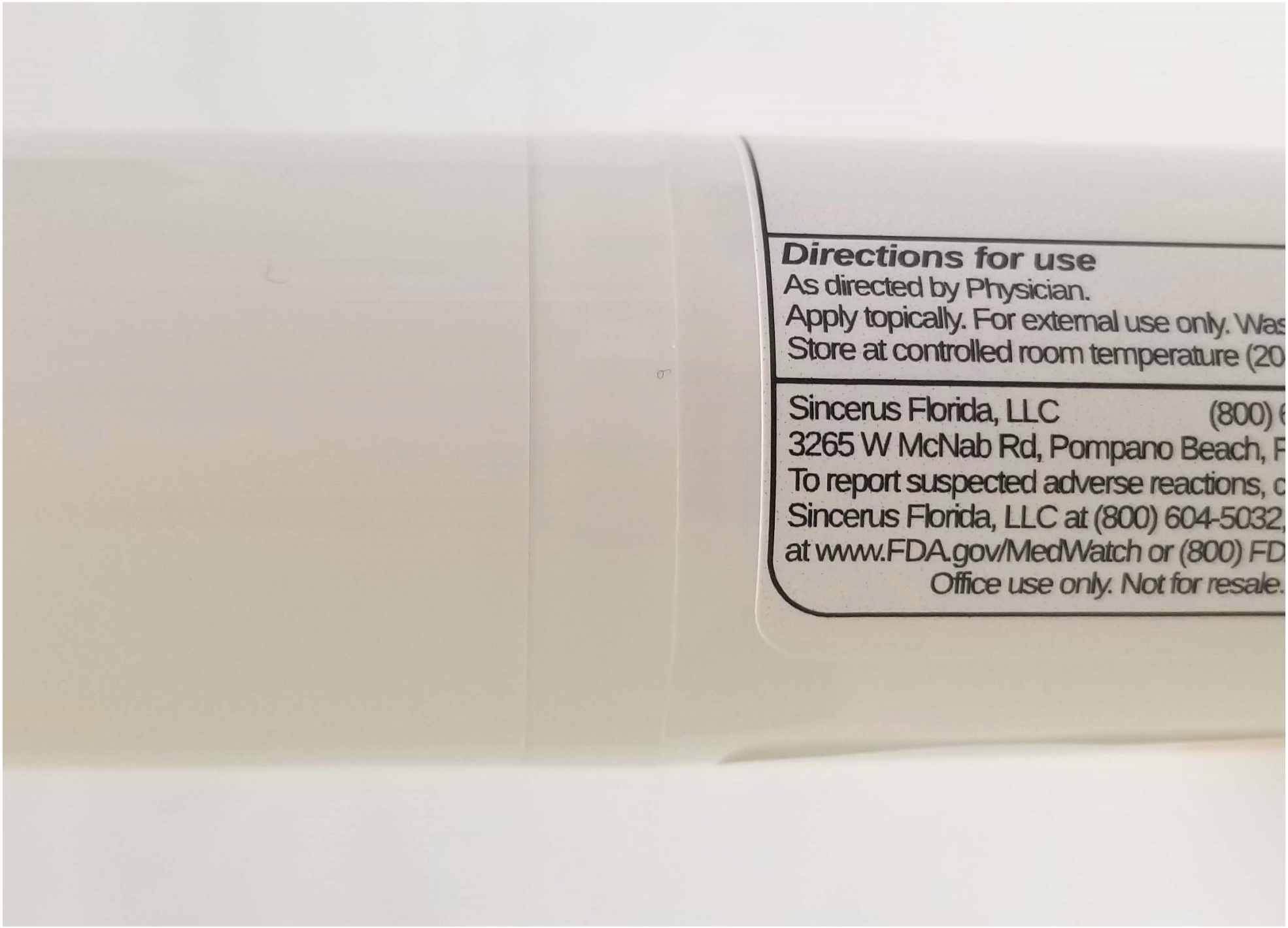
HYDROQUINONE 8% / TRETINOIN 0.025%- hydroquinone 8% / tretinoin 0.025% emulsion
Sincerus Florida, LLC

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.

HYDROQUINONE 8% / TRETINOIN 0.025%

Directions for use

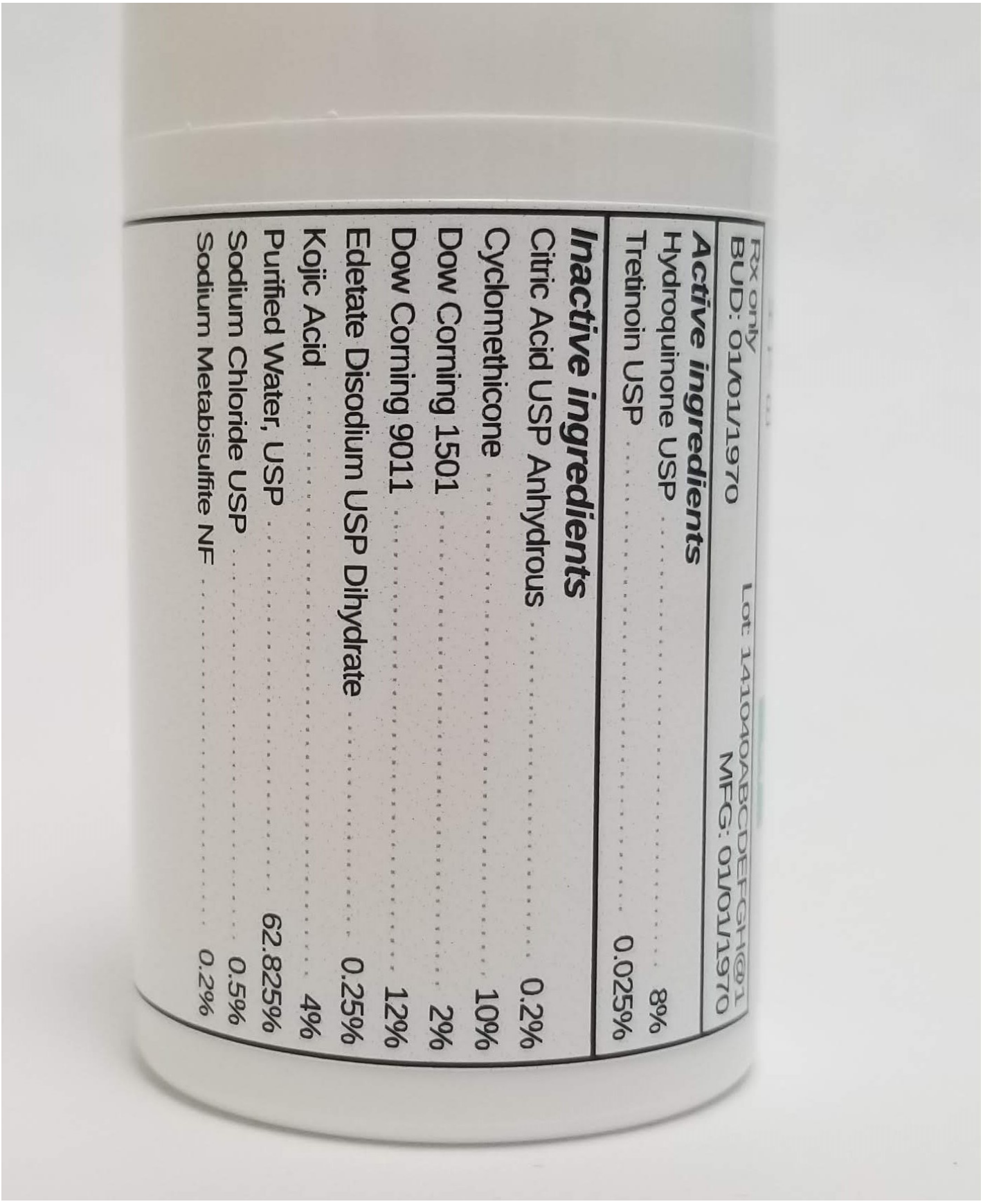
Sincerus Florida, LLC adverse reactions.



Directions for use
As directed by Physician.
Apply topically. For external use only. Was
Store at controlled room temperature (20

Sincerus Florida, LLC (800) 6
3265 W McNab Rd, Pompano Beach, F
To report suspected adverse reactions, c
Sincerus Florida, LLC at (800) 604-5032,
at www.FDA.gov/MedWatch or (800) FD
Office use only. Not for resale.

Active, inactive



NDC 72934-6123-2
HYDROQUINONE 8% /TRETINOIN 0.025%
Emulsion 30gm.


NDC 72934-6123-2
HYDROQUINONE USP 8%
TRETINOIN USP 0.025%
EMULSION 30gm
Rx only
BUD: 01/01/1970

HYDROQUINONE 8% / TRETINOIN 0.025%

hydroquinone 8% / tretinoin 0.025% emulsion

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-6123
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.025 g in 100 g
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	8 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:72934-6123-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/20/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-6123)

Revised: 5/2019 Sincerus Florida, LLC