

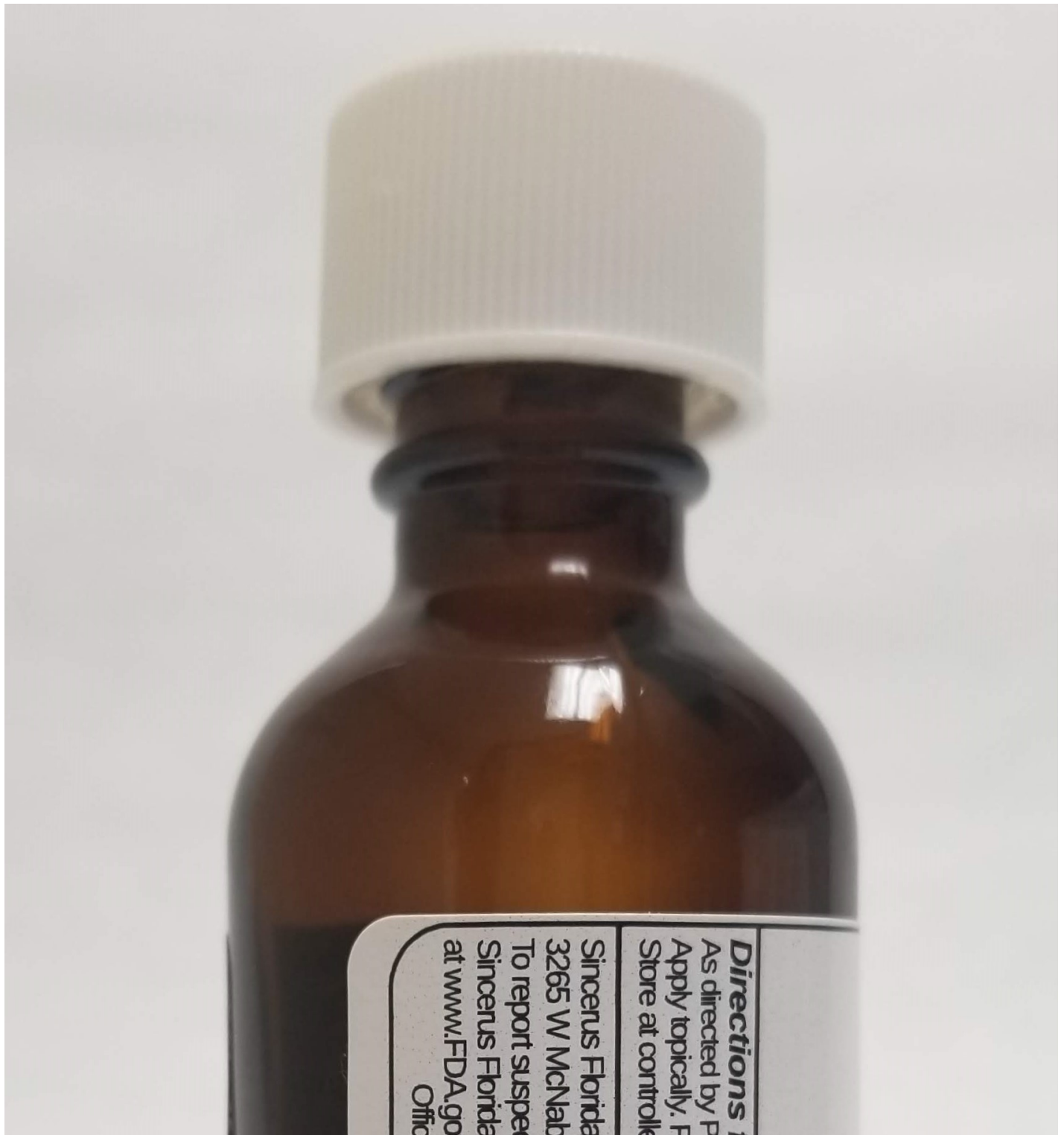
**FLUOCINOLONE ACETONIDE 0.01% / MINOXIDIL 7% / PROGESTERONE 0.1% -  
fluocinolone acetonide 0.01% / minoxidil 7% / progesterone 0.1% solution**

**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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**FLUOCINOLONE ACETONIDE 0.01% / MINOXIDIL 7% / PROGESTERONE 0.1%**

**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](http://www.FDA.gov/MedWatch) or (800) FDA-1088.  
Office use only. Not for resale.



Active, inactive



NOI  
FLUOCINON  
USP 0.01%  
MINOXIDIL  
PROGESTE  
SOLUTION

Rx only  
BUD: 01/01/1970

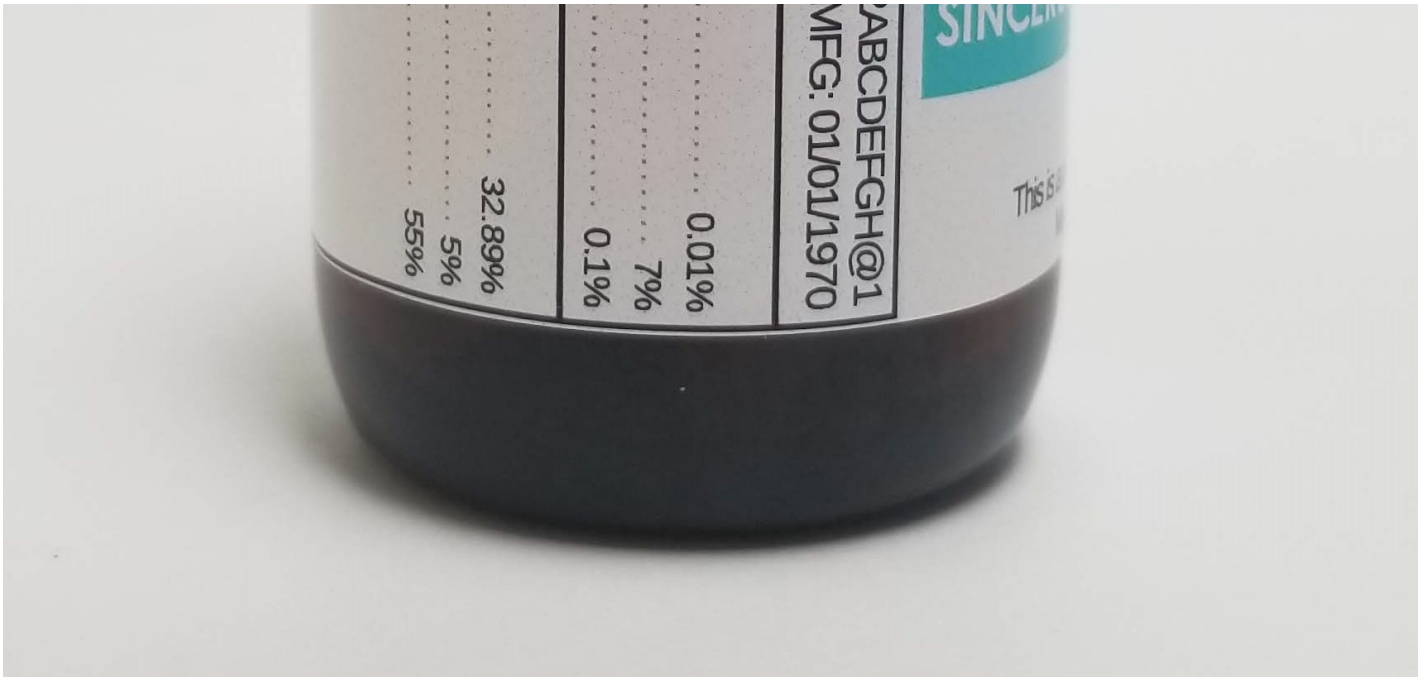
Lot: 031092

**Active ingredients**

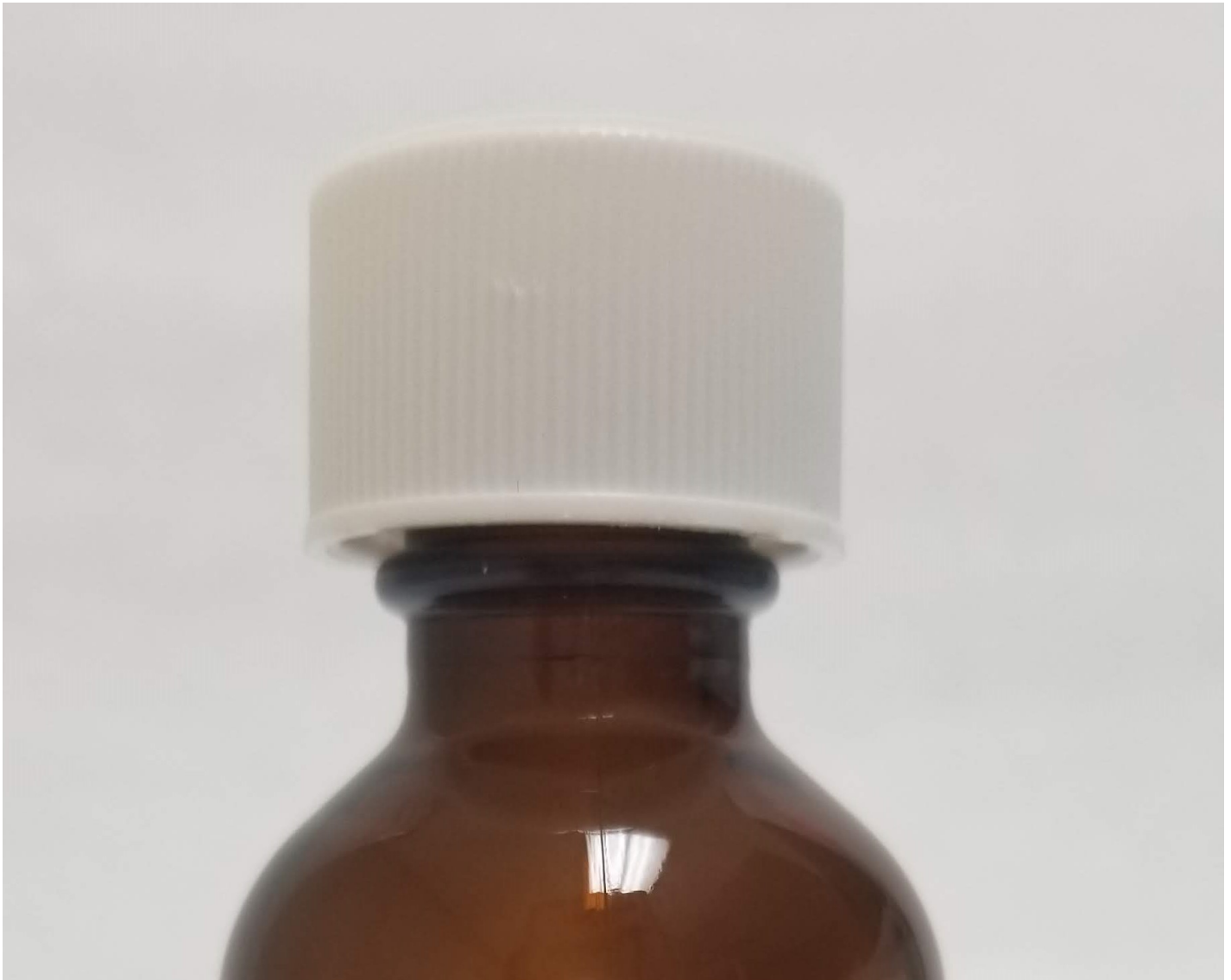
Fluocinolone Acetonide USP .....  
Minoxidil USP .....  
Progesterone USP .....

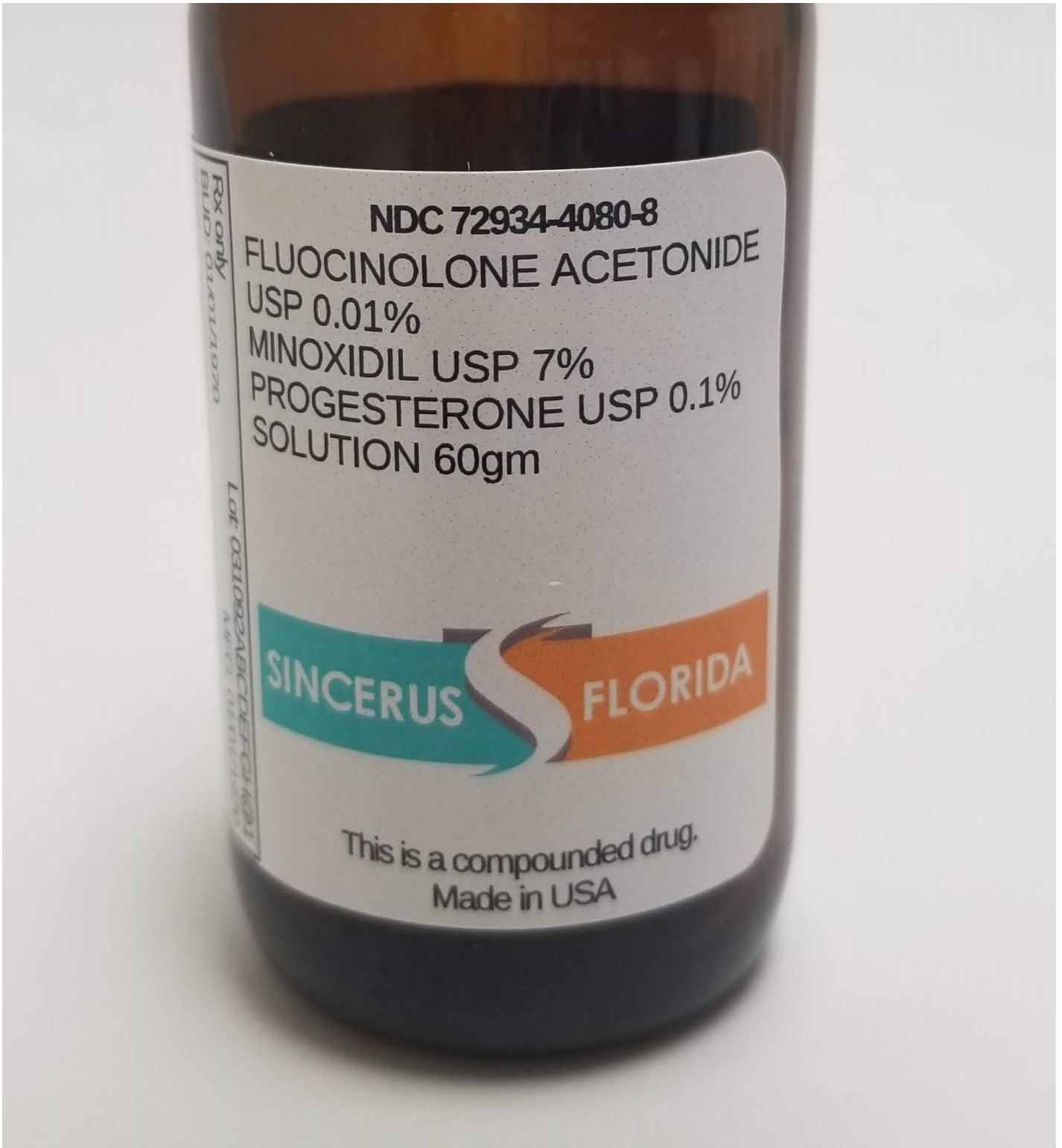
**Inactive ingredients**

Ethyl Alcohol USP .....  
Lactic Acid USP .....  
Propylene Glycol USP .....



**NDC 72934-4080-8. FLUOCINOLONE ACETONIDE USP 0.01% / MINOXIDIL USP 7% /  
PROGESTERONE USP 0.1%. Solution 60 gm.**





**FLUOCINOLONE ACETONIDE 0.01% / MINOXIDIL 7% / PROGESTERONE 0.1%**

fluocinolone acetonide 0.01% / minoxidil 7% / progesterone 0.1% solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-4080
Route of Administration	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>FLUOCINOLONE ACETONIDE</b> (UNII: 0CD5FD6S2M) (FLUOCINOLONE ACETONIDE - UNII:0CD5FD6S2M)	FLUOCINOLONE ACETONIDE	0.01 g in 100 g
<b>MINOXIDIL</b> (UNII: 5965120SH1) (MINOXIDIL - UNII:5965120SH1)	MINOXIDIL	7 g in 100 g
<b>PROGESTERONE</b> (UNII: 4G7DS2Q64Y) (PROGESTERONE - UNII:4G7DS2Q64Y)	PROGESTERONE	0.1 g in 100 g

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:72934-4080-8	60 g in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	05/09/2019	

**Marketing Information**

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

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

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Sincerus Florida, LLC		080105003	manufacture(72934-4080)

Revised: 5/2019

Sincerus florida, LLC