

**FINASTERIDE 0.1% / MINOXIDIL 5% - finasteride 0.1% / minoxidil 5% solution**  
**Sincerus Florida, LLC**

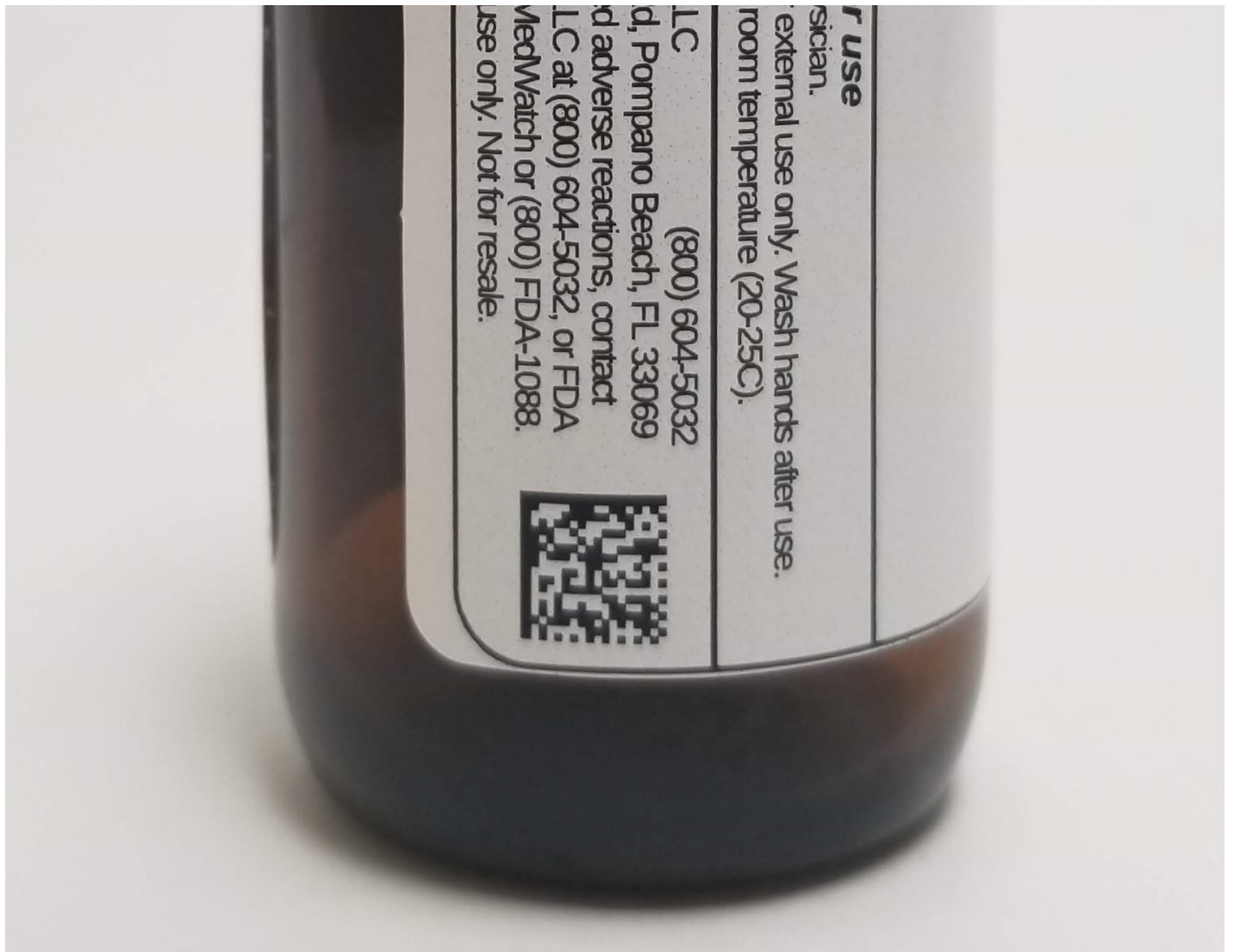
*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click [here](#).*

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**FINASTERIDE 0.1% / MINOXIDIL 5%**

**Directions for use**





**Sincerus Florida, LLC**



**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](http://www.FDA.gov/MedWatch) or (800) FDA-1088.

Office use only. Not for resale.





Active, inactive



FINASTERIDE  
MINOXIDIL  
SOLUTION

Rx only  
BUD: 01/01/1970

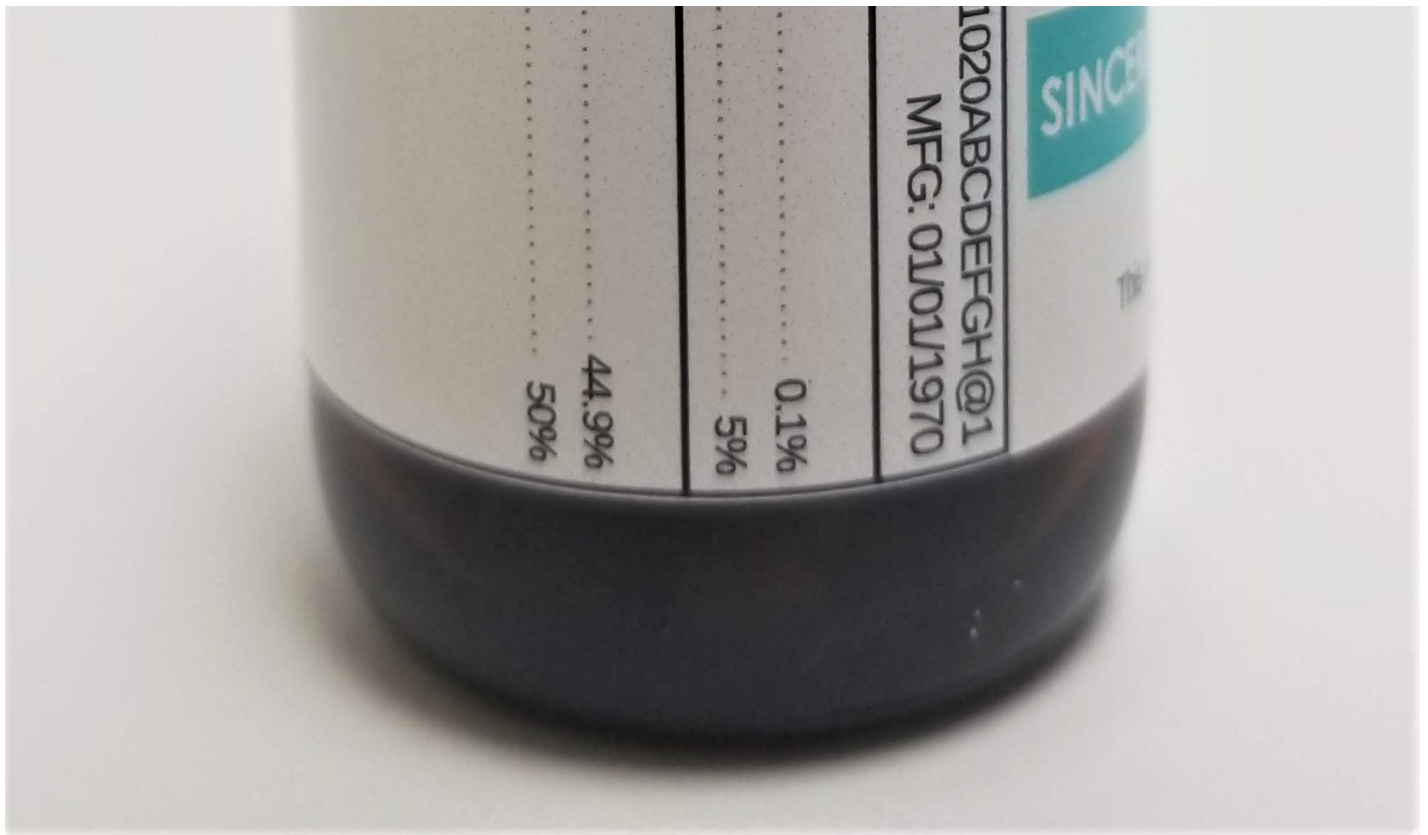
Lot: 03

**Active ingredients**

Finasteride USP .....  
Minoxidil USP .....

**Inactive ingredients**

Ethyl Alcohol USP .....  
Propylene Glycol USP .....



**NDC 72934-4074-8. FINASTERIDE USP 0.1% / MINOXIDIL USP 5%. Solution 60 gm**



NDC 72934-4074-8

FINASTERIDE USP 0.1%  
MINOXIDIL USP 5%  
SOLUTION 60gm

Rx only  
BUD: 01/01/1970

Lot: 031020ABCDEF@1  
MFG: 01/01/1970

SINCERUS

FLORIDA

This is a compounded drug.  
Made in USA



## FINASTERIDE 0.1% / MINOXIDIL 5%

finasteride 0.1% / minoxidil 5% solution

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-4074
<b>Route of Administration</b>	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

### Product Characteristics

<b>Color</b>	white (clear solution)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-4074-8	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(72934-4074)