

LIDOCAINE 23% / TETRACAINE 7% - lidocaine 23% / 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 23% / TETRACAINE 7%

Directions for use



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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 or (800) FDA-1088.

Office use only. Not for resale.



Sincerus Florida, LLC. Adverse reactions





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Active, inactive



LIDOCAIN
TETRACA
OINTMENT

Rx only
BUD: 01/01/1970

Lot 20

Active ingredients

Lidocaine USP
Tetracaine USP

Inactive ingredients

Butylated Hydroxytoluene NF (B)
Oleabase Plasticized

SINCE

1019ABCDEF@1
MFG: 01/01/1970

..... 23%
..... 7%

HT) 0.1%
..... 69.9%

NDC 72934- 5040-4 LIDOCAINE USP 23% / TETRACAINE USP 7%. Ointment 120gm



NDC 72934-5140-4

**LIDOCAINE USP 23%
TETRACAINE USP 7%
OINTMENT 120gm**



**This is a compounded drug.
Made in USA**

Rx only
BUD: 01/01/1970
Lot 201019ABCDEF@1
MFG: 01/01/1970
Active Ingredients
Lidocaine USP



LIDOCAINE 23% / TETRACAINE 7%

lidocaine 23% / tetracaine 7% ointment

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-5140
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	23 g in 100 g
TETRACAINE (UNII: 0619F35CGV) (TETRACAINE - UNII:0619F35CGV)	TETRACAINE	7 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:72934-5140-4	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(72934-5140)

Revised: 5/2019

Sincerus Florida, LLC