Change in Peak Systolic Velocity of Cavernous Artery in Response to Low Intensity Shock Waves Therapy in Diabetic Polyneuropathy Patients with Erectile Dysfunction: A Randomized Controlled Trial

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Received: 06/01/2020 Accepted: 13/04/2020 Published: 20/09/2020

Abstract

Although several reports have documented the subjective improvement of erectile function after low-intensity extracorporeal shockwave therapy (LI-ESWT) in patients with vasculogenic erectile dysfunction (ED), objective assessment data of penile hemodynamics are lacking. The aim of this research is to detect the Change in peak systolic velocity of cavernous artery in response to low intensity shock waves therapy in diabetic patients with erectile dysfunction. This study is a randomized controlled trial. Overall forty male individuals with mean age (48.6±5.52 years) who have erectile dysfunction after diabetes were equally divided into two separate groups. 1st Group received low intensity extracorporeal shock wave therapy (ESWT). 2nd Group received sham therapy by put head of shock wave while machine is turned off. 1st Group received low intensity extracorporeal shock wave with the following parameters: - 3000 SWs (energy intensity of 0.09mJ/mm2) to each of five different sites of application: three areas along the penile shaft and two areas at the crural level for six weeks as a total period of treatment. Measurements of peak systolic velocity by penile duplex were performed before the treatment and after 3 months. There were change in peak systolic velocity of the right and left cavernous arteries were low at baseline, indicating arterial insufficiency. After treatment, PSV significantly in the two groups; however, the post-treatment PSV was significantly higher in the SW group compared to the PFE group (p < 0.001, for both arteries). LIESWT is lead to significant changes in patients suffering from erectile dysfunction after diabetes as evidenced by increase in the peak systolic volume of the both right and left cavernous arteries of the shock wave group.

Keywords: Peak systolic velocity, Low intensity shock wave therapy, Erectile Dysfunction and Diabetic patients

1 Introduction

Erectile dysfunction (ED) is defined as the continuous inability to obtain and maintain an erection sufficient to allow satisfactory sexual intercourse. Although ED is a benign disorder, it may affect physical and psychosocial health and may have a significant impact on the quality of life (QoL) of sufferers and their partners [1]. According to the underlying causes, ED can be classified as psychological, endocrinological, neurological, and vascular. Vascular erectile dysfunction (ED) is defined as inability to obtain or keep an erection firm enough for sexual performance due to diseases such as diabetes mellitus and atherosclerotic vascular occlusive disease [2]. The onset of ED usually occurred within the first 10 years of diagnosis of type 1 and type 2 diabetes in > 50% of men affected by ED [3]. The etiology of ED in diabetic patients can be vascular or neurogenic and any disruption of the primary hemodynamic events controlled by the central or peripheral neural networks that promote an erectile response will impair erectile function so both the endothelium-dependent and neural mechanisms that mediate relaxation of the smooth muscle of the corpus cavernosa impaired in diabetic patients with ED [4]. ED is one of the most frequent complications of diabetes mellitus (DM), which leads to a marked decrease in the quality of life and is usually treated is difficult because of a combination of microangiopathy and peripheral diabetic neuropathy, which adversely affect the mechanism of erection [5]. ED in men with diabetes has also been associated with increased age, poor glycemic control, smoking, increased alcohol intake, depression, use of specific type of medications and micro-vascular diabetic complications [6]. It has been proven that ED and coronary artery disease (CAD) share pathways. Endothelial dysfunction is one of the basic mechanisms has effect in the pathophysiology of vasculogenic ED, which develops as a result of decrease in the synthesis and bioavailability of nitric oxide (NO) and subsequent atherosclerosis. Atherosclerosis leads to an impairment of the blood flow required for normal erection [7].
progressive impotence related directly to diabetes is the most frequent form. This form of impotence occurs years after the diagnosis of the patient. This form is progressive, irreversible and is accepted as a natural result (neuropathic, vascular or both) of the diabetes [8]. Patients with diabetic and neuropathic ED have been noted to have similar frequencies of somatic and autonomic neuropathies, suggesting that neuropathy contributes significantly to diabetic ED [9]. Low-intensity extracorporeal SW therapy (Li-ESWT) was used both in vitro and in vivo studies and the results show that shock wave energy can stimulate angiogenesis. The idea of application of Li-ESWT to the penis came from animal studies in which Li-ESWT was applied to the myocardium of pigs, where it has been found that there was an improvement in ischemia that induced myocardial dysfunction [10]. Low-intensity extracorporeal SW therapy (LIESWT) of the penis would improve penile blood flow and endothelial function by stimulating angiogenesis in the corpora [11]. Consequently, it will be of importance to do a randomized controlled trial using valid and reliable measures to investigate the effect of Li-ESWT in treatment of erectile function in diabetic polyneuropathic patients.

2 Materials and Methods
2.1 Design of the study
The study was designed as a prospective, randomized, controlled trial. It was performed between July 2018 and January 2020.

2.2 Participants
The study was carried out on forty outpatient male individuals with mean age (48.6±5.52 years) who have erectile dysfunction after diabetes. Their age was ranging from 35 to 60 years. They were selected from South valley University hospitals. The body mass index (BMI) of those individuals not more than 32 they were diagnosed with type 2 diabetes with secondary complication of erectile dysfunction. These individuals reported no other pathological conditions lead to erectile dysfunction. They showed the necessary cooperation needed to enable the investigator to secure the required data. Patients were equally divided into two separate groups. 1st Group (the study group) received the low intensity extracorporeal shock wave therapy (ESWT). 2nd Group (control group) received only shame therapy.

2.3 Inclusive criteria
All patients had the following characteristics; their ages ranged between 35 and 60 years, they had the body mass index (BMI) of those individuals not more than 32 they were diagnosed with type 2 diabetes with secondary complication of erectile dysfunction was confirmed by using penile duplex. All patients take their medications described by their andrologist physicians.

2.4 Exclusive criteria
Patients with a history of past radical prostatectomy or extensive pelvic surgery, recovering from cancer in the past 5 years, any unstable medical, psychiatric disorder, spinal cord injury and another neurological disease, penile anatomical abnormalities, clinically significant chronic hematological disease, patients with untreated hypogonadism and cardiovascular conditions that prevent sexual activity (heart attack, stroke or life-threatening arrhythmia within the previous 6 months were excluded.

2.5 Randomization
Every individual was informed about the nature, purpose, benefits of the research and their right to withdraw or refuse at any time. The patients were randomly assigned into 2 equal groups (control group and study group) with the use of a computer-based randomization program. No subject dropped out from the study after randomization. The patients were blinded about which group they were allocated.

2.6 Ethical approval
The research has complied with all the relevant national regulations and institutional policies and has followed the tenets of the Declaration of Helsinki and the Consolidated Standards of Reporting Trials and has been approved by the institutional review board at Faculty of Physical Therapy, Cairo University. No.P.T.REC/012/001983.

2.7 Procedures
2.7.1 Assessment device
The equipment used in the measurement on this research was the Penile color-coded duplex scanning machine (ultrasound with 8.4 MHz linear vascular probe (VF 15-3 tradsucer, ACSON X 300 SIEMENS ultrasound) that used to get an unbiased assessment of the penile haemodynamics.

2.7.2 Treatment devices
The therapeutic equipment and tools used on this research were Shock wave device; ESWT device (MASTERPULS MP200, StorzMedical, Tägerwilen, Switzerland).

3.3 Procedures
3.3.1 Assessment procedures
Penile color-coded duplex scanning (Color-coded Duplex scanner Transducer: 5-10 MHz)
- Patient’s position: supine
- Probe position: placed on the back of the penis
- Transversal scanning to assess the echo structure and dimensions of the cavernous bodies.
- Longitudinal scanning to identify the cavernous arteries, their patency and their hemodynamic through these parameters. (I) Peak systolic volume for right cavernous artery and (II) peak systolic volume for left cavernous artery.

Treatment Procedure: The 1st group received low intensity shock wave while the 2nd group received shame therapy. Shock wave therapy application: The patient was placed in supine lying position receive low intensity extracorporeal shock wave with the following parameters: - 3000 SWs (energy intensity of 0.09 ml/mm2) to each of five different areas: three along the shaft of penis and two at the crural level [12]. The protocol consists of 6 treatment sessions, once per week. The shock waves delivered through the applicator covering the corpora cavernosa of the penis along the penile shaft and the crura. Before the experiment, recording of the peak systolic velocity of right and left cavernous arteries for every individual were done in both groups. Treatment of ESWT application was applied one session every week for six weeks after which the second (final) penile duplex parameter (PSV) recording was taken. The pre and post experimental measurements were performed during the same time of the day to decrease the variability. The step-by-step procedure for data recording was identical to that followed during pre-experimental measurement of the penile duplex parameter (peak systolic velocity) taken before the treatment as an initial record and then after six weeks as a second last record in the two groups.
3.3.2 Statistical analysis

Data collected were admitted into a personal computer to analyze the statistics; the statistical analysis was conducted by using t-test to compare the patient characteristics of both groups. The Shapiro-Wilk test revealed that the data were normally distributed for all dependant variables. Levene’s test was conducted to test the homogeneity between groups. To compare the mean values of peak systolic velocity between both groups and before and after the treatment in both groups use paired t test. Level of significance for all statistical tests was set at P < 0.05. with using the statistical package for social studies (SPSS) version 23 and window (IBM SPSS, CHICAGO IL USA) for data analysis.

3.3.3 Informed consent

Informed consent has been obtained from all individuals included in this study.

4 Results

4.1 Demographic characteristics

Table 1 showed demographic characteristics of both groups. There was no significant difference between both groups in the mean age, weight, height and body mass index (p<0.05). The mean values of peak systolic velocity for right and left cavernous arteries showed a statistically significant improvement (p < 0.05) With in both groups (A and B). The post-treatment comparison of both groups revealed a statistically significant increase in the mean values of peak systolic velocity for right and cavernous arteries in favor of group (A) (Table 2).

5 Discussion

This study demonstrated the positive effect of Li-ESWT on erectile function in patients with ED caused by diabetic polyneuropathy three months after treatment. Li-ESWT induced a significant increase of penile hemodynamics, evidenced by a significant rise of PSV. Generally, the treatment was safe and well-tolerated, with no serious adverse events. In the current study, all patients were confirmed to have diabetic polyneuropathy. Doppler assessment before treatment indicated arterial insufficiency (a PSV < 25 cm/s) in less than half of the patients. This observation indicates that neurogenic, rather than a vasculogenic element, is operating as a cause of ED in the current series. In fact, DM-related ED is a multifactorial process that involves vascular and neurogenic causes in addition to endothelial dysfunction and nitric-oxide system disturbance [13]. However, diabetic neuropathy is believed to be a major contributing factor in the mechanism of ED in patients with type-2 DM [14].

<table>
<thead>
<tr>
<th>Study group</th>
<th>Control group</th>
<th>t-value</th>
<th>p-value</th>
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<tbody>
<tr>
<td>Age (years)</td>
<td>48.6±5.52</td>
<td>47.5±5.68</td>
<td>0.56</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>84.5±8.98</td>
<td>85.4±8.89</td>
<td>0.080</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>167.5±3.58</td>
<td>169.8±6.73</td>
<td>0.814</td>
</tr>
<tr>
<td>Body mass index</td>
<td>25.2±2.46</td>
<td>26.2±2.46</td>
<td>0.232</td>
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<table>
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<th>Pre treatment</th>
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<tr>
<td>Study group</td>
<td>Control group</td>
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<tr>
<td>PSV OF RT cavernous artery cm/sec</td>
<td>26.58±6.94</td>
</tr>
<tr>
<td>PSV OF LT cavernous artery cm/sec</td>
<td>26.59±6.65</td>
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*: Mean, SD: Standard deviation; p-value, Level of significance.
The neurogenic element in DM exerts its effect by inefficient nerve signaling to the corpora cavernosa. This functional alteration results in the reduction in nitric oxide (NO) load in the smooth muscles [15]. The mechanism of action of Li-ESWT is still hypothetical. Physically, shockwaves exert two effects, mechanical stress due to exposure to high-pressure waves and cavitations bubbles formed in liquids. These bubbles result from the vaporization of the liquid. Consequently, these cavities collapse when exposed to high-pressure causing local trauma and neovascularization[16]. In vitro and in vivo studies confirmed the process of neovascularization in response to shockwave, which is believed to be a principal therapeutic mechanism in the treatment of ED[17]. Other mechanisms are hypothesized, including NO induction [18], nerve regeneration, and stem cell proliferation[19]. In diabetic rat model, Li-ESWT was found to affect penile neural tissue with the enhancement of neuronal NO synthase positive cells[20].

The present study was performed in an animal model. Shock wave therapy enhanced nerve stimulated erection in diabetic rats, increase the smooth muscle- collagen ratio, increased the endothelial content of penile tissue and up-regulated the expression of growth factors, Qiu, X et al. [25]. Found that LiESWT are good for penile tissue regenerating by mesenchymal stem cells activation and nerve regeneration (via Schwann cells activation), and vessels, with the consequent release of pro-angiogenetic growth factors. In addition, LIESWT also enhance erectile function via nitric oxide/cGMP-nondependent mechanisms.Rizk et al. [26] included randomized controlled trials, meta-analyses, and select single-arm studies on the use of Li-ESWT in the treatment of at least mild ED, with some data supporting efficacy in moderate-to-severe ED also demonstrated some benefit in specific subsets of men with vasculogenic ED (including patients with DM). Kalyvianakis et al. [27] assessed the efficacy and safety of 6- and 12-treatment sessions within a 6-week treatment period and also investigated the effect of repeat treatment after a 6-month period in a 2-phase study in patients with vasculogenic ED. The results demonstrated that re-treating patients after 6 Months could further improve EF without side effects. In addition, it was demonstrated that 12 sessions can be delivered within 6 weeks without a 3-week break period with similar clinical outcome. After discussion of the results and according to reports of the previous investigators in fields related to this study, it can be claimed that the application of Li-ESWT in erectile dysfunction in diabetic polyneuropathy patients increase peak systolic velocities of right and left cavernous arteries which result in improved erectile functions.

6 Limitations
The study was limited by emotional state of the patients, and the psychological condition of the patients at the time of performance which might affect the results, also, other limitation are small sample size and possible errors in measuring penile haemodynamics. So, more extensive studies assigning the efficacy of ESWT on the erectile function in diabetic polyneuropathy patients with larger sample are needed. Follow-up studies would be of great interest to detect the long-term effect of ESWT and the recurrence of erectile dysfunction.

7 Conclusion
Application of ESWT in erectile dysfunction in diabetic patients improve peak systolic velocity for both right and left cavernous arteries. This study demonstrated that using the ESWT is beneficial in treating diabetic patients suffering from erectile dysfunction by improving peak systolic velocity in right and left cavernous arteries.

Acknowledgment
We would like to represent our gratitude to our patients for participation in this study.

Ethical issue
Authors are aware of, and comply with, best practice in publication ethics specifically with regard to authorship (avoiadance of guest authorship), dual submission, manipulation of figures, competing interests and compliance with policies on research ethics. Authors adhere to publication requirements that submitted work is original and has not been published elsewhere in any language.

Competing interests
The present study was performed in absence of any conflict of interest which was declared by the authors.

Authors’ contribution
AA, WS, ME, OE and MS conceived of the study, designed the study protocol and drafted the manuscript. WE are the corresponding author and supervisor of the research. AA helped us in drafting the revised manuscript and substantively helped us to revise the manuscript. All authors have reviewed the final version of the manuscript and approve it for publication.

Disclosure statement
No author has any financial interest or received any financial benefit from this research.

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