LIDOCAINE 7% / TETRACAINE 7%- lidocaine 7% / tetracaine 7% ointment
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.

LIDOCAINE 7% / TETRACAINE 7%

Directions for use

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As directed by Physician.
Apply topically. For external use only. Wash hands
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 or (800) FDA-1088.

Office use only. Not for resale.

Sincerus Florida, LLC. Adverse reactions

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Active, inactive

NDC 72

LIDOCAINE US
TETRACAINE
OINTMENT 1.2

Rx only
BUD: 01/01/1970

Lot: 201010

Active ingredients

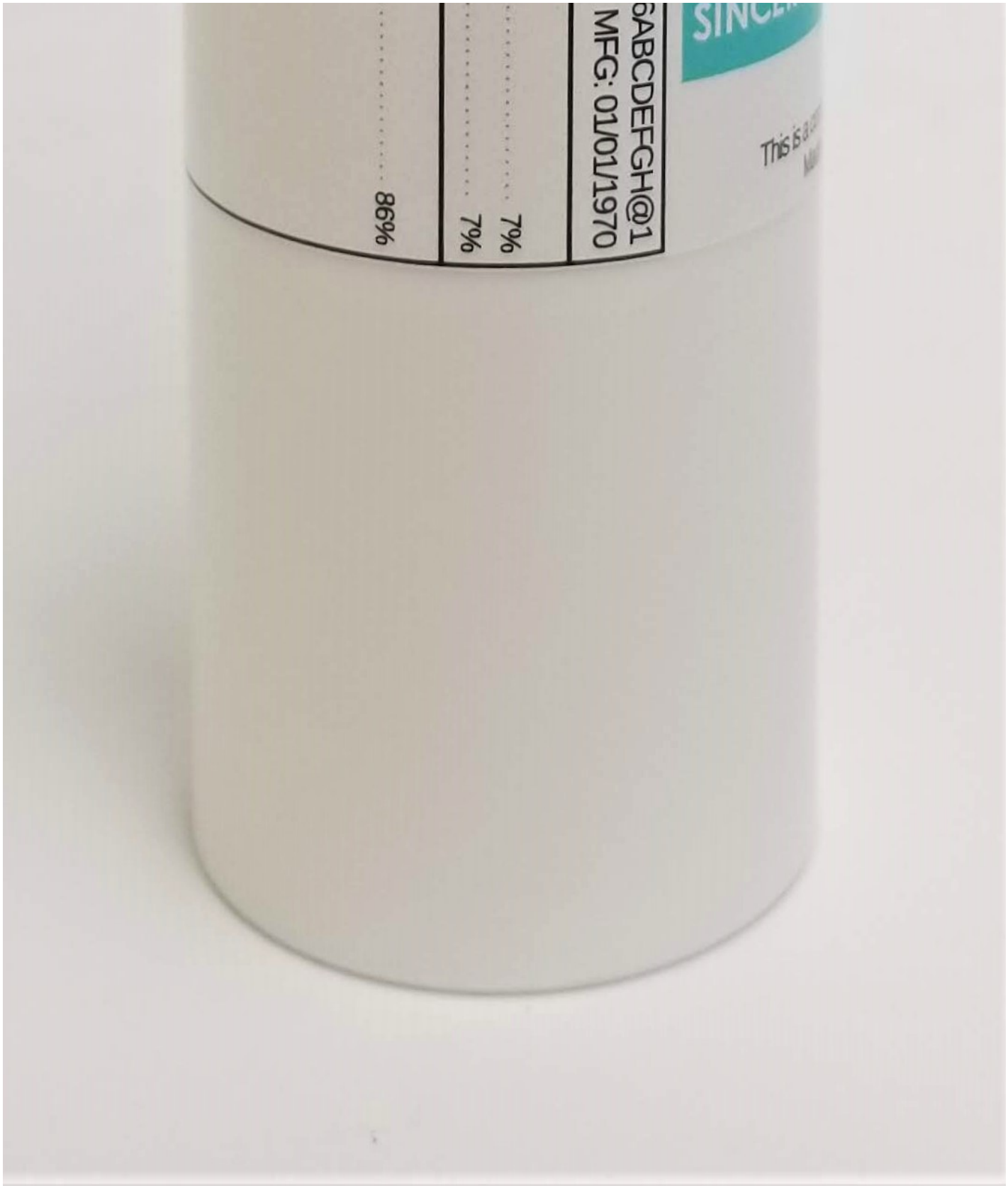
Lidocaine USP .....

Tetracaine USP .....

Inactive ingredients

Oleabase Plasticized .....

DANGERUS



NDC 72934- 5142-4 LIDOCAINE 7% / TETRACAINE 7%. Ointment 120gm

Rx only
BUD: 01/01/1970
Lot: 20101 6ABCDEFGH@1
MFG: 01/01/1970
Active ingredients
Lidocaine USP
Tetracaine USP

NDC 72934-5142-4

LIDOCAINE USP 7%
TETRACAINE USP 7%
OINTMENT 120gm

SINCERUS FLORIDA

This is a compounded drug.
Made in USA

LIDOCAINE 7% / TETRACAINE 7%

lidocaine 7% / tetracaine 7% ointment

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-5142
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	7 g in 100 g
TETRACAINE (UNII: 0619F35CGV) (TETRACAINE - UNII:0619F35CGV)	TETRACAINE	7 g in 100 g

Product Characteristics			
Color	white (clear ointment)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-5142-4	120 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/17/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/17/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-5142)

Revised: 5/2019 Sincerus Florida, LLC