

**SALICYLIC ACID 2% /SODIUM SULFACETAMIDE MONOHYDRATE 8% - salicylic acid 2% /sodium sulfacetamide monohydrate 8% suspension**

**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](#).*

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**SALICYLIC ACID 2% /SODIUM SULFACETAMIDE MONOHYDRATE 8%**

**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. Adverse Reactions**



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**Active, Inactive**





NO  
SALICYLIC  
SULFACET  
MONOHYD  
SUSPENS

Rx only

Lot:

BUD: 01/01/1970

**Active ingredients**

Salicylic Acid USP .....

Sulfacetamide Sodium Monor

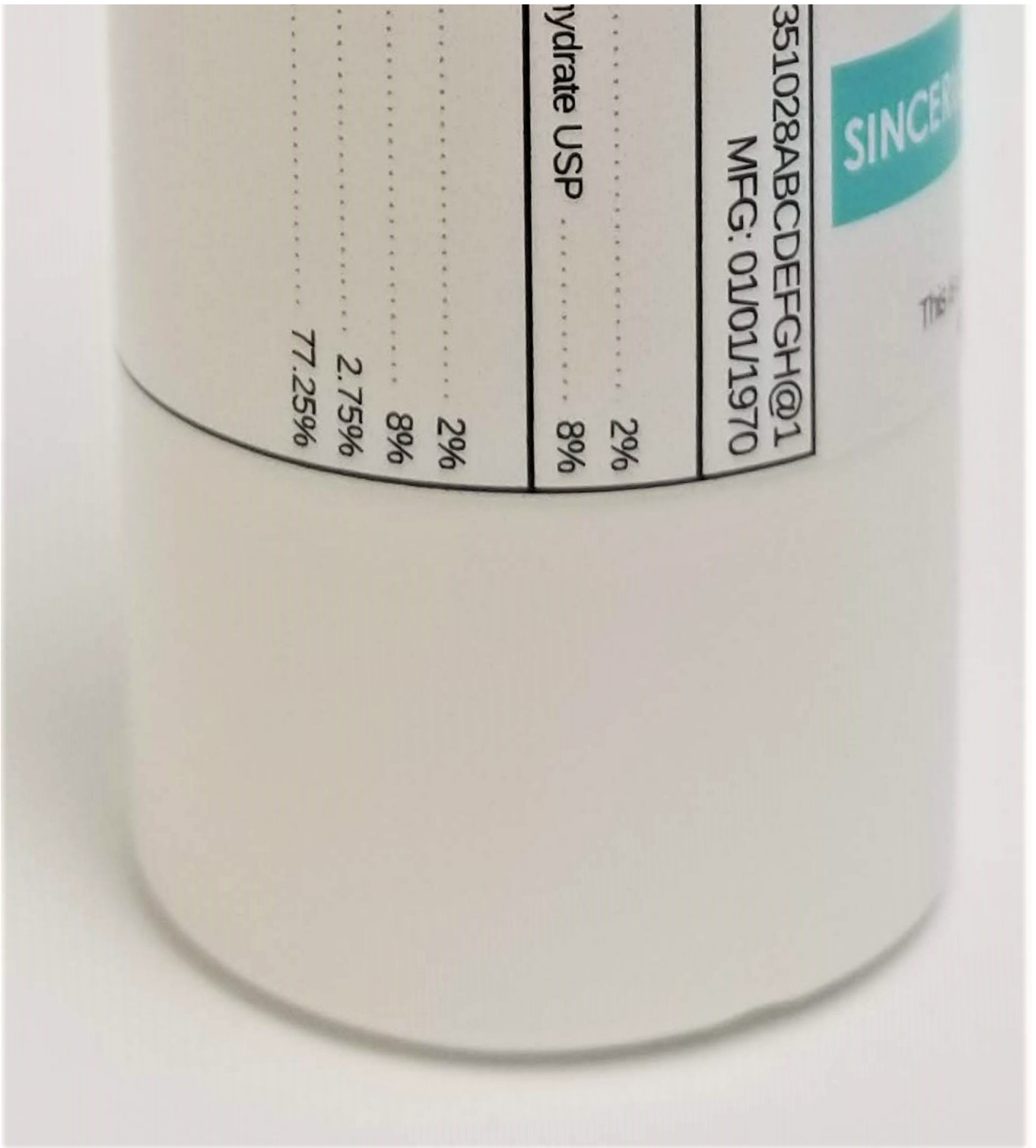
**Inactive ingredients**

Cocamide Dea .....

Propylene Glycol USP .....

Sulfur Precipitated USP .....

Suspendisse Shampoo .....



**NDC 72934-8170-6 SALICYLIC ACID 2% /SODIUM SULFACETAMIDE MONOHYDRATE  
8% Suspension 120 gm**



Rx only  
BUD: 01/01/1970

**NDC 72934-8170-6**

SALICYLIC ACID USP 2%  
SULFACETAMIDE SODIUM  
MONOHYDRATE USP 8%  
SUSPENSION 120gm





**SALICYLIC ACID 2% /SODIUM SULFACETAMIDE MONOHYDRATE 8%**

salicylic acid 2% /sodium sulfacetamide monohydrate 8% suspension

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-8170
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
SULFACETAMIDE SODIUM (UNII: 4NRT660KJQ) (SULFACETAMIDE - UNII:4965G3J0F5)	SULFACETAMIDE SODIUM	8 g in 100 g
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 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:72934-8170-6	120 g in 1 CYLINDER; 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(72934-8170)

Revised: 4/2019

Sincerus Florida, LLC