

HYDROCORTISONE 0.5% / HYDROQUINONE 6% / TRETINOIN 0.025% - hydrocortisone 0.5% / hydroquinone 6% / 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](#).

HYDROCORTISONE 0.5% / HYDROQUINONE 6% / TRETINOIN 0.025%

Directions for use



Kojic Acid 0.29%
Purified Water, USP
Sodium Chloride USP 0.59%
Sodium Metabisulfite NF 0.29%

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.



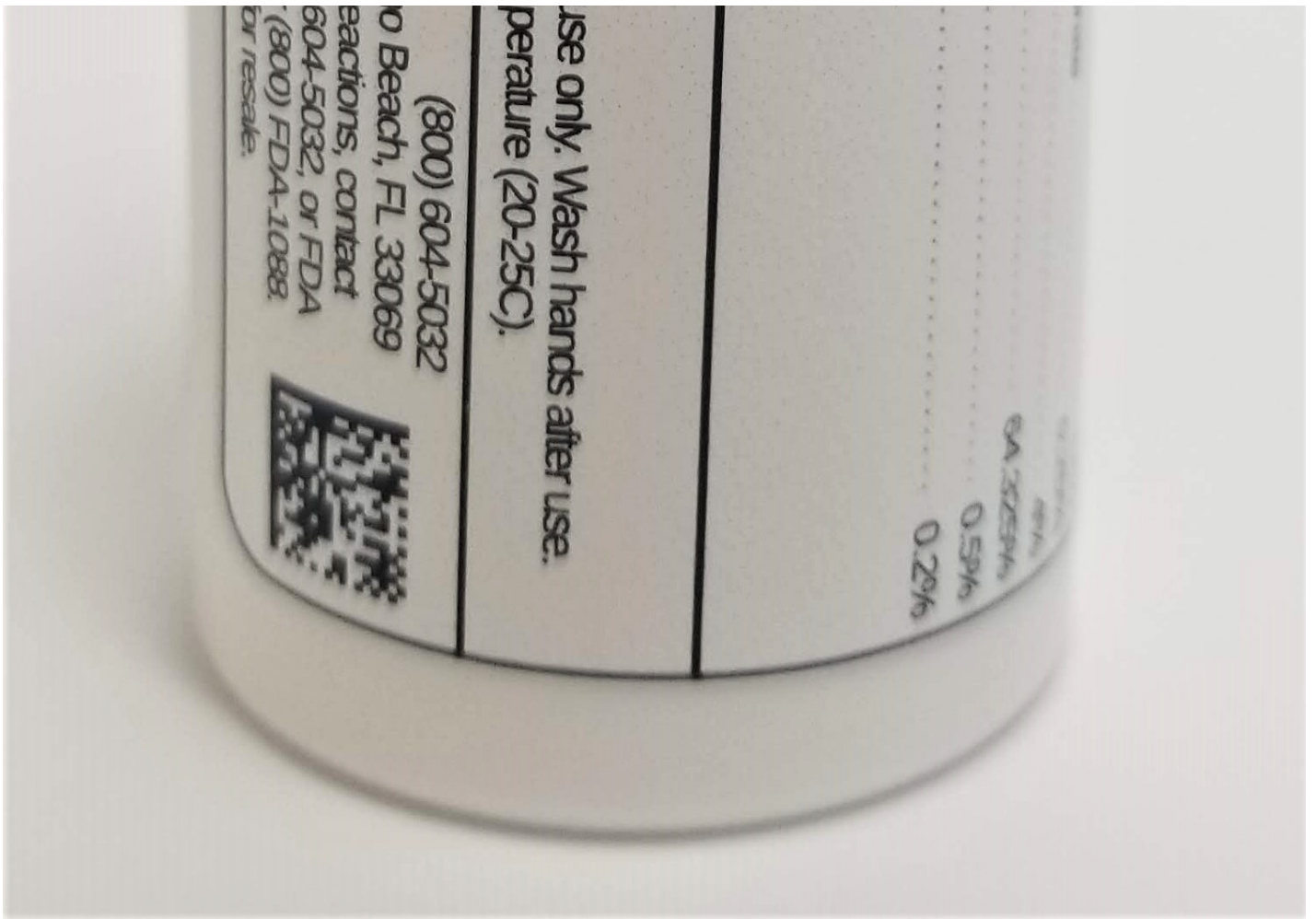
Sincerus Florida, LLC adverse reactions.

Chlorine Dioxide USP
Kojic Acid
Purified Water, USP
Sodium Chloride USP
Sodium Metabisulfite NF

Directions for use

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

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Active, inactive



EXP. DATE: 01/01/2019
BUD: 01/01/2019
USE: VAGINAL CREAM
MFG: GSK

Active ingredients

Hydrocortisone USP 0.5%

Hydroquinone USP 9%

Tretinoin USP 0.025%

Inactive ingredients

Citric Acid USP Anhydrous 0.1%

Cyclomethicone 1%

Dow Corning 1501 1%

Dow Corning 9011 1%

Edetate Disodium USP Dihydrate 0.2%

Kojic Acid 4%

Purified Water, USP 64.325%

Sodium Chloride USP 0.5%

Sodium Metabisulfite NF 0.2%



**NDC 72934-6104-2 HYDROCORTISONE USP 0.5% / HYDROQUINONE USP 6% /
TRETINOIN USP 0.025%. Emulsion 30 gm**

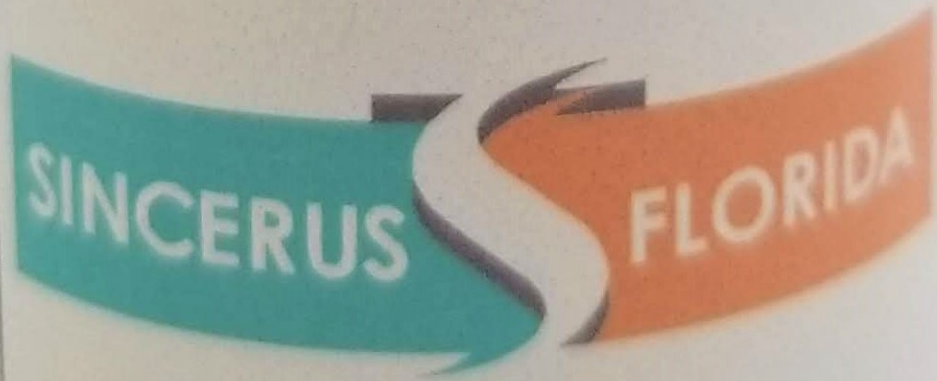


NDC 72934-6104-2

HYDROCORTISONE USP 0.5%
HYDROQUINONE USP 6%
TRETINOIN USP 0.025%
EMULSION 30gm

Rx only
BUD: 01/01/1970

Lot: 1A101ZABCF0H01
NDC 72934-6104-2



This is a compounded drug.
Made in USA

HYDROCORTISONE 0.5% / HYDROQUINONE 6% / TRETINOIN 0.025%

hydrocortisone 0.5% / hydroquinone 6% / tretinoin 0.025% emulsion

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-6104
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	6 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

Color	yellow	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-6104-2	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(72934-6104)