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LASER TREATMENT OF THE PROSTATE

Abstract

A method of treating prostatic tissue includes trans-perineally introducing at least one energy delivery device in the prostate of a patient. Once the energy delivery device has been introduced, energy is delivered to a volume of tissue of the prostate until the volume is vaporized or sublimated and a cavity is formed in the prostate tissue. The energy delivery device can then be removed from the prostate.

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Background/Summary

CROSS REFERENCE TO RELATED APPLICATIONS [0001] This application is continuation under 37 U.S.C. 1.53 (b) of pending prior U.S. patent application Ser. No. 17/160,016 filed Jan. 27, 2021, which is a continuation-in-part under 37 U.S.C. 1.53 (b) of pending prior U.S. patent application Ser. No. 15/947,292 filed Apr. 6, 2018, the entire contents of each application are incorporated herein by reference.

TECHNICAL FIELD

[0002] Disclosed herein are ablative treatments of human body. Embodiments disclosed herein specifically concern the treatment of Benign Prostatic Hyperplasia or Benign Prostatic Hypertrophy (shortly here on referred to also as BPH) by ablation therapy. Some embodiments specifically refer to laser ablation therapy. Embodiments disclosed herein also relate to treatment of benign or malignant tumoral tissues from an organ, in particular from the prostatic gland.

BACKGROUND

[0003] Benign Prostatic Hyperplasia (BPH), also known as Benign Prostatic Hypertrophy, is a benign pathology of the prostate characterized by proliferation of prostate cellular elements.

[0004] Cellular accumulation and gland enlargement may result from both epithelial and stromal proliferation and/or reduced cell death (apoptosis). The enlarged gland has been proposed to contribute to the overall lower urinary tract symptoms (LUTS).

[0005] BPH/LUTS prevalence rates ranges from 50% to 75% among men 50 years of age and older to 80% among men 70 years of age and older. The overall incidence rates ranges from 8.5 to 41 cases/1000 person per years.

[0006] The presence of moderate-to-severe LUTS was also associated with the development of acute urinary retention (AUR) as a symptom of BPH progression, increasing from a prevalence of 6.8 episodes per 1000 patient years of follow-up in the overall population to a high of 34.7 episodes in men aged 70 and older with moderate to severe LUTS.

[0007] Enlargement of the prostate involves a series of compression symptoms that also affect the dynamics of urination and greatly alter the quality of life of the affected patient. Although LUTS secondary to BPH (LUTS/BPH) is not often a life-threatening condition, the impact of LUTS/BPH on quality of life (QoL) can be significant and should not be underestimated. Some symptoms include: hesitation, weak and/or intermittent urinary stream, urinary retention, burning sensation and/or pain during urination, dysuria, urgency incontinence (urgent need to urinate), urination frequency, urine leakage, incontinence, dribbling at the end of urination, nocturia, incomplete emptying of the bladder, straining.

[0008] TURP (transurethral resection of prostate) is today the most common technique in the treatment of BPH. This technique represents an evolution of open surgery achieving improvement in LUTS symptoms with lower hospitalization time. The technique, which nowadays represents the gold standard, consists in the removal of the enlarged tissue by means of a surgical instrument called a resectoscope inserted into the penis through the urethra, such that no external incision is required, as instead in the case of open surgery. In this way, the hypertrophic tissues of the prostatic gland, which creates compression on the urethra, is sliced in small pieces and removed by the resectoscope. TURP is performed in general anesthesia or spinal anesthesia and requires from one to two hospital recovery after the procedure. A catheter is needed due to swelling that blocks urine flow in post treatment (generally left in place for at least 24 to 48 hours).

[0009] TURP is still considered an aggressive surgery involving several acute and chronic complications that can greatly affect the life of the patient in post-surgery. Bleeding especially in patients receiving an anticoagulant therapy (heparin, coumadin related compounds, antiplatelet agents) remains a concern and may require, although rarely, blood transfusion.

[0010] Additional several post-treatment urinary symptoms exist, due to the fact that the urethra epithelium has been completely removed. For instance urination may be painful; a sense of urgency or frequent urination 6 to 8 weeks after TURP has also been reported by patients. One of the possible permanent side effects of TURP is retrograde ejaculation which results when the tiny sphincter muscle that usually blocks off the bladder during ejaculation is damaged during the procedure. This side effect can force patient to sterility. The main complications of TURP and their incidence are: [0011] temporary difficulty urinating (urinary retention): 3% [0012] urinary tract infection: 1.7% [0013] erectile dysfunction: 2.1-11% [0014] heavy bleeding: transfusion rate 0.4% [0015] difficulty holding urine (early urge incontinence): 30-40% [0016] need for retreatment due to urethral strictures (2.2-9.8%). Math. [0017] bladder neck contractures (0.3-9.2%)

[0018] In order to improve the patient's quality of life, a number of mini-invasive techniques have been developed to achieve symptomatic remission with reduced hospitalization time and less complication rates. Based on the different approaches on which they are based, these techniques can be grouped as follows: [0019] A. Techniques aimed at immediately removing an excess hypertrophic tissue (as TURP but with less complications and side effects). In this sense, PVP (Photo Selective Vaporization, HoLAP Holmium Laser Ablation, HoL RP holmium laser resection, HoLEP-TuLEP Holmium Laser Enucleation, Thulium Laser Enucleation) have been developed. All of these techniques are performed by a trans-urethral approach using a cystoscope and include the use of power laser that is brought into the working area by contact or non-contact optical fibers to vaporize or resect hypertrophic tissue. Among techniques of this first group, the following can be mentioned: [0020] a. Visual Laser Ablation of the Prostate (VLAP): This procedure is performed by trans-urethral approach and consists of lasing prostatic tissue in a noncontact fashion to create an area of heat-induced coagulative necrosis that extends about 10 mm into the tissue. Edema and prolonged sloughing of the coagulated tissue leads to irritation at the level of the lower urinary tract and to urinary retention requiring long periods of catheterization, up to 3 months, in up to 30% of cases; [0021] b. Photo Selective Vaporization (PVP): According to this technique, the urethral and periurethral tissue that causes obstruction is vaporized with the purpose of reopening the urinary duct immediately. A KTP laser (wavelength of 532 nm, visible green light) with a side firing fiber in a non-contact mode is used for this application. This wavelength is strongly absorbed by hemoglobin and therefore has a very short absorption depth in well-vascularized tissue such as the prostate; [0022] c. Holmium Laser Ablation (HoLA P): An Holmium laser is used to produce vaporization of the tissue; [0023] d. Holmium Laser Resection of Prostate (HoLRP): This technique expects to use high-density laser energy to vaporize/incising exceeding tissue which is cut into small pieces that fall into the bladder and are then subsequently expelled; [0024] e. Holmium-Thulium Laser Enucleation (HoLEP-TuLEP): Resection of the entire hypertrophic lobe is achieved through a contact optical fiber that vaporizes and separates tissues. The large removed lobes fall into the bladder and are removed by morcellation, which takes considerable time especially for big prostates; [0025] f. Trans-Urethral Incision of the Prostate (TUIP): a contact optical fiber and a high energy laser are used to produce an incision in the periurethral tissue that behaves as an alternative urine channel, [0026] B. Techniques aimed at preserving the prostatic urethra and its urothelium. These techniques usually exploit a minimally invasive approach and consist of placing an energy applicator directly into the prostatic adenoma by a cystoscope. Radiofrequency, microwave, or laser energy are released into the adenoma and, due to tissue absorption mechanisms, are transformed into heat with increasing local temperature. Temperature and exposure time induce irreversible cell damage and coagulation of small vessels with subsequent cell death in the surrounding tissue. Necrotic tissue is subsequently reabsorbed by the

physiological wound healing mechanism of the human body and a prostatic volume reduction takes place, with consequent remission of BPH symptoms. There is also the possibility of obtaining necrosis of the tissues by the use of cryoablation, applied to the hypertrophic area resulting from several freezing-unfreezing cycles. Thermal energy techniques are reported below: [0027] a) The radiofrequency technique has been named Transurethral Needle Ablation (TUNA). It involves positioning electric needles through the urethra in the hypertrophic part and generating alternating electric current between the needle tips and a recovery plate placed on the patient's skin (usually behind the legs). The current heats the prostatic tissue by Joule effect and determines cell death due to increased temperature. Urethra burns and consequent damages may occur due to incorrect needle positioning and to poor flow control of the current circulating in low impedance paths. These effects are unpredictable and thus unavoidable; [0028] b) The procedure using microwaves on the prostate is called Transurethral Microwave Therapy (TUM T) and consists of a catheter with an antenna that is placed in the urethra through a blistered bladder balloon. The portion of the urethral tissue facing the antenna undergoes the highest temperatures of the generated thermal field and this usually causes irreversible necrosis of the urethral and surrounding tissues; [0029] c) The technique using laser radiation to obtain tissue coagulation goes under the name of Interstitial Laser Coagulation (ILC). The procedure is performed by placing laser-diffusing fibers directly into the prostatic adenoma, either via the transurethral cystoscopic approach, or the perineal approach. The ILC laser technique uses particular optical diffusers to avoid overheating the tissues close to the tip of the flat type fiber optic. These fiber tip diffusers allow treating extended tissue portions by maintaining optical density at levels such as to induce tissue coagulation, with temperatures below 100° C. Some systems have a temperature feedback control to maintain the temperature within the desired range of 80-100° C. (Indigo® Optima Laser Treatment System; Ethicon Endo-Surgery, Cincinnati, OH). Optical diffusers generally have a higher caliper than bare optical fibers, because they are equipped with a protective optically transparent dome. The procedure induces substantial tissue edema and hence necessitates prolonged (7-21 days) postoperative catheterization. Retreatment rates are problematic: as high as 20% at 2 years, 41% at 3 years and 50% at 54 months.

[0030] Summarizing, trans-urethral laser techniques involving vaporization or vapor-enucleation of the exceeding tissue cause destruction of the urethral portion and it takes several weeks for new epithelization and wound healing. This type of approach implies much of the side effects and complications of treatment such as prolonged bleeding, difficulty urinating, burning, infections.

[0031] Techniques that produce tissue coagulation by a transurethral (with a cystoscope) or trans-perineal (under the ultrasound transcriptional guide) approach are aimed at preserving the functional anatomical structures of the gland and induce a coagulative necrosis in the central part of the adenoma. Anyway, the transurethral approach maintains a relative high risk of urethra damage and transperineal approach is preferred as safer and prone to less complications. However, the main drawback remains the slow improvement of symptoms which takes several weeks or months to take place.

[0032] A need therefore exists, for more efficient, less invasive treatments of BPH, which also may result in faster recovery and less post-operative discomfort for the patient.

SUMMARY

[0033] Disclosed herein is a method for treating benign prostatic hypertrophy, benign tumor tissue or malignant tumor tissue by tissue removal, which exploits the advantages of the trans-perineal approach and energy-tissue interaction in order to remove enlarged tissues of the adenoma leaving intact all the critical prostatic and peri-prostatic structures (gland capsule, urethra, neurovascular bundles). Enlarged tissue removal by local energy application leads to the formation of cavities with an immediate reduction of gland compression on adjacent structures and urethra followed by the remission of LUTS symptoms.

[0034] According to one aspect, the subject matter disclosed herein concerns a method of treating

benign prostatic hyperplasia in a patient in need of said treatment, comprising the following steps: [0035] trans-perineally introducing at least one energy delivery device through a perineal area of the patient in a first position in a prostate of the patient; [0036] delivering energy from an energy source through the energy delivery device to a first volume of tissue of said prostate, until said first volume is vaporized or sublimated and a cavity is formed in the prostate tissue; [0037] removing the energy delivery device from the prostate; and [0038] reducing the volume of the cavity such that compression of the urethra by prostate tissue surrounding the urethra is relieved.

[0039] While several embodiments disclosed here on use laser energy, the option of using other sources of energy is not excluded. Laser beam can be easily conveyed through optical fibers, which may be beneficial since it results in less invasive treatments.

[0040] Optical fibers with very small diameter can be introduced in the adenomatous tissue with very fine needles or introducers in order to deliver laser energy. The number of treatment sites (i.e. cavity formation by removal of hypertrophic tissue) can be arbitrary chosen. One, two or more needles and relevant fibers can be introduced in several positions of the prostate, to perform simultaneous treatment in different volumes of the prostate. The needles can be arranged symmetrically with respect to a sagittal plane.

[0041] In some embodiments, one optical fiber is introduced through each needle or introducer. Each optical fiber can be coupled to a respective energy source, in particular a respective laser source. Thus, in some embodiments, a number of laser sources equal to the number of needles and optical fibers simultaneously introduced in the prostate for multiple simultaneous treatments can be used. This renders the treatment faster and reduces patient's discomfort.

[0042] The energy delivered in the prostatic tissue is modulated to provoke vaporization of water contained in the tissues. The energy can also interact with carbon or other substances contained in the tissues and provoke sublimation thereof. Measures can be taken to remove gaseous or vapor material resulting from the energy/tissue interaction. The vaporization and possible sublimation generate cavities in the tissue, which can result in immediate reduction of the bulk of the gland, and therefore immediate relieve of the compression provoked by swollen adenomatous tissue on the urethra. The more the tissue removal the more effective the treatment is. Single cavities generated by several energy applicators, such as optical fibers, can also merge in a larger cavity depending on the distance between the tips of the optical fibers used during treatment and treatment parameters.

[0043] The trans-perineal approach is less traumatic than trans-urethral or trans-rectum approach and allows a lower risk of infections and is performed under local anesthesia, or even without anesthesia, if thin needles are used. As a consequence, the described treatment can be carried out in outpatient setup without general or spinal anesthesia with very short recovery times for the patient and less or no hospitalization costs. Moreover, due to the respect of critical prostate structure also catheterization can be reduced or avoided.

[0044] Presently preferred embodiments involve the use of optical fibers which are introduced through thin needles that are inserted into the gland under image monitoring (ultrasound or magnetic resonance imaging). Optical fibers may have a core diameter preferably ranging from 100 to 500 micrometers. Thin optical fibers can be introduced through very thin needles, e.g. 25G-20G caliber needles, which can be used without local anesthesia.

[0045] According to a further aspect, disclosed herein is a method of removing tissue from an organ of a patient, comprising the following steps: [0046] introducing at least one energy delivery device in a first position in an organ of the patient; [0047] delivering energy from an energy source through the energy delivery device to a first volume of tissue of said organ, until said first volume is vaporized or sublimated and a cavity is formed in the organ tissue; [0048] removing the energy delivery device from the organ.

[0049] The various features of novelty which characterize the invention are pointed out with particularity in the claims annexed to and forming a part of this disclosure. For a better understanding of the invention, its operating advantages and specific objects attained by its uses,

reference is made to the accompanying drawings and descriptive matter in which preferred embodiments of the invention are illustrated.

Description

BRIEF DESCRIPTION OF THE DRAWINGS

[0050] In the drawings:

[0051] FIG. 1 is a schematic cross-sectional view of a prostate affected by BPH;

[0052] FIG. 2 is another schematic cross-sectional view, according to a sagittal plane, of a prostate affected by BPH;

[0053] FIG. 3A is a sagittal cross-sectional view according to line III-III of FIG. 2 during a subsequent treatment step;

[0054] FIG. 3B is a sagittal cross-sectional view according to line III-III of FIG. 2 during a subsequent treatment step;

[0055] FIG. 3C is a sagittal cross-sectional view according to line III-III of FIG. 2 during a subsequent treatment step;

[0056] FIG. 4 is a schematic of a cavity growing during tissue vaporization;

[0057] FIG. 5 is a cross-sectional view similar to FIGS. 1 and 2, in a further embodiment, during or prior to the treatment;

[0058] FIG. 6 is the same view of FIG. 5 after treatment;

[0059] FIG. 7 is a diagram showing the cavity volume increase during time;

[0060] FIG. 8 is a view similar to FIGS. 3A-3B-3C of a further embodiment;

[0061] FIG. 9 is a sagittal cross-sectional view of a prostate during treatment with trans-rectal ultrasound (US) imaging guidance;

[0062] FIG. 10 is a schematic cross-sectional view in a sagittal plane of a needle and optical fiber arrangement with improved gas removing facilities, aimed at removing gaseous products resulting from the tissue/laser interaction and resulting tissue vaporization and sublimation;

[0063] FIG. 10A is an enlargement of a detail of FIG. 10;

[0064] FIG. 11 is a schematic cross-sectional view similar to FIG. 10 of a further embodiment with improved gas removing facilities;

[0065] FIG. 12 is a schematic cross-sectional view similar to FIG. 10 of a further embodiment with improved gas removing facilities;

[0066] FIG. 13 is a cross-sectional view according to line XIII-XIII of FIG. 12 in an exemplar embodiment;

[0067] FIG. 14 is a cross-sectional view according to line XIII-XIII of FIG. 12 in an exemplar embodiment;

[0068] FIG. 15A is a cross-sectional view according to a sagittal plane in a further embodiment;

[0069] FIG. 15B is a cross-sectional view according to a sagittal plane in a further embodiment;

[0070] FIG. 15C is a cross-sectional view according to a sagittal plane in a further embodiment;

[0071] FIG. 16 is a flow-chart summarizing the main steps of BPH treatment methods according to the present disclosure;

[0072] FIG. 17 is a flow-chart summarizing the main steps of BPH treatment methods according to the present disclosure;

[0073] FIG. 18 is a flow-chart summarizing the main steps of BPH treatment methods according to the present disclosure;

[0074] FIG. 19 is a cross-sectional view according to a sagittal plane of a prostate under treatment with the use of temperature sensors;

[0075] FIG. 20 is a cross-sectional view according to line XX-XX of FIG. 19;

[0076] FIG. 21A is a view illustrating steps of a further method according to the present disclosure;

[0077] FIG. **21B** is a view illustrating steps of a further method according to the present disclosure; [0078] FIG. **21C** is a view illustrating steps of a further method according to the present disclosure; [0079] FIG. **21D** is a view illustrating steps of a further method according to the present disclosure; [0080] FIG. **22** is a view illustrating a modified step of the method of FIGS. **21A-21D**; and [0081] FIG. **23** is a view illustrating a further modified step of the method of FIGS. **21A-21D**.

DESCRIPTION OF PREFERRED EMBODIMENTS

[0082] The following detailed description of the exemplary embodiments refers to the accompanying drawings. The same reference numbers in different drawings identify the same or similar elements. Additionally, the drawings are not necessarily drawn to scale. Also, the following detailed description does not limit the invention. Instead, the scope of the invention is defined by the appended claims.

[0083] Reference throughout the specification to “one embodiment” or “an embodiment” or “some embodiments” means that the particular feature, structure or characteristic described in connection with an embodiment is included in at least one embodiment of the subject matter disclosed. Thus, the appearance of the phrase “in one embodiment” or “in an embodiment” or “in some embodiments” in various places throughout the specification is not necessarily referring to the same embodiment(s). Further, the particular features, structures or characteristics may be combined in any suitable manner in one or more embodiments.

[0084] In embodiments disclosed herein, BPH is treated with a mini-invasive procedure using energy applicators introduced into the prostate through the transperineal route. Embodiments disclosed herein use laser energy conveyed in situ by means of optical fibers. The laser radiation parameters are selected such that water contained in the tissue is vaporized and other substances contained in the prostatic tissue can sublime. The resulting gaseous by-products of the tissue/laser interaction can be removed, possibly with the aid of ad-hoc removing devices. Contrary to treatments of the current art, which are mainly based on tissue denaturation and subsequent tissue debulking, an immediate reduction in volume of the gland is achieved, which results in immediate relieve of the BPH symptoms, mainly linked to compression of the urethral lumen.

[0085] Turning now to the figures, FIGS. **1** and **2** illustrate two schematic sectional views of exemplary prostatic glands **1** affected by BPH along a transverse plane, i.e. orthogonal to the sagittal plane. The two glands **1** differ from one another as far as their total dimension is concerned. The prostate of FIG. **1** is larger than the one of FIG. **2**. Both prostatic glands **1** contain adenomatous tissue **3**, which has increased the overall dimension of the gland and caused swelling thereof. The adenoma causes compression of the urethral lumen (urethra) **5** which extends through the two lateral lobes **1A**, **1B** of the prostate. The prostate **1** is surrounded by a capsule **7**. Neuro-vascular bundles **9** are arranged on the exterior of the capsule **7** on two sides of the prostatic gland **1**.

[0086] Any surgical treatment of BPH shall prevent damages to the capsule **7** and the neurovascular bundles **9**, and possibly preserve the urethra for fast post-surgical recovery.

[0087] In FIGS. **1** and **2** reference number **10** designates cavities which are formed in the prostate **1** by vaporizing and sublimating adenomatous prostatic tissue using laser energy. The laser treatment can be performed by introducing one or more hollow needles or introducers in both lobes **1A**, **1B** of the prostate and by introducing an optical fiber through each needle, such that laser energy can be delivered to the interior of the adenomatous tissue causing vaporization thereof.

[0088] The needles and the optical fibers are introduced through the perineum. The number of needles introduced in each lobe of the prostate **1** can depend upon the dimension of the prostate and upon the amount of adenomatous tissue to be removed. Two or more needles or introducers can be introduced at the same time in each lobe **1A**, **1B** of the prostate **1**, such that adjacent or neighboring prostatic tissue volumes can be treated simultaneously. In other embodiments, one or more needles or introducers can be introduced in sequence in the prostatic tissue, for treating neighboring or adjacent volumes of the adenoma in timely shifted manner. This second approach will require a

longer treatment time.

[0089] The number of simultaneously introduced needles can depend, inter alia, upon the number of laser sources available. It can be beneficial to provide as many independent laser sources as there are simultaneously operating optical fibers.

[0090] One or more needles or introducers and relevant optical fibers can be moved during treatment along the needle axis, such that subsequent tissue volumes can be illuminated with laser energy in a so-called pull-back procedure. With continuing reference to FIGS. 1 and 2, FIG. 3A, 3B, 3C illustrate a sectional view according to line III-III of FIG. 2, i.e. along a sagittal plane, in three different laser treatment steps of the prostate 1. In the exemplary embodiment of FIGS. 3A, 3B, 3C two introducers or needles 11, each guiding a respective optical fiber 13, are introduced in each lobe 1A, 1B of the prostate 1 through the transperineal route. The needles 11 are introduced through the perineum 15, i.e. through the area extending between the scrotum (not shown) and the anus 17. According to the embodiment shown in FIGS. 3A, 3B, 3C, each needle or introducer 11 and relevant optical fiber 13 are fully introduced in the prostate up to the starting position of FIG. 3A, where the treatment will initiate. This is the position where the needle tips 11T are farther away from the prostate apex 1C and nearest to the prostate base 1D and to the bladder floor. Laser energy generated by a laser source is conveyed through the optical fibers 13 to the tips thereof, which are located at or near the tips 11T of the needles 11.

[0091] As mentioned, according to some embodiments, independent laser sources can be provided for different optical fibers. In FIG. 3A one laser source 19A, 19B is shown for each optical fiber 13. Each laser source can be controlled independently from the others, such that for example each laser source can be turned on or off and the laser emission thereof can be adjusted independently from the others. For example the emission power, the emission time, the energy dose, the pulse frequency (in case of pulsed laser) can be adjusted for each source independently. In other embodiments each source can be coupled to multiple optical fibers.

[0092] Insertion of the needles or introducers 11 and optical fibers 13, as well as their subsequent movement in the prostate can be performed with the aid of ultrasound imaging (US) using an ultrasound probe, for instance a rectal probe, as described in greater detail later on. In other embodiments, the insertion of the needles can be performed under magnetic resonance imaging in combination with non-magnetic introducers or needles 11, or using any other suitable imaging method.

[0093] Laser emission can be controlled by a control unit 21, which can be functionally connected to the laser source 19 and to a user interface 23. As will be described in more detail here on, a controlled amount of laser energy is delivered by the laser source through the optical fibers 13 to cause vaporization and/or sublimation of tissue in a volume surrounding the tip of the optical fiber and/or in front of said tip. In FIG. 3A, V1 indicates the volume of adenomatous tissue which can be vaporized and sublimated by the laser radiation while the optical fiber tip is maintained in the position of FIG. 3A.

[0094] In order to remove a larger amount of tissue, the optical fiber 13 and the relevant hollow needle or introducer 11 can be gradually moved out of the patient's body. For instance, once the tissue volume V1 has been vaporized and/or sublimated by laser energy delivered through the optical fibers 13 in the first position of FIG. 3A, the needles 11 and relevant optical fibers 13 therein can be pulled back stepwise in an outward direction f11, in the position of FIG. 3B. The hollow needles 11 and optical fibers 13 can then be maintained in the new position of FIG. 3B for a given amount of time, during which laser radiation generated by the laser source 19 irradiates the tissue in volume V2 and causes vaporization or sublimation thereof. Once the tissue in volume V2 has been vaporized and/or sublimated, the needles 11 and optical fibers 13 are moved a further step outwardly, until the position of FIG. 3C is achieved, where a third volume V3 of adenomatous tissue is vaporized or sublimated.

[0095] As can be appreciated from FIG. 3C, in a three-step process, two elongate volumes of

adenomatous tissue have been removed by vaporization/sublimation along the trajectory of pull-back motion of the two needles **11** in lobe **1A**. The same operation can be performed simultaneously in lobe **1B**, such that at the end of the process, four volumes of tissue have been removed by vaporization and/or sublimation around the urethra **5**. These four volumes are shown in a cross-sectional view in FIG. **2** and labeled **V**. The cavities thus formed will contract and thus immediately relieve pressure on the urethra **5**. In contrast to current art so-called laser ablation techniques, which are based on tissue denaturation through laser radiation and subsequent removal of the denaturated/desiccated tissue through subsequent resorption, the treatment method of the present disclosure involves an immediate tissue volume reduction, with consequent immediate, real-time relieve of compressive forces exerted by the adenomatous tissues on the urethral lumen. The real-time effect of laser ablation by vaporization and/or sublimation of tissues through the transperineal route results in very fast recovery and short or no hospitalization.

[0096] As mentioned above, while in some embodiments all the needles **11** (four in the exemplary embodiment of FIGS. **2** and **3**) are introduced at substantially the same time, such that four cavities **10** corresponding to the removed volumes **V1**, **V2**, **V3** will form at substantially the same time, in other embodiments, the needles **11** and optical fibers **13** can be introduced in sequence, thus forming the cavities in two or more steps. For instance, the needles in one lobe **1A**, **1B** can be introduced first, and only upon completion of the treatment of the first lobe, needles will be introduced in the other lobe **1B**, **1A**. In other embodiments, a first step may involve insertion of one needle per lobe and a second step, to be performed upon completion of the first step, may involve insertion of the other needles in the two lobes.

[0097] Depending upon the dimension of the prostate **1** to be treated, a different number of hollow needles or introducers **11** and of optical fibers **13** can be used. In FIG. **1** two cavities **10** are formed by using just two needles **11**, one per lobe **1A**, **1B**.

[0098] The dynamics of laser ablation by vaporization and/or sublimation with naked optical fibers involves the formation of a cavity of sublimated/vaporized tissue, surrounded by a small layer of vacuolated and dehydrated tissue. For a frontal-emission optical fiber **13** the cavity of vaporized tissue grows with respect to energy delivery time, i.e. as a direct function of the energy dose.

[0099] FIG. **4** pictorially illustrates cavity formation from the beginning to final saturated dimension. If the laser radiation intensity is above the tissue vaporization and sublimation threshold, a cavity **C1** starts to form and grows in front of the fiber tip **13T**. The laser energy is absorbed by the water contained in the tissue and causes vapor formation at the tip of the optical fiber **13**. In FIG. **4** the shape of the cavity is the one obtained with an optical fiber having a flat fiber tip, which produces a cavity of elliptic shape. As laser emission continues, the laser emission will further propagate through the empty volume formed by vaporization of the tissue such that the front of the tissue moves forward. Tissue is, dehydrated and vaporized and finally sublimated in a cyclic way leading to a well-defined cavity in front of the fiber tip. As time proceeds, the cavity enlarges from **C1** to **C6**.

[0100] As the volume of the cavity increases, the front velocity, i.e. the speed at which the surface of the cavity advances in front of the stationary fiber tip **13T**, decreases as pictorially represented by cavities **C3**, **C4**, **C5**, **C6**. The reason for this is that the energy per surface which impinges on the advancing cavity surfaces reduces with the square of the distance from the fiber tip. The speed of ablation by vaporization slows down until an asymptotic limit is achieved. When the cavity achieves the dimension represented by **C6**, whose size (length and the maximum diameter) depends on the chosen dose (duration of the treatment multiplied by the mean power value), the power density at the cavity margin is not able to sustain the process of vaporization and a stable condition is achieved in terms of removed volume, i.e. no further tissue can be vaporized and removed, such that the dimension of the cavity remains constant and the fiber is to be switched off.

[0101] When the dimension **C6** is achieved, the needle or introducer **11** and the optical fiber **13** therein can be withdrawn stepwise with a so-called pull-back maneuver, such that an adjacent

tissue volume can be treated in the next treatment step.

[0102] Since the parameters of the laser emission are chosen such that the irradiated tissue is removed by vaporization and sublimation, the relief of the compression exerted by the hypertrophic prostatic tissue on the urethra **5** is rapidly obtained as can be understood from FIGS. **5** and **6**. An additional brief rectal internal massage of the prostate operated by a finger can improve the effect by promoting collapsing of the tissue around the cavity thus formed.

[0103] In FIG. **5** a schematic sectional view of the prostate **1** during or immediately after laser application is shown. One or more cavities **10** are generated by tissue vaporization and sublimation. In the exemplary embodiment of FIG. **5** five cavities **10** have been formed in each lobe **1A**, **1B** of the prostate **1**. The tissue around the cavities **10** will collapse spontaneously or helped by a short rectal massage, such that the cavities **10** will shrink, as shown in FIG. **6**. Shrinkage of the cavities causes a reduction of the external volume of the prostatic gland **1** and a reduction of the urethra compression. The urethra **5** re-opens as a direct consequence of the resulting pressure relieve thereon.

[0104] From an experimental work ex vivo with an optical camera it was possible to measure the length (i.e. longitudinal dimension) of the cavity during its formation and the cavity dimension versus time could be plotted. FIG. **7** illustrates the cavity length vs. time. The dotted line represents the position (distance from the fiber tip) of the active front of the cavity. The continuous line is an exponential fitting curve of the raw data. Lasing starts at time 0. The data of FIG. **7** were obtained with a continuous laser beam with a power of 5 W, 1064 nm laser wavelength, conveyed through a quartz optical fiber with flat tip and a core diameter of 300 micrometer. The experimental data were obtained using a fresh porcine liver sample at room temperature.

[0105] At the beginning of the laser emission, for about 40 seconds, there is no movement in the forward direction, i.e. no cavity is formed in front of the fiber. This time is necessary to heat the tissue above 100° C. and activate intra and inter cellular water boiling and subsequent vapor formation in front of the fiber. If treatment stops before the vapor formation threshold is achieved, a spherical coagulated thermal lesion is created in the tissue, centered on the fiber tip. Once the vaporization threshold is achieved and exceeded, a cyclic action of dehydration, vaporization and sublimation occurs on new tissue layers facing the laser radiation emitted by the fiber tip. The active front of the cavity being formed moves forward in an exponential way (as described by the continuous fitting line in the graph of FIG. **7**, up to saturation of the cavity length. This means that there is no further cavity enlargement even if energy delivery continues. This saturation phenomenon is dependent upon the slight divergence of the optical fiber emission. This means that the power density i.e. the radiation intensity decreases as the distance of the cavity surface from the fiber tip increases. In the asymptotic condition the power density decreases to a value that is under the vaporization threshold and hence the laser radiation is unable to induce further tissue removal. Once this condition is achieved, further energy delivered by the fiber is dissipated by thermal conduction to surrounding tissue and blood perfusion.

[0106] The vaporization and sublimation threshold depends on the power intensity at the output of the fiber optic and therefore depends on the input power, the emission surface, and the dose, i.e. the amount of energy needed to give rise to the phenomenon. The wavelength also becomes important because it determines the type of interaction between laser radiation and tissue. In particular the radiation absorption coefficient varies with the wavelength.

[0107] For instance, for a laser radiation having a wavelength of 1064 nm, 2W power (continuous wave) is enough to achieve tissue sublimation and cavity formation with an optical fiber having a core diameter of 300 micrometers.

[0108] The lower power threshold to achieve vaporization and sublimation on a specific tissue depends on the power, energy delivery mode (continuous or pulsed), fiber tip (dimension and shape), absorption and scattering coefficients, which in turn depend on tissue and wavelength combination. The higher the power used, the faster the cavity formation speed. At 1064 nm the

preferred power values are from 3 to 7 W in continuous wave mode.

[0109] The ratio between power and fiber tip surface defines the emitted radiation intensity. For 5 W delivered by a 300 micrometer core fiber diameter, the intensity is 17.7 W/mm². Intensities should be above 1 W/mm² to excite cavity formation with a 1064 nm wavelength and continuous wave with a dose of 1200-1800 J. With other laser wavelengths intensities are different and can easily be re-assessed.

[0110] Doses range from a few hundreds to a few thousands Joule. The optimal results are obtained from 600 J to 1800 J. There is a linear relationship between cavity volume and dose in this range. Doses greater than 1800 J induce a slight and not interesting increase in cavity enlargement.

[0111] According to some embodiments, for instance using a 5 W laser power and a wavelength of 1064 nm, the time required to start formation of a cavity in the tissue can be around 40 seconds, which correspond to an energy dose of 200 J. A first heating step precedes the actual tissue vaporization and cavity formation. The initial heating step is needed to bring the vaporization of water contained in the tissues. The first heating step is followed by cavity formation with a cavity volume which increases asymptotically. After a dose of 1800 J has been delivered (approximately 6 minutes) the maximum cavity volume has been achieved. Continued laser radiation with the fiber tip in the same position will not lead to any significant increase in the cavity volume.

[0112] The cavity formation process can be monitored by the ultrasound probe, exploiting the variable echogenicity of the tissues. During the first treatment step, preceding vaporization, the lased tissues will have a negligible variation of echogenicity. The echogenicity will increase when vapor bubbles start developing. The echogenicity variations can be detected through the ultrasound probe. According to other embodiments, cavity formation can be monitored by Magnetic Resonance Imaging (MRI) or Computer Tomography (CT) imaging and can be detected as change in the tissue density. A combination of various imaging techniques is not ruled out.

[0113] Laser wavelengths that can be employed can be for instance those, which can be guided by optical fibers or wave guides usually in the visible and near infrared region. In some embodiments UV radiation can be used as well. According to the absorption spectrum, different wavelengths interact with different chromophores or water contained in the tissue. Thus, for each laser wavelength a set of lasing parameters should be evaluated in order to obtain tissue removal by vaporization and sublimation (i.e. vaporization and sublimation threshold should be evaluated for each configuration of lasing parameters in order to produce a cavity in a specific tissue). I.e. ablation parameters can be adjusted based upon the wavelength used. In some embodiments, a 10.6 microns laser radiation generated by a CO₂ laser source can also be used as waveguide are now available, which can guide also this radiation wavelength. According to other embodiments the following laser sources can be used: Nd:YAG laser emitting at 1064 nm; thulium laser emitting at 2010 nm; holmium lasers emitting at 2100 nm. Other suitable laser sources can include erbium laser at 2940 nm, using a hollow waveguide.

[0114] In some embodiments, different laser sources can be used in combination.

[0115] In some embodiments, the laser radiation can be continuous. In other embodiments a pulsed laser source can be used. Tests ex-vivo have been carried out with a pulsed holmium laser source using porcine liver tissue maintained at 20° C. (+/-5° C.) in a thermostatic bath. This temperature value was chosen with reference to the physiological in vivo value (about 37° C.). The operating temperature used during ex-vivo tests was reduced to 20° C. in order to take into consideration the absence of local blood perfusion which, in a living body, enables local heat removal during laser treatment. A holmium laser source with a wavelength of 2100 nm and pulsed emission was used in combination with a quartz optical fiber with a 550 micrometer core diameter.

[0116] Different pulse repetition frequencies and pulse amplitudes have been tested. Specifically, pulse repetition frequencies from 1 Hz to 20 Hz have been used, in combination with pulse amplitudes ranging from 0.2 Joule to 2 Joule. Different combinations of pulse repetition frequencies (PRF) and pulse amplitudes result in different power values. The following table

summarizes different trials performed with different pulse repetition frequencies, pulse amplitude values, power, energy doses and treatment times (value in bold are constant for subgroup of experimental tests). The last column reports the dimensions of the cavity obtained in the liver tissue (length and width in millimeters):

TABLE-US-00001		Expo-	Cav-	sure	ity	Cavity	Pulse	Dose	Time	forma-	dimension	Sample	PRF																																																																																																																															
Energy	Power [Joule]	[s]	tion	[mm × mm]	#1	20 Hz	0.5	10 W	1200	120	Yes	14 × 6	#2	20 Hz	0.5	10 W	900	90	Yes	16 × 5	#3	20 Hz	0.5	10 W	600	60	Yes	10 × 3	#4	20 Hz	0.5	10 W	300	30	Yes	9 × 3	#5	20 Hz	0.5	10 W	100	10	Yes	7 × 3	#6	20 Hz	0.4	8 W	100	12.5	Yes	5 × 2	#7	20 Hz	0.3	6 W	100	16.7	Yes	5 × 2	#8	20 Hz	0.2	4 W	100	25	Yes	5 × 2	#9	20 Hz	0.4	8 W	80	10	Yes	8 × 3	#10	20 Hz	0.3	6 W	60	10	Yes	7 × 2	#11	20 Hz	0.2	4 W	40	10	No	NA	#17	5 Hz	2	10 W	100	10	Yes	8 × 3	#13	15 Hz	1	15 W	100	6.7	Yes	6 × 2	#12	15 Hz	0.5	7.5 W	100	13.3	Yes	7 × 2	#14	10 Hz	0.5	5 W	100	20	Yes	8 × 2	#16	5 Hz	0.5	2.5 W	100	40	Yes	6 × 2	#15	1 Hz	0.5	0.5 W	100	200	No	NA

[0117] According exemplary embodiments of the laser ablation method disclosed herein, the emission parameters may be maintained constant during the entire treatment time. However, this may not always be possible or preferred. In some embodiments, the laser ablation treatment can be performed with a gradually increasing emission power, for instance to prevent tissue explosion due to abrupt vapor formation. In other embodiments, higher power values can be used at the beginning of the treatment, before the tissue cavity starts forming, followed by lower power emission, which can be used to promote hemostasis.

[0118] When pulsed laser emissions are used, the pulse repetition frequency, or pulse repetition rate, and the energy per pulse can be used as further selectable and adjustable parameters. The same mean power can be indeed achieved with different combinations of pulse repetition rates and energy per pulse. However highly energetic pulses with a low pulse repetition rate will have a different effect on the treated tissue than pulses having a lower energy but a higher repetition rate.

[0119] As mentioned, the method involves the use of one or more fibers that are trans-perineally introduced in the prostate by means of thin needles or introducers. The more fibers are placed inside the lobes **1A**, **1B** of the prostate **1** the quicker and more effective the treatment will be. The optimal trade-off between effectiveness and time duration consists of simultaneous energy delivery, which involves one or two fibers **13** and respective introducers **11** per lobe (thus from two to four optical fibers **13** for the whole gland) as shown in FIG. 2. However, a smaller number of fibers (FIG. 1) or a larger number of fibers (FIGS. 5 and 6) can be used.

[0120] Two or more adjacent cavities may merge into a single cavity, if so desired. The treatment can also be performed with a single fiber and with multiple illuminations and repositioning.

[0121] According to the cavity shape, which depends upon several parameters, such as shape of the fiber tip, power, dose, absorption coefficient, it is possible to define safety criteria for the fiber tip positioning. As a matter of fact, the fiber tip can be placed at a safety distance from critical structures inside, around and adjacent the prostate **1**, such as the urethral lumen **5** inside the prostate **1**, the capsule **7** around the prostate **1** and the two neurovascular bundles **9** outside and adjacent the prostate **1**.

[0122] According to some embodiments, using a 3 W continuous wave power with a flat tip optical fiber the following safety rules can be applied: lateral distance equal to or larger than 5 mm (optimal distance 10 mm); front distance between the fiber tip and the capsule **7**, at least 15 mm. Higher power can be used with redefinition of the safety criteria. Higher power levels generate longer cavities. Thus, a larger safety distance between the fiber tip and the prostate base **1D** (front distance between capsule and fiber tip) should be adopted.

[0123] The mutual distance between fibers **13** and needles **11** depends on how many cavities **C** are planned to be formed in the tissue. Usually said mutual distance ranges between 5 mm and 15 mm depending upon total prostate volume and other constraints imposed by the above mentioned safety criteria (distance from capsule and urethra, for instance).

[0124] At 5 W power, safety distances should be re-assessed, especially as far as the front distance, i.e. the distance between the capsule and the fiber tip is concerned).

[0125] Depending on the prostate volume and in particular the longitudinal length (from base to apex), it is possible to gradually and stepwise withdraw the needles **11** and optical fibers **13** and perform a laser radiation at each newly reached position (pull-back maneuver), as described above in connection with FIGS. **3A-3C**. This allows maximizing volume removal with a single needle insertion. The extension of each withdrawal movement, i.e. the length of the needle **11** and the fiber **13** which is extracted from the prostate **1** at each pull-back maneuver, can be assessed by the aid of an ultrasound or magnetic resonance imaging system or by ticks on the needle cannula.

[0126] In some embodiments, the treatment can start with the fiber tips positioned at 15 mm from the capsule (prostate base) in the front direction and after a first energy delivery, if the apex-to-base distance allows it, the needle can be pulled back and a new dose of laser energy can be delivered in the new fiber position. The number of pull-back movements is related to the apex-base dimension of the gland **1**. The greater the number of pull-back steps, the greater the amount of tissue removed from the gland **1**. For each insertion of the fibers the treatment stops when it is not possible to withdraw the fiber further, based on the safety rule (minimum distance between capsule and fiber tip must be respected).

[0127] The method can be carried out with optical fibers having a flat tip. In other embodiments, however, special optical fibers, having a different geometry, can be used. Special optical fibers may have a special tip shape obtained by chemical etching, mechanical lapping, thermal fusion or other chemical, physical or mechanical treatment.

[0128] In some embodiments, the optical fiber may be adapted to achieve side firing or globular firing. As used herein, the term “side firing” optical fiber can be understood as an optical fiber which emits at least one optical beam from the side surface thereof. As used herein, the term “globular firing” optical fiber can be understood as a fiber having a tip shaped such as to generate a substantially globular or spherical emission.

[0129] In some embodiments, globular firing can be obtained with an optical fiber having a conical tip, rather than a flat, planar tip. A globular firing optical fiber can generate substantially spherical cavities in the tissue. FIG. **8** schematically illustrates a cross sectional view according to a sagittal plane of a prostate **1** under treatment, similar to FIGS. **3A-3C**. The same elements are labeled with the same reference numbers as in FIGS. **3A-3C** and will not be described again. In the embodiment of FIG. **8** the optical fiber tip **13T** is shaped such as to have a globular, i.e. substantially spherical emission, which generates a substantially isotropic energy distribution on a spherical surface. For instance, a conical fiber tip can be used. This results in a spherical volume **V1** of tissue vaporization and sublimation and consequently generates a substantially spherical cavity. **V2**, **V3**, **V4** indicate ablation volumes which are obtained in subsequent pull-back steps, during which the introducers **11** and the optical fibers **13** guided therein are gradually withdrawn from the prostate **1**.

[0130] The mechanism of tissue removal remains the same as described above. The cyclic dehydration, vaporization and sublimation process takes place on enlarging spherical surfaces due to isotropic energy deliver energy. Spherical cavities are more suitable for treatments in small organs and close to critical vital structures, because of the geometrical shape of the spherical cavity formed, the surface whereof is maintained at substantially the same distance from the fiber tip **13T** in all directions. In other words, the shape of the cavity generated by laser vaporization or sublimation of the tissues is not related to the direction of insertion of the needles, contrary to what happens when optical fibers with a flat tip are used, which generate elliptic cavities.

[0131] Also with globular firing optical fibers laser power and energy can be set at 3 W and 1800 J, respectively, for a 1064 nm wavelength laser source operating in a continuous wave mode. In this case, the above mentioned safety rules setting the minimum distances between the tips of the optical fibers and the critical structures inside, around and outside the prostate must be adapted to the new shape of the cavity that is produced. In some embodiments using a globular (spherical)

firing optical fiber with a 5 W laser source, the tip **13T** of each optical fiber **13** should be at least 1 cm from the critical structures (urethra, capsule) even in the transverse direction.

[0132] The use of isotropic or quasi-isotropic radiators (optical fiber tips **13T**) allows using higher power ranges than those which can be applied using free handle flat-tip optical fibers. The safety distances between fiber tip and critical structures of the prostate should be reassessed for each case.

[0133] As mentioned above, the positioning of the needles or introducers **11** can be performed under imaging guidance, for instance under ultrasound guidance preferably with a trans-rectal probe and preferably with a bi-plane probe. A bi-plane probe allows displaying images according to transverse and longitudinal (sagittal) planes of the prostatic gland.

[0134] FIG. **9** shows a sectional view according to a sagittal plane of a prostate during insertion of two needles **11** in one of the lobes **1A**, **1B** of the prostate **1**, using US (ultrasound) imaging guidance. An ultrasound (US) trans-rectal probe **31** can be inserted into the rectum and allows imaging the portion of interest. The US trans-rectal probe **31** can be connected to an ultrasound imager **33** which can be provided with a display **35**. The trans-perineal insertion of the needles or introducers **11** can be performed freehand or through a needle guide device **37** connect to the trans-rectal probe **31**.

[0135] The US trans-rectal probe **31** can be maintained in place also after insertion of the needles **11** and optical fibers **13**, during part of the treatment or during the entire treatment. Ultrasound imaging can be used to control that tissue vaporization or sublimation and the formation of the cavity in the gland tissue proceeds correctly. Additionally, in case of pull-back maneuver, as described above in connection with FIGS. **3A**, **3B**, **3C**, US imaging can assist the operator to retract the needles **11** by the required length at each pull-back step, such that subsequently removed tissue volumes merge in a single, longitudinally extending empty cavity formed in the gland tissue.

[0136] During treatment a catheter **36** can be placed into the urethra up to the bladder **38** to provide a reference point in the US image shown on the display **35**.

[0137] In some embodiments of the method disclosed herein, removal of gaseous by-products, generated by vaporization and/or sublimation of the lased gland tissues may be particularly beneficial. Removing gaseous by-product generated by interaction of the laser radiation with the gland tissue may promote and facilitate the entire tissue removal process. Vapor or other side products which are not removed are otherwise slowly absorbed by microcirculation and blood perfusion, a process which takes some time until complete elimination is obtained. Removal of the vapor or other side products through the needles during lasing or after lasing but prior to removing the needles from the prostate can substantially accelerate the reduction of the prostate volume and thus the relief on the urethra **5**.

[0138] During lasing and gas formation following tissue vaporization or sublimation, a positive pressure is generated inside the cavity formed by the laser radiation. The gas can flow in a gap between the optical fiber **13** and the inner surface of the needle **11**, thus escaping from the needle hub. For this reason, according to some embodiments, the coupling between needle hub and optical fiber hub is kept opened. In other embodiments, selectively closable and openable ports can be provided, for gas or vapor venting purposes. A possible coupling system with grooves for gas discharge is disclosed in U.S. Pat. No. 8,265,446, the content whereof is entirely incorporated herein by reference.

[0139] According to some embodiments, improved removal capability of the needle shaft or introducer cannula **11** can be achieved by providing side holes or ports in the wall of the needle close to the tip thereof. These side holes or ports can prevent obstruction of the main channel of the needle in case solid particles and debris are present in the cavity generated by laser radiation.

[0140] FIGS. **10** and **10A** schematically illustrate a needle or introducer **11** with a needle hub **11A** and a needle tip **11T**. An optical fiber **13** is introduced through the needle **11** and the emitting fiber tip **13T** is located in a cavity being formed by lasing the surrounding prostatic tissue. The cavity **C** contains pressurized vapor or gaseous side-products generated by the vaporization or sublimation

of the tissue. The gas escapes through the annular gap between the optical fiber **13** and the inner surface of the needle **11**. The fiber hub is shown at **13A** and at least one venting port is available between the fiber hub **13A** and the needle hub **11A**, such that gas G can escape from the cavity C. In FIG. **10A**, which shows an enlargement of the needle and fiber tip area, side holes **11H** near the needle tip **11T** are provided, which facilitate gas or vapor removal should the end of the needle become clogged by solid or high-density liquid debris.

[0141] In some embodiments, improved efficiency can be achieved by promoting gas and vapor extraction from the cavity being formed in the prostate tissue. According to some embodiments, a pressure reduction can be generated in the interior of the needles **11**, for instance by means of a suction device, such as a vacuum pump or the like.

[0142] With continuing reference to FIG. **10**, in FIG. **11** a vacuum pump **41** is fluidly coupled to the interior of the needle **11**. The pump **41** is adapted remove gas or vapor with a negative pressure from outside. The negative pressure in the needle **11** can be enough to aspirate gas or vapor from the cavity C being formed by the laser radiation in the prostate tissue, such that the pressure in the cavity C is maintained under control. A first duct **43** can fluidly couple the interior of the needle **11** to a collecting tank **46**. The latter can be fluidly coupled through a duct **45** to the vacuum pump **41**. Particles or condensed vapor can be collected in the tank **44** and prevented from reaching the vacuum pump **41**.

[0143] According to further exemplary embodiments, a double way introducer or needle **11** can be used, in order to apply suction to a first way and refill the cavity under formation with fresh air through the other way. This can be beneficial in order to fully remove from the cavity C the gases or vapors that can interfere with the laser radiation decreasing the effectiveness of laser-to-tissue interaction.

[0144] With continuing reference to FIGS. **10** and **11**, FIG. **12** illustrates an embodiment wherein a two-way introducer or needle **11** is used in combination with suction and fresh air feeding inside the cavity C being formed during laser vaporization or sublimation of the lased prostate tissue. The same reference numbers used in the previous figures designate the same or corresponding parts or elements, which are not described again. FIGS. **13** and **14** illustrate cross-sectional views according to line XIII-XIII of FIG. **12**, of two exemplary embodiments of a dual pathway needle or inserter **11**. The needle **11** may comprise an external cannula **11X** and an internal tubular element **11Y**. The optical fiber **13** can be housed in the internal tubular element **11Y** in a substantially coaxial position therewith. In some embodiments (FIG. **13**) distancing and centering members **55** can be arranged between the cannula **11X** and the inner tubular element **11Y**. A first annular passageway **51** is thus formed between the cannula **11X** and the inner tubular element **11Y**. A second annular passageway **53** is formed between the inner tubular element **11Y** and the optical fiber **13**.

[0145] Referring to FIG. **12**, a duct **57** can be in fluid communication with the environment and with the first pathway **53**. An air filter **59** can prevent pollutants, and pathogens, such as micro-organisms, from entering the needle passageway **53**. The second passageway **53** is fluidly coupled to the duct **43** and thus, through the collector tank **44** and the duct **45**, with the suction or vacuum pump **41**.

[0146] The device illustrated in FIGS. **12**, **13**, **14** operates as follows. Once the needle **11** has been placed in the correct position inside the prostate, possibly with the aid of US imaging or other imaging facilities, the laser source is activated and laser radiation interacts with the tissue surrounding the tip **13T** of the optical fiber **13**, causing vaporization and/or sublimation of the tissue, forming a cavity C of gradually increasing volume. The vacuum pump **41** removes vapors or gases from the cavity C under formation through the second pathway **53** of the needle, and the negative pressure generated in the cavity C causes fresh air to enter the cavity through the air filter **59**, the duct **57** and the first passageway **51** of the needle **11**. The vapor removal and air circulation can continue for the entire cavity formation step, such that the cavity C will be free or substantially free of laser absorbing gaseous side products generated by laser-tissue interaction. This may be

beneficial in terms of speed of treatment and dimension of the cavity C obtained for each position of the needle or introducer **11**.

[0147] According to yet further embodiments, enhanced BPH treatment results can be achieved by combining tissue ablation through vaporization with a mechanical action through the urethra **5**. In some exemplary embodiments, an inflatable balloon can be introduced in the urethral lumen **5** of the patient prior, during or after insertion of the needles **11**.

[0148] With continuing reference to FIGS. **1** to **14**, FIGS. **15A**, **15B** and **15C** illustrate a cross sectional view along a sagittal plane of the prostate during treatment with optical fibers introduced trans-perineally in the prostate and one or more inflatable balloons placed in the urethra **5**.

[0149] In FIG. **15A** an elongated inflatable balloon **61** has been placed in the urethra **5** of the patient by means of a catheter **62** and inflated. The balloon **61** can be inflated, for instance, with a physiologic solution at suitable pressure and temperature conditions. The balloon **61** can be introduced in the urethra **5** with any suitable means, such as the catheter **62**. Inflatable balloons for angioplasty or similar surgical applications are known, and they can be adapted for the use disclosed herein.

[0150] Upon inflation, the balloon **61** causes compression of the surrounding tissue of the prostatic gland and dilation of the urethra **5** at its physiologically normal conditions, or even larger. Compression of the surrounding tissues can facilitate evacuation of the gaseous or vapor by-products generated by laser-tissue interaction and tissue vaporization or sublimation, as disclosed above.

[0151] In addition, due to a thermo-plastic effect, once the balloon **61** is deflated and removed through the urethra **5**, the surrounding tissues will at least partly maintain their compressed condition, thus providing more efficient relief to the urethra **5** immediately after treatment.

[0152] In some embodiments, the filling fluid used to inflate the balloon **61** can be circulated, to maintain the fluid temperature under control. In some embodiments, the fluid temperature can be maintained above basal temperature, e.g. around 40-42° C. In other embodiments, the fluid temperature can be maintained at a lower temperature, even below the body temperature. The temperature control and fluid circulation can be used to cool the urethra **5** and the tissue immediately surrounding the urethra, preventing damages due to over-heating.

[0153] In the embodiment of FIG. **15A** a single balloon **61** with an enlarged head end **61A** is retained in the correct position by introducing the balloon **61** in the urethra **5** until the head end **61A** thereof reaches the bladder **38**.

[0154] In FIG. **15B**, with continuing reference to FIGS. **1-15A**, an inflatable balloon **61** is used, which has an elongated central body and two expanded ends **61A**, **61B**. These latter provide for a precise and correct positioning of the balloon **61** in the urethra **5** and contribute in maintaining the inflated balloon **61** in the correct position during lasing of the prostate **1**.

[0155] In FIG. **15C**, with continuing reference to FIGS. **1-15B**, two inflatable balloons **61X**, **61Y** are used in combination with one another. The first balloon **61X** can be introduced in the bladder **38** and the second balloon **61Y** can be positioned in the urethra **5**. The two balloons **61X**, **61Y** can be introduced by means of the same catheter **61** and the balloon **61X** provides for safe positioning of the second balloon **61Y** in the urethra **5**.

[0156] Balloons **61** of different shapes and/or dimensions can be selected by the operator based upon the dimension of the prostate **1** to be treated, on the anatomic features of the patients, or upon other factors.

[0157] FIGS. **16**, **17** and **18** show three flow-charts which summarize some of the methods of treating BPH disclosed herein.

[0158] In some embodiments, treatment safety can be increased by adding for instance temperature control facilities, aimed at preventing over-heating of critical structures inside or around the prostate **1** under treatment.

[0159] Referring to FIGS. **19** and **20**, with continuing reference to FIGS. **1** to **15C**, temperature

sensing arrangements **71** can be introduced in the prostate **1** under treatment in combination with needles **11** and optical fibers **13**. Thermocouples, thermoresistors or temperature measuring fibers can be introduced through needles or cannulae in the prostate. In FIGS. **19** and **20** reference number **71** indicates any possible kind of temperature measurement device, apparatus or arrangement. For instance, a pervious needle or a cannula can be used to introduce a thermocouple or a thermoresistor near critical areas of the prostate **1**. In some embodiments an optical fiber with a Bragg grating can be used instead of, or in combination with thermocouples and/or thermoresistors. [0160] In some embodiments, the temperature sensitive portion of the temperature sensing arrangement can be located at or near the tip of the device **71**. In other embodiments, temperature sensing areas can be located in several positions along the axial extension of the device **71**, for example if optical fibers with a Bragg grating are used. These latter will be able to detect the temperature at different depths in the prostate.

[0161] In FIGS. **19** and **20** two temperature sensing arrangements **71** are introduced in the prostate **1**, one between the urethra **5** and the area where a first cavity **10** will be generated by a first optical fiber **13**. A second temperature sensing arrangement **71** is arranged between the neuro-vascular bundle **9** and the area where a second cavity **10** will be generated by a second optical fiber **13**. Thermal damages of the urethra **5** and of the neuro-vascular bundles **9**, due to over-heating or excessive laser radiation, can thus be prevented. The temperature sensors can be functionally coupled to the control unit **21**, such that the emission parameters of the laser sources **19A**, **19B** can be modulated automatically to maintain the tissue temperature under control and prevent thermal damages of critical structures in and around the prostatic gland **1**. For instance the control unit **21** can be adapted to reduce the mean power, or the peak pulse energy, or other laser emission parameters, if the temperature measured by the temperature sensing arrangement **71** increases above a given safety threshold. In some embodiments, one or more temperature sensing arrangements **71** can be combined with some or each one of the optical fibers **13**, such that each or some of the laser sources **19A**, **19B** can be automatically controlled through data from the temperature sensing arrangements **71** combined with the respective optical fibers **13**.

[0162] In other embodiments, temperature information from the temperature sensing arrangements **71** can be made available to the operator, for instance through a suitable user interface, such as a display or monitor. The operator will then manually modify the laser emission parameters based on the temperature information from the temperature sensing arrangements.

[0163] According to a further aspect, the disclosure also concerns a method of reducing a volume of a benign or malignant tumor in an organ of a patient in need of said treatment. According to a yet further aspect, the disclosure concerns a method for removing tissue from an organ of a patient according to a mini-invasive technique.

[0164] According to some embodiments the method comprises the following steps. A first step involves the introduction of at least one energy delivery device in a first position in an organ of the patient. The energy delivery device can include an optical fiber coupled to a laser source. The optical fiber can be introduced through a needle or introducer. Once the energy delivery device has been placed in the correct position, possibly with the aid of a ultrasound or other imaging device, the method comprises the step of delivering energy from an energy source coupled to the energy delivery device. The energy is delivered through the energy delivery device to a first volume of tissue of the organ where the energy delivery device has been positioned. Energy is delivered until at least a portion of the first volume is vaporized or sublimated and a cavity is formed in the organ tissue. According to some embodiments, during or after energy delivery, gaseous side-products generated by tissue vaporization or sublimation can be removed, e.g. by suction, as described above. The method further includes the step of removing the energy delivery device from the organ. If needed, prior to removing or after removing, the energy delivery device can be re-positioned in a different position inside the organ, to repeat the above steps and remove by vaporization and/or sublimation a further portion of tissue.

[0165] The tissue can be hard tissue, such as bone or the like, or soft tissue, such as liver, thyroid, pancreas, brain, or other organs which may need treatment. In particular if soft tissue is treated, the volume of the cavity thus formed can be reduced, e.g. by massage on the organ.

[0166] According to some embodiments, the method can further include the step of introducing in the cavity formed by tissue vaporization or sublimation, at least one medically active element. In some embodiments, the method includes the step of introducing in the cavity thus formed at least one of the following: a medicament, a drug, a slow-absorption drug, a radioactive seed, such as a seed for brachytherapy, a chemotherapeutic agent, a combination thereof. The method allows forming a cavity in the organ to be treated without resorting to cutting instruments, which would destroy also portions of healthy tissue surrounding the area where the tumor to be treated is positioned.

[0167] While in some embodiments the tissue to be removed can be a benign or malignant tumoral tissue, the method is not limited to removal of tumoral tissue. More in general, a mini-invasive surgical method is disclosed, aimed at generating a cavity in an organ in a body of a patient, for instance an organ which is difficult to reach or which is usually reached through cutting of tissue surrounding the area where the cavity is to be formed. The method disclosed herein suggests a new way of mini-invasively reaching the site where tissue is to be removed, reducing as far as possible any impact on the surrounding tissue. This can be of paramount importance, for instance, when the need exists to preserve the integrity of surrounding tissue, such as in brain surgery.

[0168] FIGS. **21A**, **21B**, **21C**, **21D** pictorially illustrate steps of a method involving tissue removal in a generic organ **101** through an introducer **11** and an energy delivery device **13**. In the exemplary embodiment of FIGS. **21A-21D** the introducer **11** includes a hollow needle and the energy delivery device **13** includes an optical fiber. The energy source may include a laser source. In FIG. **21A** the introducer **11** and the optical fiber **13** have been introduced through surrounding tissue **102** in the organ **101** to be treated. In FIG. **21B** a cavity **103** has been generated by energy-tissue interaction and tissue vaporization and/or sublimation. Resulting gaseous by-products can be removed as described above, through the introducer **11**.

[0169] In FIG. **21C** the optical fiber **13** has been removed, while the introducer **11** is still in place. The cavity **103** has been generated in the organ **101**. In a further step of the method a basket, for example a shape memory basket **105**, **105A** can be introduced through the introducer **11** and the head thereof **105A** can be expanded in the cavity **103**, to prevent collapsing of the cavity **103**. Expandable baskets useful in the present method are known in the art. For instance, the basket **105**, **105A** can be configured as disclosed in US20180042625. FIG. **21D** illustrates the expanded basket **105**, **105A**.

[0170] The basket **105**, **105A** can be used to maintain the cavity **103** in an expanded condition and temporarily prevent collapsing of the cavity. For instance, the cavity **103** can be maintained in the expanded status in order to facilitate introduction therein of a medium, for instance a liquid or semi-liquid medium, such as a drug carrier.

[0171] In some embodiments, a different medium, such as a solid medium can be introduced in the cavity. For instance, radioactive seeds for brachytherapy can be placed in the expanded cavity **103**.

[0172] The basket can be recoverable. The method can thus include a step of introducing basket in the cavity generated by tissue vaporization or sublimation and a subsequent step of removing the basket from the cavity. According to other embodiments, the basket can be left in the cavity and can be made of absorbable materials.

[0173] According to some embodiments, nanoparticles, for instance gold or iron-based nanoparticles as drug carriers can be introduced in the cavity **103**. Drugs, such as monoclonal drugs or monoclonal antibodies can be introduced in the cavity **103**, possibly through suitable carriers, such as nanoparticles.

[0174] Drugs or medicaments can be conveyed in a liquid or semi-liquid suspension. An expandable basket **105**, **105A** can prevent cavity reduction and facilitate the insertion of the

suspension. In some embodiments, the basket can be retained in place until the drug has been absorbed.

[0175] In other embodiments, for instance if a semi-liquid or highly viscous carrier is used, a basket may be dispensed with or may be removed soon after inoculation of the drug suspension.

[0176] FIG. 22 illustrates a cross sectional view similar to FIGS. 21A-21D, wherein the cavity 103 has been filled with a liquid, semi-liquid or gel substance 107. The substance 107 may contain a drug or medicament, and may in turn comprise particles, such as nanoparticles used as drug carriers, such as monoclonal antibodies or drugs. The filling substance can be introduced through the introducer 11 or through a cannula inserted into the organ 101. Introducer 11 can be used also for the insertion of the basket 105, 105A. In some embodiments, the basket 105, 105A can be removed along with the introducer 11 from the organ 101 upon injection of the substance 107. The tissue wall surrounding the introducer or needle 11 can collapse after removal of the needle 11, such that reverse flow of the substance 107 through the perforation can be prevented.

[0177] FIG. 23 illustrates a cross sectional view similar to FIGS. 21A-21D, 22, wherein a solid substance, such as radioactive seeds, has been placed in the cavity 103. While in FIG. 23 no basket is shown in the cavity 103, in other embodiments, the cavity 103 may be maintained in its expanded condition by a basket. The use of a basket or other effectors aimed at preserving the inner volume of the cavity 103 may be beneficial when the cavity 103 has been formed in a soft tissue, or is surrounded by soft tissue. In some embodiments, the cavity 103 may be formed in harder tissue, such as bone tissue or cartilaginous tissue. An effector, such as a basket, aimed at maintaining the cavity wide spread could then be dispensed with.

[0178] While the invention has been described in terms of various specific embodiments, it will be apparent to those of ordinary skill in the art that many modifications, changes, and omissions are possible without departing from the spirit and scope of the claims. In addition, unless specified otherwise herein, the order or sequence of any process or method steps may be varied or re-sequenced according to alternative embodiments.

[0179] For instance, while the above described exemplary embodiments use laser sources and laser energy to obtain tissue ablation through vaporization, the option is not ruled out of using a different power source, such as a radiofrequency power source, and respective energy delivery device, to deliver the energy in the tissue volumes where ablation is required.

[0180] While specific embodiments of the invention have been shown and described in detail to illustrate the application of the principles of the invention, it will be understood that the invention may be embodied otherwise without departing from such principles.

Claims

1. A method of treating prostatic tissue in a patient affected by a prostate disease, comprising the following steps: (a) trans-perineally introducing at least one energy delivery device reaching at least one prostate lobe requiring treatment; (b) delivering energy from an energy source through the at least one energy delivery device to a first volume of tissue of said prostate, until said first volume is vaporized or sublimated and a cavity is formed in the prostate tissue; (c) pulling back the energy delivery device from the first position to a second position; (d) delivering energy from the energy source through the energy delivery device to a second volume of tissue of said prostate while the energy delivery device is in the second position, until said second volume is vaporized or sublimated and the cavity is enlarged; and (e) repeating steps (c) and (d), if needed; (f) removing the energy delivery device from the prostate.
2. The method of claim 1, wherein the energy source comprises a laser source.
3. The method of claim 1, further comprising the step of removing vapor or gas resulting from the tissue vaporization or sublimation while energy is delivered through the at least one energy delivery device.

4. The method of claim 1, further comprising the step of removing vapor or gas resulting from the tissue vaporization or sublimation through the at least one energy delivery device.
 5. The method of claim 4, wherein the step of removing vapor or gas comprises the step of applying a negative pressure through the at least one energy delivery device.
 6. The method of claim 4, wherein the step of removing vapor or gas comprises the step of circulating a clean fluid medium in the cavity.
 7. The method of claim 4, wherein the step of removing vapor or gas comprises the step of fluidly coupling a vacuum pump to a pathway of the at least one energy delivery device.
 8. The method of claim 1, further comprising the steps of: introducing an inflatable balloon in the urethra of the patient; inflating said inflatable balloon; deflating said inflatable balloon after sublimation or vaporization of the tissue; and removing the inflatable balloon from the urethra.
 9. The method of claim 8, further comprising the step of circulating a fluid in the balloon.
 10. The method of claim 9, further comprising the step of controlling a temperature of the fluid circulating in the balloon.
 11. The method of claim 8, wherein said balloon has at least one enlarged terminal end adapted to maintain the balloon in position during treatment.
 12. The method of claim 1, further comprising the steps of: introducing a trans-rectal imaging probe in the rectum of the patient; and performing the treatment under imaging control through the trans-rectal imaging probe.
 13. The method of claim 1, further comprising the steps of: introducing at least one temperature sensing device in the prostate; and detecting the temperature in at least one location inside the prostate during energy delivering.
 14. The method of claim 13, further comprising the step of controlling the energy source based upon temperature information from the temperature sensing device.
 15. The method of claim 13, wherein the step of introducing at least one temperature sensing device comprises the step of introducing said temperature sensing device between a volume of action of the energy delivery device and a critical structure of the prostate.
 16. The method of claim 15, wherein the critical structure of the prostate is one of a urethra, a prostate capsule and neuro-vascular bundles around the prostate.
 17. The method of claim 13, wherein the temperature sensing device is introduced trans-perineally in the prostate.
 18. The method of claim 1, further comprising the step of performing a trans-rectal massage of the prostate after formation of said cavity, promoting cavity collapse.
 19. A method of removing tissue from an organ of a patient, the method comprising the following steps: introducing at least one energy delivery device in a first position in an organ of the patient; delivering energy from an energy source through the energy delivery device to a first volume of tissue of said organ, until said first volume is vaporized or sublimated and a cavity is formed in the organ tissue; removing the energy delivery device from the organ.
 20. The method of claim 19, further comprising the step of reducing the volume of the cavity.
 21. The method of claim 19, further comprising the step of introducing in said cavity at least one of: a medicament, a drug, a slow-absorption drug, a radioactive seed, a chemotherapeutic agent, nanoparticles, a drug carrier, an expandable basket.
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