

**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](#).*

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**HYDROCORTISONE 0.5% / HYDROQUINONE 8% / TRETINOIN 0.025%**

**Directions for use**



Purified Water, USP  
Sodium Chloride USP  
Sodium Metabisulfite NF  
Vitamin E Acetate USP Liquid

0.5%  
0.2%  
1%

**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.



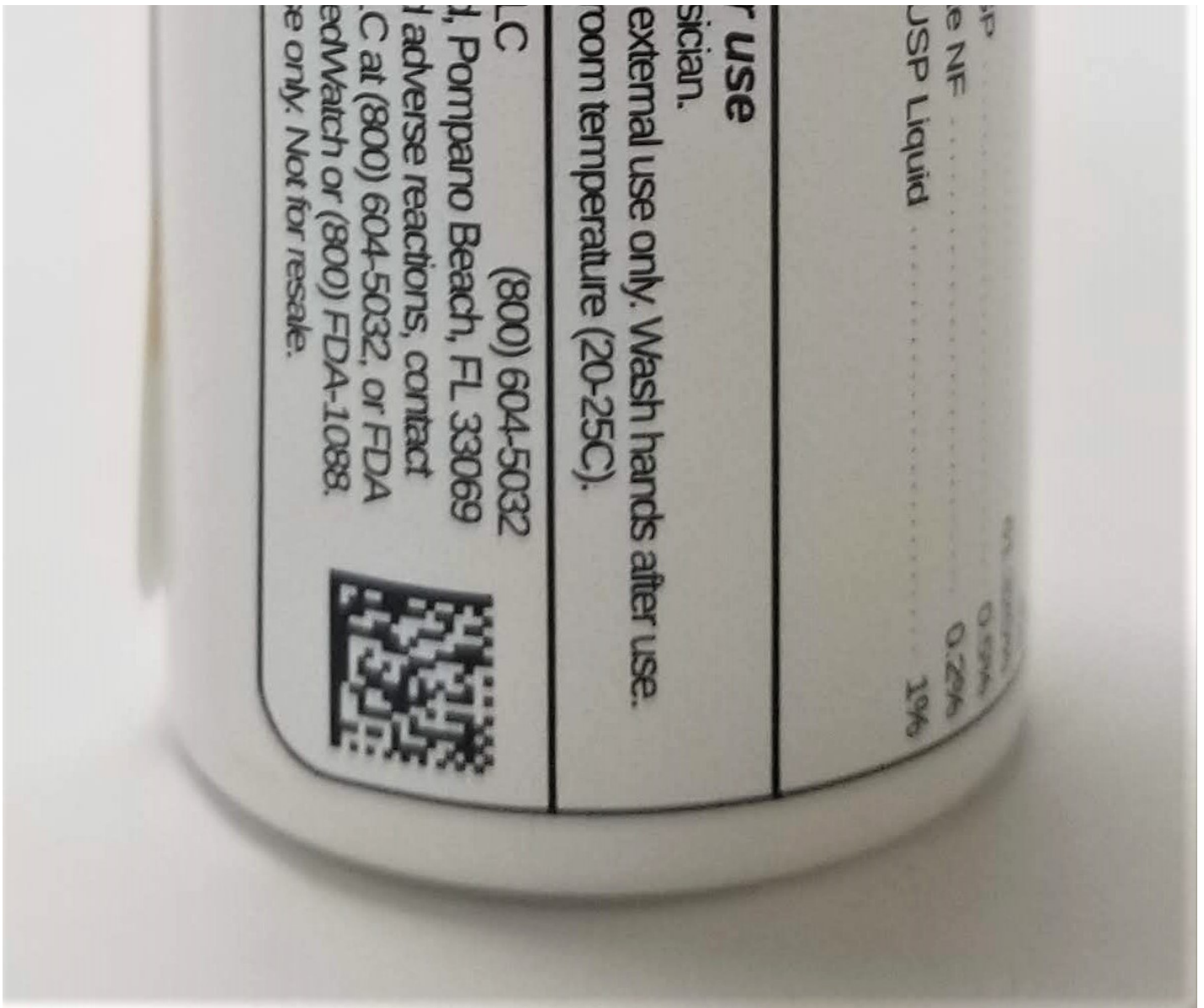
Sincerus Florida, LLC adverse reactions.

Purified Water, USP  
Sodium Chloride US  
Sodium Metabisulfite  
Vitamin E Acetate 1

**Directions for**

As directed by Phys  
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Sincerus Florida, LL  
3265 W McNab Rd  
To report suspectec  
Sincerus Florida, LL  
at [www.FDA.gov/M](http://www.FDA.gov/M)  
Office us



Active, Inactive



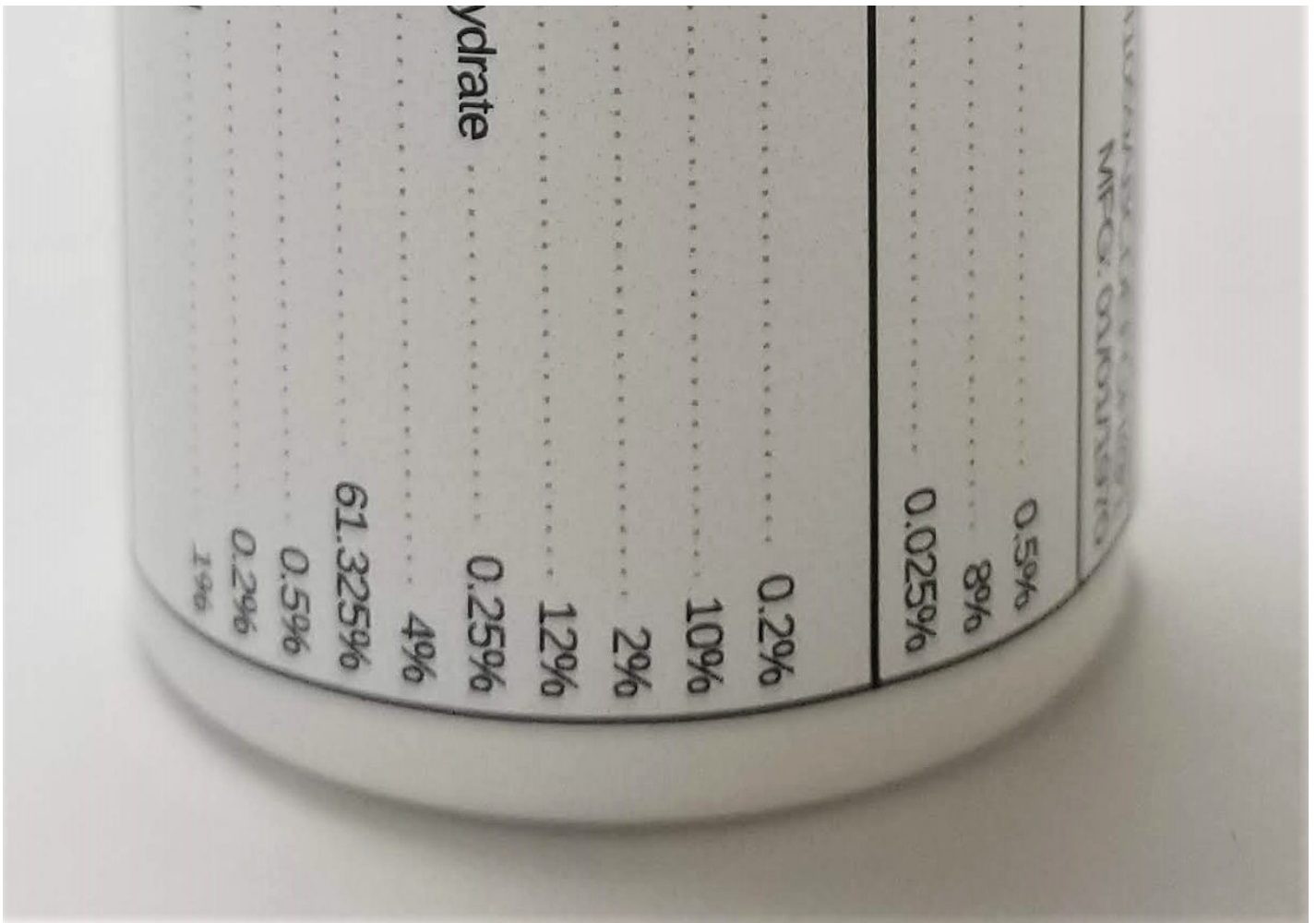
FOR ORAL USE  
BUD: 01/01/1970

**Active ingredients**

- Hydrocortisone USP
- Hydroquinone USP
- Tretinoin USP

**Inactive ingredients**

- Citric Acid USP Anhydrous
- Cyclomethicone
- Dow Corning 1501
- Dow Corning 9011
- Edetate Disodium USP Dihydrate
- Kojic Acid
- Purified Water, USP
- Sodium Chloride USP
- Sodium Metabisulfite NF
- Vitamin E Acetate USP Liquid



**NDC 72934-6106-2 HYDROCORTISONE USP 0.5% / HYDROQUINONE USP 8% / TRETINOIN USP 0.025%. Emulsion 30 gm**

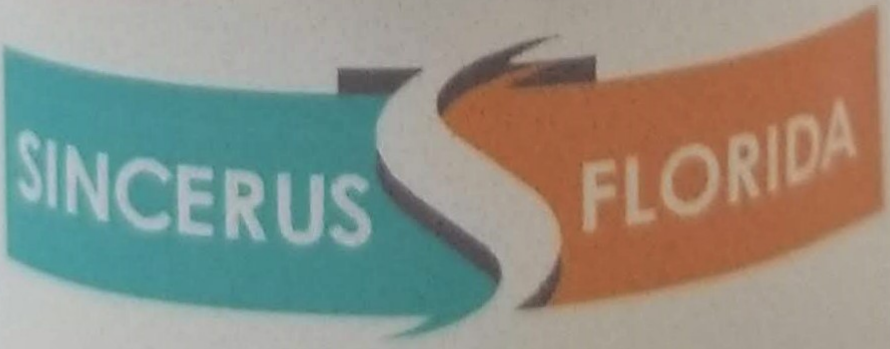


**NDC 72934-6106-2**

**HYDROCORTISONE USP 0.5%  
HYDROQUINONE USP 8%  
TRETINOIN USP 0.025%  
EMULSION 30gm**

Rx only  
EUD: 01/01/1970

Lot 141059A  
ABCDEFGHI  
MFG 01/01/1970



This is a compounded drug.  
Made in USA

## HYDROCORTISONE 0.5% / HYDROQUINONE 8% / TRETINOIN 0.025%

hydrocortisone 0.5% / hydroquinone 8% / tretinoin 0.025% emulsion

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-6106
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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
HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE	0.5 g in 100 g

### Product Characteristics

<b>Color</b>	yellow	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-6106-2	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(72934-6106)

Revised: 4/2019

Sincerus Florida, LLC