HYALURONIC ACID SODIUM SALT 0.5% / TAZAROTENE 0.1%- hyaluronic acid sodium salt 0.5% / tazarotene 0.1% 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.

HYALURONIC ACID SODIUM SALT 0.5% / TAZAROTENE 0.1%

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

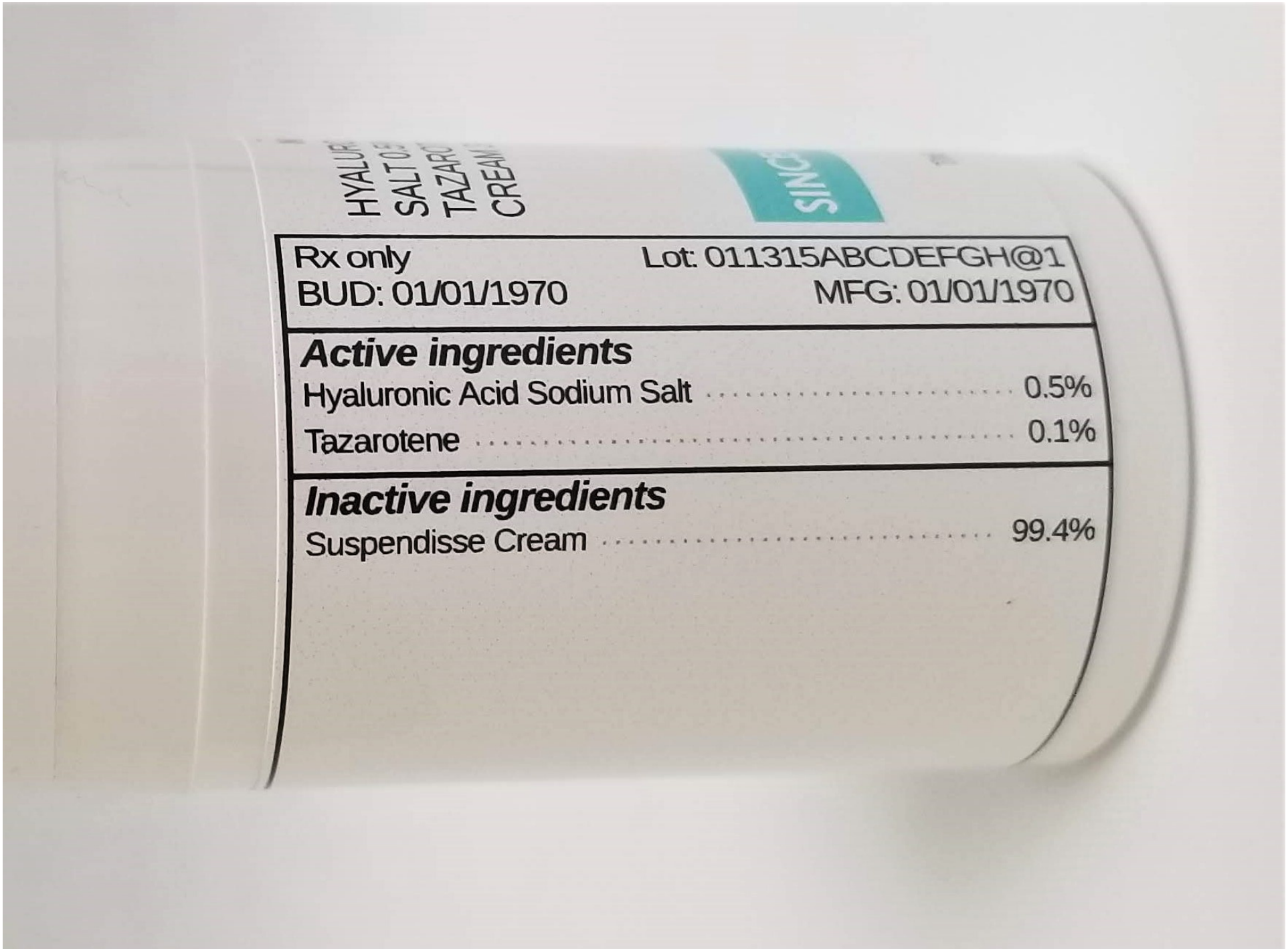
To report suspected adverse reactions

Sincerus Florida, LLC at (800) 604-503

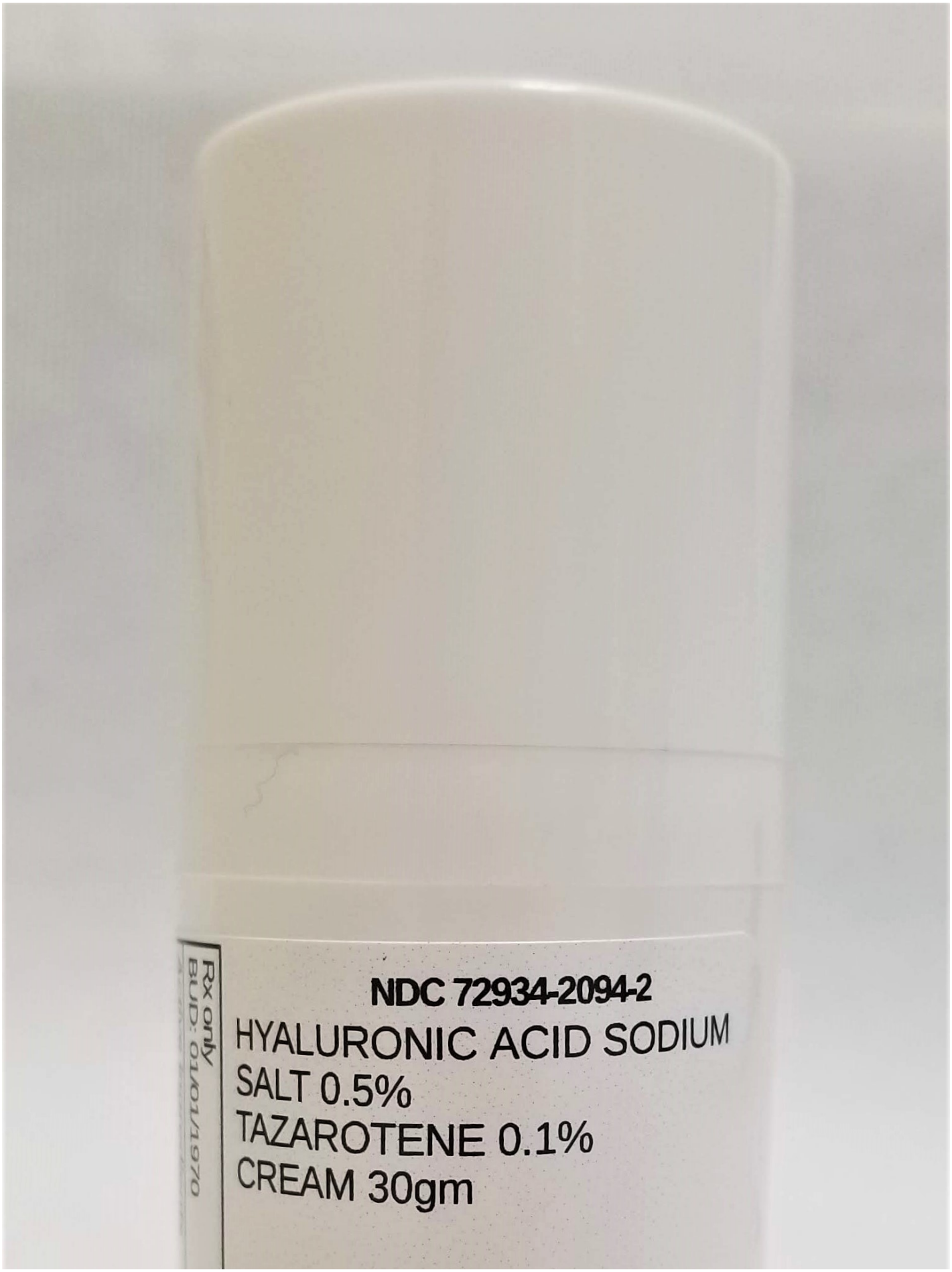
at www.FDA.gov/MedWatch or (800) F

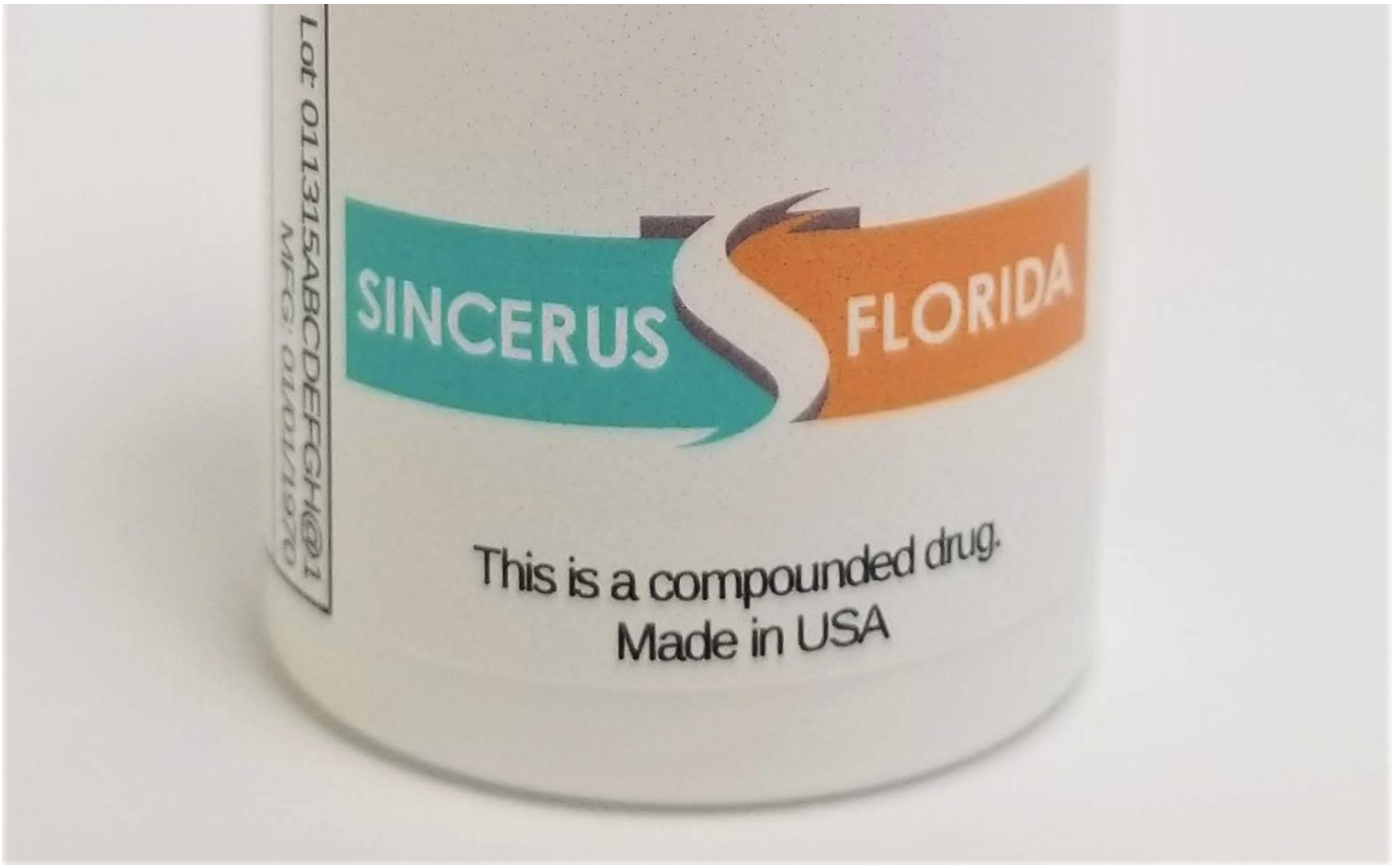
Office use only. Not for resa

Active, inactive


HYALUR
SALT 0.5
TAZARO
CREAM
SING
Rx only
Lot: 011315ABCDEFGH@1
BUD: 01/01/1970
MFG: 01/01/1970
Active ingredients
Hyaluronic Acid Sodium Salt 0.5%
Tazarotene 0.1%
Inactive ingredients
Suspendisse Cream 99.4%

HYALURONIC ACID SODIUM SALT 0.5% / TAZAROTENE 0.1%. Cream 30 gm





HYALURONIC ACID SODIUM SALT 0.5% / TAZAROTENE 0.1%

hyaluronic acid sodium salt 0.5% / tazarotene 0.1% cream

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-2094
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYALURONATE SODIUM (UNII: YSE9PPT4TH) (HYALURONIC ACID - UNII:S270N0TRQY)	HYALURONATE SODIUM	0.5 g in 100 g
TAZAROTENE (UNII: 81BDR9Y8PS) (TAZAROTENE - UNII:81BDR9Y8PS)	TAZAROTENE	0.1 g in 100 g

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:72934-2094-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/07/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/07/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2094)

Revised: 5/2019 Sincerus Florida, LLC