LIDOCAINE 15% / PHENYLEPHRINE HCL 0.25% / PRILOCAINE 5% - lidocaine 15% / phenylephrine hcl 0.25% / prilocaine 5% 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 USP 15% / PHENYLEPHRINE HCL USP 0.25% / PRILOCAINE USP 5%.

Director use



Directions for use

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







Active, inactive



PHENYL HYDRO PRILOC OINTNE

Rx only BUD: 01/01/1970

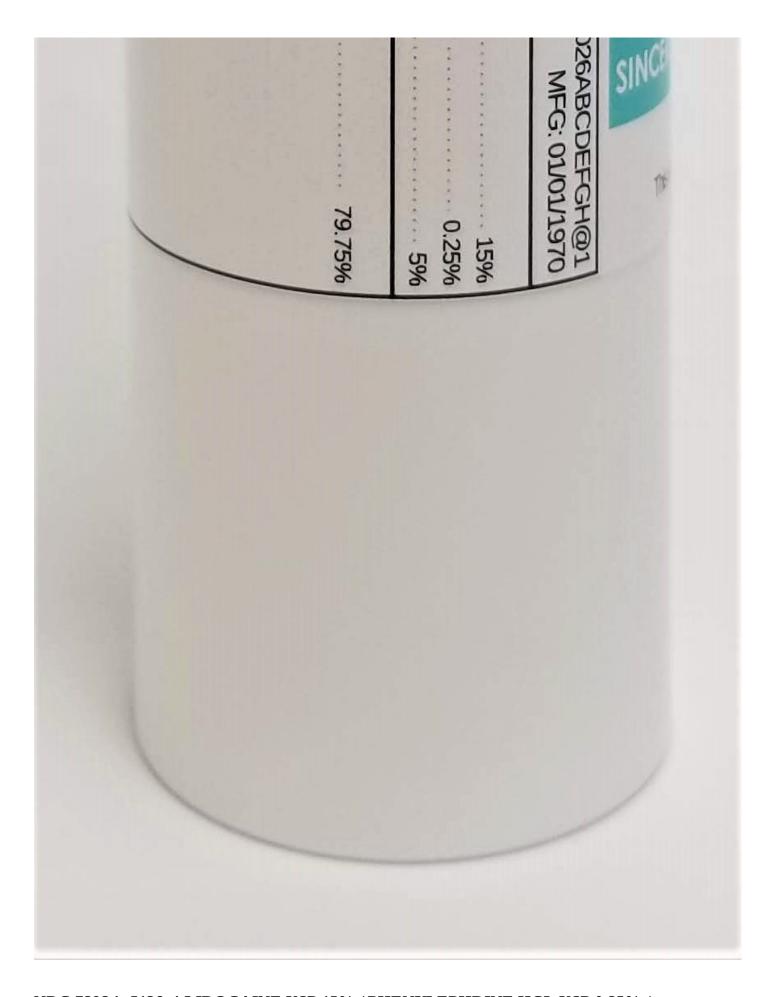
Lot: 2010

Active ingredients Lidocaine USP

Phenylephrine Hydrochloride USP Prilocaine USP

Inactive ingredients

Oleabase Plasticized ...



NDC 72934- 5138-4 LIDOCAINE USP 15% / PHENYLEPHRINE HCL USP 0.25% /

PRILOCAINE USP 5%. Ointment 120gm



BUD: 01/01/1970

Lot: 201026

NDC 72934-5138-4
LIDOCAINE USP 15%
PHENYLEPHRINE
HYDROCHLORIDE USP 0.25%
PRILOCAINE USP 5%
OINTMENT 120gm



LIDOCAINE 15% / PHENYLEPHRINE HCL 0.25% / PRILOCAINE 5%

lidocaine 15% / phenylephrine hcl 0.25% / prilocaine 5% ointment

Product Information

	Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-5138
l	Route of Administration	TOPICAL		

ı	Active Ingredient/Active Moiety		
ı	Ingredient Name	Basis of Strength	Strength
ı	LIDO CAINE (UNII: 98PI200987) (LIDO CAINE - UNII:98PI200987)	LIDOCAINE	15 g in 100 g

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:72934-5138-	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(72934-5138)	

Revised: 5/2019 Sincerus Florida, LLC