# Titan® Penile Implant Devices

## Physician Dictation Reference Sheet

### Measurement Information

<table>
<thead>
<tr>
<th></th>
<th>Right</th>
<th>Left</th>
</tr>
</thead>
<tbody>
<tr>
<td>Proximal</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Distal</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Date of Implant

- **Approach (circle one)**: Infrapubic, Penoscrotal
- **Model Number**
- **Serial Number**
- **Cylinder Size**
- **Rear Tip Extenders**: Right: _________ Left: _________
- **Reservoir Size (circle one)**: 75 mL, 125 mL
- **Reservoir Fill Volume**: ___________ cc/mL
- **Reservoir Location**

### Recommended Fill Volumes (cc)

<table>
<thead>
<tr>
<th>Cylinder Size</th>
<th>Fill Volumes (cc)</th>
<th>Tubing Length Scrotal (cm/in)</th>
<th>Tubing Length Infrapubic (cm/in)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Standard (45°)/Zero Degree (0°)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>14cm Standard (45°)</td>
<td>36 – 41</td>
<td>8.5/3.35</td>
<td>16/6.30</td>
</tr>
<tr>
<td>16cm Standard (45°)/Zero Degree (0°)</td>
<td>44 – 49</td>
<td>8.5/3.35</td>
<td>16/6.30</td>
</tr>
<tr>
<td>18cm Standard (45°)/Zero Degree (0°)</td>
<td>54 – 59</td>
<td>10.5/4.13</td>
<td>16/6.30</td>
</tr>
<tr>
<td>20cm Standard (45°)/Zero Degree (0°)</td>
<td>68 – 73</td>
<td>10.5/4.13</td>
<td>16/6.30</td>
</tr>
<tr>
<td>22cm Standard (45°)/Zero Degree (0°)</td>
<td>91 – 96</td>
<td>10.5/4.13</td>
<td>16/6.30</td>
</tr>
</tbody>
</table>

### Cylinders Only

- 24cm (45°)/Zero Degree (0°): 107 – 112
- 26cm (45°)/Zero Degree (0°): 114 – 119
- 28cm (45°)/Zero Degree (0°): 120 – 125

### Narrow base (N)

- **11cm (N) Zero Degree (0°)**: 18 – 23
- **14cm (N) Zero Degree (0°)**: 27 – 32
- **16cm (N) Standard (22.5°)**: 20 – 25
- **14cm (N) Standard (22.5°)**: 27 – 32
- **16cm (N) Standard (22.5°)**: 32 – 37
- **18cm (N) Standard (22.5°)**: 48 – 53

### Tubing Length

- Tubing cut to length
TITAN® BRIEF STATEMENT

Indications
The Titan® family of Inflatable Penile Prosthesis is indicated for male patients suffering from erectile dysfunction (impotence) who are considered to be candidates for implantation of a penile prosthesis.

Contraindications
The Titan® device is contraindicated in patients who have one or more of the following conditions: Patients with an active infection present anywhere in the body, especially urinary tract or genital infection. Patients with unresolved problems affecting urination, such as an elevated residual urine volume secondary to bladder outlet obstruction or neurogenic bladder. Patients unwilling to undergo any further surgery for device revision.

Warnings
Implantation of the device eliminates natural erections, as well as other related treatment options, impossible. Men with diabetes or spinal cord injuries, as well as immunocompromised patients, may have an increased risk of infection associated with a prosthesis. Failure to evaluate and promptly treat erosion may result in a substantial worsening of the condition, leading to infection and loss of tissue. Implantation of a penile prosthesis may result in penile shortening, curvature or scarring. Pre-existing abdominal or penile scarring or contracture may make surgical implantation more complicated or impractical. The risks and benefits of implanting this device in patients with lupus, scleroderma, myasthenia gravis, or documented sensitivity to silicone should be carefully considered.

Precautions
A thorough preoperative consultation should include a discussion between the patient and physician of all available treatment options and their risks and benefits.

Potential Complications
Scrotal swelling, auto-inflation, discomfort, angulation/curvature, edema, device malfunction/deflation, pain, difficulty with ejaculation, transient urinary retention, fever, migration, patient dissatisfaction, infection, hematoma wound leakage, bleeding, delayed wound healing, phimosis, sensory loss cylinder aneurysm, fibrous capsule formation, over/under inflation, erosion, scrotal erythema, genital change, inguinal hernia

See the device manual for detailed information regarding the implant procedure, contraindications, warnings, precautions, and potential complications/adverse events. For further information, call Coloplast Corp. at 1-800-258-3476 and/or consult the company website at www.coloplast.us.

Caution: Federal law (USA) restricts this device to sale by or on the order of a physician.