REVIEW

The Impact Of Pelvic Floor Muscle Training On Urinary Incontinence In Men After Radical Prostatectomy (RP) – A Systematic Review

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Department of Physiotherapy, Nicolaus Copernicus University in Torun Collegium Medicum in Bydgoszcz, Bydgoszcz, Poland **Purpose:** The purpose of this study was to determine the efficacy of pelvic floor muscle training (PFMT) in the treatment of urinary incontinence (UI) in men after radical prostatectomy (RP).

Methods: PubMed, ScienceDirect, and Cochrane Library databases were searched for studies published in years 2000–2019. We included randomized controlled trials in English which compare clinic-based vs home-based PFMT, preoperative and postoperative PFMT, supervised vs unsupervised PFMT, and PFMT alone vs no treatment at all.

Results: Eight articles were included in the final review. There was a total of 1078 patients aged 45–75 in all study groups. The study participants received radical retropubic prostatectomy or radical prostatectomy. Included studies assessed the following interventions: preoperative and postoperative PFMT, supervised vs home-based PFMT, unsupervised PFMT vs no treatment at all, and PFMT combined with resistance and flexibility exercises vs PFMT alone.

Conclusion: PFMT is an effective treatment for urinary incontinence in men after radical prostatectomy. PFMT improves not only physical parameters but also the quality of life of men after RP.

Keywords: pelvic floor muscle training, prostatectomy, urinary incontinence

Introduction

Prostate cancer (PCa) is a malignant tumor commonly diagnosed among men in the USA and Europe.¹ Age is strongly associated with the risk of prostate cancer. PCa is rare in men under 40, whereas the incidence of prostate cancer is increasing rapidly after the age of 55 years.¹ Epidemiological data show that in 350 men, only one person under the age of 50 will be diagnosed with prostate cancer, and the incidence increases to 1 in 52 men aged 50 to 59. In contrast, the incidence is almost 60% in men older than 65 years.² About 81% of cases of prostate cancer are detected early on, which allows patients to receive effective treatment. The choice of treatment depends on tumor stage and patient age. Although until recently surgical treatment was reserved for patients with prostate cancer limited to the prostate, nowadays criteria for surgical or radiotherapeutic treatment options are broadened with the intent to cure the illness.³ Prostatectomy may be performed classically as open surgery or as laparoscopic or robotic-assisted laparoscopic surgery. High-risk cancer patients may undergo radical prostatectomy which includes the removal of the entire prostate gland (in the section between the external sphincter and the bladder neck),

Correspondence: Magdalena Weber-Rajek Department of Physiotherapy, Nicolaus Copernicus University in Torun Collegium Medicum in Bydgoszcz, ul. Jagiellońska 13-15, Bydgoszcz 85-067, Poland Email magdawr69@gmail.com



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The Purpose Of The Systematic Review

The purpose of this study was to determine the efficacy of pelvic floor muscle training (PFMT) in the treatment of urinary incontinence (UI) in men prior to and after radical prostatectomy (RP). Randomized controlled trials (RCTs) on the use of PFMT after RP were included in this systematic review.

Methods

Data Sources

The present study is a systematic review of controlled clinical trials (RCTs). We identified studies from the following three databases: PubMed, Cochrane Library, and ScienceDirect from years 2000 to 2019.

Study Selection

The electronic medical databases were searched in May 2019 for studies from years 2000 to 2019. Three independent reviewers (K.S., Z.P., and A.S.) conducted the database searches to avoid missing relevant publications. Each investigator analyzed only RCTs published in English that examined the effect of pelvic floor muscle training on men with urinary incontinence pre- and post-prostatectomy. Identified reports were critically appraised for quality and relevance. Several thousand articles were found using the

following search keywords: nonsurgical treatments OR conservative treatment OR conservative management OR pelvic floor muscle training AND prostatectomy OR/AND urinary incontinence. RCTs that compared clinic-based vs home-based PFMT, PFMT prior to vs after the prostatectomy, supervised vs unsupervised PFMT, and PFMT alone vs no treatment at all were included in the systematic review. In addition, the following study exclusion criterion was added: combined therapy (electrostimulation, biofeedback, vibration, magnetic stimulation, and device therapy) and surgery type (robot assisted and TURP) to improve the quality of relevant searches. Only articles that met the following specific criteria were included in the analysis: prostate cancer patients received RP; the experimental or control groups received preoperative and postoperative PFMT or no treatment. Studies which contained insufficient information about the research methodology were excluded.

Data Extraction

Two reviewers (M.W.R. and A.R.) independently extracted the data to ensure data reliability and accuracy. The following information was extracted from all qualified studies: first author's name, country and year of publication, age of participants, sample size, intervention time, surgery and intervention type, study design, outcome measures, and main results. Moreover, the description of the study design contained details about the duration of observation and patient assessment time points. Any differences in opinion were resolved through discussion.

Quality Assessment

Systematic review methods were based on the PRISMA checklist (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) to strengthen reporting (Figure 1).⁶

Methodological quality of the selected studies was assessed using the PEDro scale which is commonly employed in systematic review papers (Table 1). The PEDro scale scores 10 items: eligibility, randomization, allocation of subjects, similarity at baseline, subject blinding, therapist blinding, assessor blinding, >85% follow-up for at least one key outcome, intention of treatment, statistical comparison between groups, estimated points, and variability measures for at least one key outcome. The final score is calculated by adding up all points. Two evaluators (Z.P. and A.S.) independently classified the data using the PEDro scale. In our systematic review, we included only high-quality studies which scored at least 6 points on the scale.⁷



Figure I Flow diagram of the study selection process.

Results And Discussion

Record Results

Consequently, 237 publications were found in 3 databases. Then, 30 duplicated articles were removed. The three researchers involved in data analysis read 207 abstracts and excluded 142 articles that did not focus on patients with UI following prostatectomy. After prior analysis of the full texts, another 57 articles were excluded based on the exclusion criteria. Ultimately, eight articles were included in this systematic review. According to the methodological quality assessment performed using the PEDro scale, 6 studies were carried out at a high level and only two at a satisfactory level.

Study Characteristics

The electronic medical databases were searched in May 2019. The studies were selected using a very detailed flowchart (Figure 1). Eight articles were included in the final review (Table 1). There was a total of 1078 patients (EG = 554, CG = 524) aged 45–75 in all study groups. In 6 studies, the

Study/Criteria Of The PEDro Scale	Centemero /2012/	Dubbelman /2012/	Filocamo /2005/	Manassero /2007/	Nilssen /2012/	Overgard /2008/	Park /2012/	Patel /2013/
Eligibility	+	+	+	+	+	+	+	+
Randomization	+	+	?	+	+	+	+	+
Allocation of subjects	+	+	?	?	?	?	+	?
Similar groups at baseline in terms of the most important prognostic indicators	+	+	+	?	+	+	+	+
Blinded subjects	-	_	_	_	-	_	+	_
Blinded therapist	-	-	-	-	+	+	-	-
Blinded evaluators	+	-	-	+	-	-	+	-
Adequate follow-up	+	+	+	+	+	+	+	?
Intention of treatment	+	+	+	+	+	+	+	+
Comparison between groups	+	+	+	+	+	+	+	+
Estimated points and variability	+	+	+	+	+	+	+	+
Total score	9	8	6	7	8	8	10	6

 Table I Methodological Quality Of The Included Studies Evaluated With The PEDro Scale

Notes: +The criteria are clearly satisfactory; -The criteria are clearly not satisfactory; ?The criteria are unclear.

participants received radical retropubic prostatectomy, and in 2 studies the participants received radical prostatectomy. Included studies assessed the following interventions: preoperative vs postoperative PFMT (2 studies), supervised vs home-based PFMT (3 studies), unsupervised PFMT vs no treatment at all (2 studies), and PFMT combined with resistance and flexibility exercises vsPFMT alone (1 study).

The Study Groups Were Also

Characterized Using The Following Criteria Body Mass Index (BMI)

Patient BMI was assessed in four studies.^{8,12–14} In all these studies, BMI results suggested that patients are overweight. Obesity is a risk factor for prostate cancer progression, and it may worsen the treatment results. However, there is little research to be found on the relationship between obesity and UI in male patients after radical prostatectomy. Wei et al.¹⁶ carried out a meta-analysis of studies on the relationship between the BMI and urinary incontinence in men after radical prostatectomy. The findings showed that obesity may be a urinary incontinence risk factor in male patients 12 and 24 months after robotic-assisted laparoscopic radical prostatectomy (RLRP). Nevertheless, no significant association was observed

between obesity and urinary incontinence in male patients 24 months after laparoscopic radical prostatectomy (LRP).

Clinical T-Stage

The stage of prostate cancer was determined in seven studies.^{7,9–14} The American Joint Committee on Cancer TNM system (T – tumor; N – nodes; M – metastasized) is a standard tool used by healthcare professionals for pathological classification of tumors, commonly used to define prostate cancer stage. The TNM most recent update was is in January 2018.¹⁷

The Gleason Score

The aggressiveness of prostate cancer was determined in five studies.^{8,12–15} This grading system may be used to choose appropriate treatment options. The Gleason Score ranges from 1 to 5, and it describes how closely a biopsy sample resembles healthy tissue (lower score) or abnormal tissue (higher score).¹⁸

Prostate-Specific Antigen (PSA)

Patient PSA was determined in seven studies.^{8,12–15} Prostate-specific antigen is a glycoprotein produced in the prostate gland. Elevated PSA concentration in blood is an indicator of prostatic hypertrophy or prostate cancer. The PSA antigen is considered a sensitive marker for detecting and treating prostate cancer. PSA concentration up to 4 ng/mL is regarded as normal.¹⁹

Outcome Measures

The following diagnostic tests were used in the research studies covered in this systematic review:

Muscle Strength

In their studies, Nilssen et al assessed muscle strength during a palpation exam.¹² Whereas Overgard et al assessed pelvic floor muscle strength using anal pressures during pelvic floor muscle contractions.¹³ For this purpose, the authors used a balloon catheter attached to a pressure transducer.

Urodynamic Examination, Pad Test, And Bladder Diary

The 24-hr pad test was conducted in subjects from seven studies,^{8–11,13–15} and the 1-hr pad test was performed only in one study.¹⁰ The 3-day bladder diary was used in three studies.^{8,10,14} Subjects underwent urodynamic examination in two studies.^{9,10}

Health-Related Quality Of Life (HRQOL)

The following tools were used to assess the HRQOL: University of California, Los Angeles Prostate Cancer Index;¹²

Short Form-12¹² International Continence Society Male Short Form;^{8,14} Medical Outcomes Study 36-item Short Form;¹⁴ and the Health-Related Quality of Life Questionnaire.¹²

Other Questionnaires

In their studies, Centemero et al examined patients' cognitive abilities using the Mini Mental State Examination.⁸ Park et al used the Beck Depression Inventory to assess the presence of depressive syndrome symptoms.¹⁴ Whereas Manaserro et al used the Visual Analogue Scale in their studies.¹¹

The Experimental Group Versus The Control Group

In the studies of Filocamo et al.¹⁰ and Manaserro et al.¹¹ the experimental groups received PFMT, whereas the control groups did not receive any therapeutic intervention. In both of these studies, better results were observed in the experimental group. Five studies^{8,9,12,13} compared physiotherapist-guided PFMT (EG) with unsupervised PFMT performed after receiving verbal training instructions (CG). In all referenced studies, better results were observed in patients that underwent physiotherapist-guided PFMT.

In the studies of Patel et al.¹⁵ the experimental group started physiotherapist-guided PFMT 4 weeks prior to the surgery, and the control group received verbal training instructions on PFMT from a surgeon. All patients were offered physiotherapist-guided PFMT postoperatively. The study results demonstrated that a physiotherapist-guided PFMT started 4 weeks prior to the surgery significantly reduced urinary incontinence duration and severity. Whereas Park et al.¹⁴ compared a combined exercise intervention (resistance, flexibility, PFMT) (EG) with PFMT alone (CG). No statistically significant differences in UI severity between the groups were observed during the basic assessment.

Treatment Start Point

In two studies,^{8,15} the experimental groups started a physiotherapist-guided PFMT a month prior to the surgery and continued treatment after the surgery. Whereas the control group patients received verbal training instructions on PFMT. The authors of these studies suggest that preoperative PFMT may improve incontinence and quality of life outcomes after RP.

Treatment Duration

In the analyzed studies, the treatment lasted for 1–12 months. Furthermore, five studies analyzed the treatment efficacy at different time intervals. In their studies, Centemero et al.⁸ demonstrated that patients who underwent preoperative PFMT had a 0.41-fold lower risk of being incontinent 1 month after RP and a 0.38-fold lower risk of being incontinent 3 months after RP. In the studies of Patel et al.¹⁵ the primary outcome measure was a 24-hr pad weight test conducted at 6 weeks and 3 months postoperatively. Overgard et al.¹³ assessed treatment efficacy in two intervention groups at 6 weeks, 3, 6, and 12 months after the surgery. Both groups received training instructions on PFMT. However, one of the groups was offered additional follow-up training instructions from a physiotherapist during a year. No statistically significant differences in continence status between the groups were found at 3 months postoperatively; however, at 12 months postoperatively, better results were observed in the group that was offered additional follow-up training instructions. Filocamo et al.¹⁰ assessed results of the experimental group after PFMT and the control group that was not offered any therapeutic intervention. In the EG, 19% of patients achieved continence after 1 month, and 94.6% after 6 months. In the CG, 8% of patients achieved continence after 1 month, and 65% after 6 months. In their studies, Manassero et al.¹¹

rials Included In The Review	Age Of Sample Time Of Type Of Intervention And Outcome Measures Results Participants Size Interventions Surgery Study Design (yr)	46-68 N=118; 1-month pre-RP; RP EG - preoperative and BMI; s-PSA; clinical Preoperative PFMT improved EG=59; 1-month post-RP 2 1-month post-RP 7 stage; Gleason score; early continence and QoL CG=59 CG=59 2 4.rs pad test; 3d Bladder Dlary; outcomes after RP Anoth 30 mins and 30 mins ICS male SF; MMSE; assessment in at home, maximal and lub lub spot-RP 1,3 months post-RP Anoth EG - postoperative PFMT I.3 months post-RP early continence and QoL Anoth EG - postoperative I.3 months post-RP early continence and QoL Anoth I.3 months post-RP bub submaximal contrib; euthomes after RP Bub submaximal contrib; CG - postoperative PMT early continence and QoL Bub submaximal contrib; Bub submaximal contrib; early contrib early continence and QoL Bub submaximal contrib; Bub submaximal contrib; Bub submaximal contrib; Bub submaximal contrib; Bub submaximal contrib; Bub submaximal contrib; Bub submaximal contrib; Bub submaximal contrib; Bub submaximal contrib; Bub submaximal contrib; Bub submaximal contrib; Bub su	60-67 N=66; 26 weeks post-RP RRP EG - PFMT 24-hr pad test; UD; DOA More intensive PFMT might have a nds EG=33; (supervised, 2x week, assessment 1,2,3,6,12 months lowering effect on bladder outflow CG=33 CG=33 30 mins); post-RP resistance after RRP CG unsupervised PFMT PFMT PFMT	45-75 N=300: 12 months post-RP RRP EG-FMT 1 hr and 24 hrs pad test; UD; After RRP an early supportive EG=150: EG=150: (unsupervised, at bladder diary: rehabilitation program like PFMT CG=150 A. Nome, 10 contr 5 sec s-PSA; clinical T stage; assessment significantly reduces continence 3X daily in sitting, 1,3,6,12 months post-RP recovery time etanding, and squatting position and going up and down staris); cG- no treatment staris);	67.3 (range N=94; 3 months post-RP RRP EG - PFMT 24-hr pad test; s-PSA; clinical T stage; The early intensive prolonged not reported) EG=54; (unsupervised, at VAS; PFMT can further increase the not reported) EG=40 a single question of QoL; assessment number of continent patients, and CG=40 CG=40 1,3,6,12 months post-RP this improvement carries on in the sitting, standing 1,3,6,12 months post-RP first 12 months.
Included In The Review	Age OfSampleTime OfParticipantsSizeInterventions(yr)	46–68 N=I I8; I-month pre-RP, EG=59; I-month post-RP CG=59	60–67 N=66; 26 weeks post-RP EG=33; CG=33	45–75 N=300; I2 months post-RI EG=I50; CG=I50	67.3 (range N=94; 3 months post-RP not reported) EG=54; CG=40
able 2 Characteristics Of Trials	Authors/Year Country	Centemero A, et al, Italy 2010 ⁸	Dubbelman et al, The 2012 ⁹ Netherlands	Filocamo MT, et al, Italy 2005 ¹⁰	Manassero F, et al. Italy 2007''

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Nilssen SR, et al, 2012 ¹²	Norway	48-72	N=85; EG=42; CG=43	12 months post-RP	RP	EG - PFMT (supervised, 1x week 45 mins) 3x 10 contr. daily x 6-8 s and last 3-4 fast contr. at home in supine, sitting, and standing position; CG - unsupervised PFMT	BMI; s-PSA; clinical T stage; Gleason score; digital and palpation exams to assess muscle strength; HRQoL (urinary, sexual and mental function); UCLA-PCI; SF-12: assessment in 6 weeks, 3,6,12 months post-RP	PFMT after RP improved postoperative urinary incontinence significantly compared to those patients receiving standard care/training, this was not reflected in better outcome in HRQoL parameters
Overgard M, et al, 2008 ¹³	VSN	>65	N=51; EG=26 CG=25	12 weeks post-RP	RP	EG - (resistance, flexibility and PFMT; 2x week 60 mins); CG - PFMT	BMI; s-PSA; clinical T stage; Gleason score; 24-hr pad test; 3d Bladder Diary; ICS male SF; MMSE; ICIQ, Beck Depression Inventory, Study 36-item short-form health survey (SF - 36); assessment in 1 month before and 3, 15 weeks post-RP	Exercise intervention improves the continence rate and the quality of life after RP. These exercise intervention effects could contribute to achieving prompt recovery of daily living
Park SW, et al, 2012 ¹⁴	ASU	> 65	N=51; EG=26 CG=25	12 weeks post-RP	Ч	EG - (resistance, flexibility and PFMT; 2x week 60 mins); CG - PFMT	BMI; s-PSA; clinical T stage; Gleason score; 24-hr pad test; 3 d Bladder Diary; ICS male SF; MMSE; ICIQ, Beck Depression Inventory, Study 36-item short-form health survey (SF - 36); assessment in 1 month before and 3, 15 weeks post-RP	Exercise intervention improves the continence rate and the quality of life after RP. These exercise intervention effects could contribute to achieving prompt recovery of daily living
Patel MI, et al, 2013 ¹⁵	Australia		N=284; EG=152; CG=132	4 weeks pre-RP; 3 months post-RP	RP	EG - preoperative and postoperative PFMT (supervised, 10 contr 10 s in each sitting, standing, and lying position); CG - postoperative unsupervised PFMT;	24-hr pad test; 6 weeks and 3 months post-RP, s-PSA; clinical T stage; Gleason score	Postoperative FFMT commenced 4 weeks before RRP has a clear advantage in improving both the severity and duration of PPUI

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assessed the EG (PFMT treatment) and CG (no treatment) results at 1, 3, 6, and 12 months postoperatively. At each time point, better results were reported for all measured variables for the EG.

Pelvic Floor Muscle Training Regimen

The PFMT regimens vary considerably across the analyzed studies. Significant differences were observed between each PFMT regimen when it comes to contraction number and duration as well as the PFMT session frequency per day. Pelvic floor muscle training regimens are summarized in Table 2.

Filocamo et al.¹⁰ instructed the patients to perform PFMT exercises 3 times a day (10 contractions lasting 5 s with 10 s of muscular relaxation). The PFMT regimen of Manassero et al.¹¹ comprised a series of 15 contractions repeated 3 times a day, whereas Patel et al proposed¹⁵ 10 contractions lasting 10 s. Additionally, in the two latter studies, the PFMT treatment was divided into a training of slow-twitch fibers and a training of fast-twitch fibers. Nilssen et al.¹² and Overgard et al.¹³ instructed their patients to perform additional 3-4 fast contractions following each PFMT session (10 contractions of 6-8 s). Moreover, in most of the studies, PFMT was performed in supine, sitting, and standing positions. Dubbelman et al.⁹ and Park et al.¹⁴ described PFMT treatment frequency (twice a week) and session duration (30-60 mins) in the methodology sections. Whereas Centemero et al.⁸ also provided additional information on contraction types (maximal and submaximal) but did not include their number.

Limitations

Methodological differences and a small amount of research.

Conclusion

Findings of this review highlighted that PFMT is an effective treatment for UI in men after radical prostatectomy. PFMT improves not only physical parameters but also the quality of life of men after RP. There is a need for further research to determine the exact procedure for pelvic floor muscle exercise in men after radical prostatectomy.

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Author contributions

All authors contributed toward data analysis, drafting and critically revising the paper. The authors gave final approval of the version to be published and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

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