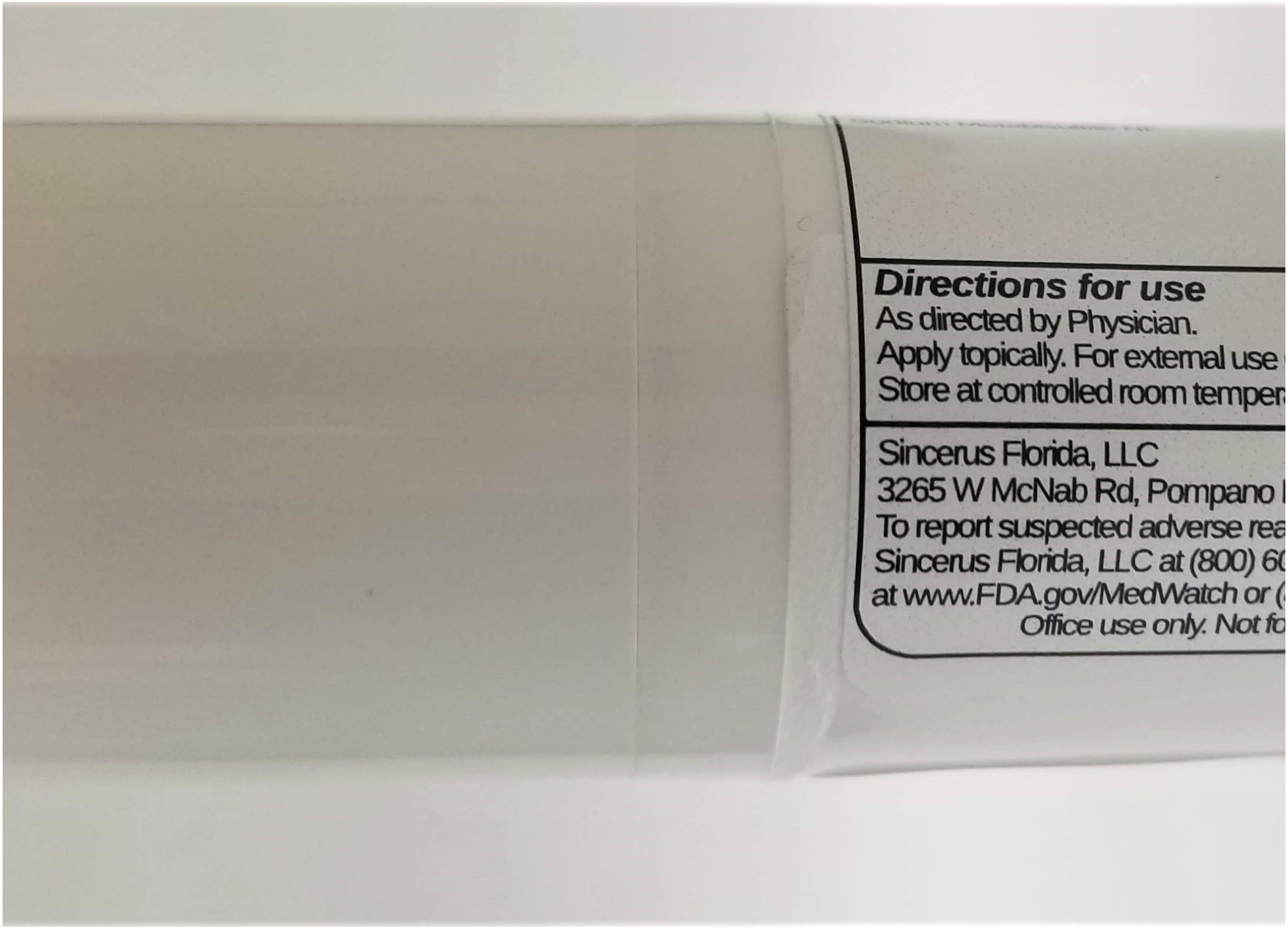
HYDROQUINONE 4%- hydroquinone 4% emulsion
Sincerus Florida, LLC

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 4%

Directions for use

Sincerus Florida, LLC adverse reactions.


Directions for use
As directed by Physician.
Apply topically. For external use
Store at controlled room temper
Sincerus Florida, LLC
3265 W McNab Rd, Pompano
To report suspected adverse rea
Sincerus Florida, LLC at (800) 6
at www.FDA.gov/MedWatch or
Office use only. Not fo

Active, inactive

Rx only
BUD: 01/01/1970

Lot: 141010ABCDEFGH
MFG: 01/01/1

Active ingredients

Hydroquinone USP 4%

Inactive ingredients

Citric Acid USP Anhydrous .....

Cyclomethicone .....

Dow Corning 1501 .....

Dow Corning 9011 .....

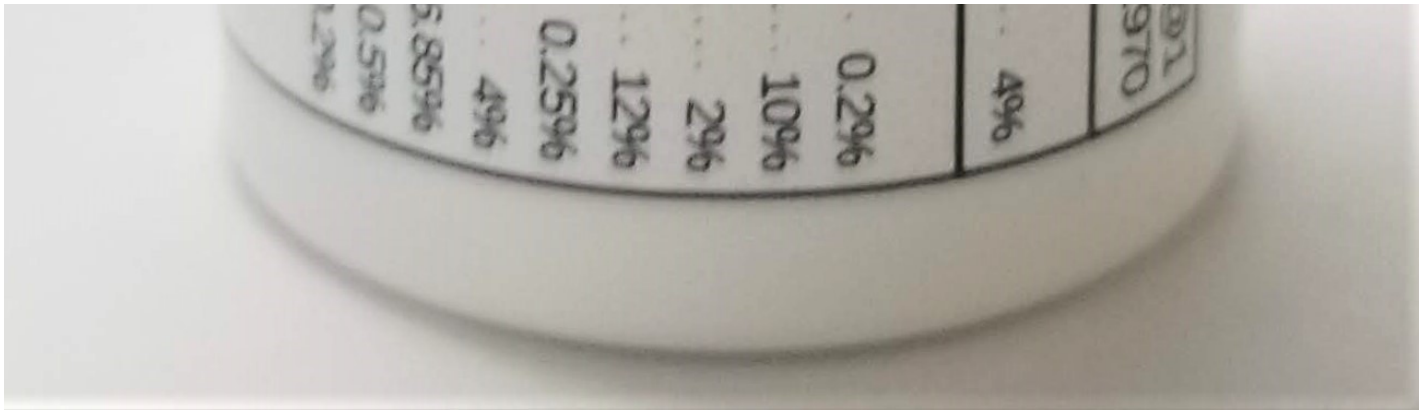
Edetate Disodium USP Dihydrate .....

Kojic Acid .....

Purified Water, USP 6

Sodium Chloride USP .....

Sodium Metabisulfite NF .....



NDC 72934-6114-2 HYDROQUINONE USP 4%, Emulsion 30 gm


Rx only
Lot 141010ABCDEFGH@1
NDC 72934-6114-2
HYDROQUINONE USP 4%
EMULSION 30gm
SINCERUS FLORIDA
This is a compounded drug.
Made in USA

HYDROQUINONE 4%

hydroquinone 4% emulsion

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-6114
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	4 g in 100 g

Product Characteristics			
Color	yellow (BEIGE OPAQUE)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72934-6114-2	30 g in 1 BOTTLE, PUMP; 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(72934-6114)

Revised: 4/2019 Sincerus Florida, LLC