TACROLIMUS 0.1% - tacrolimus 0.1% solution
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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TACROLIMUS 0.1%

Directions for use
Sincerus Florida, LLC. Adverse reactions
NDC 72934-4173-8. TACROLIMUS MONOHYDRATE USP 0.1%. Solution 60 gm.
NDC 72934-4173-8
TACROLIMUS MONOHYDRATE
USP 0.1%
SOLUTION 60gm

SINCERUS FLORIDA

This is a compounded drug.
Made in USA
TACROLIMUS 0.1%
tacrolimus 0.1% solution

Product Information
Product Type: HUMAN PRESCRIPTION DRUG
Item Code (Source): NDC:72934-4173
Route of Administration: TOPICAL

Active Ingredient/Active Moiety
<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>TACROLIMUS</td>
<td>TACROLIMUS ANHYDROUS</td>
<td>0.1 g in 100 g</td>
</tr>
<tr>
<td>(UNII: WM0HAQ4WNM)</td>
<td>(TACROLIMUS ANHYDROUS - UNII:Y5L2157C4J)</td>
<td></td>
</tr>
</tbody>
</table>

Product Characteristics
Color: white (clear solution)
Shape:
Size:
Flavor:
Imprint Code:
Contains:

Packaging
<table>
<thead>
<tr>
<th>#</th>
<th>Item Code</th>
<th>Package Description</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>NDC:72934-4173-8</td>
<td>60 g in 1 BOTTLE, GLASS; Type 0: Not a Combination Product</td>
<td>05/10/2019</td>
<td></td>
</tr>
</tbody>
</table>

Marketing Information
Marketing Category: unapproved drug other
Application Number or Monograph Citation:
Marketing Start Date: 05/10/2019
Marketing End Date:

Labeler - Sincerus Florida, LLC (080105003)

Establishment
<table>
<thead>
<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
<th>Business Operations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sincerus Florida, LLC</td>
<td>080105003</td>
<td>manufacture(72934-4173)</td>
<td></td>
</tr>
</tbody>
</table>