BENZOCAINE 20% / LIDOCAINE 10% / TETRACAINE 10%- benzocaine 20% / lidocaine 10% / tetracaine 10% 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.

BENZOCAINE 20% / LIDOCAINE 10% / TETRACAINE 10%

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

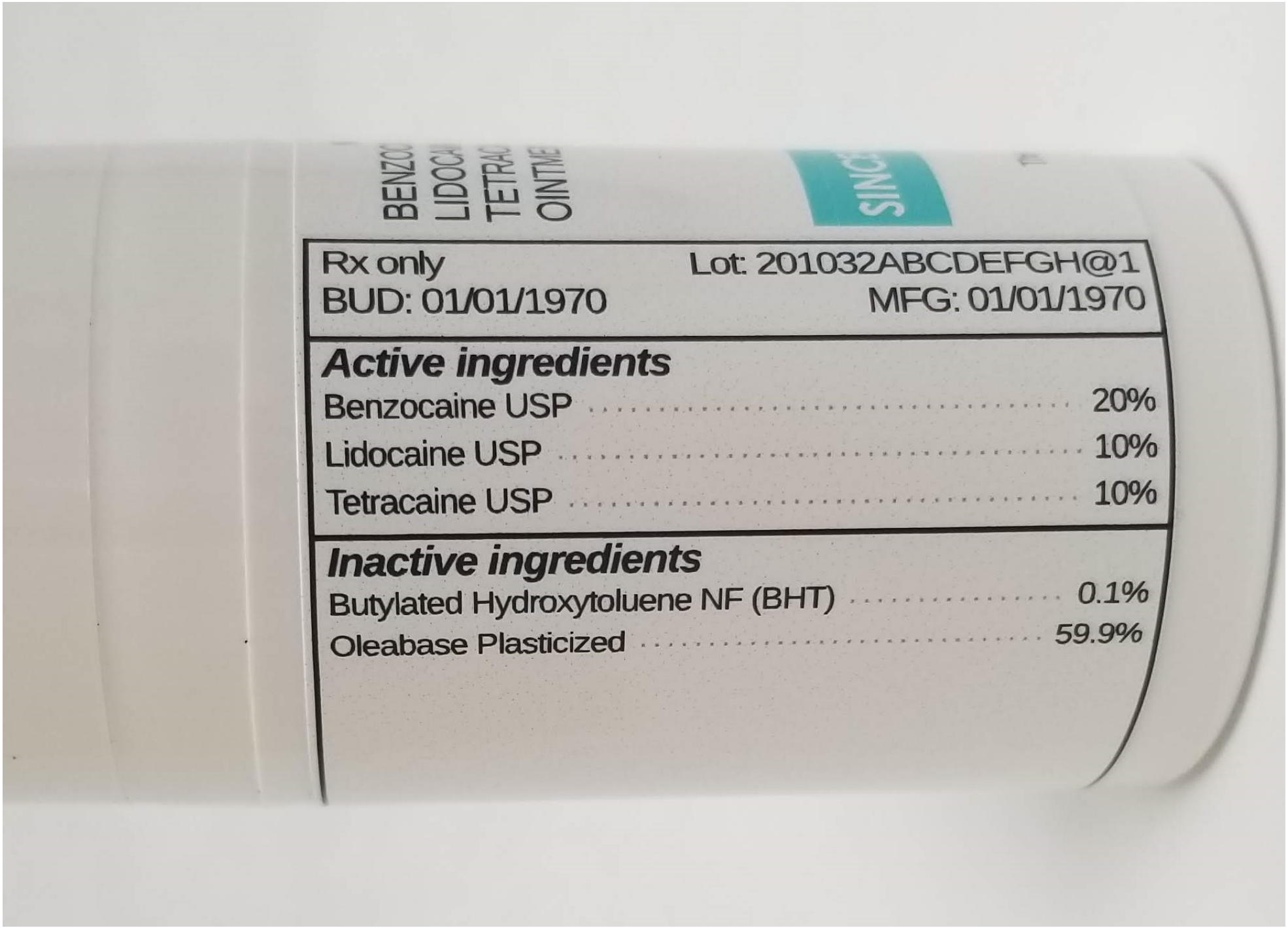
To report suspected adverse reactions

Sincerus Florida, LLC at (800) 604-503

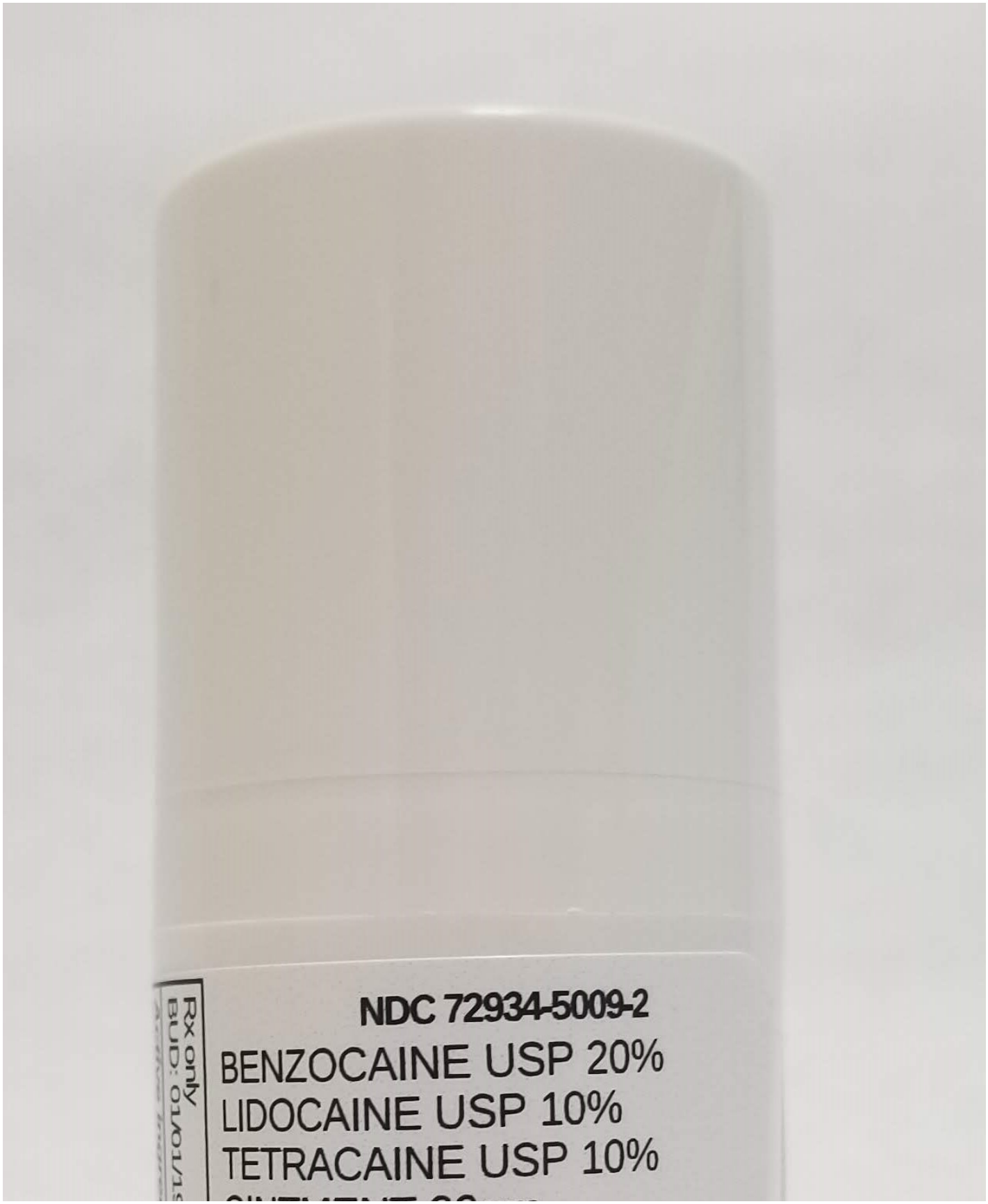
at www.FDA.gov/MedWatch or (800) F

Office use only. Not for resa

Active, inactive


BENZO
LIDOCA
TETRAC
OINTME
SINC
Rx only
Lot: 201032ABCDEFGH@1
BUD: 01/01/1970
MFG: 01/01/1970
Active ingredients
Benzocaine USP 20%
Lidocaine USP 10%
Tetracaine USP 10%
Inactive ingredients
Butylated Hydroxytoluene NF (BHT) 0.1%
Oleabase Plasticized 59.9%

NDC 72934- 5040-2 BENZOCAINE USP 20% / LIDOCAINE USP 10% / TETRACAINE USP 10%. Ointment 120gm



BENZOCAINE 20% / LIDOCAINE 10% / TETRACAINE 10%

benzocaine 20% / lidocaine 10% / tetracaine 10% ointment

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-5009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TETRACAINE (UNII: 0619F35CGV) (TETRACAINE - UNII:0619F35CGV)	TETRACAINE	10 g in 100 g
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	20 g in 100 g
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	10 g in 100 g

Product Characteristics			
Color	yellow (off white ointment)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-5009-2	30 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(72934-5009)

Revised: 5/2019 Sincerus Florida, LLC