

Is Laser Disinfection an Effective Adjunctive Treatment to Bone Augmentation for Peri-Implantitis? A Review of Current Evidence

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Focused Clinical Question:

In patients with endosseous dental implants that demonstrate peri-implantitis, does surgical bone augmentation with adjunctive laser implant surface disinfection have an effect on implant survival rates, and do these rates differ based on laser treatment modality?

Clinical Scenario:

A 55-year-old female presents 10 years after implant placement at sites #18 and #20 (Fig. 1). She demonstrates a 9-mm probing depth mesially and distally at implant #20. Bleeding on probing is present at all six sites around implant #20. The patient has not noted any discomfort, and suppuration has not been noted on clinical examination. Her medical history is significant for osteoarthritis, gastroesophageal reflux disease, and anxiety. She reports taking ibuprofen as needed for pain, 150 mg ranitidine twice daily, and 20 mg citalopram daily. The patient is concerned about the possibility of implant loss and states that she wants to save and treat the implant, if possible. During flap reflection, a circumferential defect at implant #20 is noted intrasurgically (Fig. 2).

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Key Words: Alveolar bone grafting; dental implants; lasers; peri-implantitis; periodontal diseases; review.

Background

A large number of dental implants are placed currently, and although they have been shown to be a predictable treatment modality to replace missing teeth,¹ reports indicate that up to 12% to 43% of dental implants will develop clinical signs and symptoms of peri-implantitis and crestal bone loss over 5 or more years after functional loading.^{2,3} Treatment strategies for management and treatment of peri-implantitis have been varied, and there is little evidence of treatment efficacy.⁴ A consensus report of the 6th European Workshop on Periodontology stated that non-surgical therapy of peri-implantitis has not been found to be effective.⁵ Conversely, surgical outcomes with guided bone regeneration (GBR) have demonstrated improved clinical and radiographic outcomes for peri-implantitis treatment.⁶⁻⁹ The adjunctive use of laser and photodynamic therapy (PDT) has also been considered to improve implant surface decontamination with both surgical and non-surgical therapy.¹⁰⁻¹²

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Search Strategy

The following terms were searched in the PubMed database (in which mh is the MeSH term, tw is the text word, pt is the publication type, and sb is the subset): Peri-Implantitis OR peri-implantitis OR perimplantitis OR peri-implant OR peri-implant OR perimplant OR (Dental Implantation OR Dental Implants) AND (Prosthesis-Related Infections OR Alveolar Bone Loss OR Periodontal Diseases OR Periodontitis OR Gingivitis OR Pericoronitis OR infection OR infections OR microbiology OR Bacteria OR Dental Plaque OR Mucositis OR Stomatitis OR Periodontal Debridement OR Debridement OR debrid* OR Dental Scaling OR Dental Air Abrasion OR “air abrasive” [tw] OR Dental Disinfectants OR Dental Etching OR Dental Prophylaxis OR Disinfection OR Disinfectants OR disinfect* OR Decontamination OR decontaminat* OR Prosthesis Failure OR Dental Restoration Failure) AND Laser Therapy OR Lasers OR laser OR Er-YAG OR eryag [tw] OR erl [tw] OR Nd-YAG OR ndyag [tw] OR photodynamic OR Photochemotherapy OR semiconductors OR diode OR co2 OR “co 2” [tw] OR Carbon Dioxide AND (cohort studies [mh] OR meta-analysis [pt] OR meta-analysis as topic [mh] OR randomized controlled trial [pt] OR randomized controlled trials as topic [mh] OR systematic [sb] OR cohort [tw] OR “meta analysis” [tw] OR “randomized controlled” [tw] OR systematic [tw]).

Search Outcome

Ninety-six abstracts were hand reviewed, and 19 full-text articles were reviewed. Four articles are included in this review. Original human participant case series, cohort, and randomized controlled trials were included. Papers were eliminated for the following reasons: 1) study design; 2) lack of published data on peri-implantitis treatments; 3) non-unique patient populations; and 4) surgical access only with laser debridement. Data for guided tissue regeneration (GTR) with laser implant surface debridement were extracted from datasets if the following were true: 1) additional treatment modalities were included in the manuscripts; 2) findings were reported; and 3) those specific data are reported in this report. The findings of the relevant current literature¹³⁻¹⁶ are reviewed in Table 1.

Discussion

Commercially available lasers that have been used for implant surface disinfection include carbon dioxide (CO₂), diode, erbium:yttrium-aluminum-garnet (Er:YAG), and

neodymium:yttrium-aluminum-garnet (Nd:YAG). Current evidence demonstrates inconsistent data regarding laser reduction of bacterial loads on tooth surfaces beyond that achieved with non-surgical periodontal therapy alone.^{17,18} However, in vitro and in vivo trials have demonstrated elimination of bacterial smear layer and viable bacteria with the use of laser therapy.¹⁹⁻²² Although non-surgical treatment of peri-implantitis with the adjunctive use of laser treatment was not found to be effective in a previous review,⁵ surgical access with surface debridement and bone augmentation has been shown to be effective.^{6,7} Variability within the treatment outcomes may be influenced by implant surface characteristics, bone graft substitute qualities, and defect characteristics.^{8,23}

One article reviewed¹⁶ was a randomized controlled trial, but the manuscript only presented clinical data at baseline and follow-up. Surgical reentry data were available in one article reviewed,¹⁴ although patient assignment to treatment groups was not stated to be randomized in this study. A total of 86 implants were treated with GBR and laser implant surface decontamination in all articles reviewed, and the total follow-up time ranged from 0 to 60 months. Considerable heterogeneity exists in evaluation procedures for resolution of peri-implantitis in the articles reviewed, and none of the articles evaluated microbiologic outcomes. All articles reviewed demonstrated an improvement in clinical and/or radiographic outcomes from baseline to follow-up for failing implants treated with laser and GBR protocols. Furthermore, the two studies that demonstrated comparative results for implants treated with GBR without laser debridement demonstrated improved clinical and/or radiographic outcomes for the laser decontamination group when compared with standard decontamination.^{14,16} Because the articles reviewed used varied laser protocols, including CO₂,^{14,15} Er:YAG,¹⁶ and soft light laser at 906 nm,¹³ no meaningful conclusions can be made about the effectiveness of individual laser types for decontamination purposes from these studies. In vitro and animal studies have suggested that low-level laser with dye may improve implant surface disinfection.^{12,24,25} Furthermore, in vitro studies have shown that diode, CO₂, Er:YAG, and Nd:YAG lasers at appropriate wavelength settings may be sufficient to reduce and/or eliminate surface bacteria without damaging implant surface characteristics if appropriate published parameters are used.^{26,27}

Implant failure has been classified into clinically distinct types with differing microbiota based on the underlying etiology.²⁸ Early implant failure may be associated with lack of primary stability, surgical trauma, or postoperative infection, whereas late failure has been shown to be associated with occlusal overload and peri-implantitis.^{29,30} In one article reviewed,¹⁵ 15 of the 19 implants treated demonstrated bone loss before Phase II uncover procedures and prosthetic restoration. These implants may represent a different subset of microbiota than those with late and/or infection-related failures and may respond differently to therapy. Evaluation of specific microbiota associated with peri-implantitis, specifically *Tannerella forsythia*, *Campylobacter*

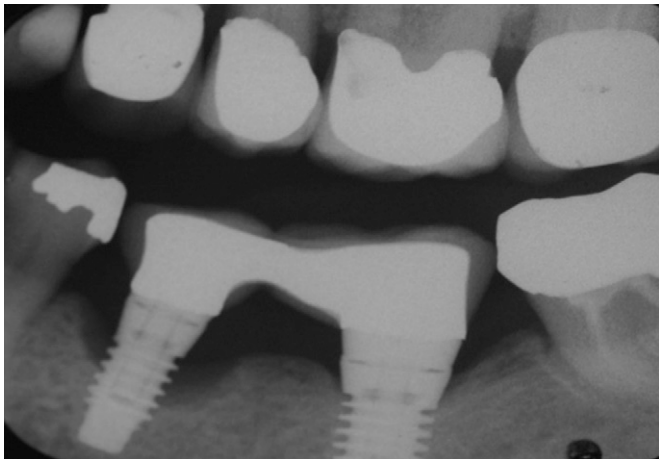


FIGURE 1 Bitewing radiograph demonstrating radiographic bone loss of 40% to 60% of implant length at both the mesial and distal surfaces of implant #20.



FIGURE 2 Intrasurgical clinical photograph taken after debridement demonstrating a circumferential bony defect at implant #20.

TABLE 1 Adjunctive Laser Use With Bone Grafting Around Implants: A Summary of the Published Literature

Reference	Implant Data	Laser Type	Study Type (Level of Evidence)	Methods	Key Results	Comments
Haas et al., 2000 ¹³	24 flame-sprayed cylindrical implants.	Soft light laser at 906 nm for 1 minute after toluidine blue O (100 µg/mL) treatment for 1 minute; preprocedural saline rinse.	Case series	Patients with peri-implantitis were defined by an infrabony pocket of ≥6 mm and progressive bone loss over the past 12 months. Sites were accessed surgically, and the implant surfaces were treated with toluidine blue O (100 µg/mL), soft light laser at 906 nm, and bone augmentation with autogenous bone and ePTFE membrane. Membrane removal was scheduled 3 months after placement but was removed immediately if exposed and demonstrating signs of inflammation. Radiographic bone gain was evaluated at 9.5 months postoperatively.	Two implants failed and were explanted. Mean defect depth decreased statistically significantly from preoperative levels. Defect depths preoperatively were 5.5 ± 2.0 mm, and a statistically significant mean bone gain of 2.0 ± 1.9 mm was seen over the study period. Early membrane retrieval was associated with decreased bone gain.	A large percentage of membranes were exposed early, and this may have altered treatment effectiveness. Furthermore, there was no control group that did not receive adjunctive implant laser disinfection, so the additive benefit of laser treatment is unable to be determined.
Deppe et al., 2007 ¹⁴	32 implants; 17 received bone augmentation and laser debridement, 15 received traditional disinfection and bone augmentation. Resective surgical treatment with or without laser disinfection was performed in the other groups. Two implant types were included.	CO ₂ laser at 10.6 µm wavelength and a maximum of 7 W. No irrigation was noted.	Case series	Patients who were identified with progressive vertical bone loss, PD ≥5 mm, or BOP received surgical access, bone augmentation treatment using β-tricalcium phosphate and autogenous bone with laser or traditional (cotton pellet, saline, plastic curet) implant surface decontamination. Non-resorbable membranes were used. Surgical reentry and 5-year follow-up were performed.	Implants receiving laser decontamination demonstrated statistically significant improvements in CAL when compared with those receiving conventional disinfection at reentry. Statistically significantly more vertical bone height (DIB) was demonstrated in laser-debrided implants at surgical reentry.	All implants were treated with air-powder abrasive disinfection, and, in 22, adjunctive CO ₂ laser treatment was performed. Although the differences between laser-debrided and non-debrided groups receiving bone augmentation are clinically significant, both bone augmentation treatment modalities demonstrated more vertical bone height and CAL gain than implants receiving resective therapy.

TABLE 1 (Continued) Adjunctive Laser Use With Bone Grafting Around Implants: A Summary of the Published Literature

Reference	Implant Data	Laser Type	Study Type (Level of Evidence)	Methods	Key Results	Comments
Romanos and Nentwig, 2008 ¹⁵	19 implants; all received laser debridement and grafting with either autogenous or xenograft material. Multiple implant types were included.	CO ₂ laser at 2.84 ± 0.83 W for 1 minute. No irrigation was noted.	Case series	Patients were identified with deep infrabony defects, bone loss over two thirds of the implant length, and no implant mobility. Four implants were identified after restoration and 15 after initial placement before implant uncover. Surgical access and debridement was performed, and a CO ₂ laser was used to disinfect implant surfaces. Grafting was performed with autogenous or xenograft bone material. Resorbable collagen membranes were used. Implants were observed up to 27 months, and radiographic analysis was performed.	Statistically significant improvements were seen in the sulcus bleeding index and PD postoperatively compared with preoperatively, despite no significant differences in plaque index and width of keratinized gingiva at surgical sites. The number and extent of peri-implant defects with vertical bone loss improved postoperatively as well.	This treatment protocol proved to be effective in increasing radiographic bone fill and decreasing PD in implants with vertical bone loss. The majority of these implants must be considered early failures because they were not exposed to the oral environment before peri-implantitis diagnosis and treatment. Also, this study did not include a control group, so the effect of the CO ₂ laser used for surface disinfection compared with the effectiveness of augmentation procedures is unknown.
Schwarz et al., 2012 ¹⁶	26 implants; all received flap access and surgical debridement. Patients were then randomly assigned to receive adjunctive debridement with Er:YAG laser or standard disinfection before bone augmentation. Various implant types were included.	Er:YAG laser with pulse radiation at 2.940 nm with 100 mJ/pulse and 10 Hz. Continuous water irrigation was used.	Randomized controlled trial	Patients who were identified with infrabony and supracrestal defects with PD >6 mm and infrabony component >3 mm were defined as having peri-implantitis. All patients received surgical access and degranulation of the surgical sites and then were randomized to receive implant surface decontamination with either Er:YAG laser or standard disinfection protocols (cotton pellet, saline, plastic curet). All participants then received bone augmentation	Statistically significant differences were seen at 12-month follow-up within each group from baseline for BOP, PD, mucosal recession, and CAL. At 24 months, intragroup differences in BOP were seen. There were no statistically significant differences between groups at either time point.	Site-level analysis demonstrates a high level of heterogeneity in the responses to both treatment modalities. All patients received additional peri-implantitis treatment after the 24-month observation period was over.

TABLE 1 (Continued) Adjunctive Laser Use With Bone Grafting Around Implants: A Summary of the Published Literature

Reference	Implant Data	Laser Type	Study Type (Level of Evidence)	Methods	Key Results	Comments
				grafting with xenograft bone graft and resorbable collagen membrane. Clinical follow-up was performed to 24 months.		

ePTFE = expanded polytetrafluoroethylene; PD = probing depth; DIB = distance from implant platform to first implant bone contact radiographically.

spp., and *Parvimonas micra*, and evaluation of loss of radiographic lamina dura³¹ in future investigations may allow for more certainty in the diagnosis of late implant failure, i.e., true peri-implantitis.

GBR alone has been used in both animal and human models to treat peri-implant defects.^{8,9,32-37} Furthermore, bone replacement grafting has demonstrated differing success on various implant surface types.³⁷ Bone substitute materials, defect morphology, and implant surface characteristics may all influence clinical outcomes of peri-implantitis treatment with GBR. Participants in the studies reviewed received grafting with autogenous bone, xenograft, and alloplast materials, and numerous implant surface types were included in the treated implant group. The heterogeneity of implant systems and surface types, bone replacement graft materials used, and possible varied etiologies for peri-implant bone loss are confounders when evaluating the current literature.

One study included in this review demonstrated long-term follow-up with a randomized controlled trial design.⁹ Previous data were published on the short-term (6-month) results after Er:YAG laser surface decontamination.²³ Although 6-month data demonstrated higher reduction in bleeding on probing (BOP) and clinical attachment level (CAL) at the laser disinfected sites, at 24 months, there were no statistically significant differences between the groups, and neither group demonstrated CAL levels that differed from baseline findings. These results may indicate that adjunctive laser disinfection may result in short-term decontamination and improved clinical findings, but the longevity of these effects may not allow long-term improvements at implants with peri-implantitis.

There have also been adverse events associated with implant treatment with lasers. Nd:YAG lasers have been shown in some studies to boil additive implant surfaces, yielding porosities, which could serve as a niche for additional bacterial proliferation.^{38,39} Additionally, the use of CO₂ lasers at pulsed and continuous settings has been demonstrated to raise the temperature of implants in vitro 9.5°C to 12.2°C, respectively.⁴⁰ This type of increase in

temperature from physiologic 37°C could yield local circumimplant temperatures in the range of bone necrosis. Furthermore, variability in wavelength, pulse, and irrigation have been shown to have demonstrative effects on decontamination effectiveness and alterations to surface chemistry.⁴¹ Some of the risks of laser therapy may be mitigated by the use of PDT, which uses low-level laser therapy to perform surface decontamination. A recent in vitro study indicated that PDT may be more efficient than standard laser disinfection protocols without many of the associated risks.¹² Therefore, it is very important to note the time, wavelength, presence of cooling, and power of lasers used for peri-implantitis treatment when adapting a published laser treatment protocol for clinical application. ■

Clinical Bottom Line

Adjunctive use of laser disinfection protocols with GTR for the treatment of peri-implantitis may improve clinical and radiographic findings up to 5 years after therapy. Significant heterogeneity exists in implant decontamination protocols. Care must be taken to develop standardized, ideal laser decontamination protocols that fully characterize the wavelength, time, power, and presence of cooling that may yield ideal results for peri-implantitis treatment. Additional longitudinal studies are necessary to allow for the development of standardized protocols for surgical treatment of peri-implant bone loss and clinical inflammation.

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The authors report no conflicts of interest related to this study.

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