

HYDROCORTISONE 0.5% / HYDROQUINONE 8% / TRETINOIN 0.025% - hydrocortisone 0.5% / 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](#).

HYDROCORTISONE 0.5% / HYDROQUINONE 8% / TRETINOIN 0.025%

Directions for use



Sodium Chloride USP
Sodium Metabisulfite NF
Vitamin E Acetate USP Liquid

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 or (800) FDA-1088.

Office use only. Not for resale.



Sincerus Florida, LLC adverse reactions



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Active, inactive

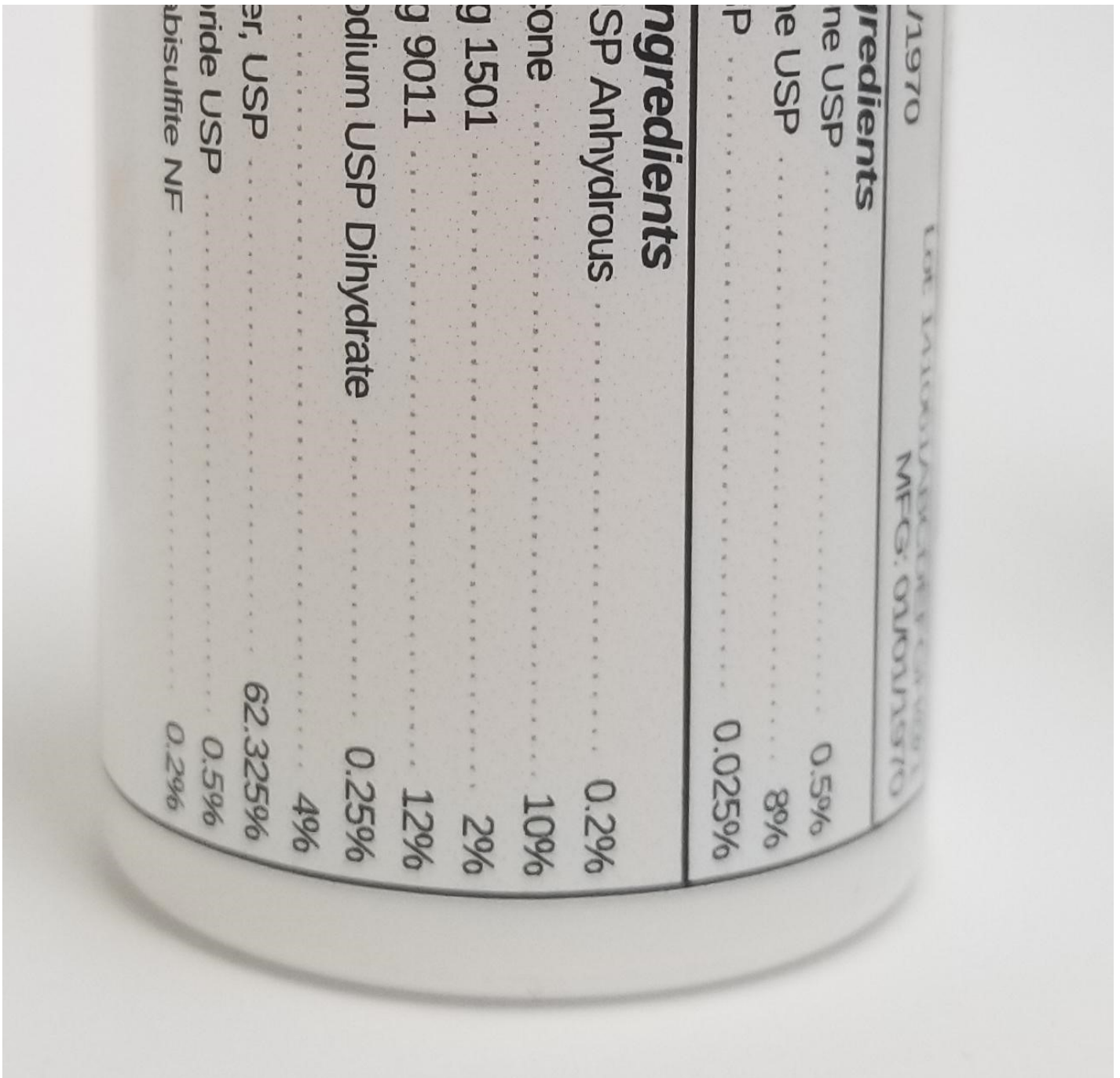
RX only
BUD: 01/01

Active ing

Hydrocortiso
Hydroquinor
Tretinoin US

Inactive i

Citric Acid U
Cyclomethic
Dow Cornin
Dow Cornin
Edetate Disc
Kojic Acid
Purified Wate
Sodium Chlo
Sodium Meta



NDC 72934-6107-2 HYDROCORTISONE USP 0.5% / HYDROQUINONE USP 8% / TRETINOIN USP 0.025%. Emulsion 30 gm



NDC 72934-6107-2

HYDROCORTISONE USP 0.5%

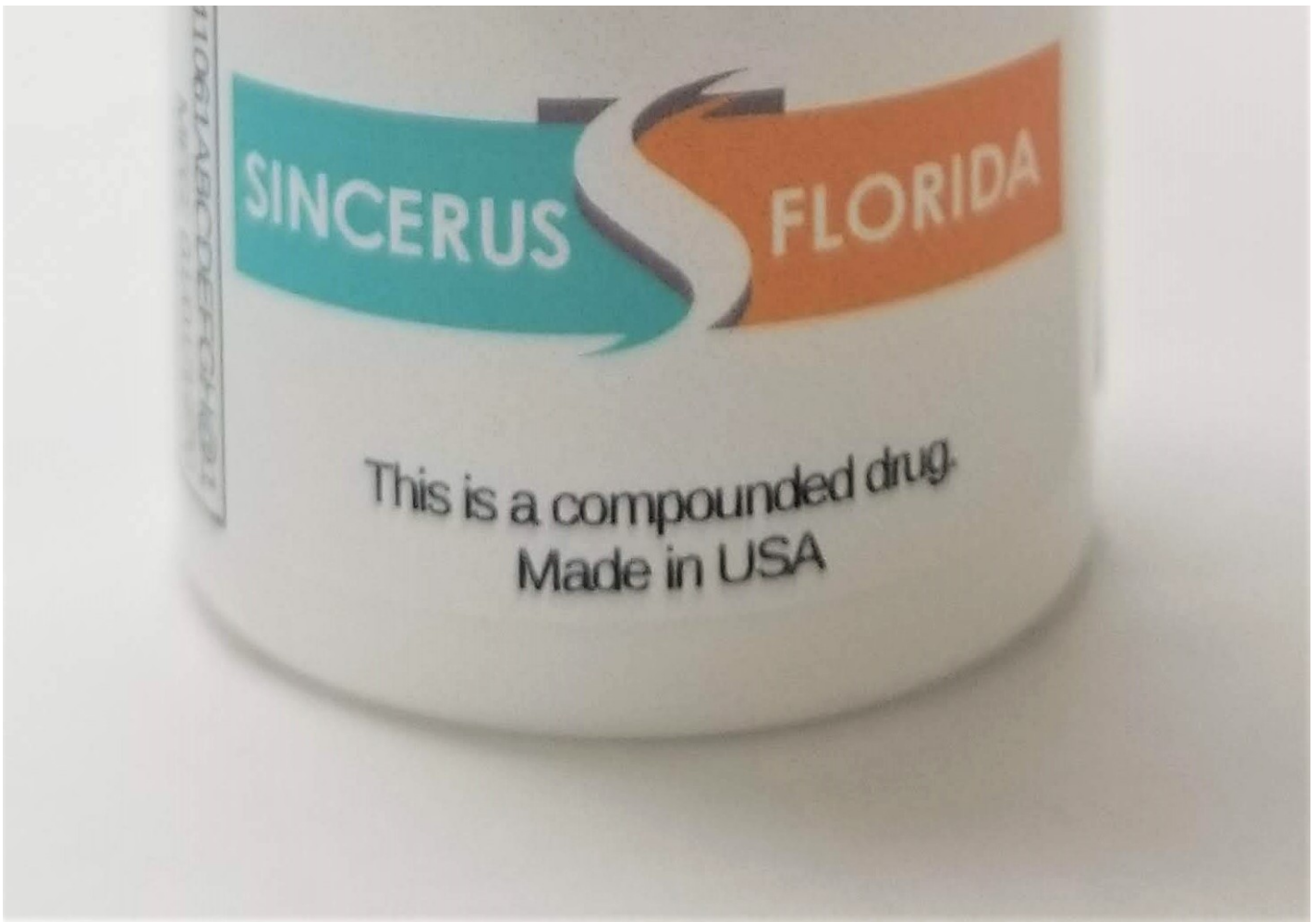
HYDROQUINONE USP 8%

TRETINOIN USP 0.025%

EMULSION 30gm

Rx only
NDC 72934-6107-2

Lot 1



HYDROCORTISONE 0.5% / HYDROQUINONE 8% / TRETINOIN 0.025%
 hydrocortisone 0.5% / hydroquinone 8% / tretinoin 0.025% emulsion

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-6107
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE (UNII: W14X0X7BPJ) (HYDROCORTISONE - UNII:W14X0X7BPJ)	HYDROCORTISONE	0.5 g in 100 g
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	8 g in 100 g
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.025 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:72934-6107-2	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(72934-6107)

Revised: 4/2019

Sincerus Florida, LLC