

**BRIMONIDINE TARTRATE 0.25% / POTASSIUM AZELOYL DIGLYCINATE 8% -  
brimonidine tartrate 0.25% / potassium azeloyl diglycinate 8% 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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**BRIMONIDINE TARTRATE 0.25% / POTASSIUM AZELOYL DIGLYCINATE 8%**

**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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Sincerus Florida, LLC (800

3265 W McNab Rd, Pompano Beach

To report suspected adverse reactions

Sincerus Florida, LLC at (800) 604-503

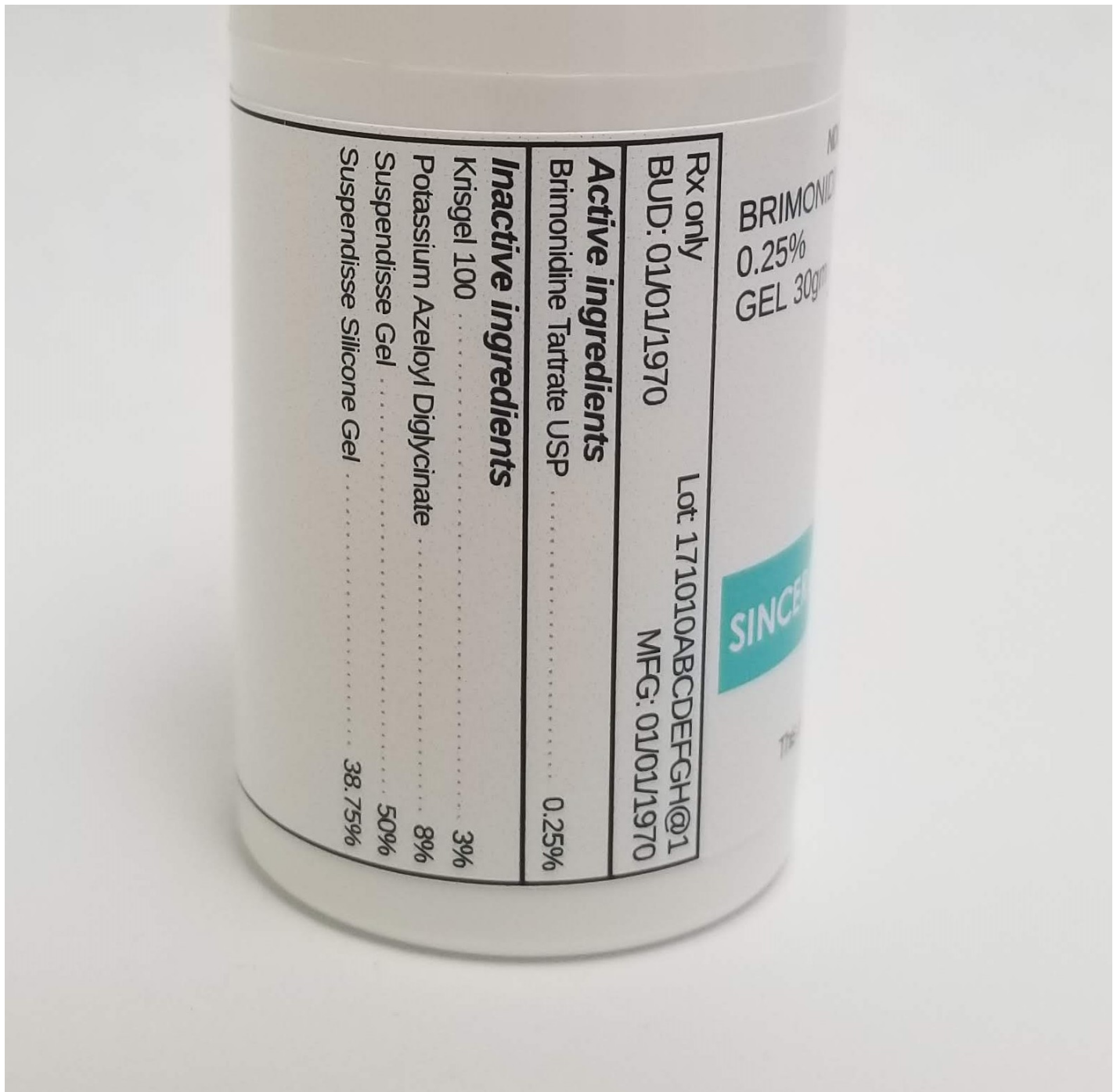
at [www.FDA.gov/MedWatch](http://www.FDA.gov/MedWatch) or (800) F

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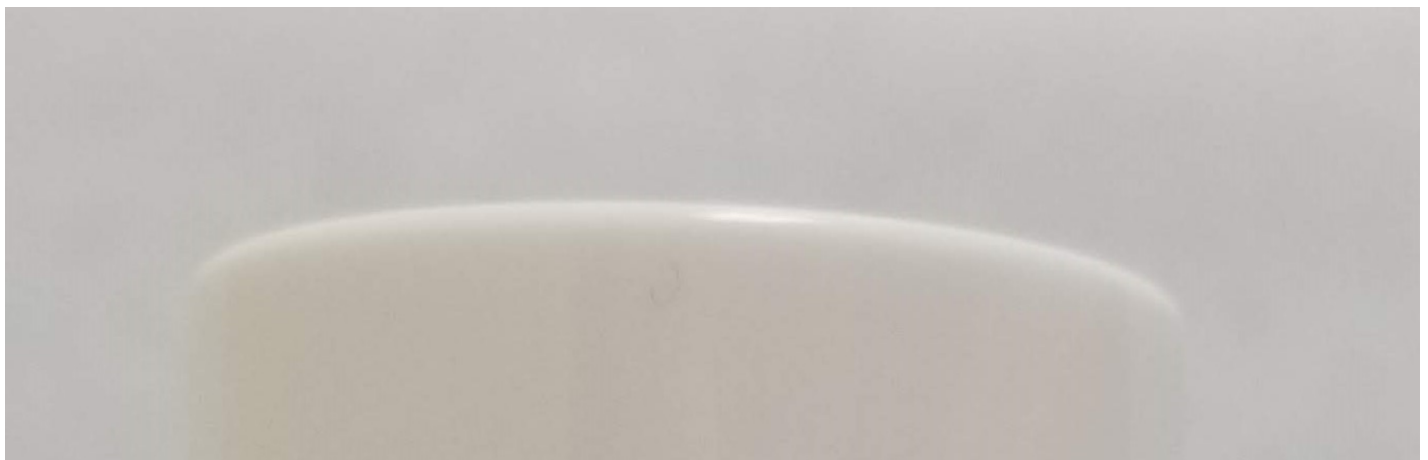


**Active, inactive**





NDC 72934- 1027-2 BRIMONIDINE TARTRATE USP 0.25% . Gel 30gm



Rx only  
NDC 72934-1027-2

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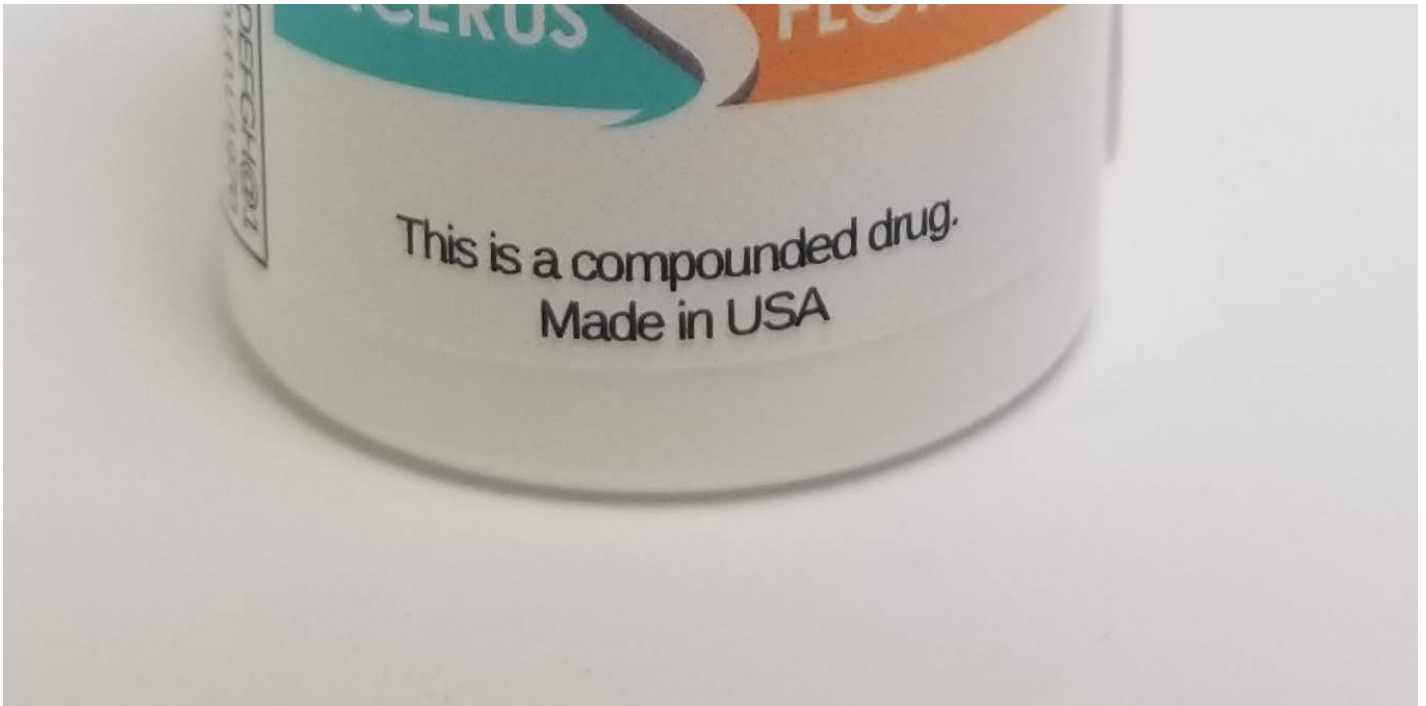
**BRIMONIDINE TARTRATE USP**

0.25%

GEL 30gm

Lot 171010AEC  
EXP 03/2023





**BRIMONIDINE TARTRATE 0.25% / POTASSIUM AZELOYL DIGLYCINATE 8%**

brimonidine tartrate 0.25% / potassium azeloyl diglycinate 8% gel

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72934-1027
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
BRIMONIDINE TARTRATE (UNII: 4S9CL2DY2H) (BRIMONIDINE - UNII:E6GNX3HHTE)	BRIMONIDINE TARTRATE	0.25 g in 100 g

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-1027-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/16/2019	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/16/2019	

**Labeler** - Sincerus Florida, LLC (080105003)

## Establishment

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-1027)

Revised: 5/2019

Sincerus Florida, LLC