

HYDROCORTISONE 2.5% / HYDROQUINONE 4% / TRETINOIN 0.025%, - hydrocortisone 2.5% / hydroquinone 4% / 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 2.5% / HYDROQUINONE 4% / TRETINOIN 0.025%,

Directions for use



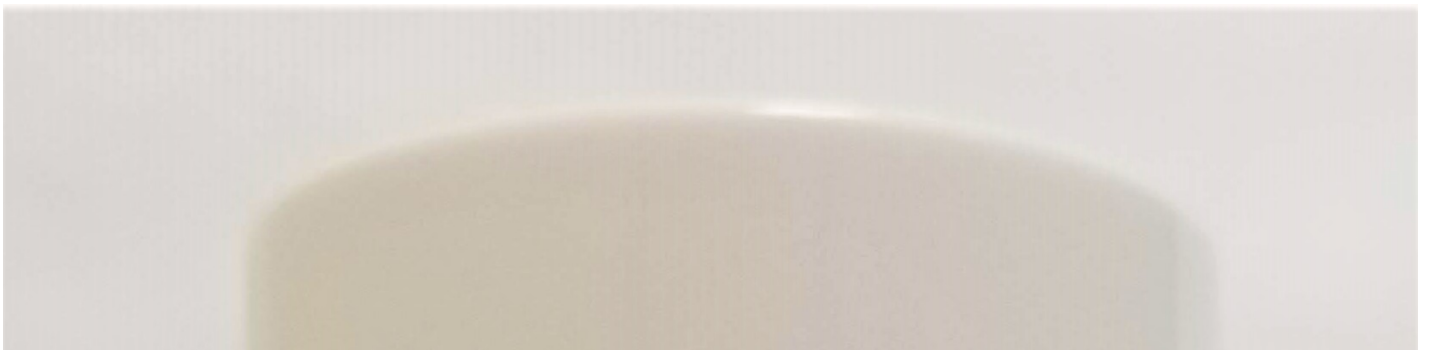
Directions for use

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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Store at controlled room temperature (2

Sincerus Florida, LLC (800

3265 W McNab Rd, Pompano Beach

To report suspected adverse reactions

Sincerus Florida, LLC at (800) 604-503

at www.FDA.gov/MedWatch or (800) F

Office use only. Not for resal



Active, inactive



Active ingredients	
Hydrocortisone USP	2.5%
Hydroquinone USP	4%
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%
Emulsifx	2%
Krisigel 100	2%
Purified Water, USP	64.325%
Sodium Chloride USP	0.5%
Sodium Metabisulfite NF	0.2%

Directions for use
As directed by physician

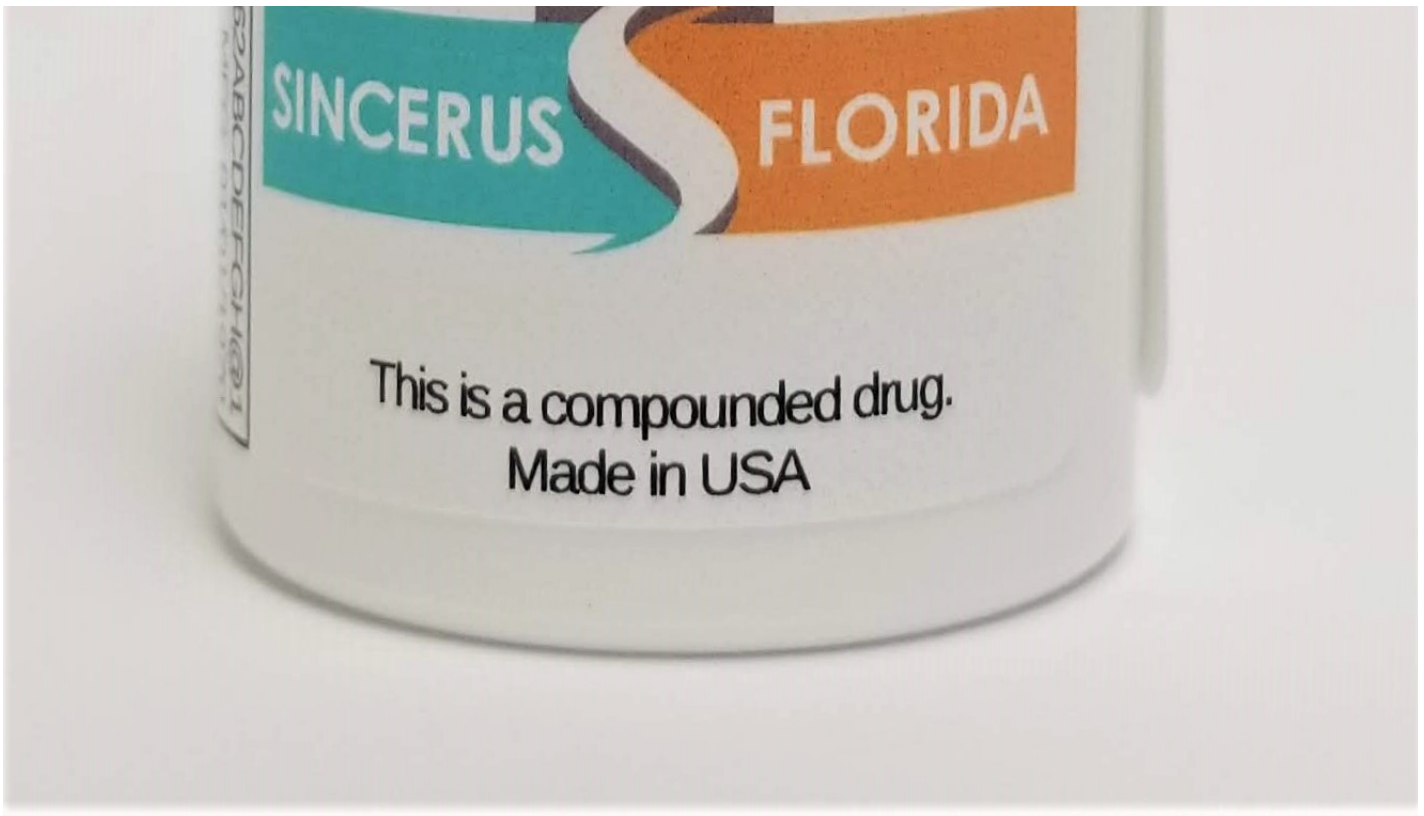
NDC 72934- 6110-2 HYDROCORTISONE USP 2.5% / HYDROQUINONE USP 4% / TRETINOIN USP 0.025%. Emulsion 30 gm

NDC 72934-6110-2

HYDROCORTISONE USP 2.5%
HYDROQUINONE USP 4%
TRETINOIN USP 0.025%
EMULSION 30gm

RX ONLY

LOT 1410



HYDROCORTISONE 2.5% / HYDROQUINONE 4% / TRETINOIN 0.025%,
 hydrocortisone 2.5% / hydroquinone 4% / tretinoin 0.025%, emulsion

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-6110
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	4 g in 100 g
HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE	2.5 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:72934-6110-2	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(72934-6110)

Revised: 5/2019

Sincerus Florida, LLC