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.

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 or (800) FDA-1088.

Office use only. Not for resale.

Sincerus Florida, LLC. Adverse reactions

Active, inactive

NDC 72

SALICYLIC AC
SULFACETAMI
MONOHYDRAT
SUSPENSION

Rx only
BUD: 01/01/1970

Active ingredients

Salicylic Acid USP . . . . . .

Sulfacetamide Sodium Mo

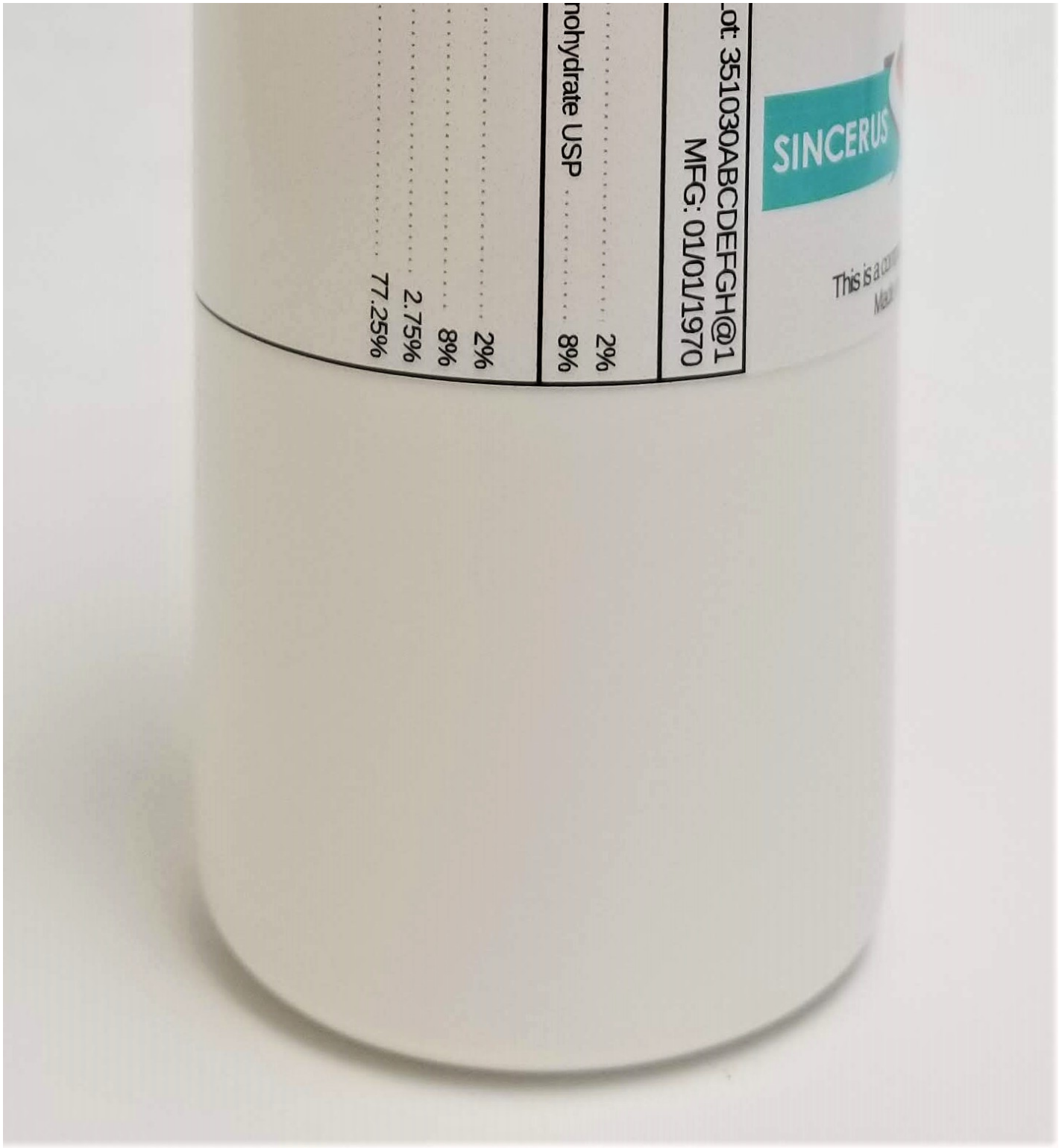
Inactive ingredients

Cocamide Dea . . . . . .

Propylene Glycol USP . . . .

Sulfur Precipitated USP . . .

Suspendisse Shampoo . . . .



NDC 72934- 8170-7 SALICYLIC ACID USP 2% / SODIUM SULFACETAMIDE MONOHYDRATE USP 8% . Suspension 240gm


NDC 72934-8170-7
SALICYLIC ACID USP 2%
SULFACETAMIDE SODIUM
MONOHYDRATE USP 8%
SUSPENSION 240gm
SINCERUS FLORIDA
Rx only
BUD: 01/01/1970
Lot: 351030ABCDEFG
MFG: 01/01
Active ingredients
Salicylic Acid USP
Sulfacetamide Sodium Monohydrate USP
Inactive Ingredients

SALICYLIC ACID 2% /SODIUM SULFACETAMIDE MONOHYDRATE 8%

salicylic acid 2% /sodium sulfacetamide monohydrate 8% suspension

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-8170
Route of Administration	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:72934-8170-7	240 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/17/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/17/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-8170)

Revised: 5/2019 Sincerus Florida, LLC