

**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](#).*

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**CICLOPIROX 3%**

**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](http://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**



CICLOPIROX  
3%  
GEL 30g

SIN

Rx only  
BUD: 01/01/1970

Lot: 071012ABCDEF@1  
MFG: 01/01/1970

**Active ingredients**

Ciclopirox Olamine USP ..... 3%

**Inactive ingredients**

Ethyl Alcohol USP ..... 4%

Krisigel 100 ..... 2%

Suspendisse Gel ..... 91%

Rx only  
NDC 72934-1042-2

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**CICLOPIROX OL AMINE 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