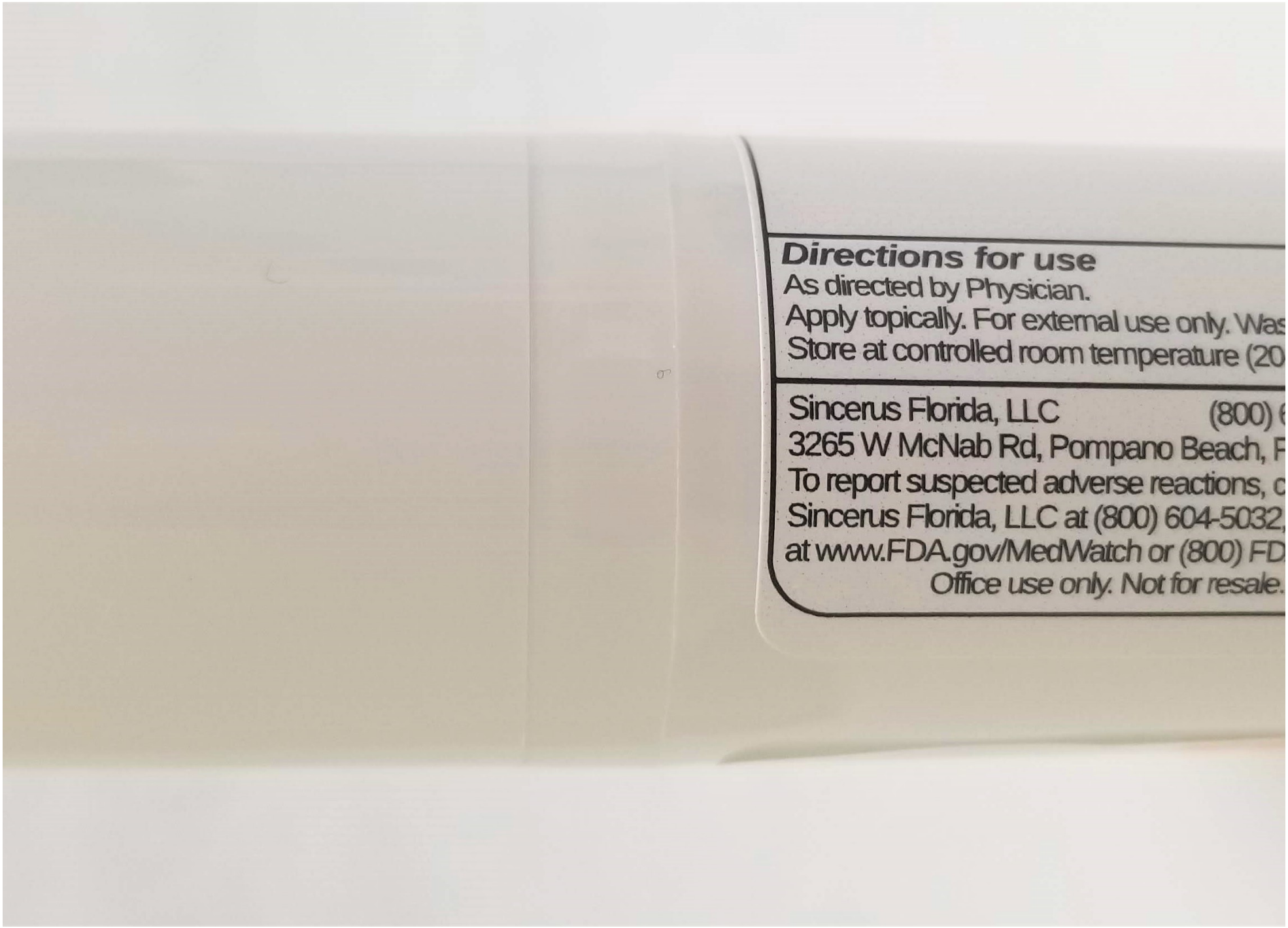
CICLOPIROX 3%- ciclopirox 3% gel
Sincerus Florida, LLC

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.

CICLOPIROX 3%

Directions for use

Sincerus Florida, LLC. Adverse reactions



Directions for use
As directed by Physician.
Apply topically. For external use only. Was
Store at controlled room temperature (20

Sincerus Florida, LLC (800) 6
3265 W McNab Rd, Pompano Beach, F
To report suspected adverse reactions, c
Sincerus Florida, LLC at (800) 604-5032,
at www.FDA.gov/MedWatch or (800) FD
Office use only. Not for resale.

Active, inactive

NDC 72934- 1042-2 CICLOPIROX OLAMINE USP 3%. Gel 30 gm


NDC 72934-1042-2
CICLOPIROX OLAMINE USP

CICLOPIROX 3%

ciclopirox 3% gel

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-1042
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CICLOPIROX OLAMINE (UNII: 50MD4SB4AP) (CICLOPIROX - UNII:19W019ZDRJ)	CICLOPIROX	3 g in 100 g

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72934-1042-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/20/2019	

Labeler - Sincerus Florida, LLC (080105003)

Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-1042)

Revised: 5/2019 Sincerus Florida, LLC