ABSTRACT: Benign Prostatic Hyperplasia (BPH) occurs in up to 80% of men older than 70 years. The prevalence of BPH increases with age, from approximately 25% of men at 50 years of age to 50 to 90% of individuals in their 8th decade of life. Treatment options for lower urinary tract symptoms (LUTS) secondary to BPH have been long restricted to drug therapy and, after failure of the former, surgical treatment. However, prostate surgery is associated with risks such as bleeding, necessity of transfusion and ejaculatory dysfunction. Several novel and minimally invasive methods for treatment of BPH have arisen in the past years. Common to all of these techniques are a more favorable safety profile and varying rates of success and clinical improvement, however frequently inferior to standard surgical techniques. Patients who will benefit the most from these procedures are those who would not wish to undergo surgery, those at prohibitively high risk for surgery, and those wishing to avoid ejaculatory dysfunction secondary to trans-urethral resection of the prostate (TURP) or open resection. Mechanical devices such as Temporary implantable nitinol device and Urolift appear to provide satisfying results with a favorable safety profile. Persistence of improvement is still questionable since follow-up longer than 5 years is not available for these methods. Of note, these methods may be of limited performance for larger prostates, in special, Urolift for those with a large median lobe or a prostate larger than 100 grams. Similarly, novel techniques for tissue ablation, including convective water vapor energy, appear to offer promising and safe results, yet with limited follow-up. Prostate artery embolization (PAE) is another safe procedure and an option for patients who are not suited for surgery, with satisfying short and medium-term follow-up but unknown results at long-term.

Keywords: Prostatic hyperplasia; Prostate; Minimally invasive surgical procedures.
INTRODUCTION

Benign Prostatic Hyperplasia (BPH), as a histological phenomenon, occurs in up to 80% of men older than 70 years. Prostatic growth may manifest as different lower urinary tract symptoms (LUTS), including storage and voiding symptoms. The prevalence of BPH increases with age, from approximately 25% of men at 50 years of age to 50 to 90% of individuals in their 8th decade of life. LUTS resulting from BPH may cause significant impairment in quality of life in older men, provoking at least mild modifications in lifestyle in up to 80% of patients with BPH.

Treatment options for LUTS secondary to BPH have been long restricted to drug therapy or surgical treatment. Use of alpha adrenoceptor antagonists can be successful in reducing LUTS in short to mid term, while inhibitors of the enzyme 5α-reductase may decrease long term risk of urinary retention. When medical treatment fails, surgery is generally offered as the gold standard for treatment of BPH, including endoscopic resection with either laser or cautery energy, laparoscopic or open surgery. However, in recent years, several minimally invasive techniques for treatment of BPH have arisen, presenting promising functional with lesser risks for complications and a shorter hospital stay.

To date, there is evidence for short-term results for most of such techniques, although long-term results are still expected. In general, results appear to be slightly inferior in terms of urinary obstruction, with favorable outcomes in terms of morbidity and complications. Among these, Prostate Artery Embolization (PAE), mechanical devices and alternate sources for tissue resection are the main groups of techniques proposed. Interestingly, these techniques also avoid postoperative ejaculatory dysfunction, a complication that up to 77% of patients would like to avoid if given the chance.

In this review, we will outline the most relevant minimally invasive options for treatment of BPH, the description of the techniques, the preliminary results, and limitations for each modality.

Mechanical devices

Several mechanical devices, being permanent or temporary, have been developed in recent years, aiming to keep the patency of the prostatic urethra without need for hospitalization or invasive procedures. These models have been perfected, with resolution of issues such as rejection to foreign body, post-procedure pain and device migration. Below, we will detail principles of each existing technique and current evidence for each method, respectively.

Temporary implantable nitinol device (TIND)

This is a temporary device called Temporary implantable nitinol device (TIND, Medi-tate, Israel). It features a nitinol structure which is installed in the prostate through a transurethral approach with a 22 French cystoscope. The device is composed of an anchorage leaflet and radial structures which exert centrifugal force over the prostatic tissue to expand the prostatic urethra diameter (Figure 1). The installation of the device only requires light sedation and local anesthesia. This device is 50 millimeters long and covers the whole extension of the prostate. Device withdrawal is usually programed within 5 to 7 days from installation and is achieved with a bladder catheter, without need for hospitalization. The mechanism of action consists in induction of ischemia in the prostatic tissue, in the areas which are under pressure by the nitinol structure, hence remodeling the prostate and allowing for a larger urine flow through the urethra.

Figure 1. Model representation of the position of the Temporary implantable nitinol device (TIND) in the prostate. Figure obtained from website of the manufacturer: http://www.medi-tate.com/patients/tind/
A prospective study has evaluated the use of TIND in 32 patients with 1 year follow-up, assessing functional results, complications, feasibility and safety of use of this device. Mean duration of the procedure was 5.8 minutes, with same day discharge being adopted after initial experience with the first 20 cases. Short-term complications occurred in 12.5% of patients (urinary tract infection, prostatic abscess, acute urinary retention, transient urinary incontinence). However, there were no cases of late or persistent complications after 1 years; also of note, re-interventions were not needed in any case to the end of follow-up. After 12 months, the authors noted a 67% raise of urinary flow rate and a 45% improvement in LUTS as evaluated with the International Prostate Symptom Score (IPSS). These authors concluded that the use of TIND is safe, easy, feasible and provides clinical efficacy in selected cases. There are currently 2 ongoing prospective studies further evaluating safety and performance of this device. TIND is not currently approved by the United States Food and Drug Administration, but it is approved for use in Brazil.

**Urolift**

Urolift is a permanent device (Urolift, Neotract, United States of America) that creates a continuous path through the prostatic urethra, from the bladder neck to the verumontanum, by means of implants applied bilaterally to the prostatic lobes through the urethra, pushing the lobes laterally and thus permitting increased flow through the urethra (Figure 2). Each implant is composed of an urethral end made of stainless steel, a non-absorbable polyethylene monofilament and a capsular end made of nitinol. These implants are installed using a cystoscope; usually 2 or 3 pieces are required for each prostatic lobe. These implants compress and push the prostatic lobes laterally, so as to free the prostatic urethra. The application of implants requires only local anesthesia, without necessity for hospitalization. Furthermore, the present of the implants does not preclude later surgical interventions eventually needed for benign or malignant disease. However, the efficacy of the procedure is limited if the prostate weight exceeds 100 grams; also of note, results are deficient if there is a large median prostatic lobe. Urolift has been approved by the United States Food and Drug Administration; to date, it is not approved for use in Brazil.

Urolift is the most studied minimally invasive mechanical device to date for the treatment of BPH. In a multicentric prospective randomized and controlled study including 206 patients, symptom improvement reached 50% as per IPSS evaluation; urinary flow rate increased by 64% 3 months after the procedure. Regarding the safety profile of the procedure, there were no complications or persistent side effects after 2 weeks from intervention; similarly, there was no impairment in sexual function as evaluated with the International Index of Erectile Function (IIEF). Patients from the same cohort, when re-evaluated after a 5 year follow-up, had maintained the short-term benefits previously demonstrated. Patients presented symptom improvement, as compared to baseline, of 44% as per IPSS evaluation; quality of life improvement of 42% and an increase in urinary flow rate of 41%. Other multicentric, prospective and randomized studies confirmed the effect of Urolift for the treatment of LUTS related to BPH, however less efficiently than Transurethral Resection of the Prostate (TURP). Notwithstanding, Urolift presented superior results in regard to post procedure recovery and sexual function, notably for not causing retrograde ejaculation. Such evidence have been corroborated by a recent meta-analysis including the currently existing data in the literature.

**Tissue ablation**

The process of prostatic tissue ablation by means of minimally invasive procedures has been tried in the past, for instance with transurethral ablation with radiofrequency and microwave treatment. However, long-term follow-up of such techniques did not reveal persistent benefit of these procedures, with a frequent necessity for surgery months after primary intervention.

In recent years, the development of thermal tissue ablation with water steam, through the use of convective radiofrequency, has allowed for new possibilities. A new system for the treatment of BPH with prostate ablation, called Rezum, consists in the use of radiofrequency to
heat up water stored in the system turning it into steam; steam is then injected into the prostatic tissue and rapidly condensed, thus liberating thermal energy. This energy, in turn, leads to cell denaturation, tissue necrosis and vascular occlusion. This procedure is performed transurethrally with a cystoscope, requiring only local transrectal and urethral, with optional sedation. Steam applications are performed with a retractile needle through the cystoscope, with 1 centimeter of distance between injections (Figure 3). Patients are discharged on the same day of the procedure and should keep an indwelling bladder catheter for 1 week.

Available studies to date have demonstrated that, after procedure, there is an increase in urinary flow rate of up to 6 milliliters per second, as well as a 11-point improvement of symptoms as evaluated with the IPSS\textsuperscript{18,19}. So far, efficacy of the results has been demonstrated at the end of a 2-year follow-up\textsuperscript{19}. Furthermore, prostate volume in patients undergoing Rezum ablation has decrease by 17 to 28%\textsuperscript{18,20}. Significantly, sexual function is well preserved after this procedure, with ejaculatory dysfunction occurring in only 3% of subjects; erectile function impairment was noted only among patients with a previous dysfunction\textsuperscript{18,21}. Among main complications, urinary tract infection occurred in 17% of patients and a transient urinary retention after catheter withdrawn happened in 14% of cases, while other complications such as urethral stenosis occurred in less than 4% of patients\textsuperscript{18}. Efficacy of the procedure was not related to prostate volume.

**Prostatic Artery Embolization (PAE)**

Prostatic Artery Embolization (PAE) was originally conceived as a treatment for refractory hematuria from prostatic etiology; however, it was later noticed that patients undergoing this therapy also improved in terms of LUTS. It subsequently started to be tested as a treatment for BPH among patients who were at high risk for a surgical procedure. PAE is performed by interventional radiologists through the femoral arteries with local anesthesia and allows for same day discharge. Major complications listed are transient ischemic proctitis, urethral burning, nausea and vomiting, with the short-term complications being identified as the “post-PAE syndrome”\textsuperscript{22}. Also of note, is the risk of renal injury from contrast toxicity, as an angiography is needed in this procedure.

Data available to date indicate that PAE presents high technical success rates and provides benefits in the treatment of LUTS secondary to BPH. Overall, symptom improvement and urinary flow rate increases are inferior to those of standard surgical treatment; however, for certain patients who are not suited for surgery due to a prohibitive risk, PAE can be a valuable option. In a prospective trial at University of Sao Paulo including 30 patients randomized to TURP or PAE, both procedures were considered efficient and safe; however, symptom improvement was higher for TURP but the complication profile was safer for PAE\textsuperscript{23}. A retrospective study including 630 patients undergoing PAE...
demonstrated clinical success at medium and long-term follow-up of 81% and 76%, with major complications in only 2 patients\(^2\). Yet, a recent meta-analysis of data available on PAE concluded that this technique should still be considered experimental, despite of safety evidence available\(^2^5\). Patients who are candidates for this procedure and not for surgery should be carefully selected and informed.

**CONCLUSIONS**

Several novel and minimally invasive methods for treatment of BPH have arisen in the past years. Common to all of these techniques are a more favorable safety profile and varying rates of success and clinical improvement, however frequently inferior to standard surgical techniques. Patients who will benefit the most from these procedures are those who would not wish to undergo surgery; those at prohibitively high risk for surgery and those wishing to avoid ejaculatory dysfunction secondary to TURP or open resection. Mechanical devices such as TIND and Urolift appear to provide satisfying results with a favorable safety profile. Persistence of improvement is still questionable since follow-up longer than 5 years is not available for these methods. Of note, these methods may be of limited performance for larger prostate, in special, Urolift for those with a large median lobe or a prostate larger than 100 grams. Similarly, novel techniques for tissue ablation, including convective water vapor energy, appear to offer promising and safe results, yet with limited follow-up. Prostate artery embolization is another safe procedure and not for surgery should be carefully selected and an option for patients who are not suited for surgery, with satisfying short and medium-term follow-up but unknown results at long-term.

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