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Autologous platelet concentrates for treating periodontal infrabony defects (Review)

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[Intervention Review]

Autologous platelet concentrates for treating periodontal infrabony defects

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ABSTRACT

Background

Periodontal disease is a condition affecting tooth supporting tissues (gingiva, alveolar bone, periodontal ligament, and cementum), with the potential of introducing severe adverse effects on oral health. It has a complex pathogenesis which involves the combination of specific micro-organisms and a predisposing host response. Infrabony defects are one of the morphological types of alveolar bone defects that can be observed during periodontitis. Recent approaches for the treatment of infrabony defects, combine advanced surgical techniques with platelet-derived growth factors. These are naturally synthesized polypeptides, acting as mediators for various cellular activities during wound healing. It is believed that the adjunctive use of autologous platelet concentrates to periodontal surgical procedures produces a better and more predictable outcome for the treatment of infrabony defects.

Objectives

To assess the effects of autologous platelet concentrates (APC) used as an adjunct to periodontal surgical therapies (open flap debridement (OFD), OFD combined with bone grafting (BG), guided tissue regeneration (GTR), OFD combined with enamel matrix derivative (EMD)) for the treatment of infrabony defects.

Search methods

Cochrane Oral Health's Information Specialist searched the following databases: Cochrane Oral Health's Trials Register (to 27 February 2018); the Cochrane Central Register of Controlled Trials (CENTRAL; 2018, Issue 1) in the Cochrane Library (searched 27 February 2018); MEDLINE Ovid (1946 to 27 February 2018); Embase Ovid (1980 to 27 February 2018); and LILACS BIREME Virtual Health Library (from 1982 to 27 February 2018). The US National Institutes of Health Ongoing Trials Register (ClinicalTrials.gov) and the World Health Organization International Clinical Trials Registry Platform were searched for ongoing trials on 27 February 2018. No restrictions were placed on the language or date of publication when searching the electronic databases.

Selection criteria

We included randomised controlled trials (RCTs) of both parallel and split-mouth design, involving patients with infrabony defects requiring surgical treatment. Studies had to compare treatment outcomes of a specific surgical technique combined with APC, with the same technique when used alone.

Data collection and analysis

Two review authors independently conducted data extraction and risk of bias assessment, and analysed data following Cochrane methods. The primary outcomes assessed were: change in probing pocket depth (PD), change in clinical attachment level (CAL), and change in radiographic bone defect filling (RBF). We organised all data in four groups, each comparing a specific surgical technique when applied with the adjunct of APC or alone: 1. APC + OFD versus OFD, 2. APC + OFD + BG versus OFD + BG, 3. APC + GTR versus GTR, and 4. APC + EMD versus EMD.

Main results

We included 38 RCTs. Twenty-two had a split-mouth design, and 16 had a parallel design. The overall evaluated data included 1402 defects. Two studies were at unclear overall risk of bias, while the remaining 36 studies had a high overall risk of bias.

1. APC + OFD versus OFD alone

Twelve studies were included in this comparison, with a total of 510 infrabony defects. There is evidence of an advantage in using APC globally from split-mouth and parallel studies for all three primary outcomes: PD (mean difference (MD) 1.29 mm, 95% confidence interval (CI) 1.00 to 1.58 mm; P < 0.001; 12 studies; 510 defects; very low-quality evidence); CAL (MD 1.47 mm, 95% CI 1.11 to 1.82 mm; P < 0.001; 12 studies; 510 defects; very low-quality evidence); and RBF (MD 34.26%, 95% CI 30.07% to 38.46%; P < 0.001; 9 studies; 401 defects; very low-quality evidence).

2. APC + OFD + BG versus OFD + BG

Seventeen studies were included in this comparison, with a total of 569 infrabony defects. Considering all follow-ups, as well as 3 to 6 months and 9 to 12 months, there is evidence of an advantage in using APC from both split-mouth and parallel studies for all three primary outcomes: PD (MD 0.54 mm, 95% CI 0.33 to 0.75 mm; P < 0.001; 17 studies; 569 defects; very low-quality evidence); CAL (MD 0.72 mm, 95% CI 0.43 to 1.00 mm; P < 0.001; 17 studies; 569 defects; very low-quality evidence); and RBF (MD 8.10%, 95% CI 5.26% to 10.94%; P < 0.001; 11 studies; 420 defects; very low-quality evidence).

3. APC + GTR versus GTR alone

Seven studies were included in this comparison, with a total of 248 infrabony defects. Considering all follow-ups, there is probably a benefit for APC for both PD (MD 0.92 mm, 95% CI -0.02 to 1.86 mm; P = 0.05; very low-quality evidence) and CAL (MD 0.42 mm, 95% CI -0.02 to 0.86 mm; P = 0.06; very low-quality evidence). However, given the wide confidence intervals, there might be a possibility of a slight benefit for the control. When considering a 3 to 6 months and a 9 to 12 months follow-up there were no benefits evidenced, except for CAL at 3 to 6 months (MD 0.54 mm, 95% CI 0.18 to 0.89 mm; P = 0.003; 3 studies; 134 defects). No RBF data were available.

4. APC + EMD versus EMD

Two studies were included in this comparison, with a total of 75 infrabony defects. There is insufficient evidence of an overall advantage of using APC for all three primary outcomes: PD (MD 0.13 mm, 95% CI -0.05 to 0.30 mm; P = 0.16; 2 studies; 75 defects; very low-quality evidence); CAL (MD 0.10 mm, 95% CI -0.13 to 0.32 mm; P = 0.40; 2 studies; 75 defects; very low-quality evidence), and RBF (MD -0.80%, 95% CI -6.21% to 5.01%; P = 0.83; 1 study; 49 defects; very low-quality evidence).

All studies in all groups reported a survival rate of 100% for the treated teeth. No complete pocket closure was reported. No quantitative analysis regarding patients' quality of life was possible.

Authors' conclusions

There is very low-quality evidence that the adjunct of APC to OFD or OFD + BG when treating infrabony defects may improve probing pocket depth, clinical attachment level, and radiographic bone defect filling. For GTR or EMD, insufficient evidence of an advantage in using APC was observed.

PLAIN LANGUAGE SUMMARY

Autologous platelet concentrates for treating periodontal infrabony defects

Review question

Does the addition of autologous platelet concentrates (APC) improve surgical treatment outcomes of bone defects in gum disease?

Background

Teeth are maintained in their position by soft and hard tissues (gums and surrounding bone). Gum disease or periodontitis, is an inflammatory condition of all these tissues caused by the bacteria present in the dental plaque. If left untreated, gum disease can cause teeth to loosen and eventually lead to tooth loss. The destruction of jaw bone around teeth (called the alveolar bone) during gum disease, can be horizontal (where the whole level of bone around the root is reduced) or vertical, forming a bone defect within the bone (infrabony defect). There are several available surgical treatments for infrabony defects, including: 1. open flap debridement in which the gum is lifted back surgically in order to clean the deep tartar; 2. bone graft in which a portion of natural or synthetic bone is placed in the area of bone loss; 3. guided tissue regeneration in which a small piece of membrane-like material is placed between the bone and gum tissue in order to keep the gum tissue from growing into the area where the bone should be; and 4. the use of enamel matrix derivative, a gel-like material which is placed in the area where bone loss has occurred and promotes its regeneration. In order to accelerate the healing process, autologous platelet concentrates have been recently used. They are concentrates of the platelets of patient's own blood containing growth factors that are thought to promote tissue regeneration. The aim of this review was to assess if the addition of APC brings any benefits in the treatment of infrabony defects when combined with different surgical treatments.

Study characteristics

Authors from Cochrane Oral Health carried out this review and the evidence is up to date to 27 February 2018. We included 38 studies and a total of 1042 infrabony defects. We considered four different types of surgical treatments and compared each technique with the same one when APC was added. Overall we considered three comparisons: open flap debridement with APC versus without APC; open flap debridement and bone graft with APC versus without APC; guided tissue regeneration with APC versus without APC; and enamel matrix derivative with APC versus without APC.

Key results

There is very low-quality evidence that the addition of APC to two types of treatment: open flap debridement and open flap debridement with bone graft, may bring some advantages in the treatment of infrabony defects. However, for the other two types of treatment, guided tissue regeneration and enamel matrix derivative, there is insufficient evidence of a benefit.

Quality of evidence

We judged the quality of the evidence to be very low due to problems with the design of the studies.

SUMMARY OF FINDINGS FOR THE MAIN COMPARISON [Explanation]

APC + OFD compared to OFD (9-12 months follow-up) for treating periodontal infrabony defects

Patient or population: patients affected by infrabony defects requiring surgical treatment
Settings: tertiary care
Intervention: APC + OFD
Comparison: OFD

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	Number of participants/defects (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	OFD	APC + OFD				
Change in probing depth (PD) (mm) (9-12 months follow-up)	Mean PD change (gain) across control groups ranged from 2.40 to 3.68 (2.36) mm Mean PD baseline value was 7.92 mm (95% CI 6.25 to 9.54)	The mean PD change (gain) in the intervention groups was **1.29 mm higher** (1.00 to 1.58 higher)	Mean difference 1.29 (1.00 to 1.58) mm	510 (12 studies)	⊕◯◯◯ **very low**[1,2]	There is evidence of an advantage in using APC
Change in clinical attachment level (CAL) (mm) (9-12 months follow-up)	Mean CAL change (gain) across control groups ranged from 1.27 to 4.14 (2.03) mm Mean CAL baseline value was 6.78 mm (95% CI 5.56 to 7.54)	The mean CAL change (gain) in the intervention groups was **1.47 mm higher** (1.11 to 1.82 higher)	Mean difference 1.47 (1.11 to 1.82) mm	510 (12 studies)	⊕◯◯◯ **very low**[1,2]	There is evidence of an advantage in using APC
Change in radiographic bone defect filling (RBF) (%) (9-12 months follow-up)	Mean RBF change (gain) across control groups ranged from -3.60% to 54.20% (16.90%)	The mean RBF change (gain) in the intervention groups was **34.26% higher** (30.07 to 38.46 higher)	Mean difference 34.26% (30.07 to 38.46)	401 (9 studies)	⊕◯◯◯ **very low**[1,2]	There is evidence of an advantage in using APC

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% CI) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).
APC: autologous platelet concentrates; **CAL:** clinical attachment level; **CI:** confidence interval; **OFD:** open flap debridement; **PD:** probing depth; **RBF:** radiographic bone defect filling.

GRADE Working Group grades of evidence
High quality: further research is very unlikely to change our confidence in the estimate of effect.
Moderate quality: further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
Low quality: further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: we are very uncertain about the estimate.

[1]Downgraded by 2 levels for high risk of performance bias.
[2]Downgraded by 2 levels for high heterogeneity.

BACKGROUND

Description of the condition

Periodontitis is a disease of the periodontium characterized by the irreversible loss of connective tissue attachment and supporting alveolar bone (Pihlstrom 2005). For its onset, the presence of specific micro-organisms together with an altered response of the host, are necessary. Despite its many variations, a typical course of periodontitis starts with pocket formation induced by bacterial plaque and a subsequent alveolar bone destruction typical of chronic periodontitis. Bone destruction during periodontitis can be of different morphological patterns including suprabony (horizontal) defects and infrabony (vertical) defects (Kinane 2001). An infrabony defect represents the anatomic sequelae resulting from the apical advancement of the dental plaque during the progression of the disease (Waerhaug 1979). Such defects, if left untreated, easily promote periodontitis progression and further loss of attachment (Papapanou 1991). Because infrabony defects are common in periodontitis (Vrotsos 1999), there is a considerable interest in approaches that will convert such defects, at risk for disease progression, to easily maintainable shallow probing sites (Crea 2014).

Description of the intervention

The ultimate goal of periodontal therapy is to preserve the natural dentition for as long as possible and enhance patient's comfort and aesthetic features by maintaining and improving the health and function of all tooth-supporting tissues (gingiva, periodontal ligament, cementum, alveolar bone). Conventional treatment of periodontal disease may arrest bone destruction but usually does not restore the already lost alveolar bone or periodontal connective tissue. Various surgical techniques have been developed as an attempt to provide an efficient treatment to periodontitis. Open flap debridement (OFD) is among the earliest and most promising procedures to be used (Caffesse 1986; Cortellini 1996). Its main objective is to reduce the presence of micro-organisms which develop and maintain the inflammatory process. By doing so, it consequently promotes the regeneration properties of the host, despite not being a regenerative procedure. Later, the combination of conventional OFD with various biomaterials such as bone grafts, enamel matrix derivative or membranes (guided tissue regeneration), resulted in the development of regenerative treatment protocols which introduced significant clinical benefits (Cochran 2003; Cortellini 1996; Esposito 2009; Hoidal 2008; Needleman 2006).

Despite advances in surgical procedures and materials, a complete and predictable regeneration, defined as the development of new bone, periodontal ligament and cementum on a root surface previously exposed to periodontal disease, remains a challenge (AAP 1992). Consequently, the concept of tissue engineering (Rose 2002) which requires the presence of cells, scaffold and signalling molecules, gained particular attention in terms of periodontal regeneration. Bone grafts and membranes used in guided tissue regeneration (GTR) can serve as scaffolds but there always exists a need of signalling molecules.

Recently, polypeptide growth factors have been investigated as possible signalling factors for enhancing periodontal regeneration. As preliminary evidence for their potential applications in periodontal wound healing, several polypeptide growth factors have been identified in the human periodontal tissues by immuno-histochemistry and in-situ hybridization (Giannobile 1996). An abundant source of such growth factors are platelets, easily utilisable in the form of autologous platelet concentrates (APC). Therefore, the adjunctive use of APC in combination with periodontal surgery has emerged as a possible tool to enhance the predictability of infrabony defects treatment.

APC, based on their preparation protocol, can be of various types, including platelet-rich plasma (PRP) (Marx 1998), platelet-rich fibrin (PRF) (Choukroun 2001), and plasma-rich growth factors (PRGF) (Anitua 2001). Several commercial techniques for obtaining platelet concentrates are available. However, their indication of use has been confusing because each method leads to a different product with different biological properties and possible applications. PRP represents the first generation of platelet concentrate, and shows a release of an array of growth factors for 7 days, with a peak release on its first day of application (Dohan Ehrenfest 2009). PRF represents the second generation APC, and its technique of preparation is simplified when compared to PRP. Moreover, PRF showed a sustained growth factors release for a period of 21 days with a peak release at 7 days (Carroll 2005). PRGF is also a second generation platelet concentrate, whose main difference when compared to PRP is the absence of leucocytes and the small blood volume required for its preparation (Anitua 2001). Following an upgrade in their classification (Dohan Ehrenfest 2009), platelet concentrates can be divided into four categories, based on the presence of leucocyte and fibrin: P-PRP (pure PRP, without leucocytes, which includes PRGF), L-PRP (leucocyte and platelet-rich plasma), P-PRF (pure PRF), and L-PRF (leucocyte PRF).

How the intervention might work

The contribution of blood-derived platelets to the bone healing process is thought to be based on the growth factors stored in their granules and released upon activation. The main growth factors released from platelet aggregates are the following: platelet derived growth factor (PDGF), transforming growth factor-beta (TGF-β), vascular endothelial growth factor (VEGF), epithelial growth factor (EGF), insulin-like growth factor-1 (IGF-1), and basic fibroblast growth factor (bFGF), as well as three blood proteins known to act as cell adhesion molecules for osteo-conduction (fibrin, fibronectin and vitronectin). The set of these factors serve as

biological mediators with the ability to regulate cell proliferation, chemotaxis, and differentiation.

Why it is important to do this review

The considerably increased interest in combining APC with surgical techniques for better outcomes in the treatment of infrabony defects, has made it necessary a thorough investigation of the actual benefits that can be obtained. The first systematic review that evaluated the effect of PRP on clinical applications in dentistry reported beneficial effects of PRP in the treatment of periodontal defects (Plachokova 2008). Another systematic review that evaluated the effect of a PRP adjunct in treatment of intraosseous defects, underlined the limits and the heterogeneity of available data and cautiously concluded that the specific selection of the graft type and the surgical procedures combined with PRP may be important (Kotsovilis 2010). A subsequent systematic review also evaluated the effect of platelet rich plasma in various regenerative procedures of periodontal defects, and concluded that PRP may be advantageously used as an adjunct to grafting procedures treatment for infrabony defects (Del Fabbro 2011). Such review also suggested that the use of PRP is ineffective when GTR procedure is used for treating infrabony defects.

Despite the numerous reports on the adjunctive use of autologous platelet concentrate to periodontal surgical procedures, its efficacy remains controversial. This is partly due to a large heterogeneity among different studies (Del Fabbro 2011; Del Fabbro 2013), concerning methods, study design, protocols for platelet concentrate preparation, participants selection criteria, outcome variables assessed, etc. Therefore, a review of the current state of the evidence is crucial in order to clarify whether the clinical use of APCs eventually produces better outcomes in the treatment of infrabony defects, and if their effect is particularly enhanced when combined with a specific surgical technique. By doing so, clear and relevant guidelines can be addressed to clinicians.

OBJECTIVES

To assess the effects of autologous platelet concentrates used as an adjunct to periodontal surgical therapies (open flap debridement (OFD), OFD combined with bone grafting, guided tissue regeneration, OFD combined with enamel matrix derivative) for the treatment of infrabony defects.

METHODS

Criteria for considering studies for this review

Types of studies

Randomised controlled trials, of both parallel and split-mouth design.

Types of participants

Patients affected by infrabony defects requiring surgical treatment, regardless of their age or gender.

Types of interventions

Experimental intervention: autologous platelet concentrates (APCs) (irrespective of the type: platelet-rich plasma (PRP), plasma-rich growth factors (PRGF), or platelet-rich fibrin (PRF)) used in conjunction with a specific surgical technique (open flap debridement (OFD), OFD + bone grafts (BG), guided tissue regeneration (GTR), enamel matrix derivative (EMD)).

Comparison (control) intervention: the same surgical techniques when used alone (without the adjunct of APCs).

Types of outcome measures

Primary outcomes

Change in probing depth (PD), change in clinical attachment level (CAL), and change in radiographic bone defect filling (RBF).

Secondary outcomes

Tooth survival, pocket closure, and oral health-related quality of life.

Search methods for identification of studies

Electronic searches

Cochrane Oral Health's Information Specialist conducted systematic searches in the following databases for randomised controlled trials and controlled clinical trials. There were no language, publication year or publication status restrictions:

- Cochrane Oral Health's Trials Register (searched 27 February 2018) (Appendix 1);
- Cochrane Central Register of Controlled Trials (CENTRAL; 2018, Issue 1) in the Cochrane Library (searched 27 February 2018) (Appendix 2);
- MEDLINE Ovid (1946 to 27 February 2018) (Appendix 3);
- Embase Ovid (1980 to 27 February 2018) (Appendix 4);
- LILACS BIREME Virtual Health Library (Latin American and Caribbean Health Science Information database; 1982 to 27 February 2018) (Appendix 5).

Subject strategies were modelled on the search strategy designed for MEDLINE Ovid. Where appropriate, they were combined with subject strategy adaptations of the highly sensitive search strategy designed by Cochrane for identifying randomised controlled trials and controlled clinical trials as described in the *Cochrane Handbook for Systematic Reviews of Interventions* Chapter 6 (Lefebvre 2011).

Searching other resources

The following trial registries were searched for ongoing studies:

- US National Institutes of Health Ongoing Trials Register ClinicalTrials.gov (clinicaltrials.gov; searched 27 February 2018) (Appendix 6);
- World Health Organization International Clinical Trials Registry Platform (apps.who.int/trialsearch; searched 27 February 2018) (Appendix 7).

An adjunctive search was performed on the reference lists of the included articles and reviews retrieved.

Moreover, a handsearch was performed on the issues since January 2010 (including the 'early view' or equivalent section) of the following journals: *International Journal of Periodontics and Restorative Dentistry, Journal of Clinical Periodontology, Journal of Periodontal Research, Journal of Periodontology, Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology and Endodontology* (last search was performed on 2 March 2018). Two review authors independently performed the searches (Saurav Panda (SP), Cristina Bucchi (CB)).

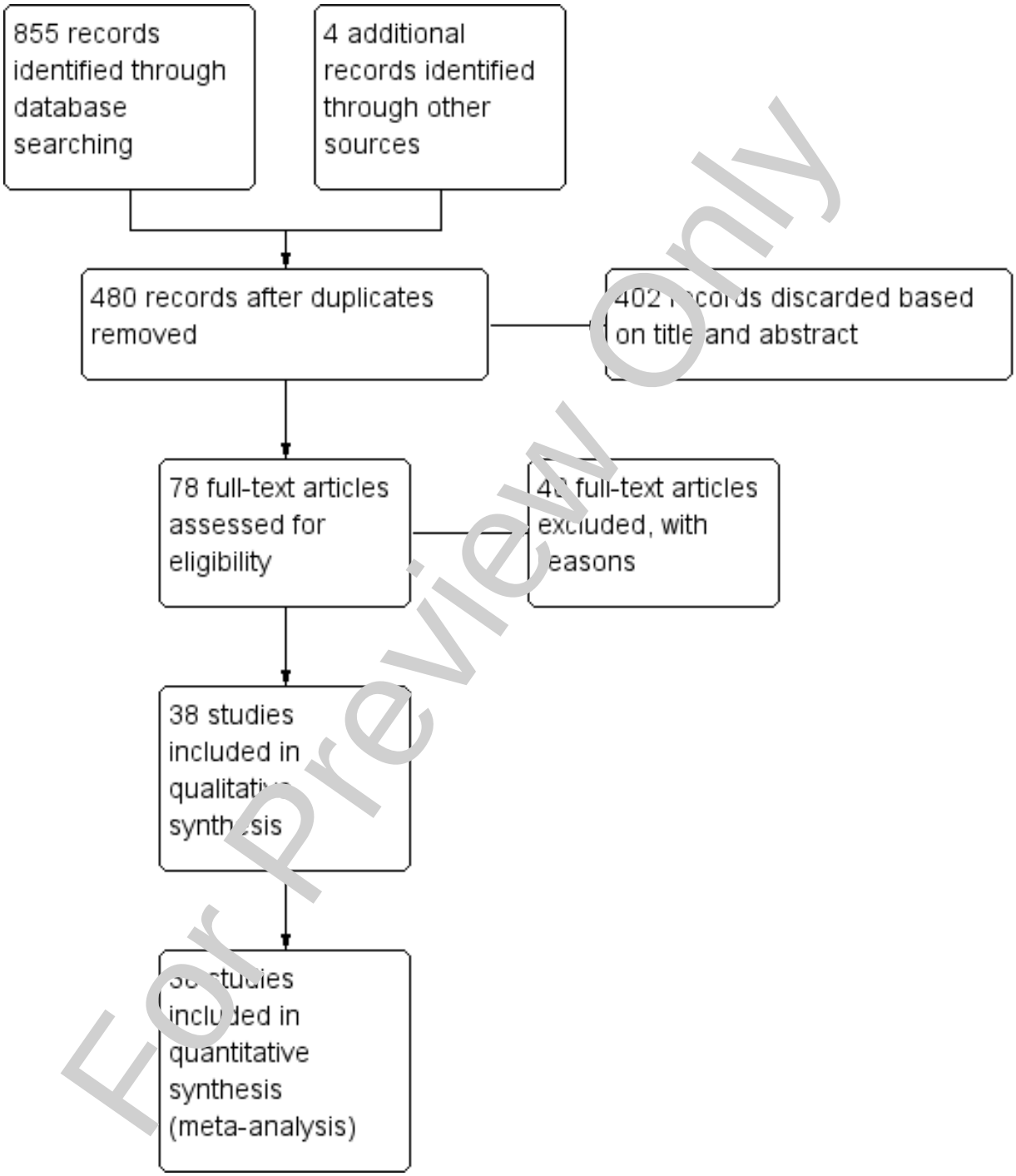
We also searched for grey literature, such as conference abstracts, proceedings and theses on the following databases: www.greylit.org; www.opengrey.eu (last search was performed on 2 March 2018, see Appendix 8).

Data collection and analysis

Selection of studies

Following the electronic search, two review authors (Jayakumar Nadathur Doraiswamy (JND), Malaiappan Sankari (MS)) independently screened the titles and abstracts (if available) to exclude all articles clearly not meeting the inclusion criteria. The search was designed to be sensitive and include controlled clinical trials, these were filtered out early in the selection process if they were not randomized. Of all the remaining articles, full texts were obtained and assessed independently by two review authors (JND, MS) and only articles fully meeting the inclusion criteria were considered. In case of disagreement between the two review authors, a third review author (Massimo Del Fabbro (MDF)) was consulted. Detailed reasons were stated for all excluded studies. This process is summarised in Figure 1.

Figure 1. Study flow diagram.



Data extraction and management

Three review authors (SP, Lorena Karanxha (LK), CB) independently extracted and recorded data on ad hoc forms. Any disagreement was solved through discussion, or a third review author was consulted (MDF). In case of missing or unclear information, we contacted the authors of the included reports by email to provide clarification or missing information. In case of missing or incomplete data and absence of further clarification by study authors we excluded the report from the analysis.

We recorded the following data for each included report:

- demographic characteristics of the population;
- defect characteristics (PD, CAL, RBF);
- type of platelet concentrate used (PRP, PRF, PRGF);
- outcome characteristics (outcome variables assessed such as CAL and PD, follow-up duration);
- when possible, we also recorded the expertise of the clinician (years of experience with using platelet concentrates); and
- source of funding.

Assessment of risk of bias in included studies

Three review authors (LK, SP, CB) independently assessed the risk of bias in the included studies. In case of disagreement a fourth review author (MDF) was consulted. Since some of the authors of one of the randomised controlled trials included (Trombelli 2016) are also authors of this review (SP, MDF, Silvia Tonelli (ST)), the risk of bias assessment for that study was carried out by other review authors not involved in the study (LK, CB).

The assessment was conducted following the instructions and the approach described in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). For each study, the following domains were considered: selection bias (random sequence generation and allocation concealment), performance bias (blinding of participants and personnel), detection bias (blinding of outcome assessment), attrition bias (incomplete outcome data addressed), and reporting bias (selective reporting).

For each domain the risk was judged either low, unclear or high. If one study had low risk for all domains, the study was judged at low risk of bias. If it had an unclear risk for at least one domain, the study was judged at unclear risk of bias. If it had a high risk for at least one domain, the study was judged at high risk of bias. It was considered that blinding of patient and clinician might be difficult/impossible as for many studies involving surgical procedures where interventions are quite different from each other.

We categorised the overall risk of bias of individual studies. Studies were categorised as being at low, high, or unclear risk of bias according to the following criteria:

- low risk of bias (plausible bias unlikely to seriously alter the results) if all domains were at low risk of bias;
- high risk of bias (plausible bias that seriously weakens confidence in the results) if one or more domains were at high risk of bias; or
- unclear risk of bias (plausible bias that raises some doubt about the results) if one or more domains were at unclear risk of bias.

These assessments are reported in the Characteristics of included studies table and also graphically.

Measures of treatment effect

For continuous outcomes (e.g. PD, CAL, RBF), mean differences (change score) along with 95% confidence intervals (CIs) were used to summarise data for each treatment group. We expressed the data in mm for PD and CAL and in percentage for RBF, as they were reported in the studies.

Unit of analysis issues

The statistical unit of analysis in parallel studies was the patient, unless the study provided data only for defects. We considered one infrabony defect per patient in studies with parallel design. In the case of split-mouth studies, the unit of analysis was the defect; a single defect per patient per group was considered.

Dealing with missing data

In case of missing data, we contacted the corresponding author of the article through e-mail to obtain complete data. In case of no response, the same e-mail was sent to co-authors for a maximum of three times. If no answer was obtained, the study was excluded from the analysis. When feasible, missing standard deviations were estimated using the methods described in Section 7.7.3 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011).

Assessment of heterogeneity

Heterogeneity among studies was assessed with Cochran's test for heterogeneity, with a significance threshold of $P < 0.1$. The quantification of the heterogeneity was calculated with I^2 statistic. For the interpretation of I^2 the ranges suggested in Section 9.5.2 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011) were considered.

Assessment of reporting biases

We assessed publication bias by testing for funnel plot asymmetry, as described in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). If asymmetry was evident, we investigated this and described possible causes.

Data synthesis

The meta-analysis was performed only with studies with similar comparisons reporting the same outcome measures. We combined mean differences for continuous data, using random-effects models if at least four studies were included in the meta-analysis, while if there were less than four studies a fixed-effect model was chosen. The software RevMan 5 (Review Manager 2014) was used for meta-analysis computations. Data from split-mouth and parallel-group studies were combined (Elbourne 2002). The appropriate standard errors were estimated where they were not present in the trial reports (Follmann 1992). For the split-mouth studies the standard error was calculated assuming an intraclass correlation coefficient of 0. The generic inverse variance procedure in RevMan 5 was used to combine these two subgroups in the analyses.

Subgroup analysis and investigation of heterogeneity

In addition to the different surgical protocols for different types of infrabony defects, duration of the follow-up was investigated a factor possibly affecting the outcome. The subgroups included data up to 6 months (3 to 6 months) and longer than 6 months (9 to 12 months).

Sensitivity analysis

Sensitivity analysis was performed in order to evaluate the effect of risk of bias and source of funding on the overall effects (e.g. omitting studies at unclear or high risk of bias or those sponsored by the manufacturer of the product under investigation). The effect of excluding specific studies that eventually appeared to be outliers was also investigated.

Summary of findings

We produced a 'Summary of findings' table for each comparison in which there was more than one study. We included the change in PD, CAL and RBF of the all follow-up periods of each comparison group. We used GRADE methods, and GRADEpro software (GRADEpro GDT 2015) for developing 'Summary of findings' tables. We assessed the quality of the body of evidence for each comparison and outcome by considering the overall risk of bias of the included studies, the directness of the evidence, the inconsistency of the results, the precision of the estimates, and the risk of publication bias. We categorised the quality of each body of evidence as high, moderate, low, or very low.

RESULTS

Description of studies

Results of the search

The electronic search retrieved 855 records, four trials were identified by handsearching and none by searching the grey literature. After discarding the duplicates, two review authors (Jayakumar Nadathur Doraiswamy (JND), Malaiappan Sankari (MS)) screened 480 titles and abstracts and rejected 402. The full text was obtained for 78 potentially eligible articles and of these, 40 were excluded with reasons (see Characteristics of excluded studies table). Finally, after agreement among the review authors 38 studies were included in this review (Figure 1).

Included studies

Design

Of the 38 included studies, 22 had a split-mouth design, reporting for a total of 371 participants and 701 teeth (Agarwal 2014; Agarwal 2015; Agarwal 2016; Arabaci 2017; Aydemir 2016; Camargo 2009; Christgau 2006; Elgendy 2015; Gupta 2014; Hanna 2004; Hassan 2012; Kaushick 2011; Khosropanah 2015; Naqvi 2017; Ozdemir 2012; Panda 2016; Patel 2017; Ravi 2017; Rosamma Joseph 2012; Sezgin 2017; Shukla 2016; Thorat 2017); 16 studies had a parallel design with a total of 645 patients and 721 teeth (Chandradas 2016; Demir 2007; Döri 2007a; Döri 2007b; Döri 2008a; Döri 2008b; Döri 2009; Garg 2017; Kanoriya 2016; Martande 2016; Okuda 2005; Piemontese 2008; Pradeep 2015; Pradeep 2016; Sharma 2011; Thorat 2011). Of the 38 included studies only one was a multicentric study (Elgendy 2015). Finally, two studies declared that they were supported in part by companies whose products were used in the trials (Döri 2008a; Döri 2008b).

Sample size calculation was reported only by 15 studies (Döri 2007a; Döri 2007b; Döri 2008a; Döri 2008b; Döri 2009; Kanoriya 2016; Panda 2016; Patel 2017; Pradeep 2015; Pradeep 2016; Ravi 2017; Rosamma Joseph 2012; Sezgin 2017; Sharma 2011; Thorat 2011), meaning that in almost 60% of cases there was no rationale regarding the choice of the sample size.

Participants

The age range of the participants of included studies was between 17 and 74 years. However, four studies did not report the age of the participants (Agarwal 2016; Gupta 2014; Naqvi 2017; Shukla 2016) and 10 studies (Agarwal 2015; Aydemir 2016; Chandradas 2016; Demir 2007; Elgendy 2015; Hassan 2012; Khosropanah

2015; Okuda 2005; Ozdemir 2012; Sezgin 2017) reported only mean ages, ranging from 36.03 and 55.5 years.

35 studies included both men and women, but with different proportions, and three studies did not report this information (Gupta 2014; Elgendy 2015; Kaushick 2011). Finally, most of the studies did not include smokers (Agarwal 2014; Agarwal 2015; Agarwal 2016; Arabaci 2017; Aydemir 2016; Chandradas 2016; Döri 2007a; Döri 2007b; Döri 2008a; Döri 2008b; Döri 2009; Garg 2017; Gupta 2014; Hassan 2012; Kanoriya 2016; Kaushick 2011; Khosropanah 2015; Martande 2016; Naqvi 2017; Okuda 2005; Ozdemir 2012; Panda 2016; Patel 2017; Piemontese 2008; Pradeep 2015; Pradeep 2016; Ravi 2017; Rosamma Joseph 2012; Sezgin 2017; Sharma 2011; Shukla 2016; Thorat 2011; Thorat 2017).

Interventions

The general comparison was between a group that received autologous platelet concentrates (APC) as an adjunct to surgical treatment (experimental group), and a group that received surgical treatment alone (control group). Four different types of comparisons were assessed, based on the treatment type:

1. APC + open flap debridement (OFD) versus OFD alone (12 trials): Agarwal 2016; Arabaci 2017; Chandradas 2016; Kanoriya 2016; Martande 2016; Patel 2017; Pradeep 2015; Pradeep 2016; Rosamma Joseph 2012; Sharma 2011; Thorat 2011; Thorat 2017
2. APC + OFD + bone graft (BG) versus OFD + BG (17 trials): Agarwal 2014; Agarwal 2015; Demir 2015; Döri 2009; Elgendy 2015; Garg 2017; Gupta 2014; Handa 2014; Hassan 2012; Kaushick 2011; Khosropanah 2015; Naqvi 2017; Okuda 2005; Ozdemir 2012; Piemontese 2008; Sezgin 2017; Shukla 2016
3. APC + guided tissue regeneration (GTR) versus GTR (7 trials): Camargo 2009; Christgau 2006; Döri 2007a; Döri 2007b; Döri 2008a; Panda 2016; Ravi 2017
4. APC + enamel matrix derivative (EMD) versus EMD (2 trials): Aydemir 2016; Döri 2008b.

Outcomes

Primary outcomes

- Change in probing depth (PD), reported by all 38 included studies.
- Change in clinical attachment level (CAL), defined relative attachment level (RAL) in some studies, reported by all 38 included studies.
- Change in radiographic bone defect filling (RBF), reported by 31 studies.

Secondary outcomes

All articles in all groups reported a survival rate of 100% for the treated teeth. No complete pocket closure was reported. No quantitative analysis regarding patients' quality of life was possible.

Excluded studies

We excluded 21 studies from the review, for the following reasons (see Characteristics of excluded studies table):

- no randomisation (Aleksić 2008; Jovicić 2013; Saini 2011)
- no control group (Camargo 2002; Camargo 2005; Lekovic 2012)
- gingival recession, not infrabony defects (Aroca 2009; Dogan 2015; Huang 2005; Jankovic 2010; Padma 2013; Shepherd 2009; Shivakumar 2016; Thamaraiselvan 2015)
- same patients reported in a previous study (Cetinkaya 2014; Döri 2013; Moder 2012; Yajamanya 2017)
- non-independence of analysing unit (Gupta 2014b; Pradeep 2012a)
- incomplete data (Cieplik 2018; Harnack 2009; Keceli 2008; Keles 2006; Menezes 2012; Shah 2015; Yassibag-Berkman 2007; Yen 2007)
- no APCs (fibrin glue) (Cortellini 1995; Trombelli 1995; Trombelli 1996)
- APC not the only difference between groups (Cheung 2004; Eren 2014; Jankovic 2012)
- studies with mixed (parallel/split-mouth) design (Agarwal 2017; Bajaj 2017; Chatterjee 2017; Ouyang 2006; Pradeep 2017; Qiao 2016).

Risk of bias in included studies

The risk of bias in included studies is summarized in Figure 2 and Figure 3. Two studies were at unclear overall risk of bias (Ravi 2017; Rosamma Joseph 2012). The remaining 36 studies had a high overall risk of bias.

Figure 2. Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.

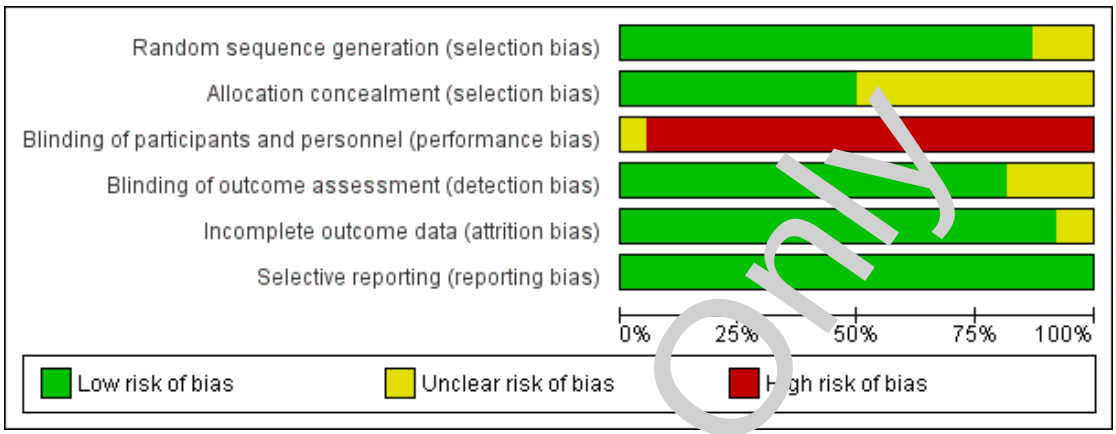
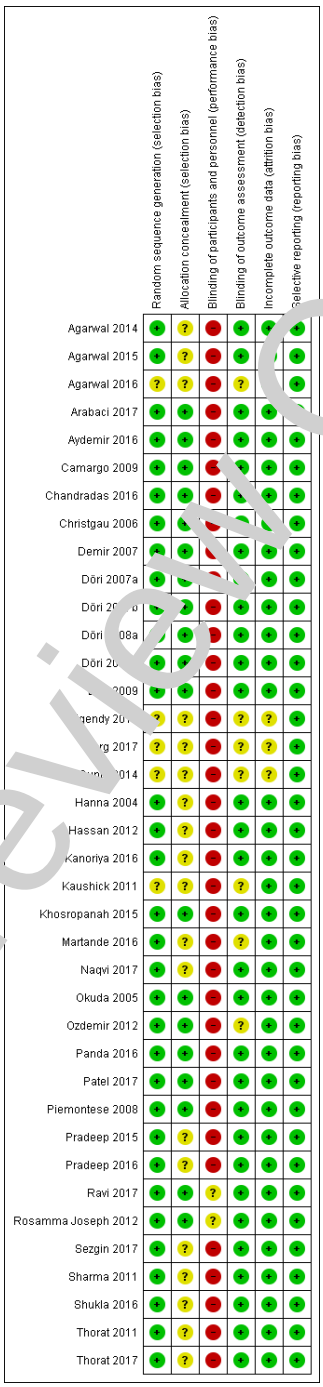


Figure 3. Risk of bias summary: review authors' judgements about each risk of bias item for each included study.



Allocation

Random sequence generation

The randomisation was performed correctly in most of the studies. The methods used were the tossing of a coin (Agarwal 2014; Agarwal 2015; Camargo 2009; Demir 2007; Gupta 2014; Hanna 2004; Khosropanah 2015; Okuda 2005; Ozdemir 2012; Panda 2016; Patel 2017; Piemontese 2008; Ravi 2017; Rosamma Joseph 2012; Sezgin 2017; Sharma 2011; Thorat 2011), the block approach (Döri 2007a; Döri 2007b; Döri 2008a; Döri 2008b; Döri 2009), the use of a freeware link (Chandradas 2016), computerized generated scheme (Aydemir 2016; Kanoriya 2016; Martande 2016; Pradeep 2016; Shukla 2016; Thorat 2011), biased coin randomisation (Hassan 2012), lottery method (Naqvi 2017), and a table of random numbers (Christgau 2006; Pradeep 2015). The randomisation method was not described in five articles, which were considered to be an unclear risk of bias (Agarwal 2016; Elgendy 2015; Garg 2017; Gupta 2014; Kaushick 2011).

Allocation concealment

The concealment of the allocation was correctly done in 19 studies (Arabaci 2017; Aydemir 2016; Camargo 2009; Chandradas 2016; Christgau 2006; Demir 2007; Döri 2007a; Döri 2007b; Döri 2008a; Döri 2008b; Döri 2009; Khosropanah 2015; Okuda 2005; Ozdemir 2012; Panda 2016; Patel 2017; Piemontese 2008; Ravi 2017; Rosamma Joseph 2012). In the remaining 19 studies, insufficient information was provided regarding the exact method used for allocation concealment (Agarwal 2014; Agarwal 2015; Agarwal 2016; Elgendy 2015; Garg 2017; Gupta 2014; Hanna 2004; Hassan 2012; Kanoriya 2016; Kaushick 2011; Martande 2016; Naqvi 2017; Pradeep 2015; Pradeep 2016; Sezgin 2017; Sharma 2011; Shukla 2016; Thorat 2011; Thorat 2017).

Blinding

Blinding of participants and personnel (performance bias)

Being the intervention surgical in nature, blinding of participants and treating clinicians is almost unfeasible either in a parallel or split-mouth design: 36 out of 38 studies had a high risk of performance bias. For two studies an unclear risk of performance bias was assigned given that it was stated in the paper that blinding of the operator was performed but without specifying how (Ravi 2017; Rosamma Joseph 2012). The blinding of the personnel was also evaluated, which was reported in most of the studies except for eight studies (Agarwal 2016; Christgau 2006; Elgendy 2015; Garg 2017; Gupta 2014; Kaushick 2011; Okuda 2005; Ozdemir 2012). However, again for the fact that the intervention has a surgical nature, it is unlikely that blinding or not of the personnel could influence the outcome. Therefore such parameter did not influence the assignment of the risk of performance bias.

Blinding of outcome assessment (detection bias)

The blinding of the outcome assessor was done in most of the studies. However, it was not reported in seven studies, which were considered to be at unclear risk of detection bias (Agarwal 2016; Elgendy 2015; Garg 2017; Gupta 2014; Kaushick 2011; Martande 2016; Ozdemir 2012).

Incomplete outcome data

The completeness of outcome data was adequate in all but three studies in which the number of subjects that finished the study was not clear (Elgendy 2015; Garg 2017; Gupta 2014).

Selective reporting

All studies properly reported data for all patients.

Effects of interventions

See: **Summary of findings for the main comparison** APC + OFD compared to OFD (9-12 months follow-up) for treating periodontal infrabony defects; **Summary of findings 2** APC + OFD + BG compared to OFD + BG (all follow-ups) for treating periodontal infrabony defects; **Summary of findings 3** APC + GTR compared to GTR (all follow-ups) for treating periodontal infrabony defects; **Summary of findings 4** APC + EMD compared to EMD (all follow-ups) for treating periodontal infrabony defects

For the meta-analyses of all follow-ups, where the study presented multiple follow-ups, we used the longest one.

1. Autologous platelet concentrates (APC) + open flap debridement (OFD) versus OFD

Summary of findings for the main comparison.

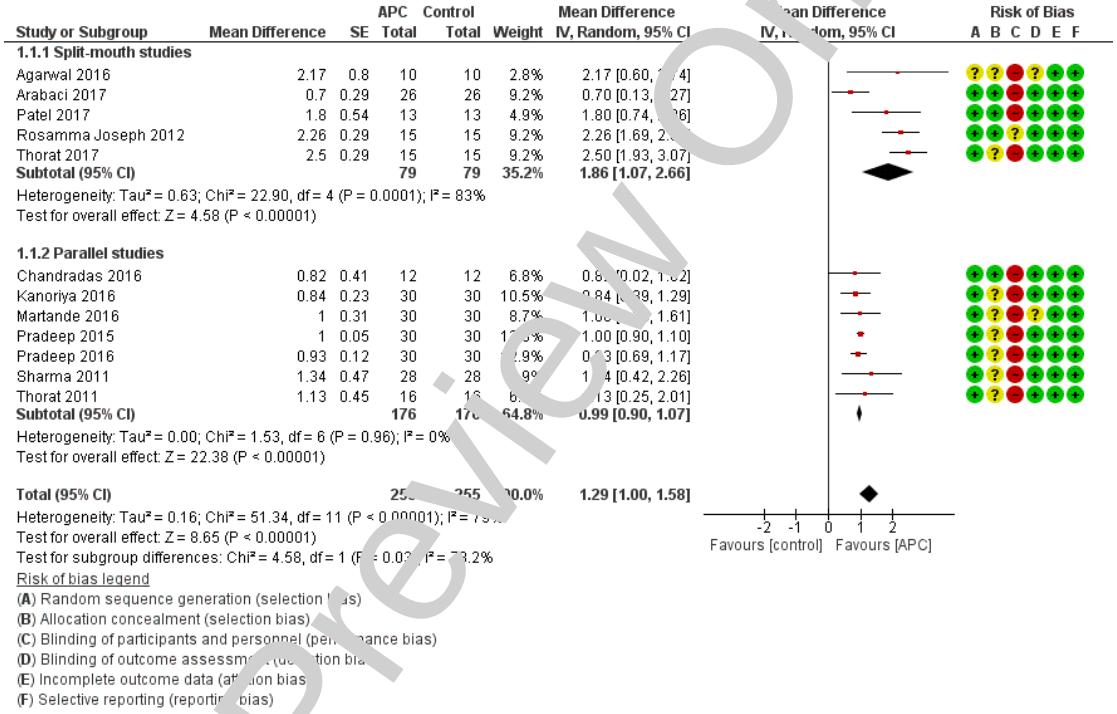
In this comparison we did not divide the data according to the follow-up duration, because all studies had a follow-up duration between 9 and 12 months.

Change in probing depth (PD) (mm)

Follow-up between 9 and 12 months

There is evidence of an advantage in using APC from both split-mouth studies (mean difference (MD) 1.86, 95% confidence interval (CI) 1.07 to 2.66; P < 0.001; 5 studies; 158 participants) and parallel studies (MD 0.99, 95% CI 0.90 to 1.07; P < 0.001; 7 studies, 352 participants). Overall, there is evidence of an advantage in using APC (MD 1.29, 95% CI 1.00 to 1.58; P < 0.001) (Figure 4; Analysis 1.1).

Figure 4. Forest plot of comparison: 1 APC + OFD versus OFD (9-12 mo[illegible]s follow[illegible]up); outcome: 1.1 Probing depth (mm).



Change in clinical attachment level (CAL) (mm)

Follow-up between 9 and 12 months

There is evidence of an advantage in using APC from split-mouth studies (MD 2.36, 95% CI 1.19 to 3.54; P < 0.001; 5 studies; 158 participants) and parallel studies (MD 0.99, 95% CI 0.84 to 1.14; P < 0.001; 7 studies; 352 participants). Overall, there is evidence of an advantage in using APC (MD 1.47, 95% CI 1.11 to 1.82; P < 0.001) (Analysis 1.2).

Change in radiographic bone defect filling (RBF) (%)

Follow-up between 9 and 12 months

There is evidence of an advantage in using APC from split-mouth studies (MD 27.32%, 95% CI 20.92% to 33.72%; P < 0.001; 2 studies; 49 participants) and parallel studies (MD 35.77%, 95% CI 31.20% to 40.35%; P < 0.001; 7 studies; 352 participants). Overall, there is evidence of an advantage in using APC (MD 34.26%, 95% CI 30.07% to 38.46%; P < 0.001) (Analysis 1.3).

2. APC + OFD + bone graft (BG) versus OFD + BG

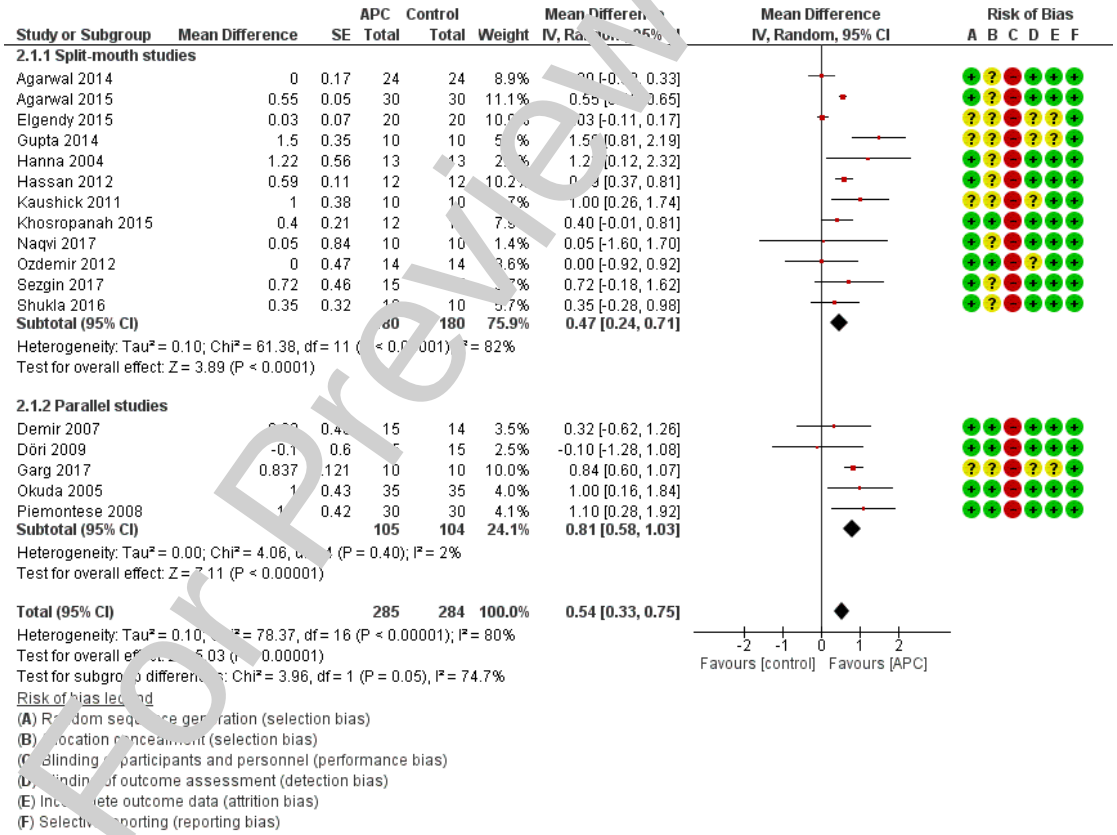
Summary of findings 2.

Change in PD (mm)

All follow-ups

There is evidence of an advantage in using APC from split-mouth studies (MD 0.47, 95% CI 0.24 to 0.71; P < 0.001; 12 studies; 360 participants) and from parallel studies (MD 0.81, 95% CI 0.58 to 1.03; P < 0.001; 5 studies; 209 participants). Overall, there is evidence of an advantage in using APC (MD 0.54, 95% CI 0.33 to 0.75; P < 0.001) (Figure 5; Analysis 2.1).

Figure 5. Forest plot of comparison: 2 APC + OFD + BG versus OFD + BG (all follow-ups); outcome: 2.1 Probing depth (mm).



Follow-up between 3 and 6 months

There is evidence of an advantage in using APC from split-mouth studies (MD 0.58, 95% CI 0.25 to 0.92; P = 0.0007; 10 studies; 252 participants). However, there is only one study to consider of parallel design (MD 0.84, 95% CI 0.60 to 1.07; P < 0.001; 20 participants). Overall, there is evidence of an advantage in using APC with a shorter follow-up duration (MD 0.62, 95% CI 0.30 to 0.94; P = 0.0002) (Analysis 3.1).

Follow-up between 9 and 12 months

There is evidence of an advantage in using APC from split-mouth studies (MD 0.49, 95% CI 0.26 to 0.72; P < 0.001; 6 studies; 192 participants), and from parallel studies (MD 0.58, 95% CI 0.09 to 1.06; P = 0.02; 4 studies; 189 participants). Overall, there is evidence of an advantage in using APC (MD 0.50, 95% CI 0.31 to 0.69; P < 0.0001) (Analysis 4.1).

Change in CAL (mm)

All follow-ups

There is evidence of an advantage in using APC from split-mouth studies (MD 0.67, 95% CI 0.35 to 0.99; P < 0.001; 12 studies; 360 participants) and from parallel design studies (MD 0.89, 95% CI 0.49 to 1.29; P < 0.001; 5 studies; 209 participants). Overall, there is evidence of an advantage in using APC (MD 0.72, 95% CI 0.43 to 1.00; P < 0.001) (Analysis 2.2).

Follow-up between 3 and 6 months

There is evidence of an advantage in using APC from split-mouth studies (MD 0.40, 95% CI 0.02 to 0.77; P = 0.04; 10 studies; 252 participants). However, there is only one study to consider of parallel design (MD 1.00, 95% CI 0.93 to 1.07; P < 0.001; 20 participants). Overall, there is evidence of an advantage in using APC (MD 0.47, 95% CI 0.11 to 0.83; P = 0.01) (Analysis 3.2).

Follow-up between 9 and 12 months (only split-mouth studies)

There is evidence of an advantage in using APC (MD 0.84, 95% CI 0.62 to 1.06; P < 0.001; 6 studies; 192 participants) (Analysis 4.2).

Change in RBF (%)

All follow-ups

There is evidence of an advantage in using APC from both split-mouth studies (MD 7.73%, 95% CI 4.50% to 10.97%; P < 0.001; 8 studies; 270 participants) and parallel studies (MD 9.66%, 95% CI 5.39% to 13.94%; P < 0.001; 3 studies; 150 participants). Overall, there is evidence of an advantage in using APC (MD 8.10%, 95% CI 5.26% to 10.94%; P < 0.001) (Analysis 2.3).

Follow-up between 3 and 6 months

There is evidence of an advantage in using APC from split-mouth studies (MD 3.59%, 95% CI 0.13% to 7.05%; P = 0.04; 5 studies; 142 participants) and from one parallel study (MD 10.00%, 95% CI 4.90% to 15.10%; P = 0.0001; 20 participants). Overall, there is evidence of an advantage in using APC (MD 4.76%, 95% CI 1.27% to 8.25%; P = 0.008) (Analysis 3.3).

Follow-up between 9 and 12 months

There is evidence of an advantage in using APC from split-mouth studies (MD 10.16%, 95% CI 6.18% to 14.14%; P < 0.001; 4 studies; 152 participants), and from parallel studies (MD 8.87%, 95% CI 1.03% to 16.71%; P = 0.03; 2 studies; 130 participants). Overall, there is evidence of an advantage in using APC (MD 9.99%, 95% CI 6.44% to 13.55%; P < 0.001) (Analysis 4.3).

3. APC + guided tissue regeneration (GTR) versus GTR

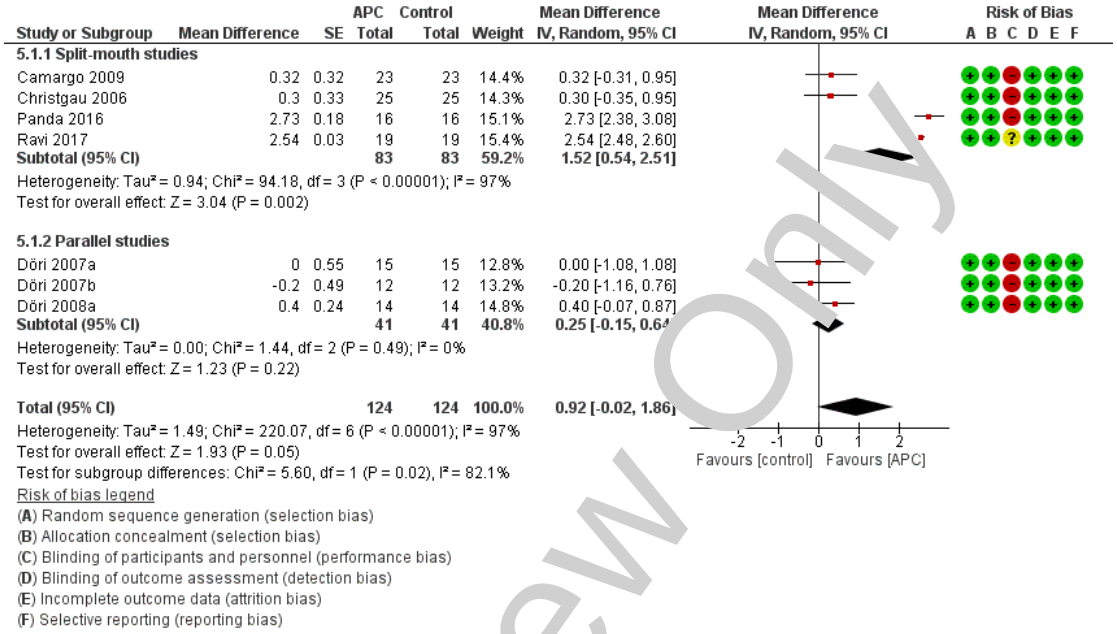
Summary of findings 3.

Change in PD (mm)

All follow-ups

There is evidence of an advantage in using APC from split-mouth studies (MD 1.52, 95% CI 0.54 to 2.51; P = 0.002; 4 studies; 166 participants) but not from parallel studies (MD 0.25, 95% CI -0.15 to 0.64; P = 0.22; 3 studies, 82 participants). Overall, there is evidence of an advantage in using APC (MD 0.92, 95% CI -0.02 to 1.86; P = 0.05). However, given the wide confidence intervals, there is a possibility of an advantage for the control group (Figure 6; Analysis 5.1).

Figure 6. Forest plot of comparison: 5 APC + GTR versus GTR (all follow-ups), outcome: 5.1 Probing depth (mm).



Follow-up between 3 and 6 months (only split-mou studies)

There is insufficient evidence of an advantage in using AP MD 1.07, 95% CI -0.71 to 2.86; P = 0.24; 3 studie 134 participants) (Analysis 6.1).

Follow-up between 9 and 12 mo

There is insufficient evidence of an adv tage in using APC from both split-mouth studies (MD 5% CI -0.85 to 3.91; P = 0.21; 2 studies; 82 participants) and arallel studies (MD 0.25, 95% CI -0.15 to 0.64; P = 0.22; 3 studies; 82 participants). Overall, there is insufficient evidence of an advantage in using APC (MD 0.68, 95% CI -0.66 to 02; P = 0.32) (Analysis 7.1).

Change in CAL (m

All follow-u

There is evidence of an advantage in using APC from split-mouth studies (MD 0.67, 95% CI 0.20 to 1.14; P = 0.005; 4 studies; 166 participants) but not from parallel studies (MD 0.09, 95% CI -0.32 to 0.50; P = 0.66; 3 studies; 82 participants). Overall, there is insufficient evidence of an advantage in using APC (MD 0.42, 95% CI -0.02 to 0.86; P = 0.06) (Analysis 5.2).

Follow-up between 3 and 6 months (only split-mouth studies)

There is evidence of an advantage in using APC (MD 0.54, 95% CI 0.18 to 0.89; P = 0.003; 3 studies; 134 participants) (Analysis 6.2).

Follow-up between 9 and 12 months

There is insufficient evidence of an advantage in using APC from both split-mouth studies (MD 0.51, 95% CI -0.72 to 1.73; P = 0.42; 2 studies; 82 participants) and parallel studies (MD 0.09, 95% CI -0.32 to 0.50; P = 0.66; 3 studies; 82 participants). Overall, there is no evidence of an advantage in using APC (MD 0.27, 95% CI -0.39 to 0.93; P = 0.42) (Analysis 7.2).

4. APC + enamel matrix derivative (EMD) versus EMD

Summary of findings 4.

Change in PD (mm)

All follow-ups

Only one study had a split-mouth design and showed insufficient evidence of an advantage in using APC (MD 0.13, 95% CI -0.05 to 0.31; P = 0.15; 49 participants). Equally only one study had a parallel design which showed insufficient evidence of an advantage in using APC (MD -0.10, 95% CI -1.32 to 1.12; P = 0.87; 26 participants). Overall, there is insufficient evidence of an advantage in using APC (MD 1.13, 95% CI -0.05 to 0.30; P = 0.16) (Analysis 8.1).

Change in CAL (mm)

All follow-ups

Only one study had a split-mouth design and showed insufficient evidence of an advantage in using APC (MD 0.12, 95% CI -0.12 to 0.36; P = 0.32; 49 participants). The only one study with a parallel design also showed insufficient evidence of an advantage in using APC (MD -0.20, 95% CI -1.06 to 0.66; P = 0.65; 26 participants). Overall, there is insufficient evidence of an advantage in using APC (MD 0.10, 95% CI -0.13 to 0.32; P = 0.40) (Analysis 8.2).

Change in RBF (%)

All follow-ups

Only one split-mouth study provided data and showed insufficient evidence of an advantage in using APC (MD -0.60%, 95% CI -6.21% to 5.01%; P = 0.83; 49 participants) (Analysis 8.3).

Secondary outcomes

All the studies in all groups reported a survival rate of 100% for the treated teeth. No complete pocket closure was reported. No quantitative analysis regarding patients' quality of life was possible.

ADDITIONAL SUMMARY OF FINDINGS [*Explanation*]

APC + OFD + BG compared to OFD + BG (all follow-ups) for treating periodontal infrabony defects						
Patient or population: patients affected by infrabony defects requiring surgical treatment **Settings:** tertiary care **Intervention:** APC + OFD + BG **Comparison:** OFD + BG						
Outcomes	**Illustrative comparative risks* (95% CI)**		**Relative effect (95% CI)**	**Number of participants/defects (studies)**	**Quality of the evidence (GRADE)**	**Comments**
	Assumed risk	**Corresponding risk**				
	OFD + BG	**APC + OFD + BG**				
Change in probing depth (PD) (mm) (All follow-ups)	Mean PD change (gain) across control groups ranged from 1.90 to 5.30 (3.54) mm Mean PD baseline value was 7.32 mm (95% CI 5.94 to 8.65)	The mean PD change (gain) in the intervention groups was **0.54 mm higher** (0.33 to 0.75 higher)	Mean difference 0.54 (0.33 to 0.75) mm	569 (17 studies)	⊕◯◯◯ **very low**[1,2]	There is evidence of an advantage in using APC
Change in clinical attachment level (CAL) (mm) (All follow-ups)	Mean CAL change (gain) across control groups ranged from 1.30 to 4.70 (3.20) mm Mean CAL baseline value was 7.34 mm (95% CI 5.21 to 9.82)	The mean CAL change (gain) in the intervention groups was **0.72 mm higher** (0.43 to 1.00 higher)	Mean difference 0.72 (0.43 to 1.00) mm	569 (17 studies)	⊕◯◯◯ **very low**[1,2]	There is evidence of an advantage in using APC
Change in radiographic bone defect filling (RBF) (%) (All follow-ups)	Mean RBF change (gain) across control groups ranged from 9.20% to 57.20% (40.54%)	The mean RBF change (gain) in the intervention groups was **8.10% higher** (5.26 to 10.94 higher)	Mean difference 8.10% (5.26 to 10.94)	420 (11 studies)	⊕◯◯◯ **very low**[1,2]	There is evidence of an advantage in using APC

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% CI) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).
APC: autologous platelet concentrates; **BG:** bone graft; **CAL:** clinical attachment level; **CI:** confidence interval; **OFD:** open flap debridement; **PD:** probing depth; **RBF:** radiographic bone defect filling.

GRADE Working Group grades of evidence
High quality: further research is very unlikely to change our confidence in the estimate of effect.
Moderate quality: further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
Low quality: further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: we are very uncertain about the estimate.

[1]Downgraded by 2 levels for high risk of performance bias.
[2]Downgraded by 2 levels for high heterogeneity.

APC + GTR compared to GTR (all follow-ups) for treating periodontal infrabony defects

Patient or population: patients affected by infrabony defects requiring surgical treatment
Settings: tertiary care
Intervention: APC + GTR
Comparison: GTR

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	Number of participants/defects (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	GTR	APC + GTR				
Change in probing depth (PD) (mm) (All follow-ups)	Mean PD change (gain) across control groups ranged from 3.19 to 6.00 mm (4.40 mm) Mean PD baseline value was 8.67 mm (95% CI 6.29 to 10.31)	The mean PD change (gain) in the intervention groups was **0.92 mm higher** (-0.02 lower to 1.86 higher)	Mean difference 0.92 mm (-0.02 to 1.86)	248 (7 studies)	⊕◯◯◯ **very low**[1,2,3]	There is insufficient evidence of an advantage in using APC
Change in clinical attachment level (CAL) (mm) (All follow-ups)	Mean CAL change (gain) across control groups ranged from 3.38 to 5.20 mm (4.38 mm) Mean CAL baseline value was 9.40 mm (95% CI 5.97 to 11.40)	The mean CAL change (gain) in the intervention groups was **0.42 mm higher** (-0.02 lower to 0.86 higher)	Mean difference 0.42 mm (-0.02 to 0.86)	248 (7 studies)	⊕◯◯◯ **very low**[1,2,3]	There is insufficient evidence of an advantage in using APC

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% CI) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).
APC: autologous platelet concentrates; **CAL:** clinical attachment level; **CI:** confidence interval; **GTR:** guided tissue regeneration; **OFD:** open flap debridement; **PD:** probing depth; **RBF:** radiographic bone defect filling.

GRADE Working Group grades of evidence
High quality: further research is very unlikely to change our confidence in the estimate of effect.
Moderate quality: further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
Low quality: further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: we are very uncertain about the estimate

[1]Downgraded by 2 levels for high risk of performance bias.
[2]Downgraded by 2 levels for high heterogeneity.
[3]Downgraded by 2 levels for imprecision (wide confidence interval and small sample size).

APC + EMD compared to EMD (all follow-ups) for treating periodontal infrabony defects

Patient or population: patients affected by infrabony defects requiring surgical treatment
Settings: tertiary care
Intervention: APC + EMD
Comparison: EMD

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	Number of participants/defects (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	EMD	APC + EMD				
Change in probing depth (PD) (mm) (All follow-ups)	Mean PD change (gain) across control groups ranged from 3.87 to 5.90 mm (4.89 mm)	The mean PD change (gain) in the intervention groups was **0.13 mm higher** (-0.05 lower to 0.30 higher)	Mean difference 0.13 mm (-0.05 to 0.30)	75 (2 studies)	⊕◯◯◯ **very low**[1,2]	There is insufficient evidence of an advantage in using APC
Change in clinical attachment level (CAL) (mm) (All follow-ups)	Mean CAL change (gain) across control groups ranged from 3.30 to 5.00 mm (4.15 mm)	The mean CAL change (gain) in the intervention groups was **0.10 mm higher** (-0.13 lower to 0.32 higher)	Mean difference 0.10 mm (-0.13 to 0.32)	75 (2 studies)	⊕◯◯◯ **very low**[1,2]	There is insufficient evidence of an advantage in using APC
Change in radiographic bone defect filling (RBF) (%) (All follow-ups)	Only 1 study reported RBF outcome with a mean change in control groups of 18.30%	The mean RBF change (gain) in the intervention group was **0.60% lower** (-6.21 lower to 5.01 higher)	Mean difference -0.60 (-6.21 to 5.01)	49 (1 study)	⊕◯◯◯ **very low**[1,2]	There is insufficient evidence of an advantage in using APC

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% CI) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).
APC: autologous platelet concentrates; **CAL:** clinical attachment level; **CI:** confidence interval; **EMD:** enamel matrix derivative; **OFD:** open flap debridement; **PD:** probing depth; **RBF:** radiographic bone defect filling.

GRADE Working Group grades of evidence
High quality: further research is very unlikely to change our confidence in the estimate of effect.
Moderate quality: further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
Low quality: further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: we are very uncertain about the estimate.

[1]Downgraded by 2 levels for high risk of performance bias.
[2]Downgraded by 2 levels for imprecision (wide confidence interval and small sample size).

DISCUSSION

Summary of main results

We included 38 studies in this review. These studies assessed the effects of autologous platelet concentrates (APC) used as an adjunct to periodontal surgical therapies for the treatment of infrabony defects. We assessed the quality of the body of evidence using GRADE criteria, and our assessment is presented in Summary of findings for the main comparison (for APC + open flap debridement (OFD) versus OFD alone); Summary of findings 2 (for APC + OFD + bone graft (BG) versus OFD + BG); Summary of findings 3 (for APC + guided tissue regeneration (GTR) versus GTR); and Summary of findings 4 (for APC + enamel matrix derivative (EMD) versus EMD).

All data were analysed separately by subgroups and for specific parameters. In an overall assessment of outcomes, there is evidence that the presence of APC brings advantages in the change of probing depth and clinical attachment level in two types of interventions (APC + OFD and APC + OFD + BG) but it did not show any benefit for probing depth for the APC + GTR and the APC + EMD groups. For the radiographic bone defect filling outcome, there is evidence that the adjunct of APC brings benefits in two types of treatment (APC + OFD and APC + OFD + BG) but it showed insufficient advantage when associated to the treatment with EMD, and no data were available for the GTR group. In the second comparison group (APC + OFD + BG versus OFD + BG), there was evidence of an advantage of APC in all follow-up and for all three parameters: probing depth, clinical attachment level, and radiographic bone defect filling. Conversely, when APC are used in combination with GTR or EMD insufficient benefits were observed at any follow-up period except for clinical attachment level at the 3 to 6 months follow-up. This could suggest that potential benefits of APC are masked by the well known advantages of gold standard treatments for infrabony defects such as GTR and EMD.

Regarding secondary outcomes, all the studies in all groups reported a survival rate of 100% for the treated teeth. No complete pocket closure was reported. No quantitative analysis regarding patients' quality of life was possible.

Overall completeness and applicability of evidence

Even though most of the studies were conducted by experienced professionals in university settings, we believe that with the adequate training the techniques are applicable in general everyday practice and therefore the generalisation of the results of this review is feasible.

Except for the radiographic bone filling, all other clinical parameters have some level of subjectivity in terms of measurements. However, the procedure for their assessment is generally well standardized and with basic training the result can be reproducible from one practitioner to another.

The follow-up periods of the studies were, in general, adequate for each of the outcomes. All the included studies had a follow-up period of at least 3 months for clinical outcomes (probing depth and clinical attachment level), which is adequate for this type of outcome. The radiographic bone defect filling, which requires a longer time in order to be detected, was measured in the majority of the studies between 9 and 12 months.

The vast majority of the patients completed the follow-up periods in their respective studies and the dropouts never exceeded 20%. Furthermore, all 38 included studies reported the numerical data for the main clinical outcomes (probing depth and clinical attachment level), which made it possible to perform the meta-analysis with a fair number of studies.

Quality of the evidence

Even though all studies included in this review were randomised controlled trials, 36 of them had a high risk of bias and 2 had an uncertain risk of bias. Consequently, to all of our study groups a high risk of bias was assigned because more than 50% of the studies included in each group had at least one domain rated at high risk of bias. This led to a downgrade of GRADE assessments for all groups.

The body of evidence for APC + OFD versus OFD was assessed as having a very low quality for all three parameters (probing depth, clinical attachment level, and radiographic bone filling). There was evidence of high heterogeneity, however, the study population was larger than 400.

The body of evidence for APC + OFD + BG versus OFD + BG was assessed as being of very low quality for all three parameters (probing depth, clinical attachment level, and radiographic bone filling). They had an adequate study population (larger than 400) but a high heterogeneity.

The body of evidence for APC + GTR versus GTR was assessed as being of very low quality for probing depth and clinical attachment level. There was evidence of imprecision for both parameters despite a good consistency.

The body of evidence for APC + EMD versus EMD was assessed as being of very low quality for probing depth, clinical attachment level, and radiographic bone filling. There was evidence of a high imprecision for all parameters.

Potential biases in the review process

A sensitive electronic search of multiple databases was conducted to identify suitable studies for this review. We did not apply restriction of language or date of publication. For the ongoing studies that met our inclusion criteria and for already published studies with missing data, we directly contacted the corresponding au-

thors, but we were not always able to have a response from them. This led to an exclusion of all missing data from our review. One of the present review authors (Massimo Del Fabbro) is also among the authors of one of the reviews used as a comparative for the outcomes of the current review. We addressed this bias by not involving this author at the evaluation process of the 'Agreements and disagreements with other studies or reviews' session.

This review was aimed at analysing the effect of any type of autologous platelet concentrate for enhancing healing of infrabony defects, and no separate analysis was done for each type of APC. It is possible that the effect of different APCs is different in different subgroups, but since no study was found that compared two or more APCs among them and with a control group, we abandoned the idea of a comparison between APCs.

Agreements and disagreements with other studies or reviews

In general our results were concordant with those of previous systematic reviews.

A systematic review published in the *Journal of Periodontology* (Del Fabbro 2011) included 16 studies that evaluated treatme outcomes of infrabony defects and gingival recession with our without the adjunct of platelet-rich plasma (PRP). They foun significant positive effect of the adjunct of PRP to OFD the clinical attachment level parameter of infrabony defects On t other hand, no significant difference was found between roup with or without PRP in infrabony defects treated with G ese results are in agreement with the results of our curre t review.

Another review (Roselló-Camps 2015) evalua d 2 stu es about the use of PRP for periodontal regenerat n co d to other regenerative procedures such as GTR. Si r to our results they found that APC significantly im ove clinica ttachment level and radiographic bone filling. H owever ney did not find additive benefits of APC for probing dep re uction.

Finally, a recent review (Castro 2017) nalysed 21 articles about the use of leukocyte- and platelet-rich fibrin (L-PRF). Similar to our systematic review Ca o et al found that APC was beneficial for probing depth reductio clinical attachment level gain and radiographic bone fi ing, hen comparing to OFD alone. However, they did not fi d differe ces on these outcomes when L-PRF was compar to treat nts consisting of a connective tissue graft utilisatio

AUTHORS' CONCLUSIONS

Implications for practice

This review found very low-quality evidence that the adjunct of autologous platelet concentrates (APC) to specific surgical techniques such as open flap debridement (OFD) and OFD + bone graft (BG) when treating infrabony defects, may improve probing pocket depth, clinical attachmen level, and radiographic bone defect filling outcomes. For guided ssue regeneration (GTR) and enamel matrix derivati e (E D) in ventions, insufficient evidence of an advantage in ing APC w s observed. The number of studies concerning t e tec iques as very limited (only two studies for EMD) an their ality was assessed as very low. Consequently, these asse ents cannot be conclusive.

Implic tions for esearch

The main roblem w encountered while performing this review, was the high of bias for almost all included studies. Even thou h we very well understand the many difficulties in carrying ou randomised controlled trial, such a standard of evidence i da y in order to come to conclusive results and clinical gui lines. Furthermore, for some specific interventions such as d EMD, there are few studies available that can be consulted in order to formulate conclusions. Therefore, we encourage in estigators to further investigate this argument and to increase e quality of the evidence with attention paid to allocation concealment and blinding of the personnel which were not correctly performed in the majority of studies. Additionally, we advise authors of future studies to follow the CONSORT Statement, to clearly detail baseline and follow-up data for the clinical outcomes and to always perform a sample size calculation.

Lastly, because of very few data available, we could not include in this review a comparison among different types of APC. Therefore, we encourage authors of future studies, to compare in the same study, different types of APC in combination with different surgical interventions in order to assess if one type of APC is more beneficial than another one when used as an adjunct to a specific surgical technique.

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* *Indicates the major publication for the study*

CHARACTERISTICS OF STUDIES

Characteristics of included studies *[ordered by study ID]*

Agarwal 2014

Methods	Trial design: randomised, split-mouth trial Location: Aligarh, India Number of centres: 1: Department of Periodontics, Dental College, Aligarh, India Recruitment period: not stated Source of funding: not stated Ethical approval: yes, ethical committee of Aligarh Muslim University Number of surgeons: 1
Participants	Inclusion criteria: absence of any systemic disease, not taking any medication, no pregnancy or lactation, non-smokers, no previous treatment for periodontal reasons, no furcation involved, matched pairs of intrabony defects with PD ≥ 6 mm following initial therapy Exclusion criteria: failure to satisfy inclusion criteria Age at baseline: mean not specified, range 30 to 65 years Gender: F 10/M 14 Smokers: excluded Teeth treated: maxillary, mandibular premolars and first/second molars Number randomised (participants/teeth): 24/48 Number evaluated (participants/teeth): 24/48
Interventions	Comparison: PRP + DFDBA versus DFDBA + saline solution Test group: PRP + DFDBA (n = 24 defects) Control group : DFDBA + saline (n = 24 defects) Surgical technique: OFD with the adjunct of a graft with DFDBA + PRP in test and saline in control Follow-up duration: 12 months
Outcomes	Clinical: PD, CAL Radiographic: CEJ-AC, AC-BD, CEJ-BD, defect width
Notes	Sample size calculation not reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, but not specified if it was assessed Complications reported: yes (no complications)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "defects were randomly divided into 2 groups by the flip of a coin" Comment: correct method for random sequence generation

Agarwal 2014 (*Continued*)

Allocation concealment (selection bias)	Unclear risk	Quote: "defects were randomly divided into 2 groups by the flip of a coin" Comment: not sufficient information provided for allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the clinician given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "Clinical parameters were recorded preoperatively and at 12 months postoperatively by one trained examiner who was blind to the treatment assignments. Radiographs were assessed on a light box by a single experienced clinician who was blind to the treatment used" Comment: blinding likely to have been done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised participants were included in the analyses
Selective reporting (reporting bias)	Low risk	Data for outcomes of this review were reported appropriately

Agarwal 2015

Methods	Trial design: randomised, split-mouth trial Location: Aligarh, India Number of centres: 1: Department of Periodontics, Dental College, Aligarh, India Recruitment period: not specified Source of funding: not reported Ethical approval: ethical committee of Dr ZA Dental College, Aligarh Number of surgeons: 1
Participants	Inclusion criteria: presence of a matched pair of interproximal, intrabony defects with PD $\geq$ 6 mm with defect depth $\geq$ 4 mm, in asymptomatic posterior teeth. Osseous defects needed to have 2 and/or 3 walls. The plaque and gingival indices, associated with interested tooth, achieved following re-evaluation of initial therapy had to be $\leq$ 1. Radiographic evidence of intrabony defects Exclusion criteria: presence of any systemic disease, patients taking any medication, pregnancy or lactation, smokers, previously treated for periodontal reasons, 1-wall defects and furcation involvement Age at baseline: mean age = 52 ± 7 years Gender: F 14/M 18 Smokers: excluded Teeth treated: 64 Number randomised (participants/teeth): 32/64

Agarwal 2015 (*Continued*)

	Number evaluated (participants/teeth): 30/60
Interventions	Comparison: PRF + DFDBA versus DFDBA + saline solution Test group: PRF + DFDBA Control group: DFDBA + saline Surgical technique: open flap debridement wit[illegible]he adjun[illegible]f a graft with DFDBA + PRP in test and saline in control Follow-up duration: 12 months
Outcomes	Clinical: PD, CAL, measured from [illegible] CEJ Radiographic: CEJ-AC, AC-BD, CEJ-BD. Differences between pre- and postoperative RBL measurements were consid[illegible]red as the ra[illegible]iographic bone loss/gain
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "The study used a split-mouth design, in which 2 interproximal sites were randomly (toss of a coin, performed by the study therapists) assigned to the DFDBA with saline or DFDBA with the PRF group" Comment: correct method for random sequence generation
Allocation concealment (selection bias)	Unclear risk	Quote: "The study used a split-mouth design, in which 2 interproximal sites were randomly (toss of a coin, performed by the study therapists) assigned to the DFDBA with saline or DFDBA with the PRF group" Comment: insufficient information for allocation concealment
Blinding of parti[illegible]pants [illegible]d personnel (performan[illegible] bias) All outco[illegible]es	High risk	Impossible to blind the clinician given the surgical nature of the treatment
Blinding of o[illegible]come assessment (detection bias) All outcomes	Low risk	Quotes: "The research was designed as a randomized, double-blinded, parallel, controlled clinical trial.." and "One operator (AA) performed all the surgeries, whereas another operator (NDG) performed all the clinical and radiographic measurements without knowledge of the groups" Comment: blinding likely to have been

Agarwal 2015 (*Continued*)

		done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	Only 2 patients (4 sites) did not return for follow-up examinations, 1 of the test group and 1 of the control group
Selective reporting (reporting bias)	Low risk	Data for outcomes of this review were reported appropriately

Agarwal 2016

Methods	Trial design: randomised, split-mouth trial Location: Institute of Dental Sciences, Bareilly, India Number of centres: 1 Recruitment period: not specified Source of funding: nil Ethical approval: Ethics Committee of MJP Rohilkhand University, Bareilly, India Number of surgeons: not reported
Participants	Inclusion criteria: adult patients in good general health and diagnosed with chronic advanced periodontitis, presence of 3 deep intrabony defects (3-walled) with a PD > 5 mm located in the interproximal area in maxillary or mandibular posterior teeth in 3 different quadrants. Radiographic evidence of the defects should exist Exclusion criteria: smoking, antibiotic, or anti-inflammatory treatment or the known use of any medication with the potential to affect periodontal tissues within the preceding 3 months and pregnancy Age at baseline: not reported Gender: F 3/M 7 Smokers: excluded Teeth treated: not specified Number randomised (participants/teeth): 10/30 Number evaluated (participants/teeth): 8/28
Interventions	Comparison: the control group (C) consisted of sites treated with OFD alone. Whereas, test group A consisted of sites treated with PRP alone and test group B received PRP in combination with DFDBA Test group: OFD + PRP and PRP + DFDBA Control group: OFD Surgical technique: OFD Follow-up duration: 12 months
Outcomes	Clinical: PI, GI, PD, and CAL Radiographic: defect depth reduction and defect resolution. Defect fill was assessed by measuring distance between CEJ and base of the defect. The distance between alveolar crest and base of the defect depicted defect resolution. Change in alveolar crest level was also seen as a measurement of distance between CEJ and alveolar crest
Notes	

Agarwal 2016 (*Continued*)

Risk of bias		
Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Unclear risk	Quote: "The defects were assigned randomly to 3 groups" Comment: insufficient information to determine method of random sequence generation
Allocation concealment (selection bias)	Unclear risk	Quote: "The defects were assigned randomly to 3 groups" Comment: insufficient information to determine method of allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the clinician given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No information was provided
Incomplete outcome data (attrition bias) All outcomes	Low risk	Only 2 patients (2 defects) did not return for follow-up examination
Selective reporting (reporting bias)	Low risk	Data for outcomes of this review were reported appropriately

Arabaci 2017

Methods	Trial design: randomised, split-mouth trial Location: Atatürk University, Department of Periodontics, Faculty of Dentistry, Erzurum, Turkey Number of centres: 1 Recruitment period: October 2013 to September 2015 Source of funding: the Scientific Research Fund of Atatürk University (AtaUni BAP-2011/300) Ethical approval: ethics committee of Atatürk University Faculty of Dentistry, Turkey Number of surgeons: 1
Participants	Inclusion criteria: patients with moderate to severe chronic periodontitis with PD $\geq$ 5 mm and horizontal bone loss of at least 2 quadrants of the jaws after phase I therapy (SRP) Exclusion criteria: smoking or tobacco use in any form; medications known to affect periodontal treatment and blood coagulation; systemic conditions known to affect periodontal status; pregnancy/lactation; and disagreeable oral hygiene (PI > 1.5). Patients with teeth with 3-wall deep intrabony defects, gingival recession, endodontic lesion, or

Arabaci 2017 (*Continued*)

	furcation involvement were also excluded Age at baseline: 29 to 46 years (mean age = 36.49 ± 7.03 years) Gender: F 9/M 17 Smokers: excluded Teeth treated: tooth type was not specified Number randomised (participants/teeth): 26/52 Number evaluated (participants/teeth): 26/52
Interventions	Comparison: OFD + PRF versus OFD alone Test group: OFD + PRF (n = 26 defects) Control group: OFD alone (n = 26 defects) Surgical technique: full-thickness mucoperiosteal flap with PRF in test site and full-thickness mucoperiosteal flap alone in control site Follow-up duration: 9 months
Outcomes	Clinical: PI, modified sulcus bleeding index, PD, relative attachment level, gingival margin level Radiographic: not reported Other: levels of growth factors (fibroblast growth factor-2 (FGF-2), platelet-derived growth factor-BB (PDGF-BB), and transforming growth factor-beta (TGF-β)) in the gingival crevicular fluid
Notes	Sample size calculation: not reported Full-mouth radiographs were taken only for diagnostic purpose Comparability at baseline: assessed for biochemical parameters, not reported for clinical parameters Communications: not reported Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "The chosen sites were disunited fortuitously (using a coin toss method) into test groups and control" Comment: correct method for random sequence generation
Allocation concealment (selection bias)	Low risk	Quote: "The chosen sites were disunited fortuitously (using a coin toss method) into test groups and control" comment: correct method for allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment. The patients were blinded to their treatment group allocation. However blinding of the pa-

Arabaci 2017 (*Continued*)

		tients is unlikely to influence treatment outcome again because of the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quotes: "The study, conducted from October 2013 to September 2015, was planned as a randomized double-blinded, controlled clinical trial that used a split-mouth design test" and "A single periodontal surgeon (TA) carried out all the surgical procedures and a second operator (AK) performed all clinical measurements without information of the groups" Comment: blinding of outcomes assessment done correctly
Incomplete outcome data (attrition bias) All outcomes	Low risk	There were no dropouts and outcomes were reported for all patients
Selective reporting (reporting bias)	Low risk	Data for outcomes of this review were reported appropriately

Aydemir 2016

Methods	Trial design: randomised, split-mouth trial Location: Kirikkale University, Periodontology Department, Turkey Number of centres: 1 Recruitment period: February to August 2014 Source of funding: authors' institution Ethical approval: yes Number of surgeons: 1
Participants	Inclusion criteria: existence of chronic periodontitis showing similar bilateral defects with minimum width of 2 mm and a maximum width of 4 mm in a radiographic evaluation at least 6 weeks after phase I therapy (consisted of SRP, oral hygiene instructions and occlusal adjustment, if necessary); existence of at least 2 mm keratinised gingiva; absence of caries and/or untreated endodontic problems; and full mouth plaque and bleeding scores $\leq$ 20 after phase I therapy Exclusion criteria: defects extending to the furcation area were not included. Systemic conditions, such as diabetes mellitus, rheumatoid arthritis, pregnancy, or lactation, that may affect the periodontal state or healing; antibiotic use in the last 6 months; and smoking (current, occasional or former) Age at baseline: mean = 38.5 ± 9.24 years Gender: F 14/M 14 Smokers: excluded Teeth treated: not specified Number randomised (participants/teeth): 28/56 Number evaluated (participants/teeth): 24/49

Aydemir 2016 (*Continued*)

Interventions	Comparison: EMD + PRF and EMD Test group: EMD + PRF (25 defects) Control group : EMD (24 defects) Surgical technique: OFD Follow-up duration: 6 months
Outcomes	Clinical: GI, PI, PD, CAL, GR Radiographic: total defect depth from the CEJ to the base of the defect at a line tangent to the adjacent root surface; suprabony defect depth from the CEJ to the alveolar crest; defect width: the horizontal distance from the alveolar crest to the root surface; defect angle: the angle between the line connecting the CEJ to the base of the defect and the lateral border of the defect; linear bone growth and bone fill percentage (BF%)
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "To assign the defects into 2 groups, EMD + PRF (28 defects - test) and EMD (28 defects - control), a computer-generated randomization scheme (without blocking) was utilized by 1 author (AD)" Comment: correct method for random sequence generation
Allocation concealment (selection bias)	Low risk	Quote: "The use of opaque, numbered envelopes that contained the assigned intervention concealed the allocation" Comment: correct method for allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "The author who performed the measurements on the participants (HGK) and the statistician (AD) were blinded to the surgical procedures and measurements" Comment: blinding done correctly
Incomplete outcome data (attrition bias) All outcomes	Low risk	Only 3 patients (6 defects) did not return for follow-up examination and the reason was provided. (Data of 1 patient from the EMD group was removed from the study due to an acute mechanical trauma 7 days

Aydemir 2016 (*Continued*)

		after surgery)
Selective reporting (reporting bias)	Low risk	Data for outcomes of this review were reported appropriately

Camargo 2009

Methods	Trial design: randomised, split-mouth trial Location: School of Dentistry, University of Belgrade, Belgrade, Republic of Serbia Number of centres: 1 Recruitment period: 15 May 1999 to 20 March 2000 Source of funding: not stated Ethical approval: yes, University Institutional Review Board Number of surgeons: 2
Participants	Inclusion criteria: patients having 2 similar interproximal defects with PD > 6 mm after initial therapy. Radiographic evidence of intrabony defects had to exist. Upon surgical exposure, defects needed to have a minimum depth of 3 mm and present with 2 or 3 walled defects Exclusion criteria: systemic illnesses, compromised immune system, pregnant and/or lactating women, and patients taking any drug known to cause gingival enlargement. Patients allergic or sensitive to any of the medications to be used, teeth non-responsive to cold or endodontically treated Age at baseline: 34 to 67 years (mean age = 47 ± 10 years) Gender: F 14/M 9 Smokers: 11 smokers/12 non-smokers Jaws treated: maxillary and mandibular posteriors Number randomised (participants/teeth): 23/46 Number evaluated (participants/teeth): 23/46
Interventions	Comparison: PRP/BPBM/GTR versus BPBM/GTR Test group: PRP/BPBM/GTR (n = 23) Control group: BPBM/GTR (n = 23) Surgical technique: intrabony defects treated with PRP/BPBM/GTR for test group and BPBM/GTR for control group Follow-up duration: 6 months
Outcomes	Clinical: PD, CAL, defect fill (re-entry surgery) Radiographic: none Other: alveolar crest resorption
Notes	Sample size calculation: not reported Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Camargo 2009 (*Continued*)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "T[illegible] study used a split-mouth design, and 2 [illegible]terproximal sites were ran[illegible]mly ([illegible] of [illegible]coin) assigned to the control a[illegible] experime[illegible]tal groups" C[illegible]ment: [illegible]rrec[illegible] method for random sequence [illegible]eneration
Allocation concealment (selection bias)	Low risk	Quo[illegible]: "The study used a split-mouth design, and 2 interproximal sites were ran[illegible]omly (toss of a coin) assigned to the control and experimental groups" Comment: correct method for allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low ris[illegible]	Quote: "An examiner other than the surgeons performed all clinical measurements without knowledge of the treatment groups" Comment: blinding of outcome assessment done properly
Incomplete outcome data (attrition bias) All outcomes	Lo[illegible]ris[illegible]	All randomised patients completed the study
Selective reporting (reporting [illegible]ias)	Low risk	Data for outcomes of this review were reported appropriately

Chandradas 2016

Methods	Trial design: randomised, parallel trial Location: Department of Periodontics, Sree Mookambika Institute of Dental Sciences, India Number of centres: 1 Recruitment period: not specified Source of funding: nil Ethical approval: institutional ethics committee Number of surgeons: 1
Participants	Inclusion criteria: systemically healthy patients diagnosed with chronic periodontitis based on the international workshop for the classification of periodontal disease, having ≥ 20 teeth and ≥ 30% of sites with > 4 mm clinical attachment loss, PD ≥ 5 mm, and presence of intrabony defect ≥ 3 mm (measured from alveolar crest to the base of the

Chandradas 2016 (*Continued*)

	defect on intraoral periapical radiograph) Exclusion criteria: patients with use of tobacco or tobacco-related products; systemic or local application of antibiotics within the previous 6 months; patients with poor oral hygiene (PI ≥ 3) after the revaluation of cause-related th. rapy Age at baseline: 44.4 years Gender: F 18/M 18 Smokers: excluded Teeth treated: maxilla and mandible Number randomised (participants/teeth): 3 /36 Number evaluated (participants/teeth): 36/3
Interventions	Comparison: group A, PRF + D M; group B PRF alone; and group C, control (OFD) Test groups: PRF + DBM (n = 1), PRF alo e (n = 12) Control group: OFD (n = 12) Surgical technique: OFD Follow-up duration: 9 month
Outcomes	Clinical: GI, GR, PD, rela ve attachment level was measured from apical border of the stent to the base of the p Radiographic: li ear one growth and percentage in bone fill
Notes	

Risk of bias

Bias	A s' judgement	Support for judgement
Random sequence generation (selectio bias)	L risk	Quote: "Allotment of participants within the groups was performed randomly by creating a randomization list by means of a freeware link (http://www.graphad.com/quickcalcs/randomize1.cfm)" Comment: likely to have been done properly
Allocation concealment (s tion bias)	Low risk	Quote: "The treatment allocation of the patients was prepared and sealed in the numbered opaque envelopes and were opened during surgery immediately after completing the defect debridement. Allocation protocol was unavailable to the periodontal examiner (RS) throughout the study" Comment: correct method for allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment. The patients were blinded to their treatment group allocation. However blinding of the pa-

Chandradas 2016 (*Continued*)

		tients is unlikely to influence treatment outcome again because of the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "The pre- and postoperative assessments were performed by another examiner (...) without knowledge of the nature of intervention" Comment: blinding done correctly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised patients completed the study
Selective reporting (reporting bias)	Low risk	Data for outcomes of this review were reported appropriately

Christgau 2006

Methods	Trial design: randomized, split-mouth trial Location: Department of Operative Dentistry and Periodontology, University of Regensburg, Germany Number of centres: 1 Recruitment period: not stated Source of funding: reported, Robert Mathys Foundation, Bettlach, Schweiz Ethical approval: ethical committee of the medical facility of University of Regensburg Number of surgeons: 1
Participants	Inclusion criteria: patient having 1 pair of contralateral deep, intrabony, inter-proximal periodontal defects with a PPD of at least 6 mm, radiographic evidence of angular bone loss of at least 4 mm at baseline, none of the defects to show furcation involvement Exclusion criteria: not meeting the inclusion criteria Age at baseline: 26 to 62 years (median 42 years) Gender: F 15/M 10 Smokers: 5 patients (smoking 8 cigarettes per day) Teeth treated: not specified Number randomised (participants/teeth): 25/50 Number evaluated (participants/teeth): 25/50
Interventions	Comparison: β-TCP/GTR + APC versus β-TCP/GTR Test group: β-TCP/GTR + APC (n = 25) Control group : β-TCP/GTR (n = 25) Surgical technique: intrabony defects were treated with β-TCP/GTR bioresorbable barrier membrane at control site and APC was additionally applied on test group Follow-up duration: 12 months
Outcomes	Clinical: papillary bleeding index, approximal plaque index, CAL, gingival recession, PPD, depth of osseous defect Radiographic: digital subtraction radiography - bone density

	Other: vertical relative attachment gain
Notes	Sample size calculation: not reported Radiographs were taken at baseline and at end of follow-up to analyse by digital subtraction radiography Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "For randomized treatment allocation, a randomizing table was created by our mathematician (K-AH) using the SPSS software (Version 13.0, SPSS Inc., Chicago, IL, USA)" Comment: likely to have been done properly
Allocation concealment (selection bias)	Low risk	Quote: "The randomization table comprised the patient numbers (1-25) and the corresponding defect numbers (1 and 2) per patient. The therapy methods (test or control) were randomly allocated to the defect numbers. By entering the study, the patient numbers were consecutively allocated to the patients and the defect numbers were allocated to the 2 teeth to be treated. Treatment allocation was concealed to the surgeon until the beginning of the surgery" Comment: likely to have been done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "Clinical examination was performed by 2 masked examiners .." Comment: blinding of outcome assessment done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised patients completed the study
Selective reporting (reporting bias)	Low risk	All data were properly reported

Demir 2007

Methods	Trial design: randomised, parallel trial Location: Department of Periodontology, Faculty of Dentistry, Hacettepe University, Ankara, Turkey Number of centres: 1 Recruitment period: not stated Source of funding: mentioned, The Research Foundation of Hacettepe University Ethical approval: yes, Faculty of Medicine, Ethical Committee of Medical, Surgical and Drug Research, Hacettepe University Number of surgeons: 1
Participants	Inclusion criteria: patient with no systemic disease, having a good level of oral hygiene, mobility < 1 mm in total, radiographic evidence of vertical alveolar bone loss at the mesial aspect of the tooth, presence of a mesial inter-proximal probing pocket depth > 6 mm following initial therapy, no prosthetic restoration or endodontic treatment on the related tooth, any medications affecting the coagulation mechanism Exclusion criteria: failing to meet the inclusion criteria Age at baseline: mean = 36.03 ± 12.02 years Gender: F 16/M 13 Smokers: yes (9 smokers - 6 to 10 cigarettes per day) Teeth treated: maxillary and mandibular anterior and posterior teeth Number randomised (participants/teeth): 29/29 Number evaluated (participants/teeth): 29/29
Interventions	Comparison: PRP/BG versus BG alone Test group: PRP/BG (n = 15) Control group: BG alone (n = 14) Surgical technique: OFD + intrabony defects treated with BG in control group and addition of PRP with BG in test group Follow-up duration: 9 months
Outcomes	Clinical: PI, GI, BOP, PD, GR, CAL Radiographic: none reported Other: surgical re-entry (CEJ-BD, CEJ-CD, intrabony defect depth)
Notes	Sample size calculation: not reported Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Patients included in the study were divided into 2 groups randomly by the flip of a coin" Comment: correct method for random sequence generation

Demir 2007 (*Continued*)

Allocation concealment (selection bias)	Low risk	Quote: "Patients included in the study were divided into 2 groups randomly by the flip of a coin" Comment: correct method for allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "All clinical and intrasurgical measurements were performed by a single examiner (author AB) at baseline and 9 months after the surgical procedure without knowledge of the treatment groups" Comment: blinding done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised patients completed the study
Selective reporting (reporting bias)	Low risk	Data were properly reported

Döri 2007a

Methods	Trial design: randomised, parallel trial Location: Semmelweis University, Budapest, Hungary Number of centres: 1 Recruitment period: not stated Source of funding: Semmelweis University Ethical approval: yes, University Ethical Board Number of surgeons: 1
Participants	Inclusion criteria: no systemic diseases that could influence the outcome of the therapy, good level of oral hygiene - plaque index, compliance with the maintenance program and presence of 1 intrabony defect with a PD of at least 6 mm and an intrabony component of at least 3 mm as detected on the radiographs, non-smoker Exclusion criteria: failing to meet inclusion criteria Age at baseline: 28 to 56 years Gender: F 16/M 14 Smokers: excluded Teeth treated: maxillary and mandibular anterior, premolars and molars Number randomised (participants/teeth): 30/30 Number evaluated (participants/teeth): 30/30
Interventions	Comparison: PRP + NBM/GTR versus NBM/GTR Test group: PRP + NBM/GTR (n = 15/15) Control group: NBM/GTR (n = 15/15) Surgical technique: intrabony defects treated with NBM/GTR in control group and with

Döri 2007a (*Continued*)

	addition of PRP in test group Follow-up duration: 1 year
Outcomes	Clinical: PI, GI, BOP, PD, GR, CAL Radiographic: not reported Other: INTRA (defined as the distance from the alveolar bone crest to the bottom of the defect) (before surgery)
Notes	Sample size calculation: reported Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropout

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "The defects were randomly assigned before surgery to the 2 treatment groups with the randomized block approach. Blocking was performed to control for the effects of the prognostic variables INTRA and CAL to decrease outcome variability (Fleiss 1986). For allowing randomization, INTRA (defined as the distance from the alveolar bone crest to the bottom of the defect) was estimated before surgery on pre-operative radiographs and by performing trans-gingival bone sounding" Comment: random sequence generation likely to have been done properly
Allocation concealment (selection bias)	Low risk	Quote: "The defects were randomly assigned before surgery to the 2 treatment groups with the randomized block approach. Blocking was performed to control for the effects of the prognostic variables INTRA and CAL to decrease outcome variability (Fleiss 1986). For allowing randomization, INTRA (defined as the distance from the alveolar bone crest to the bottom of the defect) was estimated before surgery on pre-operative radiographs and by performing transgingival bone sounding. In each case, the surgeon was informed of the assigned treatment option after completion of flap elevation and defect debridement. Also, blood samples were collected from all

Döri 2007a (*Continued*)

		patients regardless of the subsequent PRP application" Comment: allocation concealment likely to have been done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "The examiner was not aware, in any of the cases, of the type of treatment rendered" Comment: blinding done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Döri 2007b

Methods	Trial design: randomised, parallel trial Location: Department of Periodontology, Semmelweis University, Budapest, Hungary Number of centres: 1 Recruitment period: July 2002 to September 2003 Source of funding: not stated Ethical approval: yes, Semmelweis University Ethical Board Number of surgeons: 1
Participants	Inclusion criteria: patient having no systemic diseases that could influence the outcome of the therapy; having good level of oral hygiene (PI < 1); having compliance with the maintenance program; with presence of 1 intrabony defect with PD > 6 mm and an intrabony component (INTRA) > 3 mm as detected on the radiographs and measured at bone sounding; no intrabony defects extending into a furcation area; and no teeth presenting furcation involvements Exclusion criteria: patients failing to meet the inclusion criteria Age at baseline: 26 to 55 years Gender: F 14/M 10 Smokers: none of the patients were smokers Teeth treated: maxillary and mandibular anterior, premolars and molars Number randomised (participants/teeth): 24/24 Number evaluated (participants/teeth): 24/24
Interventions	Comparison: PRP + ABBM + GTR versus ABBM + GTR Test group: PRP + ABBM + GTR (n = 12/12) Control group: ABBM + GTR (n = 12/12) Surgical technique: intrabony defects were treated with ABBM + GTR in control group and PRP was additionally applied in test group

Döri 2007b (*Continued*)

	Follow-up duration: 1 year
Outcomes	Clinical: PI, GI, BOP, PD, GR, and CAL Radiographic: preoperative non-standardized radiographs were taken with the long cone parallel technique for the purpose of baseline defect characteristics for inclusion Other: none reported
Notes	Sample size calculation: reported Radiographs were taken without a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "Using a randomized block approach, the defects were randomly assigned before surgery to the 2 treatment groups. Blocking to control for the effects of the prognostic variables, the distance from the alveolar bone crest to the bottom of the defect (INTRA) and CAL was used to decrease outcome variability. 34 INTRA was estimated before surgery based on radiographs and transgingival bone sounding recordings" Comment: random sequence generation likely to have been done properly
Allocation concealment (selection bias)	Low risk	Quote: "Using a randomized block approach, the defects were randomly assigned before surgery to the 2 treatment groups. Blocking to control for the effects of the prognostic variables, the distance from the alveolar bone crest to the bottom of the defect (INTRA) and CAL was used to decrease outcome variability. 34 INTRA was estimated before surgery based on radiographs and transgingival bone sounding recordings" Comment: allocation concealment likely to have been done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment

Döri 2007b (*Continued*)

Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "The examiner was not aware, in any of the cases, of the type of treatment rendered" Comment: blinding done correctly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Döri 2008a

Methods	Trial design: randomised, parallel trial Location: Department of Periodontology, Semmelweis University, Budapest, Hungary Number of centres: 1 Recruitment period: June 2002 and November 2003 Source of funding: Department of Periodontology, Semmelweis University. Part of the grafting material was provided by Curasan, Kleinostheim, Germany Ethical approval: yes, Semmelweis University Ethical Board Number of surgeons: 1
Participants	Inclusion criteria: patient having no systemic diseases that could influence the outcome of the therapy; having good level of oral hygiene (PI < 1); having compliance with the maintenance program; with presence of 1 intrabony defect with PD > 6 mm and an intrabony component > 3 mm as detected on the radiographs and measured at bone sounding; no intrabony defects extending into a furcation area; and no teeth presenting furcation involvements Exclusion criteria: patients failing to meet the inclusion criteria Age at baseline: 28 to 58 years Gender: F16/M 12 Smokers: none of the patients were smokers Teeth treated: maxillary and mandibular anterior, premolars and molars Number randomised (participants/teeth): 28/28 Number evaluated (participants/teeth): 28/28
Interventions	Comparison: PRP + β-TCP + GTR versus β-TCP + GTR Test group: PRP + β-TCP + GTR (n = 14/14) Control group: β-TCP + GTR (n = 14/14) Surgical technique: intrabony defects were treated with β-TCP + GTR in control group and PRP was additionally applied in test group Follow-up duration: 1 year
Outcomes	Clinical: PI, GI, BOP, PD, GR, and CAL Radiographic: preoperative non-standardized radiographs were taken with the long cone parallel technique for the purpose of baseline defect characteristics for inclusion Other: none reported

Döri 2008a (*Continued*)

Notes	Sample size calculation: reported Radiographs were taken without a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "Using a randomized block approach, the defects were assigned to the 2 treatment groups before surgery. Blocking to control for the effects of the prognostic variables, INTRA (the distance from the alveolar bone crest to the bottom of the defect) and CAL were used to decrease outcome variability. 42 INTRA was estimated before surgery based on radiographs and transgingival bone sounding recordings" Comment: random sequence generation likely to have been done properly
Allocation concealment (selection bias)	Low risk	Quote: "Using a randomized block approach, the defects were assigned to the 2 treatment groups before surgery. Blocking to control for the effects of the prognostic variables, INTRA (the distance from the alveolar bone crest to the bottom of the defect) and CAL were used to decrease outcome variability. 42 INTRA was estimated before surgery based on radiographs and transgingival bone sounding recordings" Comment: allocation concealment likely to have been done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "The examiner was not aware of the type of treatment rendered" Comment: blinding done correctly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Döri 2008b

Methods	Trial design: randomised, parallel trial Location: Department of Periodontology, Semmelweis University, Budapest, Hungary Number of centres: 1 Recruitment period: September 2004 and September 2005 Source of funding: the study was funded by the author's own institution. Part of the graft material was kindly provided by Geistlich, Wolhusen, Switzerland Ethical approval: yes, Semmelweis University Ethical Board Number of surgeons: 1
Participants	Inclusion criteria: patient having no systemic diseases that could influence the outcome of the therapy; having good level of oral hygiene (PI < 1); having compliance with the maintenance program; with presence of 1 intrabony defect with PD at least 6 mm and an intrabony component > 4 mm as detected on the radiographs Exclusion criteria: patients failing to meet the inclusion criteria Age at baseline: 32 to 56 years Gender: F 14/M 12 Smokers: none of the patients were smokers Teeth treated: maxillary and mandibular anterior, premolars and molars Number randomised (participants/teeth): 26/26 Number evaluated (participants/teeth): 26/26
Interventions	Comparison: EMD + NBM + PRP versus EMD + NBM Test group: EMD + NBM + PRP (n = 13/13) Control group: EMD + NBM (n = 13/13) Surgical technique: intrabony defects were treated with EMD + NBM in control group and PRP was additionally applied in test group Follow-up duration: 1 year
Outcomes	Clinical: PI, GI, BOP, PD, GR, and CAL Radiographic: preoperative non-standardized radiographs were taken with the long cone parallel technique for the purpose of baseline defect characteristics for inclusion Other: none reported
Notes	Sample size calculation: reported Radiographs were taken without a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "The defects were randomly assigned before surgery to the 2 treatment groups with the randomized block approach. Blocking to control for the effects of the prognostic variables INTRA and CAL was used to decrease outcome vari-

Döri 2008b (*Continued*)

		ability (Fleiss 1986). To allow randomization, INTRA (defined as the distance from the alveolar bone crest to the bottom of the defect) was estimated before surgery on pre-operative radiographs and by performing transgingival bone sounding" Comment: random sequence generation [illegible] properly
Allocation concealment (selection bias)	Low risk	Quote: "The defects were randomly assigned before surgery to the 2 treatment groups with the randomized block approach. Blocking to control for the effects of the prognostic variables INTRA and CAL was used to decrease outcome variability (Fleiss 1986). To allow randomization, INTRA (defined as the distance from the alveolar bone crest to the bottom of the defect) was estimated before surgery on pre-operative radiographs and by performing transgingival bone sounding" Comment: allocation concealment likely to have been done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "The examiner was not aware, in any of the cases, of the type of treatment administered" Comment: blinding done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Döri 2009

Methods	Trial design: randomised, parallel trial Location: Department of Periodontology, Semmelweis University, Budapest, Hungary Number of centres: 1 Recruitment period: June 2006 and May 2007 Source of funding: stated, Department of Periodontolog[y] and Oral and Maxillofacial Surgery, Semmelweis University Ethical approval: yes, Semmelweis University Eth[ical] Board Number of surgeons: 1
Participants	Inclusion criteria: patient having no systemic diseases that could influence the outcome of the therapy; having good level of oral hygiene (PI < 1); having compliance with the maintenance program; with presence of 1 intrabony defect with PD > 6 mm and an intrabony component (INTRA) ≥ 3 mm as detected on the radiographs and measured at bone sounding; no intrabony defects extending into a furcation area; and no teeth presenting furcation involvements Exclusion criteria: patients failing to meet the inclusion criteria Age at baseline: 28 to 65 years Gender: F 21/M 9 Smokers: none of the patients were smokers Teeth treated: maxillary and mandibular anterior, premolars and molars Number randomised (participants/teeth): 30/30 Number evaluated (participants/teeth): 30/30
Interventions	Comparison: PRP + ABBM versus ABBM alone Test group: PRP + ABBM (n = 15/15) Control group: ABBM alone (n = 15/15) Surgical technique: CAF + intrabony defects were treated with ABBM alone in control group and PRP was additionally applied in test group Follow-up duration: 1 year
Outcomes	Clinical: PI, GI, BOP, PD, GR, and CAL Radiographic: preoperative non-standardized radiographs were taken with the long cone parallel technique for the purpose of baseline defect characteristics for inclusion Other: none reported
Notes	Sample size calculation: reported Radiographs were taken without a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Using a randomized block approach, the defects were randomly assigned before surgery to the 2 treatment groups... "

Döri 2009 (*Continued*)

		Comment: random sequence generation likely to have been done properly
Allocation concealment (selection bias)	Low risk	Quote: "U..ng a randomized block approach the de..cts were randomly assigned be..e surgery ..the 2 treatment groups... " Co..ent: allocation concealment likely to have bee.. done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the ..rgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "The examiner was not aware of the type of treatment rendered" Commet: blinding of outcomes assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Elgendy 2015

Methods	..y desi..a: randomised, split-mouth trial Loca..ion: Department of Periodontology, Faculty of Dentistry, October 6 University ..nd Tanta University, Egypt Nu..ber of centres: 2 Recruitment period: February to December 2013 Source of funding: not stated Ethical approval: Research Ethical Committee of Tanta University, Egypt Number of surgeons: not stated
Participants	Inclusion criteria: presence of 2 almost identical interproximal intrabony defects, 1 on either side of the arch based on radiographic observations with clinical probing depth ≥ 6 mm in teeth Exclusion criteria: any systemic disease that affect the periodontium and contraindicate for periodontal surgery; patients having insufficient platelet count for PRF preparation; patients with coagulation defect or anticoagulation treatment; pregnant or lactating mothers; postmenopausal women; people who take anti-inflammatory drugs, antibiotics or vitamins within the previous 3 months; people who use mouthwashes regularly; heavy smoking (> 10 cigarettes/day); history of alcohol abuse; unacceptable oral hygiene after the re- evaluation of phase I therapy Age at baseline: group I 44.25 ± 8.45 years, group II 39.70 ± 6.36 years Gender: not stated Smokers: heavy smokers (> 10 cigarettes/day) were excluded

Elgendy 2015 (*Continued*)

	Teeth treated: not reported Number randomised (participants/teeth): 20/40 Number evaluated (participants/teeth): 20/40
Interventions	Comparison: PRF + NcHA bone graft versus NcHA bone graft alone Test group: PRF + NcHA bone graft (n =20) Control group: NcHA bone graft alone (n = 20) Surgical technique: OFD + intrabony defect were treated with NcHA bone graft alone in control group and PRF was additionally applied in test group Follow-up duration: 6 months
Outcomes	Clinical: PI, GI, PPD, CAL Radiographic: bone density Other: none
Notes	Sample size calculation: reported Comparability at baseline: yes, assessed Complications reported: no Dropouts: not reported, reasons not given

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Unclear risk	Quote: "Selected sites were randomly divided into 2 groups" Comment: insufficient information regarding the random sequence generation method
Allocation concealment (selection bias)	Unclear risk	Quote: "Selected sites were randomly divided into 2 groups" Comment: insufficient information regarding allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No information is provided
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	It is unclear wether or not all patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Garg 2017

Methods	Trial design: randomised, parallel trial Location: Department of Periodontics, Azamgarh Dental College, Azamgarh, India Number of centres: 1 Recruitment period: March 2013 to February 2014 Source of funding: not stated Ethical approval: Institutional Ethical Committee and Review Board of the Government Dental College and Research Institute, Bangalore, India Number of surgeons: 1
Participants	Inclusion criteria: good general health with no history of allergy, presence of moderate to severe periodontitis, presence of a 3-wall intrabony defect with PD > 5 mm and CAL > 5 mm with radiographic angular defect depth > 3 mm, located in the interproximal area Exclusion criteria: medically compromised patients, smokers, generalized aggressive periodontitis, pregnant and lactating women, and teeth with grade III mobility Age at baseline: 28 to 47 yea Gender: F 15/M 9 Smokers: excluded Teeth treated: not stat Number randomised (participants/teeth): 24/24 Number evaluated (participants/teeth): 24/24
Interventions	Comparison: OFD + HA/β-TCP + PRF versus OFD + HA/β-TCP alone Test group: OFD + HA/β-TCP + PRF (n = 12/12) Control group: OFD + HA/β-TCP alone (n = 12/12) Surgical technique: a mucoperiosteal flap was elevated with sulcular incisions. After complete debridement of the defect, scaling and root planing, the defect were filled with PRF and HA/β-TCP in test site and HA/β-TCP alone in control site Follow-up duration: 9 months
Outcomes	Clinical: BOP, PD, CAL Radiographic: radiographic bone filling Other: none
Notes	Sample size calculation: not stated Standardized parallel cone technique with grid mount was used to take radiographs Comparability at baseline: yes, assessed Complications reported: no complications Dropouts: no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Quote: "Patients who met all criteria for entry into surgical phase of the study were then randomized to 'test Group-I' (PRP + HA and β-TCP) and 'control Group-II' (saline + HA and β-TCP)"

Garg 2017 (*Continued*)

		Comment: insufficient information provided on the method used for random sequence generation
Allocation concealment (selection bias)	Unclear risk	Insufficient information provided on the method used for allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Insufficient information provided
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No information provided
Selective reporting (reporting bias)	Low risk	All results properly reported

Gupta 2014

Methods	Trial design: randomised, split-mouth trial Location: not reported Number of centres: not reported Recruitment period: not stated Source of funding: reported, no funding Ethical approval: yes, Institutional Review Board Number of surgeons: not stated
Participants	Inclusion criteria: patients selected were in good general health, having an intrabony defect ≥ 2 mm with PD ≥ 6 mm Exclusion criteria: patients with abnormal platelet count, smokers, and pregnant women Age at baseline: not stated Gender: not stated Smokers: excluded Teeth treated: maxillary and mandibular arch Number randomised (participants/teeth): 10/20 Number evaluated (participants/teeth): 10/20
Interventions	Comparison: PRP/HA versus HA alone Test group: PRP/HA (n = 10) Control group: HA alone (n = 10) Surgical technique: OFD + intrabony defects were treated with HA bone graft in control group and PRP was additionally applied in test group Follow-up duration: 1 year

Gupta 2014 (*Continued*)

Outcomes	Clinical: plaque control record, BOP, PD, and relative attachment level Radiographic: INFRA (size of the defect) Other: none
Notes	Sample size calculation: not reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: no Dropouts: not reported

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Unclear risk	Quote: "10 L-AgP [localized aggressive periodontitis] patients having bilateral intrabony defect ≥ 2 mm and probing depth (PD) ≥ 6 mm were randomly treated either with the PRP/HA graft or HA graft alone" Comment: not sufficient information provided regarding the method of random sequence generation
Allocation concealment (selection bias)	Unclear risk	Quote: "10 L-AgP patients having bilateral intrabony defect ≥ 2 mm and probing depth (PD) ≥ 6 mm were randomly treated either with the PRP/HA graft or HA graft alone" Comment: not sufficient information provided regarding the method of allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No information provided
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	It is not reported wether all patients concluded the study or not
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Hanna 2004

Methods	Trial design: randomised, split-mouth, double-blinded trial Location: The University of Texas Health Science Center at Houston, Texas, USA Number of centres: 1 Recruitment period: not reported Source of funding: not stated Ethical approval: yes, Committee for the Protection of Human Subjects at the University of Texas Health Science Center at Houston, Texas, USA Number of surgeons: 1
Participants	Inclusion criteria: patients between 35 to 75 years of age; exhibited plaque score of 20% or less prior to the surgical phase; teeth with mobility less than Miller's Class III or mobile teeth requiring splinting; and teeth responding normally to vitality testing or with stable endodontic therapy Exclusion criteria: known systemic diseases and/or drug therapy known to interfere with wound healing; known drug allergies to any of the medications used in the study; using systemic antibiotics or having received antibiotic therapy in the last 3 months; abnormal platelet counts disclosed by a complete blood count (CBC) test performed within 1 month prior to surgery; and participation in other dental clinical trials Age at baseline: 35 – 74 years Gender: F 8/M 5 Smokers: yes, 1 heavy smoker (> 20 cigarettes/day) Number randomised (participants/teeth): 13/26 Number evaluated (participants/teeth): 13/26
Interventions	Comparison: BDX + PRP versus BDX alone Test group: BDX + PRP (n = 13 defects) Control group: BDX alone (n = 13 defects) Surgical technique: OFD + intrabony defects treated with BDX alone in control group and additionally PRP was applied in test group Follow-up duration: 6 months
Outcomes	Clinical: GI, PI, PD, CAL, recession as the position of the gingival margin from the CEJ, and BOP Radiographic: none reported Other: none
Notes	Sample size calculation: not reported Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Randomization was performed immediately following defect debridement by the flip of a coin"

Hanna 2004 (*Continued*)

		Comment: correct method for random sequence generation
Allocation concealment (selection bias)	Unclear risk	Quote: "Randomization was performed immediately following defect debridement by the flip of a coin" Comment: not sufficient information provided regarding the method of allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "13 patients were enrolled in a randomized, split-mouth, double-masked clinical trial" Comment: blinding of outcomes likely to have been done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Hassan 2012

Methods	Trial design: randomised, split-mouth trial Location: University of Dammam, College of Dentistry, Kingdom of Saudi Arabia Number of centres: 1 Recruitment period: not stated Source of funding: self-funded Ethical approval: yes, Ethical Committee of the College of Dentistry, Dammam University, Kingdom of Saudi Arabia Number of surgeons: 1
Participants	Inclusion criteria: patient free from any systemic diseases, non-smokers, not pregnant (female cases), had a good level of oral hygiene, and had infrabony 2 osseous walls defect with PPD 6 mm and CAL = 5 mm Exclusion criteria: failing to meet the inclusion criteria Age at baseline: mean age = 41.4 + 2.61 years Gender: F 5/M 7 Smokers: excluded Teeth treated: not stated Number randomised (participants/teeth): 12/24 Number evaluated (participants/teeth): 12/24

Hassan 2012 (*Continued*)

Interventions	Comparison: Torus mandibularis bone chips with PRP versus Torus mandibularis bone chips alone Test group: Torus mandibularis bone chips with PRP (n = 12 defects) Control group: Torus mandibularis bone chips alone (n = 12 defects) Surgical technique: OFD + intrabony defects were surgically treated using Torus mandibularis bone chips alone in control group and Torus mandibularis bone chips with PRP in test group Follow-up duration: 1 year
Outcomes	Clinical: PI, GI, PPD, CAL Radiographic: bone density, marginal bone levels Other: none
Notes	Sample size calculation: not reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: no Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "24 sites were selected by using a split-mouth design for each patient determined randomly through a biased coin randomization" Comment: random sequence generation done correctly
Allocation concealment (selection bias)	Unclear risk	Quote: "24 sites were selected by using a split-mouth design for each patient determined randomly through a biased coin randomization" Comment: not enough information to understand if allocation concealment was done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "Blinded clinical and radiological assessments were performed at baseline and after 3, 6, 9 and 12 months" Comment: blinding of outcomes likely to have been done properly

Hassan 2012 (*Continued*)

Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Kanoriya 2016

Methods	Trial design: randomised, parallel trial Location: Department of Periodontology, Government Dental College and Research Institute (GDCRI), Bengaluru, Karnataka, India Number of centres: 1 Recruitment period: October 2014 to June 2015 Source of funding: not stated Ethical approval: Institutional Ethical Committee, GDCRI, Bengaluru, Karnataka, India Number of surgeons: 1
Participants	Inclusion criteria: presence of 3-walled intrabony defects ≥ 3 mm deep (distance measured on intraoral periapical radiographs between alveolar crest and defect base) and interproximal probing depth ≥ 5 mm after etiotropic phase in asymptomatic teeth Exclusion criteria: participants with aggressive periodontitis; known systemic conditions that affect periodontal status; blood disorders and inadequate platelet count (< 200,000/mm^3); known medications that affect periodontal therapy outcomes; pregnancy or lactation; smokers and tobacco users; immunodeficient patients; allergies to bisphosphonates; and/or systemic bisphosphonate therapy. Patients with poor oral hygiene (PI > 1.5) after etiotropic phase re-evaluation were also excluded. Apart from this, furcation involved non-vital teeth, carious teeth indicated for restorative therapy, and grade II mobile teeth were also eliminated Age at baseline: mean age = 39 years Gender: F 55/M 53 (for all 3 groups) Smokers: excluded Teeth treated: 36 were maxillary and mandibular single-rooted teeth and 54 were maxillary and mandibular multirooted teeth sites Number randomised (participants/teeth): 64/64 Number evaluated (participants/teeth): 60/60
Interventions	Comparison: OFD + PRF versus OFD alone Test group: OFD + PRF (n = 30/30) Control group: OFD alone (n = 30/30) Surgical technique: full thickness mucoperiosteal flap with PRF in test site and full thickness mucoperiosteal flap alone in control site Follow-up duration: 9 months
Outcomes	Clinical: PI, modified sulcus bleeding index, PD, CAL Radiographic: radiographic bone filling Other: none

Kanoriya 2016 (*Continued*)

Notes	Sample size calculation: yes For radiographs, single customized bite blocks and paralleling technique were used. Radiographs were taken with a scanner of 6400 dots per inch Comparability at baseline: yes, assessed Complications reported: no complications Dropouts: 4 dropouts 3rd group data (PRF + 1% alendronate gel) not included in this review

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "These patients were divided into 3 groups randomly using a computer" Comment: correct method for random sequence generation
Allocation concealment (selection bias)	Unclear risk	Not enough information is provided regarding the allocation concealment method
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment. For the same reason, even though the patient was blinded, it does not influence the outcome of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "One operator (DK) performed all surgeries and a different operator (ARP) performed all parameter measurements without information about the groups" Comment: blinding of outcome assessment done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	6 patients did not complete the study (4 patients for the 2 groups considered in this review)
Selective reporting (reporting bias)	Low risk	All results properly reported

Kaushick 2011

Methods	Trial design: randomised, split-mouth trial Location: Department of Periodontics, Saveetha Dental College and Hospitals Chennai, India Number of centres: 1 Recruitment period: not stated Source of funding: nil Ethical approval: Instutional Review Board Number of surgeons: 1
Participants	Inclusion criteria: males and females aged between 20 and 50 years; attachment loss > 3 mm as assessed by periodontal probe with diagnosis of chronic periodontitis; presence of infrabony defects (2/3 wall confirmed upon surgical exposure); patients with a minimum of 2 intrabony defects in different quadrants; vital teeth; teeth with mobility less than grade I; patients willing to comply with multiple recall schedules Exclusion criteria: patients with systemic illness such as diabetes, hypertension, bleeding disorders, epilepsy, or abnormal blood picture; pregnant/lactating women; patients on medications known to cause gingival overgrowth or interfere with wound healing; patients allergic to routine medications prescribed following surgery; mucogingival problems; aggressive periodontitis; smokers; trauma from occlusion Age at baseline: 20 - 50 years Gender: not stated Smokers: excluded Teeth treated: not stated Number randomized (participants/teeth): 10/20 Number evaluated (participants/teeth): 10/20
Interventions	Comparison: PRP + bone graft (HA + ß-TCP) versus saline + bone graft (HA + ß-TCP) Test group: PRP + bone graft (HA + ß-TCP) (n = 10) Control group: saline + bone graft (HA + ß-TCP) (n = 10) Surgical technique: OFD+ intrabony defects were treated with PRP + bone graft (HA + ß-TCP) on test group sites and saline + bone graft (HA + ß-TCP) on control group Follow-up duration: 6 months
Outcomes	Clinical: PI (Silness and Löe), GI (Löe and Silness), PD; relative attachment levels (distance between the most apical portion of the stent and the base of the pocket), relative gingival margin levels (distance between the apical most part of the stent and the coronal limit of the gingival margin) Radiographic: radiographic measurements. Radio density Other: none
Notes	Sample size calculation: not reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement

Kaushick 2011 (*Continued*)

Random sequence generation (selection bias)	Unclear risk	Quote: "Patients were then randomized into the designated study groups" Comment: insufficient information provided regarding the method for random sequence generation
Allocation concealment (selection bias)	Unclear risk	No information provided regarding the method for allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No information provided regarding the blinding of outcome assessment
Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Khosropanah 2015

Methods	Trial design: randomised, split-mouth trial Location: Department of Periodontology, Shiraz Dental School, Iran Number of centres: 1 Recruitment period: not specified Source of funding: Vice-Chancellery of Research of Shiraz University, Iran Ethical approval: ethical approval (CT-90-5834) Number of surgeons: not reported
Participants	Inclusion criteria: moderate to advanced periodontitis, at least 2 intrabony defects with > 4 mm depth based on clinical examination, and at least 3 mm of keratinized tissue Exclusion criteria: systemic diseases or pregnancy, tobacco use, antibiotic intake in the past 3 months, taking anticoagulants for any reason, and history of periodontal therapy Age at baseline: 45 ± 10.7 years Gender: F 7/M 5 Smokers: excluded Number randomised (participants/teeth): 12/24 Number evaluated (participants/teeth):12/24
Interventions	Comparison: DFDBA + PRP versus DFDBA Test group: DFDBA + PRP (n = 12) Control group: DFDBA (n = 12) Surgical technique: OFD Follow-up duration: 6 months

Outcomes	Clinical: PI, BOP, PD, CAL, recession Radiographic: defect height, defect width and angle and hard tissue fills
Notes	Radiographs were taken with cone beam computed tomography (CBCT) Dropouts: reported, no dropouts

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "In this study, randomization was done using a 2-step coin tossing method. The first step of coin tossing was performed to choose the right side (tails) versus the left side (heads) and in the second step of coin tossing, the tails indicated controls and the heads indicated the test group. This way, location and type of intervention were both randomized" Comment: random sequence generation properly done
Allocation concealment (selection bias)	Low risk	Quote: "In this study, randomization was done using a 2-step coin tossing method. The first step of coin tossing was performed to choose the right side (tails) versus the left side (heads) and in the second step of coin tossing, the tails indicated controls and the heads indicated the test group. This way, location and type of intervention were both randomized" Comment: allocation concealment done correctly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "All measurements, including defect height, defect width and angle at baseline and 6 months later were recorded by an expert radiologist who was blinded to the type of surgical procedure" Comment: correct method for blinding of outcome assessment
Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients concluded the study

Khosropanah 2015 (*Continued*)

Selective reporting (reporting bias)	Low risk	Outcomes properly reported

Martande 2016

Methods	Trial design: randomised, parallel trial Location: Department of Periodontics, Government Dental College and Research Institute, Bangalore, India Number of centres: 1 Recruitment period: March 2013 to February 2014 Source of funding: not stated Ethical approval: Institutional Ethical Committee and Review Board of the Government Dental College and Research Institute, Bangalore, India Number of surgeons: 1
Participants	Inclusion criteria: patients with moderate-to-severe chronic periodontitis, based on the 1999 consensus classification of periodontal diseases; and presence of a 3-walled infrabony defect ≥ 3 mm deep in which depth was measured radiographically from the alveolar crest to the base of the defect on an intraoral periapical radiograph (IOPA) and the architecture of the 3-walled infrabony defect confirmed upon surgical exposure of the defect Exclusion criteria: patients with aggressive periodontitis; patients with systemic diseases affecting periodontal condition; those who had received periodontal therapy during the previous 6 months and/or are taking antibiotics for any chronic inflammatory conditions; smokers; pregnant and/or lactating females. Individuals with unacceptable oral hygiene (PI > 1.5) after re-evaluation of phase I therapy, and teeth with questionable to poor prognosis including: furcation defects; gingival recessions; carious teeth requiring extensive restorations; and non-vital teeth were also excluded. In addition, 1-walled and combined 1- and 2-walled defects confirmed upon surgical exposure were also excluded from the study Age at baseline: 30 to 50 years; mean age = 37.6 years Gender: F 48/M 48 (for all 3 groups) Smokers: excluded Teeth treated: 42 sites were from maxillary and mandibular single-rooted teeth, and the remaining 48 sites were from maxillary and mandibular multirooted teeth Number randomised (participants/teeth): 64/64 (96/96 for all 3 groups) Number evaluated (participants/teeth): 60/60 (90/90 for all 3 groups)
Intervention	Comparison: OFD + PRF versus OFD alone Test group: OFD + PRF (n = 30) Control group: OFD alone (n = 30) Surgical technique: full thickness mucoperiosteal flap with PRF in test site and full thickness mucoperiosteal flap alone in control site Follow-up duration: 9 months
Outcomes	Clinical: PI, modified sulcus bleeding index, PD, relative attachment level, gingival margin level Radiographic: radiographic bone filling Other: none

Notes	Sample size calculation: yes Radiographs were standardized using customized bite blocks and parallel angle technique and scanned with a scanner of 6400 dots per inch Comparability at baseline: yes, assessed Complications reported: no complications Dropouts: reported, 4 dropouts (6 for all 3 groups, 2 for each group) 3rd group data (OFD + PRF + 1.2% atorvastatin gel) not included in this review

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "Selected sites were divided randomly (computer-generated tables) into control and test groups (PRF or PRF + 1.2% ATV [atorvastatin])" Comment: random sequence generation likely to have been done properly
Allocation concealment (selection bias)	Unclear risk	Quote: "Patients were masked regarding their allocation to specific group and treatment" Comment: insufficient information is provided for the allocation concealment method
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment. For the same reason, blinding of the patient, even though it was done, does not influence the outcome of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Quote: "To avoid interoperative and inter-examiner bias, all surgical procedures were performed by a single operator (SSM) and all clinical and radiographic measurements were performed by a single examiner (ARP)" Comment: insufficient information is provided regarding blinding of the outcome assessment
Incomplete outcome data (attrition bias) All outcomes	Low risk	Only 6 patients did not complete the study (4 patients for the 2 groups considered in this review)
Selective reporting (reporting bias)	Low risk	All results properly reported

Naqvi 2017

Methods	Trial design: randomised, split-mouth, double-blinded clinical trial Location: Department of Periodontics and Oral Implantology, Santosh Dental College and Hospital, Santosh University, Ghaziabad, Uttar Pradesh, India Recruitment period: not reported Source of funding: not stated Ethical approval: the institutional ethical committee Number of surgeons: not stated
Participants	Inclusion criteria: presence of moderate to severe localised chronic periodontitis, having radiographic evidence of 1 or more vertical defects (2- or 3-walled) and probing pocket depth of 5 mm or more at the experimental site Exclusion criteria: patients with systemic diseases, on anticoagulants, those with habit of smoking and alcohol, with known history of allergy to graft material and who have undergone periodontal surgical treatment for chronic periodontitis within 12 months for the same defects. Pregnant and lactating females as well as patients on antibiotic therapy Age at baseline: 20 to 50 years Gender: F 3/M 7 Smokers: excluded Teeth treated: not stated Number randomised (participants/teeth): 10/20 Number evaluated (participants/teeth): 10/20
Interventions	Comparison: OFD + bioactive glass putty + PRF versus OFD + bioactive glass putty alone Test group: OFD + bioactive glass putty + PRF (n = 10) Control group: OFD + bioactive glass putty alone (n = 10) Surgical technique: full thickness mucoperiosteal flap with OFD + intrabony defects treated with bioactive glass putty alone in control group and additionally PRF was applied in test group Follow-up duration: 9 months
Outcomes	Clinical: PD, CAL Radiographic: radiographic bone filling Other: none
Notes	Sample size calculation: not reported Standardized intraoral periapical radiographs of the defects were taken using a paralleling technique Comparability at baseline: yes, assessed Complications reported: no complications Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement

Naqvi 2017 (*Continued*)

Random sequence generation (selection bias)	Low risk	Quote: "The intrabony defects were randomly assigned to either control group (bioactive glass putty alone) and test group (bioactive glass putty and PRF) by draw of chits" Comment: random sequence generation properly done
Allocation concealment (selection bias)	Unclear risk	Quote: "The intrabony defects were randomly assigned to either control group (bioactive glass putty alone) and test group (bioactive glass putty and PRF) by draw of chits" Comment: allocation concealment likely to have been done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment. For the same reason, blinding of the patient, even though it was done, does not influence the outcome of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "Neither the patients nor the investigator was aware of the group assignment, thereby assuring double blindness" Comment: blinding of outcome assessment done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients concluded the study
Selective reporting (reporting bias)	Low risk	All results properly reported

Okuda 2005

Methods	Trial design: randomised, parallel trial Location: Niigata University Medical and Dental Hospital, Japan Number of centres: 1 Recruitment period: not stated Ethical approval: ethical committee for human subject use at Niigata University Medical and Dental Hospital in accordance with the Helsinki Declaration of 1975 as revised in 1983 Number of surgeons: 3
Participants	Inclusion criteria: individuals who were non-smoking, free of systemic complications, and without a history of allergies; had not used antibiotics within the previous 6 months prior to treatment; had not been treated for periodontitis during the previous 2 years; had 1 intrabony defect

Okuda 2005 (*Continued*)

	with PD) ≥ 6 mm, CAL loss ≥ 6 mm, and an osseous defect depth estimated from radiographic evaluation as ≥ 3 mm; and had at least 2 mm of keratinized gingiva on the facial aspect of the selected tooth Exclusion criteria: failing to meet inclusion criteria Age at baseline: mean age = 55.5 ± 8.2 years Gender: F 49/M 21 Smokers: excluded Number randomised (participants/teeth): 70/70 Number evaluated (participants/teeth): 70/70
Interventions	Comparison: PRP + HA versus saline + HA Test group: PRP + HA (n = 35/35) Control group: saline + HA (n = 35/35) Surgical technique: OFD + intrabony defects were treated with PRP + HA on test group sites and saline + HA on control group Follow-up duration: 1 year
Outcomes	Clinical: PD, CAL, GR, vertical relative attachment gain Radiographic: intrabony defect depth fill Other: none
Notes	Sample size calculation not reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Patients who met all criteria for entry into the surgical phase of the study were then randomized by a coin toss to the test (PRP + HA) or control (saline + HA) study groups" Comment: random sequence generation done properly
Allocation concealment (selection bias)	Low risk	Quote: "Patients who met all criteria for entry into the surgical phase of the study were then randomized by a coin toss to the test (PRP + HA) or control (saline + HA) study groups" Comment: allocation concealment likely to have been done properly

Okuda 2005 (*Continued*)

Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "All radiographs were evaluated by a single examiner (author KT) who was masked to the treatment group to which a patient was assigned" Comment: blinding of outcome assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Ozdemir 2012

Methods	Trial design: randomized, split-mouth trial Location: Department of Periodontology, Faculty of Dentistry, Gazi University, Turkey Number of centres: 1 Recruitment period: not reported Source of funding: not reported Ethical approval: Ethical Board of Gazi University School of Medicine, Turkey Number of surgeons: 1
Participants	Inclusion criteria: patients with no periodontal treatment and consumption of medicine 6 months before the study; a good level of oral hygiene (PI < 1) at re-evaluation sessions; no orthodontic treatment; compliance with the maintenance program; the involved teeth were vital and had no mobility, occlusal trauma, endodontic treatment, or prosthetic restoration; at least 2 similar 3-walled intrabony defects with 6 mm PD at interproximal region, which was supported by periapical radiographs; intrabony defects that were not on the same tooth or at the same interproximal region and were localized to the interproximal region of mandibular and maxillary anterior and premolar teeth and mesial root of the first mandibular molars; and keratinized gingival width of at least 2 to 3 mm in the defect region Exclusion criteria: pregnant and/or lactating women; smokers; abnormal platelet counts disclosed by a complete blood count test performed within 2 weeks before surgery; and participation in other dental clinical trials at the time of this trial Age at baseline: mean = 48.9 + 6.6 years Gender: F 5/M 9 Smokers: excluded Teeth treated: not reported Number randomised (participants/teeth): 14/28 Number evaluated (participants/teeth): 14/28

Ozdemir 2012 (*Continued*)

Interventions	Comparison: PRP/ß-TCP versus ß-TCP alone Test group: PRP/ß-TCP (n = 14) Control group: ß-TCP alone (n = 14) Surgical technique: OFD intrabony defects were treated with ß-TCP alone in control group and additionally PRP was applied to the test group Follow-up duration: 6 months
Outcomes	Clinical: PI, GI, PPD, CAL, BOP, and GR measured between CEJ and gingival margin Radiographic: radiographic intrabony defect depth Other: none
Notes	Sample size calculation: not reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "ß-TCP (n = 14) and PRP/ß-TCP groups (n = 14) were selected randomly by the toss of a coin and each patient had 1 pair of both ß-TCP and PRP/ß-TCP group defects" Comment: random sequence generation done properly
Allocation concealment (selection bias)	Low risk	Quote: "ß-TCP (n = 14) and PRP/ß-TCP groups (n = 14) were selected randomly by the toss of a coin and each patient had 1 pair of both ß-TCP and PRP/ß-TCP group defects" Comment: allocation concealment likely to have been done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Insufficient information is provided regarding the blinding of outcomes
Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients concluded the study

Ozdemir 2012 (*Continued*)

Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Panda 2016

Methods	Trial design: randomised, split-mouth trial Location: Department of Periodontics, Saveetha Dental College and Hospitals Saveetha University, Tamil Nadu, India Number of centres: 1 Recruitment period: March to December 2012 Source of funding: no funding Ethical approval: Institutional Human Ethics Committee of Saveetha Dental College and Hospitals, Chennai, India Number of surgeons: 1
Participants	Inclusion criteria: presence of interproximal intrabony defects ≥ 2 mm (distance between alveolar crest and base of the defect evaluated on intraoral periapical radiographs) along with an interproximal PD ≥ 5 mm following phase I therapy (SRP) Exclusion criteria: presence of past systemic illnesses known to affect the outcomes of periodontal therapy; immunocompromised status; tobacco use in any form; current medications that may interfere with periodontal therapy; haematologic disorders, or insufficient platelet count (< 200,000/mm^3) and poor oral hygiene after the re-evaluation of phase I therapy (PI ≥ 1.5); pregnancy and lactation; teeth with furcation defects; mobility of at least Grade II, and carious lesions needing restorations; 2- and 1-wall defects and interdental craters Age at baseline: mean = 38.12 ± 2.06 years Gender: F 8/M 10 Smokers: excluded Teeth treated: not stated Number randomised (participants/teeth): 18/36 Number evaluated (participants/teeth): 16/32
Interventions	Comparison: GTR + PRF versus GTR alone Test group: GTR + PRF (n = 16) Control group: GTR alone (n = 16) Surgical technique: GTR and PRF in test site and GTR alone in control site Follow-up duration: 9 months
Outcomes	Clinical: PI, modified sulcus bleeding index, PPD, CAL, and gingival marginal level Radiographic: radiographic bone filling Other: none
Notes	Sample size calculation: yes PD and the CAL were measured using customized acrylic stents with grooves to ensure a reproducible placement of the probe Radiographs were made using customized bite blocks and long cone paralleling angle technique Comparability at baseline: yes, assessed Complications reported: no complications

Panda 2016 (*Continued*)

	Dropouts: 2 dropouts	
Risk of bias		
Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "A simple randomization (coin toss) scheme was used by 1 of the authors (MDF) to assign the patients with an allocation ratio of 1:1 into 2 study groups: PRF + GTR (18 patients, test) and GTR alone (18 patients, control)" Comment: random sequence generation properly done
Allocation concealment (selection bias)	Low risk	Quote: "Allocations were concealed by using number-labelled opaque envelopes containing the name of the assigned intervention" Comment: allocation concealment properly done
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment. For the same reason, blinding of the patient, even though it was done, does not influence the outcome of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "Since only 1 examiner (SM) measured the clinical and radiographic parameters in the study, intra-examiner reliability assessment was done to validate the ability of the examiner to constantly replicate the quantitative outcome measurements of the parameters used." "The examiner was blinded to treatment" (information provided by the author) Comment: blinding of outcome assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	Only 2 patients did not conclude the study
Selective reporting (reporting bias)	Low risk	All results properly reported

Patel 2017

Methods	Trial design: randomised, split-mouth trial Location: Department of Periodontology, Jagadguru Sri Shivarathreshwara (JSS) Dental College and Hospital, Mysore, India Number of centres: 1 Recruitment period: from October 2010 to (not stated) Source of funding: not stated Ethical approval: Institutional Review Board of the JSS University governing the use of human patients in clinical experimentation Number of surgeons: 1
Participants	Inclusion criteria: the presence of 2 similar mandibular interproximal, 3-walled intrabony defects with PD ≥ 6 mm and radiographic evidence of ≥ 3 mm distance between alveolar crest and base of the defect. PI and GI achieved after initial therapy had to be < 1. Only vital teeth were included in the study Exclusion criteria: individuals with underlying systemic illnesses and those taking any drug known to affect the outcome of periodontal therapy and/or drugs effecting platelets; smokers, immunocompromised individuals; and pregnant or lactating individuals. Defect sites which were found to be 1-walled on flap reflection were also excluded Age at baseline: mean 44 ± 9 years Gender: F 9/M 4 Smokers: excluded Teeth treated: lower single-rooted and multirooted teeth Number randomised (participants/teeth): 13/26 Number evaluated (participants/teeth): 13/26
Interventions	Comparison: OFD + PRF versus OFD alone Test group: OFD + PRF (n = 13) Control group: OFD alone (n = 13) Surgical technique: full thickness mucoperiosteal flap and debridement + PRF in test site and full thickness mucoperiosteal flap and debridement alone in control site Follow-up duration: 12 months
Outcomes	Clinical: PI, GI, reduction in PD, gain in CAL Radiographic: radiographic bone filling Other: wound healing index
Notes	Sample size calculation: yes PD and CAL were measured by a manual periodontal probe using customized acrylic stents Radiographic evaluation was done using digital radiography/radiovisiography with the long cone parallel technique Comparability at baseline: yes, assessed Complications reported: no complications Dropouts: no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement

Patel 2017 (*Continued*)

Random sequence generation (selection bias)	Low risk	Quote: "Randomization of the selected sites (i.e. 2 similar interproximal sites in each indivi ual) was done by toss of a coin by the study herapist (GP)" C mi rat om sequence generation done roperly
Allocation concealment (selection bias)	Low risk	Quote: "Randomization of the selected ites (i.e. 2 similar interproximal sites in eac ndividual) was done by toss of a coin y the study therapist (GP)" omment: allocation concealment likely to nave been done properly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "It was a double-masked, single-center, prospective study of 12 months duration" Comment: blinding of the outcome assessment likely to have been done properly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients concluded the study
Selective reporting (reporting bias)	Lo ri	All results properly reported

Piemontese 2008

Methods	Trial design: randomised, parallel trial Location: Polytechnic University of Marche, Ancona, Italy Number of centres: 1 Recruitment period: 2002 to 2003 Source of funding: study supported by the Polytechnic University of Marche, Ancona, Italy Ethical approval: Committee for the Protection of Human Subjects at the Polytechnic University of Marche Number of surgeons: 1
Participants	Inclusion criteria: individuals who were non-smoking, free of systemic complications, and without a history of allergies; had not used antibiotics within the previous 6 months prior to treatment; had not had abnormal platelet counts disclosed by a complete blood cell count performed within 1 month prior to surgery; had not been treated for periodontitis during the previous 2 years; had radiographic and clinical evidence of 1 defect with PD > 6 mm, CAL > 6 mm, osseous defect depth estimated from radiographic evaluation as > 3 mm, and 2 or 3 osseous walls; had no intrabony defects extending into

Piemontese 2008 (*Continued*)

	a furcation area; and had no teeth presenting furcation involvement Exclusion criteria: patients failing to meet inclusion criteria Age at baseline: 47 to 72 years Gender: F 29/M 31 Smokers: excluded Teeth treated: maxillary and mandibular incisors and premolars and maxillary molar Number randomised (participants/teeth): 60/60 Number evaluated (participants/teeth): 60/60
Interventions	Comparison: PRP + DFDBA versus DFDBA + saline Test group: PRP + DFDBA (n = 30/30) Control group: DFDBA + saline (n = 30/30) Surgical technique: OFD intrabony defects were treated with PRP/DFDBA in test group and saline/DFDBA in control group Follow-up duration: 1 year
Outcomes	Clinical: PI, GI, BOP, PD, CAL, REC Radiographic: CEJ-BD , AC-BD, CEJ-AC Other: none
Notes	Sample size calculation: not reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, no dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Randomization was performed by the toss of a coin immediately following defect debridement" Comment: random sequence generation done correctly
Allocation concealment (selection bias)	Low risk	Quote: "Randomization was performed by the toss of a coin immediately following defect debridement" Comment: allocation concealment likely to have been done correctly
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quotes: "The study was designed as a randomized, double-masked, clinical trial comparing the periodontal outcomes ..."

Piemontese 2008 (*Continued*)

		and "On the day of the surgical procedure, baseline clinical measurements were recorded by the same calibrated examiner (SDA) masked to the treatment" Comment: blinding of outcome assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients concluded the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Pradeep 2015

Methods	Trial design: randomised, longitudinal, triple-masked, parallel trial Location: Department of Periodontics, Government Dental College and Research Institute, Bangalore, India Number of centres: 1 Recruitment period: November 2013 to July 2014 Source of funding: not stated Ethical approval: Institutional Ethical Committee and Review Board of the Government Dental College and Research Institute, Bangalore, India Number of surgeons: 1
Participants	Inclusion criteria: presence of intrabony defect ≥ 3 mm deep (distance between alveolar crest and base of the defect on an intraoral periapical radiograph (IOPA)) along with an interproximal PD ≥ 5 mm after phase I therapy (scaling and root planing) in asymptomatic maxillary/mandibular molar teeth Exclusion criteria: aggressive periodontitis patients; patients with systemic conditions known to affect the periodontal status; medications known to affect the outcomes of periodontal therapy; haematological disorders and insufficient platelet count (< 200, 000/mm^3); pregnancy/lactation; smoking and tobacco use in any form; and immunocompromised individuals. Those having unacceptable oral hygiene (PI > 1.5) after re-evaluation of phase I therapy were also excluded. In addition, teeth with furcation defects, non-vital teeth, carious teeth warranting restorations and mobility of at least grade II were also excluded Age at baseline: mean = 41 years Gender: F 68/M 68 (for all 4 groups) Smokers: excluded Teeth treated: maxillary and mandibular molar Number randomised (participants/teeth): 64/64 (126/126 for all 4 groups; 136 eligible but 10 excluded at time of surgery) Number evaluated (participants/teeth): 60/60 (120/120 for all 4 groups)
Interventions	Comparison: OFD alone versus OFD + PRF Group 1: OFD alone (n = 30) Group 2: OFD + PRF (n = 30) Surgical technique: in group 1, only OFD was done, without addition of any regenerative

Pradeep 2015 (*Continued*)

	material into the bone defect; in group 2, PRF of the required size was filled into the intrabony defect after OFD Follow-up duration: 9 months
Outcomes	Clinical: site specific PI, modified sulcus bleeding index, relative attachment level, gingival marginal level Radiographic: radiographic intrabony defect depth Other: none
Notes	Sample size calculation: reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, 4 dropouts (6 in all groups) 3rd and 4th group data (OFD + 1% metformin and OFD + PRF + 1% metformin) not included in this review

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "These patients were divided randomly (computer generated tables) into 4 groups" Comment: random sequence generation properly done
Allocation concealment (selection bias)	Unclear risk	Insufficient information provided for allocation concealment method
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment. For the same reason, blinding of the patient, even though it was done, does not influence the outcome of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quotes: "This was a randomized, single-centre, longitudinal, triple-masked (investigators, individuals and statistician), parallel arm design study" and "One operator (KN) performed all the surgeries, whereas another operator (ARP) performed all the clinical and radiographic measurements without knowledge of the groups" Comment: blinding of outcome assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	4 dropouts (6 for all 4 groups)

Pradeep 2015 (*Continued*)

Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Pradeep 2016

Methods	Trial design: randomised, parallel trial Location: Periodontics Clinic, GDCRI, Bengaluru, Karnataka, India Number of centres: 1 Recruitment period: January to October 2015 Source of funding: not stated Ethical approval: ethical approval from the Institutional Ethical Committee and Review Board of Government Dental College and Research Institute (GDCRI), Bengaluru, Karnataka, India Number of surgeons: 1
Participants	Inclusion criteria: systemically healthy with diagnosis of chronic periodontitis; PD ≥ 5 mm; CAL ≥ 3 mm; and 2 or 3-walled intrabony defect on at least 1 mandibular molar; vertical bone loss ≥ 3 mm on intraoral periapical radiographs and no antibiotic or periodontal therapy in 6 months before study Exclusion criteria: statin allergy; statin therapy; any systemic condition or medication altering periodontal condition; an immunocompromised state; haematologic disorders; insufficient platelet count (< 200,000/mm^3); aggressive periodontitis; substance/tobacco abuse; lactating or pregnant females. Age at baseline: mean age = 35 years Gender: F/M 45 (for all 3 groups) Smokers: excluded Teeth treated: 42 sites were from maxillary and mandibular single-rooted teeth, and the remaining 48 sites were from maxillary and mandibular multirooted teeth Number randomised (participants/teeth): 60/60 (90/90 for all 3 groups) Number evaluated (participants/teeth): 60/60 (90/90 for all 3 groups)
Interventions	Comparison: OFD + PRF versus OFD alone Test group: OFD + PRF (n = 30/30) Control group: OFD alone (n = 30/30) Surgical technique: full thickness mucoperiosteal flap with PRF in test site and full thickness mucoperiosteal flap alone in control site Follow-up duration: 9 months
Outcomes	Clinical: PI, modified sulcus bleeding index, PD, CAL Radiographic: radiographic bone filling Other: none
Notes	Sample size calculation: yes Reproducible parallel-angle radiographs of concerned sites using customized bite blocks Comparability at baseline: yes, assessed Complications reported: yes, no complications Dropouts: no dropouts 3rd test group data (OFD + PRF + 1.2% rosuvastatin gel) not included in this review

Pradeep 2016 (*Continued*)

Risk of bias		
Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "...power calculations-based (90% confidence at $P \leq 0.05$) enrolment, ...computer-assisted random allocation of the 90 patients was done into 3 treatment groups" Comment: random sequence generation done correctly
Allocation concealment (selection bias)	Unclear risk	Insufficient information provided for allocation concealment method
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator given the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "90 patients ..... were enrolled for this placebo-controlled, triple-masked, single-center randomized controlled clinical trial from January 2015 to October 2015 (9-month study)" Comment: blinding of outcome assessment done correctly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients completed the study
Selective reporting (reporting bias)	Low risk	All results properly reported

Ravi 2017

Methods	Trial design: randomised, split-mouth trial Location: Department of Periodontology, Saveetha University, India Number of centres: 1 Recruitment period: September 2015 to September 2016 Source of funding: no funding Ethical approval: Institution Human Ethics Committee Number of surgeons: 1
Participants	Inclusion criteria: presence of generalized chronic periodontitis (on the basis of the 1999 consensus classification of periodontal diseases); presence of bilateral intrabony defect $\geq$ 3 mm deep (distance between alveolar bone crest and base of defect on intraoral periapical radiograph); presence of interproximal PD $\geq$ 5 mm after phase I periodontal therapy (scaling and root planing); systemically healthy condition Exclusion criteria: history of periodontal surgical treatment within the last 6 months,

Ravi 2017 (*Continued*)

	smokers, pregnant or lactating women Age at baseline: mean age = 43.26 ± 9.45 years Gender: F 9/M 5 Smokers: excluded Teeth treated: premolars and molars Number randomised (participants/teeth): 14/[illegible] Number evaluated (participants/teeth): 12/38
Interventions	Comparison: GTR + PRGF versus GTR [illegible]ne Test group: GTR + PRGF (n = 19 sit[illegible]) Control group : GTR alone (n= [illegible] sites) Surgical technique: GTR and P[illegible]GF in test s[illegible]e and GTR alone in control site Follow-up duration: 6 months
Outcomes	Clinical: GI, PD, CAL Radiographic: radiographic b[illegible]e filling Other: none
Notes	Sample size calculation: [illegible] Customized pur[illegible]y bi[illegible] blocks were made for each patient to standardize positioning of the sensor and [illegible]gl[illegible] with which radiographs were taken Comparabil[illegible]ty at [illegible]eli[illegible]e: yes, assessed Complic[illegible]ions [illegible]ported: no complications Dropouts: [illegible]patients, 4 sites

Risk of bias

Bias	**[illegible]nors' judgement**	**Support for judgement**
Random sequence generation [illegible]ion bias)	[illegible]w risk	Quote: "Selected sites were randomly assigned to 1 of the following groups: 1) PRGF plus GTR or 2) GTR alone by using the coin toss method for each patient (NJ) " Comment: random sequence generation properly done
Allocation conceal[illegible]ent (sel[illegible]tion bias)	Low risk	Quote: "Selected sites were randomly assigned to 1 of the following groups: 1) PRGF plus GTR or 2) GTR alone by using the coin toss method for each patient (NJ) " Comment: allocation concealment likely to have been done properly
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Quote: "The present study was a split-mouth randomized control trial in which the operator and assessor were masked" It is stated that the operator was blinded

Ravi 2017 (*Continued*)

		but no further information is provided on the exact method in which it was done
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quotes: "The present study was a split-mouth randomized control trial in which the operator and assessor were masked" and "The examiner, however, was not aware, in any of the cases, of the type of treatment rendered (SV and SM)" Comment: blinding of outcome assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	Only 2 patients failed to complete the study
Selective reporting (reporting bias)	Low risk	All results properly reported

Rosamma Joseph 2012

Methods	Trial design: randomised split-mouth trial Location: Department of Periodontics, Government Dental College, Kozhikode, Kerala, India Number of centres: 1 Recruitment period: September 2009 to October 2010 Source of funding: not stated Ethical approval: Institutional Ethics Committee, Government Dental College, Kozhikode, in accordance with the Helsinki Declaration of 1975 as revised in 2000 Number of surgeons: 1
Participants	Inclusion criteria: patients had paired, contralateral interproximal infrabony defect with a probing PD > 6 mm, CAL loss > 5 mm, and an osseous defect depth estimated from radiographic evaluation as > 4 mm; were systemically healthy without a history of allergies; and had at least 2 mm of keratinized gingiva on the facial aspect of the selected tooth Exclusion criteria: haematological or immunological disorders; pregnancy or lactation; smoking or the use of other tobacco products; those taking drugs known to interfere with wound healing; had used antibiotics within the previous 1 year; had been treated for periodontitis during the previous 2 years; those with unacceptable oral hygiene (PI) after the re-evaluation of phase I therapy; were not willing to sign an informed consent Age at baseline: mean = 29.47 + 7.65 years (range 17 to 44 years) Gender: F 9/M 6 Smokers: excluded Teeth treated: not reported Number randomised (participants/teeth): 15/30 Number evaluated (participants/teeth): 15/30
Interventions	Comparison: OFD + PRFm versus OFD alone Test group: OFD + PRFm (n = 15/15) Control group: OFD alone (n = 15/15)

	Surgical technique: test group was treated by placement of platelet-rich fibrin matrix following OFD and control group was treated by OFD alone Follow-up duration: 1 year
Outcomes	Clinical: PD, recession/enlargement, CAL, PI, modified G Radiographic: the vertical dimension between th projection the bone crest on the root surface (BCP) and the most coronal level along the ot surfa where the periodontal ligament space was considered to have a nor al dth (BoBD-base of bone defect) was measured and designated as infrabony def depth (D = BCP - BoBD). The distance from the crest of remaining alveolar b e to C J was also recorded (CEJ-BC) Other: a visual analogue scale (V 1) was d d to assess the patient experience with the 2 treatment modalities. Anothe isual analo e scale (VAS2) was designed and used to assess the initial soft tissue heali
Notes	Sample size calculation: rep rted Radiographs were taken with bite block for ensuring reproducibility Comparability at baseli s a essed Complications reported: y Dropouts: reported, no d ts

Risk of bias

Bias	Authors' dgei nt	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Either right sided or maxillary defects were operated first and whether the site belonged to experimental or control group was determined by a simple lottery method by the toss of a coin" Comment: random sequence generation done correctly
Allocation concealment (selection b)	Low risk	Quote: "The sites were divided into experimental and control groups at the time of periodontal surgery. Either right sided or maxillary defects were operated first and whether the site belonged to experimental or control group was determined by a simple lottery method by the toss of a coin" Comment: allocation concealment likely to have been done correctly
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	It is stated that the operator was blinded but no further information is provided on the exact method in which it was done
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "All radiographs were evaluated by a single examiner (RJ) who was masked to the treatment group to which a patient was

Rosamma Joseph 2012 (*Continued*)

		assigned and also to whether the radiograph was taken at baseline or re-evaluation" Comment: blinding of outcome assessment done correctly
Incomplete outcome data (attrition bias) All outcomes	Low risk	All enrolled patients completed the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Sezgin 2017

Methods	Trial design: randomised, split-mouth trial Location: Department of Periodontology, Gazi University, Turkey Number of centres: 1 Recruitment period: not reported Source of funding: Gazi University Research Grant, Turkey Ethical approval: approved by the ethics board at the Faculty of Dentistry, Gazi University, Turkey Number of surgeons: 2
Participants	Inclusion criteria: no systemic diseases; a good level of oral hygiene (PI < 0.15); presence of 2 paired, 2- or 3-walled intrabony defects with PD ≥ 6 mm and an intrabony component of ≥ 3 mm, as detected on radiographs; no intrabony defects extending into the furcation area; tooth mobility ≤ 1; tooth and adjoining teeth testing vital and without symptoms or signs of endodontic involvement; and tooth and adjoining teeth free of caries or inadequate restorations Exclusion criteria: patients with compromised immune systems; pregnant and/or lactating women; patients taking any drug known to affect the periodontal status or the coagulation system; and smokers Age at baseline: 38 to 61 years Gender: F 7/M 8 Smokers: excluded Teeth treated: all Number randomised (participants/teeth): 21/42 Number evaluated (participants/teeth): 15/30
Interventions	Comparison: ABBM + PRF versus ABBM alone Test group: ABBM + PRF (n = 15) Control group: ABBM alone (n = 15) Surgical technique: OFD Follow-up duration: 6 months
Outcomes	Clinical: PI, GI, PD, CAL and GR Radiographic: vertical bone loss, depth of intrabony defect, radiographic defect angle
Notes	Sample size calculation: reported Radiographs were taken using long cone parallel and direct digital radiography

Sezgin 2017 (*Continued*)

	Comparability at baseline: yes, assessed Complications reported: no complications Dropouts: reported, 1 dropout (5 patients excluded because the defects did not meet the study criteria at surgery)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "The selected sites were randomly (coin toss) divided into control (ABBM alone) and test (ABBM-PRF) groups" Comment: correct method of random sequence generation
Allocation concealment (selection bias)	Unclear risk	Insufficient information provided for the method of allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "One examiner other than the surgeons performed all clinical measurements, and another examiner performed all radiographical measurements. Both examiners were blinded to the study groups" Comment: blinding of outcome assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	Only 1 patient failed to complete the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Sharma 2011

Methods	Trial design: randomised, parallel trial Location: Department of Periodontics, Government Dental College and Research Institute, Bangalore, India Number of centres: 1 Recruitment period: June 2009 to March 2010 Source of funding: nil Ethical approval: Institutional Ethical Committee and Review Board, Government Dental College and Research Institute, Bangalore, India Number of surgeons: 1

Sharma 2011 (*Continued*)

Participants	Inclusion criteria: presence of 3-walled intrabony defects > 3 mm deep (the distance between the alveolar crest and base of the defect on an intraoral periapical radiograph (IOPA)) along with an interproximal PD > 5 mm after phase 1 therapy (scaling and root planing) in an asymptomatic tooth Exclusion criteria: patients with aggressive periodontitis with known systemic illness and taking any medications known to affect the outcomes of periodontal therapy; an insufficient platelet count (< 200,000/mm^3); pregnancy or lactation; use of any form of tobacco; patients who had unacceptable oral hygiene (PI > 1.5) after the re-evaluation of phase 1 therapy; teeth with furcation defects, non-vital teeth or teeth with mobility > grade II Age at baseline: 30 to 50 years; mean = 35.5 + 6.45 years Gender: F 18/M 24 Smokers: excluded Teeth treated: 17 of the 56 sites were from upper and lower single-rooted teeth, and the remaining 39 sites were from upper and lower multirooted teeth Number randomised (participants/teeth): 42/69 Number evaluated (participants/teeth): 35/56
Interventions	Comparison: PRP/OFD versus OFD alone Test group: PRP/OFD (n = 18/28) Control group: OFD alone (n = 17/28) Surgical technique: intrabony defects treated with OFD alone in control group and additional PRP was added in test group Follow-up duration: 9 months
Outcomes	Clinical: site specific PI, modified sulcus bleeding index, PD, periodontal attachment level, gingival margin level Radiographic: radiographic intrabony defect depth Other: none
Notes	Sample size calculation: reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, reasons given, 7 patients, 13 sites did not return for follow-up examinations

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "The selected sites were divided randomly (by using a coin-toss method) into control and test groups" Comment: correct method for random sequence generation
Allocation concealment (selection bias)	Unclear risk	Insufficient information provided for allocation concealment method

Sharma 2011 (*Continued*)

Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "One operator (AS) performed all surgeries whereas another operator (ARP) performed all clinical and radiographic measurements without knowledge of the groups" Comment: blinding of outcome assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	7 patients out of 42 failed to complete the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Shukla 2016

Methods	Trial design: randomised split-mouth trial Location: outpatient service on a teaching dental institute in North India (no further details given) Number of centres: 1 Recruitment period: not specified Source of funding: not stated Ethical approval: Institutional Ethical Committee and Review Board of the Government Dental College and Research Institute, India Number of surgeons: 1
Participants	Inclusion criteria: presence of intrabony defects > 3 mm deep (distance between alveolar crest and base of the defect on intraoral periapical radiograph (IOPA)) and an interproximal PD > 5 mm Exclusion criteria: known systemic illness; taking any medications known to affect the outcomes of periodontal therapy; pregnancy/lactation; use of any form of tobacco; allergy to calcium phosphosilicate putty Age at baseline: mean = 40 + 10.5 years Gender: F 7/M 13 Smokers: excluded Teeth treated: not stated Number randomised (participants/teeth): 20/40 Number evaluated (participants/teeth): 20/40
Interventions	Comparison: OFD + calcium phosphosilicate (CPS) + PRF versus OFD + CPS alone Test group: OFD + CPS + PRF (n = 20) Control group: OFD + CPS alone (n = 20) Surgical technique: full thickness mucoperiosteal flap with PRF and CPS in test site and full thickness mucoperiosteal flap with CPS alone in control site Follow-up duration: 9 months

Shukla 2016 (*Continued*)

Outcomes	Clinical: PI, PD, CAL, GI Radiographic: radiographic bone filling Other: none	
Notes	Sample size calculation: yes Comparability at baseline: yes, assessed Complications reported: no complications Dropouts: no dropouts	
Risk of bias		
Bias	**Authors' judgement**	**Support for judgement**
Random sequence generation (selection bias)	Low risk	Quote: "randomization was performed using a computer-generated randomization list" Comment: correct method for random sequence generation
Allocation concealment (selection bias)	Unclear risk	Insufficient information provided on the allocation concealment method
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "All the evaluations were performed by an independent trained observer not involved in the study" Comment: blinding of outcome assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients completed the study
Selective reporting (reporting bias)	Low risk	All results properly reported

Thorat 2011

Methods	Trial design: randomised, parallel trial Location: Department of Periodontics, Government Dental College and Research Institute, Bangalore, India Number of centres: 1 Recruitment period: April 2009 to January 2010 Source of funding: self-funded Ethical approval: Institutional Review Board, Ind Number of surgeons: 1
Participants	Inclusion criteria: presence of interproximal trabony defects > 3 mm deep (distance between alveolar crest and base of the defect on intraoral periapical radiograph (IOPA)) along with an interproximal PD > 5 mm following phase I therapy (scaling and root planing in vital, asymptomatic first and second mandibular molars without furcation involvement Exclusion criteria: patients with present or past systemic illness that were known to affect the outcomes of periodontal therapy; insufficient platelet count (< 200,000/mm ^{3}); immunocompromised patients; pregnancy/lactation; smoking (any other tobacco products); patients taking medications that may interfere with wound healing; those allergic to other medication and having unacceptable oral hygiene (PI > 3) after the re-evaluation of phase I therapy Age at baseline: 25 to 45 years; mean = 31.1 + 2.06 years Gender: F 18/M Smokers: excluded Teeth treated: first and second mandibular molars Number randomised (participants/teeth): 40/40 Number evaluated (participants/teeth): 32/32
Interventions	Comparison: PRF + OFD versus OFD alone Test group: PRF + OFD (n = 16/16) Control group: OFD alone (n = 16/16) Surgical technique: intrabony defects treated with OFD alone in control group and additionally PRF was added in test group Follow-up duration: 9 months
Outcomes	Clinical: PI, sulcus bleeding index, PD, CAL, and gingival marginal level Radiographic: bone defect fill Other: none
Notes	Sample size calculation: reported Radiographs were taken with a bite block for ensuring reproducibility Comparability at baseline: yes, assessed Complications reported: yes Dropouts: reported, reasons given, 8 dropouts

Risk of bias

Bias	Authors' judgement	Support for judgement

Thorat 2011 (*Continued*)

Random sequence generation (selection bias)	Low risk	Quote: "The selected sites were divided randomly (coin toss) into the control and test groups" Comment: correct method for random sequence generation
Allocation concealment (selection bias)	Unclear risk	Insufficient information provided regarding the method of allocation concealment
Blinding of participants and personnel (performance bias) All outcomes	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quotes: "A review of all the radiographs was performed in a single reference center by a blind evaluator" and "An examiner (ARP) other than the operator performed all clinical measurements without knowledge of the treatment groups" Comment: blinding of outcome assessment properly done
Incomplete outcome data (attrition bias) All outcomes	Low risk	8 out of 40 patients failed to complete the study
Selective reporting (reporting bias)	Low risk	All outcomes properly reported

Thorat 2017

Methods	Trial design: randomised, split-mouth trial Location: Department of Periodontics, Government Dental College and Research Institute, Bangalore, India Number of centres: 1 Recruitment period: not stated Source of funding: not stated Ethical approval: Institutional Ethical Committee and registered with Clinical Trials Registry India REF/12/006069) Number of surgeons: 1
Participants	Inclusion criteria: localized aggressive periodontitis; presence of at least 2 contralateral interproximal intrabony defects; intrabony defect $\geq$ 3 mm (vertical distance between alveolar crest and base of the defect on standardized intraoral periapical radiographs) with corresponding PD $\geq$ 5 mm following phase I therapy; individual PI score $\leq$ 2; asymptomatic first/second molars without furcation involvement Exclusion criteria: present or past systemic illness known to affect the outcomes of periodontal therapy; insufficient platelet counts (< 200,000/mm^3); immunocompromised status; pregnancy/lactation; taking medications that might interfere with wound healing; and tobacco habits

	Age at baseline: mean = 25 ± 1.5 years Gender: F 10/M 8 (3 did not receive surgery, their gender not specified) Smokers: excluded Teeth treated: first/second molars Number randomised (participants/teeth): 15/30 Number evaluated (participants/teeth): 15/30
Interventions	Comparison: OFD + PRF versus OFD alon Test group: OFD + PRF (n = 15) Control group: OFD alone (n = 15) Surgical technique: Kirkland mo ined flap peration Follow-up duration: 12 months
Outcomes	Clinical: gain in CAL, reduction in change in gingival margin level Radiographic: radiographic bone filling Other: none
Notes	Sample size calculation: ye PD and the CAL were me d by a manual periodontal probe using customized acrylic stents Radiographic e alu tion was done on intraoral periapical radiographs using long cone paralleling ngle chni ue and individualized bite blocks with a positioning device Compara ility baseline: yes, assessed Complicati s repo ted: no complications Drop : no dropouts

Risk of bias

Bias	**Authors' judgement**	**Support for judgement**
Random sequence generatio (selec on bias)	Low risk	Quote: "Sites were assigned using a computer-generated randomization process" Comment: correct method for random sequence generation
Allocation concealment (se tion bias)	Unclear risk	Insufficient information regarding allocation concealment method was provided
Blinding o partici nts nd personnel (perform ce bia All outco es	High risk	Impossible to blind the operator due to the surgical nature of the treatment
Blinding of outcom e assessment (detection bias) All outcomes	Low risk	Quote: "The preoperative and postoperative clinical parameters were checked by a single blinded examiner. Another blinded and calibrated examiner (radiologist) recorded the radiographic parameters" Comment: blinding of outcome assessment properly done

Thorat 2017 (*Continued*)

Incomplete outcome data (attrition bias) All outcomes	Low risk	All patients completed the study
Selective reporting (reporting bias)	Low risk	All results properly reported

ABBM = anorganic bovine bone mineral; AC = alveolar crest; APC = autologous platelet concentrate; BD = base of the defect; BDX = bovine derived xenograft; BG = bone graft; BOP = bleeding on probing; BPBM = bovine porous bone mineral; ß-TCP = beta-tricalcium phosphate; CAF = coronally advanced flap; CAL = clinical attachment level; CD = crest of the defect; CEJ = cemento-enamel junction; DBM = demineralized bone matrix; DFDBA = demineralized freeze-dried bone allograft; EMD = enamel matrix derivative; F = female; GI = gingival index; GR = gingival recession; GTR = guided tissue regeneration; HA = hydroxyapatite; M = male; n = number; NBM = natural bone mineral; NcHA = nanocrystalline hydroxyapatite; OFD = open flap debridement; PD = probing depth; PI = plaque index; PPD = probing pocket depth; PRF = platelet-rich fibrin; PRFm = platelet-rich fibrin matrix; PRGF = plasma-rich growth factors; PRP = platelet-rich plasma; RBL = radiographic bone loss; REC = gingival recession; SRP = scaling and root planing; TDD = total defect depth.

Characteristics of excluded studies *[ordered by study ID]*

Study	Reason for exclusion
Agarwal 2017	Mixed design - randomised controlled trial
Aleksic 2008	No randomisation
Aroca 2009	Gingival recession (not infrabony defects)
Bajaj 2017	Mixed design - randomised controlled trial
Camargo 2002	No control group
Camargo 2005	No control group
Cetinkaya 2014	Same participants of Keles 2006
Chatterjee 2017	Mixed design - randomised controlled trial
Cheung 2004	Autologous platelet concentrates not the only difference between groups
Cieplik 2018	Incomplete data
Cortellini 1995	No platelet concentrate (fibrin glue)
Dogan 2015	Gingival recession (not infrabony defects)

(*Continued*)

Döri 2013	Same participants of Döri 2008b
Eren 2014	Autologous platelet concentrates not the only difference between groups
Gupta 2014b	Non-independence of analysis unit
Harnack 2009	Incomplete data
Huang 2005	Gingival recession (not infrabony defects)
Jankovic 2010	Gingival recession (not infrabony defects)
Jankovic 2012	Autologous platelet concentrates not the only difference between groups
Jovicic 2013	No randomisation
Keceli 2008	Incomplete data
Keles 2006	Incomplete data
Lekovic 2012	No control group
Menezes 2012	Incomplete data
Moder 2012	Same participants as Christgau 2006
Ouyang 2006	Mixed design randomised controlled trial
Padma 2013	Gingival recession (not infrabony defects)
Pradeep 2012a	Non-independence of analysis unit
Pradeep 2017	Mixed design - randomised controlled trial
Qiao 2016	Mixed design - randomised controlled trial
Saini 2011	No randomisation
Shah 2015	Incomplete data
Shepherd 2009	Gingival recession (not infrabony defects)
Shivakumar 2016	Gingival recession (not infrabony defects)
Thamaraiselvan 2015	Gingival recession (not infrabony defects)
Trombelli 1995	No platelet concentrate (fibrin glue)

(*Continued*)

Trombelli 1996	No platelet concentrate (fibrin glue)
Yajamanya 2017	Same participants of Chatterjee 2017
Yassibag-Berkman 2007	Incomplete data
Yen 2007	Incomplete data

DATA AND ANALYSES

Comparison 1. APC + OFD versus OFD (9-12 months)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Probing depth (mm)	12	510	Mean Difference (Random, 95% CI)	1.29 [1.00, 1.58]
1.1 Split-mouth studies	5	158	Mean Difference (Random, 95% CI)	1.86 [1.07, 2.66]
1.2 Parallel studies	7	352	Mean Difference (Random, 95% CI)	0.99 [0.90, 1.07]
2 Clinical attachment level (mm)	12	510	Mean Difference (Random, 95% CI)	1.47 [1.11, 1.82]
2.1 Split-mouth studies	5	158	Mean Difference (Random, 95% CI)	2.36 [1.19, 3.54]
2.2 Parallel studies	7	352	Mean Difference (Random, 95% CI)	0.99 [0.84, 1.14]
3 Radiographic bone defect filling (%)	9	401	Mean Difference (Random, 95% CI)	34.26 [30.07, 38.46]
3.1 Split-mouth studies	2	49	Mean Difference (Random, 95% CI)	27.32 [20.92, 33.72]
3.2 Parallel studies	7	352	Mean Difference (Random, 95% CI)	35.77 [31.20, 40.35]

Comparison 2. APC + OFD + BG versus OFD + BG (all follow-ups)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Probing depth (mm)	17	569	Mean Difference (Random, 95% CI)	0.54 [0.33, 0.75]
1.1 Split-mouth studies	12	360	Mean Difference (Random, 95% CI)	0.47 [0.24, 0.71]
1.2 Parallel studies	5	209	Mean Difference (Random, 95% CI)	0.81 [0.58, 1.03]
2 Clinical attachment level (mm)	17	569	Mean Difference (Random, 95% CI)	0.72 [0.43, 1.00]
2.1 Split-mouth studies	12	360	Mean Difference (Random, 95% CI)	0.67 [0.35, 0.99]
2.2 Parallel studies	5	209	Mean Difference (Random, 95% CI)	0.89 [0.49, 1.29]
3 Radiographic bone defect filling (%)	11	420	Mean Difference (Random, 95% CI)	8.10 [5.26, 10.94]
3.1 Split-mouth studies	8	270	Mean Difference (Random, 95% CI)	7.73 [4.50, 10.97]
3.2 Parallel studies	3	150	Mean Difference (Random, 95% CI)	9.66 [5.39, 13.94]

Comparison 3. APC + OFD + BG versus OFD + BG (3-6 months)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Probing depth (mm)	11	272	Mean Difference (Random, 95% CI)	0.62 [0.30, 0.94]
1.1 Split-mouth studies	10	252	Mean Difference (Random, 95% CI)	0.58 [0.25, 0.92]
1.2 Parallel studies	1	20	Mean Difference (Random, 95% CI)	0.84 [0.60, 1.07]
2 Clinical attachment level (mm)	11	272	Mean Difference (Random, 95% CI)	0.47 [0.11, 0.84]
2.1 Split-mouth studies	10	252	Mean Difference (Random, 95% CI)	0.40 [0.02, 0.77]
2.2 Parallel studies	1	20	Mean Difference (Random, 95% CI)	1.0 [0.93, 1.07]

3 Radiographic bone defect filling (%)	6	162	Mean Difference (Random, 95% CI)	4.76 [1.27, 8.25]
3.1 Split-mouth studies	5	142	Mean Difference (Random, 95% CI)	3.59 [0.13, 7.05]
3.2 Parallel studies	1	20	Mean Difference (Random, 95% CI)	10.0 [4.90, 15.10]

Comparison 4. APC + OFD + BG versus OFD + BG (9-12 months)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Probing depth (mm)	10	381	Mean Difference (Random, 95% CI)	0.50 [0.31, 0.69]
1.1 Split-mouth studies	6	192	Mean Difference (Random, 95% CI)	0.49 [0.26, 0.72]
1.2 Parallel studies	4	189	Mean Difference (Random, 95% CI)	0.58 [0.09, 1.06]
2 Clinical attachment level (mm)	6	192	Mean Difference (Random, 95% CI)	0.84 [0.62, 1.06]
2.1 Split-mouth studies	6	192	Mean Difference (Random, 95% CI)	0.84 [0.62, 1.06]
3 Radiographic bone defect filling (%)	6	282	Mean Difference (Random, 95% CI)	9.99 [6.44, 13.55]
3.1 Split-mouth studies	4	152	Mean Difference (Random, 95% CI)	10.16 [6.18, 14.14]
3.2 Parallel studies	2	130	Mean Difference (Random, 95% CI)	8.87 [1.03, 16.71]

Comparison 5. APC + GTR versus GTR (all follow-ups)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Probing depth (mm)	7	248	Mean Difference (Random, 95% CI)	0.92 [-0.02, 1.86]
1.1 Split-mouth studies	4	166	Mean Difference (Random, 95% CI)	1.52 [0.54, 2.51]
1.2 Parallel studies	3	82	Mean Difference (Random, 95% CI)	0.25 [-0.15, 0.64]
2 Clinical attachment level (mm)	7	248	Mean Difference (Random, 95% CI)	0.42 [-0.02, 0.86]
2.1 Split-mouth studies	4	166	Mean Difference (Random, 95% CI)	0.67 [0.20, 1.14]
2.2 Parallel studies	3	82	Mean Difference (Random, 95% CI)	0.09 [-0.32, 0.50]

Comparison 6. APC + GTR versus GTR (3-6 months)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Probing depth (mm)	3	134	Mean Difference (Random, 95% CI)	1.07 [-0.71, 2.86]
1.1 Split-mouth studies	3	134	Mean Difference (Random, 95% CI)	1.07 [-0.71, 2.86]
2 Clinical attachment level (mm)	3	134	Mean Difference (Random, 95% CI)	0.54 [0.18, 0.89]
2.1 Split-mouth studies	3	134	Mean Difference (Random, 95% CI)	0.54 [0.18, 0.89]

Comparison 7. APC + GTR versus GTR (9-12 months)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Probing depth (mm)	5	164	Mean Difference (Random, 95% CI)	0.68 [-0.66, 2.02]
1.1 Split-mouth studies	2	82	Mean Difference (Random, 95% CI)	1.53 [-0.85, 3.91]
1.2 Parallel studies	3	82	Mean Difference (Random, 95% CI)	0.25 [-0.15, 0.64]
2 Clinical attachment level (mm)	5	164	Mean Difference (Random, 95% CI)	0.27 [-0.39, 0.93]
2.1 Split-mouth studies	2	82	Mean Difference (Random, 95% CI)	0.51 [-0.72, 1.73]
2.2 Parallel studies	3	82	Mean Difference (Random, 95% CI)	0.09 [-0.32, 0.50]

Comparison 8. APC + EMD versus EMD (all follow-up)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Probing depth (mm)	2	75	Mean Difference (Random, 95% CI)	0.13 [-0.05, 0.30]
1.1 Split-mouth studies	1	49	Mean Difference (Random, 95% CI)	0.13 [-0.05, 0.31]
1.2 Parallel studies	1	26	Mean Difference (Random, 95% CI)	-0.10 [-1.32, 1.12]
2 Clinical attachment level (mm)	2	75	Mean Difference (Random, 95% CI)	0.10 [-0.13, 0.32]
2.1 Split-mouth studies	1	49	Mean Difference (Random, 95% CI)	0.12 [-0.12, 0.36]
2.2 Parallel studies	1	26	Mean Difference (Random, 95% CI)	-0.2 [-1.06, 0.66]
3 Radiographic bone defect filling (%)	1	49	Mean Difference (Random, 95% CI)	-0.6 [-6.21, 5.01]
3.1 Split-mouth studies	1	49	Mean Difference (Random, 95% CI)	-0.6 [-6.21, 5.01]

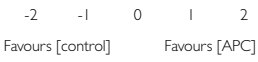
Analysis 1.1. Comparison 1 APC + OFD versus OFD (9-12 months), Outcome 1 Probing depth (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 1 APC + OFD versus OFD (9-12 months)

Outcome: 1 Probing depth (mm)

Study or subgroup	APC N	Control N	Mean Difference (SE)	Mean Difference IV,Random,95% CI	Weight	Mean Difference IV,Random,95% CI
1 Split-mouth studies						
Agarwal 2016	10	10	2.17 (0.8)		2.8 %	2.17 [0.60, 3.74]
Arabaci 2017	26	26	0.7 (0.29)		9.2 %	0.70 [0.13, 1.27]
Patel 2017	13	13	1.8 (0.54)		4.9 %	1.80 [0.74, 2.86]
Rosamma Joseph 2012	15	15	2.26 (0.29)		9.2 %	2.26 [1.69, 2.83]
Thorat 2017	15	15	2.5 (0.29)		9.2 %	2.50 [1.93, 3.07]
Subtotal (95% CI)	**79**	**79**			**35.2 %**	**1.86 [1.07, 2.66]**
Heterogeneity: Tau² = 0.63; Chi² = 22.90, df = 4 (P = 0.00013); I² =83%						
Test for overall effect: Z = 4.58 (P < 0.00001)						
2 Parallel studies						
Chandradas 2016	12	12	0.82 (0.[illegible])		6.8 %	0.82 [0.02, 1.62]
Kanoriya 2016	30	30	0.84 (0.23)		10.5 %	0.84 [0.39, 1.29]
Martande 2016	30	30	[illegible]0.31)		8.7 %	1.00 [0.39, 1.61]
Pradeep 2015	30	30	1 (0.05)		13.8 %	1.00 [0.90, 1.10]
Pradeep 2016	30	30	[illegible].12)		12.9 %	0.93 [0.69, 1.17]
Sharma 2011	28	28	1.34 (0.47)		5.9 %	1.34 [0.42, 2.26]
Thorat 2011	16	[illegible]	1.13 (0.45)		6.2 %	1.13 [0.25, 2.01]
Subtotal (95% CI)	**176**	**17[illegible]**			**64.8 %**	**0.99 [0.90, 1.07]**
Heterogeneity: Tau² = 0.0; Chi² = 1.53[illegible] = 6 [illegible] = 0.96), I² =0.0%						
Test for overall effect: Z = 22.38 (P [illegible].00001)						
Total (95% CI)	**255**	**255**			**100.0 %**	**1.29 [1.00, 1.58]**
Heterogeneity: Tau² = 0.16; Chi² = 51.34, df [illegible] (P<0.00001); I² =79%						
Test for overall effect: Z = 8.65 (P < 0.00001)						
Test for subgroup differences: [illegible]² = 4.58, df = 1 (P = 0.03), I² =78%						

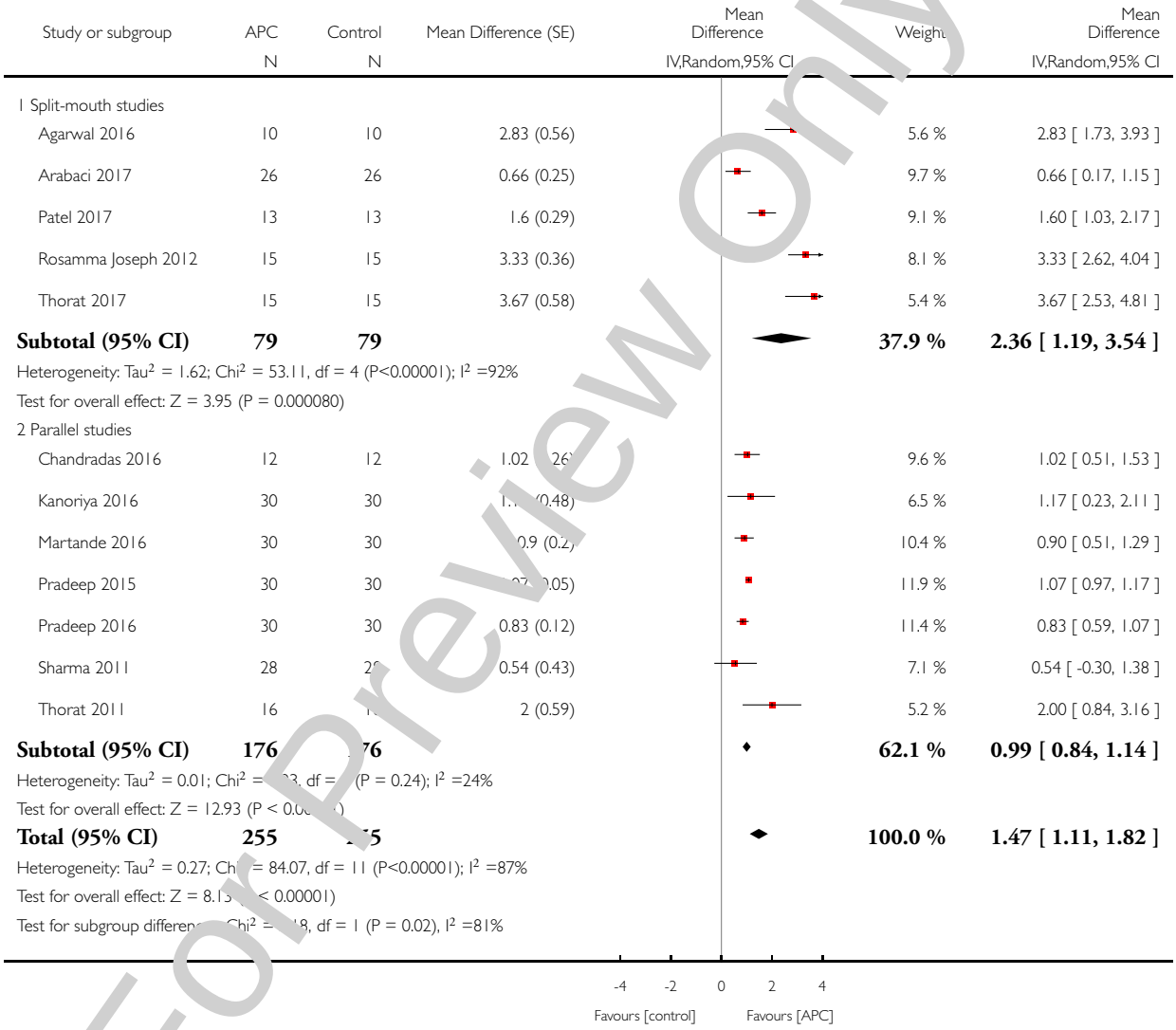


Analysis 1.2. Comparison 1 APC + OFD versus OFD (9-12 months), Outcome 2 Clinical attachment level (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 1 APC + OFD versus OFD (9-12 months)

Outcome: 2 Clinical attachment level (mm)

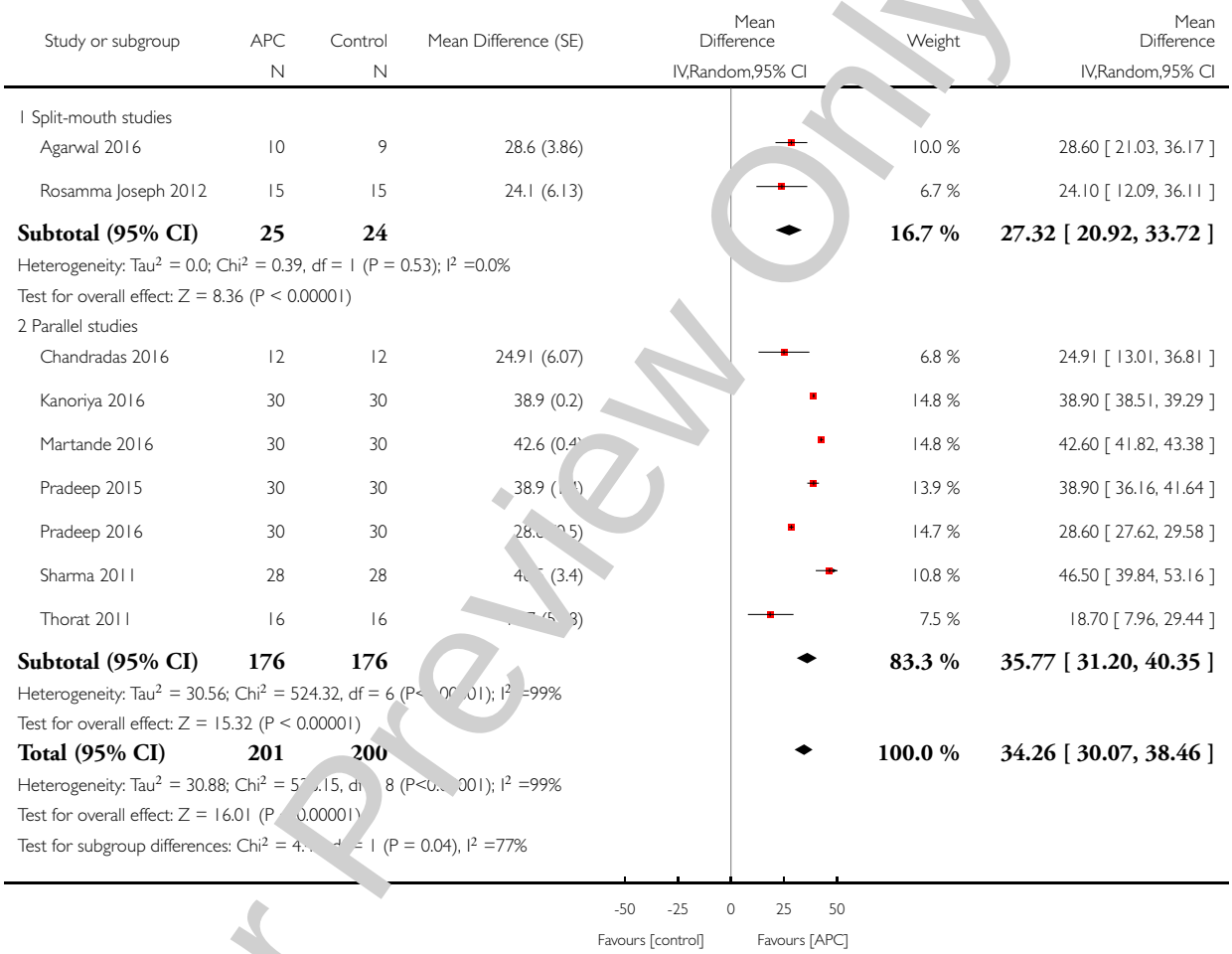


Analysis 1.3. Comparison 1 APC + OFD versus OFD (9-12 months), Outcome 3 Radiographic bone defect filling (%).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 1 APC + OFD versus OFD (9-12 months)

Outcome: 3 Radiographic bone defect filling (%)

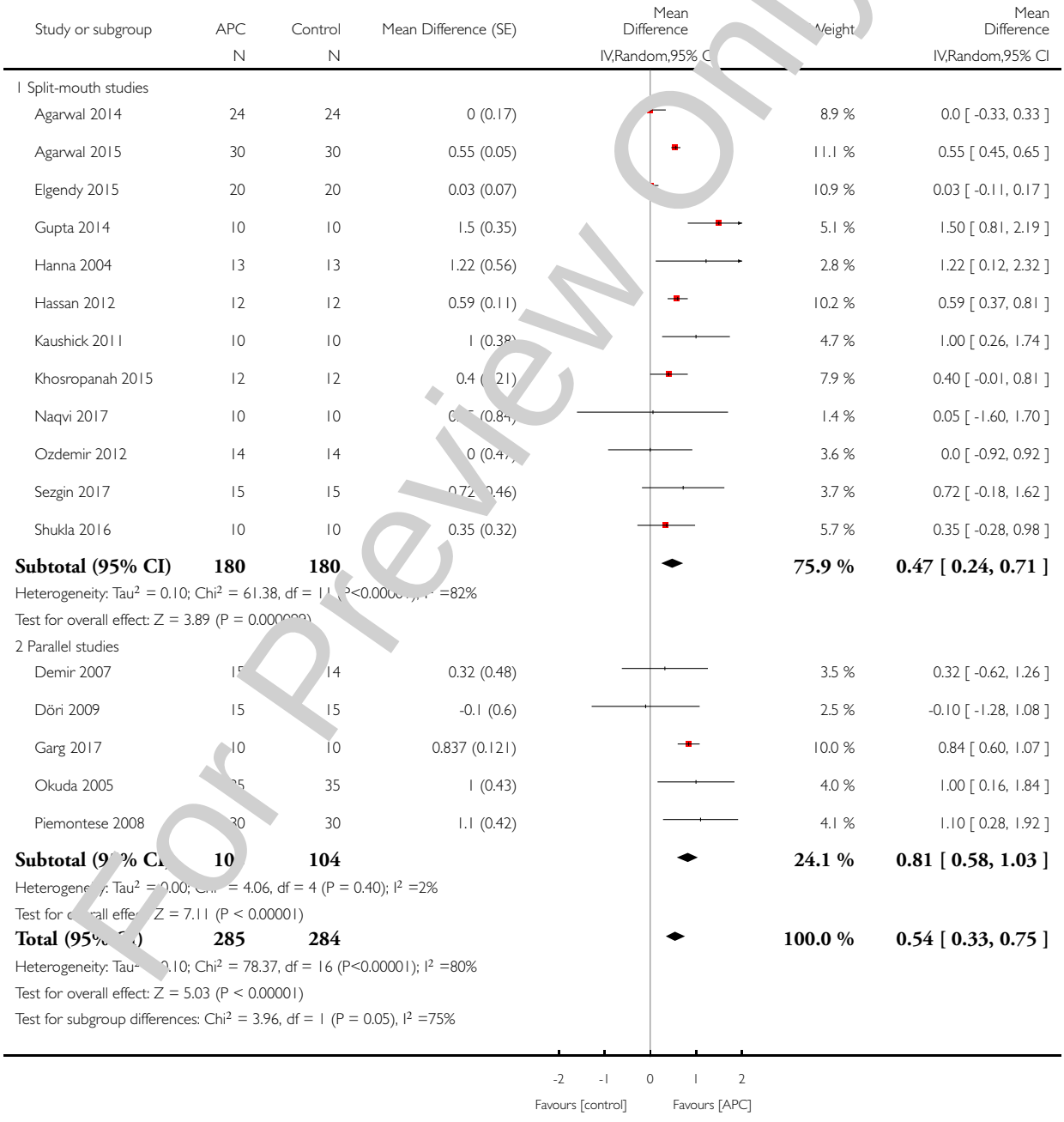


Analysis 2.1. Comparison 2 APC + OFD + BG versus OFD + BG (all follow-ups), Outcome 1 Probing depth (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 2 APC + OFD + BG versus OFD + BG (all follow-ups)

Outcome: 1 Probing depth (mm)

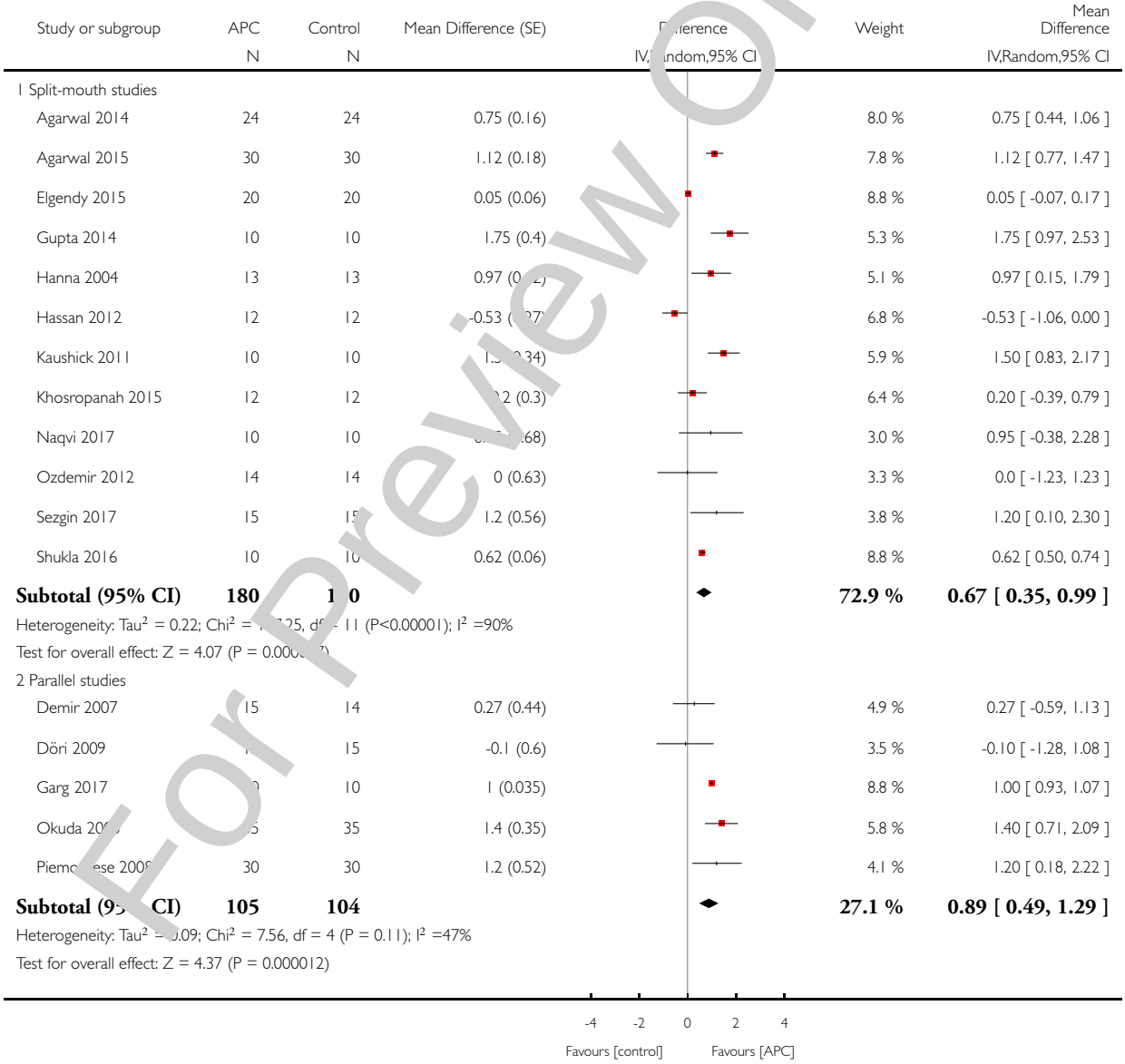


Analysis 2.2. Comparison 2 APC + OFD + BG versus OFD + BG (all follow-ups), Outcome 2 Clinical attachment level (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 2 APC + OFD + BG versus OFD + BG (all follow-ups)

Outcome: 2 Clinical attachment level (mm)



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Study or subgroup	APC N	Control N	Mean Difference (SE)	Mean Difference IV,Random,95% CI	Weight	Mean Difference IV,Random,95% CI
Total (95% CI)	**285**	**284**		◆	**100.0 %**	**0.72 [0.43, 1.00]**
Heterogeneity: Tau² = 0.23; Chi² = 242.48, df = 16 (P<0.00001); I² =93%						
Test for overall effect: Z = 4.98 (P < 0.00001)						
Test for subgroup differences: Chi² = 0.75, df = 1 (P = 0.39), I² =0.0%						

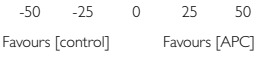
Analysis 2.3. Comparison 2 APC + OFD + BG versus OFD + BG (all follow-ups), Outcome 3 Radiographic bone defect filling (%).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 2 APC + OFD + BG versus OFD + BG (all follow-ups)

Outcome: 3 Radiographic bone defect filling (%)

Study or subgroup	APC N	Control N	Mean Difference (SE)	Mean Difference IV,Random,95% CI	Weight	Mean Difference IV,Random,95% CI
1 Split-mouth studies						
Agarwal 2014	24	24	5.5 (0.59)	■	15.8 %	5.50 [4.34, 6.66]
Agarwal 2015	30	30	9.9 (0.92)	■	15.2 %	9.90 [8.10, 11.70]
Elgendy 2015	20	20	2.7 (0.75)	■	15.5 %	2.70 [1.23, 4.17]
Gupta 2014	10	10	12.2 (20.25)	■	0.5 %	12.20 [-27.49, 51.89]
Hassan 2012	12	12	15 (2.24)	■	11.7 %	15.00 [10.61, 19.39]
Kaushick 2011	10	10	15.4 (3.39)	■	8.6 %	15.40 [8.76, 22.04]
Ozdemir 2012	14	14	-8.4 (8.89)	■	2.3 %	-8.40 [-25.82, 9.02]
Sezgin 2013	15	15	4.4 (2.51)	■	10.9 %	4.40 [-0.52, 9.32]
Subtotal (95% CI)	**135**	**135**		◆	**80.4 %**	**7.73 [4.50, 10.97]**
Heterogeneity: Tau² = 13.66; Chi² = 64.96, df = 7 (P<0.00001); I² =89%						
Test for overall effect: Z = 4.69 (P < 0.00001)						
2 Parallel studies						



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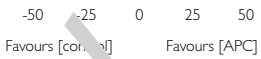
Study or subgroup	APC N	Control N	Mean Difference (SE)	Mean Difference IV,Random,95% CI	Weight	Mean Difference IV,Random,95% CI
Garg 2017	10	10	10 (2.6)		10.6 %	10.00 [4.90, 15.10]
Okuda 2005	35	35	13.7 (7.58)		0 %	13.70 [-1.16, 28.56]
Piemontese 2008	30	30	7 (4.71)		6.0	7.00 [-2.23, 16.23]
Subtotal (95% CI)	**75**	**75**			**19.6 %**	**9.66 [5.39, 13.94]**
Heterogeneity: Tau^2 = 0.0; Chi^2 = 0.62, df = 2 (P = 0.73); I^2 =0.0%						
Test for overall effect: Z = 4.43 (P < 0.00001)						
Total (95% CI)	**210**	**210**			**100.0 %**	**8.10 [5.26, 10.94]**
Heterogeneity: Tau^2 = 12.93; Chi^2 = 68.44, df = 10 (P<0.00001); I^2 =85%						
Test for overall effect: Z = 5.60 (P < 0.00001)						
Test for subgroup differences: Chi^2 = 0.50, df = 1 (P = 0.48), I^2 =0.0%						

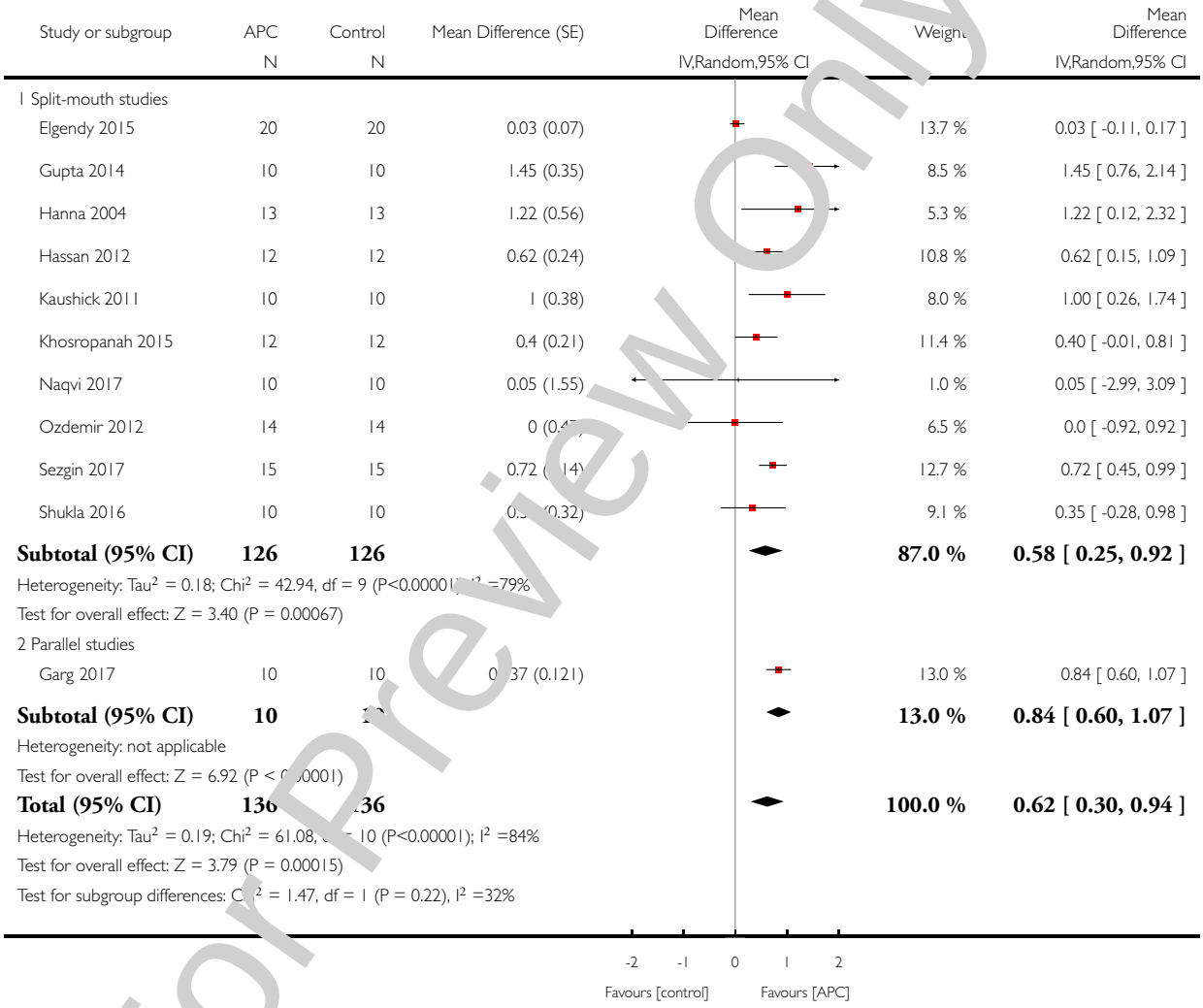


Analysis 3.1. Comparison 3 APC + OFD + BG versus OFD + BG (3-6 months), Outcome 1 Probing depth (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 3 APC + OFD + BG versus OFD + BG (3-6 months)

Outcome: 1 Probing depth (mm)

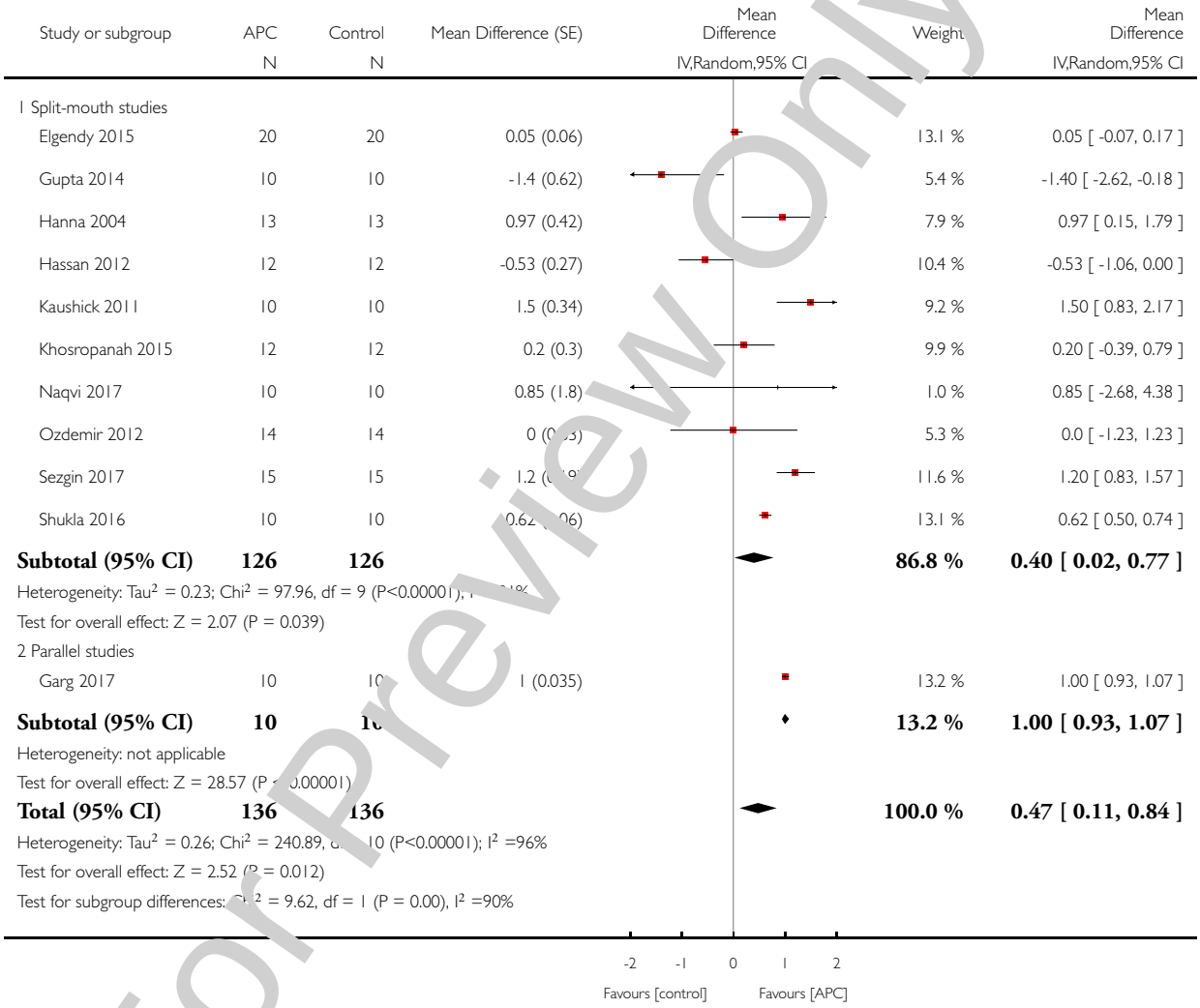


Analysis 3.2. Comparison 3 APC + OFD + BG versus OFD + BG (3-6 months), Outcome 2 Clinical attachment level (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 3 APC + OFD + BG versus OFD + BG (3-6 months)

Outcome: 2 Clinical attachment level (mm)

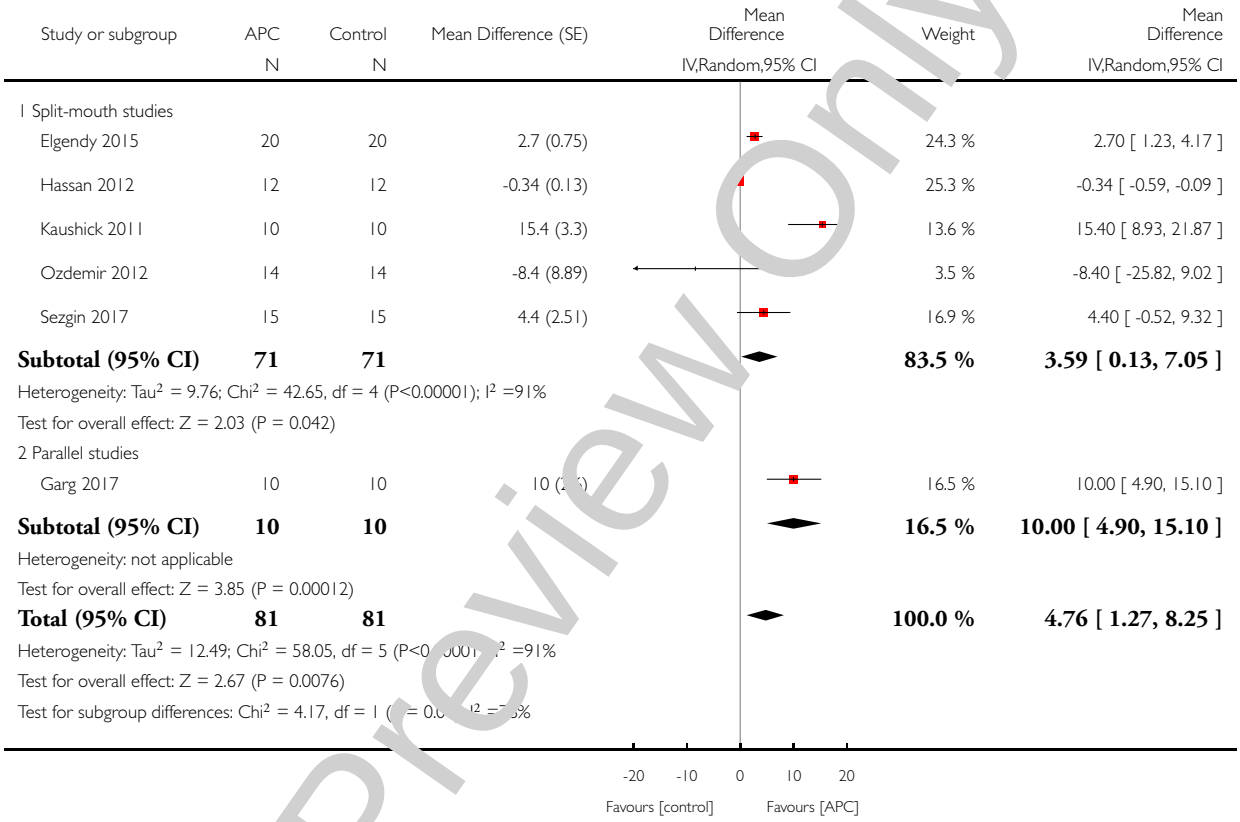


Analysis 3.3. Comparison 3 APC + OFD + BG versus OFD + BG (3-6 months), Outcome 3 Radiographic bone defect filling (%).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 3 APC + OFD + BG versus OFD + BG (3-6 months)

Outcome: 3 Radiographic bone defect filling (%)

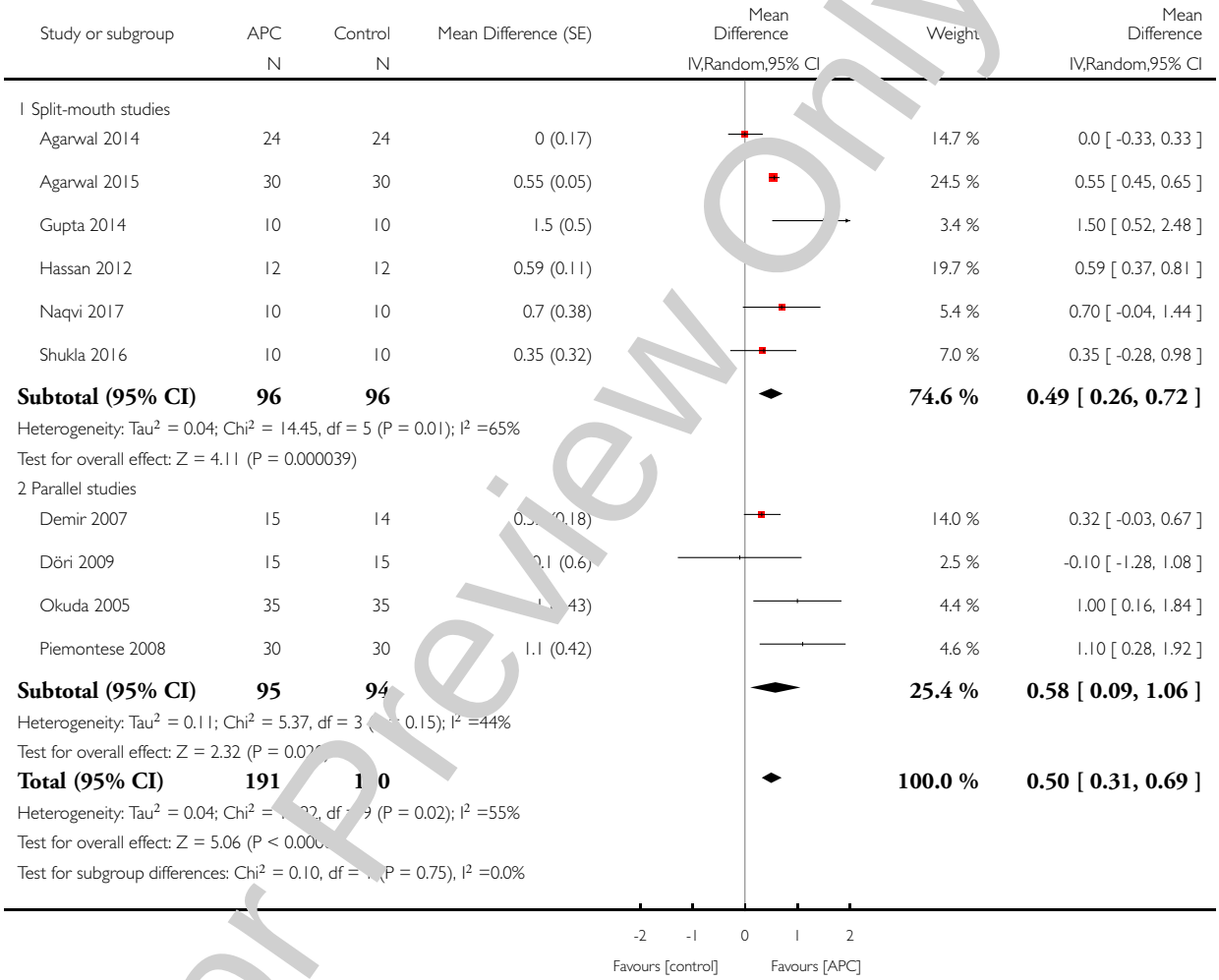


Analysis 4.1. Comparison 4 APC + OFD + BG versus OFD + BG (9-12 months), Outcome 1 Probing depth (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 4 APC + OFD + BG versus OFD + BG (9-12 months)

Outcome: 1 Probing depth (mm)



Analysis 4.2. Comparison 4 APC + OFD + BG versus OFD + BG (9-12 months), Outcome 2 Clinical attachment level (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 4 APC + OFD + BG versus OFD + BG (9-12 months)

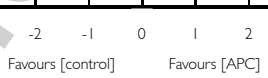
Outcome: 2 Clinical attachment level (mm)

Study or subgroup	APC N	Control N	Mean Difference (SE)	Mean Difference IV,Random,95% CI	Weight	Mean Difference IV,Random,95% CI
1 Split-mouth studies						
Agarwal 2014	24	24	0.75 (0.16)		20.4 %	0.75 [0.44, 1.06]
Agarwal 2015	30	30	1.12 (0.18)		18.3 %	1.12 [0.77, 1.47]
Gupta 2014	10	10	1.75 (0.58)		3.3 %	1.75 [0.61, 2.89]
Hassan 2012	12	12	0.87 (0.12)		25.1 %	0.87 [0.63, 1.11]
Naqvi 2017	10	10	0.95 (1.23)		0.8 %	0.95 [-1.46, 3.36]
Shukla 2016	10	10	0.62 (0.06)		32.2 %	0.62 [0.50, 0.74]
Total (95% CI)	**96**	**96**			**100.0 %**	**0.84 [0.62, 1.06]**

Heterogeneity: $Tau^2 = 0.03$; $Chi^2 = 12.50$, df = 5 (P = 0.03); $I^2 = 60\%$

Test for overall effect: Z = 7.62 (P < 0.00001)

Test for subgroup differences: Not applicable

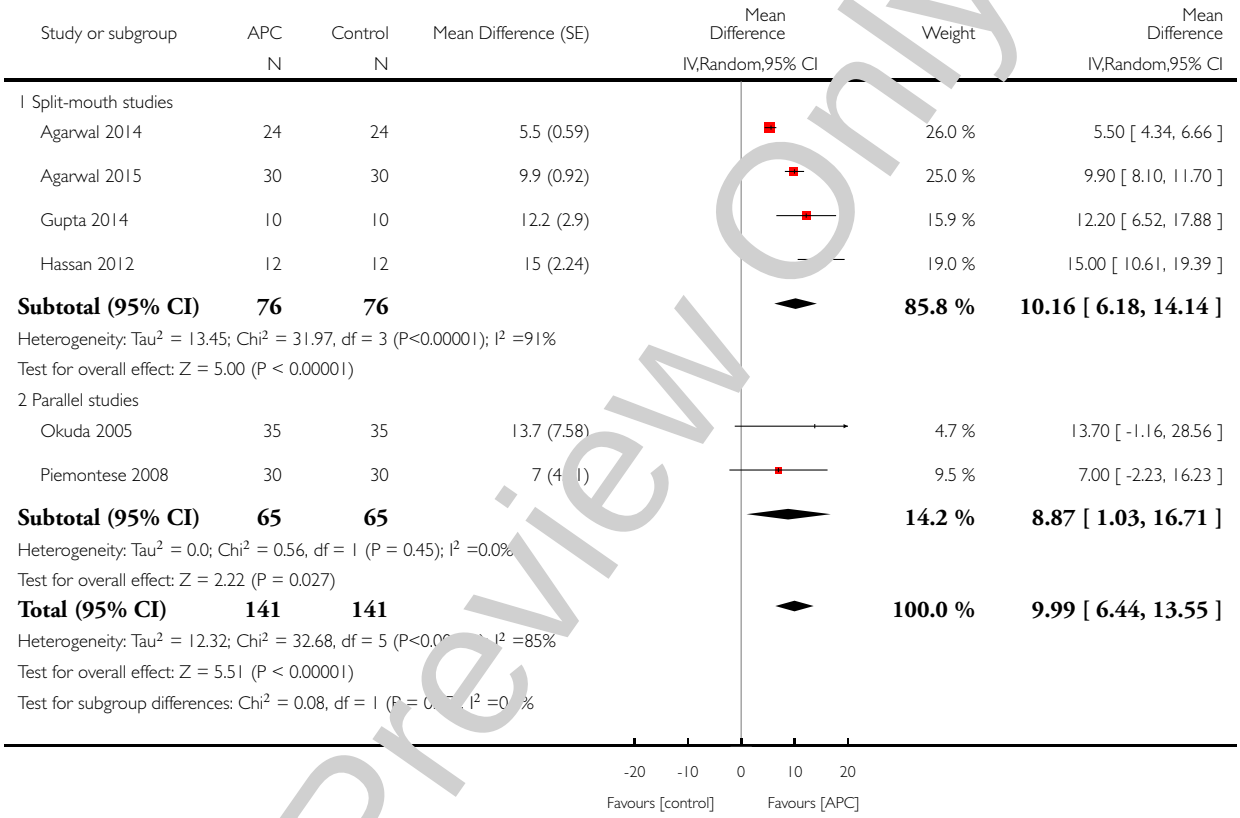


Analysis 4.3. Comparison 4 APC + OFD + BG versus OFD + BG (9-12 months), Outcome 3 Radiographic bone defect filling (%).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 4 APC + OFD + BG versus OFD + BG (9-12 months)

Outcome: 3 Radiographic bone defect filling (%)

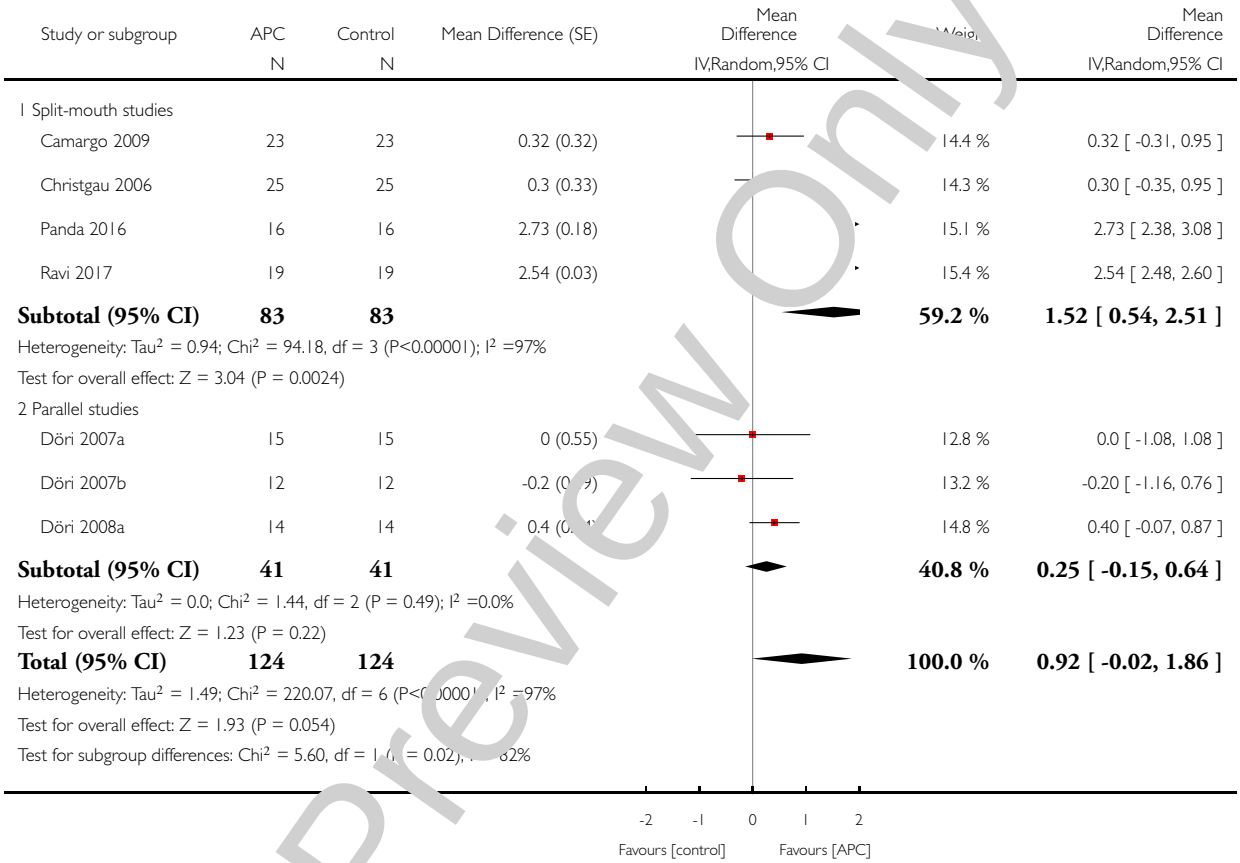


Analysis 5.1. Comparison 5 APC + GTR versus GTR (all follow-ups), Outcome 1 Probing depth (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 5 APC + GTR versus GTR (all follow-ups)

Outcome: 1 Probing depth (mm)

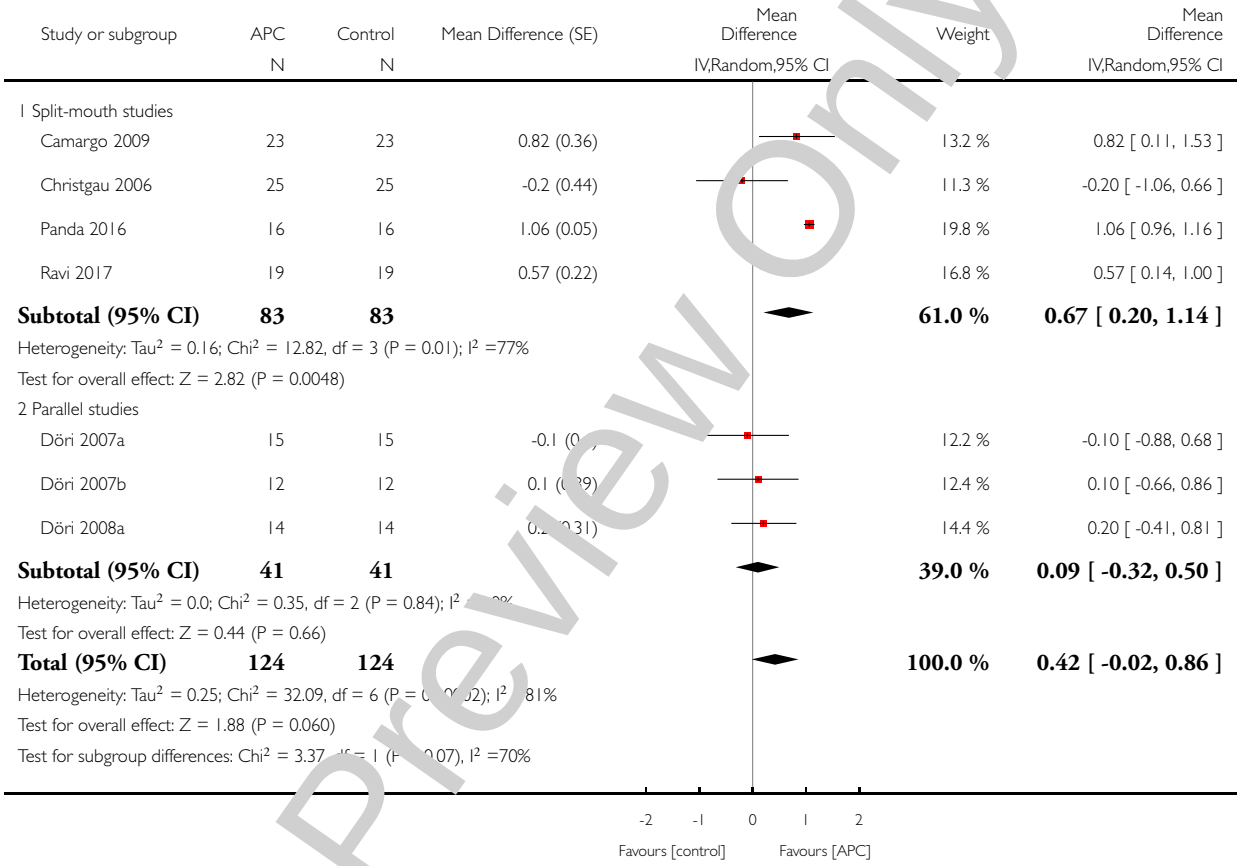


Analysis 5.2. Comparison 5 APC + GTR versus GTR (all follow-ups), Outcome 2 Clinical attachment level (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 5 APC + GTR versus GTR (all follow-ups)

Outcome: 2 Clinical attachment level (mm)

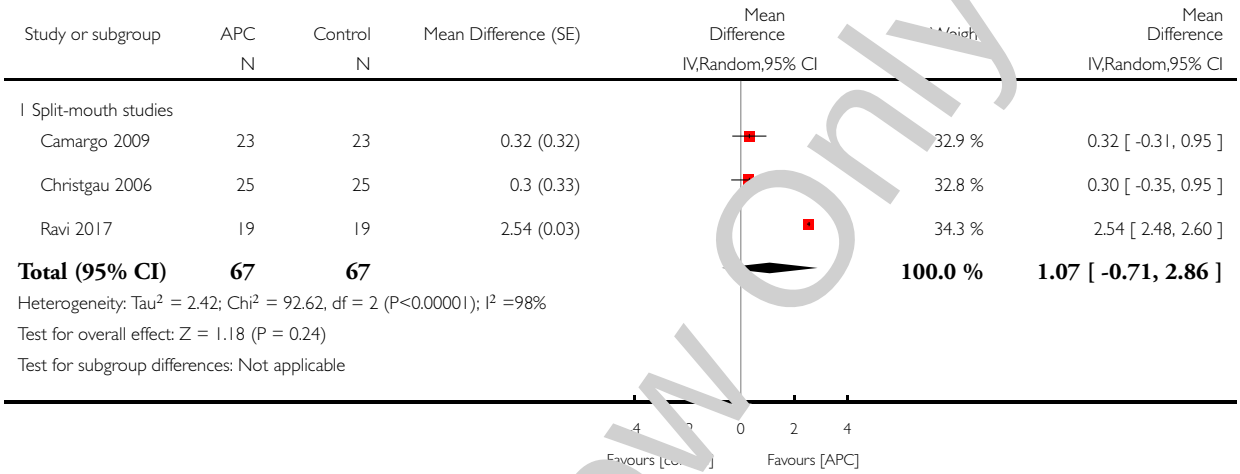


Analysis 6.1. Comparison 6 APC + GTR versus GTR (3-6 months), Outcome 1 Probing depth (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 6 APC + GTR versus GTR (3-6 months)

Outcome: 1 Probing depth (mm)



Analysis 6.2. Comparison 6 APC + GTR versus GTR (3-6 months), Outcome 2 Clinical attachment level (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 6 APC + GTR versus GTR (3-6 months)

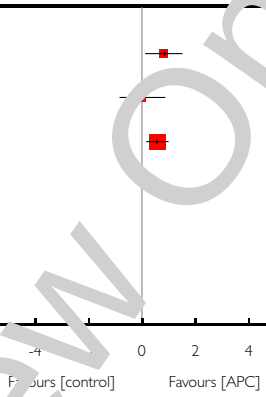
Outcome: 2 Clinical attachment level (mm)

Study or subgroup	APC N	Control N	Mean Difference (SE)	Mean Difference IV,Random,95% CI	Weight	Mean Difference IV,Random,95% CI
1 Split-mouth studies						
Camargo 2009	23	23	0.82 (0.36)		24.1 %	0.82 [0.11, 1.53]
Christgau 2006	25	25	0.001 (0.44)		16.4 %	0.00 [-0.86, 0.86]
Ravi 2017	19	19	0.57 (0.22)		59.5 %	0.57 [0.14, 1.00]
Total (95% CI)	**67**	**67**			**100.0 %**	**0.54 [0.18, 0.89]**

Heterogeneity: $Tau^2 = 0.01$; $Chi^2 = 2.12$, df = 2 (P = 0.35); $I^2 = 6\%$

Test for overall effect: Z = 2.96 (P = 0.0031)

Test for subgroup differences: Not applicable

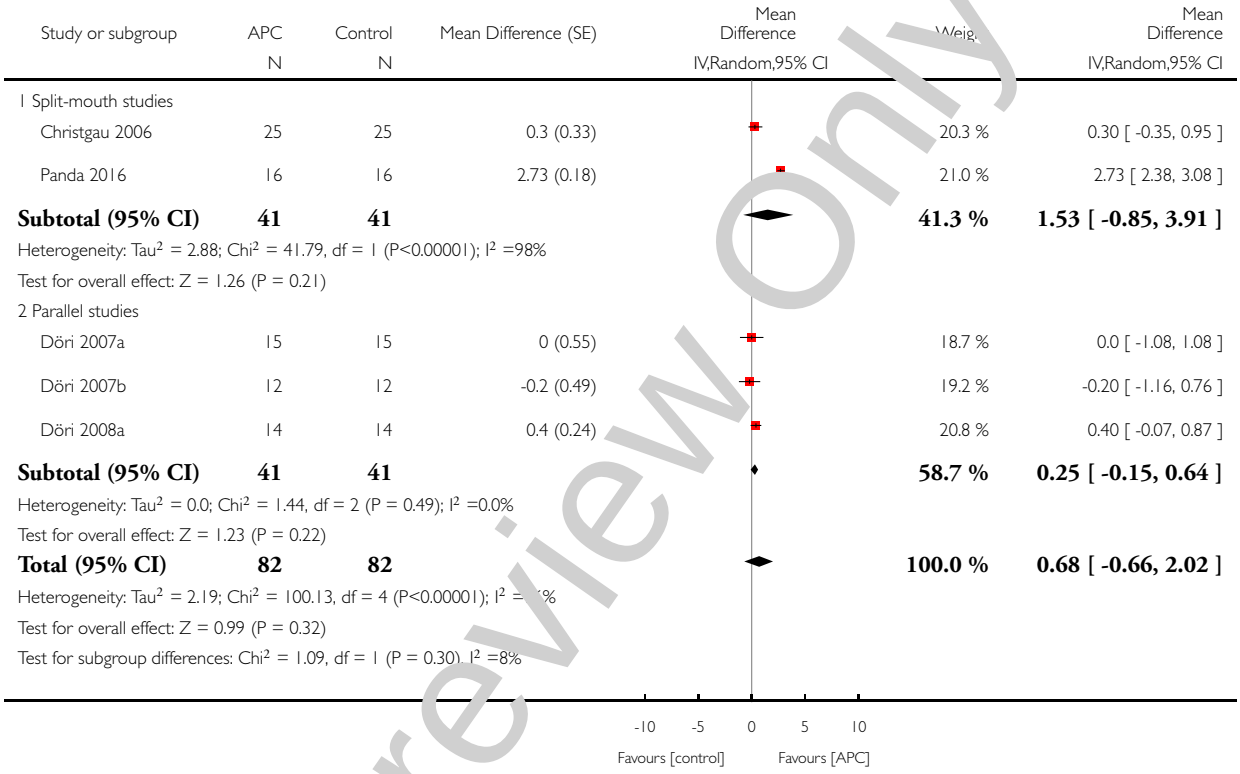


Analysis 7.1. Comparison 7 APC + GTR versus GTR (9-12 months), Outcome 1 Probing depth (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 7 APC + GTR versus GTR (9-12 months)

Outcome: 1 Probing depth (mm)



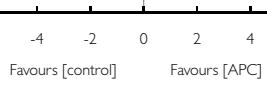
Analysis 7.2. Comparison 7 APC + GTR versus GTR (9-12 months), Outcome 2 Clinical attachment level (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 7 APC + GTR versus GTR (9-12 months)

Outcome: 2 Clinical attachment level (mm)

Study or subgroup	APC N	Control N	Mean Difference (SE)	Mean Difference IV,Random,95% CI	Weight	Mean Difference IV,Random,95% CI
1 Split-mouth studies						
Christgau 2006	25	25	-0.2 (0.44)		17.5 %	-0.20 [-1.06, 0.66]
Panda 2016	16	16	1.06 (0.05)		24.9 %	1.06 [0.96, 1.16]
Subtotal (95% CI)	**41**	**41**			**42.3 %**	**0.51 [-0.72, 1.73]**
Heterogeneity: Tau^2 = 0.70; Chi^2 = 8.10, df = 1 (P = 0.004); I^2 =88%						
Test for overall effect: Z = 0.81 (P = 0.42)						
2 Parallel studies						
Döri 2007a	15	15	-0.1 (0.4)		18.4 %	-0.10 [-0.88, 0.68]
Döri 2007b	12	12	0.1 (0.39)		18.7 %	0.10 [-0.66, 0.86]
Döri 2008a	14	14	0.2 (0.3[illegible])		20.6 %	0.20 [-0.41, 0.81]
Subtotal (95% CI)	**41**	**41**			**57.7 %**	**0.09 [-0.32, 0.50]**
Heterogeneity: Tau^2 = 0.0; Chi^2 = 0.35, df = 2 (P = 0.84); I^2 =0.0%						
Test for overall effect: Z = 0.44 (P = 0.66)						
Total (95% CI)	**82**	**82**			**100.0 %**	**0.27 [-0.39, 0.93]**
Heterogeneity: Tau^2 = 0.45; Chi^2 = 28.40, df = 4 (P = 0.0000[illegible]); I^2 =86%						
Test for overall effect: Z = 0.81 (P = 0.42)						
Test for subgroup differences: Chi^2 = 0.40, df = 1 (P = [illegible]53), I^2 =0.0%						

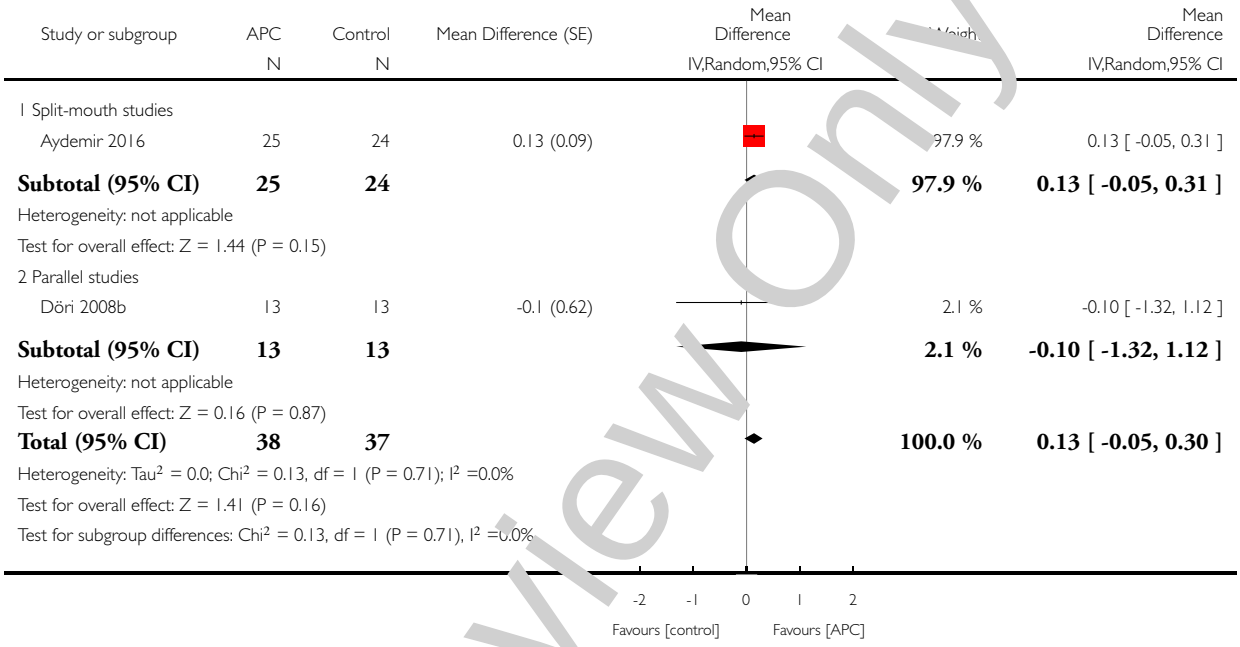


Analysis 8.1. Comparison 8 APC + EMD versus EMD (all follow-ups), Outcome 1 Probing depth (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 8 APC + EMD versus EMD (all follow-ups)

Outcome: 1 Probing depth (mm)



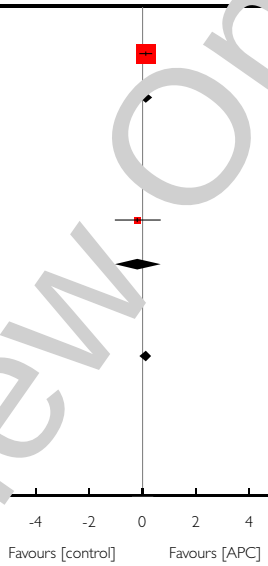
Analysis 8.2. Comparison 8 APC + EMD versus EMD (all follow-ups), Outcome 2 Clinical attachment level (mm).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 8 APC + EMD versus EMD (all follow-ups)

Outcome: 2 Clinical attachment level (mm)

Study or subgroup	APC N	Control N	Mean Difference (SE)	Mean Difference IV,Random,95% CI	Weight	Mean Difference IV,Random,95% CI
1 Split-mouth studies						
Aydemir 2016	25	24	0.12 (0.12)		93.1 %	0.12 [-0.12, 0.36]
Subtotal (95% CI)	**25**	**24**			**93.1 %**	**0.12 [-0.12, 0.36]**
Heterogeneity: not applicable						
Test for overall effect: Z = 1.00 (P = 0.32)						
2 Parallel studies						
Döri 2008b	13	13	-0.2 (0.44)		6.9 %	-0.20 [-1.06, 0.66]
Subtotal (95% CI)	**13**	**13**			**6.9 %**	**-0.20 [-1.06, 0.66]**
Heterogeneity: not applicable						
Test for overall effect: Z = 0.45 (P = 0.65)						
Total (95% CI)	**38**	**37**			**100.0 %**	**0.10 [-0.13, 0.32]**
Heterogeneity: Tau^2 = 0.0; Chi^2 = 0.49, df = 1 (P = 0.48); I^2 =0.0%						
Test for overall effect: Z = 0.85 (P = 0.40)						
Test for subgroup differences: Chi^2 = 0.49, df = 1 (P = 0.48), I^2 =0.0%						

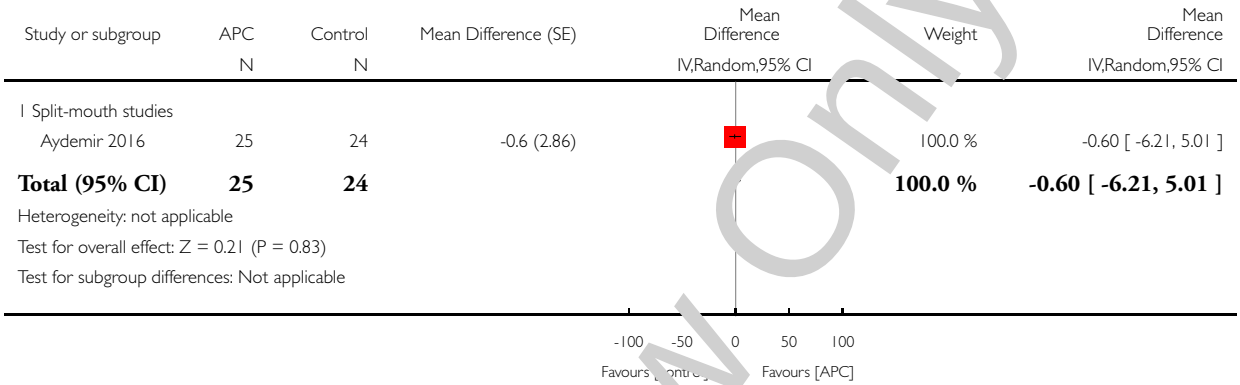


Analysis 8.3. Comparison 8 APC + EMD versus EMD (all follow-ups), Outcome 3 Radiographic bone defect filling (%).

Review: Autologous platelet concentrates for treating periodontal infrabony defects

Comparison: 8 APC + EMD versus EMD (all follow-ups)

Outcome: 3 Radiographic bone defect filling (%)



APPENDICES

Appendix 1. Cochrane Oral Health's Trials Register search strategy

1. (periodont*:ti,ab) AND (INREGISTER)
2. ((infrabony or "infra bony" or intrabony or "intra bony" or infraosseous or "infra osseous" or endosseous or apicomarginal or "apico marginal" or interproximal or "inter proximal"):ti,ab) AND (INREGISTER)
3. (("vertical bone" and defect*):ti,ab) AND (INREGISTER)
4. ((bone and resorp*):ti,ab) AND (INREGISTER)
5. ((intraalveolar or "intra alveolar"):ti,ab) AND (INREGISTER)
6. (#1 or #2 or #3 or #4 or #5) AND (INREGISTER)
7. ((platelet* and (plasma* or fibrin* or concentrat*))) AND (INREGISTER)
8. ((PRP or PRGF or PRF or L-PRF):ti,ab) AND (INREGISTER)
9. (#7 or #8) AND (INREGISTER)
10. (#6 and #9) AND (INREGISTER)

Appendix 2. Cochrane Central Register of Controlled Clinical Trials (CENTRAL) search strategy

#1 [mh "Platelet-rich plasma"]
#2 [mh Fibrin]
#3 (platelet* near/5 (plasma* or fibrin* or concentrat*))
#4 (PRP or L-PRP or PRF or L-PRF):ti,ab
#5 {or #1-#4}
#6 [mh "periodontal diseases"]
#7 periodont*
#8 (infrabony or "infra bony" or intrabony or "intra bony" or infraosseous or "infra osseous" or endosseous or apicomarginal or "apico marginal" or interproximal or "inter proximal")
#9 ("vertical bone" and defect*)
#10 (bone near/3 resorp*)
#11 (intraalveolar or "intra alveolar")
#12 {or #6-#11}
#13 #5 and #12

Appendix 3. MEDLINE Ovid search strategy

1. Platelet-rich plasma/
2. exp Fibrin/
3. (platelet$ adj5 (plasma$ or fibrin$ or concentrat$)).mp.
4. (PRP or L-PRP or PRF or L-PRF).ti,ab.
5. or/1-4
6. exp Periodontal diseases/
7. periodont$.mp.
8. (infrabony or "infra bony" or intrabony or "intra bony" or infraosseous or "infra ossesous" or endosseous or apicomarginal or "apico marginal" or interproximal or "inter proximal").ti,ab.
9. ((vertical adj bone) and defect$).ti,ab.
10. (bone adj3 resorp$).ti,ab.
11. (intraalveolar or "intra alveolar").ti,ab.
12. or/6-11
13. 5 and 12

This subject search was linked to the Cochrane Highly Sensitive Search Strategy (CHSSS) for identifying randomised trials (RCTs) in MEDLINE: sensitivity-maximising version (2008 revision) as referenced in Chapter 6.4.11.1 and detailed in box 6.4.c of the*Cochrane Handbook for Systematic Reviews of Interventions,* Version 5.1.0 (updated March 2011) (Lefebvre 2011).

1. randomized controlled trial.pt.
2. controlled clinical trial.pt.
3. randomized.ab.
4. placebo.ab.
5. drug therapy.fs.
6. randomly.ab.
7. trial.ab.
8. groups.ab.
9. or/1-8
10. exp animals/ not humans.sh.
11. 9 not 10

Appendix 4. Embase Ovid search strategy

1. Thrombocyte rish plasma/
2. Fibrin/
3. (platelet$ adj5 (plasma$ or fibrin$ or concentrat$)).mp.
4. (PRP or L-PRP or PRF or L-PRF).ti,ab.
5. or/1-4
6. exp Periodontal disease/
7. periodont$.mp.
8. (infrabony or "infra bony" or intrabony or "intra bony" or infraosseous or "infra ossesous" or "[illegible]dosseous or apicomarginal or "apico marginal" or interproximal or "inter proximal").ti,ab.
9. ((vertical adj bone) and defect$).ti,ab.
10. (bone adj3 resorp$).ti,ab.
11. (intraalveolar or "intra alveolar").ti,ab.
12. or/6-11
13. 5 and 12

This subject search was linked to an adapted version of the Cochrane Embase [illegible]ject filt[illegible] for identifying RCTs in Embase Ovid (see www.cochranelibrary.com/help/central-creation-details.html for information).

1. Randomized controlled trial/
2. Controlled clinical study/
3. Random$.ti,ab.
4. randomization/
5. intermethod comparison/
6. placebo.ti,ab.
7. (compare or compared or comparison).ti.
8. ((evaluated or evaluate or evaluating or assessed or asses[illegible]and ([illegible]mpare or compared or comparing or comparison)).ab.
9. (open adj label).ti,ab.
10. ((double or single or doubly or singly) adj (blind or bl[illegible]ded or blindly)).ti,ab.
11. double blind procedure/
12. parallel group$1.ti,ab.
13. (crossover or cross over).ti,ab.
14. ((assign$ or match or matched or allo[illegible]atio[illegible]adj5 (alternate or group$1 or intervention$1 or patient$1 or subject$1 or participant$1)).ti,ab.
15. (assigned or allocated).ti,ab.
16. (controlled adj7 (study or d[illegible]sign or [illegible]rial)).ti,ab.
17. (volunteer or volunteers).ti,[illegible]
18. trial.ti.
19. or/1-18
20. (exp animal/ or animal[illegible]nw. or nonhuman/) not (exp human/ or human cell/ or (human or humans).ti.)
21. 19 not 20

Appendix 5. LILACS BIREME Virtual Health Library (Latin American and Caribbean Health Science Information database) search strategy

(Mh Platelet-Rich Plasma or "platelet rich plasma" or "Plasma Rico en Plaquetas" or "Plasma Rico em Plaquetas" or Mh Fibrin or fibrin$)
AND
periodont$

Appendix 6. US National Institutes of Health Ongoing Trials Register (ClinicalTrials.gov) search strategy

periodontal and platelet rich plasma
periodontal and fibrin

Appendix 7. World Health Organization International Clinical Trials Registry Platform search strategy

periodontal and platelet rich plasma
periodontal and fibrin

Appendix 8. Grey literature (www.greylit.org; www.opengrey.eu) search strategy

periodontal and platelet-rich plasma
periodontal and fibrin

CONTRIBUTIONS OF AUTHORS

Draft the protocol	Massimo Del Fabbro, Saurav Panda
Develop a search strategy	Saurav Panda, Surendar Ramamoorthi
Search for trials	Saurav Panda, Cristina Bucchi
Obtain copies of trials	Massimo Del Fabbro
Select which trials to include (two + one arbiter)	Jayakumar Nadathur Doraiswamy, Malaiappan Sankari, Massimo Del Fabbro
Extract data from trials (three people)	Lorena Karanxha, Saurav Panda, Cristina Bucchi
Enter data into Review Manager	Massimo Del Fabbro, Lorena Karanxha, Cristina Bucchi
Carry out the analysis	Massimo Del Fabbro, Lorena Karanxha, Saurav Panda, Sheeja Varghese, Cristina Bucchi
Interpret the analysis	Sheeja Varghese, Jayakumar Nadathur Doraiswamy, Silvio Taschieri
Draft the final review	Massimo Del Fabbro, Lorena Karanxha, Malaiappan Sankari, Cristina Bucchi
Update the review	Massimo Del Fabbro, Lorena Karanxha, Saurav Panda

DECLARATIONS OF INTEREST

The review authors declare they have no conflicts of interest.

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- Cochrane Oral Health Global Alliance, Other.

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DIFFERENCES BETWEEN PROTOCOL AND REVIEW

We did not perform a global comparison between the gro using autologous platelet concentrates and the control groups, because the differences in the surgical protocols amon subg oups were consistent, and preferred to directly perform subgroup analyses.

The primary outcomes of the protocol are the ondar outcomes in the review and vice versa. We did not consider other participant-reported outcomes (including preferenc ain and cost-effectiveness).