

**HYDROQUINONE 8% - hydroquinone 8% 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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**HYDROQUINONE 8%**

**Directions for use**



SODIUM CHLORIDE USP  
Sodium Metabisulfite NF

0.25%

**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.*



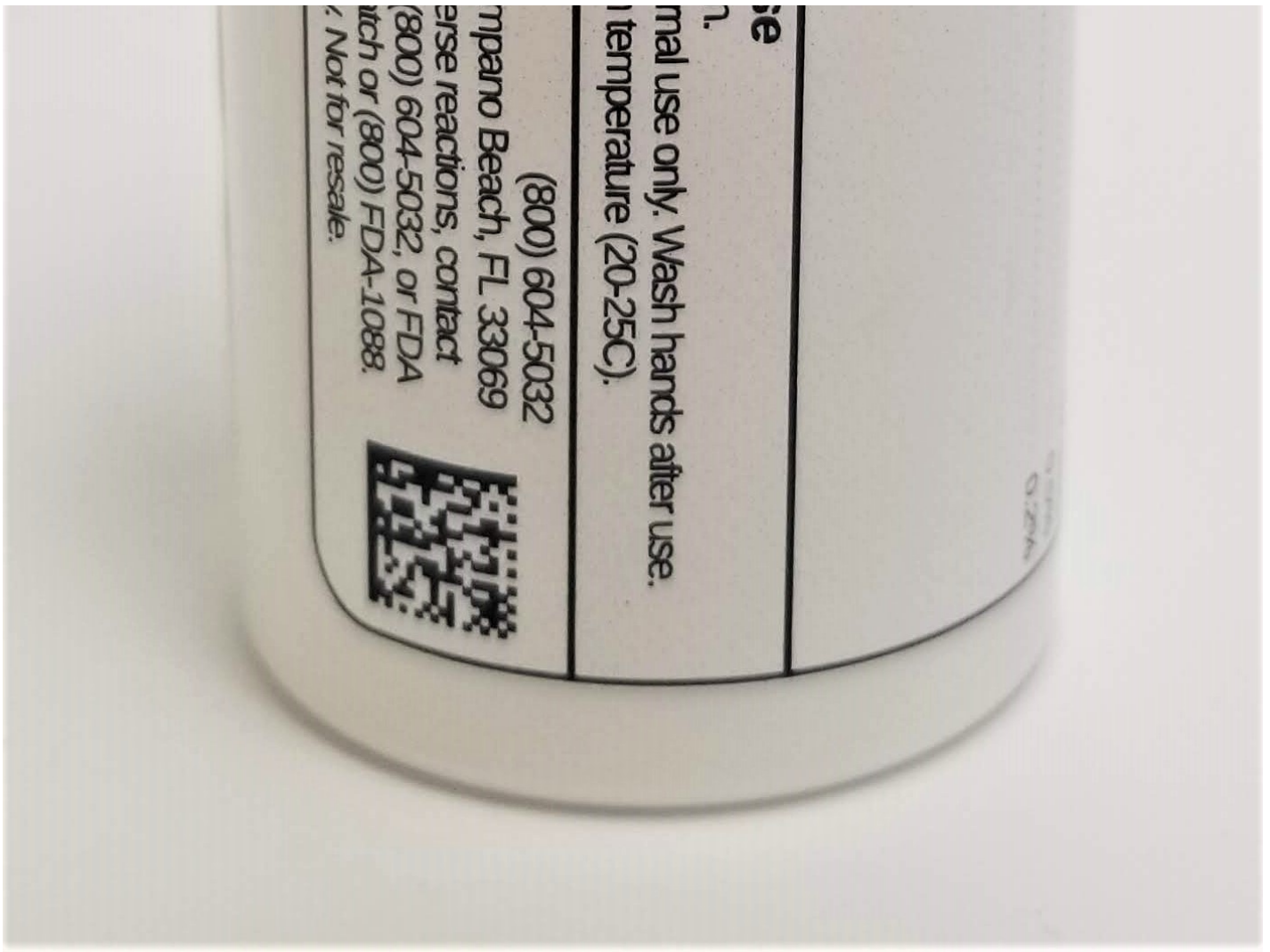
Sincerus Florida, LLC adverse reactions.

Sodium Citrate USP  
Sodium Metabisulfite NF

**Directions for use**

As directed by Physician  
Apply topically. For external use only.  
Store at controlled room temperature.

Sincerus Florida, LLC  
3265 W McNab Rd, Port St. Lucie, FL 34952  
To report suspected adverse events, contact  
Sincerus Florida, LLC at 1-800-368-7273 or  
at [www.FDA.gov/medwatch](http://www.FDA.gov/medwatch)  
Office use only



**Active, inactive**



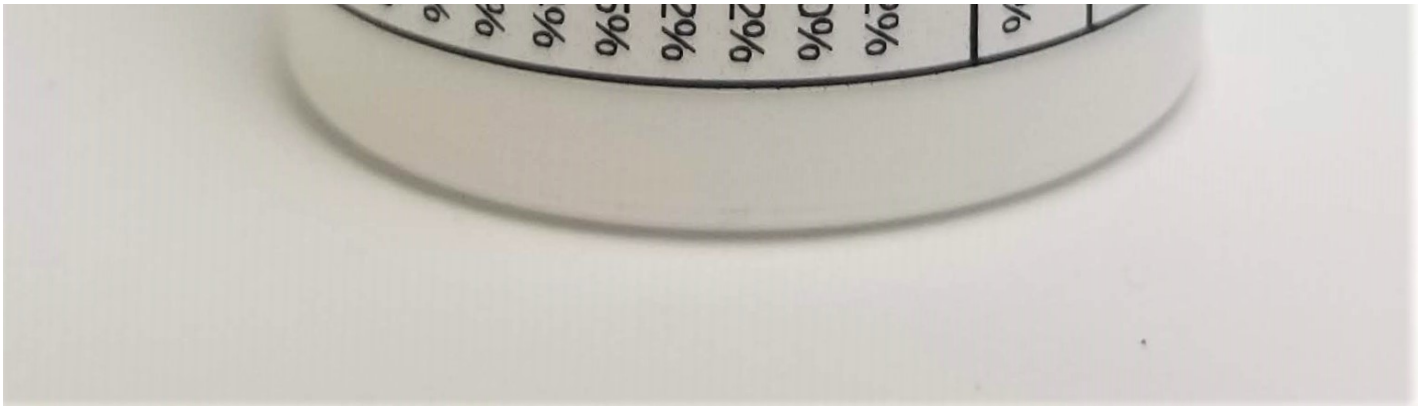


Rx only  
BUD: 01/01/1970  
Lot 141030A B C D E F G H I J  
MFG: 01/01/1970

**Active ingredients**  
Hydroquinone USP ..... 80

**Inactive ingredients**

Citric Acid USP Anhydrous ..... 0.2  
Cyclomethicone ..... 10  
Dow Corning 1501 ..... 2  
Dow Corning 9011 ..... 12  
Edetate Disodium USP Dihydrate ..... 0.25  
Kojic Acid ..... 4  
Purified Water, USP ..... 62.85  
Sodium Chloride USP ..... 0.59  
Sodium Metabisulfite NF ..... 0.2%



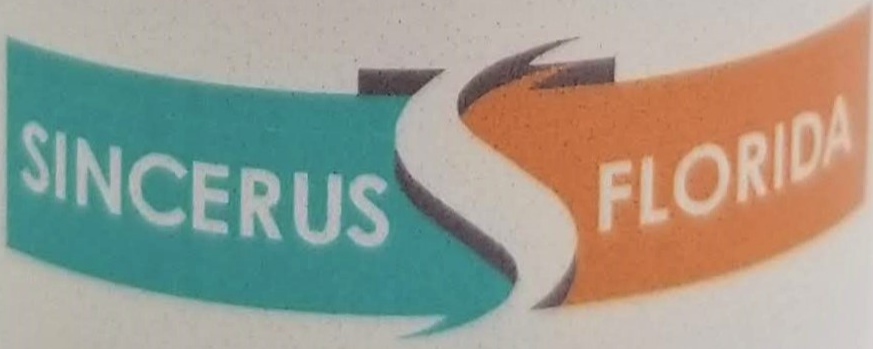
**NDC 72934-6121-2 Hydroquinone USP 8%.  
Emulsion 30gm**



Rx only  
BUD: 01/01/1970

Lot: 141030ABCDEFH01  
MFG: 01/01/1970

**NDC 72934-6121-2**  
**HYDROQUINONE USP 8%**  
**EMULSION 30gm**



This is a compounded drug.  
Made in USA

# HYDROQUINONE 8%

hydroquinone 8% emulsion

## Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-6121
<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

## Product Characteristics

<b>Color</b>	yellow (BEIGE OPAQUE)	<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:72934-6121-2	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(72934-6121)

Revised: 4/2019

Sincerus Florida, LLC