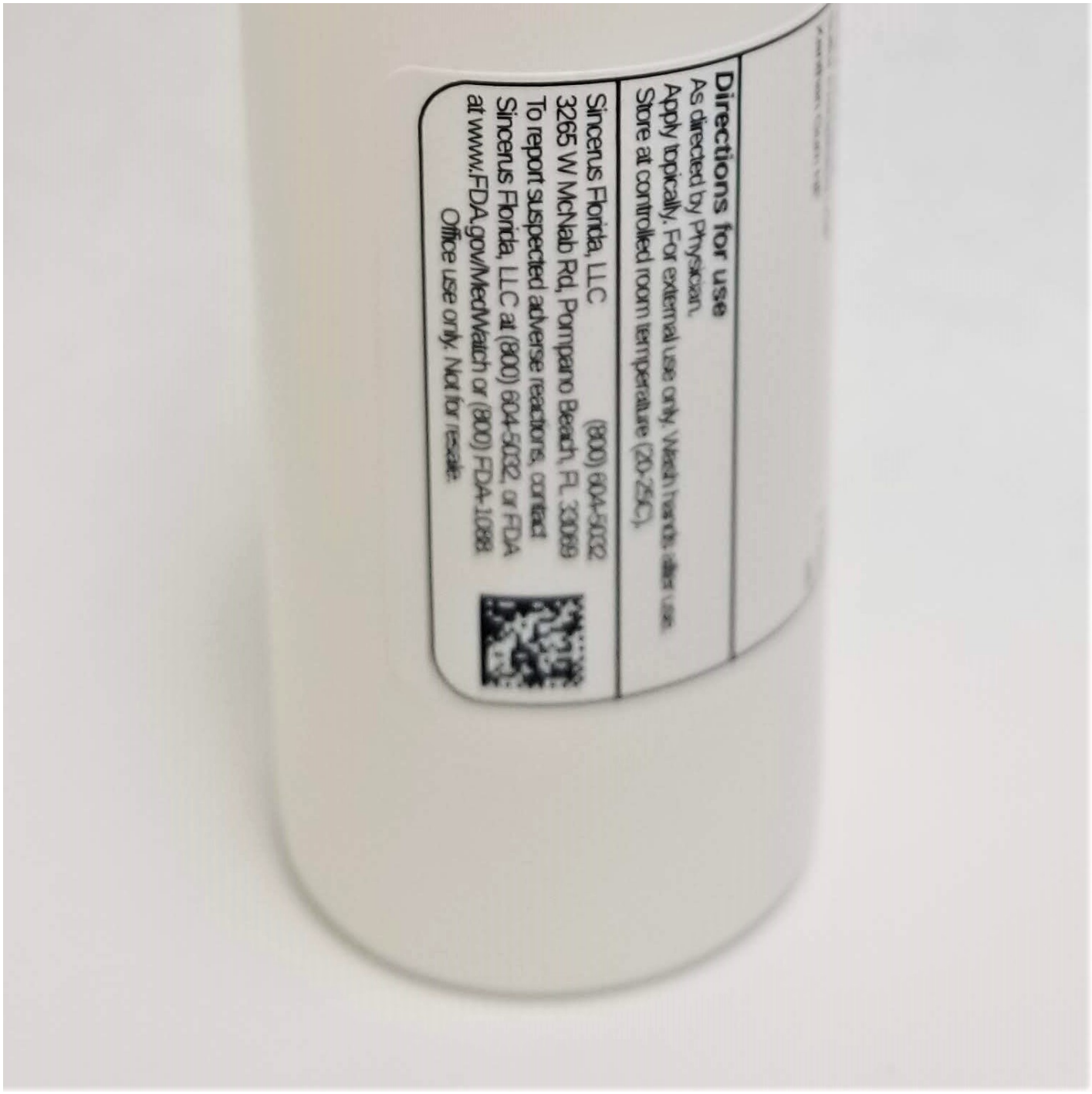
SALICYLIC ACID 2% / SODIUM SULFACETAMIDE MONOHYDRATE 8%- salicylic acid 2% / sodium sulfacetamide monohydrate 8% suspension
Sincerus Florida, LLC

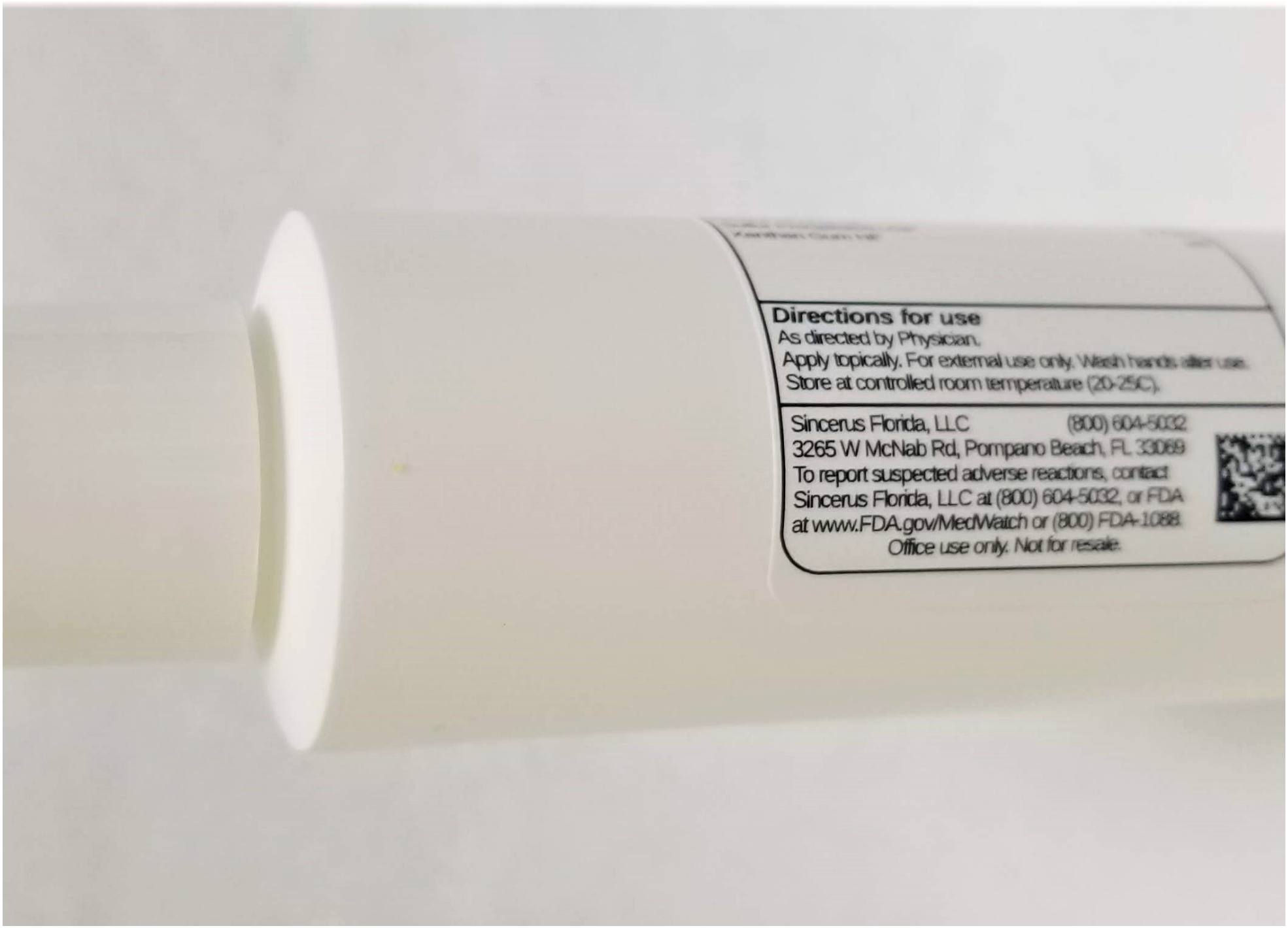
Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

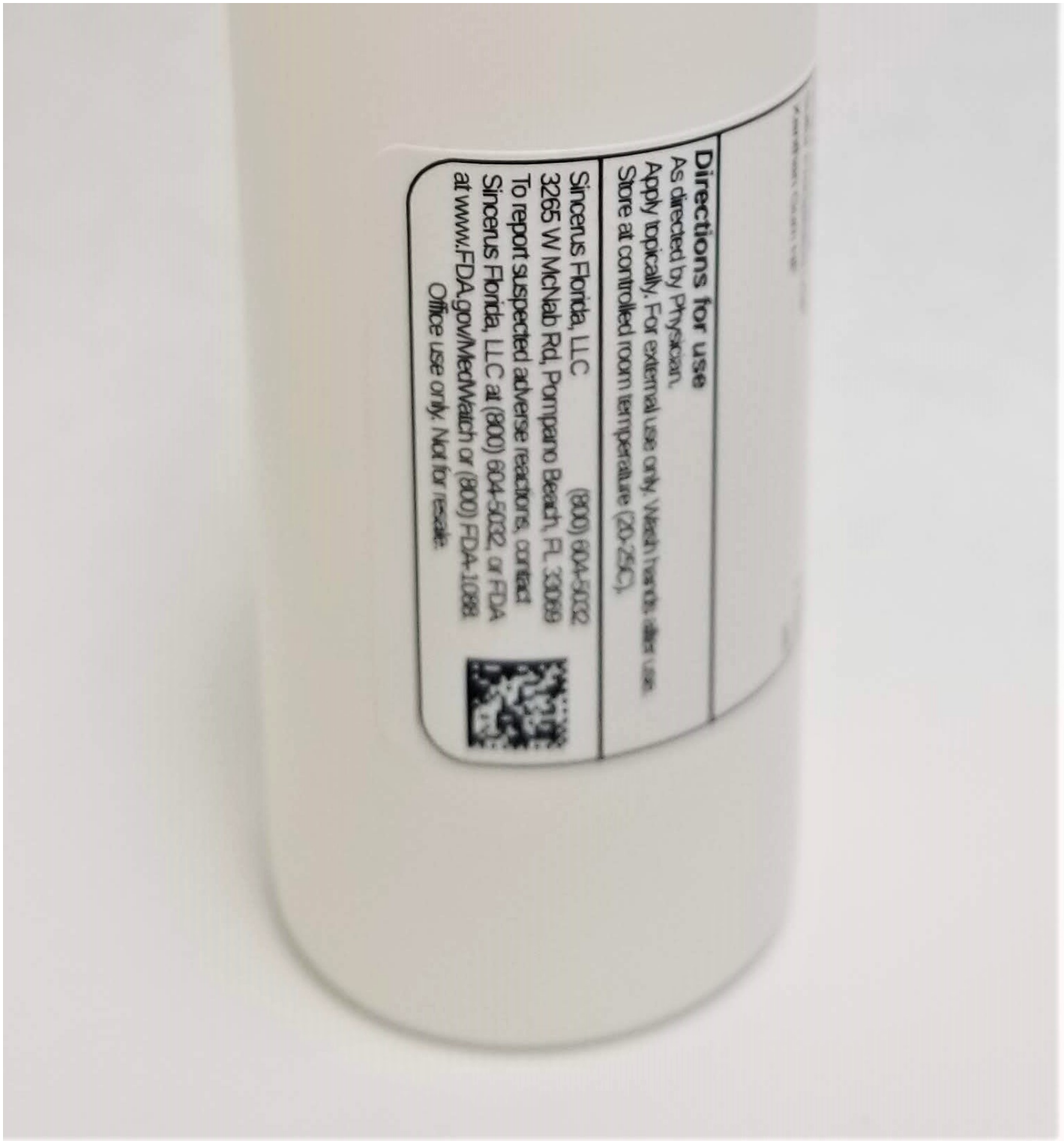
SALICYLIC ACID 2% / SODIUM SULFACETAMIDE MONOHYDRATE 8%

Directions for use

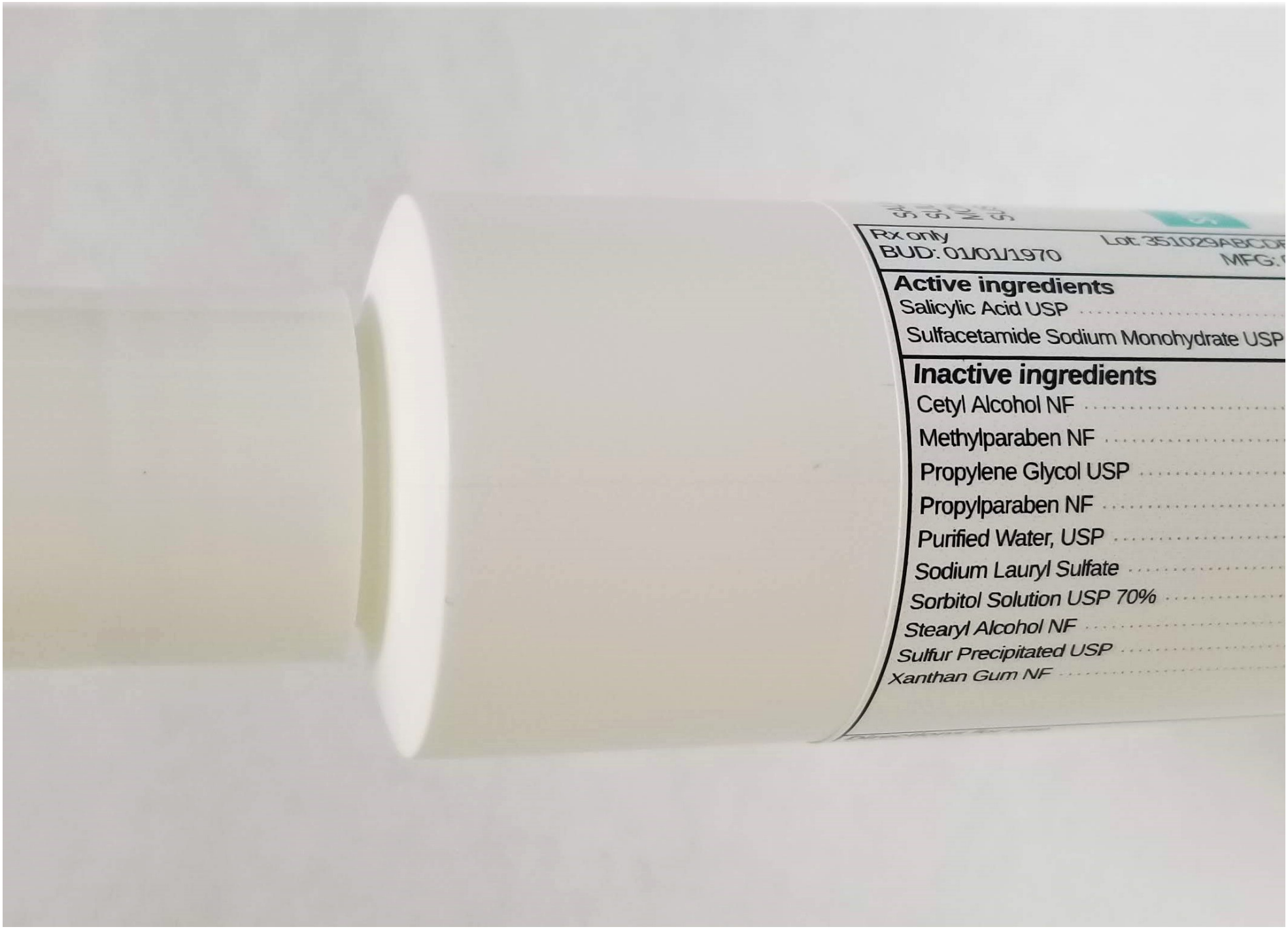


Sincerus Florida, LLC. Adverse reactions

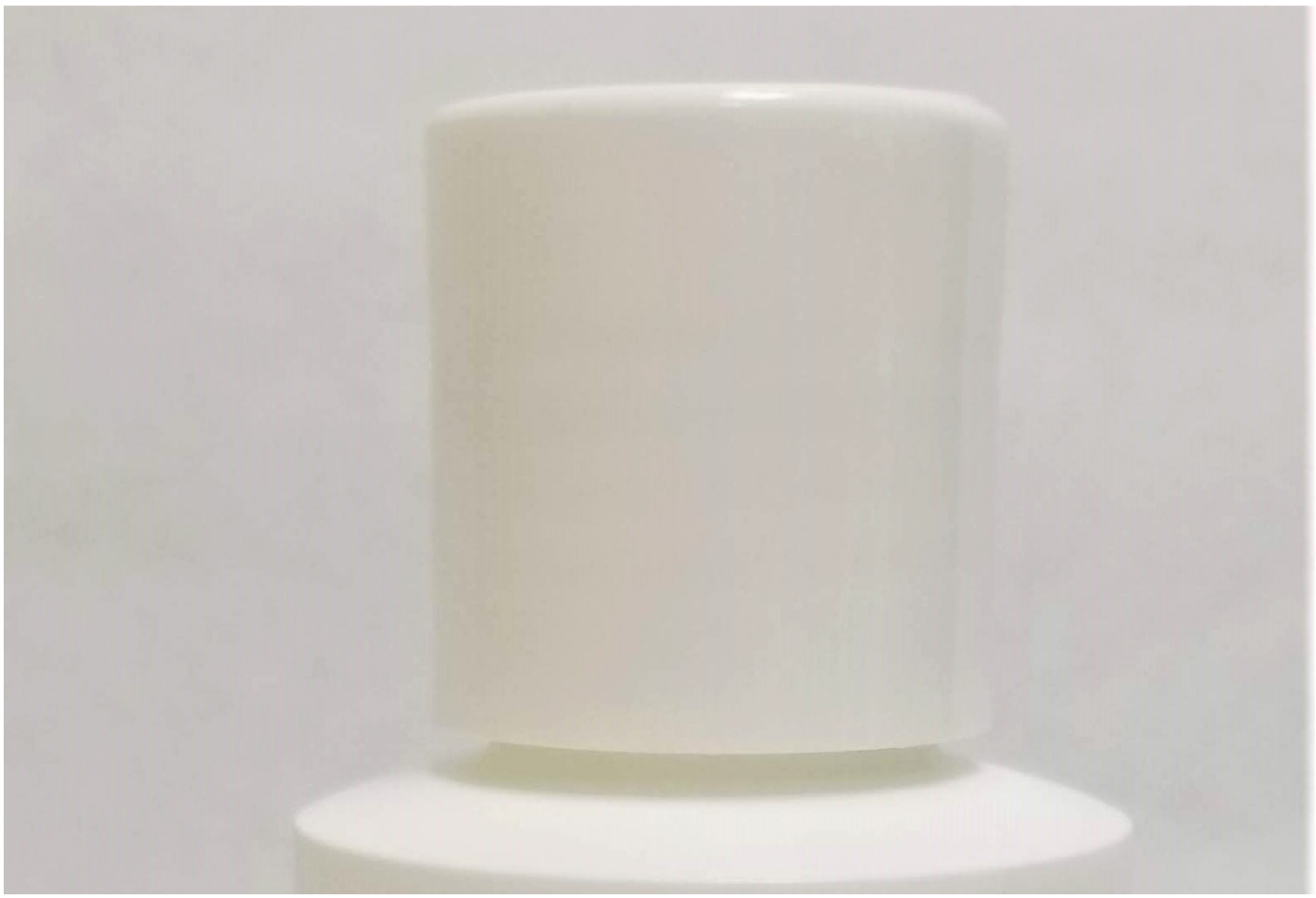

Directions for use
As directed by Physician.
Apply topically. For external use only. Wash hands after use.
Store at controlled room temperature (20-25C).
Sincerus Florida, LLC (800) 604-5032
3265 W McNab Rd, Pompano Beach, FL 33069
To report suspected adverse reactions, contact
Sincerus Florida, LLC at (800) 604-5032, or FDA
at www.FDA.gov/MedWatch or (800) FDA-1088.
Office use only. Not for resale.



Active, inactive


Rx only
BUD: 01/01/1970
Active ingredients
Salicylic Acid USP
Sulfacetamide Sodium Monohydrate USP
Inactive ingredients
Cetyl Alcohol NF
Methylparaben NF
Propylene Glycol USP
Propylparaben NF
Purified Water, USP
Sodium Lauryl Sulfate
Sorbitol Solution USP 70%
Stearyl Alcohol NF
Sulfur Precipitated USP
Xanthan Gum NF

NDC 72934- 8171-6 SALICYLIC ACID USP 2% / SODIUM SULFACETAMIDE MONOHYDRATE USP 8%.




Rx only
BUD: 90 days
NDC 72934-8171-6
SALICYLIC ACID USP 2%
SULFACETAMIDE SODIUM
MONOHYDRATE USP 8%
SUSPENSION 120gm
SINCERUS FLORIDA
This is a compounded drug.
Made in USA

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:72934-8171
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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72934-8171-6	120 g in 1 CYLINDER; 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(72934-8171)

Revised: 5/2019 Sincerus Florida, LLC