141060 HYDROCORTISONE 0.5% / HYDROQUINONE 4% / TRETINOIN 0.025%- 141060 hydrocortisone 0.5% / 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.

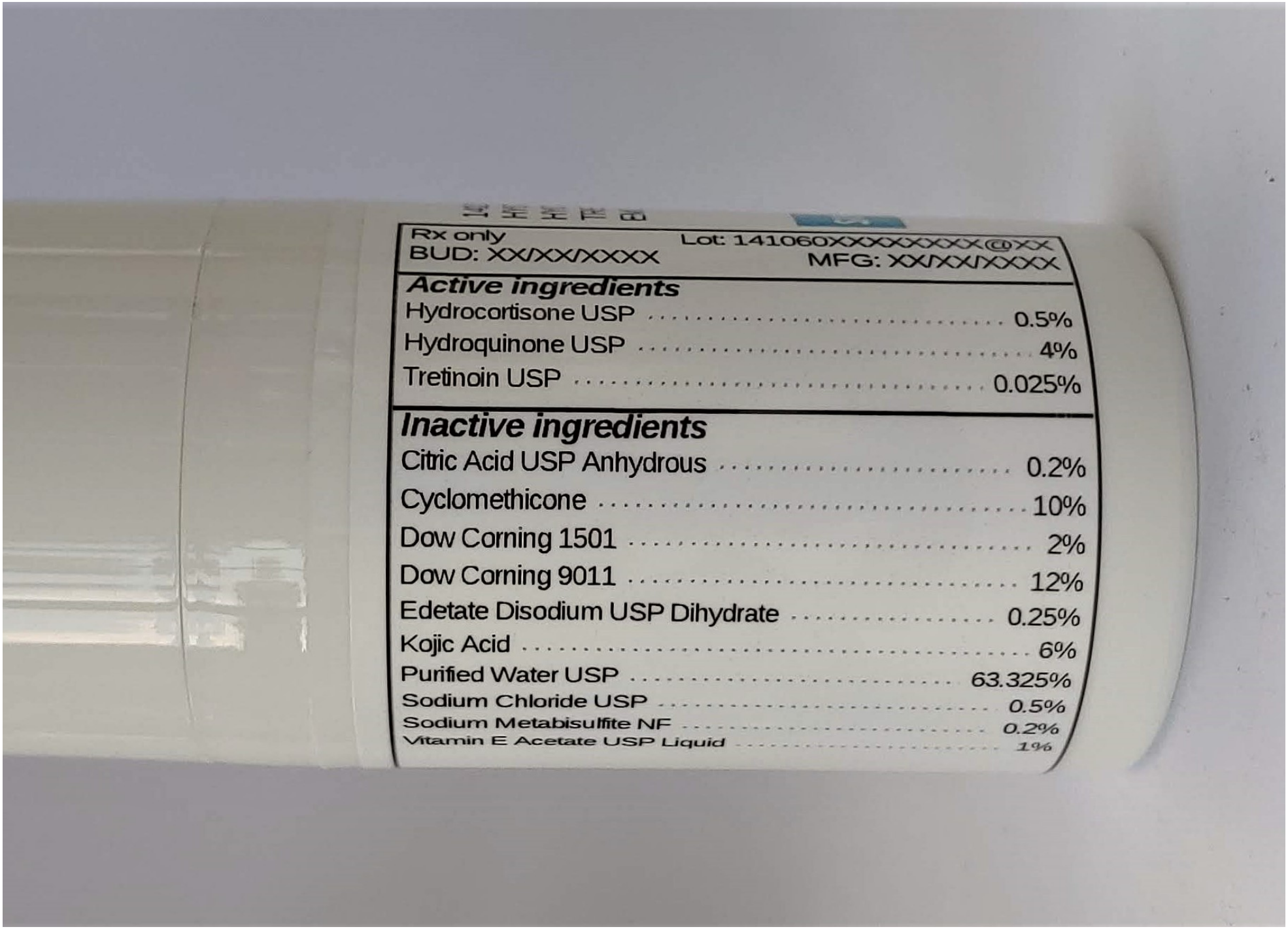
141060 HYDROCORTISONE 0.5% / HYDROQUINONE 4% / TRETINOIN 0.025%

Directions for use

Sincerus Florida, LLC adverse reactions


Directions for use
As directed by Physician.
Apply topically. For external use o
Store at controlled room tempera
Sincerus Florida, LLC
3265 W McNab Rd, Pompano B
To report suspected adverse reac
Sincerus Florida, LLC at (800) 60
at www.FDA.gov/MedWatch or (8
Office use only. Not for

Active, inactive


Rx only
Lot: 141060XXXXXXXXX@XX
BUD: XX/XX/XXXX
MFG: XX/XX/XXXX
Active ingredients
Hydrocortisone USP 0.5%
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%
Kojic Acid 6%
Purified Water USP 63.325%
Sodium Chloride USP 0.5%
Sodium Metabisulfite NF 0.2%
Vitamin E Acetate USP Liquid 1%

NDC 72934-6232-2 141060 HYDROCORTISONE 0.5% / HYDROQUINONE 4% / TRETINOIN 0.025% Emulsion 30 gm

141060 HYDROCORTISONE 0.5% / HYDROQUINONE 4% / TRETINOIN 0.025%

141060 hydrocortisone 0.5% / hydroquinone 4% / tretinoin 0.025% emulsion

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-6232
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
HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE	0.5 g in 100 g
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	4 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:72934-6232-2	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(72934-6232)

Revised: 7/2020 Sincerus Florida, LLC