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.

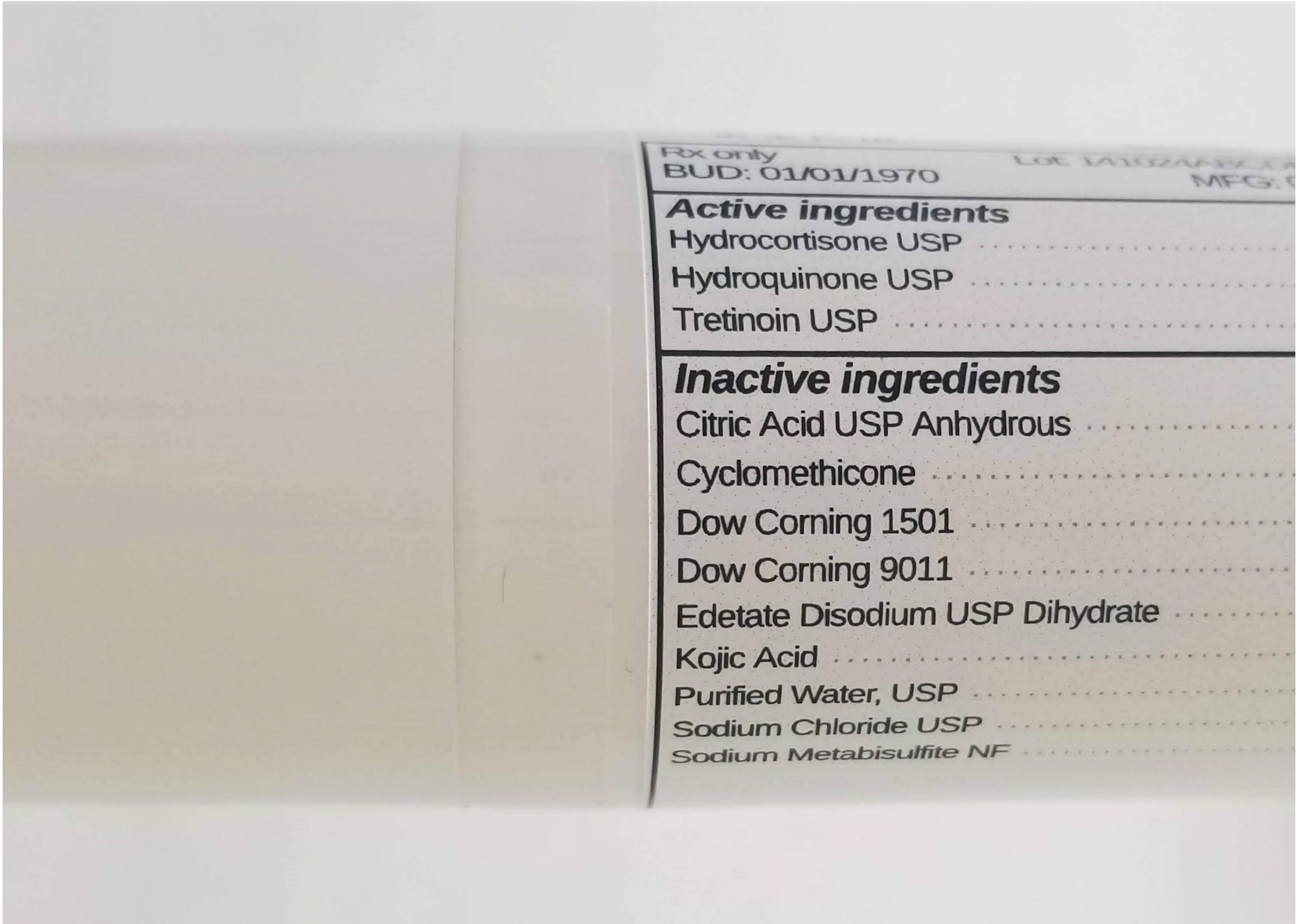
HYDROCORTISONE 0.5% / HYDROQUINONE 6% / TRETINOIN 0.025%

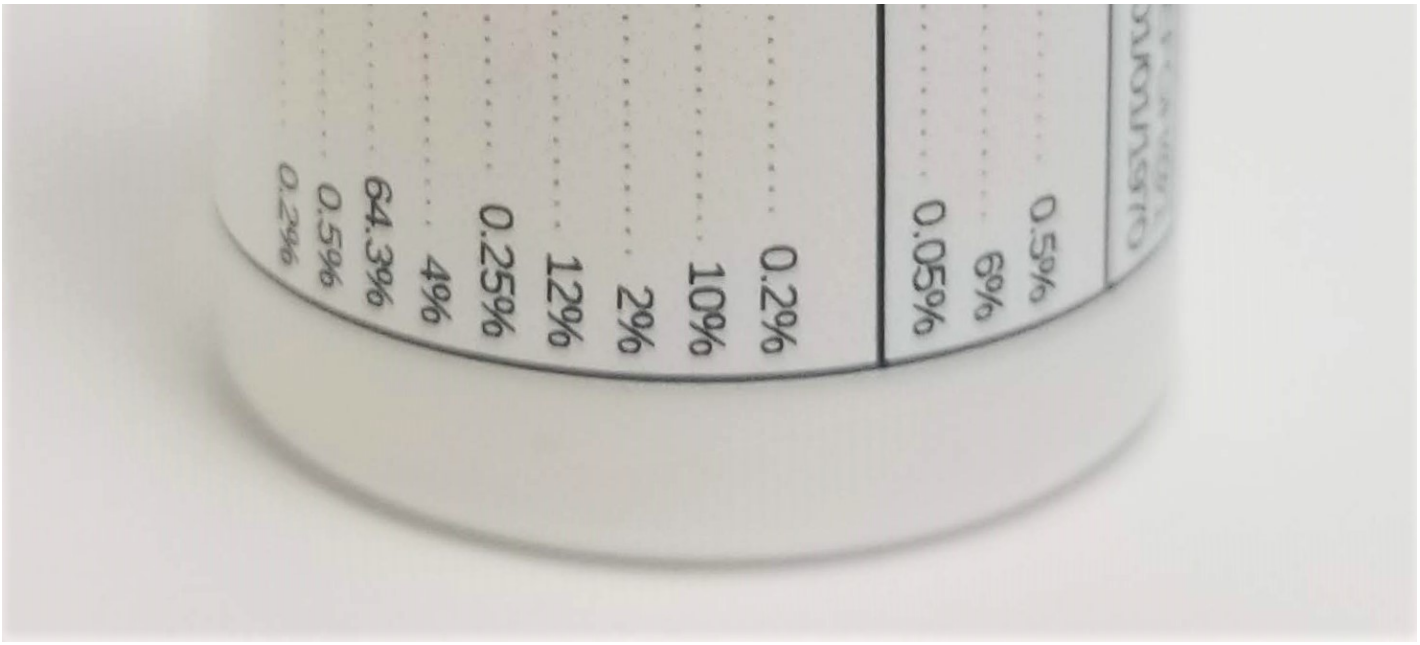
Directions for use

Sincerus Florida, LLC adverse reactions.


Directions f
As directed by Ph
Apply topically. F
Store at controlle
Sincerus Florida
3265 W McNab
To report suspec
Sincerus Florida
at www.FDA.gov
Office

Active, Inactive


Rx only
BUD: 01/01/1970
Active ingredients
Hydrocortisone USP
Hydroquinone USP
Tretinoin USP
Inactive ingredients
Citric Acid USP Anhydrous
Cyclomethicone
Dow Corning 1501
Dow Corning 9011
Edetate Disodium USP Dihydrate
Kojic Acid
Purified Water, USP
Sodium Chloride USP
Sodium Metabisulfite NF



NDC 72934-6105-2 HYDROCORTISONE USP 0.5% / HYDROQUINONE USP 6% / TRETINOIN USP 0.05%. Emulsion 30 gm

Rx only
BUD: 01/01/1970

Lot: 1410244ABCDEFGH@1
MFG: 01/01/1970

NDC 72934-6105-2

HYDROCORTISONE USP 0.5%
HYDROQUINONE USP 6%
TRETINOIN USP 0.05%
EMULSION 30gm

SINCERUS FLORIDA

This is a compounded drug.

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:72934-6105
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	0.05 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:72934-6105-2	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(72934-6105)

Revised: 4/2019 Sincerus Florida, LLC