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Abstract: BACKGROUND To selectively review the existing literature on post-interventional maintenance protocols in patients with periodontal disease receiving either non-surgical or surgical periodontal treatment. METHODS Three systematic reviews with different periodontal interventions, i.e. scaling and root planing (SRP), SRP with adjunctive antibiotics or regenerative periodontal surgery were evaluated focusing on their post-interventional maintenance care. Due to the early publication of one review an additional literature search update was undertaken. The search was executed for studies published from January 2001 till March 2015 through an electronic database to ensure the inclusion of resent studies on SRP. Two reviewers guided the study selection and assessed the validity of the three reviews found. RESULTS Within the group of scaling and root planing alone there have been nine studies with more than three appointments for maintenance care and five studies with more than two appointments in the first 2 months after the intervention. Chlorhexidine was the most frequently used antiseptic agent used for 2 weeks after non-surgical intervention. Scaling and root planing with adjunctive antibiotics showed a similar number of visits with professional biofilm debridement, whereas regenerative studies displayed more studies with more than three visits in the intervention group. In addition, the use of antiseptics was longer and lasted 4 to 8 weeks after the regenerative intervention. The latter studies also showed more stringent maintenance protocols. CONCLUSIONS With increased interventional effort there was a greater tendency to increase frequency and duration of the maintenance care program and antiseptic agents.

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Conclusions: With increased interventional effort there was a greater tendency to increase frequency and duration of the maintenance care program and antiseptic agents.

Keywords: Debridement, Flap surgery, Regeneration, Maintenance, Antiseptics

Background

Colonization by a pathogenic biofilm is recognized as the primary etiologic factor for the initiation and progression of periodontitis [1]. Despite the fact that host and environmental factors may significantly contribute to the resulting inflammatory process [2], it has been convincingly shown that professional supra- and subgingival biofilm control is able to control disease initiation and progression [3]. The effective control and management of the supra- and subgingival biofilm is traditionally performed by mechanical means, such as hand instruments and/or ultrasonic debridement [4]. Further methods include air-polishing devices with various inserts and powders, the latter preferably being effective in removing biofilms but low-abrasive dental hard tissues [5, 6].

Thorough non-surgical scaling and root planing (SRP) was demonstrated as an important part of successful periodontal treatment, especially on deeper periodontal pockets [7]. The results of such treatment may only be maintained in the long-term when an
effective supragingival plaque control is performed, and a regular supportive periodontal treatment (SPT) is applied [1, 8]. In addition, a body of evidence shows the benefit of systemically administered antibiotics as an adjunctive to SRP, particularly in patients with aggressive periodontitis and in those with advanced chronic disease [9, 10].

However, in distinct clinical situations with local defects, e.g., in teeth with furcation involvement or in single-rooted teeth with vertical bone defects, residual increased probing pocket depth (PPD) might persist after non-surgical therapy and require further treatment, e.g., surgical interventions, in order to prevent ongoing loss of attachment and tooth loss [11, 12].

Different studies have analyzed the effects of supervised maintenance care after periodontal therapy e.g. subgingival scaling and root planing or surgical interventions. Such maintenance programs included the adjunctive use of antiseptic rinsing followed by professional supragingival cleanings [13, 14]. These supervised maintenance care recommendations are mostly given after elaborate, regenerative periodontal surgery.

However, there is no comparative study or systematic review available, which evaluates the influence of different approaches on clinical outcomes. Therefore, the purpose of this study was to assess post-interventional maintenance protocols in terms of frequency and adjunctive antiseptic infection control for three different treatment modalities concerning infectious periodontal conditions: after non-surgical periodontal therapy with and without systemic antibiotics and regenerative surgical interventions.

The following specific questions were addressed as follows:

1. In a patient population with chronic periodontal disease or periodontal disease with infrabony defects, who underwent different periodontal interventions, which frequency of post interventional maintenance was applied?
2. Is there a difference in pocket depth reduction among the same groups of periodontal therapy with different recall maintenance protocols?

Methods
Protocol
The present article merged and screened three already existing systematic reviews, that assessed three different treatment options: SRP [15], systemic antibiotics (amoxicillin and metronidazole) as adjunctive to SRP [16] and regenerative periodontal surgery [17]. All three reviews covered different periodontal therapeutic procedures. The intention of this article was not to compare these primary therapeutic concepts but to expose the measures that were taken after each of these therapies, to put a specific light on the post-interventional protocols and to elaborate any potential differences between the different therapeutic approaches. All studies within the reviews did not show any overlap in the articles the authors have chosen.

Two [16, 17] of the three reviews were fitted to match the current PRISMA (Preferred Reporting Items for Systematic Review and Meta-analyses) criteria for reviews [18]. An older systematic review did not follow up to date protocol recommendations [15] due to its earlier publishing date. To assure up-to-dateness and to avoid missing current articles a new literature search was undertaken as described below:

Eligibility criteria for additional search
Following parameters of the publications needed to be presented in order to be eligible for inclusion:

1. The articles needed to be randomized controlled trials (RCT) or controlled clinical trials on periodontal treatment with a follow-up of at least 12 months or more, written in English. (2) Patients with chronic periodontitis by the age of at least 20 years. (3) A recording of maintenance care plan of at least 2 months post intervention.

Outcome measures
The main focus of this study was to filter out different maintenance strategies after any periodontal intervention such as frequency of appointments after SRP and periodontal surgery. In addition changes in periodontal probing depth (PPD) were extracted as primary parameter outcome for meta-analysis. Secondary parameter outcome such as recession (REC), clinical attachment level (CAL) or plaque index (PI) were not part of this meta-analysis due to the non-homogenous data presentation in the single studies. Since the data of probing depth at the requested time points were missing in the non-surgical interventions, only guided tissue regeneration (GTR) studies, which adequately reported on this parameter could be included in the forest-plot.

Additional analysis and information sources
Due to one study’s early publishing date, the literature search was updated and the electronic databases MEDLINE and Cochrane (Oral Health Group Specialist Trials Register) were consulted again for studies published from January 2001 to March 2015, while the search strategy was re-formulated based on the suggested three complexes “non-surgical therapy” AND “surgical therapy” AND types of studies. Two independent reviewers (ID and PRS) screened for additional titles written in English and searched for possible inclusion criteria, which would match this
Influence of maintenance on therapy
In order to assess the influence of different maintenance protocols, the probing depth reduction served as a clinical outcome. The data on mean and standard deviation of probing depth reduction were extracted from each of the included studies for meta-analysis. Because of differences in the observation period across studies, only those studies were pooled that had somewhat similar follow-up frequencies. Due to a large amount of heterogeneity between studies ($I^2 > 50\%$), a random effects model was necessary for pooling. All analyses were performed with R [19]. The studies were arranged in the following categories: protocol 1 (two or less recall visits within the 2 months) and protocol 2 (three or more visits within the 2 months after periodontal intervention). The duration of the use of antiseptic agents were categorized into A, B and C. CHX/A accounts for the use of antiseptics lasting for 2 weeks of rinsing after periodontal intervention. CHX/B required rinsing of 4 weeks and CHX/C of up to 8 weeks after periodontal intervention. For example protocol 2 and CHX/C displays the most vigorous maintenance protocol compared to protocol 1 and CHX/A being the least vigorous post interventional maintenance care (Fig. 2).

Quality assessment
Studies within the three systematic reviews were methodologically screened by two reviewers in order to assess the quality and a potential risk of bias [15–17].

Summary of measures
Throughout the three systematic reviews there have been a variety of different maintenance protocols. All studies, which utilized an antiseptic agent, included chlorhexidine (CHX) into their maintenance program, whereas the concentration and duration varied among the studies. If mentioned, all studies provided supra-gingival cleanings, in two cases oral hygiene instructions and motivation was given. The frequency of follow-up intervals throughout the different reviews was heterogenic.

Evaluation of the maintenance programs
The following aspects of the post-interventional maintenance protocols were analyzed. The recall frequency including mechanical re-instrumentation and/or re-motivation in the first 2 months after the intervention was arbitrarily identified, whether or not antiseptic rinsing was utilized (active ingredient, concentration, frequency and duration). The results were organized in a subgroup analysis assessing the change of pocket depth reduction. Subgroups were defined as following: (I) The recall frequency during the first 2 months with $1 =$ two or less visits ($\leq 2$) and $2 = $ three or more visits within the 2 months ($\geq 3$) and (II) the duration of adjunctive use of antiseptics for (A) two or less than 2 weeks, (B) Up to 4 weeks and (C) more than 5 weeks. According to this classification the lowest level of maintenance strategy was therefore 1A and the highest-level 2C. Based on this classification system design, further subgroup combinations were possible.

Results
Study selection
In total, three reviews were identified by the electronical database search. Since the date of the publication by Heitz-Mayfield and co-workers dating back to 2002 was not up-to-date, an additional investigation was initiated. The latter revealed another 697 publications. After the independent screening procedure by two of the authors (I.D. und P.R.S.), eight studies were included for the full-text analysis. Finally, one additional study met the inclusion criteria and was entered in Table 1 for analysis [20] (Fig. 1).

Description of study maintenance protocol
The analysis of the maintenance protocols included reviews with three different periodontal approaches. Mayfield et al. (2002) evaluated studies dealing with a non-surgical approach whereas Zandbergen and co-workers in 2013 analyzed studies treating periodontal disease with conventional non-surgical therapy and the adjunctive use of systemically administered amoxicillin and metronidazole. Further Graziani et al. (2012) focused on studies dealing with regenerative therapy. Taking into account the heterogenic study designs of 78 studies in total depending on the treatment, most of the studies listed in the reviews followed a specific
maintenance protocol after treatment. A detailed overview of the different maintenance protocols is given in Tables 1, 2, and 3. This analysis’s keynote was type and concentration of antiseptic formula as well as instruction given concerning intake frequency and duration. In addition, the type of maintenance and intervals after treatment were defined. With regard to the antiseptic rinsing, all studies used chlorhexidine (CHX) in concentrations ranging from 0.06 to 0.2 %. The latter and therefore highest concentration was used in roughly 50 % of the studies. Patients were advised to rinse twice daily in most articles, whereas the individual antiseptic rinsing period varied significantly. The minimum concentrations and time for adjunctive chemical plaque control was found in the SRP with adjunctive use of antibiotics group of one study. There, patients were advised to use 0.06 % antiseptic chlorhexidine once daily for 8 days [21–23] (Table 2). The most extensive antiseptic regimen was revealed in studies with regenerative treatment. The minimum rinsing concentrations of 0.12 % chlorhexidine and a rinsing period of up to 11 weeks going up to a concentration of 0.2 % and a rinsing duration of 10 weeks could be observed [24, 25] (Table 3). The recall duration and frequency including professional plaque control also varied throughout all the included studies as listed in the reviews. Table 4 compares the antiseptic duration and recall frequency 2 months after treatment within the different treatment groups. The non-surgical approach group chose to administer the higher chlorhexidine concentration with 0.2 %. However, only five out of 14 studies in the review by Mayfield et al. 2002 advised adjunctive antiseptic rinsing, whereas all 14 studies displayed post-interventional recall intervals of at least once every 2 months. Nine studies exhibited a recall interval of equal three if not more visits. Not listed in Table 4 are the five studies within the scaling and root planing group who merely appeared 3 months after the intervention. Whether or not patients were advised to rinse with antiseptic agents is not mention in the studies.

Table 1 List of the analyzed studies dealing with the non-surgical approach [15] (updated 2015)

| Study               | Antiseptic Galenic formulation | Application frequency | Duration category | Maintenance protocol in the first 2 months | Category |
|---------------------|--------------------------------|-----------------------|-------------------|------------------------------------------|----------|
| Pihlstrom & Ortiz 1981 | n.r.                          |                       |                   | Recall frequency varied, 3–4x/yr          | 1        |
| Lindhe & Westfeldt 1982 | CHX 0.2 %                     | 2x/d                  | 2 weeks A         | 6 month every 2nd week, then 3 month rec for 18 month | 2        |
| Pihlstrom & McHugh 1983 | n.r.                          |                       |                   | Recall 3–4x/yr                            | 1        |
| Lindhe & Westfeldt 1984 | yes                           |                       |                   | 6 month every 2nd week, then 3 month rec for 18 month | 2        |
| Pihlstrom & Oliphant 1984 | yes                           |                       |                   | Recall frequency varied, 3–4x/yr          | 1        |
| Isidor & Karring 1984  | CHX 0.2 %                     | 2x/d                  | 2 weeks A         | 2 year 1, professional prophylaxis 3-monthly year 2, 6-monthly years 3, 4 & 5 – subgingival debridement | 2        |
| Lindhe & Nyman 1985   | CHX 0.2 %                     | 2x/d                  | 2 weeks A         | every 2nd week for 12 weeks, 3 month recall | 2        |
| Kaldahl & Kalkwarf 1996 | n.r.                          |                       |                   | 3 month intervals                        | 1        |
| Isidor & Karring 1986  | CHX 0.2 %                     | 2x/d                  | 2 weeks A         | 2 year 1, professional prophylaxis 3-monthly year 2, 6-monthly years 3, 4 & 5 – subgingival debridement | 2        |
| Ramfjord & Caffesse 1987 | n.r.                          |                       |                   | 1, 2, 3, 4, then every 3 months           | 2        |
| Kaldahl et al. 1988   | n.r.                          |                       |                   | 2, 4, 7 (subgingival plaque removal)      | 2        |
| Kalkwarf et al 1988   | n.r.                          |                       |                   | 1, 2, 4, 7, then 3 month recall           | 2        |
| Kalkwarf et al. 1989  | n.r.                          |                       |                   | 1, 2, 4, 7, then 3 month recall           | 2        |
| Serino et al. 2001    | CHX 0.2 %                     | 2x/d                  | 2 months C        | 3–4x/year, re-examinations after 1, 3, 5, 13 years | 1        |
(Table 1). Seventeen out of 34 studies in the group with SRP and the use of adjunctive antibiotics listed in the review by Zandbergen and co-workers (2013) prescribed chlorhexidine mouth rinse. In this review with 34 studies in total 20 studies performed a post-interventional maintenance protocol of less than two visits. Two studies performed more than three visits within the past 2 months following the procedure. The regenerative approach showed 27 out of 30 studies with adjunctive use of antiseptic agents and 26 studies included a maintenance protocol within the 2 months. 23 studies scheduled their patients more than three times in the 2 months following the regenerative intervention. Reason for this strict maintenance protocol as described by the authors was to remove sutures after surgery, polishing for plaque control due to its tooth brushing abstention at the site of surgery and finally regular supragingival cleanings [17]. Overall, the more sophisticated a treatment intervention was the greater the tendency to increase maintenance frequency, mouth-rinse concentration and duration.

Influence of different maintenance programs on probing depth reduction
A graphical representation of the results is given in a forest plot (Fig. 2). However, to start with, it is important to highlight the nature of this analysis. It is purely based on the data of the regenerative procedure presented by the systematic review from Graziani and co-workers (2012). There have been complications along the way to extract the needed data from the other reviews. Therefore, these pooled results cannot be statistically analyzed, directly compared and interpreted.

Nevertheless, the regenerative studies with more recall interventions and longer duration of antiseptic agents after surgery displayed a greater PPD reduction compared to the studies with lower protocol interventions and antiseptic agents i.e. longer rinsing periods and/or more recall visits looking at the pooled data. The highest mean difference was observed in protocol 2/CHX = C with a mean probing pocket depth difference of 3.7 mm. In general, there has been an increase of the observed effect with increasing baseline PPD. Protocol 1/CHX = C

Fig. 1 Additional search for non-surgical therapy based on the protocol by Heitz-Mayfield and co-workers (2002) [15]
Table 2 Detailed list of analyzed studies dealing with non-surgical periodontal therapy with adjunctive use of amoxicillin and metronidazole [16]

| Study                          | Antiseptic | Galenic formulation | Application frequency | Duration | Category | Maintenance protocol in the first 2 months | Category       |
|-------------------------------|------------|---------------------|-----------------------|----------|----------|-------------------------------------------|----------------|
| Van Winkelhoff et al. 1992    | n.r.       |                     |                       |          |          | n.r.                                      | n.r.           |
| Winkel et al. 1998            | n.r.       |                     |                       |          |          | n.r.                                      | week 21        |
| Müller et al. 1998            | CHX 0.10 % | n. r.               | 2 weeks               | A        | yes      | n.r.                                      | clinical parameters at 3 months |
| Flemming et al. 1998          | CHX 0.06 % | 1x/d                | 8 days                | A        | yes      | n.r. clinical parameters at 3 months     |
| Berglundh et al. 1998         | n.r.       |                     |                       |          |          | n.r.                                      | clinical parameters at 3 months |
| Winkel et al. 2001            | n.r.       |                     |                       |          |          | yes                                       | week 6         |
| Ehmeke et al. 2003            | CHX 0.06 % | 1x/d                | 8 days                | A        | yes      | 3, 6, 9, 12, 18, 24 months               |
| Guerrero et al. 2005          | CHX 0.2 %  | 2x/d                | 2 weeks               | A        | yes      | 2 and 6 months                           |
| Mombelli et al. 2005          | n.r.       |                     |                       |          |          | yes                                       | clinical parameters at 2 and 12 months |
| Ehmeke et al. 2005            | CHX 0.06 % | 1x/d                | 8 days                | A        | yes      | 3, 6, 9, 12, 18, 24 months               |
| Xajigeorgiou et al. 2006      | n.r.       |                     |                       |          |          | yes                                       | week 6         |
| Giannopoulou et al. 2006      | n.r.       |                     |                       |          |          | n.r. clinical parameters at 10 days, 2, 6 and 12 months |
| Moeintaghavi et al. 2007      | n.r.       |                     |                       |          |          | n.r.                                      | clinical parameters at 3 months |
| Kaner et al. 2007             | CHX 0.2 %  | 2x/d                | 2 weeks               | A        | yes      | 3 and 6 months                           |
| Moreira et al. 2007           | CHX 0.12 % | 2x/d                | 2 months              | C        | yes      | 3 month                                  |
| Guerrero et al. 2007          | CHX 0.12 % | 2x/d                | 2 weeks               | A        | yes      | 2 months                                 |
| Machtet et al. 2008           | n.r.       |                     |                       |          |          | oral hygiene motivation                   | every 2nd week for 3 months |
| Johnson et al. 2008           | CHX 0.12 % | 2x/d                | 30 days               | C        | n.r.    | clinical parameters at 3 and 6 months    |
| Akincibay et al. 2008         | n.r.       |                     |                       |          |          | n.r. clinical measurements at 30, 60, 90 days |
| Valenza et al. 2009           | n.r.       |                     |                       |          |          | yes                                       | 2, 6 and 12 months |
| Del Peloso et al. 2009        | n.r.       |                     |                       |          |          | yes                                       | every month for 6 months |
| Cionca et al. 2009            | CHX 0.2 %  | 2x/d                | 10 days               | A        | yes      | 1 week, 3 and 6 months                    |
| Yek et al. 2010               | n.r.       |                     |                       |          |          | yes                                       | monthly up to 6 months |
| Cionca et al. 2010            | CHX 0.2 %  | 2x/d                | 10 days               | A        | yes      | 1 week, 3 and 6 months                    |
| Mestnik et al. 2010           | CHX 0.12 % | 2x/d                | 60 days               | C        | n.r.    | n.r. clinical measurements at 3 months   |
| Heller et al. 2011            | CHX 0.12 % | 2x/d                | 45 days               | C        | yes      | 3 months                                 |
| Rodrigues et al. 2011         | n.r.       |                     |                       |          |          | yes                                       | 1, 3 and 6 months |
| Silva et al. 2011             | n.r.       |                     |                       |          |          | n.r. clinical measurements at 3 months   |
| Varela et al. 2011            | CHX 0.12 % | 2x/d                | 45 days               | C        | yes      | 3 months                                 |
| Baltacioglu et al. 2011       | n.r.       |                     |                       |          |          | n.r. clinical measurements at 2 months   |
| Aimetti et al. 2012           | CHX 0.2 %  | 2x/d                | 2 months              | C        | yes      | every 2nd week for 6 weeks, then every 2 months up to 6 months |
| Casarin et al. 2012           | n.r.       |                     |                       |          |          | n.r.                                      | n.r.           |
and protocol 2/CHX = C – both groups having four studies with a comparable range of baseline probing pocket depths (PPD) – the mean differences greatly differed and accounted for 2.81 mm versus 3.70 mm, respectively. Both groups show different probing depth reductions with different protocols and rinsing durations due to its type of surgical intervention and patient care needs. Again the data presented and its evaluation was extracted from the forest plot done only on the studies on regenerative therapy.

Quality assessment
Graziani et al. and Zandbergen et al. demonstrated a quality assessment to estimate the risk of bias. Nine articles were using adequate methods of study design, unclear methods were used in 21 articles and inadequate methods in eight articles [17]. Out of 28 studies, 15 studies demonstrated low potential risk of bias. The remaining studies showed moderate to high risk of bias [16]. Heitz-Mayfield et al. justified missing quality assessment with a limited number of studies [15].

Discussion
Based on the premises of peer-reviewed papers, this study approach was to pool the evidence and extract the data regarding the maintenance care intervals and procedures. The aim of this article was to put a specific light on the post-interventional protocols and to elaborate any potential differences between the different therapeutic approaches, but not to compare the actual outcomes due to obvious reasons. The performed summary must not be understood as an inadmissible comparison of the clinical results from different treatment approaches. Nevertheless, the latter appraised differences of post-interventional maintenance programs and their impact on periodontal healing was done for the surgical interventions in regard to pocket depth reduction.

As an interesting main finding, the studies analyzed in the three reviews, showed different post-interventional plaque control strategies with regard to chemical and mechanical plaque control regimens among the different treatment groups. For instance, 1 to 2 weeks reflects a reasonable time span after surgical therapy until sutures are removed. Other time points were adjusted to 1 and 2 months. The regenerative surgical approach showed the highest degree of maintenance efforts after the intervention. An explanation for the continuous monitoring is the nature of regenerative therapy since this therapy is invasive and expensive. An explanation for the continuous monitoring is the nature of regenerative therapy since this therapy is invasive and expensive. Nevertheless, a prospective clinical study on patient undergoing one-stage full-mouth scaling and root planing has demonstrated a statistical significant benefit in probing depth and clinical attachment gain after 3 months of extensive use of CHX mouth rinse. [26].

Due to the fact that all studies included in this study were part of peer-reviewed reviews, the evaluation of outcome measurements such as PPD and CAL was not being weighed against each other, only the mean PPD difference on the regenerative studies presented in the forest plot. However, one systematic review did not meet the current standard requirements for systematic reviews due to an earlier publishing date. Hence, a new search was undertaken to compensate for the lack of up-to-dateness. In addition, the classification to evaluate the maintenance protocol was arbitrarily set, which might be considered a shortcoming of the present study. However, it reflects potentially relevant time frames in the course of periodontal therapy.

Postoperative success is determined by many factors, such as anatomical and technical factors, patient compliance, plaque control and cigarette smoking. All these are factors that can directly affect the predictability of periodontal regeneration [27]. Thus, low plaque scores have shown to reduce the risk of membrane exposure, infection and guarantee better complication management [25, 28]. These factors inevitably also lead to more stringent protocols, which is mainly justified by infection control and healing optimization. Common procedure such as the intake of adjunctive antibiotics or anti-inflammatory medication during regeneration could also be one factor for a favorable outcome.

The importance of postoperative plaque control in determining the outcome of periodontal surgery is well established and recognized in the literature for a long time [29].

In contrast, studies using systemic antibiotics as an adjunct to SRP disclosed an opposite tendency. Almost

| Table 2 Detailed list of analyzed studies dealing with non-surgical periodontal therapy with adjunctive use of amoxicillin and metronidazole [16] (Continued) |
|-----------------|-------|-----|-------|--------|-----------------|-----------------|-----------------|-----------------|
| Goodson et al. 2012 | CHX   | 0.12 % | 2x/d  | 2 weeks | A | yes | 3, 6, 12, 18, 24 months | 2 |
| Oliveira et al. 2012 | CHX   | 0.12 % | 2x/d  | 60 days  | C | n.r. | clinical measurements at 6 months | 1 |

clinical measurements at 3 and 6 months
Table 3 Detailed list of analyzed studies dealing with regenerative periodontal therapy, i.e. GTR [17]

| Study                  | Antiseptic | Galenic formulation | Application frequency | Duration | Category | Supragingival cleaning | Intervals (weeks)                  | Category |
|------------------------|------------|---------------------|-----------------------|----------|----------|------------------------|------------------------------------|----------|
| Yukna et al. 1985      | CHX        | n.r. n.r.           | n.r. n.r.             |          |          | yes                    | at 10, 20 and 30 days and 3, 6, 9 and 12 months | 2        |
| Cortellini et al. 1995a| CHX 0.2 %  | 2x/d 3 weeks        | B                     |          |          | yes                    | 1, 2, 3, 4, 5, 6 (and after membran removal 1, 2, 3, 4) | 2        |
| Mora & Ouhayoun 1995   | CHX 0.2 %  | n.r. 10 days        | A                     |          |          | yes                    | every 10 days for the 1st month, 2x a month for months 2 and 3 and then every 3 months for 9 months | 2        |
| Cortellini et al. 1996 | CHX 0.2 %  | 2x/d 3 weeks in the MWF group 2 months bioresorbable gr 6 weeks in the ePTFE gr |          |          |          | yes                    | 1, 2, 3, 4, 5, 6, then every month for 12 months | 2        |
| Tonetti et al. 1996   | CHX 0.2 %  | 2x/d 3 weeks        | B                     |          |          | yes                    | weekly for 6 weeks. After membran removal weekly for 4 weeks, then monthly up to 12 months | 2        |
| Mora et al. 1996       | CHX 0.2 %  | 2x/d 4 weeks        | B                     |          |          | yes                    | weekly 4 weeks, professional maintenance regimen after membran removal until 12 month | 2        |
| Zarnet et al. 1997     | CHX 0.2 %  | 2x/d 4 weeks        | B                     |          |          | yes                    | 1, 2, 3, 4, then every 4 weeks up to 3 months and then every 3 months up to 1 yr | 2        |
| Heijl et al. 1997      | CHX 0.2 %  | n.r. 4–6 weeks      | C                     | n.r. n.r.|          | yes                    | n.r. | weekly for 4 weeks, then monthly until 6 month, then every 4–6 weeks up to 1 yr | 2        |
| Mayfield et al. 1998   | CHX 0.2 %  | 2x/d 6 weeks (test) 3 w (controlgr.) |          |          |          | yes                    | 2, 4, 6, 8, 12 weeks | 2        |
| Okuda et al. 2000      | CHX 0.12 % | 3x/day 6 weeks      | C                     |          |          | yes                    | weekly for 6 weeks, then monthly up to 12 months | 2        |
| Zybutz et al. 2000     | CHX 0.12 % | 2x/d 8 weeks        | C                     |          |          | yes                    | 2, 4, 6, 8, 12 weeks | 2        |
| Ratka-Kruger et al. 2000 | CHX 0.2 % | 3x/day 6 weeks      | C                     |          |          | yes                    | at 3, 6 and 12 months | 1        |
| Sculean et al. 2001    | CHX 0.2 %  | 2x/d 6 weeks        | C                     |          |          | yes                    | every 2nd week for 2 months, then monthly up to 12 months | 2        |
| Zucchelli et al. 2002  | CHX 0.12 % | 2x/d 11 weeks      | C                     |          |          | yes                    | Weekly for 11 weeks, then monthly up to 1 yr | 2        |
| Tonetti et al. 2002    | CHX 0.12 % | 2x/d 4 weeks        | B                     |          |          | yes                    | at weeks 1, 2, 3, 4, 6. Then 3, 6, 9 month recall | 2        |
| Wachtel et al. 2003    | CHX 0.2 %  | 2x/d 4 weeks        | B                     | n.r. n.r.|          | yes                    | n.r. | | 2        |
| Sculean et al. 2003    | CHX 0.2 %  | 2x/d 4 weeks        | B                     |          |          | yes                    | every 2nd week for 2 months, then monthly up to 12 months | 2        |
| Sculean et al. 2004    | CHX 0.2 %  | 2x/d 6 weeks        | C                     |          |          | yes                    | every 2nd week for 2 months, then monthly for the 1st yr, after 1st yr for 5 yrs every 3 months | 2        |
| Tonetti et al. 2004    | CHX 0.12 % | 2x/d 4 weeks        | B                     |          |          | yes                    | at weeks 1, 2, 3, 4, 6 and 8. Then at 3, 6 and 9 months | 2        |
| Vouros et al. 2004     | CHX 0.12 % | 2x/d 4 weeks        | B                     |          |          | yes                    | monthly up to 12 months | 1        |
| Francetti et al. 2004  | CHX 0.12 % | 2x/d 6 weeks        | C                     |          |          | yes                    | recall monthly during 1 yr, then at 18 and 24 month | 1        |
no chemical plaque control was done and patients were left unsupervised until reevaluation after 3 months. Non-surgical therapy with systemic antibiotics is considered to be a more cost-effective treatment alternative in contrast to sophisticated regenerative surgery. Its aim is to reduce the need for any surgical therapy [30, 31]. In addition, less postoperative complications may be expected due to the fact that neither surgery has been performed nor foreign materials have been implanted. Quite to the contrary, patients were only under the guard of antibiotics. Overall, two different periodontal procedures with their specific therapy goals, extend of treatment site as well as different healing needs make it challenging to compare and evaluate the results. However, plaque scores after 3 months were quite high in some studies and reached a plaque index of above 30 % at reevaluation [32–34]. Some studies did not even report on plaque indices, which made a more detailed assessment of this important data impossible. Therefore, it remains unclear to what extent decreased plaque levels would have led to a better clinical outcome. In contrast, evidence suggests that the occurrence of re-established plaque may lead to recolonization, less healing and persistence of the original pathology [35, 36].

Missing quantitative data on probing depth reductions at the requested time points also made it impossible to assess and compare the results of the non-surgical interventions and include the data in a forest-plot. Therefore, only adequately reported GTR studies could be included into this analysis.

Periodontal sites, which could be influenced, were suprabony and infrabony defects as well as pockets with furcation involvement. Inarguably, the role and potential of adequate plaque control during therapy and afterwards have an impact on the subgingival microbiota [37]. The importance of an adequate maintenance protocol for the success or failure in periodontal therapy has therefore been introduced as an achievable goal for decades [38].

Table 3 Detailed list of analyzed studies dealing with regenerative periodontal therapy, i.e. GTR [17] (Continued)

| Study             | Antiseptic | Duration | Protocol | Plaque Control | Recall Frequency | Notes                                                                 |
|-------------------|------------|----------|----------|----------------|-----------------|----------------------------------------------------------------------|
| Francetti et al.  | CHX 0.12 % | 2x/d     | 6 weeks  | C n.r.         | n.r.            | 1,2,3,4, then every month for 12 months                               |
| Aimetti et al.    | CHX 0.2 %  | 2x/d     | 3 weeks  | B yes          | 1,2,3,4, then every 2nd week, then monthly up to 12 months          |
| Sculean et al.    | CHX 0.2 %  | 2x/d     | 4 weeks  | B yes          | every 2nd week for 2 months, then monthly up to 12 months          |
| Sculean et al.    | CHX 0.2 %  | 2x/d     | 4 weeks  | B yes          | every 2nd week for 2 months, then monthly up to 12 months          |
| Sculean et al.    | CHX 0.2 %  | 2x/d     | 6 weeks  | C yes          | every 2nd week for 2 months, then once per month for 1 yr, then every 3 months for 10 yrs |
| Kasaj et al. 2008 | CHX 0.2 %  | 2x/d     | 4 weeks  | B yes          | every 2 week for 2 months, after every 4 weeks                     |
| Fickl et al. 2009 | n.r.       |          |          |                | n.r.             |                                                                     |
| Stein et al. 2009 | CHX 0.2 %  | 2x/d     | 4 weeks  | B yes          | every 2nd week for 2 months, then monthly up to 12 months          |
| Cortellini et al. | CHX 0.12 % | 2x/d     | 4 weeks  | C yes          | 1,2,3,4,5,6, then 3 month recall for 1 yr                          |

Table 4 Comparisons of the different maintenance programs according to the use of antiseptic rinsing (CHX 0.06-0.2 %) and recall frequency (number of visits)

| Protocol | Duration of the use of antiseptics CHX | Number of recalls within the first 2 months |
|----------|----------------------------------------|--------------------------------------------|
|          | First 2 weeks (A) | Up to 4 weeks (B) | Up to 8 weeks (C) | \( \leq 2 \) visits (1) | \( \geq 3 \) visits (2) |
| Non-surgical only (n = 14) | 4 | - | 1 | 5 | 9 |
| non-surgical plus systemic antibiotics (n = 34) | 10 | - | 7 | 20 | 2 |
| Regenerative surgical procedures (n = 30) | 1 | 15 | 11 | 3 | 23 |

Duration of antiseptic use CHX: A = two or less than 2 weeks; B = up to 4 weeks; C = more than 5 weeks. Number of recall visits following periodontal treatment within the first 2 months; Protocol: 1 = two or less visits; 2 = three or more recall appointments within the first 2 months.
Conclusion

By tendency, regenerative studies showed a longer duration of anti-septic mouth rinse and higher quantity of maintenance protocol compared to non-surgical approaches. However, sophisticated treatment should not be a causal reason for vigorous recall intervals more than an evidence based reason. Till today there is little evidence on how elaborate a post treatment or postoperative protocol should be in order to benefit the patient. Carefully executed prospective studies on this topic are still warranted.

Abbreviations

CAL, clinical attachment level; GTR, guided tissue regeneration; PI, plaque Index; PPD, probing pocket depth; PRISMA, referred reporting items for systematic review and meta-analyses; RCT, randomized clinical control trail; REC, recession; SPT, supportive periodontal treatment; SRP, scaling and root planing

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Availability of data and materials

All relevant data supporting the conclusion of this article are within the manuscript.

Authors’ contributions

PRS planned the study. SG, ID, PS carried out the data selection and measurement methods. UH conducted the statistical analysis. SG, ID, PS, PRS and CW wrote the manuscript. All authors have read and approved the manuscript and have agreed to its submission.

Competing interests

The authors declare that they have no competing interests.

Consent for publication

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Ethics approval and consent to participate

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References

1. Teles RP, Haffajee AD, Socransky SS. Microbiological goals of periodontal therapy. Periodontol 2000. 2006;42:180–218.
2. Page RC, Kornman KS. The pathogenesis of human periodontitis: an introduction. Periodontol 2000. 1997:149–11.
3. Axelsson P, Nyström B, Lindhe J. The long-term effect of a plaque control program on tooth mortality, caries and periodontal disease in adults. Results after 30 years of maintenance. J Clin Periodontol. 2004;31:749–57.
4. Greenstein G. Nonsurgical periodontal therapy in 2000: a literature review. J Am Dent Assoc. 2000;131:1580–92.
5. Wennstrom JL, Dahlen G, Ramberg P. Subgingival debridement of periodontal pockets by air polishing in comparison with ultrasonic instrumentation during maintenance therapy. J Clin Periodontol. 2001;28:820–7.
6. Hagi TT, Kleemersberger S, Bereiter R, Nietzche G, Cosgarea R, Flury S, et al. A biofilm pocket model to evaluate different Non-surgical periodontal treatment modalities in terms of biofilm removal and reformation, surface alterations and attachment of periodontal ligament fibroblasts. PLoS One. 2015;10:e0131056.
7. Badenster A, Nileveus R, Egelberg J. Effect of nonsurgical periodontal therapy. II. Severely advanced periodontitis. J Clin Periodontol. 1984;1:63–76.

8. Lang NP. Indications and rationale for non-surgical periodontal therapy. Int Dent J. 1983;33:127–36.

9. Guerrero A, Nibali L, Lambertenghi R, Ready D, Suvan J, Griffiths GS, et al. Impact of baseline microbiological status on clinical outcomes in generalized aggressive periodontitis patients treated with or without adjunctive amoxicillin and metronidazole: an exploratory analysis from a randomized controlled clinical trial. J Clin Periodontol. 2014;41:1080–9.

10. Sgolastra F, Petrucci A, Gatto R, Monaco A. Effectiveness of systemic amoxicillin/metronidazole as an adjunctive therapy to full-mouth scaling and root planing in the treatment of aggressive periodontitis: a systematic review and meta-analysis. J Periodontol. 2012;83:731–43.

11. Matuliene G, Pjetursson BE, Salvi GE, Schmidlin K, Bragger U, Zvahnoven M, et al. Influence of residual pockets on progression of periodontitis and tooth loss: results after 11 years of maintenance. J Clin Periodontol. 2008;35:685–95.

12. Walter C, Weiger R, Zitthmann NJ. Periodontal surgery in function-involved maxillary molars revisited—a introduction of guidelines for comprehensive treatment. Clin Oral Investig. 2011;15:9–20.

13. Tonetti MS, Muller-Campenale V, Lang NP. Changes in the prevalence of residual pockets and tooth loss in treated periodontal patients during a supportive maintenance care program. J Clin Periodontol. 1998;25:1008–16.

14. Santos S, Herrera D, Lopez E, O’Connor A, Gonzalez I, Sanz M. A randomized clinical trial on the short-term clinical and microbiological effects of the adjunctive use of a 0.05 % chlorhexidine mouth rinse for patients in supportive periodontal care. J Clin Periodontol. 2004;31:45–51.

15. Heitz-Mayfield LJ, Trombelli L, Heitz F, Needleman I, Mole S. A systematic review of the effect of surgical debridement vs non-surgical debridement for the treatment of chronic periodontitis. J Clin Periodontol. 2002;29(Suppl 3):92–102. discussion 60–2.

16. Zandbergen D, Slot DE, Cobb CM, Van der Weijden FA. The clinical effect of scaling and root planing and the concomitant administration of systemic amoxicillin and metronidazole: a systematic review. J Periodontol. 2013;84:332–51.

17. Graziani F, Gennai S, Cei S, Cairo F, Baggiani A, Miccoli M, et al. Clinical performance of access flap surgery in the treatment of the intrabony defect. A systematic review and meta-analysis of randomized clinical trials. J Clin Periodontol. 2012;39:145–56.

18. Moher D, Liberati A, Tetzlaff J, Altman DG, Group P. Reprint and we will help you at every step:

19. R Development Core Team. R: a language and environment for statistical computing. Vienna: R Foundation for Statistical Computing; 2015.

20. Serino G, Rosling B, Ramberg P, Socransky SS, Lindhe J. Initial outcome and long-term effect of surgical and non-surgical treatment of advanced periodontal disease. J Clin Periodontol. 2004;29:910–2.

21. Flemmig TF, Milian E, Karch H, Klaiber B. Differential clinical treatment options and we will help you at every step:

22. Ehmke B, Beikler T, Haubitz I, Karch H, Flemmig TF. Multifactorial assessment of predictors for prevention of periodontal disease progression. Clin Oral Investig. 2003;7:217–21.

23. Ehmke B, Moter A, Beikler T, Milian E, Flemmig TT. Adjunctive antimicrobial therapy of periodontitis: long-term effects on disease progression and oral colonization. J Periodontol. 2005;76:749–59.

24. Moira F, Ouhayoun JP. Clinical evaluation of natural coral and porous hydroxyapatite implants in periodontal bone lesions: results of a 1-year follow-up. J Clin Periodontol. 1995;22:877–84.

25. Zucchelli G, Bernardi F, Montebugnoli L, De SM. Enamel matrix proteins and we will help you at every step:

26. Stratul SI, Rusu D, Didilescu A, Mesars-Anghel M, Lala C, Tion L, et al. Prospective clinical study evaluating the long-time adjunctive use of chlorhexidine after one-stage full-mouth SRP. Int J Dent Hyg. 2011;8:35–40.

27. Cortellini P, Bowers GM. Periodontal regeneration of intrabony defects: an evidence-based treatment approach. Int J Periodontics Restorative Dent. 1995;15:128–45.