Introduction

The aim of total phallic construction is to create an aesthetically acceptable and sensitivity phallus with an incorporated neo-urethra. It should allow the patient to empty into the upright position as performed by most males. However, it should have a sufficient size to accommodate a prosthesis that can guarantee sufficient rigidity for penetrative sexual relations (1-9).

Original Article

Insertion of inflatable penile prosthesis into a neophallus (single cylinder)

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Background: The goal of inserting a penile prosthesis (PP) into a neophallus is to allow the patient to achieve satisfactory erectile function. Obtaining sufficient rigidity for sexual penetration has always been a challenge due to the absence of a suitable substitute for the corpora cavernosa of the penis. In this video, we demonstrate our surgical technique for the insertion of a single cylinder inflatable PP into a neophallus in a transgender patient who underwent a phalloplasty.

Methods: After the induction of general anaesthesia and administration of antibiotic prophylaxis, a 14-Ch urethral catheter is placed. The pubic symphysis is reached through a groin incision and three stay sutures placed in the periosteum to achieve proximal stability of the prosthesis. The reservoir is placed in the extraperitoneal space using a second incision. A Dacron proximal and distal covering, the ‘cap’ and ‘sock’, is fashioned around the proximal and distal aspect of the cylinder to anchor the device to the pubic bone and to prevent displacement. The device components are prepared and one cylinder disconnected. The pump is positioned into the (right) neo-scrotum and single cylinder into the neophallus. The insertion of a single testicular prosthesis into the (left) neo scrotum is performed. Post operatively the drain and catheter are removed the following day. The device is left semi inflated until review in one week. At two weeks, patients are taught and encouraged to cycle their device regularly. Patients can use the device for sexual intercourse at 6 weeks.

Results: In our experience the overall perioperative complication rate was 20%. Primary causes of revision included infection, instability or malfunction of the device. Despite high revision rates, 80% of patients have a fully functioning prosthesis.

Conclusions: PP implantation into a neophallus is a very complex procedure with significant complication rates and should be performed only by experienced surgeons working in dedicated and high-volume centers. This surgical technique has proved to be effective and reproducible with reasonable results in terms of function and safety.

Keywords: Neophallus; penile; implant; erectile dysfunction

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One of the major challenges is to allow patients to reach a rigidity necessary for sexual penetration. This goal is made particularly difficult by the lack of an effective substitute for the corpora cavernosa in the phallic construction (10, 11).

To date there is no agreement on what is the ideal method for implanting a penile prosthesis (PP) in a neophallus. In fact, even today the choice of the best method to achieve PP stability, prevent erosion and safely anchor the PP are a source of difficulty.

In this article we will demonstrate the technique we used to perform this procedure in detail.

Female to Male gender sexual reassignment is a complex multi-stage surgical challenge based on the creation of a functional and cosmetically acceptable phallus. The first stage is fashioning the neophallus. Different flaps can be adopted, but in this case we used a radial artery based forearm free flap (RAFFF).

Once the phallus is made, the next step is the laparoscopic hysterectomy salpingo-oophorectomy, join up urethroplasty, vaginectomy, burying of the clitoris, glans sculpting, and scrotoplasty. PP implantation is performed as the last surgical procedure.

Before proceeding with a PP implantation, 1 year is the minimum time required to wait from the last operation on the phallus.

In this video we demonstrate a single cylinder PP insertion technique in RAFFF neophallus (Video 1). The PP can be inflatable or malleable and during the preoperative examination the number of cylinders to be implanted is decided based on the size and shape of the phallus and the patient’s preferences. The authors antibiotic prophylaxis preference is a single dose of Gentamicin and Amoxicillin/Clavulanic Acid 1.2 g administered at induction and a scrub of the operative field of at least 10 min with povidone-iodine is also performed.

**Methods**

**Surgical technique**

The first step of the procedure is the insertion of a urethral catheter with the help of an introducer after which an inguinal incision is performed.

**Pubis anchorage sutures positioning (00.41)**

The incision was made and dissection of subcutaneous tissues to reach the pubic bone and all subcutaneous tissues retracted medially to expose the central aspect of the symphysis pubis.

To allow for anchoring of the prosthesis, were inserted three J-needle Polyester (Ethibond™) sutures into the pubic bone (Figure 1).

The number and position of the anchoring sutures changes according to the number of cylinders implanted. In the case of a single cylinder two parallel sutures are positioned on one side and a single one on the contralateral side (as shown in the Figure 1). Two parallel rows approximately 2 cm apart for each cylinder. This can be repeated on the contralateral side if necessary.

**Neophallus dilatation (01.46)**

The space for inserting the prosthesis into the neophallus was initially formed using McIndoe’s scissors. This was done by using Hegar dilators of increasing size starting from 10 to size 22, in order to obtain sufficient space to position the cylinder and the polyethylene terephthalate (Dacron™) tip. During the dilatation, the urethral catheter should be palpable and moved securely away from the dilatation line so that the chances of performing a urethral lesion or extrusion of the prosthesis at a later date are minimized (Figure 2).

**Formation of reservoir space (02.46)**

An oblique incision is made in the lower right abdominal quadrant to perform the ectopic insertion of the reservoir. After exposing of the external obliquus muscle aponeurosis the muscles fibres are separated and an Hegar dilator is used in order to create space. An antibiotic soaked gauze is left in the space until the reservoir is ready for placement (Figure 3).
Measurement for cylinder length (03.10)

The choice of cylinder length is measured starting from the pubis up to the tip of the neophallus (Figure 4).

Preparation of the 3 pieces inflatable PP (03.16)

The neophallus lacks the native corpora, and therefore there is a risk of erosion and consequent infection or malfunction of the prosthesis cylinder that is inserted into the neophallus. This risk is reduced by fashioning a silver-coated Dacron™ covering for the tip of the cylinder and the base of the prosthesis extending up to the exit of the connection tubes to provide additional stability.

Both cylinders can be inserted but in this video, only one cylinder is used. The other is cut and the tubing is secured with a true-lock connector (Figure 5). Once the prosthesis is prepared removing all the air from the device and the Dacron™ covering is secured to the cylinders, the implant is placed once again in the antibiotic solution.

Reservoir placement (07.37)

The reservoir was inserted into the previously prepared space and the tubes are passed deep to the lateral side of the fascia and brought out through the inguinal incision and secured using two shodded clips to prevent air from entering the system whilst protecting the tubing.

Testicular prosthesis insertion (07.52)

The testicular prosthesis is inserted into the scrotum through an opposite groin incision (Figure 6). Room for the testicular implant is made through the creation of a dartos pouch obtained by guided digital dilatation.
Insertion of the cylinder (08.13)

The Furlow introducer is used to deploy the needle and position the cylinder. The Dacron™ ‘sock’ (proximal end) is anchored to the pubis using the three Ethibond™ sutures placed at the start of the procedure. The device is then cycled.

Pump placement (10.07)

A right scrotal incision is made and the pump is placed in the right scrotal pouch following manual dilatation.

Final steps (10.40)

After flushing with saline solution, to ensure no air bubbles were introduced into the system, the tubes were connected and the socks removed before closing the incisions.

Results

Postoperative period (11.17)

On the first postoperative day the bladder catheter is removed and the patient was discharged with oral co-amoxiclav 625 mg three times/day for 5 days. One week later the PP was usually deflated in clinic. Three weeks later the patient was instructed on how to cycle the prosthesis, and encouraged to have penetrative sexual intercourse after 6 weeks.

Complications

The complications rate, type of revision surgery performed and long term results are summarised in Table 1 (18). When an infection or erosion occurs, all the components of the device were removed, including the Dacron sock and cap. A broad-spectrum antibiotic cover is given to the patient and a possible replacement of a prosthesis is offered at least 6 months later. In cases of mechanical failure, the defective component was identified and replaced intraoperatively. When the prosthesis showing problems and requiring revision has been implanted more than three years before, the replacement of the entire device is proposed to the patient. This is because the life expectancy of phalloplasty implants is much lower than those implanted in men with erectile dysfunction (11-18). The main causes of mechanical failure in our experience were cylinder breakage (69%), cylinder aneurysm (19%) and breakage of the connection pipe between the cylinder and the pump (12%).

Conclusions

In conclusion, with this video we wanted to show a standardized technique for inserting a PP into a neophallus. This surgical technique proved to be effective and reproducible with reasonable results in terms of function and safety. Despite this, is a very complex procedure with significant complication rates and should be performed only by experienced surgeons working in dedicated and high-volume centres.

Acknowledgments

None.

Footnote

Conflicts of Interest: All authors have completed the ICMJE

Table 1 Complication rate, type of revision surgery and long term results in 247 patients (18)

<table>
<thead>
<tr>
<th>Type of complication &amp; long term results</th>
<th>%</th>
<th>Management</th>
</tr>
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<tbody>
<tr>
<td>Infection of the device</td>
<td>8.5</td>
<td>Removal + delayed reimplantation</td>
</tr>
<tr>
<td>Mechanical failure</td>
<td>15.4</td>
<td>Replacement of the damaged component¹</td>
</tr>
<tr>
<td>Patient dissatisfaction</td>
<td>48</td>
<td>Repositioning of the cylinder(s) or insertion of a second cylinder</td>
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<tr>
<td>Long term results</td>
<td></td>
<td></td>
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<tr>
<td>60% original prosthesis still IN-SITU at 10 years</td>
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<tr>
<td>40% 10-year revision free survival</td>
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¹, revision surgery included exchange of the affected component only if the device was <3 years old, otherwise all components were exchanged.
uniform disclosure form (available at http://dx.doi.org/10.21037/jovs.2019.07.05). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. Written informed consent was obtained from the patient for publication of this manuscript and any accompanying images.

References


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