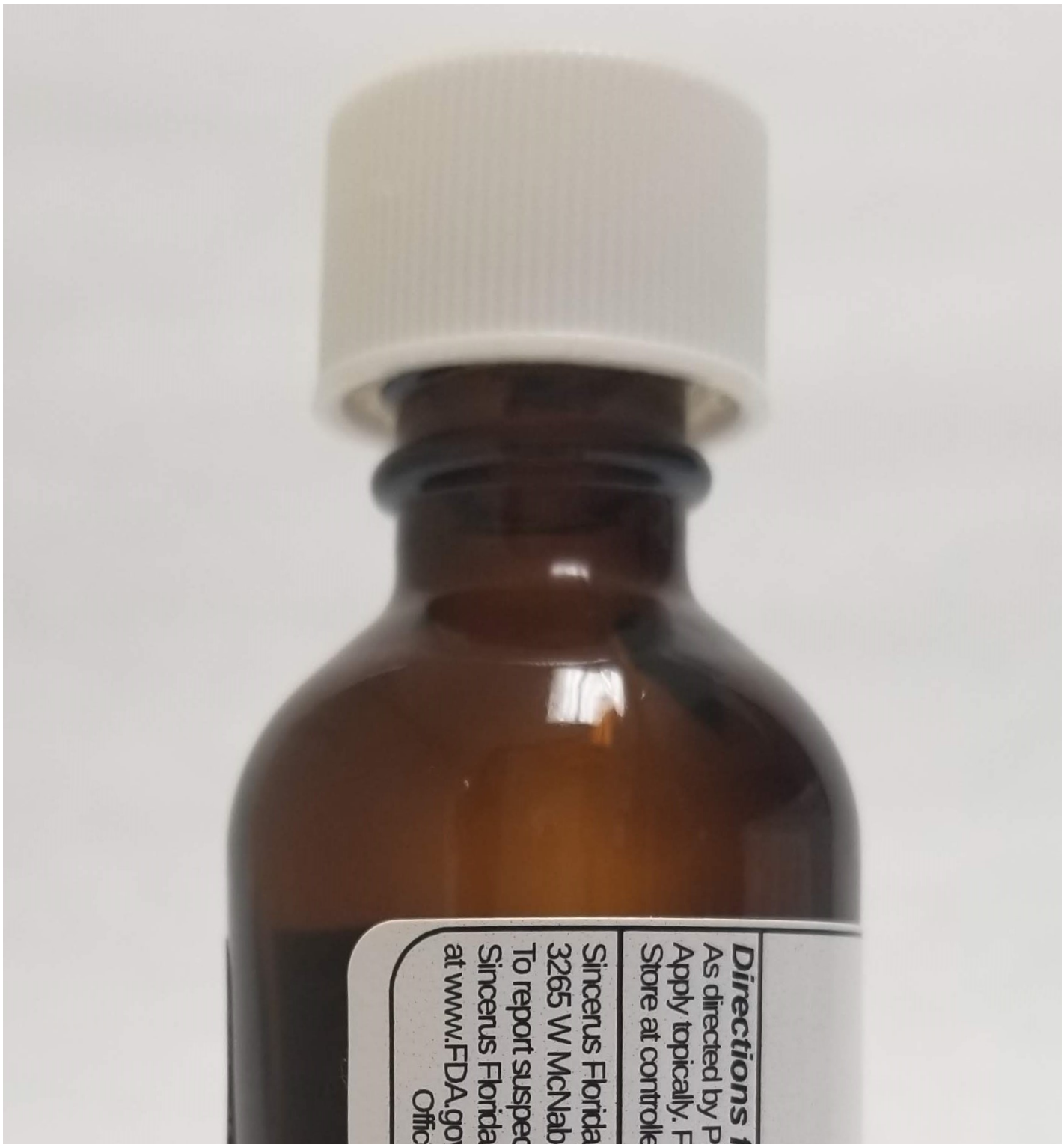
DEXAMETHASONE SODIUM PHOSPHATE 0.1% / FINASTERIDE 0.1% / MINOXIDIL 5% - dexamethasone sodium phosphate 0.1% / finasteride 0.1% / minoxidil 5% solution
Sincerus Florida, LLC

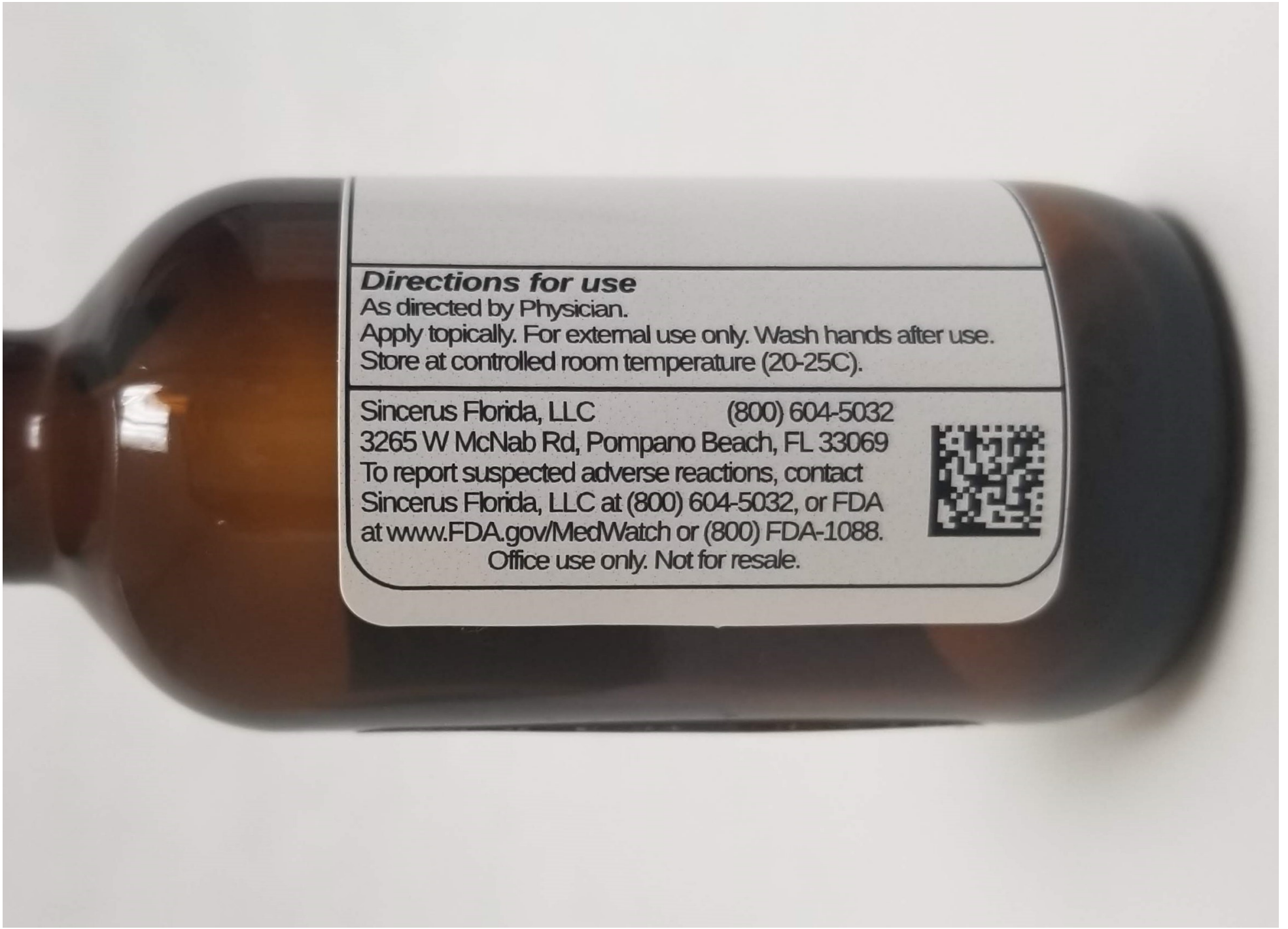
Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

DEXAMETHASONE SODIUM PHOSPHATE 0.1% / FINASTERIDE 0.1% / MINOXIDIL 5%

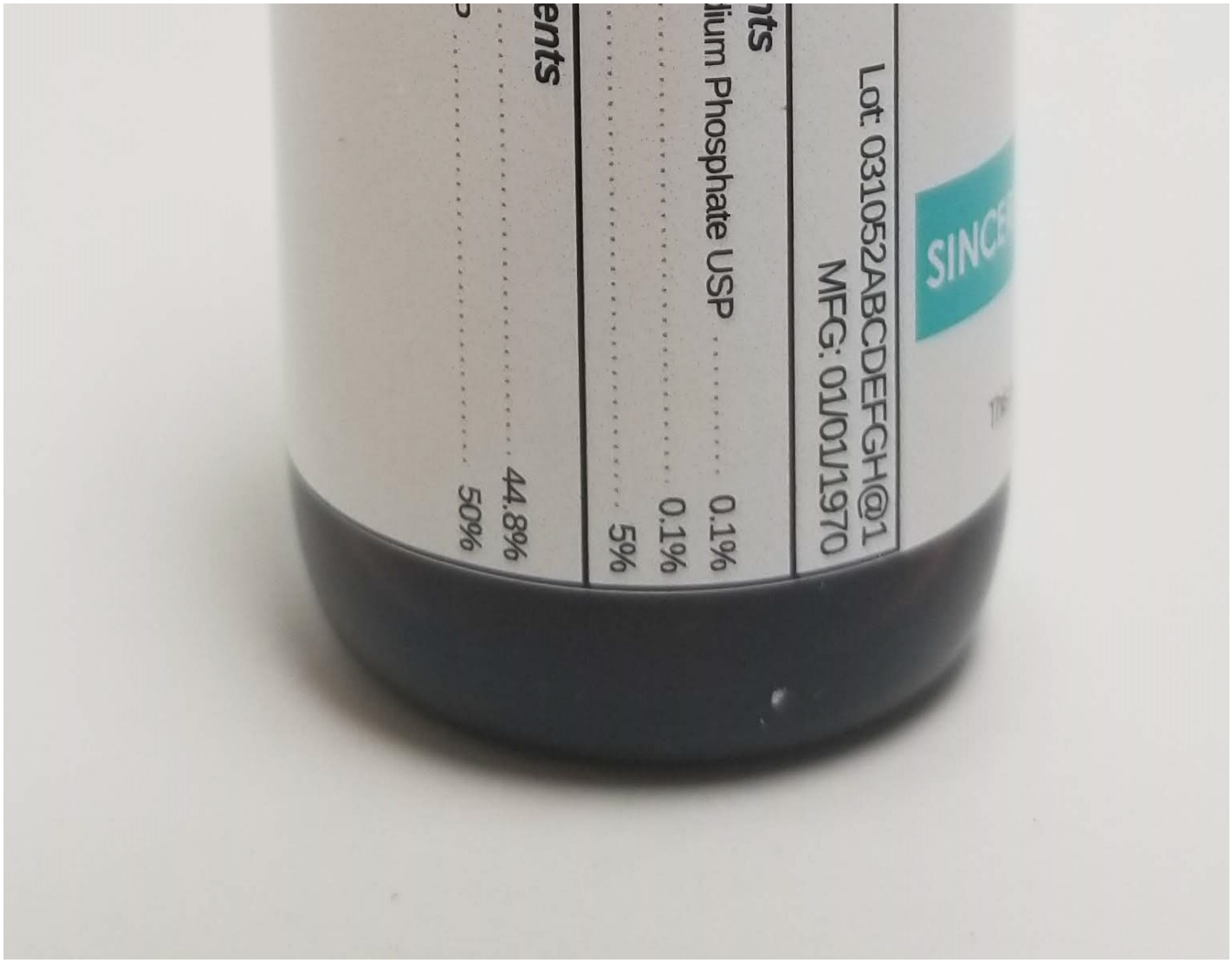
Directions for use



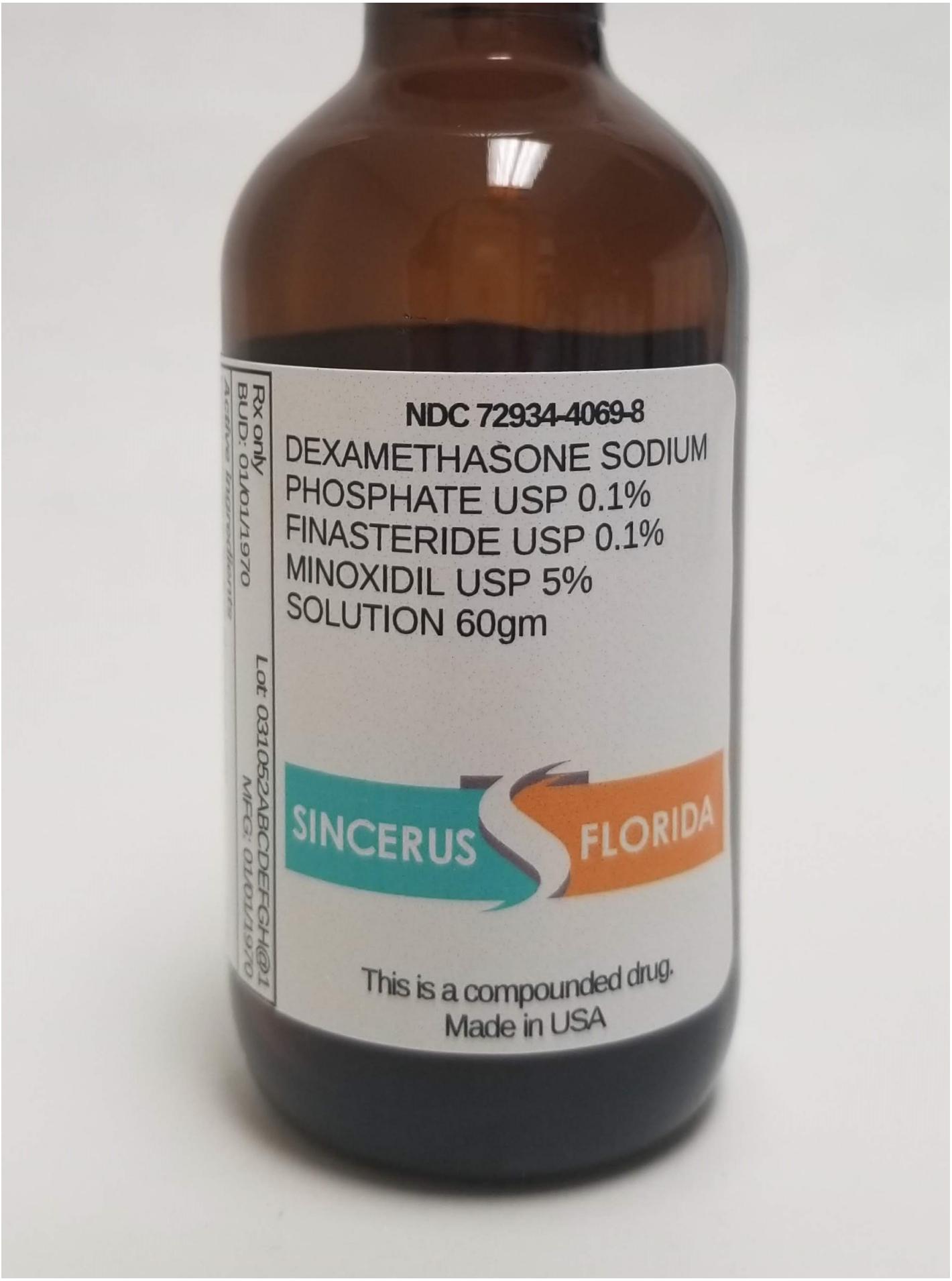
Sincerus Florida, LLC. Adverse reactions


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



NDC 72934-4069-8 DEXAMETHASONE SODIUM PHOSPHATE USP 0.1% / FINASTERIDE USP 0.1% / MINOXIDIL USP 5% . Solution 60 gm


NDC 72934-4069-8
DEXAMETHASONE SODIUM PHOSPHATE USP 0.1%
FINASTERIDE USP 0.1%
MINOXIDIL USP 5%
SOLUTION 60gm
Rx only
BUD: 01/01/1970
Active Ingredients
Lot: 031052ABCDEFGH@1
MFG: 01/01/1970
SINCERUS FLORIDA
This is a compounded drug.
Made in USA

DEXAMETHASONE SODIUM PHOSPHATE 0.1% / FINASTERIDE 0.1% / MINOXIDIL 5%

dexamethasone sodium phosphate 0.1% / finasteride 0.1% / minoxidil 5% solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-4069
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
FINASTERIDE (UNII: 57GNO57U7G) (FINASTERIDE - UNII:57GNO57U7G)	FINASTERIDE	0.1 g in 100 g
DEXAMETHASONE SODIUM PHOSPHATE (UNII: AI9376Y64P) (DEXAMETHASONE - UNII:7S5I7G3JQL)	DEXAMETHASONE PHOSPHATE	0.1 g in 100 g

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72934-4069-8	60 g in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	05/09/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/09/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-4069)

Revised: 5/2019 Sincerus Florida, LLC