

**141016 HYDROCORTISONE 0.5% / HYDROQUINONE 4% / TRETINOIN 0.025% - 141016 hydrocortisone 0.5% / hydroquinone 4% / 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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**141016 HYDROCORTISONE 0.5% / HYDROQUINONE 4% / TRETINOIN 0.025%**

**Directions for use**



**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.

LS-01-000000-00



Sincerus Florida, LLC adverse reactions

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Sincerus Florida, LLC

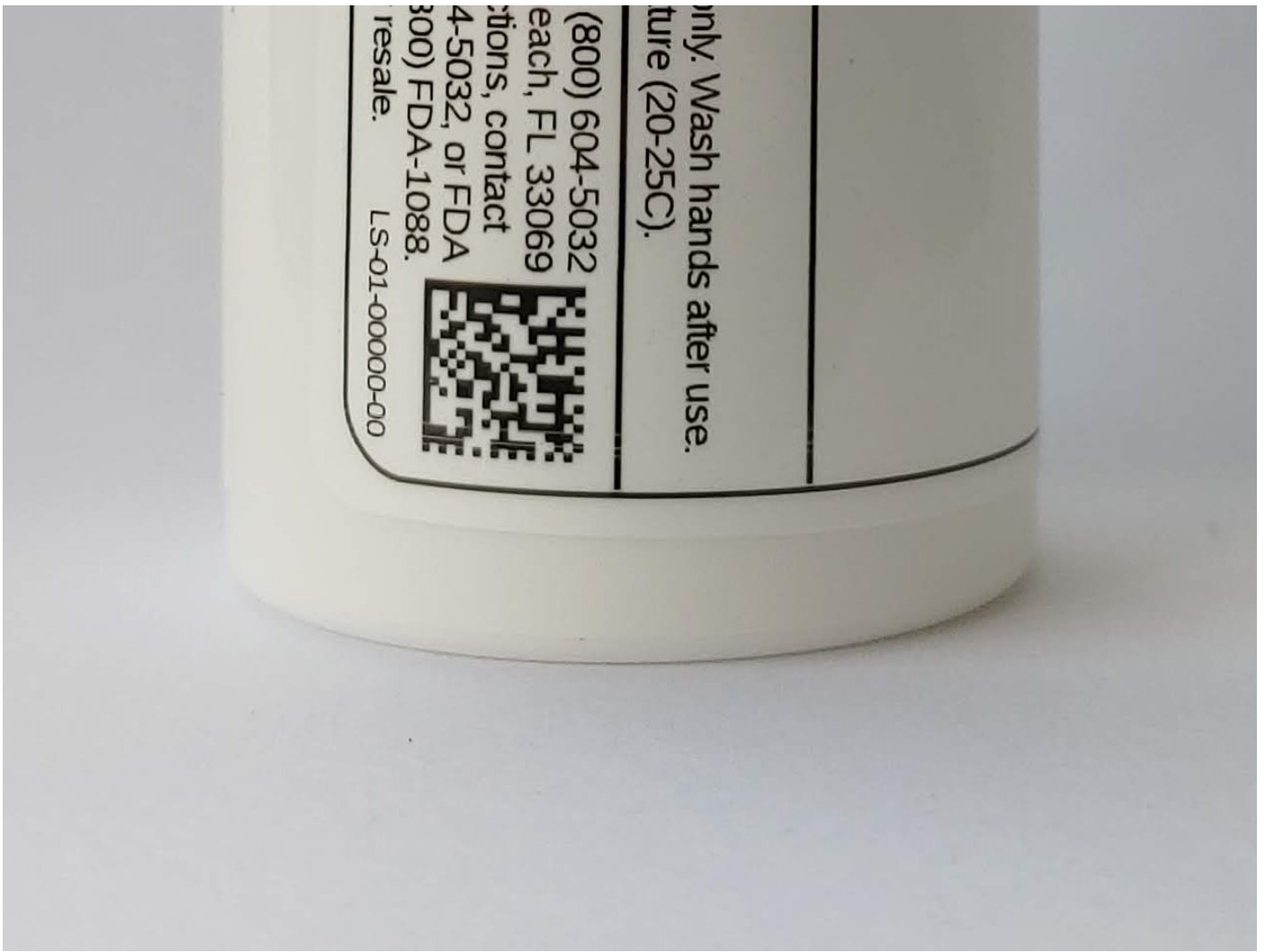
3265 W McNab Rd, Pompano B

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Sincerus Florida, LLC at (800) 60

at [www.FDA.gov/MedWatch](http://www.FDA.gov/MedWatch) or (8

Office use only. Not for



**Active, inactive**





Rx only	Lot: 141016XXXXXXXXXXXXXXX
BUD: XXXXXXXXXXXXX	MFG: XXXXXXXXXXXXX
<b>Active ingredients</b>	
Hydrocortisone USP	0.5%
Hydroquinone USP	4%
Tretinoin USP	0.025%
<b>Inactive ingredients</b>	
Citric Acid USP Anhydrous	0.2%
Cyclomethicone	10%
Dow Corning 1501	2%
Dow Corning 9011	12%
Edetate Disodium USP Dihydrate	0.25%
Kojic Acid	6%
Purified Water USP	64.325%
Sodium Chloride USP	0.5%
Sodium Metabisulfite NF	0.2%

**NDC 72934-6225-2 141016 HYDROCORTISONE 0.5% / HYDROQUINONE 4% / TRETINOIN 0.025% emulsion 30 gm**

NDC 72934-6225-2

141016

HYDROCORTISONE 0.5%

HYDROQUINONE 4%

TRETINOIN 0.025%

EMULSION 30gm

Rx only  
BUD: XXXXXXXXXX  
Active Ingredients  
Lot: 141016XXXX  
MFG: XX







**141016 HYDROCORTISONE 0.5% / HYDROQUINONE 4% / TRETINOIN 0.025%**

141016 hydrocortisone 0.5% / hydroquinone 4% / tretinoin 0.025% emulsion

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-6225
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	4 g in 100 g
HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE	0.5 g in 100 g
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.025 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:72934-6225-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/02/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/02/2020	

**Labeler** - Sincerus Florida, LLC (080105003)

**Establishment**

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-6225)

Revised: 7/2020

Sincerus Florida, LLC