**Summary for ARTG Entry:**

**ARTG entry for** Medical Device Included Class IIb  
**Sponsor** Medical Specialties Australasia Pty Ltd  
**Postal Address** PO Box 764, WILLOUGHBY, NSW, 2068 Australia  
**ARTG Start Date** 29/09/2003  
**Product category** Medical Device Class IIb  
**Status** Active  
**Approval area** Medical Devices

**Conditions**

- The inclusion of the kind of device in the ARTG is subject to compliance with all conditions placed or imposed on the ARTG entry. Refer Part 4-5, Division 2 (Conditions) of the Therapeutic Goods Act 1989 and Part 5, Division 5.2 (Conditions) of the Therapeutic Goods (Medical Devices) Regulations 2002 for relevant information.
- Breaching conditions of the inclusion related to the device of the kind may lead to suspension or cancellation of the ARTG entry; may be a criminal offence; and civil penalties may apply.

**Manufacturers**

**Name** pfm medical titanium GmbH  
**Address** Sudwestpark 42  
Nurnberg, 90449 Germany

**Products**

1. **Mesh, surgical**

<table>
<thead>
<tr>
<th>Product Type</th>
<th>Effective date</th>
<th>GMDN</th>
<th>Intended purpose</th>
<th>Exception(s) to approval</th>
<th>Exception product description</th>
<th>Reason</th>
<th>Notification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Single Device Product</td>
<td>29/09/2003</td>
<td>16048 Mesh, surgical</td>
<td>Soft tissue support</td>
<td>Active</td>
<td>12/08/2015</td>
<td>TILOOP Tape, TILOOP Two, TILOOP Total 4, TILOOP Total 6, TILOOP Fix, TILOOP Mesh, TILOOP Patch, TILOOP Clip</td>
<td>Product excluded by regulatory activity</td>
</tr>
</tbody>
</table>

**Specific Conditions**

No Specific Conditions included on Record

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