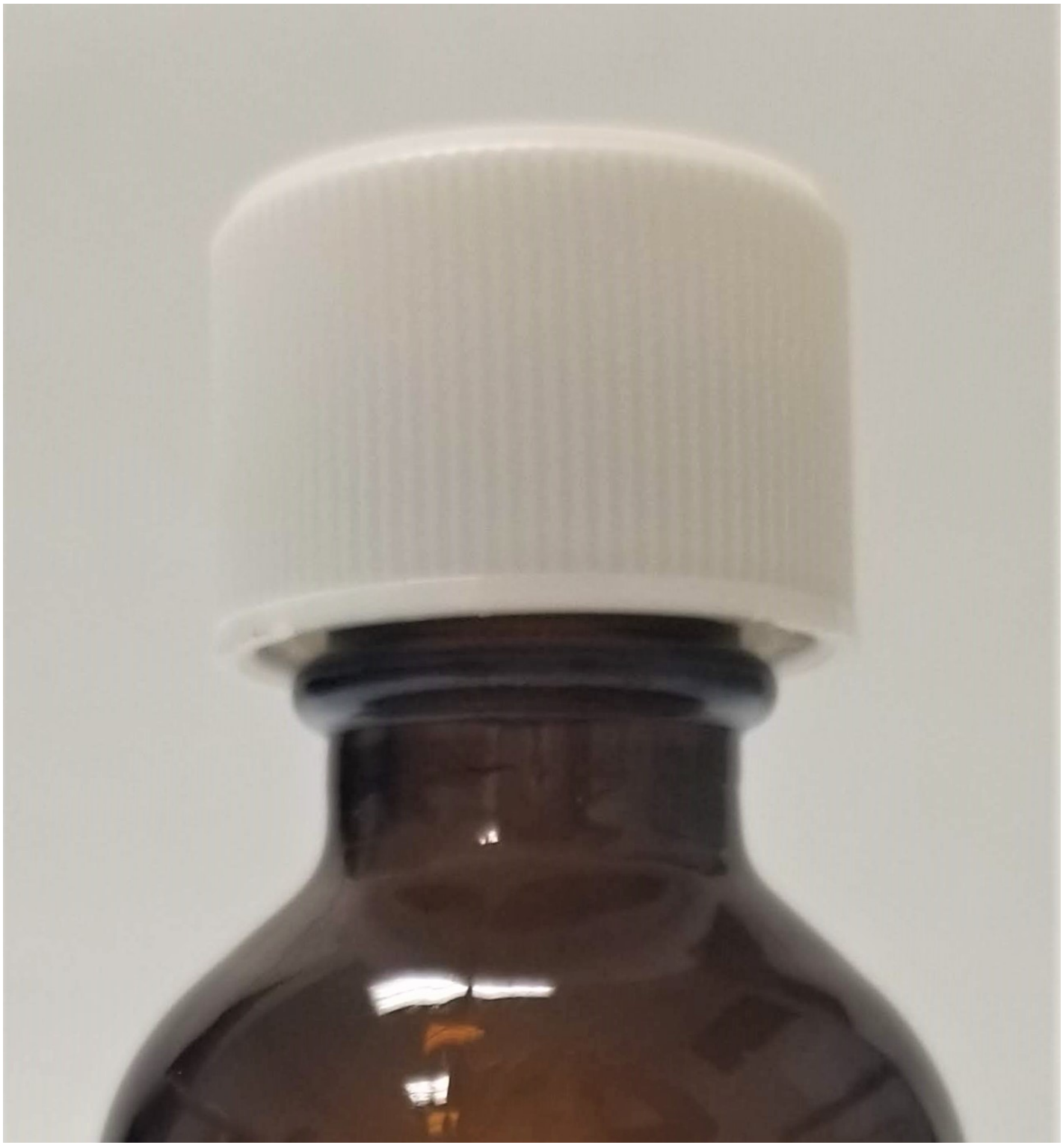
BETAMETHASONE DIPROPIONATE 0.05% / MINOXIDIL 5%- betamethasone dipropionate 0.05% / minoxidil 5% solution
Sincerus Florida, LLC

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.

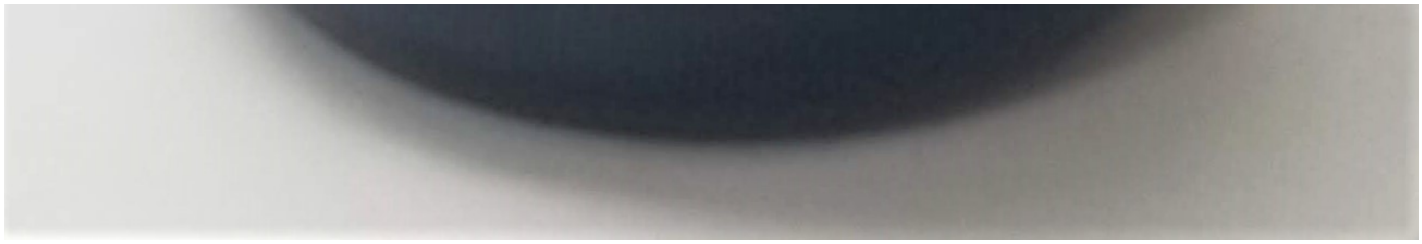
BETAMETHASONE DIPROPIONATE 0.05% / MINOXIDIL 5%

Directions for use

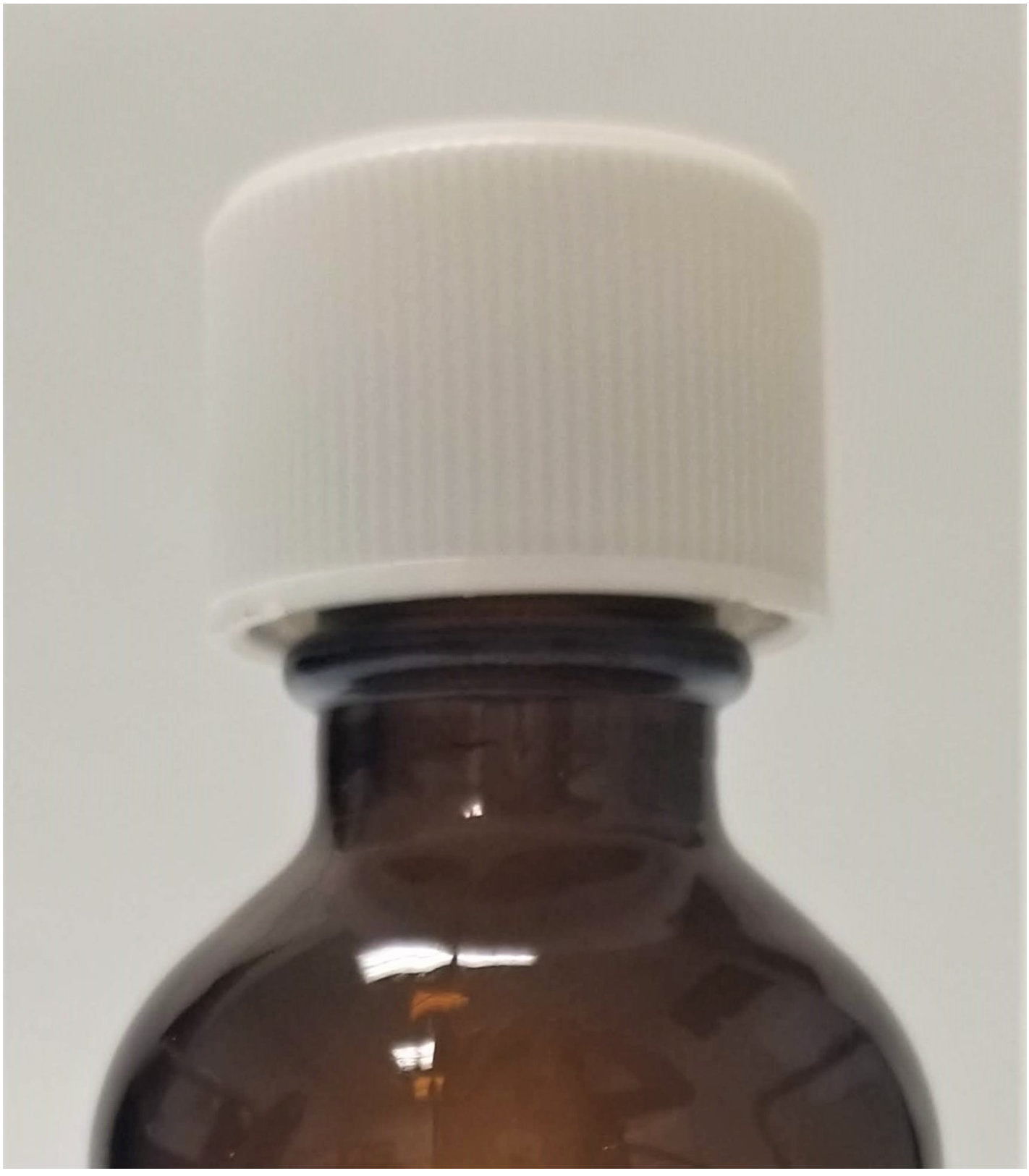


Directions for use
As directed by Physician.
Apply topically. For external use only. Wash hands after use.
Store at controlled room temperature (20-25C).

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.



Sincerus Florida, LLC adverse reactions.



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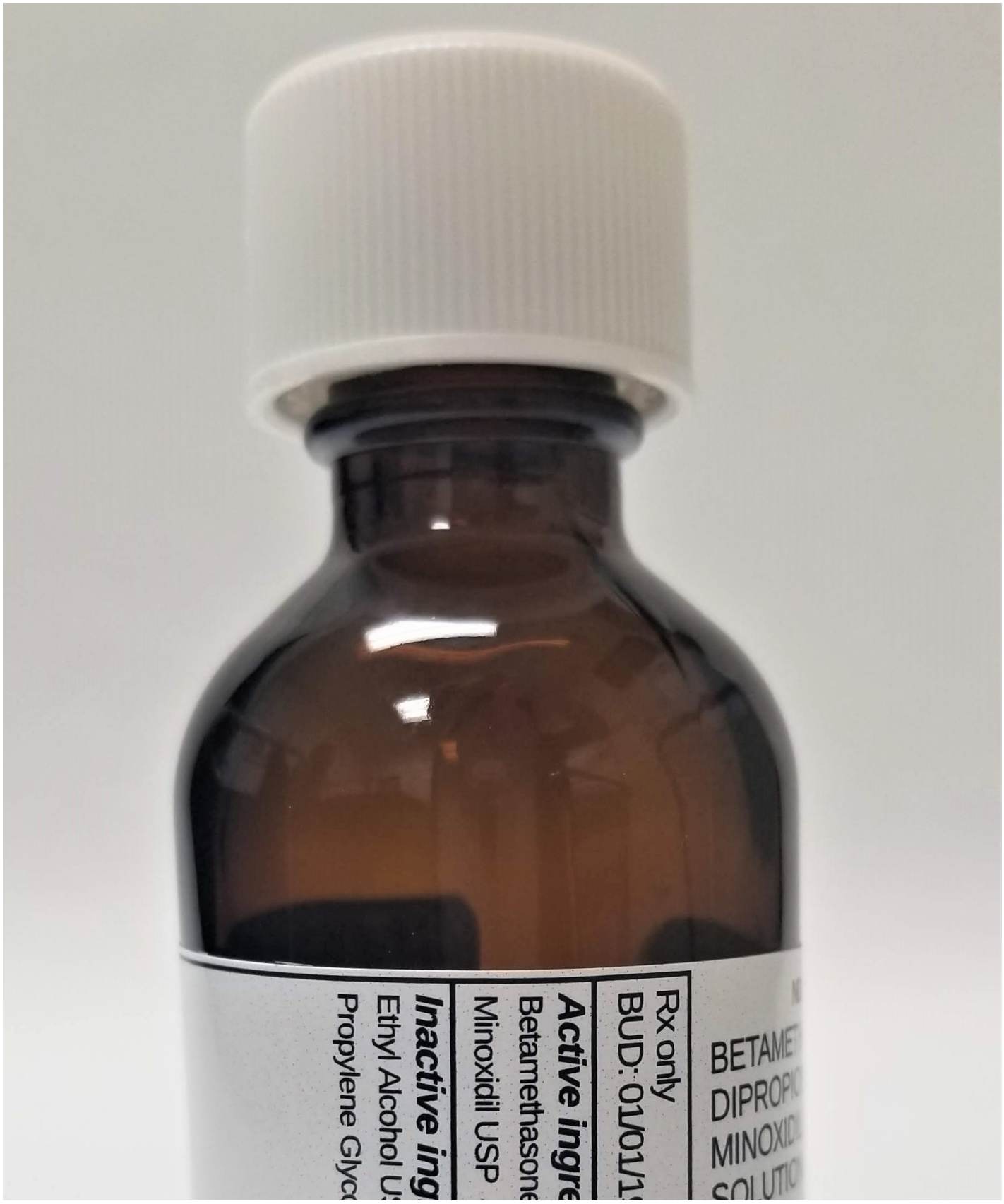
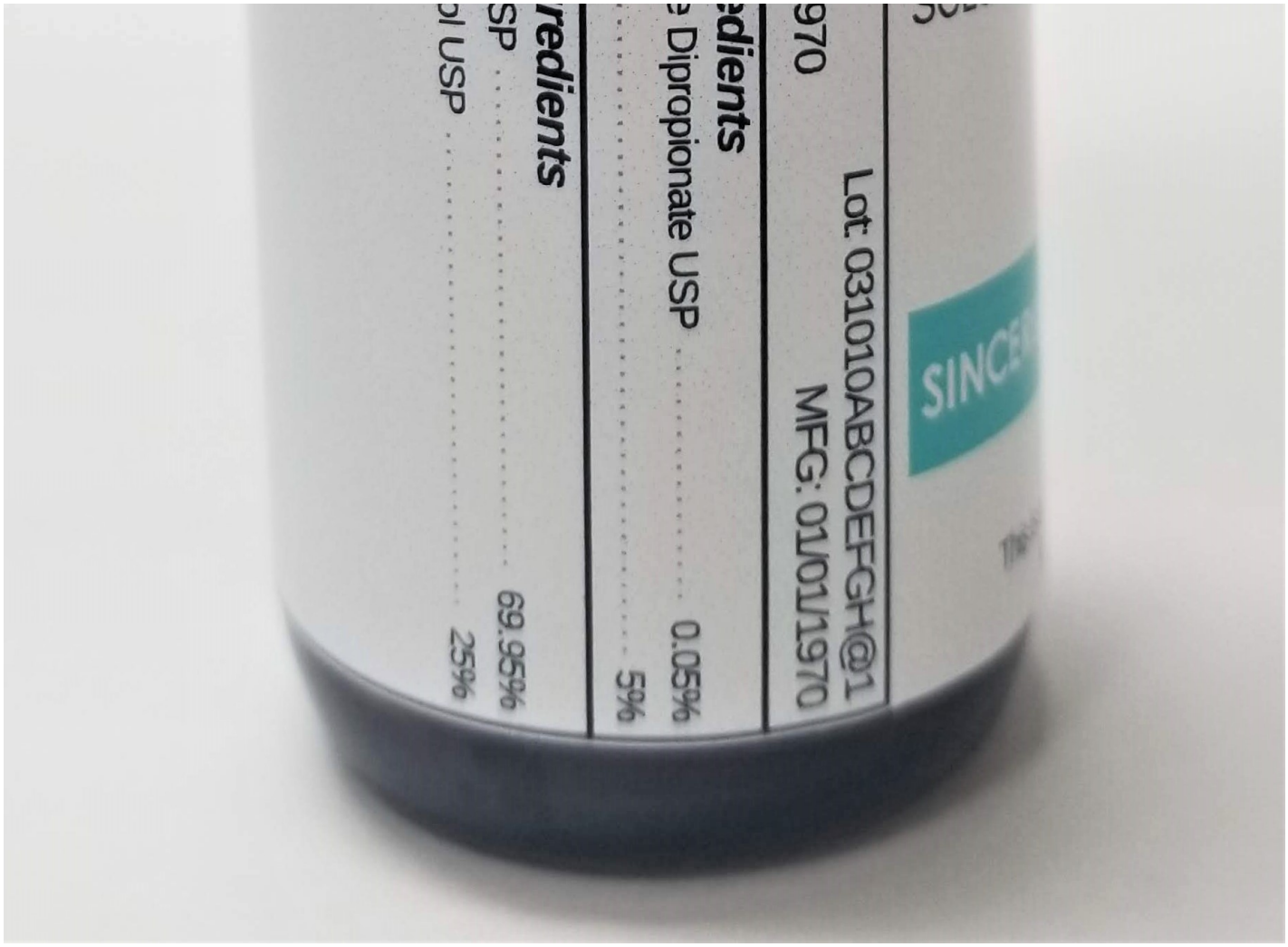
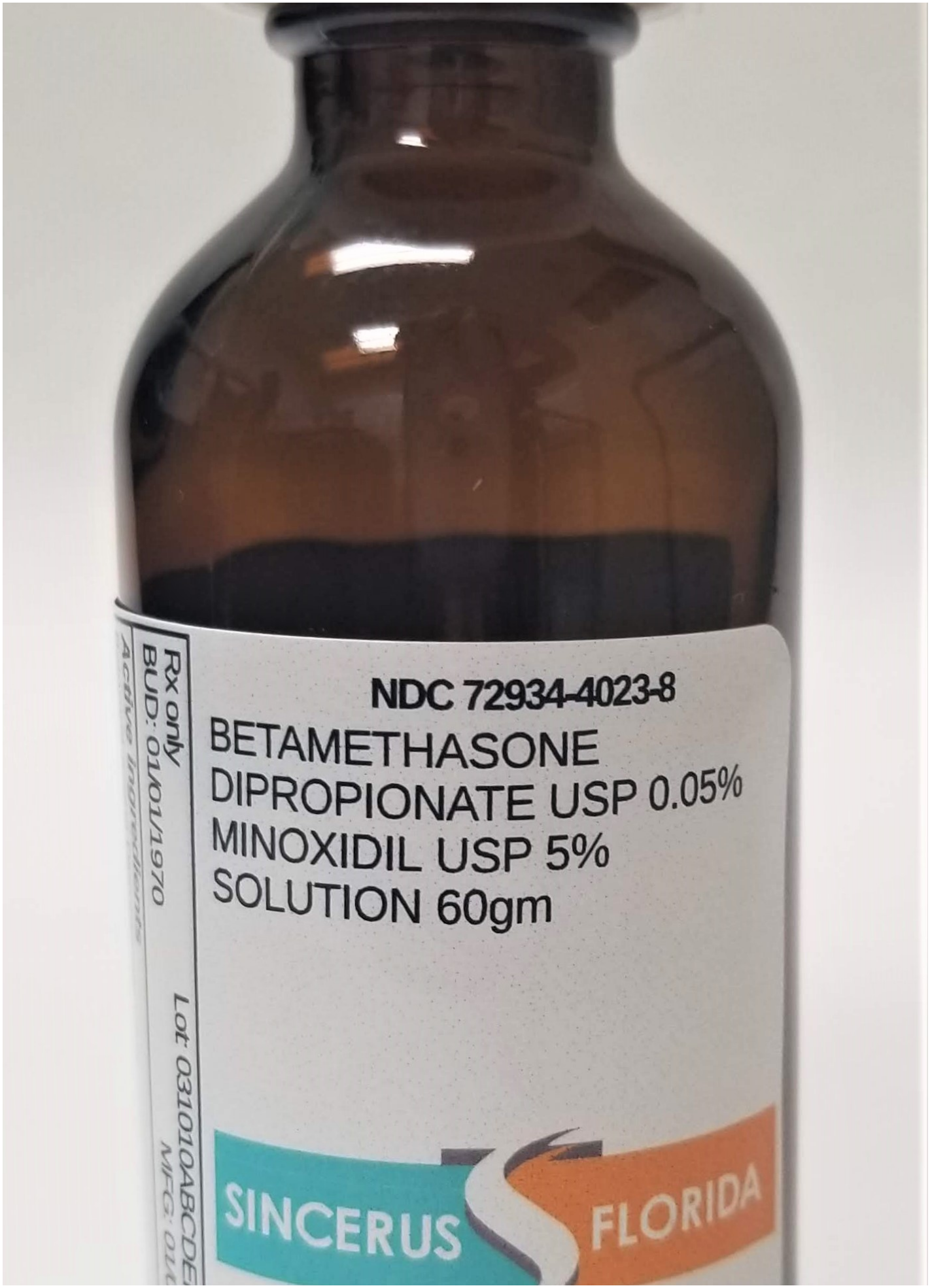
Active, inactive





NDC 72934-4023-8
BETAMETHASONE DIPROPIONATE 0.05% / MINOXIDIL 5%
Solution 60gm


NDC 72934-4023-8
BETAMETHASONE
DIPROPIONATE USP 0.05%
MINOXIDIL USP 5%
SOLUTION 60gm
Rx only
BUD: 01/01/1970
Lot: 031010ABCDE
SINCERUS
FLORIDA

BETAMETHASONE DIPROPIONATE 0.05% / MINOXIDIL 5%

betamethasone dipropionate 0.05% / minoxidil 5% solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-4023
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MINOXIDIL (UNII: 5965120SH1) (MINOXIDIL - UNII:5965120SH1)	MINOXIDIL	5 g in 100 g
BETAMETHASONE DIPROPIONATE (UNII: 826Y60901U) (BETAMETHASONE - UNII:9842X06Q6M)	BETAMETHASONE	0.05 g in 100 g

Product Characteristics			
Color	white (CLEAR SOLUTION)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72934-4023-8	60 g in 1 BOTTLE, GLASS; 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(72934-4023)