

**CIMETIDINE 10% / LIDOCAINE 5% / SALICYLIC ACID 40%- cimetidine 10% / lidocaine 5% / salicylic acid 40% 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](#).*

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**CIMETIDINE 10% / LIDOCAINE 5% / SALICYLIC ACID 40%**

**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](http://www.FDA.gov/MedWatch) or (800) FDA-1088.

Office use only. Not for resale.



**Sincerus Florida, LLC adverse reactions.**



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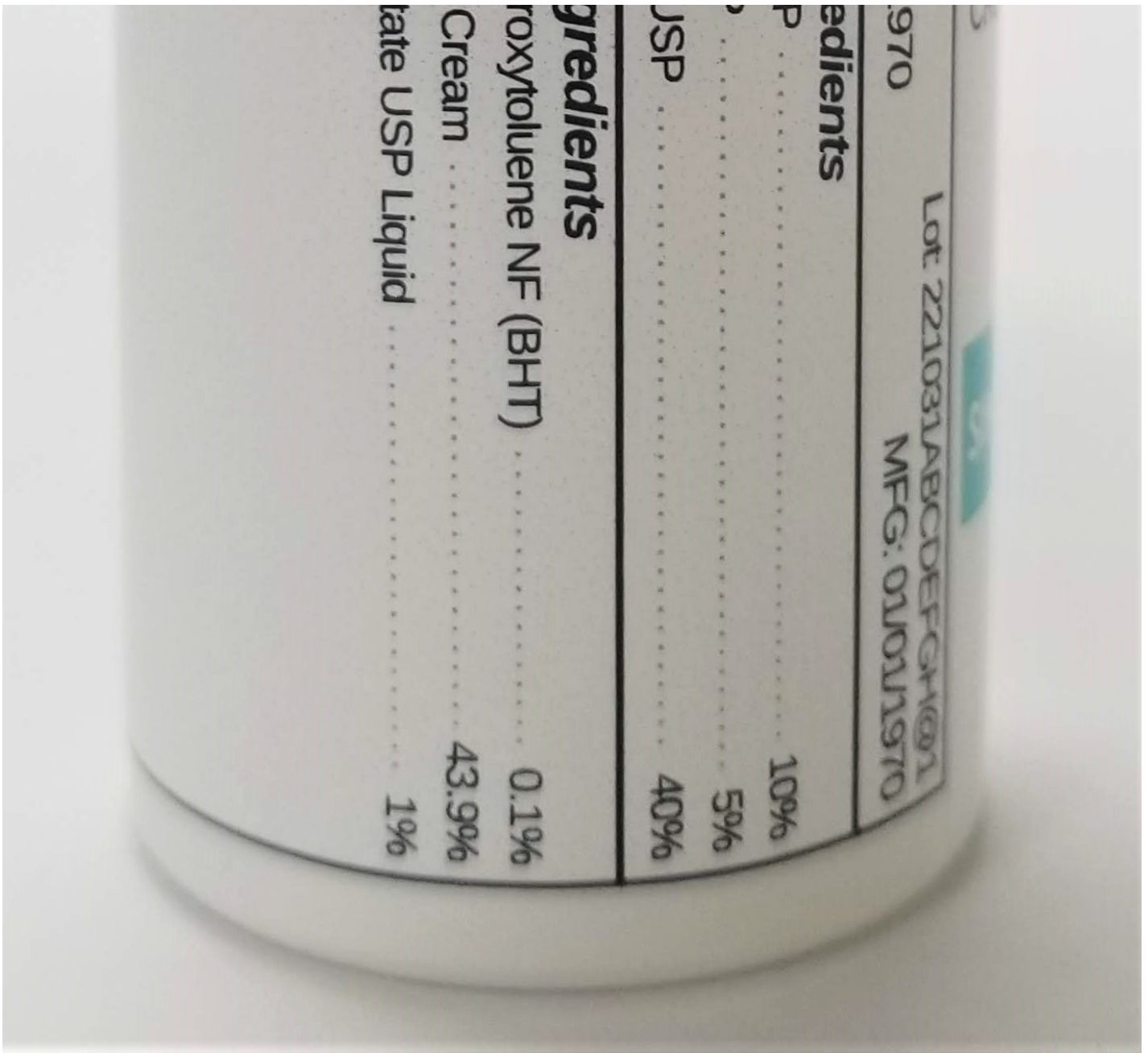
Active, inactive



Rx only  
BUD: 01/01/11

**Active ingredients**  
Cimetidine USF  
Lidocaine USF  
Salicylic Acid U

**Inactive ingredients**  
Butylated Hydroxy  
Salt Stable Ls  
Vitamin E Acetate



**NDC 72934-2046-2 CIMETIDINE USP 10% / LIDOCAINE USP 5% / SALICYLIC ACID USP 40%  
Cream 30 gm**



Rx only  
BUD: 01/01/1970

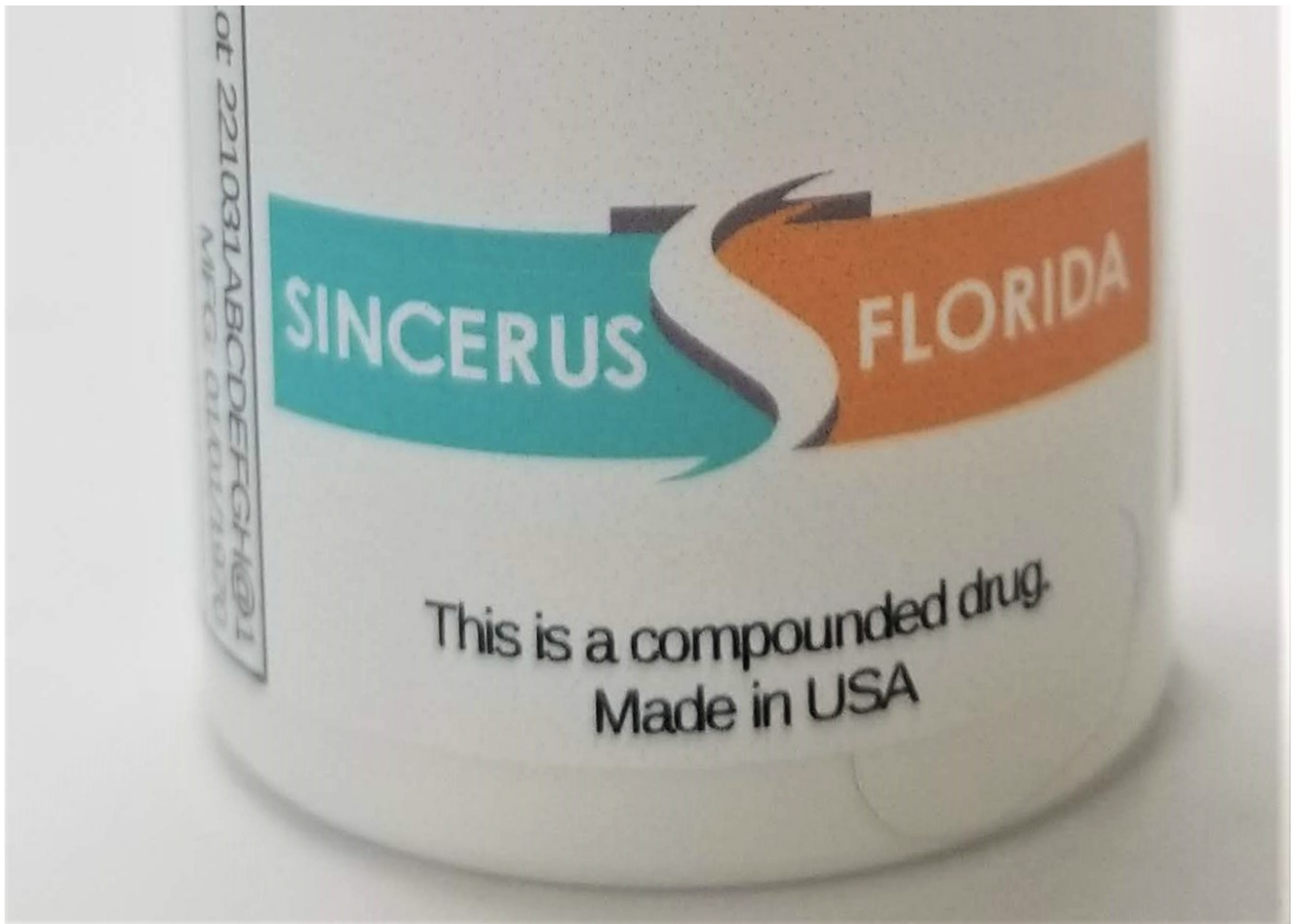
**NDC 72934-2046-2**

**CIMETIDINE USP 10%**

**LIDOCAINE USP 5%**

**SALICYLIC ACID USP 40%**

**CREAM 30gm**



## CIMETIDINE 10% / LIDOCAINE 5% / SALICYLIC ACID 40%

cimetidine 10% / lidocaine 5% / salicylic acid 40% cream

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-2046
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	40 g in 100 g
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	5 g in 100 g
CIMETIDINE (UNII: 80061L1WGD) (CIMETIDINE - UNII:80061L1WGD)	CIMETIDINE	10 g in 100 g

### Product Characteristics

<b>Color</b>	yellow	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			



**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-2046-2	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(72934-2046)

Revised: 4/2019

Sincerus Florida, LLC