

141017 HYDROCORTISONE 0.5% / HYDROQUINONE 6% / TRETINOIN 0.025% - 141017 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](#).

141017 HYDROCORTISONE 0.5% / HYDROQUINONE 6% / 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. LS-01-000000-00



Sincerus Florida, LLC adverse reactions

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Sincerus Florida, LLC

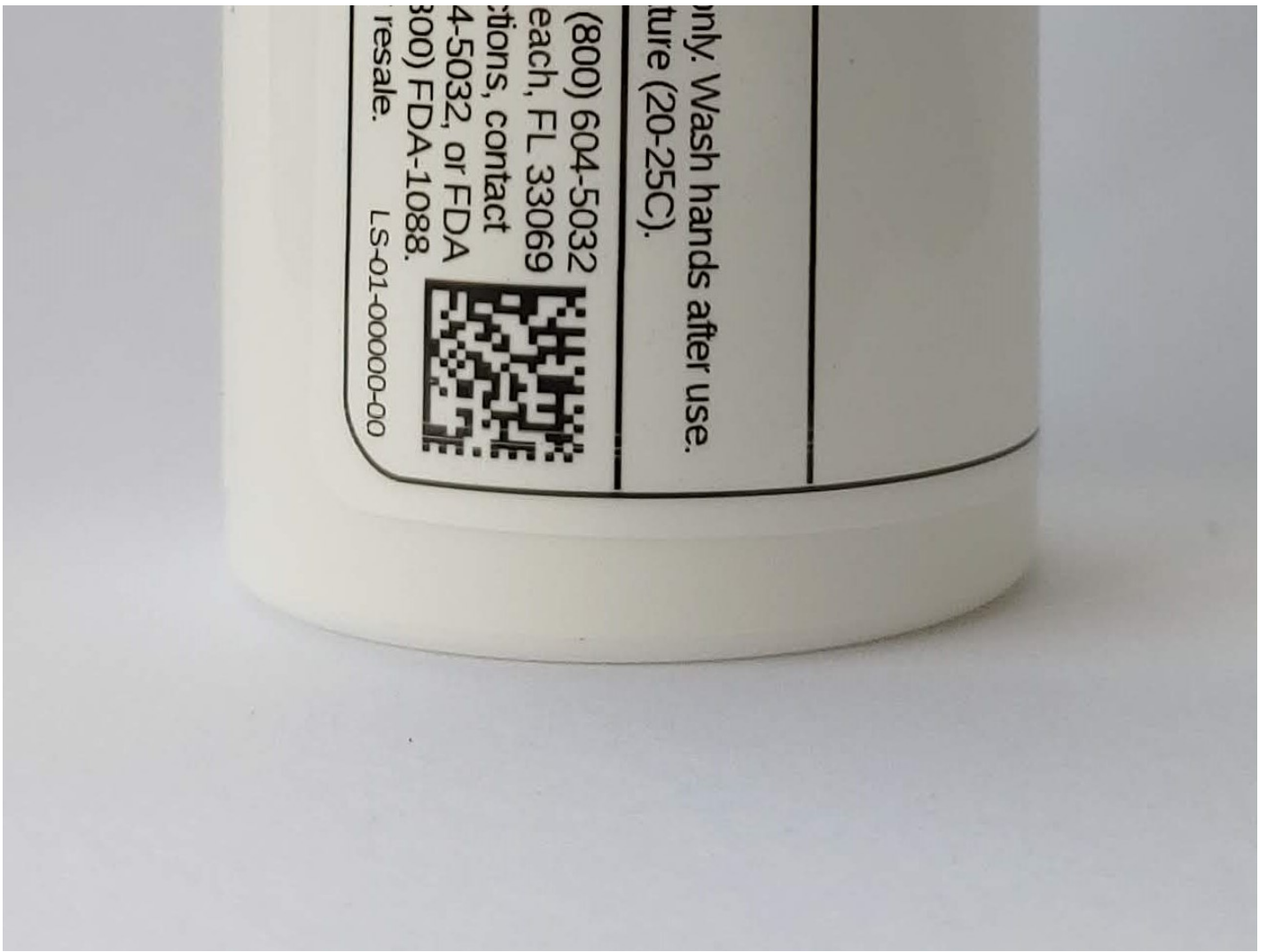
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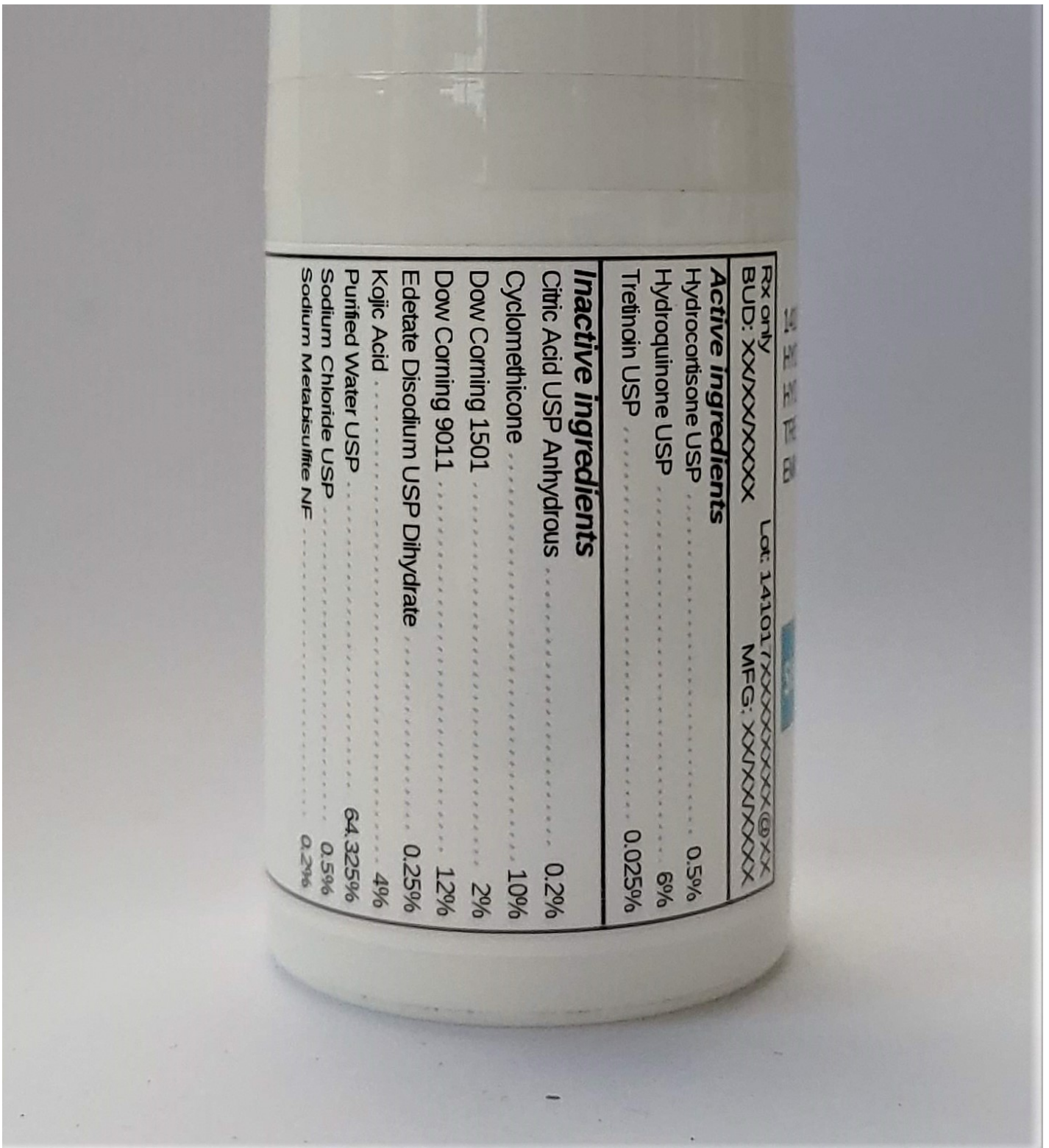
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Active, Inactive





NDC 72934-6226-2 141017 HYDROCORTISONE 0.5% / HYDROQUINONE 6% / TRETINOIN 0.025% emulsion 30gm

NDC 72934-6226-2

141017

HYDROCORTISONE 0.5%

HYDROQUINONE 6%

TRETINOIN 0.025%

EMULSION 30gm



Rx only
BUD: XXXXXXXXX
Active Ingredients
Lot: 141017XXXXXX@XX
MFG: XXXXXXXXX

This is a compounded drug.
Made in USA

141017 HYDROCORTISONE 0.5% / HYDROQUINONE 6% / TRETINOIN 0.025%

141017 hydrocortisone 0.5% / hydroquinone 6% / tretinoin 0.025% emulsion

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-6226
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE (UNII: W14X0X7BPJ) (HYDROCORTISONE - UNII:W14X0X7BPJ)	HYDROCORTISONE	0.5 g in 100 g
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	6 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:72934-6226-2	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(72934-6226)

