

# Mitigation of Postsurgical Scars Using Lasers: A Review

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**Background:** Most postsurgical scars are considered esthetically and functionally acceptable. Currently, there is no definite consensus treatment for postsurgical scarring. The purpose of this review is to shed some light on the value of scar mitigation and the efficacy of different lasers employed on postsurgical wounds.

**Methods:** A systematic literature review and computational analysis were conducted to identify relevant clinical articles that pertained to the use of lasers for mitigating postsurgical scars. Articles included the National Institutes of Health–National Center for Biotechnology Information–PubMed search and sources cited from relevant studies after 1995. Trials that attributed pre- and posttreatment scores of scar severity based on a verified scar evaluation scale (eg, Patient and Observer Scar Assessment Scale, Vancouver Scar Scale, Global Assessment Scale) were chosen. Clinical assessments varied for each study. To adequately assess the efficacy of the modalities, the final scaled scar appearance scores were realigned and normalized to a standard scale for unbiased comparison.

**Results:** After filtering through a total of 124 studies, 14 relevant studies were isolated and thus included in the review. Studied lasers were as follows: Pulsed dye laser (PDL), carbon dioxide, diode, potassium titanyl phosphate (KTP), and erbium glass (Er-Glass) lasers.

**Conclusion:** Treatment with lasers in the postsurgical wound healing phase is safe, effective, and advised in mitigation of pathologic scar formation. (*Plast Reconstr Surg Glob Open* 2020;8:e2746; doi: [10.1097/GOX.0000000000002746](https://doi.org/10.1097/GOX.0000000000002746); Published online 24 April 2020.)

## INTRODUCTION

Most postsurgical scars are considered esthetically and functionally acceptable. However, keloidal or hypertrophic scars can be symptomatic and might cause esthetic, psychological, and social distress.<sup>1</sup> There is currently no definite, consensual measure that completely prevents postsurgical scarring. Therefore, many modalities are being explored, including laser treatment, to determine the safest and most efficacious method.<sup>2</sup>

The purpose of this review is to shed some light on the value of active scar mitigation by lasers, as compared to natural, spontaneous wound healing concerning final cosmesis of scars.

## METHODS

### Search Strategy

Studies were identified through the search strategy by 2 independent reviewers.

A systematic National Institutes of Health–National Center for Biotechnology Information–PubMed search was conducted to identify relevant clinical articles that pertained to the use of lasers to mitigate postsurgical scars. Message-Subject-Headings were applied. The search algorithm used was (cicatrix OR cicatrix treatment OR scar OR scars) AND (laser OR laser treatment OR laser therapy OR fractional laser OR ablative fractional laser OR nonablative fractional laser OR dye laser OR diode laser) AND (prevention OR minimizing OR early intervention OR treatment outcome). Additional studies were acquired from review articles that appeared in the search. Only human trials that attributed pre- and posttreatment scores of scar severity based on a verified scar evaluation

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scale [eg, Patient and Observer Scar Assessment Scale, Vancouver Scar Scale (VSS), Global Assessment Scale], published in English, after January 1, 1995, were finally included.

The final studies that fit the inclusion criteria were identified using a 2-step process. First, the titles and abstracts of acquired articles were screened. Next, the complete text was reviewed. The inclusion and exclusion criteria are shown in [Figure 1](#).

#### Data Extraction

Standardized extraction of data was compiled and consolidated in a Microsoft Excel (Microsoft, Redmond, Wash., USA) spreadsheet. Acquired data included the name of the first author, year of publication, type of surgical wound, time of first treatment relative to operation date, device(s) and medications used, number of compared treatments, number of raters, number of patients subject to each treatment arm, parameters and settings on the device, number of treatments, time interval between treatments, study design, scar evaluation scale, statistical test used, *P* value, results, comments, and clarifications for future considerations.

#### Outcome Measures, Data Normalization, and Analysis

Different studies use different clinical assessment scores. To adequately assess the efficacy of the modalities, the final scaled scar appearance scores were realigned and normalized to a standard scale for unbiased comparison. This excluded studies that only reported improvement of scars and studies that did not adequately report their final results. Scales such as the VSS, the Global Assessment Scale,

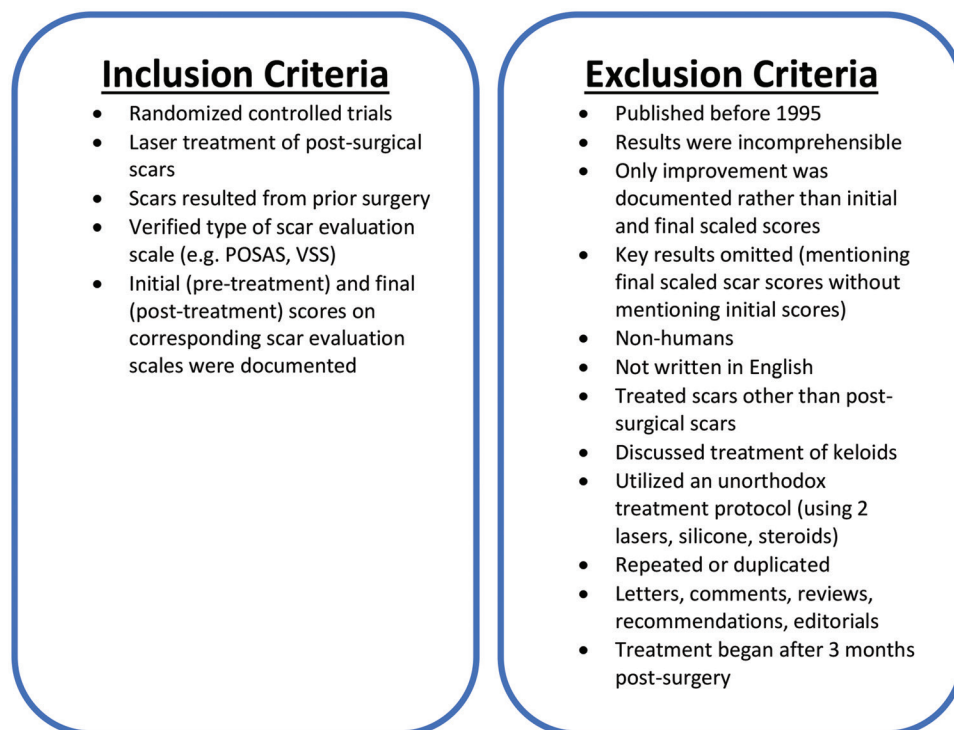
the Patient and Observer Scar Assessment Scale, Visual Analog Scale, and 4- and 5-level Likert scales were normalized to a standard 0–100 scale depicting 100 as healthy skin (= best esthetic outcome) and 0 as the worst possible scar (= worst esthetic outcome). Objective scar scores were defined as those determined by physicians, and subjective scar scores were those determined by patients.

The aligned scores were then compared: treatment versus control to calculate standard score, or *z* score, as it pertains to a single measurement, whereas standard mean difference (SMD) compares 2 groups of measurements: treatment versus control. The SMD was calculated for both objective and subjective ratings. Studies that employed more patients were weighted heavier against studies that employed fewer patients.

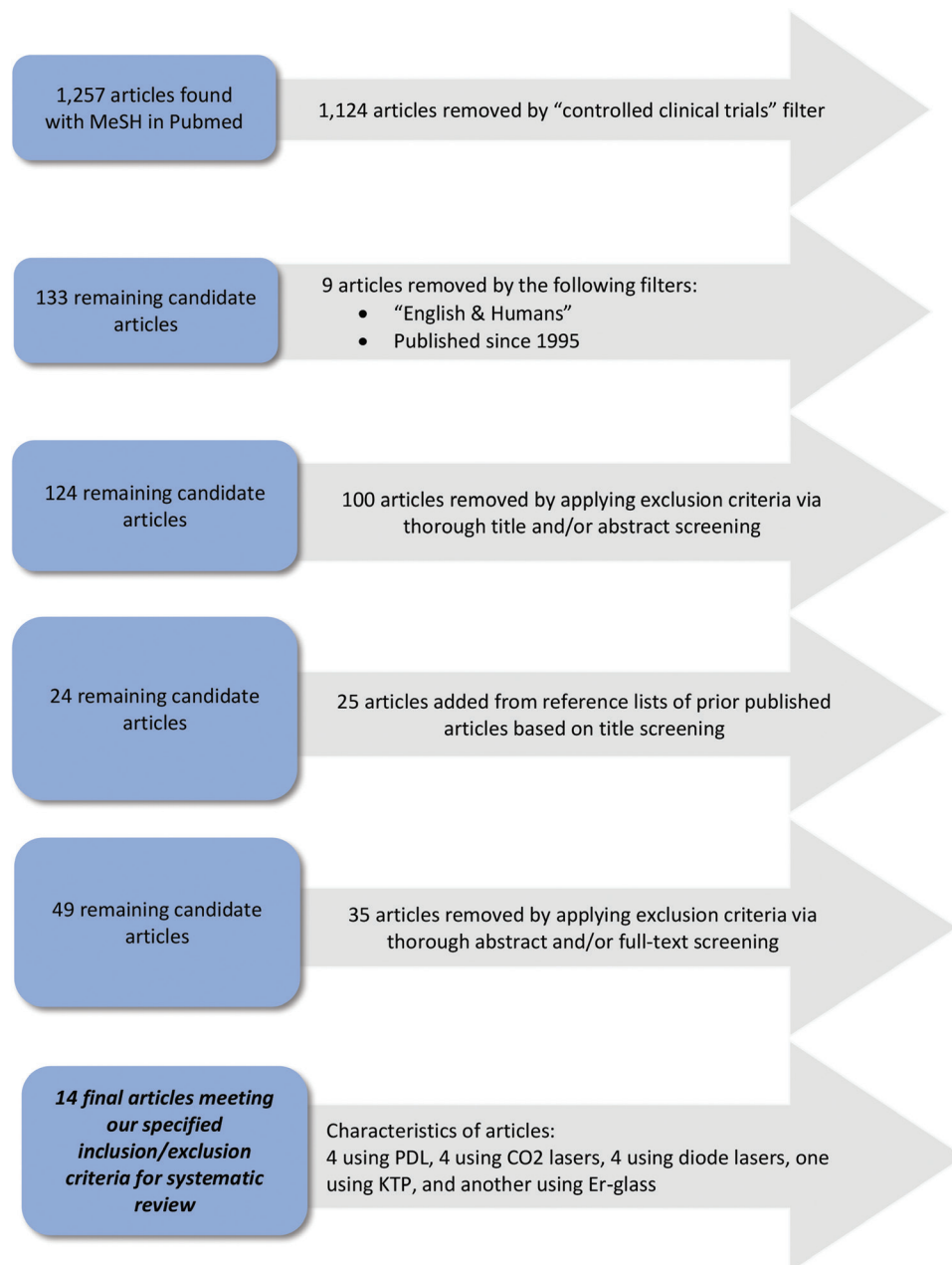
## RESULTS

The initial database search yielded 124 studies. Twenty-five studies were added from reference lists of reviews and filtered using the inclusion and exclusion criteria specified above.<sup>3,4</sup> One hundred articles were excluded following title and abstract review. Thirty-five studies were excluded following full-text screening. A total of 14 studies remained and discussed in this review ([Fig. 2](#)).

A summary of the study characteristics is depicted in [Table 1](#). Fourteen studies met the inclusion criteria of attempting to treat postsurgical scars with lasers and assessed the quality of scar healing with a final scar assessment score.<sup>5–18</sup> Nine studies used split scars, 2 used controlled cohorts, 1 used a split body (breasts) comparison, and 2 were prospective pilot studies. A total of 271 scars were



**Fig. 1.** Inclusion and exclusion criteria. POSAS, Patient and Observer Scar Assessment Scale.



**Fig. 2.** Schematic for study selection. CO<sub>2</sub>, carbon dioxide. Er, erbium.

treated with 247 matched control scars. The number of patients per study group ranged from 5 to 40 (mean = 18.5). Pulsed dye lasers (PDL), carbon dioxide (CO<sub>2</sub>) lasers, and diode lasers were the most commonly used devices followed by erbium glass (Er-Glass) and potassium titanyl phosphate (KTP) lasers.

Various protocols were employed; laser treatments were performed 2–10 times at 2- to 10-week intervals. Most study protocols included 3–4 treatments at intervals of 2–4 weeks. For all but one study,<sup>16</sup> the first laser treatment was scheduled after suture removal.

The VSS was used in 8 of the 15 studies. Most of the studies used multiple scales for each objective (physician) and subjective (patient) evaluations of the outcomes of

scars. The objective SMD between treatment and control for all cumulative studies was calculated to be 0.777 (95% CI, 0.368–1.186). Statistically significant differences were measured between treated and untreated scars among physicians and patients.

Diode lasers treatment lead to the greatest SMD of 0.624 (95% CI, 0.322–0.925) and CO<sub>2</sub> the lowest overall SMD with 0.43 (95% CI, 0.0794–0.780). The SMD for KTP, 2.669 (95% CI, 1.558–3.779), and Er-glass lasers, 0.876 (95% CI, 0.194–1.557), had the best results compared with controls, however, with low weights and a wide CIs. Objectively, the SMD of PDL, 0.45 (95% CI, 0.116–0.784), was similar to that of CO<sub>2</sub>. These results are shown in [Figure 3](#).

**Table 1. Study Characteristics**

Article	Device	Type of Control	No. Patients		Objective Scale
			Treated	Control	
Nouri et al <sup>5</sup>	PDL	Split scar	12	—	VSS
Alam et al <sup>6</sup>	PDL	Split scar	17	—	1–4 scale
Conologue and Norwood <sup>7</sup>	PDL	Split scar	13	—	VSS
Capon et al <sup>8</sup>	Diode	Split scar	5	—	1–4 scale
Choe et al <sup>9</sup>	Er-glass	Separate group	27	14	VSS
Capon et al <sup>10</sup>	Diode	Split scar	30	—	0–3 scale
Carvalho et al <sup>11</sup>	Diode	Separate group	14	14	VSS
Yun et al <sup>12</sup>	KTP	Separate group	20	8	VSS
Lee et al <sup>13</sup>	CO <sub>2</sub>	Split scar	15	—	VSS
Sobanko et al <sup>14</sup>	CO <sub>2</sub>	Split scar	20	—	VSS
Vazquez-Martinez et al <sup>15</sup>	PDL	Split scar	30	—	VSS
Buelens et al <sup>16</sup>	CO <sub>2</sub>	Split scar	9	—	GAS, POSAS
Alberti et al <sup>17</sup>	CO <sub>2</sub>	Separate group	20	21	VSS
Casanova et al <sup>18</sup>	Diode	Split chest	40	—	mOSAS

In split scar studies, the number of treated patients equals the number of controls.

GAS, Global Assessment Scale; mOSAS, modified Observer Scar Assessment Scale; POSAS, Patient Observer Scar Assessment Scale.

Patients’ evaluations reflected the same results, as shown in Figure 3C. The SMD for diode lasers was 0.844 (95% CI, 0.275–1.413). KTP lasers reflected an SMD of 1.868 (95% CI, 0.88–2.848) in a few patients. CO<sub>2</sub>, with an SMD of 0.353, had statistically insignificant results among patients (95% CI, –0.483 to 1.189).

### DISCUSSION

Postsurgical scars are ideally flat, narrow, pale, and pliable. Abnormal scars may range from hypertrophic to keloid. Abnormal scars can be unsightly, painful, functionally, and socially limiting. Various scar mitigation options exist.<sup>19</sup> This study provides a systematic review of trials performed over the last 2 decades that investigate mitigation of postoperative scars using a variety of single laser modalities and treatment protocols.

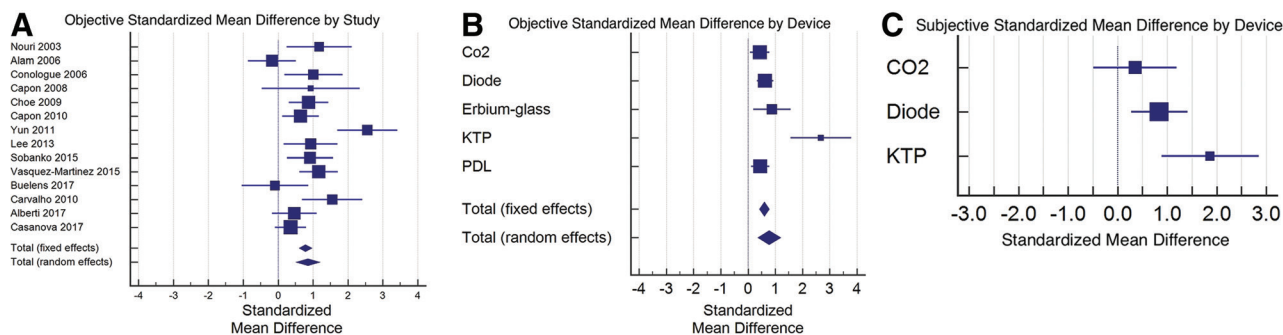
Four studies out of 14 did not demonstrate statistical significance with the treatment of postsurgical scars. Alam et al<sup>6</sup> employed a single PDL treatment immediately postsurgery and concluded that more treatments are necessary to achieve a therapeutic effect. Buelens et al<sup>16</sup> and Sobanko et al<sup>14</sup> failed to show statistically significant improvement using fractional CO<sub>2</sub> laser monotherapy. However, higher patient satisfaction was statistically significant for treated scars.<sup>16</sup>

Statistically, significant scar improvement was found in the remaining 10 studies. Diode, PDL, and CO<sub>2</sub> lasers were reported to have the best results compared with controls. Although the exact mechanics are not entirely understood, diode lasers have been shown to increase heat-shock-protein 70 induction, which is known to induce collagen proliferation and modulate transforming growth factor beta (TGF-β) expression, and remodeling.<sup>8,20,21</sup>

PDL and KTP lasers best target the oxy- and deoxyhemoglobin chromophores and are best utilized to alleviate the erythema associated with highly vascular postsurgical scars.<sup>22–25</sup> Also, PDL has been shown to upregulate p53, inhibiting cell proliferation, and reducing angiogenesis that contributes to abnormal scarring.<sup>26</sup> Optimal results were achieved after 3 PDL treatments.<sup>5–7,15</sup> Treatment intervals ranged from 2 to 10 weeks. All 4 studies started treatment on the day of suture removal and proved to be safe and efficacious.<sup>5,27</sup>

Both KTP and PDL lasers exhibited substantial improvement in similar categories of scar treatment; however, neither have had statistically significant results when compared with the other.<sup>28</sup> KTP lasers have been associated with more posttreatment pain, erythema, and edema.<sup>28,29</sup>

The chromophore targeted by CO<sub>2</sub> lasers is water, found in tissues. CO<sub>2</sub> laser treatments are considered



**Fig. 3.** Graphical depictions of standardized mean differences between treatment and controls among (A) objective results achieved by the studies, (B) objective results achieved by specific devices, and (C) subjective results achieved by specific devices. The box size of each data point depicts the weight of the device, which was deemed dependent on the number of patients treated. The whiskers of each box depict the standard error and thus the 95% CI.



more aggressive and lead to considerable dermal matrix remodeling, hopefully leading to favorable remodeling of the scar.<sup>25,30</sup> Half of the studies exploring the treatment of postsurgical scars using only CO<sub>2</sub> lasers reported statistically insignificant results with a collectively low SMD.

Combining different modalities may augment scar mitigation. A synergistic effect can be achieved from combining PDL treatment, targeting scar vascularity and pigmentation, followed by fractional CO<sub>2</sub>, and aimed at improving the texture, pliability, and height of the scar.<sup>31–33</sup> Also, combination of laser treatment with triamcinolone injection has shown promising results, perhaps by inhibiting fibroblasts and TGF-β.<sup>25,31,34–36</sup> Combining Eryttrium aluminum garnet (YAG) fractional ablation after PDL treatment improved scar pliability in addition to scar appearance.<sup>37</sup>

Our review is inherently limited by its attempt to compare different studies using various measurement scales and patient populations. This made it difficult to compare among protocols, devices, and parameters. Also, aside from 2 listed patients in one study,<sup>7</sup> only patients less prone to hypertrophic or keloid scarring are evaluated. Future studies should explore high-risk patients such as patients with abnormal scarring history or patients undergoing midline or limb incisions. Moreover, standardized objective measuring tools such as 3D, infrared cameras, and standard scar scales should be encouraged.

## CONCLUSIONS

Laser therapy is a safe and effective modality for scar mitigation. The data suggest that early intervention exhibits the best results. We recommended beginning treatment close to suture removal combining vascular and nonablative fractional resurfacing modalities for 2–4 treatments at 2- to 3-week intervals. It is imperative to monitor the wound healing process and document side effects such as erythema, discoloration, pain, and infection. Further research may help define standard treatment protocols which would benefit both evaluations of the results and clinical outcome.

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