# Systematic Review and Meta-Analysis of 16 Randomized Controlled Trials of Clinical Outcomes of Low-Intensity Extracorporeal Shock Wave Therapy in Treating Erectile Dysfunction

American Journal of Men's Health March-April 1–13 © The Author(s) 2022 Article reuse guidelines: sagepub.com/journals-permissions DOI: 10.1177/15579883221087532 journals.sagepub.com/home/jmh SAGE

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#### Abstract

We conducted a meta-analysis to evaluate the efficacy of low-intensity extracorporeal shock wave therapy (LI-ESWT) in the treatment of erectile dysfunction (ED). From July 2011 to June 2021, we finally selected 16 randomized controlled trials (RCTs) including 1,064 participants to evaluate the efficacy of LI-ESWT in the treatment of ED from PubMed, EMBASE, and Cochrane databases. The data are analyzed by Review Manager Version 5.4. Fifteen articles mentioned International Index of Erectile Function (IIEF), in the follow-up of I month (mean difference [MD] = 3.18, 95% confidence interval [CI] = [1.38, 4.98], p = .0005), 3 months (MD = 3.01, 95% CI = [2.04, 3.98], p < .00001), and 6 months (MD = 3.20, 95% CI = [2.49, 3.92], p < .00001). After treatment, the improvement of IIEF in the LI-ESWT group was better than that in the control group. Besides, eight of the I6 trials provided data on the proportion of patients with baseline Erectile Hardness Score (EHS)  $\leq$  2 improved to EHS  $\geq$  3. The LI-ESWT group was also significantly better than the placebo group (odds ratio [OR] = 5.07, 95% CI = [1.78, 14.44], p = .002). The positive response rate of Questions 2 and 3 of the Sexual Encounter Profile (SEP) was not statistically significant (SEP2: OR = 1.27, 95% CI = [0.70, 2.30], p = .43; SEP3: OR = 4.24, 95% CI = [0.67, 26.83], p = .13). The results of this meta-analysis suggest that treatment plans with an energy density of 0.09 mJ/mm<sup>2</sup> and pulses number of 1,500 to 2,000 are more beneficial to IIEF in ED patients. In addition, IIEF improvement was more pronounced in patients with moderate ED after extracorporeal shockwave therapy.

#### **Keywords**

low-intensity extracorporeal shockwave therapy, erectile dysfunction, randomized controlled trials, meta-analysis, International Index of Erectile Function, Erectile Hardness Score

Received November 24, 2021; revised January 23, 2022; accepted February 25, 2022

# Introduction

Erectile dysfunction (ED) refers to the inability of the penis to continuously achieve or maintain sufficient erection to meet a satisfactory sexual life (Burnett et al., 2018). The incidence rate of ED is increasing and the prevalence of ED is 30% to 65% among men aged 40 to 80 years (Ayta et al., 1999; Corona et al., 2010). The most common clinical treatment for ED is oral phosphodiesterase 5 inhibitor (PDE5I; Hatzimouratidis et al., 2010). However, some literatures reported that when PDE5I is used to treat ED patients, some patients say it is

ineffective, and some patients will have various side effects, such as flushing and headache (Hatzimouratidis et al., 2010; Washington & Shindel, 2010).

Extracorporeal shock wave (ESW) is a two-way sound wave carrying energy. According to the different energy density levels of ESW, ESW has different functions in clinical application (Rassweiler et al., 2011). High energy density ESW has focused on mechanical damage characteristics, so it is often used in the treatment of stones. Medium energy density ESW has anti-inflammatory function and it is often used in surgery, such as synovial bursitis and nonbinding fracture. Low energy density

Creative Commons Non Commercial CC BY-NC: This article is distributed under the terms of the Creative Commons Attribution-NonCommercial 4.0 License (https://creativecommons.org/licenses/by-nc/4.0/) which permits non-commercial use, reproduction and distribution of the work without further permission provided the original work is attributed as specified on the SAGE and Open Access pages (https://us.sagepub.com/en-us/nam/open-access-at-sage). ESW can promote angiogenesis and improve its blood supply and it is often used in chronic injury, musculoskeletal recovery, and cardiovascular disease (Nishida et al., 2004; Vardi et al., 2010; Wang et al., 2002). Studies have reported that the important mechanism of ED is vascular endothelial function injury or disorder (Gandaglia et al., 2014; Shindel et al., 2008) and low-intensity extracorporeal shock wave therapy (LI-ESWT) can stimulate the expression of angiogenesis-related factors, such as vascular endothelial growth factor (VEGF), so as to promote vascular regeneration (Cooper & Bachoo, 2018; Klomjit et al., 2020; Sundaram et al., 2018). As a result, LI-ESWT has been widely used in clinical treatment of ED (Rizk et al., 2018).

We carried out a meta-analysis of randomized controlled trials (RCTs) to systematically evaluate the efficacy of LI-ESWT in the treatment of ED.

# **Materials and Methods**

#### Search Strategy

Under the guidelines of Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA; Moher et al., 2009), we searched three databases, namely, PubMed, Embase, and Cochrane, by computer and the retrieval time was limited from July 2011 to June 2021. The retrieval strategy is to search for the following Medical Subject Headings (MeSH) terms: shock wave, erectile dysfunction, the international index of erectile function, and the erection hardness scores. In addition, the researcher also traced all references involved in the included literature to supplement and obtain relevant literature. This study only included published articles and had no restrictions on the language of the articles. All articles were read independently by two researchers. In case of disagreement, an agreement will be reached through discussion or inviting the assistance of a third researcher.

#### Inclusion Criteria and Article Selection

Included articles should meet the following inclusion criteria: (a) all RCTs describing LI-ESWT treatment for ED; (b) the content and data of any article are available; (c) all the data in the paper are true and valid; (d) no matter whether the test adopts blind method and allocation concealment or not; (e) no matter whether the patients are complicated with other complications; and (f) the severity of ED patients is not limited. Case reports, review articles, conference reports and abstracts, and some studies with incomplete data were excluded. The PRISMA flowchart of literature screening is presented in Figure 1.

#### Quality Assessment

We mainly used Cochrane bias risk assessment tool to evaluate all randomized controlled studies, supplemented by Jadad scale for reference (Cumpston et al., 2019; Moher et al., 1996). Each article was evaluated according to the following three quality evaluation criteria: (+) bias is low, (?) not mentioned or no sufficient information to judge bias, and (-) bias is high. All authors independently participated in the evaluation of each RCT and exchanged results. If there is any objection, it will be resolved through discussion and negotiation until all the results are consistent.

# Data Extraction

The two authors extracted data from the included studies according to the predetermined criteria independently, and recorded the data on the premade data extraction table. The extracted data include (a) author's name (publication time), (b) country, (c) number of participants, (d) age, (e) PDE5I response or not, (f) treatment setup, (g) control group setup, (h) follow-up time, and (i) outcome indicators. This study does not need ethical approval because it is a retrospective analysis of existing studies.

#### Statistical Analyses

This study uses Review Manager Version 5.4 (Cochrane Collaboration, London, UK) for data analysis. We use fixed effect model or random effect model for analysis. The dichotomous data are expressed in odds ratios (ORs) and 95% confidence interval (CIs), whereas the

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Figure 1. Flowchart of Selection PRISMA

Note. RCTs = randomized controlled trials; LI-ESWT = low-intensity extracorporeal shock wave therapy; ED = erectile dysfunction.

continuous outcomes are expressed in mean difference (MD) and 95% CI. We usually used *I*-square ( $I^2$ ) to assess the heterogeneity of the study. If the *p* value is greater than .05 in the *Q*-value statistic test and the  $I^2$  value is less than 50%, we believe that the study is homogeneous and can be analyzed by the fixed effect model. While the results with  $I^2$  test value are greater than 50% and significant heterogeneity, the random effect model is used for analysis. Results of this meta-analysis are presented in forest plots and the data with p < .05 were considered statistically significant.

# Results

# Study Selection, Search Results, and Characteristics of the Trials

We searched according to the above search strategy and finally retrieve 318 articles. A total of 279 articles were excluded by deleting duplicate literature and screening abstracts and titles. Of the remaining 39 articles, 23 articles were excluded because they are not RCT or lack of effective data. Finally, the remaining 16 articles were included in the study to evaluate the effectiveness of LI-ESWT in the treatment of ED (Baccaglini et al., 2020; Fojecki et al., 2017; Kalyvianakis & Hatzichristou, 2017; Kim et al., 2020; Kitrey et al., 2016; Ladegaard et al., 2021; Olsen et al., 2015; Ortac et al., 2021; Shendy et al., 2021; Sramkova et al., 2020; Srini et al., 2015; Vardi et al., 2012; Vinay et al., 2021; Yamaçake et al., 2019; Yee et al., 2014; Zewin et al., 2018). The details of each study are presented in Table 1. There was no significant difference in mean age and severity of ED between the LI-ESWT group and the placebo group.

# **Risk of Bias**

All included studies in meta-analysis were RCTs. The summary and graph of bias risk are presented in Figure 2 and Supplemental Figure S1.

Authors' name	Year	Country	No. of participants	Age	PDE5I- response or not	LI-ESWT setup	Control setup	Methodology	Follow-up time	Outcome indicators
Baccaglini et al.	2019	Brazil	77	64.6 ± 5.3	NA	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Frequency: 5 Hz</li> <li>Pulse: 600 pulses/treatment</li> <li>RENOVA® electromagnetic device</li> </ul>	A	<ul> <li>One treatment/week</li> <li>8 weeks treatment</li> </ul>	• Though	• IIEF variation scores
Fojecki et al.	2016	Denmark	81	<b>64.4</b> ± <b>8.4</b>	Yes	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Frequency: 5 Hz</li> <li>Pulse: 600 pulses/treatment</li> <li>FBL10, Richard-Wolf GmbH</li> </ul>	With a gel pad that prevent the passage of energy	<ul> <li>One treatment/week</li> <li>5 weeks treatment</li> </ul>	• month	<ul> <li>EHS response rate</li> <li>IIEF variation scores</li> </ul>
Kalyvianakis et al.	2017	Greece	46	54 (31–72)	Yes	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Frequency: 2 Hz</li> <li>Pulse: 1,500 pulses/treatment</li> <li>Omnispec ED1000</li> </ul>	With an element that block delivery of shockwaves	<ul> <li>Two treatments/week</li> <li>3 weeks treatment</li> <li>3 weeks no treatment</li> <li>3 weeks treatment</li> </ul>	<ul> <li>I month</li> <li>3 months</li> <li>6 months</li> </ul>	ILEF variation scores
Kim et al.	2019	Korea	8	<b>6</b> 4.2 ± <b>6</b> .6	ΥN	<ul> <li>Energy density: 20 m//mm<sup>2</sup></li> <li>15 m//mm<sup>2</sup></li> <li>12 m//mm<sup>2</sup></li> <li>Frequency: 5 Hz</li> <li>Pulse: 3000 pulses/treatment</li> <li>MT 2000H</li> </ul>	Sham treatment without delivering any energy	<ul> <li>Two treatments/week</li> <li>3 weeks treatment</li> <li>3 weeks treatment</li> <li>3 weeks treatment</li> </ul>	• - month • 2 months	<ul> <li>EHS response rate</li> <li>IEF variation scores</li> </ul>
Kitrey et al.	2015	Israel	55	62 (28–81)	°Z	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Frequency: 2 Hz</li> <li>Pulse: 1,500 pulses/treatment</li> <li>Omnispec ED1000</li> </ul>	Sham treatment without delivering any energy	<ul> <li>Two treatments/week</li> <li>3 weeks treatment</li> <li>3 weeks no treatment</li> <li>3 weeks treatment</li> </ul>	• month	<ul> <li>EHS response rate</li> <li>IIEF variation scores</li> </ul>
Ladegaard et al.	2021	Denmark	38	<b>62.5</b> ± 5.8	Yes	<ul> <li>Energy density: 0.15 mJ/mm<sup>2</sup></li> <li>Frequency: 5 Hz</li> <li>Pulse: 4000 pulses/treatment</li> <li>Duolith SD1</li> </ul>	With a sham pad that prevent shockwaves	<ul> <li>One treatment/week</li> <li>5 weeks treatment</li> </ul>	<ul> <li>I month</li> <li>3 months</li> </ul>	IIEF variation scores
Olsen et al.	2015	Denmark	105	60 (37–80)	Yes	<ul> <li>Energy density: 0.15 mJ/mm<sup>2</sup></li> <li>Frequency: 5 Hz</li> <li>Pulse: 3,000 pulses/treatment</li> <li>Duolith SD1</li> </ul>	With a cap used to prevent LI-ESWT	<ul> <li>One treatment/week</li> <li>5 weeks treatment</li> </ul>	<ul> <li>I month</li> <li>3 months</li> <li>6 months</li> </ul>	• EHS response rate
Ortac et al.	2021	Turkey	66	41 ± 10.7	NA	<ul> <li>Energy density: 0.2 mJ/mm<sup>2</sup></li> <li>Frequency: 5 Hz</li> <li>Pulse: 3,000 pulses/treatment</li> <li>Duolith SD1</li> </ul>	With a shock wave absorbent material	<ul> <li>One treatment/week</li> <li>4 weeks treatment</li> </ul>	<ul> <li>3 months</li> <li>6 months</li> </ul>	<ul> <li>IIEF variation scores</li> <li>SEP2 response rate</li> <li>SEP3 response rate</li> </ul>

(continued)

Table I. The details of each study.

	Outcome indicators	IIEF variation scores	<ul> <li>IIEF variation scores</li> <li>SEP2 response rate</li> <li>SEP3 response rate</li> </ul>	<ul> <li>EHS response rate</li> <li>IEF variation scores</li> </ul>	<ul> <li>EHS response rate</li> <li>IIEF variation scores</li> </ul>	<ul> <li>EHS response rate</li> <li>IEF variation scores</li> <li>SEP2 response rate</li> <li>SEP3 response rate</li> </ul>	ILEF variation scores	IIEF variation scores	<ul> <li>EHS response rate</li> <li>IIEF variation scores</li> </ul>	
Follow-up	time	• 3 months	<ul><li>I month</li><li>3 months</li></ul>	<ul> <li>I month</li> <li>3 months</li> <li>6 months</li> <li>9 month</li> <li>12 months</li> </ul>	• I month	<ul> <li>I month</li> <li>3 months</li> <li>6 months</li> </ul>	<ul><li>I month</li><li>3 months</li></ul>	• I month	<ul><li> 3 months</li><li> 6 months</li></ul>	
	Methodology	Two treatments/week 3 weeks treatment 3 weeks no treatment 3 weeks treatment	Two treatments/week 4 weeks treatment	Two treatments/week 3 weeks treatment 3 weeks no treatment 3 weeks treatment	Two treatments/week 3 weeks treatment 3 weeks no treatment 3 weeks treatment	One treatment/week 4 weeks treatment	Two treatments/week 3 weeks treatment	Two treatments/week 3 weeks treatment 3 weeks no treatment 3 weeks treatment	Two treatments/week 3 weeks treatment 3 weeks no treatment 3 weeks treatment	: : : :
		• • • •	• •	• • • •	• • • •	• •	••	• • • •	••••	
	Control setup	With an element that blocked the delivery of shock waves	With a gel head that blocked shockwaves	With a metal plate to block the transmission of the shockwave energy	With a metal plate that prevented the shock wave energy	With a probe that did not generate shockwaves	With a probe that emitted 0 energy	The energy setting was 0 during each treatment	Without any therapy	
	LI-ESWT setup	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Pulse: 3,000 pulses/treatment</li> <li>Duolith SD1</li> </ul>	<ul> <li>Energy density: 0.16 mJ/mm<sup>2</sup></li> <li>Richard Wolf GmbH</li> </ul>	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Frequency: 2 Hz</li> <li>Pulse: 1,500 pulses/treatment</li> <li>Omnispec ED1000</li> </ul>	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Frequency: 5 Hz</li> <li>Pulse: 1,500 pulses/treatment</li> <li>Omnispec ED1000</li> </ul>	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Frequency: 2 Hz</li> <li>Pulse: 5,000 pulses/treatment</li> <li>RENOVA® electromagnetic</li> <li>device</li> </ul>	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Pulse: 2,000 pulses/treatment</li> <li>Swiss Dolorclast</li> </ul>	<ul> <li>Energy density: 0.09 mJ/mm<sup>2</sup></li> <li>Frequency: 2 Hz</li> <li>Pulse: 1,500 pulses/treatment</li> <li>Omnispec ED1000</li> </ul>	Energy density: 0.09 mJ/mm <sup>2</sup> Frequency: 2 Hz Pulse: 1,500 pulses/treatment Dornier Aries device	
. 0		• • •	• •	••••	• • • •	••••	• • •	• • • •	••••	
PDE5I-	or not	Yes	Yes	Yes	Yes	°Z	AN	Yes	Ϋ́	
	Age	48 -+ 5.6	<b>54.3</b> ± <b>9.2</b>	Not mentioned	57 (27–77)	60 (53–66)	54 (46–61)	61.0 ± 7.3	52.1 ± 6.8	
No. of	participants	42	60	4	60	76	20	58	85	
	Country	Egypt	Czech	India	Israel	Spain	Brazil	China	Egypt	
	Year	2021	2019	2015	2012	2020	2018	2014	2018	
	Authors' name	Shendy et al.	Sramkova et al.	Srini et al.	Vardi et al.	Vinay et al.	Yamaçake et al.	Yee et al.	Zewin et al.	

Table I. (continued)

Note. PDE51 = phosphodiesterase 5 inhibitor; LL-ESWT = low-intensity extracorporeal shock wave therapy; NA = not available; IIEF = International Index of Erectile Function; EHS = Erectile Hardness Score.



Figure 2. The Risk of Bias Graph.

#### International Index of Erectile Function (IIEF)

Among the 16 included studies, 12 articles provided IIEF data at 1 month follow-up after treatment, eight provided data at 3 months follow-up, and four provided data at 6 months follow-up. Heterogeneity test proved that there was statistical heterogeneity among trials in each group, so random effect model was used for meta-analysis. The results reported that after 1 month (MD = 3.18, 95% CI = [1.38, 4.98], p = .0005), 3 months (MD = 3.01, 95% CI = [2.04, 3.98], p < .00001), and 6 months follow-up (MD = 3.20, 95% CI = [2.49, 3.92], p < .00001), the treatment group can significantly increase the IIEF of ED patients compared with the control group, and the results are statistically significant (Figure 3). The IIEF data analyzed are all variation values and some data with negative change values are replaced by their final values.

## **Erection Hardness Scores**

Overall, eight of the 16 articles provided data on the improvement of patients with baseline Erectile Hardness Score (EHS)  $\leq 2$  to EHS  $\geq 3$  after treatment. The random effect model was used for the meta-analysis. The results identified that there was a significant difference in the number of people of EHS improvement between the treatment group and the control group (OR = 5.07, 95% CI = [1.78, 14.44], p = .002), indicating that the treatment group can significantly improve the EHS of patients compared with the control group (Supplemental Figure S2).

#### Sexual Encounter Profile (SEP)

Questions 2 and 3 of the SEP are usually used as another evaluation criterion. These two questions were mentioned in three studies. The results identified that compared with the control group, the "yes" response rate of the LI-ESWT group was not statistically significant (SEP2: OR = 1.27, 95% CI = [0.70, 2.30], p = .43; SEP3: OR = 4.24, 95% CI = [0.67, 26.83], p = .13; Supplemental Figure S3).

## Subgroup Analysis

*IIEF Baseline*. According to the IIEF baseline value, the articles were divided into severe group (IIEF baseline value < 12), moderate group (IIEF baseline value 12–17), and mild group (IIEF baseline value > 17). We use the existing data to analysis and found that no matter in which subgroup, the improvement of IIEF in the treatment group was higher than that in the control group (severe: MD = 4.07, 95% CI = [0.49, 7.64], p = .03; moderate: MD = 4.24, 95% CI = [2.88, 5.59], p < .00001; mild: MD = 3.87, 95% CI = [3.37, 4.36], p < .00001; Figure 4).

Energy Density. Because two of the 16 RCTs did not mention specific energy density or IIEF index, only the remaining 14 experiments were analyzed. According to the set energy density, it is divided into two groups: the energy density is equal to 0.09 mJ/mm<sup>2</sup> and the energy density is between 0.1 and 0.2 mJ/mm<sup>2</sup>. The results suggested that in the two subgroups, the treatment group could significantly increase the IIEF of patients compared with the control group (0.09 mJ/mm<sup>2</sup>: MD = 3.81, 95% CI = [2.07, 5.55], p < .0001; 0.1–0.2 mJ/mm<sup>2</sup>: MD = 3.01, 95% CI = [0.89, 5.12], p = .005; Figure 5).

*Pulses.* We divided 14 RCTs into three groups according to the number of pulses per treatment: the number of pulses is equal to 600, the number of pulses is between 1,500 and 2,000, and the number of pulses is greater than 3,000. The results of the 600 pulses group reported that the treatment group could increase the IIEF of patients compared with the control group, but the difference was not statistically significant (MD = 1.50, 95% CI = [-1.44, 4.43], p = .32).

A	LI-	ESWT		C	ontrol			Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI	IV, Random, 95% Cl
Baccaglini 2019	12.39	5.02	36	9.29	3.07	41	8.1%	3.10 [1.21, 4.99]	
Fojecki 2016	13.1	3	58	13	3	60	8.6%	0.10 [-0.98, 1.18]	
<alyvianakis 2017<="" td=""><td>4.66</td><td>3.6</td><td>30</td><td>1.83</td><td>3.45</td><td>16</td><td>7.9%</td><td>2.83 [0.70, 4.96]</td><td></td></alyvianakis>	4.66	3.6	30	1.83	3.45	16	7.9%	2.83 [0.70, 4.96]	
<im 2019<="" td=""><td>20.6</td><td>3</td><td>38</td><td>14.6</td><td>3.2</td><td>43</td><td>8.4%</td><td>6.00 [4.65, 7.35]</td><td></td></im>	20.6	3	38	14.6	3.2	43	8.4%	6.00 [4.65, 7.35]	
<itrey 2015<="" td=""><td>13.35</td><td>6.9</td><td>37</td><td>8.14</td><td>3.22</td><td>18</td><td>7.4%</td><td>5.21 [2.53, 7.89]</td><td></td></itrey>	13.35	6.9	37	8.14	3.22	18	7.4%	5.21 [2.53, 7.89]	
_adegaard 2021	2.4	3.3	20	1.28	1.9	18	8.2%	1.12 [-0.57, 2.81]	
Gramkova 2019	18.7	4.75	30	16.3	4.89	30	7.6%	2.40 [-0.04, 4.84]	
Brini 2015	12.5	4.27	60	1.4	3.6	17	8.0%	11.10 [9.08, 13.12]	
/ardi 2012	6.7	4.7	40	3	3.8	20	7.8%	3.70 [1.49, 5.91]	
/inay 2020	13.13	9.23	40	11.78	10.03	36	5.8%	1.35 [-3.00, 5.70]	
ramaçake 2018	4.7	5.6	10	1.7	4.2	10	5.8%	3.00 [-1.34, 7.34]	
ree 2014	5.3	5.5	30	3.8	3.6	28	7.7%	1.50 [-0.88, 3.88]	
Zewin 2018	19.4	1.2	42	19.5	1.3	43	8.8%	-0.10 [-0.63, 0.43]	+
fotal (95% CI)			471			380	100.0%	3.18 [1.38, 4.98]	•
Heterogeneity: Tau <sup>2</sup> =	9.52; C	hi² = 18	36.87, 0	if = 12 (F	< 0.00	0001);1	²= 94%		
Fest for overall effect	Z= 3.46	i (P = 0	.0005)						-10 -5 U 5 10 Control Experimental
В									Control Experimental
	Exp	erimer	ntal	(	Control			Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Tota	I Weight	IV, Random, 95% Cl	IV, Random, 95% Cl
<alyvianakis 2017<="" td=""><td>4.66</td><td>3.5</td><td>30</td><td>1.33</td><td>3.5</td><td>18</td><td>5 11.7%</td><td>3.33 [1.21, 5.45]</td><td></td></alyvianakis>	4.66	3.5	30	1.33	3.5	18	5 11.7%	3.33 [1.21, 5.45]	
_adegaard 2021	3.45	4.01	20	0.65	2.03	18	3 12.5%	2.80 [0.81, 4.79]	
Ortac 2021	2.78	2.57	42	1.27	1.87	22	2 19.7%	1.51 [0.41, 2.61]	
Shendy 2021	4.85	2.96	21	0.65	2.73	21	14.4%	4.20 [2.48, 5.92]	
Bramkova 2019	7.7	5	30	2.5	3.63	30	11.2%	5.20 [2.99, 7.41]	
/inay 2020	15.71	10.76	40	11.84	12.35	38	3.0%	3.87 [-1.36, 9.10]	
Yamaçake 2018	6.3	5.4	10	1.6	4	10	4.5%	4.70 [0.53, 8.87]	
Zewin 2018	55.7	1.8	42	53.7	1.7	43	3 22.9%	2.00 [1.26, 2.74]	
fotal (95% CI)			235			196	5 100.0%	3.01 [2.04, 3.98]	•
-leterogeneity: Tau <sup>2</sup> =	= 0.93; C	hi² = 18	6.29, df	= 7 (P =	0.02);	1 <sup>2</sup> = 579	%		
Fest for overall effect	Z = 6.08	(P < 0	.00001	)					Control Experimental
C	Eyne	rimen	tal	C	ntrol			Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight I	V, Random, 95% Cl	IV, Random, 95% Cl
Kalwianakis 2017	5.2	3.5	30	1.52	3	16	12.9%	3.68 [1.75, 5.61]	
Ortac 2021	2.35	2.83	26	0.14	1.55	4	13.8%	2.21 [0.34, 4.08]	
/inay 2020	2.41	6.45	40	0.7	4.63	36	7.9%	1.71 [-0.80, 4.22]	
Zewin 2018	59.9	1.8	42	56.4	1.7	43	65.4%	3.50 [2.76, 4.24]	
otal (95% CI)			138			99	100.0%	3.20 [2.49, 3.92]	•
-leterogeneity: Tau <sup>2</sup> =	= 0.06; C	hi <sup>2</sup> = 3.	26, df =	3 (P = 0	0.35); I²	= 8%		-	-4 -2 0 2 4
lact for overall offect	1 = 8.78	(P < 0	10001	1					

**Figure 3.** Forest Plots Showing the Improvement of IIEF by LI-ESWT at Different Follow-Up Times After Treatment: (A) I Month Follow-Up; (B) 3 Months Follow-Up; (C) 6 Months Follow-Up

Note. LI-ESWT = low-intensity extracorporeal shock wave therapy; IIEF = International Index of Erectile Function; SD = standard deviation; IV = inverse variance; CI = confidence interval; df = degrees of freedom.

The results of the group with pulse number between 1,500 and 2,000 suggested that the treatment group could significantly increase the IIEF of patients (MD = 4.80, 95% CI = [2.61, 7.00], p < .0001). In the group with pulse number greater than 3,000, compared with the control group, the treatment group can also significantly increase the IIEF of patients and the difference is statistically significant (MD = 3.46, 95% CI = [1.89, 5.03], p < .0001; Figure 6).

# Discussion

We conducted a meta-analysis of 16 studies including 1,064 participants to compare the efficacy of LI-ESWT

and placebo in the treatment of ED. It was found that the improvement of IIEF and EHS after LI-ESWT treatment was greater than that of placebo group, but there was no significant difference in SEP2 and SEP3. These results suggest that LI-ESWT is more effective than placebo in improving the symptoms of ED patients.

At present, the clinical treatment methods of ED include oral PDE5I, injection of vasodilator into corpus cavernosum of penis, transurethral administration of prostaglandin E, penile prosthesis implantation, and vacuum assisted erection device (Salonia et al., 2021). One of the most commonly used regimes is PDE5I drug treatment, but this plan cannot correct the potential

	Exp	eriment	tal	C	ontrol	-		Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% Cl	IV, Random, 95% Cl
1.1.1 severe group						~~			
FOJECKI 2016	13.1	3	58	13	3	60	8.0%	0.10 [-0.98, 1.18]	Τ
Kitrey 2015	5.65	5	37	0.13	3.21	18	6.8%	5.52 [3.33, 7.71]	
Ladegaard 2021	3.45	4.01	20	0.65	2.03	18	7.1%	2.80 [0.81, 4.79]	
Srini 2015	12.5	4.27	60	1.4	3.6	17	7.0%	11.10 [9.08, 13.12]	
Vardi 2012	6.7	4.7	40	3	3.8	20	6.8%	3.70 [1.49, 5.91]	
Yamaçake 2018	17.2	5.7	10	16.5	5	10	4.0%	0.70 [-4.00, 5.40]	
Subtotal (95% CI)			225			143	39.7%	4.07 [0.49, 7.64]	
Heterogeneity: Tau² =	= 18.29; (	Chi² = 9	5.52, d	f= 5 (P	< 0.00	001); I²	= 95%		
Test for overall effect	Z = 2.23	P = 0.	03)						
1.1.2 moderate grou	р								
Kalyvianakis 2017	5.3	3.2	30	1.4	3.1	16	7.2%	3.90 [2.00, 5.80]	
Kim 2019	21.7	8.2	38	14.5	5.4	43	5.7%	7.20 [4.13, 10.27]	
Shendy 2021	4.85	2.96	21	0.65	2.73	21	7.4%	4.20 [2.48, 5.92]	
Sramkova 2019	7.7	5	30	2.5	3.63	30	6.8%	5.20 [2.99, 7.41]	
Vinay 2020	14.65	11.53	40	10.49	8.49	36	4.1%	4.16 [-0.36, 8.68]	
Yee 2014	5.3	5.5	30	3.8	3.6	28	6.6%	1.50 [-0.88, 3.88]	<b>_</b>
Subtotal (95% CI)			189			174	37.7%	4.24 [2.88, 5.59]	$\bullet$
Heterogeneity: Tau <sup>2</sup> =	= 1.29; C	hi² = 9.5	51, df =	5 (P = 0	.09); P	<sup>2</sup> = 47%			
Test for overall effect	Z = 6.14	(P < 0.	00001)	L.					
1.1.3 mild group					-				
Baccaglini 2019	12.39	5.02	36	9.29	3.07	41	7.2%	3.10 [1.21, 4.99]	
Ortac 2021	22.67	3.35	26	19.82	1.56	4	7.0%	2.85 [0.85, 4.85]	
Zewin 2018	60.7	1.2	42	56.7	1.3	43	8.4%	4.00 [3.47, 4.53]	
Subtotal (95% CI)			104			88	22.6%	3.87 [3.37, 4.36]	•
Heterogeneity: Tau <sup>2</sup> =	= 0.00; C	hi² = 1.8	87, df =	2(P = 0	.39); P	*= 0%			
Test for overall effect	Z = 15.2	29 (P < 0	0.0000	1)					
Total (05% CI)			510			405	100 0%	4 02 12 74 5 201	•
Hotorogonoity Tou?	5 02· 0	hiz - 14	1 6 0 4	6-14/5	~ ^ ^ ^	405	3- 070	4.02 [2.74, 5.30]	
Telefogeneity: Tau*=	= 5.03, C		1.00,0	1 = 14 (F	< 0.0	0001);1	= 87%		-10 -5 0 5 10
Test for overall effect	∠=0.15	) (P < U. . ∩ ∺?	00001)	6 0 (0	0.00	. 17 0	~		Control Experimental
l est for subaroup dif	ierences	: Chi*=	0.26.0	$\mathbf{r} = 2$ (P	= 0.88	o. r= 0	%		

Figure 4. Forest Plots Showing the Subgroup Analysis of Different IIEF Baselines

Note. IIEF = International Index of Erectile Function; SD = standard deviation; IV = inverse variance; CI = confidence interval; df = degrees of freedom.

pathophysiological mechanism of the penis, and many patients are insensitive or even ineffective to it. LI-ESWT is noninvasive and rehabilitative compared with the second-line or third-line treatment of ED, and patients who are ineffective in PDE5I treatment can also benefit from LI-ESWT treatment (Chung & Cartmill, 2015; Gruenwald et al., 2012; Kitrey et al., 2016).

The mechanism of LI-ESWT improving IIEF in the treatment of ED is not clear. In recent years, it has been identified that ESW can produce "cavitation effect," open up physiologically closed micro vessels, and accelerate capillary microcirculation (Maisonhaute et al., 2002). ESW can also promote neovascularization and the expression of angiogenic markers, so as to promote tissue remodeling (Holfeld et al., 2016; Young Academic Urologists Men's Health Group et al., 2017). However, there is controversy about whether LI-ESWT is associated with neuronal nitric oxide synthase (nNOS) synthesis. Studies have identified that LI-ESWT can promote the regeneration of in endothelial, smooth muscle, and neural expression of nNOS (Liu et al., 2013; Qiu et al., 2013), but there are also

studies reported that LI-ESWT does not rely on nitric oxide and cyclic guanosine monophosphate to improve erectile function (Assaly-Kaddoum et al., 2016). One study described that LI-ESWT can also reduce the activity of sympathetic nervous system (Sokolakis et al., 2019). Most studies only report preliminary results, but there is no clear answer to the actual mechanism of LI-ESWT.

By analyzing our results, we found that LI-ESWT had different effects on erectile function with different energy density or pulses. When the energy density is 0.09 mJ/ mm<sup>2</sup>, the improvement of IIEF is better than that in the energy density between 0.1 and 0.2 mJ/mm<sup>2</sup>. And 1,500 or 2,000 pulses per treatment bring more improvement than 600 or 3,000 pulses. The improvement of IIEF in patients with different severity of ED after LI-ESWT treatment is also different. Through our meta-analysis, we found that the improvement was more obvious in patients with moderate ED than in patients with mild or severe ED. In addition, the improvement of IIEF is different under different follow-up times. The improvement after 6 months follow-up is better than that after 1 month

	al	C	ontrol			Mean Difference	Mean Difference		
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% Cl	IV, Random, 95% Cl
1.2.1 Energy density	0.09mJ/	mm <sup>2</sup>							
Baccaglini 2019	12.39	5.02	36	9.29	3.07	41	7.6%	3.10 [1.21, 4.99]	
Fojecki 2016	13.1	3	58	13	3	63	8.5%	0.10 [-0.97, 1.17]	
Kalwianakis 2017	4.66	3.5	30	1.33	3.5	16	7.3%	3.33 [1.21, 5.45]	
Kitrey 2015	13.35	6.9	37	8.14	3.22	18	6.6%	5.21 [2.53, 7.89]	
Shendy 2021	4.85	2.96	21	0.65	2.73	21	7.8%	4.20 [2.48, 5.92]	
Srini 2015	12.5	4.27	60	1.4	3.6	17	7.4%	11.10 [9.08, 13.12]	
Vardi 2012	6.7	4.7	40	3	3.8	20	7.2%	3.70 [1.49, 5.91]	_ <b>→</b>
Vinay 2020	15.71	10.76	40	11.84	12.35	36	3.8%	3.87 [-1.36, 9.10]	
Yamaçake 2018	6.3	5.4	10	1.6	4	10	4.9%	4.70 [0.53, 8.87]	
Yee 2014	5.3	5.5	30	3.8	3.6	28	7.0%	1.50 [-0.88, 3.88]	+
Zewin 2018	55.7	1.8	42	53.7	1.7	43	8.7%	2.00 [1.26, 2.74]	
Subtotal (95% CI)			404			313	76.9%	3.81 [2.07, 5.55]	•
Heterogeneity: Tau <sup>2</sup> =	7.10; C	hi² = 10	1.86, di	f = 10 (P	< 0.00	001); I <sup>z</sup>	= 90%		
Test for overall effect:	Z = 4.30	(P < 0.)	0001)						
1.2.2 Energy density	0.1.0.2n	n.l/mm <sup>2</sup>	:						
Ladegaard 2021	3.45	4.01	20	0.65	2.03	18	7.5%	2.80 [0.81, 4.79]	
Ortac 2021	2.78	2.57	42	1.27	1.87	22	8.4%	1.51 [0.41, 2.61]	
Sramkova 2019	7.7	5	30	2.5	3.63	30	7.2%	5.20 [2.99, 7.41]	
Subtotal (95% CI)			92			70	23.1%	3.01 [0.89, 5.12]	
Heterogeneity: Tau <sup>2</sup> =	2.65; C	hi² = 8.8	1, df=	2 (P = 0	.01); I <sup>2</sup> :	= 77%			
Test for overall effect:	Z = 2.79	(P = 0.	005)						
Total (95% Cl)			496			383	100.0%	3.62 [2.26, 4.98]	•
Heterogeneity: Tau <sup>2</sup> =	5.39; C	hi <sup>2</sup> = 11	1.00, di	f = 13 (P	< 0.00	001); I <sup>z</sup>	= 88%	- / -	
Test for overall effect:	Z = 5.23	(P < 0.	00001)						-10 -5 0 5 10
		- Ohiz -	0.00 4	6 - 1 /D	- 0 5 63	13 - 00	,		Control Experimental

**Figure 5.** Forest Plots Showing the Subgroup Analysis of Different Energy Density Treatments Note. SD = standard deviation; IV = inverse variance; CI = confidence interval; df = degrees of freedom.

$\begin{array}{c c c c c c c c c c c c c c c c c c c $	Study or Subgroup	Expe	erimen	tal Total	Moan	ontrol	Total	Moight	Mean Difference	Mean Difference
The formation of the second in the formation of the second in the seco	1 3 1 600 nulses/trea	atment	30	Total	mean	30	Total	weight	1v, Random, 35% CI	14, Randolli, 35% Cl
Fojecki 2016 13.1 3 58 13 3 60 8.3% 0.10 [-0.88, 1.5] Subtotal (95% CI) 94 101 15.8% 1.50 [-1.44, 4.43] Heterogeneity: Tau <sup>2</sup> = 3.88; Chi <sup>2</sup> = 7.29, df = 1 (P = 0.007); P = 86% Test for overall effect: $Z = 1.00$ (P = 0.32) 1.3.2 1500-2000 pulses/treatment Kalyvianakis 2017 5.3 3.2 30 1.4 3.1 16 7.5% 3.90 [2.00, 5.80] Kitrey 2015 13.25 6.9 37 8.14 3.22 18 6.6% 5.21 [2.53, 7.89] Srini 2015 12.5 4.27 60 1.4 3.6 17 7.3% 11.10 [9.08, 13.12] Vardi 2012 6.7 4.7 40 3 3.8 20 7.1% 3.70 [1.09, 5.91] Yamaçake 2018 6.3 5.4 10 1.6 4 10 4.9% 4.70 [0.53, 8.87] Yee 2014 5.3 5.5 30 3.8 3.6 28 6.9% 1.50 [-0.88, 3.88] Zewin 2018 59.9 1.8 42 56.4 1.7 43 8.5% 3.50 [2.76, 4.24] Subtotal (95% CI) 249 152 48.8% 4.80 [2.61, 7.00] Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); P = 89% Test for overall effect: $Z = 4.29$ (P < 0.0001) 1.3.3 $\geqslant$ 3000 pulses/treatment Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ordra 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% CI) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); P = 62% Test for overall effect: $Z = 4.32$ (P < 0.0001) Total (95% CI) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); P = 88%	Baccaglini 2019	12.39	5.02	36	9 29	3.07	41	7.5%	3 10 [1 21 4 99]	
Subtotal (95% CI) 94 101 15.8% 1.50 [-1.44, 4.43] Heterogeneity: Tau <sup>2</sup> = 3.88; Ch <sup>2</sup> = 7.29, df = 1 (P = 0.007); P = 86% Test for overall effect Z = 1.00 (P = 0.32) 1.3.2 1500-2000 pulses/treatment Kalyvianakis 2017 5.3 3.2 30 1.4 3.1 16 7.5% 3.90 [2.00, 5.80] Kitrey 2015 12.5 4.27 60 1.4 3.6 17 7.3% 11.10 [9.08, 13.12] Vardi 2015 12.5 4.27 60 1.4 3.6 17 7.3% 11.10 [9.08, 13.12] Vardi 2012 6.7 4.7 40 3 3.8 20 7.1% 3.70 [1.49, 5.91] Yamaçake 2018 6.3 5.4 10 1.6 4 10 4.9% 4.70 [0.53, 8.87] Yee 2014 5.3 5.5 30 3.8 3.6 28 6.9% 1.50 [-0.88, 3.88] Zewin 2018 59.9 1.8 42 56.4 1.7 43 8.5% 3.50 [2.76, 4.24] Subtotal (95% CI) 249 152 48.8% 4.80 [2.61, 7.00] Heterogeneity: Tau <sup>2</sup> = 7.32; Ch <sup>2</sup> = 53.89, df = 6 (P < 0.00001); P = 89% Test for overall effect Z = 4.29 (P < 0.0001) 1.3.3 $\ge$ 3000 pulses/treatment Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ortac 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 8.45 40 0.7 4.83 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% CI) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Ch <sup>2</sup> = 10.66, df = 4 (P = 0.03); P = 62% Test for overall effect Z = 4.32 (P < 0.0001) Total (95% CI) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 10.66, df = 13 (P < 0.00001); P = 88%	Fojecki 2016	13.1	3	58	13	3	60	8.3%	0.10 [-0.98, 1.18]	
Heterogeneity: Tau <sup>2</sup> = 3.88; Chi <sup>2</sup> = 7.29, df = 1 (P = 0.007); P = 86% Test for overall effect: Z = 1.00 (P = 0.32) <b>1.3.2 f500-2000 pulses/treatment</b> Kalyvianakis 2017 5.3 3.2 30 1.4 3.1 16 7.5% 3.90 [2.00, 5.80] Kitrey 2015 13.35 6.9 37 8.14 3.22 18 6.6% 5.21 [2.53, 7.89] Srini 2015 12.5 4.27 60 1.4 3.6 17 7.3% 11.10 [9.08, 13.12] Vardi 2012 6.7 4.7 40 3 3.8 20 7.1% 3.70 [1.49, 5.91] Yamaçake 2018 6.3 5.4 10 1.6 4 10 4.9% 4.70 [0.53, 8.87] Yee 2014 5.3 5.5 30 3.8 3.6 28 6.9% 1.50 [-0.88, 3.88] Zewin 2018 59.9 1.8 42 56.4 1.7 43 8.5% 3.50 [2.76, 4.24] Subtotal (95% Cl) 249 152 48.8% 4.80 [2.61, 7.00] Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); P = 89% Test for overall effect: Z = 4.29 (P < 0.0001) <b>1.3.3</b> $\ge$ 3000 pulses/treatment Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Orda: 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% Cl) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); P = 62% Test for overall effect: Z = 4.32 (P < 0.0001) Total (95% Cl) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 10.66, df = 1 (P < 0.00001); P = 88%	Subtotal (95% CI)		-	94		-	101	15.8%	1.50 [-1.44, 4.43]	
Test for overall effect: $Z = 1.00$ (P = 0.32) <b>1.3.2 1500-2000 pulses/treatment</b> Kalyvianakis 2017 5.3 3.2 30 1.4 3.1 16 7.5% 3.90 [2.00, 5.80] Kitrey 2015 13.35 6.9 37 8.14 3.22 18 6.6% 5.21 [2.53, 7.89] Srini 2015 12.5 4.27 60 1.4 3.6 17 7.3% 11.10 [9.08, 13.12] Vardi 2012 6.7 4.7 40 3 3.8 20 7.1% 3.70 [1.49, 5.91] Yamaçake 2018 6.3 5.4 10 1.6 4 10 4.9% 4.70 [0.53, 8.87] Yee 2014 5.3 5.5 30 3.8 3.6 28 6.9% 1.50 [-0.88, 3.88] Zewin 2018 59.9 1.8 42 56.4 1.7 43 8.5% 3.50 [2.76, 4.24] Subtoal (95% Cl) 249 152 48.8% 4.80 [2.61, 7.00] Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); P = 89% Test for overall effect: Z = 4.29 (P < 0.0001) <b>1.3.3 <math>\geq</math> 3000 pulses/treatment</b> Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% Cl) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); P = 62% Test for overall effect: Z = 4.32 (P < 0.0001) Total (95% Cl) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 10.66, df = 1 (P < 0.00001); P = 88%	Heterogeneity: Tau <sup>2</sup> =	= 3.88; CI	hi² = 7.	29, df=	: 1 (P =	0.007)	; l² = 88	i%		
1.3.2 1500-2000 pulses/treatment         Kalyvianakis 2017       5.3       3.2       30       1.4       3.1       16       7.5%       3.90 [2.00, 5.80]         Kitrey 2015       13.35       6.9       37       8.14       3.22       18       6.6%       5.21 [2.53, 7.89]         Srini 2015       12.5       4.27       60       1.4       3.6       17       7.3%       11.10 [9.08, 13.12]         Vardi 2012       6.7       4.7       40       3       3.8       20       7.1%       3.70 [1.49, 5.91]         Yamagake 2018       6.3       5.4       10       1.6       4       10       4.9%       4.70 [0.53, 8.87]         Yee 2014       5.3       5.5       30       3.8       3.6       28       6.9%       1.50 [-0.88, 3.88]         Zewin 2018       59.9       1.8       42       56.4       1.7       43       8.5%       3.50 [2.76, 4.24]         Subtotal (95% CI)       249       152       48.8%       4.80 [2.61, 7.00]       Heterogeneity: Tau" = 7.32; Chi" = 53.89, df = 6 (P < 0.00001); I* = 89%	Test for overall effect:	Z=1.00	) (P = 0	.32)						
Kalyvianakis 2017 5.3 3.2 30 1.4 3.1 16 7.5% 3.90 [2.00, 5.80] Kilrey 2015 13.35 6.9 37 8.14 3.22 18 6.6% 5.21 [2.53, 7.89] Srini 2015 12.5 4.27 60 1.4 3.6 17 7.3% 11.10 [9.08, 13.12] Vardi 2012 6.7 4.7 40 3 3.8 20 7.1% 3.70 [1.49, 5.91] Yamaçake 2018 6.3 5.4 10 1.6 4 10 4.9% 4.70 [0.53, 8.87] Yee 2014 5.3 5.5 30 3.8 3.6 28 6.9% 1.50 [-0.58, 3.88] Zewin 2018 59.9 1.8 42 56.4 1.7 43 8.5% 3.50 [2.76, 4.24] Subtotal (95% Cl) 249 152 48.8% 4.80 [2.61, 7.00] Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); I <sup>2</sup> = 89% Test for overall effect: Z = 4.29 (P < 0.0001) 1.3.3 $\ge 3000$ pulses/treatment Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ortac 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% Cl) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62% Test for overall effect: Z = 4.32 (P < 0.0001) Total (95% Cl) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	1.3.2 1500-2000 puls	ses/treat	ment							
Kitrey 2015       13.35       6.9       37       8.14       3.22       18       6.6%       5.21       [2.53, 7.89]         Srini 2015       12.5       4.27       60       1.4       3.6       17       7.3%       11.10       [9.08, 13.12]         Vardi 2012       6.7       4.7       40       3       3.8       20       7.1%       3.70       [1.49, 5.91]         Yamaçake 2018       6.3       5.4       10       1.6       4       10       4.9%       4.70       [0.53, 8.87]         Yee 2014       5.5       50       3.8       3.6       28       6.9%       1.50       6.83, 8.8]         Zewin 2018       59.9       1.8       42       56.4       1.7       43       8.5%       3.50       [2.61, 7.00]         Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); I <sup>2</sup> = 89%       Test for overall effect: Z = 4.29 (P < 0.0001)	Kalwianakis 2017	5.3	3.2	30	1.4	3.1	16	7.5%	3.90 [2.00, 5.80]	
Srini 2015       12.5       4.27       60       1.4       3.6       17       7.3%       11.10 [9.08, 13.12]         Vardi 2012       6.7       4.7       40       3       3.8       20       7.1%       3.70 [1.49, 5.91]         Yamaçake 2018       6.3       5.4       10       1.6       4       10       4.9%       4.70 [0.53, 8.87]         Yee 2014       5.3       5.5       30       3.8       3.6       28       6.9%       1.50 [-0.88, 3.88]         Zewin 2018       59.9       1.8       42       56.4       1.7       43       8.5%       3.50 [2.76, 4.24]         Subtotal (95% Cl)       249       152       48.8%       4.80 [2.61, 7.00]       Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); l <sup>2</sup> = 89%       Test for overall effect: Z = 4.29 (P < 0.0001)	Kitrey 2015	13.35	6.9	37	8.14	3.22	18	6.6%	5.21 [2.53, 7.89]	
Vardi 2012 6.7 4.7 40 3 3.8 20 7.1% 3.70 [1.49, 5.91] Yamaçake 2018 6.3 5.4 10 1.6 4 10 4.9% 4.70 [0.53, 8.87] Yee 2014 5.3 5.5 30 3.8 3.6 28 6.9% 1.50 [-0.88, 3.88] Zewin 2018 59.9 1.8 42 56.4 1.7 43 8.5% 3.50 [2.76, 4.24] Subtotal (95% Cl) 249 152 48.8% 4.80 [2.61, 7.00] Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); I <sup>2</sup> = 89% Test for overall effect: Z = 4.29 (P < 0.0001) 1.3.3 ≥3000 pulses/treatment Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ortac 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% Cl) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62% Test for overall effect: Z = 4.32 (P < 0.0001) Total (95% Cl) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	Srini 2015	12.5	4.27	60	1.4	3.6	17	7.3%	11.10 [9.08, 13.12]	
Yamaçake 2018 6.3 5.4 10 1.6 4 10 4.9% 4.70 [0.53, 8.87] Yee 2014 5.3 5.5 30 3.8 3.6 28 6.9% 1.50 [-0.88, 3.88] Zewin 2018 59.9 1.8 42 56.4 1.7 43 8.5% 3.50 [2.76, 4.24] Subtotal (95% CI) 249 152 48.8% 4.80 [2.61, 7.00] Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); I <sup>2</sup> = 89% Test for overall effect: Z = 4.29 (P < 0.0001) 1.3.3 $\geq$ 3000 pulses/treatment Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ortac 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% CI) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62% Test for overall effect: Z = 4.32 (P < 0.0001) Total (95% CI) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	Vardi 2012	6.7	4.7	40	3	3.8	20	7.1%	3.70 [1.49, 5.91]	
Yee 2014 5.3 5.5 30 3.8 3.6 28 6.9% 1.50 [-0.88, 3.88] Zewin 2018 59.9 1.8 42 56.4 1.7 43 8.5% 3.50 [2.76, 4.24] Subtotal (95% Cl) 249 152 48.8% 4.80 [2.61, 7.00] Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); I <sup>2</sup> = 89% Test for overall effect: $Z = 4.29$ (P < 0.0001) 1.3.3 $\ge$ 3000 pulses/treatment Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ortac 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% Cl) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62% Test for overall effect: $Z = 4.32$ (P < 0.0001) Total (95% Cl) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	Yamaçake 2018	6.3	5.4	10	1.6	4	10	4.9%	4.70 [0.53, 8.87]	
Zewin 2018 59.9 1.8 42 56.4 1.7 43 8.5% $3.50 [2.76, 4.24]$ Subtotal (95% CI) 249 152 48.8% $4.80 [2.61, 7.00]$ Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); I <sup>2</sup> = 89% Test for overall effect: Z = 4.29 (P < 0.0001) 1.3.3 $\geq 3000 \text{ pulses/treatment}$ Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ortac 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% CI) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62% Test for overall effect: Z = 4.32 (P < 0.0001) Total (95% CI) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 10.668, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	Yee 2014	5.3	5.5	30	3.8	3.6	28	6.9%	1.50 [-0.88, 3.88]	+
Subtotal (95% Cl)       249       152       48.8%       4.80 [2.61, 7.00]         Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); I <sup>2</sup> = 89%       Test for overall effect: Z = 4.29 (P < 0.0001)	Zewin 2018	59.9	1.8	42	56.4	1.7	43	8.5%	3.50 [2.76, 4.24]	-
Heterogeneity: Tau <sup>2</sup> = 7.32; Chi <sup>2</sup> = 53.89, df = 6 (P < 0.00001); I <sup>2</sup> = 89% Test for overall effect: Z = 4.29 (P < 0.0001) <b>1.3.3</b> ≥ <b>3000 pulses/treatment</b> Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ortac 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% Cl) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62% Test for overall effect: Z = 4.32 (P < 0.0001) Total (95% Cl) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	Subtotal (95% CI)			249			152	48.8%	4.80 [2.61, 7.00]	-
Test for overall effect: $Z = 4.29$ (P < 0.0001) <b>1.3.3</b> ≥3000 pulses/treatment Kim 2019 21.7 8.2 43 14.5 5.4 38 6.2% 7.20 [4.21, 10.19] Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ortac 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% Cl) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62% Test for overall effect: Z = 4.32 (P < 0.0001) Total (95% Cl) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	Heterogeneity: Tau <sup>2</sup> =	= 7.32; C	hi² = 53	3.89, df	= 6 (P <	< 0.00	001); I <sup>z</sup>	= 89%		
1.3.3 ≥ 3000 pulses/treatment         Kim 2019       21.7       8.2       43       14.5       5.4       38       6.2%       7.20 [4.21, 10.19]         Ladegaard 2021       3.45       4.01       20       0.65       2.03       18       7.4%       2.80 [0.81, 4.79]         Ortac 2021       2.35       2.83       26       0.14       1.55       4       7.5%       2.21 [0.34, 4.08]         Shendy 2021       4.85       2.96       21       0.65       2.73       21       7.7%       4.20 [2.48, 5.92]         Vinay 2020       2.41       6.45       40       0.7       4.63       36       6.8%       1.71 [-0.80, 4.22]         Subtotal (95% Cl)       150       117       35.5%       3.46 [1.89, 5.03]         Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62%       Test for overall effect: Z = 4.32 (P < 0.0001)	Test for overall effect:	Z= 4.29	) (P < 0	1.0001)						
Kim 2019       21.7       8.2       43       14.5       5.4       38       6.2%       7.20 [4.21, 10.19]         Ladegaard 2021       3.45       4.01       20       0.65       2.03       18       7.4%       2.80 [0.81, 4.79]         Ortac 2021       2.35       2.83       26       0.14       1.55       4       7.5%       2.21 [0.34, 4.08]         Shendy 2021       4.85       2.96       21       0.65       2.73       21       7.7%       4.20 [2.48, 5.92]         Vinay 2020       2.41       6.45       40       0.7       4.63       36       6.8%       1.71 [-0.80, 4.22]         Subtotal (95% CI)       150       117       35.5%       3.46 [1.89, 5.03]         Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); l <sup>2</sup> = 62%       720 [4.24, 5.21]       +         Tost (95% CI)       493       370       100.0%       3.84 [2.46, 5.21]         Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 10.668, df = 13 (P < 0.00001); l <sup>2</sup> = 88%       -10       -5       0       5       10	1.3.3 ≥3000 pulses/	treatme	nt							
Ladegaard 2021 3.45 4.01 20 0.65 2.03 18 7.4% 2.80 [0.81, 4.79] Ortac 2021 2.35 2.83 26 0.14 1.55 4 7.5% 2.21 [0.34, 4.08] Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% Cl) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62% Test for overall effect: $Z = 4.32$ (P < 0.0001) Total (95% Cl) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	Kim 2019	21.7	8.2	43	14.5	5.4	38	6.2%	7.20 [4.21, 10.19]	
Ortac 2021       2.35       2.83       26       0.14       1.55       4       7.5%       2.21       [0.34, 4.08]         Shendy 2021       4.85       2.96       21       0.65       2.73       21       7.7%       4.20       [2.48, 5.92]         Vinay 2020       2.41       6.45       40       0.7       4.63       36       6.8%       1.71       [-0.80, 4.22]         Subtotal (95% Cl)       150       117       35.5%       3.46       [1.89, 5.03]         Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); l <sup>2</sup> = 62%       3.46       [1.89, 5.03]         Test for overall effect: Z = 4.32 (P < 0.0001)	Ladegaard 2021	3.45	4.01	20	0.65	2.03	18	7.4%	2.80 [0.81, 4.79]	
Shendy 2021 4.85 2.96 21 0.65 2.73 21 7.7% 4.20 [2.48, 5.92] Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% CI) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); I <sup>2</sup> = 62% Test for overall effect: $Z = 4.32$ (P < 0.0001) Total (95% CI) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	Ortac 2021	2.35	2.83	26	0.14	1.55	4	7.5%	2.21 [0.34, 4.08]	
Vinay 2020 2.41 6.45 40 0.7 4.63 36 6.8% 1.71 [-0.80, 4.22] Subtotal (95% CI) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); l <sup>2</sup> = 62% Test for overall effect: $Z = 4.32$ (P < 0.0001) Total (95% CI) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); l <sup>2</sup> = 88%	Shendy 2021	4.85	2.96	21	0.65	2.73	21	7.7%	4.20 [2.48, 5.92]	
Subtotal (95% CI) 150 117 35.5% 3.46 [1.89, 5.03] Heterogeneitly: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); i <sup>2</sup> = 62% Test for overall effect: $Z = 4.32$ (P < 0.0001) Total (95% CI) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneitly: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); i <sup>2</sup> = 88% -10 -5 0 5 10	Vinay 2020	2.41	6.45	40	0.7	4.63	36	6.8%	1.71 [-0.80, 4.22]	
Heterogeneity: Tau <sup>2</sup> = 1.95; Chi <sup>2</sup> = 10.66, df = 4 (P = 0.03); i <sup>2</sup> = 62% Test for overall effect: Z = 4.32 (P < 0.0001) Total (95% Cl) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); i <sup>2</sup> = 88% -10 -5 0 5 10	Subtotal (95% CI)			150			117	35.5%	3.46 [1.89, 5.03]	-
Total (95% CI) 493 370 100.0% 3.84 [2.46, 5.21] Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); I <sup>2</sup> = 88%	Heterogeneity: Tau² = Test for overall effect:	= 1.95; C Z = 4.32	hi² = 11 ? (P < 0	0.66, df 1.0001)	'= 4 (P =	= 0.03)	; l² = 62	?%		
Heterogeneity: Tau <sup>2</sup> = 5.68; Chi <sup>2</sup> = 106.68, df = 13 (P < 0.00001); l <sup>2</sup> = 88%	Total (95% CI)			493			370	100.0%	3.84 [2.46, 5.21]	•
-10 -5 0 5 10	Heterogeneity: Tau <sup>2</sup> =	5 68° C	hi <sup>2</sup> = 11	16 68	f = 13 (	′P < ∩ I	100011	12 = 88%		-+-+-+++
Test for overall effect: $7 = 5.45$ (P < 0.00001)	Tast far sus all affact	7-545	(P < 1	00001	)			. = 0070		-10 -5 0 5 10

**Figure 6.** Forest Plots Showing the Subgroup Analysis of Treatment With Different Pulse Numbers Note. SD = standard deviation; IV = inverse variance; CI = confidence interval; df = degrees of freedom.

and 3 months. However, because only one RCT mentioned the follow-up results after 12 months, the longterm effect of LI-ESWT still needs further follow-up investigation.

In the process of data extraction, some reports reported neither IIEF final average data and standard deviation nor IIEF change data. Instead, they provide data such as interquartile range (IQR), sample median, and sample size. For the consistency and comparability of statistical data, the sample mean and standard deviation were estimated using the methods provided by the researchers (Luo et al., 2018; Wan et al., 2014).

The minimum clinically important difference (MCID) is considered to be an ideal method to evaluate the real clinical efficacy of interventions. It has been determined that the MCID in IIEF score is 4 points, indicating a 4 points difference may be clinically significant to patients (Rosen et al., 2011). For the trials included in this study, the comprehensive improvement of IIEF score in some groups after LI-ESWT treatment is less than 4 points, which may not have clinical value. With the publication of more and more RCTs, MCID is very important as an evaluation standard. Therefore, it is recommended to use MCID as an accurate and meaningful tool for evaluating LI-ESWT treatment in the future.

Compared with previous meta-analyses, our study excluded studies with high heterogeneity and included many latest studies, which is more convincing. Although the articles included in this meta-analysis are high-quality RCTs, there are still some limitations as follows: (a) Some experiments did not use double-blind research in the research process, and some patients withdrew from the research because they could not tolerate the intervention measures. These bias factors will affect our final research results. (b) Some experimental data only provide median and IQR, so we must use formulas to convert them into mean and standard deviation, and there may be some errors in this process. (c) Because most study endpoints were evaluated only 1 to 6 months after treatment, we could not infer the long-term efficacy of LI-ESWT treatment. (d) Our study did not report other indicators to evaluate ED, such as quality of sexual life, peak wholebody velocity, and resistance index because only one or two RCTs reported these indicators, and the results were not convincing. Therefore, further research and relevant data are needed to help us demonstrate the impact of LI-ESWT treatment on these indicators. (e) We did not assess the potential impact of age, hypertension, diabetes, hyperlipidemia, and coronary artery disease on IIEF.

# Conclusion

This meta-analysis that contains 16 RCTs identified that LI-ESWT could significantly increase IIEF and EHS in ED patients, especially in moderate ED group, but had no significant improvement in positive response rate of SEP2 and SEP3. In general, LI-ESWT has become a popular choice for the treatment of ED because of its effectiveness and low risk, but more clinical experiments, longer follow-up, and more detailed data are still needed to support this conclusion.

#### Author contributions

J.M. and Z.Z. contributed to conceptualization; H.Y., H.L., F.S., and X.B. contributed to data curation; H.Y., X.W., G.T., and X.B. contributed to formal analysis. J.W. and J.M. contributed to funding acquisition. H.Y., X.W., H.L., and F.S. contributed to investigation. H.Y., X.W., H.L., F.S., and G.T. contributed to methodology. Z.Z. and J.M. contributed to project administration. F.S., G.T., and X.B. contributed to resources. H.Y. and X.W. contributed to software. G.T., X.B., and J.W. contributed to supervision. H.Y., X.W., and H.L. contributed to writing—original draft. Z.Z. and J.M. contributed to writing review & editing.

#### **Declaration of Conflicting Interests**

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

#### Funding

The author(s) disclosed receipt of the following financial support for the research, authorship, and/or publication of this article: This work was supported by grants from the National Nature Science Foundation of China (Nos. 81870525 and 81572835) and Taishan Scholars Program of Shandong Province (No. tsqn201909199).

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#### Supplemental Material

Supplemental material for this article is available online.

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