

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



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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		Lot: 1A10A0ABCDEF01
BUD: 01/01/1970		MFG: 01/01/1970
Active ingredients		
Hydroquinone USP	8%
Tretinoin USP	0.025%
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	62.825%
Sodium Chloride USP	0.5%
Sodium Metabisulfite NF	0.2%

NDC 72934-6123-2
HYDROQUINONE 8% /TRETINOIN 0.025%
Emulsion 30gm.

Rx only
BUD: 01/01/1970

NDC 72934-6123-2

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



HYDROQUINONE 8% / TRETINOIN 0.025%

hydroquinone 8% / tretinoin 0.025% emulsion

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-6123
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:72934-6123-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2019	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		05/20/2019		

Labeler - Sincerus Florida, LLC (080105003)

Establishment			
Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-6123)

Revised: 5/2019

Sincerus Florida, LLC