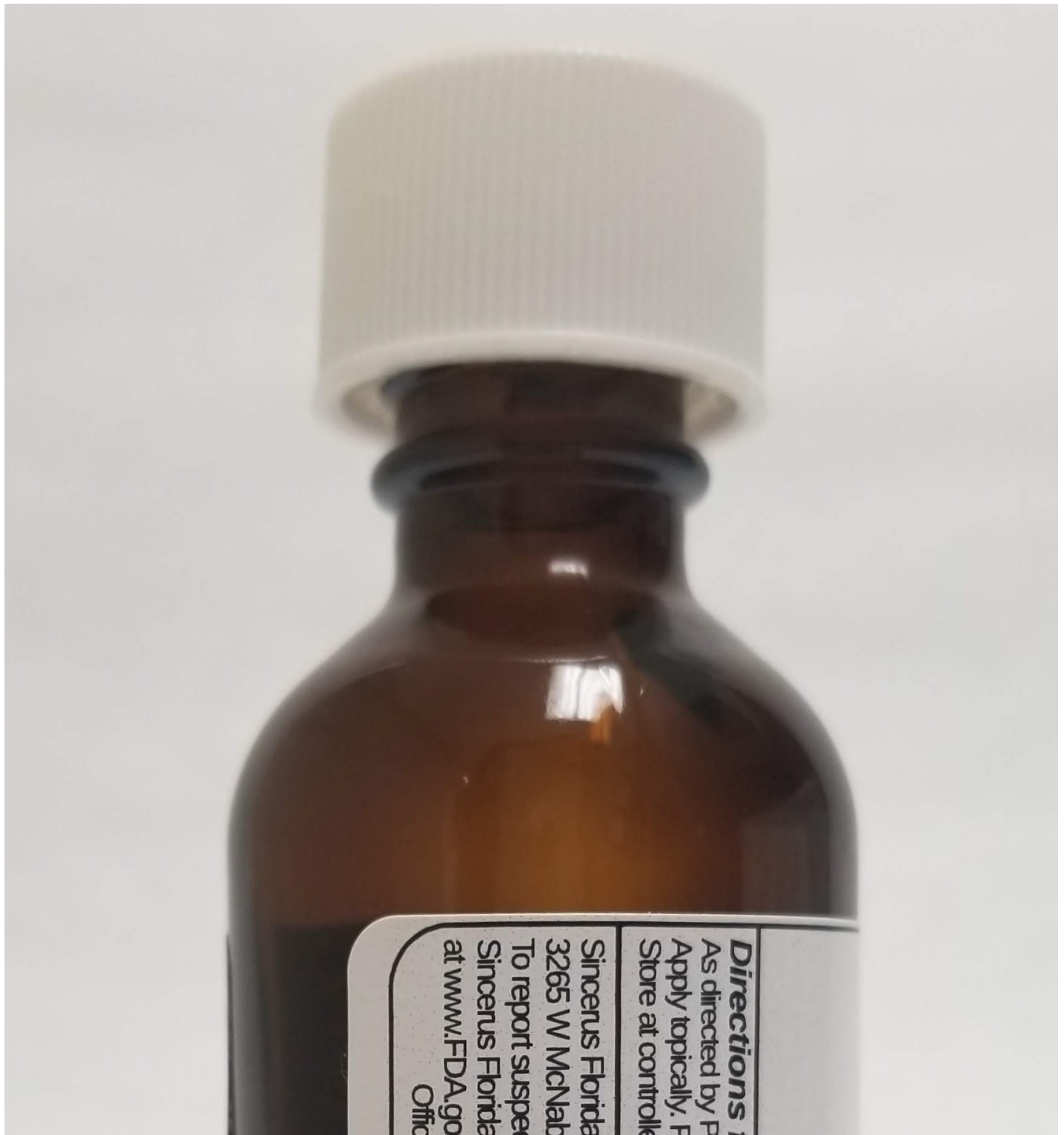


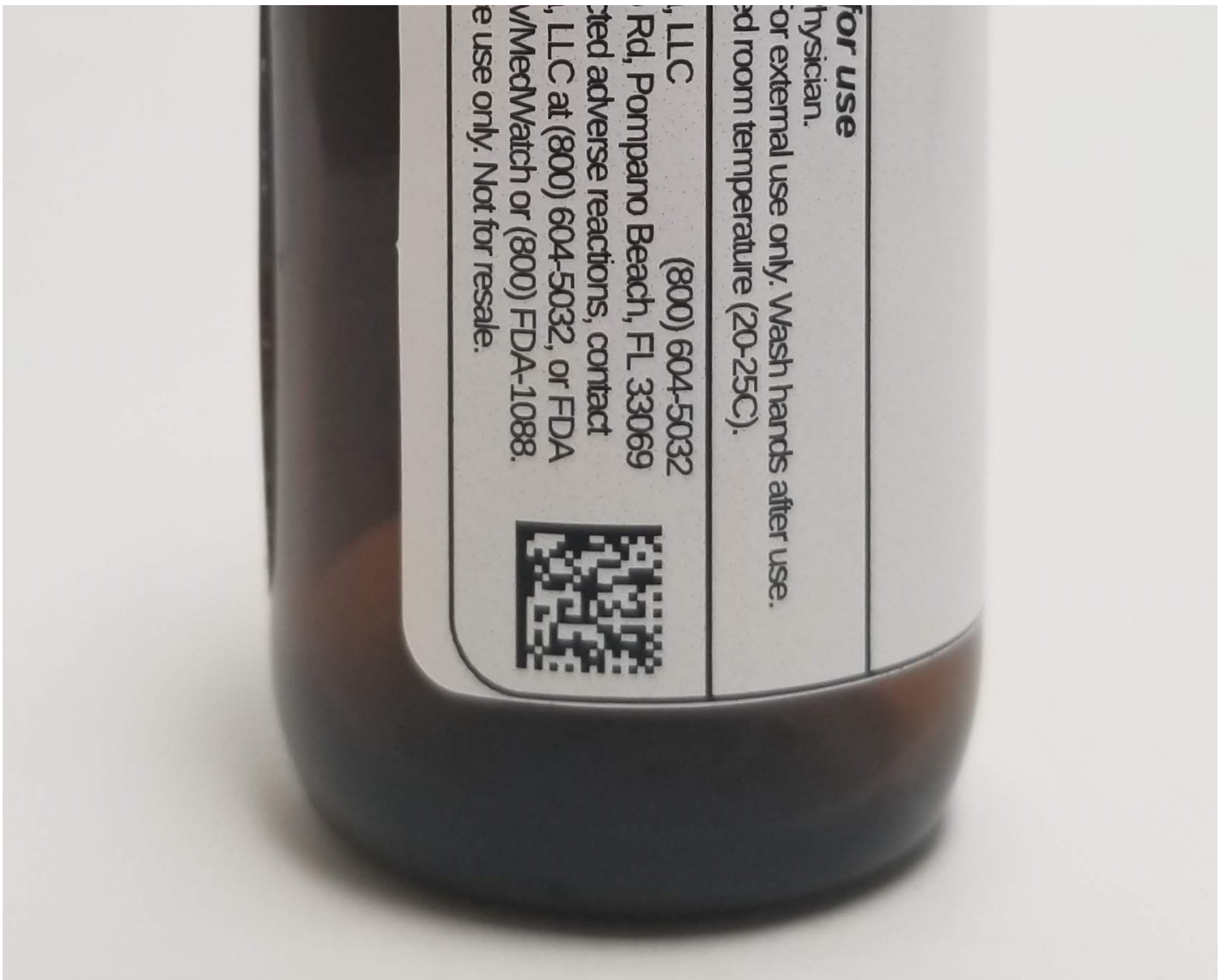
**MINOXIDIL 5% / PROGESTERONE 0.1% / TRETINOIN 0.025% - minoxidil 5% /
progesterone 0.1% / tretinoin 0.025% solution**
Sincerus Florida, LLC

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](#).

MINOXIDIL 5% / PROGESTERONE 0.1% / TRETINOIN 0.025%

Directions for use





Sincerus Florida. Adverse reactions



Directions for use

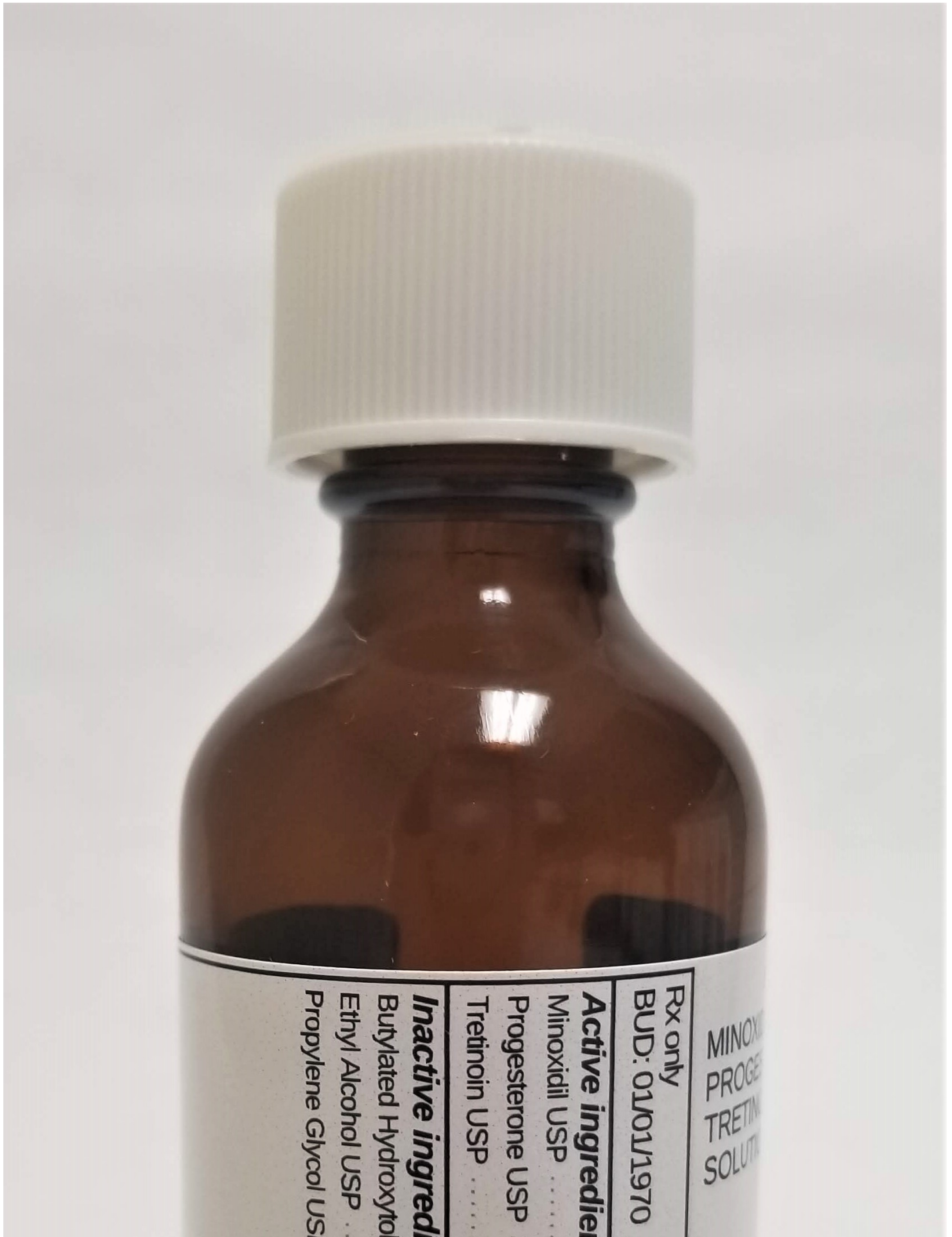
As directed by Physician.
Apply topically. For external use only. Wash hands after use.
Store at controlled room temperature (20-25C).

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



MINOXIDIL
PROGESTERONE
TRETINOIN
SOLUTION

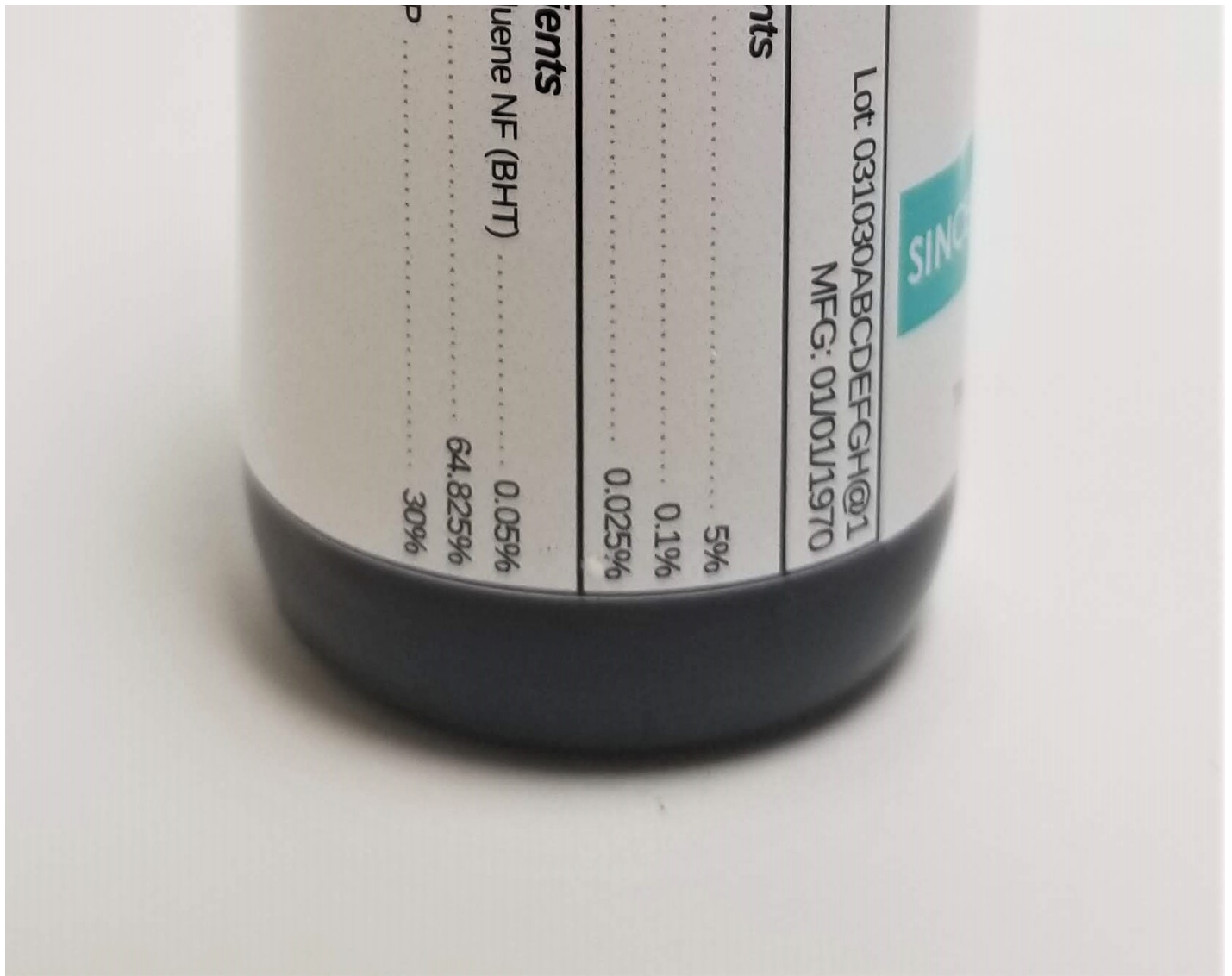
Rx only
BUD: 01/01/1970

Active ingredients

Minoxidil USP
Progesterone USP
Tretinoin USP

Inactive ingredients

Butylated Hydroxytol
Ethyl Alcohol USP ..
Propylene Glycol US



NDC 72934-4146-8 MINOXIDIL USP 5% / PROGESTERONE USP 0.1% / TRETINOIN USP 0.025%. Solution 600 gm



NDC 72934-4146-8

MINOXIDIL USP 5%
PROGESTERONE USP 0.1%
TRETINOIN USP 0.025%
SOLUTION 60gm

Rx only
BUD: 01/01/1970

Lot: 031030ABCDEF@1
MFG: 01/01/1970



This is a compounded drug.
Made in USA

MINOXIDIL 5% / PROGESTERONE 0.1% / TRETINOIN 0.025%

minoxidil 5% / progesterone 0.1% / tretinoin 0.025% solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-4146
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
PROGESTERONE (UNII: 4G7DS2Q64Y) (PROGESTERONE - UNII:4G7DS2Q64Y)	PROGESTERONE	0.1 g in 100 g
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.025 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:72934-4146-8	60 g in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	05/07/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/07/2019	

Labeler - Sincerus Florida, LLC (080105003)**Establishment**

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-4146)