% - benzoyl peroxide 8% / niacinamide 4% suspension**  
**Sincerus Florida**

*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](#).*

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**BENZOYL PEROXIDE 8% / NIACINAMIDE 4%**

**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  
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](http://www.FDA.gov/MedWatch) or (800) FDA-1088.

Office use only. Not for resale.





**Sincerus Florida, LLC. Adverse reactions**



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**Active, inactive**





BENZOYL  
NIACINAMIDE  
SUSPENSION

Rx only

BUD: 01/01/1970

**Active ingredients**

Benzoyl Peroxide USP

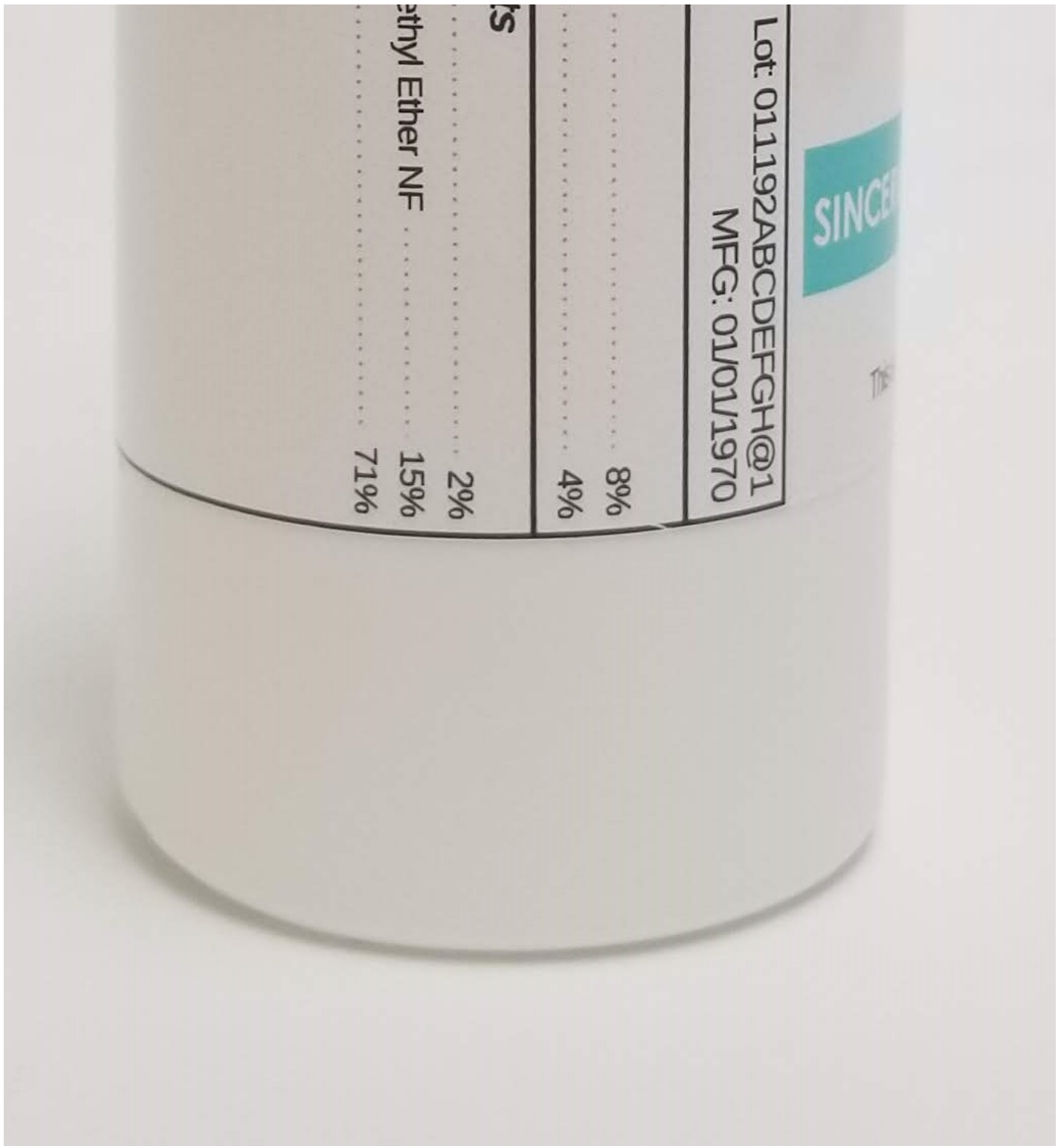
Niacinamide USP

**Inactive ingredients**

Cocamide Dea

Diethylene Glycol Mono

Suspendisse Shampoo



**NDC 72934-8022-6 BENZOYL PEROXIDE USP 8% / NIACINAMIDE USP 4%. Suspension 120gm.**



RX C  
BUD  
ACR

**NDC 72934-8022-6**

... 0.0%



BENZOYL PEROXIDE USP 8%  
NIACINAMIDE USP 4%  
SUSPENSION 120gm

only  
01/01/1970  
ve ingredients

Lot: 011192ABCD EFGH@1  
MFG: 01/01/1970



This is a compounded drug.  
Made in USA

benzoyl peroxide 8% / niacinamide 4% suspension

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-8022
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	8 g in 100 g
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	4 g in 100 g

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-8022-6	120 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/11/2019	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/11/2019	

**Labeler** - Sincerus Florida (080105003)

### Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-8022)

Revised: 5/2019

Sincerus Florida