

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



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As directed by Physician.

Apply topically. For external use

Store at controlled room temp

Sincerus Florida, LLC

3265 W McNab Rd, Pompano

To report suspected adverse re

Sincerus Florida, LLC at (800) 6

at www.FDA.gov/MedWatch or

Office use only. Not for

Use only. Wash hands after use.
Temperature (20-25C).

(800) 604-5032
Beach, FL 33069
For adverse reactions, contact
604-5032, or FDA
(800) FDA-1088.
For resale.





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3265 W McNab Rd, Pompano Beach, FL 33069

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Active, inactive







**NDC 72934- 5009-4 BENZOCAINE 20% / LIDOCAINE 10% / TETRACAINE 10%. Ointment
120gm**



NDC 72934-5009-4

**BENZOCAINE USP 20%
LIDOCAINE USP 10%
TETRACAINE USP 10%
OINTMENT 120gm**

Rx only

BUD: 01/01/1970

Active Ingredients

Benzocaine USP

Lidocaine USP

Lot: 201030ABCDEF@1
MFG: 01/01/1970



This is a compounded drug.
Made in USA

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
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	20 g in 100 g
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	10 g in 100 g
TETRACAINE (UNII: 0619F35CGV) (TETRACAINE - UNII:0619F35CGV)	TETRACAINE	10 g in 100 g

Product Characteristics

Color	yellow (OFF WHITE)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-5009-4	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(72934-5009)