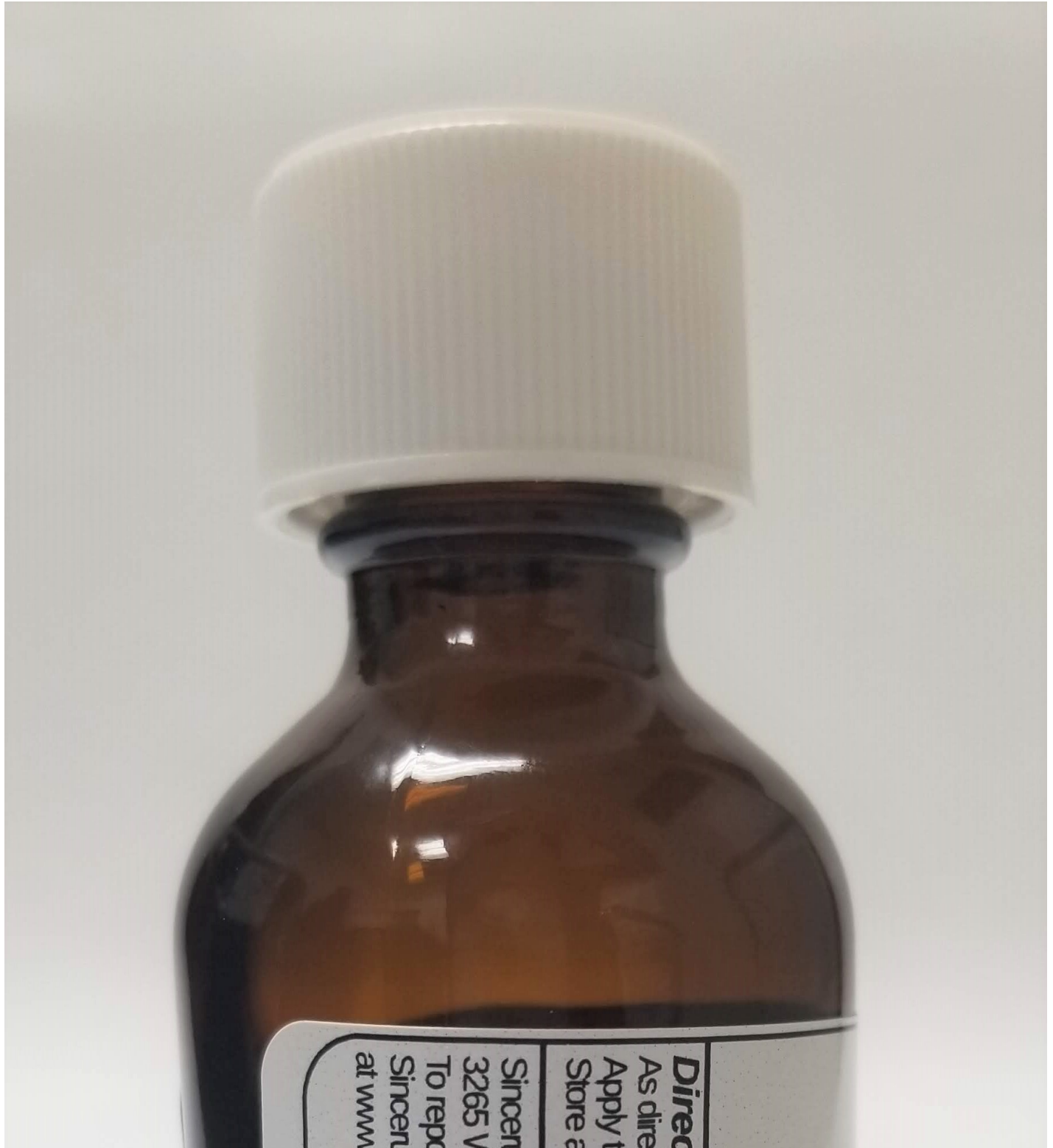


MINOXIDIL 7% / PROGESTERONE 0.1% - minoxidil 7% / progesterone 0.1% solution
Sincerus Florida, LLC

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](#).

MINOXIDIL 7% / PROGESTERONE 0.1%

Directions for use



Instructions for use

ected by Physician.

opically. For external use only. Wash hands after use.
at controlled room temperature (20-25C).

as Florida, LLC (800) 604-5032

V McNab Rd, Pompano Beach, FL 33069

rt suspected adverse reactions, contact

as Florida, LLC at (800) 604-5032, or FDA

/FDA.gov/MedWatch or (800) FDA-1088.

Office use only. Not for resale.



Sincerus Florida, LLC adverse reactions.



MINOXIDIL
PROGESTERONE
SOLUTION

Rx only

Lot 03

BUD: 01/01/1970

Active ingredients

Minoxidil USP

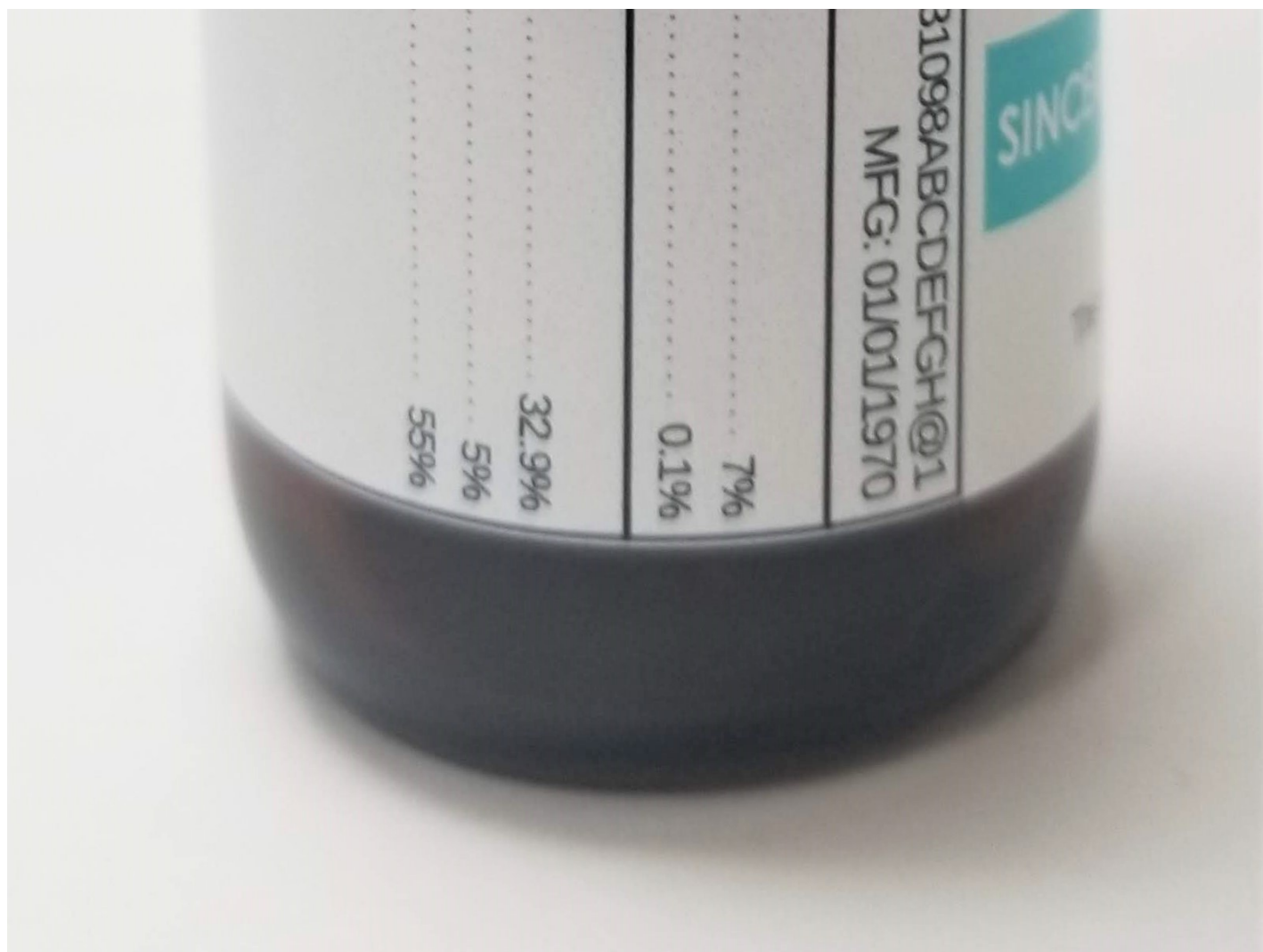
Progesterone USP

Inactive ingredients

Ethyl Alcohol USP

Lactic Acid USP

Propylene Glycol USP



Active, inactive



MINOXIDIL
PROGESTERONE
SOLUTION

SINCE

Rx only
BUD: 01/01/1970

Lot: 031098A
MFG: 01/01/19

Active ingredients

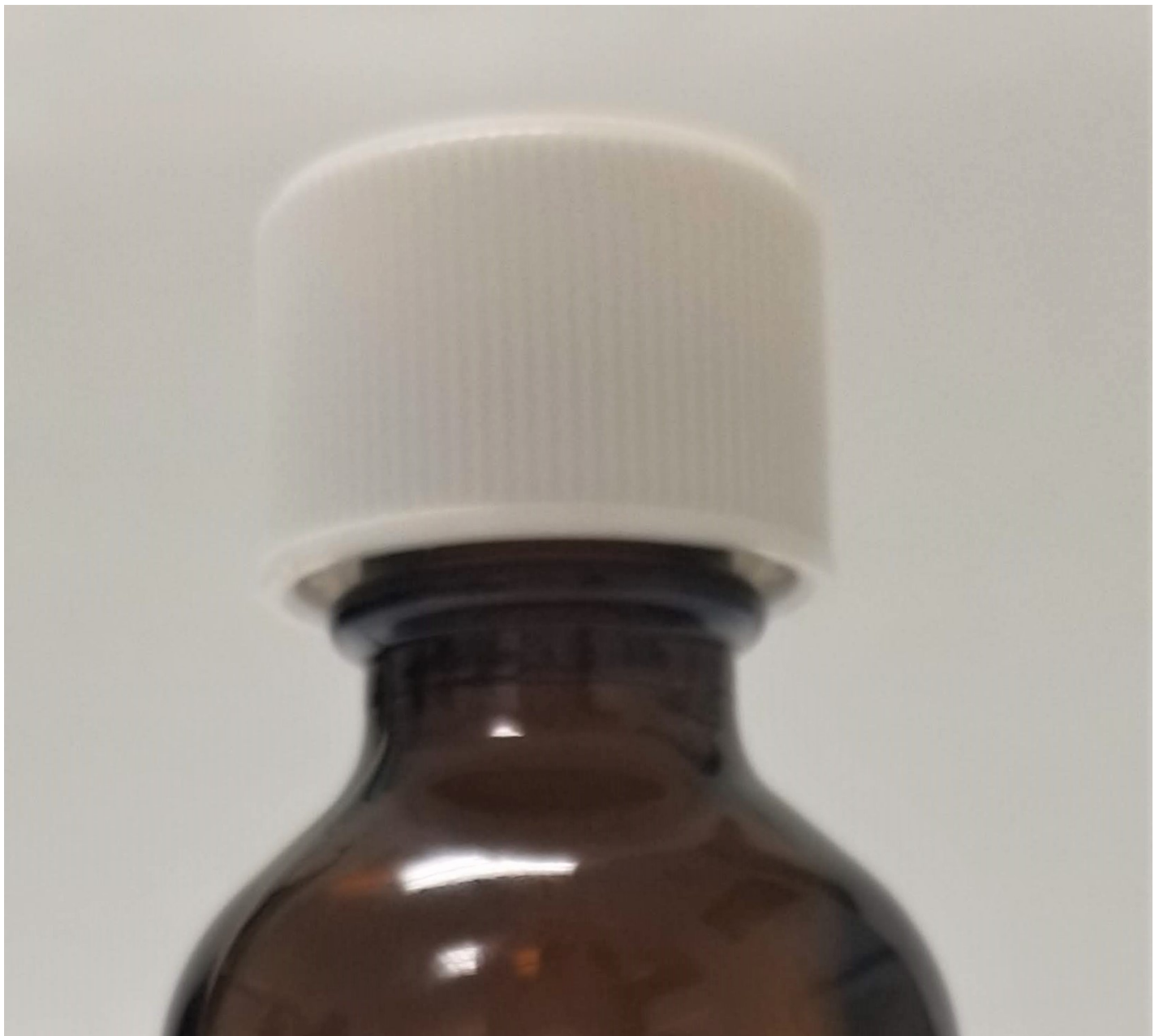
Minoxidil USP
Progesterone USP 0

Inactive ingredients

Ethyl Alcohol USP 32
Lactic Acid USP
Propylene Glycol USP 55



NDC 72934-4748-8 MINOXIDIL 7% / PROGESTERONE 0.1% Solution 60gm



NDC 72934-4748-8

MINOXIDIL USP 7%
PROGESTERONE USP 0.1%
SOLUTION 60gm

SINCERUS

FLORIDA

This is a compounded drug.
Made in USA

MINOXIDIL 7% / PROGESTERONE 0.1%

minoxidil 7% / progesterone 0.1% solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-4748
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MINOXIDIL (UNII: 5965120SH1) (MINOXIDIL - UNII:5965120SH1)	MINOXIDIL	7 g in 100 g
PROGESTERONE (UNII: 4G7DS2Q64Y) (PROGESTERONE - UNII:4G7DS2Q64Y)	PROGESTERONE	0.1 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:72934-4748-8	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(72934-4748)

Revised: 4/2019

Sincerus Florida, LLC