HYDROCORTISONE 2.5% / LEVOCETIRIZINE DIHYDROCHLORIDE 2%- hydrocortisone 2.5% / levocetirizine dihydrochloride 2% cream
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% / LEVOCETIRIZINE DIHYDROCHLORIDE 2%

Directions for use

Sincerus Florida, LLC. Adverse reactions

Directions for use

As directed by Physician.

Apply topically. For external use only. W

Store at controlled room temperature (2

Sincerus Florida, LLC (800

3265 W McNab Rd, Pompano Beach

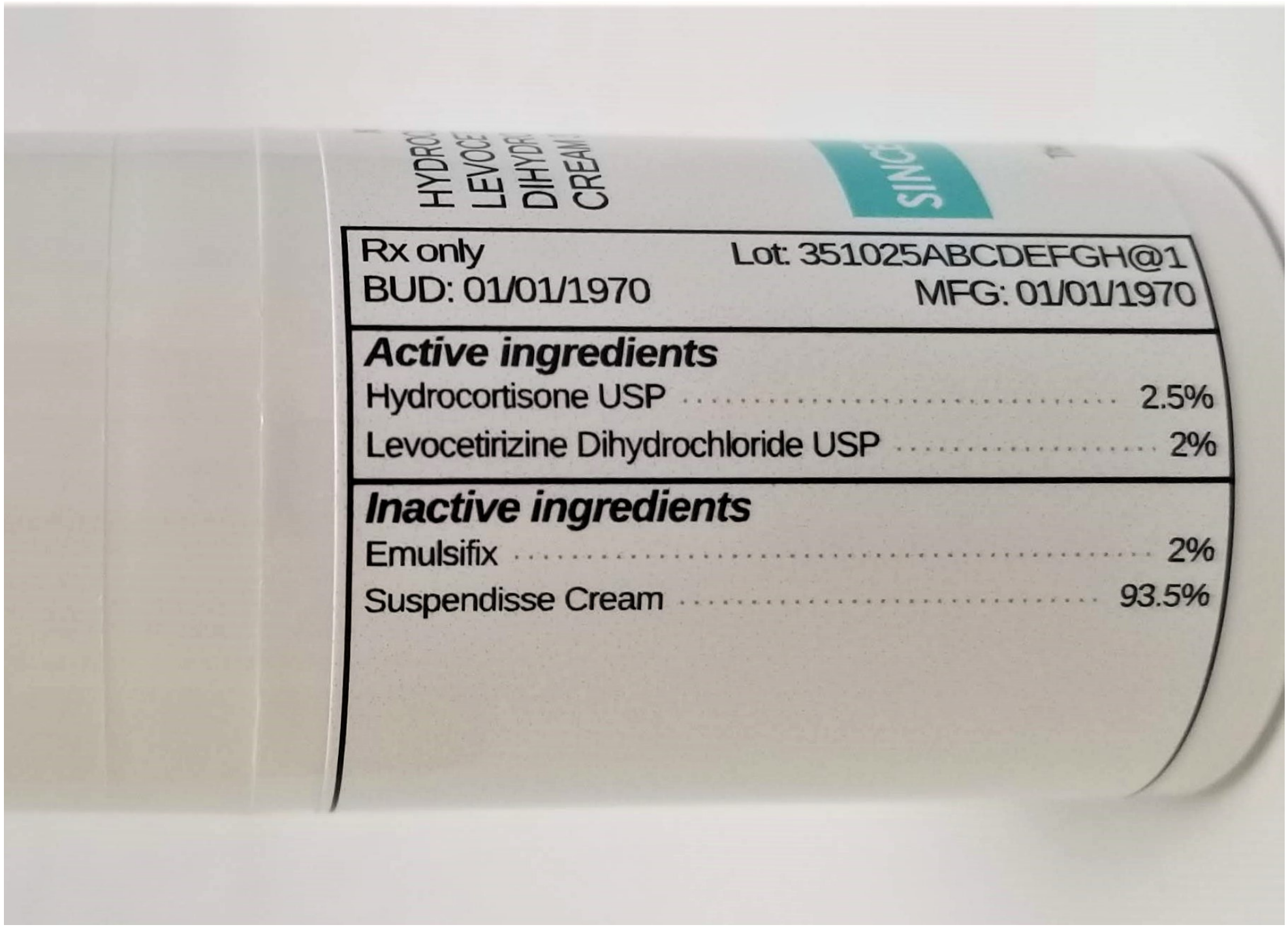
To report suspected adverse reactions

Sincerus Florida, LLC at (800) 604-503

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

Office use only. Not for resa

Active, inactive


HYDRO
LEVOCE
DIHYDR
CREAM
SINC
Rx only
BUD: 01/01/1970
Lot: 351025ABCDEFGH@1
MFG: 01/01/1970
Active ingredients
Hydrocortisone USP 2.5%
Levocetirizine Dihydrochloride USP 2%
Inactive ingredients
Emulsifix 2%
Suspendisse Cream 93.5%

NDC 72934- 2113-2 HYDROCORTISONE 2.5% / LEVOCETIRIZINE DIHYDROCHLORIDE 2%. Gel 30gm.


NDC 72934-2113-2
HYDROCORTISONE USP 2.5%
LEVOCETIRIZINE
DIHYDROCHLORIDE USP 2%
CREAM 30gm
SINCERUS FLORIDA
This is a compounded drug.
Made in USA

HYDROCORTISONE 2.5% / LEVOCETIRIZINE DIHYDROCHLORIDE 2%

hydrocortisone 2.5% / levocetirizine dihydrochloride 2% cream

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-2113
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE	2.5 g in 100 g
LEVOCETIRIZINE DIHYDROCHLORIDE (UNII: SOD6A38AGA) (LEVOCETIRIZINE - UNII:6U5EA9RT2O)	LEVOCETIRIZINE DIHYDROCHLORIDE	2 g in 100 g

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72934-2113-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/15/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/15/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2113)

Revised: 5/2019 Sincerus Florida, LLC