

141030 HYDROQUINONE 8% - 141030 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](#).

141030 HYDROQUINONE 8%

Directions for use



Directions for use

is 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
165 W McNab Rd, Pompano Beach, FL 33069
Report suspected adverse reactions, contact
Sincerus Florida, LLC at (800) 604-5032, or FDA
www.FDA.gov/MedWatch or (800) FDA-1088.



Office use only. Not for resale. LS-01-000000-00

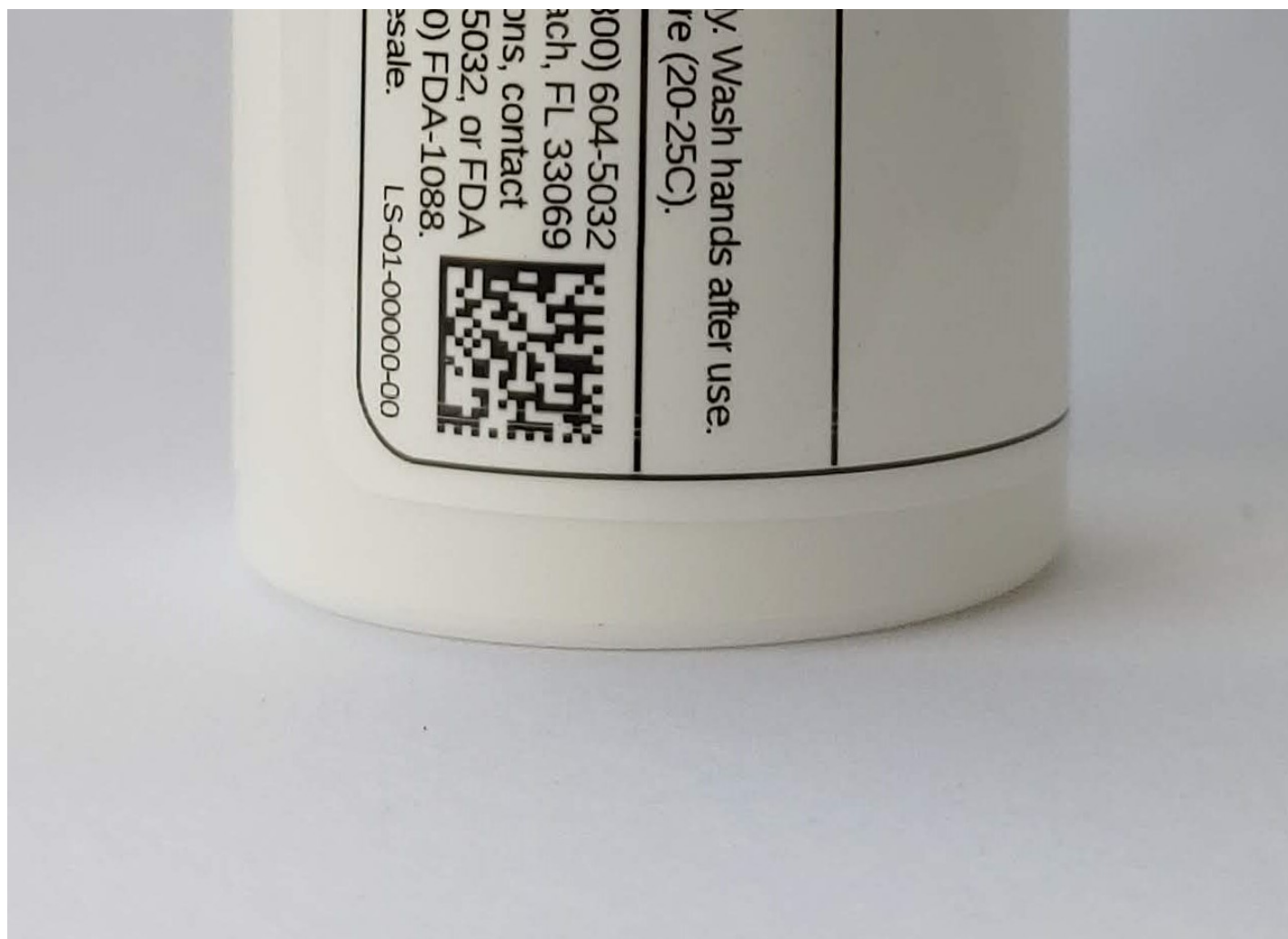
Sincerus Florida, LLC adverse reactions

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As directed by Physician.

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Sincerus Florida, LLC (8
3265 W McNab Rd, Pompano Beach
To report suspected adverse reactions,
Sincerus Florida, LLC at (800) 604-
at www.FDA.gov/MedWatch or (800)
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Active, inactive



1410
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EM

Rx only Lot: 141030XXXXXXXXXX@XX
BUD: XXXXXXXXXX MFG: XXXXXXXXXX

Active ingredients

Hydroquinone USP 8%

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 0.5%

Sodium Metabisulfite NF 0.2%

NDC 72934-6229-2 141030 HYDROQUINONE 8% emulsion 30 gm



NDC 72934-6229-2

141030
HYDROQUINONE 8%
EMULSION 30gm

Rx only

BUD: XXXXXXXXXX

Active ingredients

Hydroquinone USP



141030 HYDROQUINONE 8%

141030 hydroquinone 8% emulsion

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-6229
Route of Administration	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

Color	yellow (Beige)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-6229-2	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(72934-6229)

Revised: 7/2020

Sincerus Florida, LLC