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Implantoplasty- provoking or reducing inflammation? – a systematic scoping review

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ABSTRACT

Objectives: To evaluate clinical parameters associated with inflammation after adjunctive implantoplasty in conjunction with surgical treatment of peri-implantitis.

Materials and methods: A systematic literature search was performed in 2 databases until 29. December 2020 to find publications that report on clinical parameters after surgical peri-implantitis treatment which included adjunctive implantoplasty. Clinical studies on implantoplasty reporting on BoP as outcome were included, but other clinical or radiographic outcomes were also considered.

Results: The search resulted in 18 articles that fulfilled the inclusion criteria. The results indicated improvements of BoP and clinical parameters following surgical peri-implantitis treatment with adjunctive implantoplasty.

Conclusions: Within its limits, the findings of the present scoping review indicated that BoP is reduced following surgical peri-implantitis treatment with adjunctive implantoplasty, and that this improvement is in line with surgical peri-implantitis treatment without adjunctive implantoplasty.

ARTICLE HISTORY

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KEYWORDS

Implantoplasty; periimplantitis; implant surface modification; review

Introduction

The use of osseointegrated implants to replace missing teeth is increasing and has become a routine treatment in dentistry. Technical or biological complications may emerge following such treatment, and studies have uncovered a high prevalence of peri-implantitis [1,2]. Peri-implantitis is an inflammatory disease as a result of microbial biofilm accumulation on the implant which in turn affects the soft and hard implant-supporting tissues [3,4].

A number of approaches to treat peri-implantitis have been investigated, and the treatments proposed involve both non-surgical and surgical means. A randomised controlled study demonstrated no difference in bleeding on probing (BoP) following non-surgical mechanical debridement with titanium curets or ultrasonic devices [5]. Laser therapy may reduce BoP compared to mechanical debridement, but otherwise the treatment modes rendered similar clinical outcomes [6]. Reviews by Renvert et al. and Figuero et al. concluded that nonsurgical treatment of peri-implantitis is not effective due to limited clinical improvements and a tendency of disease recurrence [7,8].

With the limited effect of non-surgical therapy, surgical means have been considered for disease resolution. Surgical management provides direct access to the implant, facilitates removal of granulation tissue and access for implant debridement. However, studies have demonstrated modest disease resolution following surgical treatment of peri-implantitis [9–11]. Figuero et al. stated that no surface decontamination is superior to date, and there is currently no consensus on the most effective treatment [8].

Implantoplasty has been a suggested approach in conjunction with surgical treatment of peri-implantitis. Implantoplasty adjunctive to surgical treatment of periimplantitis includes the use of diamond or carbide burs to mechanically modify the implant surface, which includes thread removal and surface smoothening. This adjunctive measure may serve two purposes. The first is an effective removal of biofilm and calcified deposits on the suprabony implant surface, and the second is to render a smooth implant surface, which in turn may reduce bacterial adhesion, growth, and facilitate professional and self-performed oral hygiene. Ideally, this adjunctive treatment may result in an implant surface which impedes bacterial colonisation and facilitates soft tissue adaption. Several in vitro studies have demonstrated that smooth implant surfaces may enhance fibroblast growth compared to rough surfaces [12-15].

Clinical studies have suggested advantageous clinical outcomes following implantoplasty [16–18]. Clinical case reports have demonstrated resolution of peri-implantitis following treatment by open flap debridement with adjunctive implantoplasty [19], lower levels of planktonic microbial growth following implantoplasty [20] and that implantoplasty also can be combined with bone regeneration [21].

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On the contrary, implantoplasty is a treatment which affects the mechanical properties of implants and the procedure may lead to excess metal debris in the surgical site. A recent systematic review on complications following adjunctive implantoplasty reported only a single case of mucosal discolouration and no fractures [22], indicating that complications may be few. It has been suggested that inflammatory cytokines, inflammatory cells and osteoclast activation increase when titanium and metal debris accumulate in the soft tissue [23], which is inevitable during an implantoplasty procedure. A recent in vitro study demonstrated reduced viability of gingival fibroblasts cultured in the presence of implantoplasty debris [24]. It has also been proposed that fibroblasts exposed to titanium particles and debris may induce secretion of pro-inflammatory cytokines which in turn affects the chemotaxis and recruitment of monocytes [25]. This hypothesis may imply an aggravated inflammatory reaction following debris accumulation after implantoplasty.

Considering the contradictory suggestions in the pre-clinical literature that adjunctive implantoplasty may improve clinical parameters but also lead to aggravated inflammatory reactions in the peri-implant tissues, the aim of this study was to review the inflammatory-related clinical outcomes following such treatment.

Materials and methods

The focus question (PICO) in the present review

Does implantoplasty as adjunctive to surgical treatment of periimplantitis lead to a reduced BOP frequency?

The focus question was assessed according to the PICO strategy:

- Population: Patients with peri-implantitis.
- Intervention: Effect of surgical peri-implantitis treatment with adjunctive implantoplasty
- Comparison: Surgical peri-implantitis treatment without adjunctive implantoplasty
- Outcomes: Changes of clinical peri-implant parameters; Bleeding on probing (BoP) (primary outcome); Plaque indices (PI), Pocket probing depth (PPD), Bone level (BL), Implant survival and Clinical Attachment Level (CAL) (secondary outcomes).

Search strategy

The protocol of the present scoping review was not registered a priori, but followed the guidelines of Preferred Reporting Items for Systematic reviews or Meta-Analyses extension for Scoping Reviews (PRISMA-ScR) [26].

A systematic electronic search was performed on Medline (PubMed) and Scopus.

The database Medline was searched with the following keywords:

(periimplant* OR peri-implant*) AND (implantoplasty OR implant surface decontamination OR implant surface debridement OR

implant surface modification OR implant surface detoxification OR implant threads).

An electronic search on Scopus database was performed with the following keyword: 'implantoplasty'.

Publications not found with the specified electronic search were found manually by seeking references from previous publications or by manual search in the mentioned databases. Grey literature searches were not conducted.

The studies were included if they met the following inclusion criteria:

- English language
- Clinical studies in humans
- Subjects treated with surgical peri-implantitis treatment including adjunctive implantoplasty as at least one of the interventions
- Follow-up period of at least 6 months
- Peri-implantitis disease at baseline
- Records of BoP at baseline and at follow-up
- At least 2 subjects included in study
- Titanium dental implants

Studies that did not meet all of the criteria above were excluded.

The search was done by screening titles and abstracts. The extracted articles from abstracts were evaluated after full-text article screening. Full-text articles that met the inclusion criteria were included in the present review. When publications from the same research group described studies with the same subjects/population with follow up in multiple articles, the publications were considered the same study.

Clinical measurements

Recordings of BoP at baseline and follow-up after adjunctive implantoplasty were evaluated. As secondary outcome variables, the clinical parameters; PPD, PI, CAL, BL and implant survival and the radiologic parameter BL, were investigated to map the clinical outcomes of adjunctive implantoplasty treatment.

Results

A total of 913 (794 from PubMed and 119 from Scopus) potentially relevant titles or abstracts were yielded in the electronic search and 5 papers in the manual search. From the electronic and manual search, 39 and 5 papers were screened full text, respectively. Based on the inclusion criteria, 10 papers were excluded after full-text screening (Table 1), 3 of which were duplicates. Of the remaining 34 articles some were duplicates found in the different database searches. After removal of these duplicates, the number of included studies was 18 articles.

The publications from Romeo et al. [16,17] and those from Schwarz et al. comprise the same subjects [18,32–34] but differ in follow-up time and data presented. The study

Table 1. Excluded studies based on inclusion criteria.

Excluded studies	Reason for exclusion
Lozada et al. [19]	The clinical parameter BoP was not presented.
Geremias et al. [20]	The follow up period was less than 6 months.
Thierbach et al. [27]	The study did not present data for adjunctive implantoplasty separately.
Suh et al. [21]	The clinical parameter BoP was not presented.
Pommer et al. [28]	Did not present follow up measurements for BoP after implantoplasty treatment.
Schwarz et al. [29]	The clinical parameter BoP was not presented.
Schwarz et al. [30]	Did not have more than one patient.
Sapata et al. [31]	Did not have more than one patient.

by Ramanauskaite and co-workers [35] included some patients also participating in the studies by Schwarz et al. [18,32–34]. It was not possible to acquire clinical data for participants exclusively in this study upon contact with the authors. After restricting the same subjects in these studies (Romeo et al. [16,17], Schwarz et al. [18,32–34]) the number of included studies with unique study populations was 14.

The studies from Schwarz et al. [18,32-34] and Wang et al. [36] have a randomised controlled study design (RCT), but are randomised with respect to treatment with Er:YAG laser or control treatment in the intrabony aspect of the periimplant defect prior to regenerative therapy. In these two studies, implantoplasty was conducted in the supracrestal compartment in all subjects. In the studies by Nart et al. [37], Matarasso et al. [38], Schwarz et al. [39], Ramanauskaite et al. [35], Galarraga-Vinueza et al. [40], the supracrestal component of the implants was treated with adjunctive implantoplasty, whereas reconstructive treatment was performed in the intrabony aspect following various decontamination procedures. In Romeo et al. [16,17] and Ravida et al. [41], surgical treatment with adjunctive implantoplasty with bone recontouring was compared to surgical treatment with bone recontouring only. Englezos [42] et al. and Bianchini et al. [43,44] performed implantoplasty with some bone recontouring. In Lasserre et al. [45], implantoplasty was performed without bone recontouring and compared to treatment with glycine air-polishing. In Dalago et al. [46], open flap debridement with adjunctive implantoplasty was compared to open flap debridement with and without subepithelial connective tissue graft.

Of the 14 studies with uniqe study populations included in this scoping review, 3 were controlled studies randomised with respect to adjunctive implantoplasty; Romeo et al. [16,17]; Lasserre et al. [45]; Dalago et al. [46], 8 prospective studies [18,32–34,36–40,42,43], 2 retrospective studies [35,44], and 1 retrospective case-control study [41] (Figure 1) (Table 2).

Definition of peri-implantitis

The case definition of peri-implantitis varied among included studies, but most studies included bone loss \geq 2 mm and BoP, and many also PPD > 5 mm (Table 2).

Bleeding on probing (BoP)

BoP was graded differently in the studies. Romeo et al. [16,17] used the mBI [47], whereas the remaining studies

graded BoP dichotomously at four or six sites per implant. BoP was a requisite parameter in most peri-implantitis case definitions, but some studies included a session of non-surgical instrumentation prior to the baseline clinical assessment of BoP [18,32–35,37,38,39,42,45], which therefore could render a baseline BoP of less than 100%. All but one study presented a reduction of BoP in the follow-up visit compared to the baseline (Table 3). Five studies included more than one follow-up time [16–18,32–34,36,45,46], and these demonstrated either sustained low values or further reduction of BoP over time.

Of the studies randomised with and without adjunctive implantoplasty, a significant difference between the control and test group was only observed in the study by Romeo et al. in favour of adjunctive implantoplasty [16,17]. No differences were detected across groups in the study by Lasserre et al. [45] (2020). In the study by Dalago et al. the BoP decreased significantly from baseline to follow-up only in the group with adjunctive implantoplasty, but there was no difference between groups [46]. The retrospective case-control study by Ravida et al. did not find significant differences in BoP between the testand the control group, and moreover, no significant difference of BoP from baseline to follow-up was observed in the adjunctive implantoplasty group nor the control group [41].

Suppuration (SoP)

Seven studies reported specifically on suppuration in addition to BoP [37,40–45]. (Table 4). The only RCT recording SoP [45], demonstrated no difference between the groups with no significant reduction at 6 months. In Ravida et al., SoP was only significantly reduced in the adjunctive implantoplasty group from baseline to follow-up, but no significant difference was found as compared to the control group [41].

Periodontal probing depth (PPD)

All but one included study recorded PDD [44] (Table 5).

For the studies with multiple follow-up examinations, the mean PPD either did not change considerably after the first follow-up [16,17], or increased slightly over time [18,32–34,46].

The RCT study by Romeo et al. demonstrated significant PPD reductions in both groups [16,17], but significantly more in the group that received adjunctive implantoplasty. In the RCT by Lassere et al. the PPD change was not different between groups [45]. In the study by Dalago et al. [46], there was a significant reduction in the adjunctive implantoplasty group and in one of the two control groups from baseline to follow-up, but no differences between groups. Ravida et al.



Figure 1. Flow-chart of the literature search.

Table 2. Included studies in the present levie	Table 2.	Included	studies	in	the	present	review
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Included study	Study type	Patients	Implants	Peri-implantitis definition
Romeo et al. [16]	RCT	10	19	ВоР
				PPD > 4mm
				BL (not specified)
Romeo et al. [17]	RCT	10	20	BoP
				PPD > 4mm
				BL (not specified)
Dalago et al. [46]	RCT	8	-	BoP
				PPD > 5mm BL > 2mm
Lasserre et al. [45]	RCT	16	22	BoP/suppuration PPD \geq 5 mm
				$BL \ge 2 mm$
Galarraga-Vinueza et al. [40]	PROSPECTIVE	20	28	BoP
				$PPD \ge 6 mm BL \ge 3 mm$
Ramanskauite et al. [35]	RETROSPECTIVE	39	57	BoP
				BL > 2mm
Ravida et al. [41]	RETROSPECTIVE	19	30	Signs of inflammation BoP (suppuration)
				increased PPD (recession of mucosal margin) BL
Wang et al. [36]	PROSPECTIVE	24	24	BoP/suppuration PPD \geq 5 mm
				$BL \ge 2 mm$
Schwarz et al. [32]	PROSPECTIVE	30	35	PDD > 6mm BL > 3mm
Schwarz et al. [33]	PROSPECTIVE	24	26	PPD > 6mm BL > 3mm
Schwarz et al. [34]	PROSPECTIVE	21	21	PPD > 6mm BL > 3mm
Schwarz et al. [18]	PROSPECTIVE	15	15	PPD > 6mm BL > 3mm
Schwarz et al. [39]	PROSPECTIVE	10	13	PPD > 6mm BL > 3mm
Bianchini et al. [44]	RETROSPECTIVE	23	32	BoP
				$PPD \ge 6 mm BL \ge 3 mm$
Bianchini et al. [43]	PROSPECTIVE	4	4	BoP Suppuration PPD $>$ 5mm BL $>$ 3mm
Englezos et al. [42]	PROSPECTIVE	25	40	BoP
				$PPD \ge 6 mm BL \ge 3 mm$
Matarasso et al. [38]	PROSPECTIVE	11	11	BoP
				PPD > 5mm BL > 2mm
Nart et al. [37]	PROSPECTIVE	13	17	BoP/suppuration PPD $>$ 5mm BL $>$ 3mm

Only patients/implants receiving adjunctive implantoplasty included. Data not available (-).

reported a significant PPD reduction in both test- and control groups from baseline to follow-up, but no difference between the groups [41].

Plaque (PI, mPI, PI)

Plaque was recorded in all but three included studies [35,43,44]. Matarasso et al. reported on full-mouth plaque scores only [38].

The studies from Schwarz et al. [18,32–34,39], Lasserre et al. [45], Galarraga-Vinueza et al. [40] used PI [48]. Nart et al. [37] used the index from ÓLeary et al. [49]. Romeo et al. [16,17] and Dalago et al. [46] used the modified plaque index mPI [47]. Other studies reported plaque dichotomously [37,42].

Studies with several follow-up measurements reported decreasing plaque levels throughout the observation period [16–18,32–34,39], and one study reported an initial decrease followed by a slight increase at the 3-year follow-up [46].

Table 3. BoP from	baseline to post	t-operative m	ieasurements.												
Study	Baseline BoP	3 M	Δ 3 M	6 M	Δ6 M	12 M	Δ12 M	24 M	Δ 24 M	36 M	A 36 M	48 M	Δ 48 M	5+ Υ	Δ 5+ Y
Romeo et al [16.17]	2.8 (2.9)			0.6 (2.2)	2.2 (0.7)	0.4 (2.7)	2.5 (0.2)	0.5 (2.3)	2.3 (0.5)	0.6 (-)	2.2				
Dalago et al. [46]	4.5 (3.6, 3)					0.5 (1.7, 1.7)	4 (1.9)	0.5 (2.1, 2.3)	4 (1.5, 1.2)	1.3 (2.7, 2.4)	3.4 (0.9, 1.2)				
Lasserre et al. [45]	94.7 (88) (%)	33.4 (30.8)	61.3 (49.2)	33.3 (26.3)	61.2 (61.7)										
Galarraga-Vinueza	65 (%)			16	49										
et al. [40]															
Ramanauskaite	100 (%)														54.9 (75)
et al. [35]															
Ravida et al. [41]	88.9 (%)					88.5	0.4								
Wang et al. [36]	0.9 (0.8)	0.6 (0.5)	0.3 (0.3)	0.5 (0.5)	0.4 (0.3)										
Schwarz et al. [32]	100 (93.3) (%)			45 (45.4)	55 (47.8)										
Schwarz et al. [33]	100 (96.6) (%)					39.9 (41.6)	60.1 (55)	45.1 (21.6)	54.9 (75)						
Schwarz et al. [34]	100 (95.2) (%)											14.8 (23.5)	85.2 (71.6)		
Schwarz et al. [18]	100 (93.3) (%)													10 (6.6)	90 (86.7
Schwarz et al. [39]	92.3 (%)			17.9	73.4										
Bianchini et al. [44]	100 (%)													10.7	89.3
Bianchini et al. [43]	83 (%)									12.4	70.8				
Englezos et al. [42]	100 (%)							25	75						
Matarasso et al. [38]	19.7 (%)					6.1	13.6								
Nart et al. [37]	100 (%)					29	71								
Data from implant: the included studie	s treated with ac	djunctive imp with 1 decim	olantoplasty al nal only.	re presented,	and data fro	om control or p	barallell grou	ups are present	ed in paranthe	ses. Note that	different indice	s were used	in the studi	es. All num	bers fron

The RCT study by Romeo et al. [16,17] reported the same baseline values for both groups, which was reduced at all follow-ups but not significantly different between groups. In Lassere et al. [45], PI decreased significantly in both groups. In Dalago et al. [46], the mPI values decreased from baseline throughout the follow-ups in all groups, but only significantly in one of the control groups after 1 year. No significant differences were reported across groups (Table 6).

Clinical attachment level (CAL)

Four studies performed open flab debridement with adjunctive implantoplasty combined with reconstructive therapy [18,32–34,36,38,39], which in general led to substantial CAL gain. Other studies (Lassere et al. 2020; Romeo et al. 2005, 2007) did not include reconstructive treatment, which in general rendered limited CAL changes [16,17,45] (Table 7).

For studies with several follow-up measurements, Schwarz et al. demonstrated an initial CAL reduction which remained throughout the follow-up [18,32–34]. In Romeo et al. where no reconstructive treatment was performed, CAL was stable in the adjunctive implantoplasty group but increased successively in the control group [16,17]. The RCT by Lassere et al. presented significant CAL reductions in both groups [45].

Bone level (BL)

Ten studies measured BL changes [16,17,36–38,41–46] (Table 8). The studies that did reconstructive treatment demonstrated mean BL gain [37,38]. Studies without reconstructive therapy presented contrasting results as either mean BL loss [42,46], slight BL gain [43,45], or sustained BL values [16,17,44] were reported.

One of the studies that reported several follow-up measurements demonstrated no change of BL during the study period in the adjunctive implantoplasty group and BL loss in the control group (significant difference) [16,17]. In Dalago et al. loss of BL was observed in both test group and control groups over follow-up [46]. There was a slight BL gain in both groups in the study by Lassere et al. [45]. The retrospective case-control study by Ravida et al. showed BL loss in both groups and no difference between the groups for annual BL [41].

Implant survival

Implant survival strongly reflected the years of follow-up in the various included studies (Table 9), and the implant survival ranged from 81% to 100%.

In the RCT studies a higher implant survival in the group treated with adjunctive implantoplasty compared to the control groups was found in Romeo et al. and Dalago et al. [16,17,46], but in the study by Lassere et al. a lower implant survival was reported in the adjunctive implantoplasty group compared to the control group [45]. The retrospective case-control study by Ravida et al. reported higher implant survival in the adjunctive implantoplasty group compared to the control group after a minimum of 1-year follow-up [41].

Table 4. SoP from baseline to post-operative measurements.

Study	Baseline SoP (%)	3 M	Δ 3 M	6 M	Δ 6 M	12 M	Δ 12 M	24 M	Δ 24 M	36 M	Δ 36 M	48 M	Δ 48 M	5+ Y	Δ 5+ Y
Lasserre et al. [45]	11 (6)	4 (3)	7 (3)	4 (4)	7 (2)										
Galarraga-Vinueza	39			0	39										
et al. [40]															
Ravida et al. [41]	11					0	11								-
Bianchini et al. [44]	50													0	50
Bianchini et al. [43]	100									0	100				
Englezos et al. [42]	70							2.5	67.5						
Nart et al. [37]	88.2					0	88.2								

Implants that were treated with adjunctive implantoplasty and controls are presented.

Data from implants treated with adjunctive implantoplasty are presented, and data from control or parallell groups are presented in parantheses. All numbers from the included studies are presented with 1 decimal only.

Importantly, no significant differences were reported between groups in any of the studies for this outcome.

Post-operative peri-implant maintenance program

The frequencies and means of post-operative supportive maintenance following surgical treatment of peri-implantitis with adjunctive implantoplasty varied considerable among studies. A detailed description of the supportive maintenance administered can be found in Table 9, but not all studies disclosed this.

Implantoplasty protocols

The protocols for adjunctive implantoplasty can be found in Table 10. Diamond burs was the most frequently employed bur.

Discussion

The present scoping review considered changes in clinical parameters following surgical peri-implantitis therapy with adjunctive implantoplasty compared to baseline. In studies which included a control group of surgical peri-implantitis treatment without adjunctive implantoplasty, similar outcomes were for the most part observed.

It was not the objective of this scoping review to compare clinical outcomes following surgical peri-implantitis treatment with or without adjunctive implantoplasty, but to review the inflammatory-related clinical outcomes after adjunctive implantoplasty. Collectively, this data indicates that an aggravated inflammatory reaction as a result of titanium and metal debris was not reflected in the clinical data that has been published to date. The literature seems to suggest clinical improvements of the clinical parameters assessed, and with no pronounced difference whether adjunctive implantoplasty was performed or not. Follow-up studies for several years exist, and the findings indicate a lasting outcome. This suggests that a potential clinical effect of the suggested 'inflammatory-aggravated' situation may not be detected for the first few years after treatment. There are however few long-term studies, and the included studies vary considerably in design.

As BoP was a prerequisite for inclusion in the studies it is not surprising that BoP unequivocally decreased from baseline to the first follow-up appointment. Only one study reported similar BoP at follow-up and baseline, and this was found in both test and control groups [41]. The reason for this may be related to the retrospective case-control design which included subjects with different history of peri-implant maintenance attendance following surgical treatment with or without adjunctive implantoplasty. It was reported that clinical outcomes were influenced by the frequency of periimplant maintenance attendance and not by adjunctive implantoplasty [41]. Five studies included more than one follow-up, and the improvements in BoP was without exception maintained beyond the first follow-up. Thus, clinical signs of inflammation in peri-implant tissues did not seem to increase with time after surgical peri-implantitis treatment with adjunctive implantoplasty. Romeo et al. was the only study to find a statistically significant difference in BoP in favour of adjunctive implantoplasty over the control group [16,17].

Studies on surgical peri-implantitis treatment performed without adjunctive implantoplasty are in line with the results presented here. Sustained improved BoP values have been demonstrated over multiple follow-up appointments [50–52], but studies have also reported an increase of BoP over follow-up after the initial drop from baseline to the first follow-up [10,11,53,54].

The outcome PPD was in line with BoP. With the exception of Romeo et al. [16,17], no differences were observed between groups in studies with control groups. Mean PPD values remained low in studies with multiple follow-ups [16–18,32–34,36,39,45,46].

The same findings have been reported in studies addressing peri-implantitis surgery without adjunctive implantoplasty. Mercado et al. demonstrated stable PPD over a threeyear follow-up after peri-implantitis surgery in combination with reconstructive treatment [50]. Stable PPD has also been reported over a 12-month follow-up in studies without the use of regenerative treatment [52,54]. However, studies have also reported increased mean PPD over follow-up time both with and without reconstructive treatment in conjunction with surgical peri-implantitis treatment [10,53].

Facilitated plaque removal and impeded microbial adhesion are often advocated as rationales for adjunctive implantoplasty. The mean mPl was consecutively lower in the adjunctive implantoplasty group but not significantly in Romeo et al. [16,17]. In all the other studies with control groups, no effect of adjunctive implantoplasty on plaque indices were reported [41,45,46]. In general, mean Pl values

Study	Baseline PPD (mm)	3 M	Δ 3 M	6 M	Δ6 M	12 M	Δ 12 M	24 M	Δ 24 M	36 M	Δ 36 M	48 M	Δ 48 M	5+ Y	Δ 5+ \rangle
Romeo et al. [16,17]	5.8 (6.5)			3.4 (5.4)	2.4 (1.1)	3.4 (5.9)	2.4 (0.6)	3.6 (5.5)	2.2 (0.9)	3.2 (-)	2.6				
Dalago et al. [46]	6.4 (5.8, 5.7)					3.4 (4, 3.7)	3 (1.8, 2)	4 (4.4, 4.2)	2.4 (1.4, 1.3)	4.1 (4.4, 3.8)	2.3 (1.4, 1.9)				
Lasserre,[45]	6.7 (5.6)	3.4 (2.8)	3.3 (2.8)	2.7 (2.3)	4 (3.3)										
Galarraga-Vinueza	4.6			3.4	1.3										
et al. [40]															
Ramanauskaite et al. [35]	6.8 (6.3)														2.1 (1.3
Ravida et al. [41]	5.2					3.9	0.3								
Wang et al. [36]	6.4 (7.7)	5 (6)	1.4 (1.7)	4.6 (5.1)	1.9 (2.7)										
Schwarz et al. [32]	5.5 (5.1)			3.1 (3.4)	2.4 (1.7)										
Schwarz et al. [33]	5.2 (4.9)					3.2 (3.2)	2 (1.7)	3.7 (3.8)	1.5 (1.1)						
Schwarz et al. [34]	5.5 (5.1)											4.3 (3.8)	1.2 (1.3)		
Schwarz et al. [18]	5.8 (4.8)													3.6 (4)	2.6 (0.7
Schwarz et al. [39]	6.2			3.6	2.5										
Bianchini et al. [44]	5.8									1.3	4.5				
Englezos et al. [42]	8.7							3.3	5.4						
Matarasso et al. [38]	8.1					4	4.1								
Nart et al. [37]	6.5					3.5	ĸ								

measurements
post-operative
to
baseline
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Table

Table 6. PI from base	eline to post-ope	stative meas	surements.												
Study	Baseline Pl	3 M	Δ3 M	6 M	Δ6 M	12 M	Δ12 M	24 M	Δ 24 M	36 M	A 36 M	48 M	Δ48 M	$5+\gamma$	Δ 5+ Y
Romeo et al. [16,17]	1.5 (1.5)			0.9 (1.2)	0.6 (0.3)	0.9 (1.3)	0.7 (0.2)	0.9 (1)	0.7 (0.5)	(-) 6.0	0.7				
Dalago et al. [46]	1.6 (1.3, 1.5)					1 (0.1, 0.7)	0.6 (1.2, 0.8)	0.5 (0.7, 0.3)	1.1 (0.6, 1.2)	1 (0.8, 0.4)	0.6 (0.5, 1.1)				
Lasserre et al. [45]	0.2 (0.4)	0.4 (0.4)	0.2 (0)	0.2 (0.1)	0 (0.3)										
Galarraga-Vinueza	0.5			0.45	0.05										
et al. [40]															
Wang et al. [36]	0.6 (0.2)	0.5 (0.5)	0.1 (0.3)	0.3 (0.4)	0.2 (0.2)										
Schwarz et al. [32]	0.7 (0.7)			1.2 (1.1)	0.5 (0.4)										
Schwarz et al. [33]	0.7 (0.5)					1.1 (0.7)	0.4 (0.2)	0.7 (0.3)	0.0 (0.2)						
Schwarz et al. [34]	0.8 (0.4)											0.8 (0.8)	0.0 (0.4)		
Schwarz et al. [18]	0.8 (0.2)													0.6 (0.3)	0.2 (0.1)
Schwarz et al. [39]	0.2			0.0	0.2										
Nart et al. [37]	18 (%)					25	7								
Implants that were tr	eated with adjur	nctive impli	antoplasty ā	and controls	are presented	l. Data from i	mplants treated	i with adjunctive	e implantoplasty	r are presente	d, and data fron	n control c	r parallell g	roups are p	oresented

Study	Regenerative treatment	Baseline CAL (mm)	3 M	Δ 3 M	6 M	Δ6 M	12 M	Δ 12 M	24 M	Δ 24 M	36 M	Δ 36 M	48 M	Δ 48 M	5+ Y	Δ 5+ Y
Romeo et al. [16,17]		5.5 (6)			5.6 (6.4)	0.1 (0.4)	5.7 (7.3)	0.2 (1.3)	5.9 (7)	0.4 (1)	5.2 (-)	0.3				
Lasserre et al. [45]		7 (6.2)	4 (3.8)	3 (2.4)	3.5 (3.4)	2.5 (2.8)										
Wang et al. [36]	+	6.9 (7.4)	6 (6)	0.9 (0.4)	5.5 (5.5)	1.5 (1.9)										
Schwarz et al. [18]	+	7.1 (6.8)													4.4 (4.7)	2.7 (2.1)
Schwarz et al. [39]	+	6.7			4.6	2.1										
Matarasso et al. [38]	+	9.7					6.7	m								
Implants that were truin parantheses. All nui	eated with adjunct mbers from the in	tive implantopli	asty and cc are present	ontrols are p ted with 1 de	resented. Da ecimal only.	ta from impli	ints treated	with adjunct	ive implant	oplasty are	presented,	and data fro	om contro	l or parallell	groups are	presented

Table 7. CAL clinical values from baseline to post-operative measurements.

measurements.
post-operative
baseline to
evel) from
BL (bone k
Table 8.

	Regenerative	Baseline bone														
Study	treatment	level (mm)	3 M	Δ 3 M	6 M	Δ6 M	12 M	Δ 12 M	24 M	Δ 24 M	36 M	A 36 M	48 M	Δ48 M	5+Y 2	Δ 5+ Y
Romeo et al. [16,17]		3.9 (3.5)					3.9 (4)	0 (0.5)	3.9 (4.5)	0 (1)	3.9 (5.4)	0 (2.1)				
Dalago et al. [46]		5.5 (5, 4.3)					5.9 (5.3, 4.5)	0.4 (0.3, 0.2)	6.2 (5.4, 4.5)	0.7 (0.4, 0.2)	6.4 (5.5, 4.7)	0.9 (0.5, 0.4)				
Lasserre et al.,[45]		4.7 (5.2)	I		4.5 (4.7)	0.2 (0.5)										
Ravida et al. [41]		3.6					4.3	0.7								
Wang et al. [36]	+	I				1.1 (1.3)										
						(bone gain)										
Bianchini et al. [44]		4.4													4.5	0.1
Bianchini et al. [43]		5									4.3	0.7				
Englezos et al. [42]		5.4							5.6	0.2						
Matarasso et al. [38]	+	8					5.2	2.8								
Nart et al. [37]	+	4.3					1.2	3.1								

Implants that were treated with adjunctive implantoplasty and controls are presented. Data from implants treated with adjunctive implantoplasty are presented, and data from control or parallell groups are presented in parantheses. All numbers from the included studies are presented with 1 decimal only.

Table 9. Overview of maintenance following adjunctive implantoplasty.

Study	Implant survival (%)	Follow-up period	Maintaince program
Romeo et al. [16,17]	100	36 M	-
Dalago et al. [46]	100	36 M	Weekly plaque control in the first month and reinforcement of oral hygiene and prophylaxis every 6 months.
Lasserre et al. [45]	91	6 M	Post-operative care was provided at 1 week and 3 months prior to the final 6 months evaluation. Oral hygiene instructions and supragingival cleaning were given at 3 and 6- month evaluation.
Galarraga-Vinueza et al. [40]	100	6 M	
Ramanskauite et al. [35]	100	6 M - 10.5 Y	
Ravida et al. [41]	90	>12 M	-
Wang et al. [36]	100	6 M	Post-operative supragingival debridement around implants at 3 and 6 months after baseline.
Schwarz et al. [18,32–34,39]	81	6 Y	Every second week during the first 2 months after surgery, then monthly during the first 6 months to control oral hygiene and wound healing. Thereafter every six months after the first year. After the 2nd year annual professional cleaning and hygiene reinforcement.
Schwarz et al. [39]	100	6 M	Controls every second week during the first 2 months, and thereafter maintenance every third month.
Bianchini et al. [44]	87	24 M	
Bianchini et al. [43]	100	36 M	
Englezos et al. [42]	100	24 M	Recall between 1 to 3 months after therapy, and the frequency decreased from 2–4 times a year based on individual needs.
Matarasso et al. [38]	100	12 M	Controls were performed weekly during the first six weeks of healing, and at a 3–6 months interval in the following time based on individual risk assessment.
Nart et al. [37]	100	12 M	Post-operative care was provided every second week in the first month, and then scheduled every 2 months.

Only data for implants that received adjunctive implantoplasty treatment are presented. Data not available (-).

Table 10. Protocol used for adjunctive implantoplasty.

Protocol type	Study
A	Englezos et al. [42]
	Matarasso et al. [38]
	Romeo et al. [16,17]
	Bianchini et al. [43,44]
В	Schwarz et al. [39]
	Ramanskuaite et al. [35]
	Schwarz et al. – [18,32–34]
	Nart et al. [37]
	Gallarraga-Vinueza et al. [40]
C	Dalago et al. [46]
	Lasserre et al. [45]
D*	Ravida et al. [41]
Not specified	Wang et al. [36]

A: Diamond bur(s) + Arkansas stone + Silicone bur(s).

B: Diamond bur(s) + Arkansas stone.

C: Diamond bur(s).

D: Carbide bur(s) + Silicone bur(s).

*Not all implants were treated with silicone burs.

decreased from baseline which is to be expected after treatment and inclusion in a study.

In studies with control groups no difference in mean BL changes were reported according to adjunctive implantoplasty [41,45,46], with the exception of the study by Romeo et al. where successive loss of bone in the control group was demonstrated compared to stability in the test group [16,17]. The studies that assessed BL following peri-implantitis surgery with adjunctive implantoplasty in combination with reconstructive therapy naturally demonstrated BL gain [37,38], indicating that this treatment modality also can be combined with reconstructive therapy. This has also been demonstrated in reconstructive treatment without adjunctive implantoplasty [55]. Importantly, none of the included studies assessed BL changes over more than one time point, which points to the lack of evidence over time of outcomes after reconstructive treatment combined with adjunctive implantoplasty.

In studies that employed reconstructive treatment in conjunction with peri-implantitis surgery, but without adjunctive implantoplasty, some evidence exists. La Monaca and coworkers [53], demonstrated an increase of bone level by approximately 1.5 mm from baseline to 1-year follow-up after reconstructive peri-implantitis treatment, followed by a successive loss of BL until the 5-year follow-up with BL returning to baseline levels. However, studies have also showed BL stability over follow-up time both with and without reconstructive treatment in conjunction with peri-implantitis surgery [10,50].

None of the included case-control or RCT-studies reported a significant difference of implant survival, which may not be surprising considering the limited follow-up. The reported survival rates in the included studies between 81% and 100% corroborates studies on surgical peri-implantitis treatment without adjunctive implantoplasty [10,51–54].

Several different implantoplasty protocols were used in the included studies. The choice of protocol may be of clinical relevance as it may influence both surface roughness parameters but also the debris composition. It is therefore possible that the choice of burs for implantoplasty may be related to a potential inflammatory-aggravating effect. No study was identified that included more than one protocol, and hence, the clinical impact of different burs used for implantoplasty is not known.

The impact of peri-implant maintenance therapy frequency following surgical treatment is well documented [50]. In the studies included, the maintenance frequency interval for most studies ranged between 3–6 months, but also yearly after the first year. The frequency interval and quality of maintenance therapy may be more related to clinical outcomes related to inflammation (e.g. BoP, PPD) over time than the effect of the adjunctive implantoplasty per se, which was demonstrated in the study by Ravida et al. [41].

In the present scoping review, parameters from clinical studies on adjunctive implantoplasty has been discussed in light of a potential inflammatory-aggravated effect on periimplant tissues. This review has important limitations. A meta-analysis was not performed due to the very diverse protocols in the included studies. The search strategy included only studies in English, did not include grey literature, and there is high risk of bias in many of the included studies. Inclusion criteria, non-surgical debridement prior to surgical treatment, burs included in the implantoplasty protocols, the use of local or systemic antibiotics, screw-retained or cement-retained prostheses, were all factors that differed substantially between studies and exemplifies the very different study designs in clinical studies on implantoplasty. Not all studies provided information about every parameter discussed here except for BOP, which was an inclusion-criteria in this review. Furthermore, in this review studies including open flap debridement, but also resective and reconstructive surgical treatment approaches, were all considered. These are very different strategies which may subsequently influence the clinical outcome and parameters. The post-surgical follow-up and maintenance frequency varied considerably in the included studies, which is a limitation when considering inflammatory-related clinical parameters. Finally, one factor hardly discussed in the literature is access for the different burs required for adjunctive implantoplasty. Although burs designed particularly for implantoplasty have been developed, the osseous architecture around implants may render parts of the implant surface difficult to access for this treatment.

In this review only mean values have been discussed. It is well known that any increase or decrease of a clinical parameter very often represents deterioration or healing in few or single patients. Nevertheless, mean numbers are useful to address the aim of the study when considering a potential inflammatory-aggravating effect of implantoplasty.

Implant surface characteristics was not included in all studies and was not considered in this review. This factor further adds to the heterogeneity among the studies and points to the complexity of comparing outcomes of treatment. Although implantoplasty effectively may remove biofilm and hard deposits on implants, it is presumably hardly performed on machined implant surfaces because it would lead to a rougher surface. The study by Romeo et al. [16,17] was the only study presenting a clear advantage of adjunctive implantoplasty (significant reductions in PPD, CAL, mBI and BL) of the studies with a control group. This may be related to the fact that all implant surfaces in the study were titanium-plasma sprayed, which may suggest that adjunctive implantoplasty may be efficacious for some implant surfaces, but not for others.

Conclusions

Within the limitations of this scoping review, data from the included studies indicate that surgical peri-implantitis treatment with adjunctive implantoplasty leads to a reduction in BoP, and that this is in line with data presented in studies on peri-implantitis surgery without adjunctive implantoplasty. In general, this was also the case for the secondary outcomes PDD, BL, CAL and plaque indices. A potential clinical effect of a suggested 'inflammatory-aggravated' impact following adjunctive implantoplasty is not evident considering the existing literature, at least not for the first years after treatment. The literature demonstrates a lack of standardisation across studies. Further studies, and in particular randomised clinical studies, with uniform research protocols, sufficient sample sizes and follow-up time, are needed to establish the effect of adjunctive implantoplasty in the treatment of periimplantitis.

Disclosure statement

The authors declare no conflict of interest.

References

- Koldsland OC, Scheie AA, Aass AM. Prevalence of peri-implantitis related to severity of the disease with different degrees of bone loss. J Periodontol. 2010;81(2):231–238.
- [2] Derks J, Schaller D, Håkansson J, et al. Effectiveness of implant therapy analyzed in a Swedish population: prevalence of periimplantitis. J Dent Res. 2016;95(1):43–49.
- [3] Mombelli A, Lang NP. The diagnosis and treatment of periimplantitis. Periodontol 2000. 1998;17(1):63–76.
- [4] Zitzmann NU, Berglundh T. Definition and prevalence of periimplant diseases. J Clin Periodontol. 2008;35(8 Suppl):286–291.
- [5] Renvert S, Samuelsson E, Lindahl C, et al. Mechanical non-surgical treatment of peri-implantitis: a double-blind randomized longitudinal clinical study. I: clinical results. J Clin Periodontol. 2009; 36(7):604–609.
- [6] Schwarz F, Sculean A, Rothamel D, et al. Clinical evaluation of an Er:YAG laser for nonsurgical treatment of peri-implantitis: a pilot study. Clin Oral Implants Res. 2005;16(1):44–52.
- [7] Renvert S, Roos-Jansåker A-M, Claffey N. Claffey Non-surgical treatment of peri-implant mucositis and peri-implantitis: a literature review. J Clin Periodontol. 2008;35(8 Suppl):305–315.
- [8] Figuero E, Graziani F, Sanz I, et al. Management of peri-implant mucositis and peri-implantitis. Periodontol 2000. 2014;66(1): 255–273.
- [9] Claffey N, Clarke E, Polyzois I, et al. Surgical treatment of periimplantitis. J Clin Periodontol. 2008;35(8 Suppl):316–332.
- [10] Carcuac O, Derks J, Abrahamsson I, et al. Surgical treatment of peri-implantitis: 3-year results from a randomized controlled clinical trial. J Clin Periodontol. 2017;44(12):1294–1303.
- [11] Koldsland OC, Wohlfahrt JC, Aass AM. Surgical treatment of periimplantitis: prognostic indicators of short-term results. J Clin Periodontol. 2018;45(1):100–113.

- [12] Beheshti Maal M, Aanerød Ellingsen S, Reseland JE, et al. Experimental implantoplasty outcomes correlate with fibroblast growth in vitro. BMC Oral Health. 2020;20(1):25.
- [13] Könönen M, Hormia M, Kivilahti J, et al. Effect of surface processing on the attachment, orientation, and proliferation of human gingival fibroblasts on titanium. J Biomed Mater Res. 1992;26(10): 1325–1341.
- [14] Nothdurft FP, Fontana D, Ruppenthal S, et al. Differential behavior of fibroblasts and epithelial cells on structured implant abutment materials: a comparison of materials and surface topographies. Clin Implant Dent Rel Res. 2015;17(6):1237–1249.
- [15] Kunzler TP, Drobek T, Schuler M, et al. Systematic study of osteoblast and fibroblast response to roughness by means of surfacemorphology gradients. Biomaterials. 2007;28(13):2175–2182.
- [16] Romeo E, Ghisolfi M, Murgolo N, et al. Therapy of peri-implantitis with resective surgery. A 3-year clinical trial on rough screwshaped oral implants. Part I: clinical outcome. Clin Oral Implants Res. 2005;16(1):9–18.
- [17] Romeo E, Lops D, Chiapasco M, et al. Therapy of peri-implantitis with resective surgery. A 3-year clinical trial on rough screwshaped oral implants. Part II: radiographic outcome. Clin Oral Implants Res. 2007;18(2):179–187.
- [18] Schwarz F, John G, Schmucker A, et al. Combined surgical therapy of advanced peri-implantitis evaluating two methods of surface decontamination: a 7-year follow-up observation. J Clin Periodontol. 2017;44(3):337–342.
- [19] Lozada JL, James RA, Boskovic M, et al. Surgical repair of periimplant defects. J Oral Implantol. 1990;16(1):42–46.
- [20] Geremias TC, Montero JF, Magini RS, et al. Biofilm analysis of retrieved dental implants after different peri-implantitis treatments. Case Rep Dent. 2017;2017:1–5.
- [21] Suh JJ, Simon Z, Jeon YS, et al. The use of implantoplasty and guided bone regeneration in the treatment of peri-implantitis: two case reports. Implant Dent. 2003;12(4):277–282.
- [22] Stavropoulos A, Bertl K, Eren S, et al. Mechanical and biological complications after implantoplasty-a systematic review. Clin Oral Implants Res. 2019;30(9):833–848.
- [23] Oliveira NM, Schunemann W, Mathew M, et al. Can degradation products released from dental implants affect peri-implant tissues? J Periodont Res. 2018;53(1):1–11.
- [24] Barrak FN, Li S, Muntane AM, et al. Particle release from implantoplasty of dental implants and impact on cells. Int J Implant Dent. 2020;6(1):50.
- [25] Wu W, Wang L, Mao YQ, et al. Impaired autophagy in the fibroblasts by titanium particles increased the release of CX3CL1 and promoted the chemotactic migration of monocytes. Inflammation. 2020;43(2):673–685.
- [26] Tricco AC, Lillie E, Zarin W, et al. PRISMA extension for scoping reviews (PRISMA-ScR): checklist and explanation. Ann Intern Med. 2018;169(7):467–473.
- [27] Thierbach R, Eger T. Clinical outcome of a nonsurgical and surgical treatment protocol in different types of peri-implantitis: a case series. Quintessence Int. 2013;44(2):137–148.
- [28] Pommer B, Haas R, Mailath-Pokorny G, et al. Periimplantitis treatment: long-term comparison of laser decontamination and implantoplasty surgery. Implant Dent. 2016;25(5):646–649.
- [29] Schwarz F, John G, Becker J. Reentry after combined surgical resective and regenerative therapy of advanced peri-implantitis: a retrospective analysis of five cases. Int J Periodontics Restorative Dent. 2015;35(5):647–653.
- [30] Schwarz F, John G, Sahm N, et al. Combined surgical resective and regenerative therapy for advanced peri-implantitis with concomitant soft tissue volume augmentation: a case report. Int J Periodontics Restorative Dent 2014;34:489–495.
- [31] Sapata VM, de Souza AB, Sukekava F, et al. Multidisciplinary treatment for peri-implantitis: a 24-month follow-Up case report. Clin Adv Periodontics. 2016;6(2):76–82.

- [32] Schwarz F, Sahm N, Iglhaut G, et al. Impact of the method of surface debridement and decontamination on the clinical outcome following combined surgical therapy of peri-implantitis: a randomized controlled clinical study. J Clin Periodontol. 2011; 38(3):276–284.
- [33] Schwarz F, John G, Mainusch S, et al. Combined surgical therapy of peri-implantitis evaluating two methods of surface debridement and decontamination. A two-year clinical follow-up report. J Clin Periodontol. 2012;39(8):789–797.
- [34] Schwarz F, Hegewald A, John G, et al. Four-year follow-up of combined surgical therapy of advanced peri-implantitis evaluating two methods of surface decontamination. J Clin Periodontol. 2013;40(10):962–967.
- [35] Ramanauskaite A, Becker K, Juodzbalys G, et al. Clinical outcomes following surgical treatment of peri-implantitis at grafted and non-grafted implant sites: a retrospective analysis. Int J Implant Dent. 2018;4(1):27.
- [36] Wang C-W, Ashnagar S, Gianfilippo RD, et al. Laser-assisted regenerative surgical therapy for peri-implantitis: a randomized controlled clinical trial. J Periodontol. 2021;92(3):378–388.
- [37] Nart J, de Tapia B, Pujol À, et al. Vancomycin and tobramycin impregnated mineralized allograft for the surgical regenerative treatment of peri-implantitis: a 1-year follow-up case series. Clin Oral Investig. 2018;22(6):2199–2207.
- [38] Matarasso S, lorio Siciliano V, Aglietta M, et al. Clinical and radiographic outcomes of a combined resective and regenerative approach in the treatment of peri-implantitis: a prospective case series. Clin Oral Implants Res. 2014;25(7):761–767.
- [39] Schwarz F, Sahm N, Becker J. Combined surgical therapy of advanced peri-implantitis lesions with concomitant soft tissue volume augmentation. A case series. Clin Oral Implants Res. 2014; 25(1):132–136.
- [40] Galarraga-Vinueza ME, Obreja K, Magini R, et al. Volumetric assessment of tissue changes following combined surgical therapy of peri-implantitis: a pilot study. J Clin Periodontol. 2020; 47(9):1159–1168.
- [41] Ravidà A, Siqueira R, Saleh I, et al. Lack of clinical benefit of implantoplasty to improve implant survival rate. J Dent Res. 2020;99(12):1348–1355.
- [42] Englezos E, Cosyn J, Koole S, et al. Treatment of peri-implantitis: clinical and radiographic outcomes after 2 years. Int J Periodont Res Dent. 2018;38(5):729–735.
- [43] Bianchini MA, Galarraga-Vinueza ME, Bedoya KA, et al. Implantoplasty enhancing peri-implant bone stability over a 3year follow-up: a case series. Int J Periodontics Restorative Dent. 2020;40(1):e1–e8.
- [44] Bianchini MA, Galarraga-Vinueza ME, Apaza-Bedoya K, et al. Two to six-year disease resolution and marginal bone stability rates of a modified resective implantoplasty therapy in 32 peri-implantitis cases. Clin Implant Dent Rel Res. 2019;21(4):758–765.
- [45] Lasserre JF, Brecx MC, Toma S. Implantoplasty versus glycine air abrasion for the surgical treatment of peri-implantitis: a randomized clinical trial. Int J Oral Maxillofac Implants. 2020;35(1): 197–206.
- [46] Dalago HR, Perrotti V, Torres de Freitas SF, et al. Prospective longitudinal comparison study of surgical therapies for peri-implantitis: 3-year follow-up. Aust Dent J. 2019;64(3):237–245.
- [47] Mombelli A, Van Oosten MAC, Schürch E, et al. The microbiota associated with successful or failing osseointegrated titanium implants. Oral Microbiol Immunol. 1987;2(4):145–151.
- [48] Löe H. The gingival index, the plaque index and the retention index systems. J Periodontol. 1967;38(6 Part II):610–616.
- [49] ÓLeary TJ, Drake RB, Naylor JE. The plaque control record. J Periodontol. 1972;43(1):38.
- [50] Mercado F, Hamlet S, Ivanovski S. Regenerative surgical therapy for peri-implantitis using deproteinized bovine bone mineral with 10% collagen, enamel matrix derivative and Doxycycline- a pro-

spective 3-year cohort study. Clin Oral Impl Res. 2018;29(6): 583–591.

- [51] Heitz-Mayfield LJA, Salvi GE, Mombelli A, et al. Supportive periimplant therapy following anti-infective surgical peri-implantitis treatment: 5-year survival and success. Clin Oral Implants Res. 2018;29(1):1–6.
- [52] La Monaca G, Pranno N, Annibali S, et al. Clinical and radiographic outcomes of a surgical reconstructive approach in the treatment of peri-implantitis lesions: a 5-year prospective case series. Clin Oral Impl Res. 2018;29(10):1025–1037.
- [53] Hallström H, Persson GR, Lindgren S, et al. Open flap debridement of peri-implantitis with or without adjunctive systemic antibiotics: a randomized clinical trial. J Clin Periodontol. 2017;44(12):1285–1293.
- [54] Heitz-Mayfield LJA, Salvi GE, Mombelli A, et al. Anti-infective surgical therapy of peri-implantitis. A 12-month prospective clinical study. Clin Oral Implants Res. 2012;23(2):205–210.
- [55] Aghazadeh A, Persson GR, Renvert S. A single-centre randomized controlled clinical trial on the adjunct treatment of intra-bony defects with autogenous bone or a xenograft: results after 12 months. J Clin Periodontol. 2012;39(7):666–673.