Abstract. Radical prostatectomy is one of the most frequent therapeutic options used for the management of patients diagnosed with prostate cancer. Normal erectile function after radical prostatectomy is a great problem for numerous patients and a real challenge for urologists worldwide. The advancements that have been made over the years in terms of minimally invasive surgery, as well as in terms of surgical techniques, have reduced the incidence of erectile dysfunction, but even so, its rate remains high and the post-operative recovery of erectile function is a long and costly process. Phosphodiesterase 5 inhibitors have provided excellent results and have become the first-line treatment for these patients, followed by intracavernous injections with alprostadil. Several studies have underlined the impact of phosphodiesterase 5 inhibitors in terms of preventing the fibrotic changes that are responsible for the irreversible erectile dysfunction. The general opinion is that an erectile function recovery process should be started as soon as possible after surgery to prevent the negative effects of neuropraxia.

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1. Introduction

Prostate cancer (PCa) is one of the most frequent male malignancies worldwide. After the introduction of prostate-specific antigen (PSA) as a routine screening tool, it was noted that the incidence of advanced and metastatic PCa stages has constantly decreased, as increasing number of patients are diagnosed with localized PCa stages, which are potentially curable. This is in contrast to the pre-PSA era, when most patients were diagnosed in advanced stages, usually with metastatic disease, and associated poor prognosis (1).

Radical prostatectomy (RP) is one of the most frequent therapeutic options used for the management of patients diagnosed with PCa. This procedure offers good results in terms of disease-free survival and overall survival, but like any other major surgical procedures, this also presents several possible disadvantages, such as postoperative erectile dysfunction (ED) and urinary incontinence. Because the mean age for the time of diagnosis has decreased, postoperative erectile function (EF) recovery represents an important issue among urologists (2,3).

Although nerve-sparing (NS) techniques aim to preserve the EF, numerous patients complain of ED after surgery. Numerous factors can influence the baseline EF: Age, diabetes, obesity (5,6), smoking, chronic alcohol abuse, depression, as well as antidepressants (7), opioids, ketoconazole and numerous other drugs (8), chronic kidney disease, neurologic diseases (Parkinson's disease, multiple sclerosis, myelodysplasia, tumors, epilepsy, peripheral neuropathy), hypogonadism, hyperprolactinemia, thyroid dysfunction (hyperthyroidism) and hypocortisolism (9-11).

Diabetes mellitus is one of the most frequent causes responsible for ED, these patients presenting a much higher risk of developing ED compared with non-diabetic patients (up to
that the vein-occlusive dysfunction is time-dependent, amplifying itself with the increasing local fibrosis.

The studies published by Nehra et al (19) in 1996, and by Iacono et al (20) in 2005 reported significant cavernous structural changes following RP, that consisted of the replacement of cavernous smooth muscle fibers by collagen tissue. The authors observed that the degree of vein-occlusive dysfunction was directly influenced by the balance between smooth muscle fibers and collagen tissue.

Considering all these previously described mechanisms involved in the pathophysiology of post RP ED, numerous attempts have been made over the years to prevent these irreversible changes that significantly alter the patients' well-being.

3. The role of phosphodiesterase 5-inhibitors in EF recovery

The introduction of phosphodiesterase 5-inhibitors (PDE5-i) in the late 90's has changed the management of ED, these drugs proving to be an excellent option for patients suffering from ED. Numerous studies have evaluated the efficiency of PDE5-i in terms of post radical prostatectomy ED management (21).

The Reinvent clinical trial evaluated the efficiency of daily versus on-demand vardenafil. Over 600 patients who had undergone NSRP were randomized into three groups: group 1, daily 10 mg vardenafil and placebo on-demand; group 2, daily placebo plus on-demand vardenafil; and group 3, daily placebo and on-demand placebo. This trial had three phases: 9 months double-blind treatment, followed by a two months washout period and afterward another two months of open-label vardenafil. At the end of the study, the authors concluded that nightly vardenafil did not prove to be superior to on-demand vardenafil (22).

A similar conclusion was published by Pavlovich et al (23) in 2013. The patients enrolled in this trial have received either nightly sildenafil plus on-demand placebo or on-demand sildenafil plus daily placebo for one year. Daily sildenafil did not offer superior results compared with on-demand sildenafil, underlining that its chronic daily use is not justifiable for EF recovery in patients who have undergone bilateral NSRP.

Another study that was conducted over 12 months, followed by 1 month washout period, has aimed at identifying whether sildenafil administered daily is superior to on-demand sildenafil. At the end of this trial, the daily usage of sildenafil did not reach superior results compared with the on-demand sildenafil group (24).

The REACT clinical trial evaluated nightly versus on-demand tadalafil. The patients were randomized into three groups: 5 mg daily tadalafil, 20 mg on-demand tadalafil, and placebo. The patients received treatment for nine months, followed by a washout period of 6 weeks and afterward by a period of three months of daily tadalafil. The results achieved at the end of the first nine months have shown that daily tadalafil was superior to placebo, as well as to on-demand tadalafil. Regarding the efficiency of on-demand tadalafil versus placebo, the results did not reach a statistical significance. At the end of the drug washout period, the superiority of daily tadalafil was not maintained. After reintroducing daily tadalafil for three months, the IIEF and Sexual Encounter Profile-3 (SEP3) scores improved. In terms of penile shrinkage prevention, the daily usage of tadalafil has demonstrated superior results (25).
Kim and Sung (26) compared whether daily tadalafil administered for one year is inferior or not to tadalafil administered daily for two years. The study included 95 patients who had undergone NSRP. No statistically significant difference was noted at the end of the study between the two tadalafil groups (two years versus one year) in terms of EF recovery. The authors also reported that they did not reach any major statistical differences when they compared the patients with bilateral NSRP and those for whom unilateral NSRP was performed.

In a randomized clinical trial, Moncada et al (27) concluded that daily administration of tadalafil speeds up the EF recovery process. Their study included 423 patients who had undergone bilateral NSRP. The patients were randomized to receive either daily 5 mg tadalafil or on-demand 20 mg tadalafil or placebo, for nine months. The study also included a 6-week drug washout period and another twelve weeks of daily tadalafil for all the three groups of patients. The authors stated that the time of the treatment was too short for >50% of the patients to achieve a satisfactory erection recovery. They further reported that 25% of the patients reached an EF recovery (IIEF-EF ≥22) over a period that ranged between 5.8 and 9.3 months (5.8 months for patients with daily tadalafil versus 9 months for those with on-demand tadalafil and 9.3 months for patients who have received placebo), the daily administration of tadalafil favoring a faster EF recovery.

4. Discussion

Surgical experience and surgical approach are key elements that influence the postoperative EF recovery. Potdevin et al (28) compared the intra-fascial nerve-sparing technique with the inter-fascial technique and reported that the intra-fascial nerve-sparing approach offers superior results in terms of EF recovery. Numerous studies have reported the superiority of the robot-assisted approach over open and laparoscopic RP (ORP and LRP) in terms of nerve-sparing success and EF recovery.

Stolzenburg et al (2) analyzed the REACTT clinical trial in terms of surgical approach and its influence over EF impairment and reported that patients who had undergone nerve-sparing RARP associated superior chances (twice as high) of recovering a satisfactory EF (IIEF-EF ≥22) at the end of the drug washout period, as well as at the end of the open-label tadalafil period, compared with those for whom ORP or LRP was performed. When comparing the superiority of LRP over ORP in terms of postoperative erectile function recovery, no significant statistical difference has been achieved.

A study published in 2011, conducted on 128 patients who have undergone either RARP or LRP, concluded that the robotic approach associated with higher IIEF scores twelve months following surgery, as well as increased rates of EF recovery similar to baseline EF and a significantly faster EF recovery (29).

Greco et al (30) analyzed 457 patients who had undergone nerve-sparing LRP. The authors reported that EF assessed at twelve months following surgery was superior in patients for whom bilateral NS was achieved compared with those with unilateral NS (69 vs. 43%). On-demand 20 mg vardenafil was used after surgery, in order to improve the EF.

Another viable solution that could be used for postoperative EF recovery is represented by intracavernous injections (ICI) with alprostadil. The first mention of an EF recovery program following RP was made by Montorsi et al (31) in a study published in 1997 and it implied alprostadil ICI. This trial included 30 patients who had undergone bilateral NSRP. Following surgery, the patients were randomized to receive ICI with alprostadil three times a week, for twelve weeks, or no treatment. The authors reported that patients included in the ICI group achieved superior results in terms of EF recovery compared with the observational group (67 vs. 20%).

Numerous other studies underline the positive effect of intra-cavernous injections with alprostadil, but their use is much more limited compared with PDE5-inhibitors. This is usually related to the fact that patients must self-administer the injections and also to the discomfort/pain associated with these injections (32-35).

A study in 2015 (36) aimed at finding out if a longer alprostadil protocol could improve the postoperative erectile function in terms of spontaneous nocturnal erections. Two times a week ICIs with alprostadil were recommended, starting the first month following surgery. At approximately one year after the initiation of the alprostadil treatment, the patients were recommended to use PDE5-inhibitors and the non-responders were encouraged to continue the ICIs for another 12 months. The authors reported that alprostadil had significantly improved the patients' quality of life, but the results in terms of spontaneous nocturnal erections, between one year and two years of follow-up, were weak. The satisfaction of patients regarding the treatment efficiency had shown a decline between the 12th and 24th month.

The benefits of an EF recovery program have been underlined in numerous studies, but a clear protocol is lacking, regarding the time of treatment initiation, drug choice, administration frequency, and treatment length. According to literature, PDE5-i are the most common choice among urologists, due to their ease of use and safety, as well as their good results. The time of PDE5-i treatment initiation is not very clear, numerous studies reporting their administration several weeks after surgery, and other studies report earlier use, such as the next day following surgery or even before surgery. Another important aspect is whether chronic daily administration of PDE5 inhibitors is better than on-demand use, because the information available on this subject is, to a certain extent, contradictory. Numerous studies report the advantages of nightly administration over on-demand use, but several clinical trials have failed to prove it. In conclusion, ED after RP is an important health problem, especially because the mean age of patients diagnosed with PCa has decreased, due to the use of PSA and multiparametric MRI. Numerous studies have underlined the role of EF recovering programs, but a clear protocol, in terms of drug choice, treatment initiation and its extent, still does not exist. Nevertheless, PDE5-i have proven to be a good option for such patients. Their good results and ease of use have made them the first-line treatment for post-RP ED. Despite conflicting information whether daily usage is superior to on-demand administration, the majority of the existing studies favor their daily administration, as well as an early treatment initiation. Intra-cavernous alprostadil injections also provide good results, but due to their associated disadvantages, they remain a second option.
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Authors’ contribution

DRM, LI, CCD and ADS collected, analyzed and interpreted the patient data regarding the erectile function recovery after radical prostatectomy. CCD, OGB and DM had substantial contribution to the conception of the study and interpretation of the data. DRM, CCD and OGB drafted the manuscript and were major contributors in the writing of the manuscript. All authors read and approved the final version of the manuscript.

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Not applicable.

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Not applicable.

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