

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

HYDROQUINONE 8% / 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 or (800) FDA-1088.

Office use only. Not for resale.



Sincerus Florida, LLC. Adverse reactions



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Active, Inactive



Active Ingredients	
Hydroquinone USP	8%
Tretinoin USP	0.025%
Inactive ingredients	
Ascorbyl Palmitate FCC	2%
Citric Acid USP Anhydrous	0.2%
Cyclomethicone	10%
Dow Corning 1501	2%
Dow Corning 9011	12%
Edetate Disodium USP Dihydrate	0.25%
Green Tea Extract	2%
Kojic Acid	4%
Purified Water, USP	57.825%
Sodium Chloride USP	0.5%
Sodium Metabisulfite NF	0.2%
Vitamin E Acetate USP Liquid	1%
Directions for use	
As directed by Physician.	

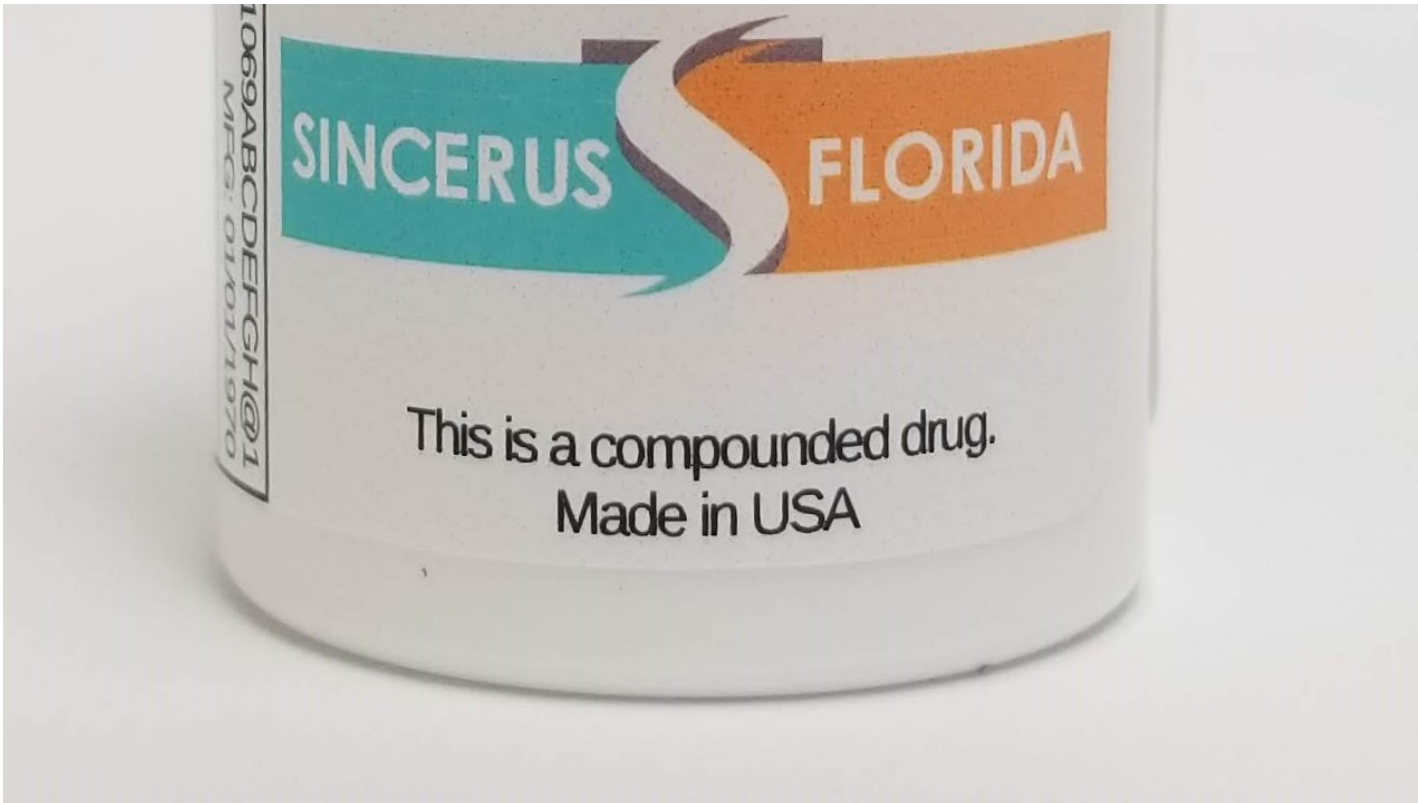
NDC 72934- 6124-2 HYDROQUINONE USP 8% / TRETINOIN USP 0.025%. Emulsion 30 gm

NDC 72934-6124-2

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

Rx only
BUD: 01/01/1970

Lot 14



HYDROQUINONE 8% / TRETINOIN 0.025%.

hydroquinone 8% / tretinoin 0.025%. emulsion

Product Information						
Product Type		HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-6124		
Route of Administration		TOPICAL				
Active Ingredient/Active Moiety						
Ingredient Name			Basis of Strength	Strength		
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)			TRETINOIN	0.025 g in 100 g		
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)			HYDROQUINONE	8 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:72934-6124-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/17/2019			

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/17/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-6124)

Revised: 5/2019

Sincerus Florida, LLC