LIDOCAINE 30%- lidocaine 30% 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 30%

Directions for use


Directions for use
As directed by Physician.
Apply topically. For externa
Store at controlled room te
Sincerus Florida, LLC
3265 W McNab Rd, Pom
To report suspected adve
Sincerus Florida, LLC at (
at www.FDA.gov/MedWa
Office use onl

l use only. Wash hands after use.
mperature (20-25C).

(800) 604-5032
mpano Beach, FL 33069
rse reactions, contact
800) 604-5032, or FDA
tch or (800) FDA-1088.
y. Not for resale.

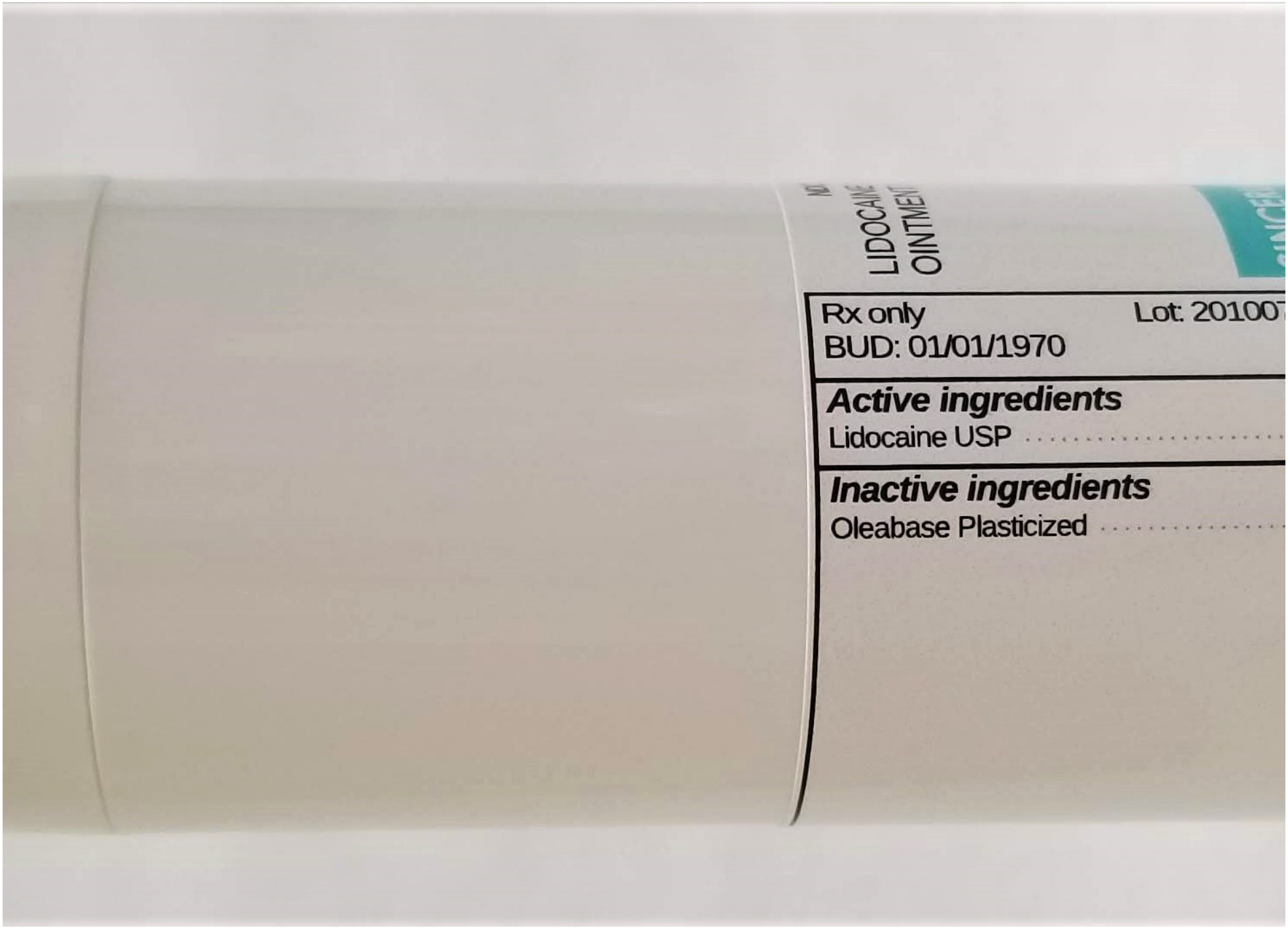
Sincerus Florida, LLC. Adverse reactions


Directions for us
As directed by Physician
Apply topically. For exter
Store at controlled room
Sincerus Florida, LLC
3265 W McNab Rd, Po
To report suspected ad
Sincerus Florida, LLC a
at www.FDA.gov/Med
Office use o

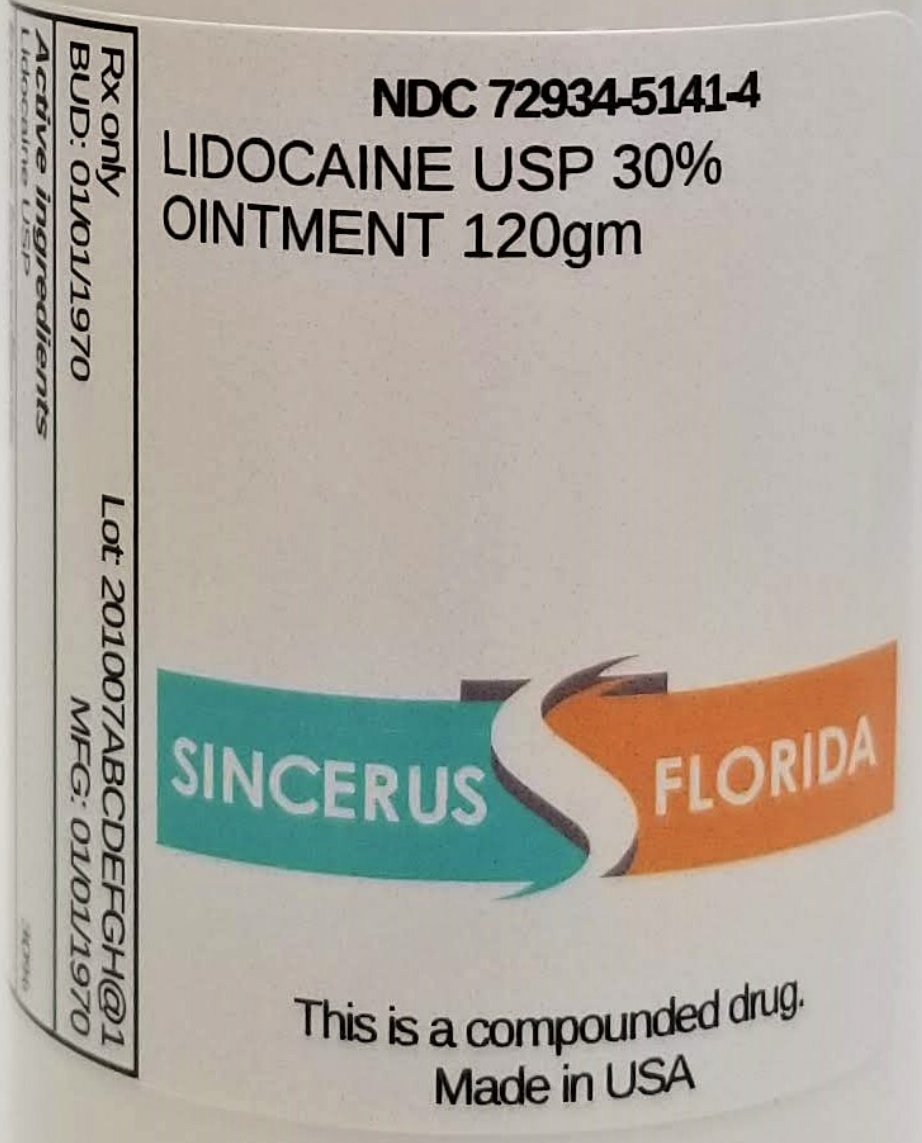
e
.
mal use only. Wash hands after use.
temperature (20-25C).

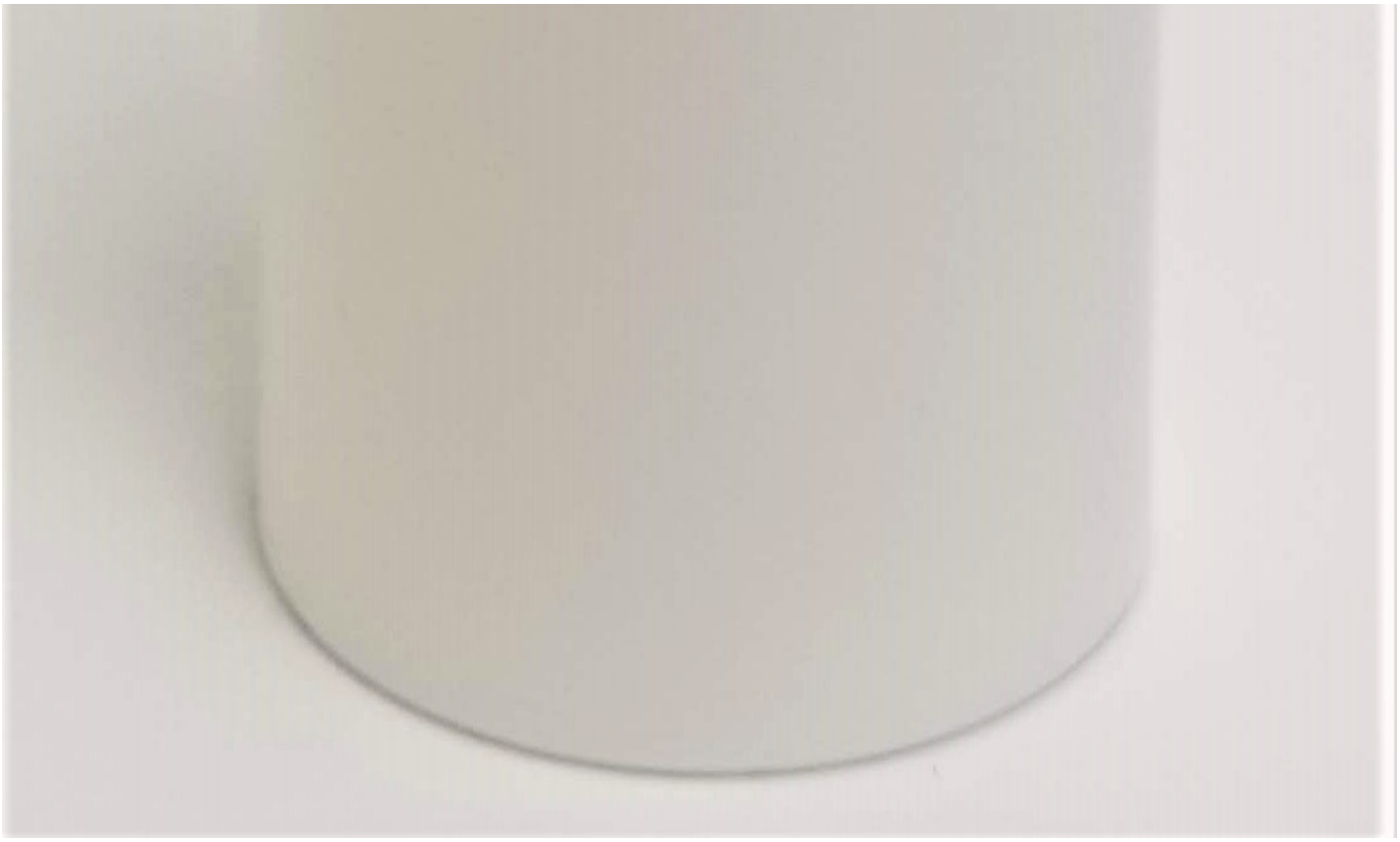
(800) 604-5032
ompano Beach, FL 33069
verse reactions, contact
at (800) 604-5032, or FDA
Watch or (800) FDA-1088.
only. Not for resale.

Active, inactive


LIDOCAINE
OINTMENT
Rx only
Lot: 201007
BUD: 01/01/1970
Active ingredients
Lidocaine USP
Inactive ingredients
Oleabase Plasticized

NDC 72934- 5141-4 LIDOCAINE 30%. Ointment 120gm


NDC 72934-5141-4
LIDOCAINE USP 30%
OINTMENT 120gm
SINCERUS FLORIDA
This is a compounded drug.
Made in USA
Rx only
BUD: 01/01/1970
Lot: 201007ABCDEFGH@1
MFG: 01/01/1970
Active ingredients
Lidocaine USP



LIDOCAINE 30%

lidocaine 30% ointment

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-5141
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	30 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:72934-5141-4	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(72934-5141)

Revised: 5/2019 Sincerus Florida, LLC