





Treatment of stage IV periodontitis: The EFP S3 level clinical practice guideline

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Abstract

Background: The recently published clinical practice guideline (CPG) for the treatment of periodontitis in stages I–III provided evidence-based recommendations for the treatment of periodontitis patients, defined according to the 2018 classification. Stage IV periodontitis shares the severity and complexity characteristics of stage III periodontitis, but includes the anatomical and functional sequelae of tooth and periodontal attachment loss (tooth flaring and drifting, bite collapse, etc.), which require additional interventions following completion of active periodontal therapy.

Aim: To develop an S3 Level CPG for the treatment of stage IV periodontitis, focusing on the implementation of inter-disciplinary treatment approaches required to treat/rehabilitate patients following associated sequelae and tooth loss.

Materials and Methods: This S3 Level CPG was developed by the European Federation of Periodontology (EFP), following methodological guidance from the Association of Scientific Medical Societies in Germany and the Grading of Recommendations Assessment, Development and Evaluation (GRADE) process. A rigorous and transparent process included synthesis of relevant research in 13 specifically commissioned systematic reviews, evaluation of the quality and strength of evidence, the formulation of specific recommendations and a structured consensus process with leading experts and a broad base of stakeholders.

Results: The S3 Level CPG for the treatment of stage IV periodontitis culminated in recommendations for different interventions, including orthodontic tooth movement,

tooth splinting, occlusal adjustment, tooth- or implant-supported fixed or removable dental prostheses and supportive periodontal care. Prior to treatment planning, it is critically important to undertake a definitive and comprehensive diagnosis and case evaluation, obtain relevant patient information, and engage in frequent re-evaluations during and after treatment. The periodontal component of therapy should follow the CPG for the treatment of periodontitis in stages I–III.

Conclusions: The present S3 Level CPG informs clinical practice, health systems, policymakers and, indirectly, the public on the available and most effective modalities to treat patients with stage IV periodontitis and to maintain a healthy dentition over lifetime, according to the available evidence at the time of publication.

KEYWORDS

clinical guideline, dental implant, orthodontic, stage IV, periodontitis, prosthodontic

Clinical Relevance

Scientific rationale for study: Patients with stage IV periodontitis share the severity and complexity features of stage III periodontitis but, in addition, suffer from the functional sequelae and consequences of tooth loss, frequently associated with advanced periodontitis. This clinical practice guideline (CPG) aimed to provide guidance on the necessary inter-disciplinary therapy required to rehabilitate the compromised dentition in such patients. The interventions described in this CPG should be derived following a rigorous evidence-based and patient-centred decision-making process.

Principal findings: This guideline was developed using strict and validated methodologies for assuring the best available evidence on the efficacy of the interventions considered, and the most appropriate recommendations based on a structured consensus process, including a panel of experts and representatives from key stakeholder groups.

Practical implications: The application of this S3 Level CPG will allow a consistent, inter-disciplinary and evidence-based approach to the management of stage IV periodontitis.

1 | INTRODUCTION

1.1 | The health problem

1.1.1 | Definition

Periodontitis is characterized by progressive destruction of the tooth-supporting apparatus (periodontium), with primary features being clinical attachment loss (CAL), radiographically assessed alveolar bone loss, presence of periodontal pockets and bleeding (Papapanou et al., 2018). Periodontitis is characterized further by defining the stage and grade of the disease: stage captures the severity, extent and distribution of the disease, as well as the anticipated complexity for its management; grade captures additional biological dimensions of the disease including the observed and/or anticipated progression rate, predicted treatment outcomes, and the risk that the disease or its treatment will adversely affect the general health of the patient (Papapanou et al., 2018; Tonetti et al., 2018).

The management of stages I–III periodontitis has already been described in a previously published clinical practice guideline (CPG) (Sanz, Herrera, et al., 2020). The present CPG focuses on the treatment

of stage IV periodontitis. The distinction between periodontitis stage III and stage IV is primarily based on the common sequelae of advanced loss of periodontal tissue support, which include: (i) tooth loss, resulting in <20 remaining teeth (<10 opposing pairs); (ii) masticatory dysfunction; (iii) tooth mobility grade ≥ 2 ; (iv) severe alveolar ridge defects; and (v) occlusal collapse (tooth drifting, flaring). These unique features of stage IV periodontitis require treatment of a higher level of complexity, but also necessitate an inter-disciplinary approach for rehabilitation of the impaired dentition (Papapanou et al., 2018; Tonetti et al., 2018; Tonetti & Sanz, 2019).

Periodontitis represents a major public health problem due to its high prevalence and associated morbidity. Severe periodontitis, in particular, may lead to disability due to impaired chewing function and aesthetics or edentulism, is a source of social inequality, and significantly impairs quality of life (Tonetti et al., 2017). In addition, severe periodontitis has a negative impact on general health and is associated with significant dental care costs (Tonetti et al., 2017).

A major difference between the treatment of periodontitis stages III and IV is the need for a stage IV patient to maintain/re-establish a functional dentition and the necessity for a rigorous supportive care

programme prior to, throughout, and following the rehabilitation phase of care. It has been shown that patients with stage IV periodontitis, when compared with stage I, have a higher risk of periodontal-related tooth loss over a period of follow-up of 10–30 years (hazard ratio 3.73) (Ravida et al., 2020) as well as a higher risk for pathological tooth migration and other functional consequences (Kwok & Caton, 2007).

1.1.2 | Pathophysiology

Periodontitis is initiated by the accumulation of a dental plaque biofilm at and below the gingival margin, which becomes increasingly dysbiotic (Meyle & Chapple, 2015) and which results in dysregulation of the host immune-inflammatory response, which further drives the dysbiosis and results in the destruction of periodontal tissues (Hajishengallis & Chavakis, 2021).

1.1.3 | Prevalence

Periodontitis is the most common chronic inflammatory non-communicable disease of humans. According to data originating from the Global Burden of Disease (GBD) database, 1.1 billion cases of severe periodontitis were prevalent globally in 2019, and an 8.44% (95% confidence interval—CI [6.62; 10.59]) increase in the age-standardized prevalence rate of severe periodontitis was observed between 1990 and 2019 (Chen et al., 2021).

In a Swedish population, the prevalence of severe masticatory dysfunction was found to be low (Salonen et al., 1990), but in a population of institutionalized older adults (65 years or more) in Italy, the prevalence of masticatory dysfunction, mainly caused by untreated edentulism, was 35% and it was associated with all-cause mortality after 9 years (Laudisio et al., 2016).

The prevalence (global age-standardized prevalence) of complete edentulism in 12-year or older individuals was 4.4% in 1990 and 2.4% in 2010, with incidences of 374 and 205 per 100,000 person-years cases, respectively, with no differences between sexes, and a gradual increase with age, peaking at 65 years (Kassebaum et al., 2014). Geographic differences were evident, with European countries presenting high incidences (Kassebaum et al., 2014). In the United States, data from NHANES 2005–2016 showed 4.5% complete edentulism with 10.3% of subjects lacking a functional dentition (i.e., having only one and up to 19 remaining teeth) (Al-Zahrani et al., 2021). Complete and partial edentulism were more frequent in persons with systemic comorbidities or with a compromised systemic condition (Parker et al., 2020). The negative impact of edentulism on quality of life, nutrition and systemic health has been well established (Griffin et al., 2012).

1.1.4 | Consequences of failure to treat

If no treatment is provided for stage IV periodontitis, or if the treatment is inadequate and/or not comprehensive (e.g., does not result in

sufficient rehabilitation or correction of the masticatory dysfunction), the risk of additional loss of tooth-supporting tissues increases and may lead to complete edentulism. Untreated severe periodontitis may result in substantial tooth loss in adults (Ramseier et al., 2017) and ranks 77th among the 100 most relevant human conditions resulting in disability (Marcenes et al., 2013).

Severe periodontitis and dental caries account for more years lost to disability than any other human disease (GBD 2017 Disease and Injury Incidence and Prevalence Collaborators, 2018).

Furthermore, periodontitis is associated with a range of systemic diseases including diabetes (Sanz et al., 2018), cardiovascular diseases (Tonetti, Van Dyke, & Working group 1 of the joint EFP/AAP Workshop, 2013; Sanz, Marco Del Castillo, et al., 2020) and adverse pregnancy outcomes (Sanz, Komman, & Working group 3 of joint EFP/AAP Workshop, 2013). It is also independently associated with premature death from all causes or cardiovascular disease (Garcia et al., 1998; Soikkonen et al., 2000; Soder et al., 2007; Linden et al., 2012), in particular in multi-morbid populations where the impact of periodontitis is equivalent to having co-morbid diabetes mellitus (Sharma et al., 2016), and also results in increased medical expenditure (Sato et al., 2016).

1.1.5 | Financial aspects

On a global scale, periodontitis of all stages is estimated to cost \$54 billion in direct treatment costs and further \$25 billion in indirect costs (GBD 2017 Disease and Injury Incidence and Prevalence Collaborators, 2018). Periodontitis, and especially severe periodontitis (including stages III and IV periodontitis), contributes significantly to the total expenditure allocated to manage dental diseases due to the need to replace lost teeth. In 2015, the total cost of dental diseases was estimated to amount to \$544.41 billion, run on billion being \$356.80 billion direct costs (dental expenditures), and \$187.61 billion indirect costs (productivity losses) (Righolt et al., 2018).

Although the economic impact of edentulism has not been clearly established, at least two factors may support its importance: on the one hand, the need of rehabilitation; on the other hand, and in case of lack of rehabilitation, the negative consequences already listed in quality of life, nutrition, systemic health, etc. In addition, it has also been concluded that individual- and community-level social inequalities have a strong impact on edentulism (Ito et al., 2015).

2 | AIM OF THE GUIDELINE

This guideline aims to highlight the importance of, and need for scientific evidence in, clinical decision-making in the treatment of patients with stage IV periodontitis. Its main objective, therefore, is to summarize the evidence-based recommendations for the individual interventions involved in the multidisciplinary management of stage IV periodontitis, based on the best available evidence and/or expert consensus. In so doing, this guideline aims to (i) inform sound multidisciplinary therapeutic approaches to the treatment of stage IV periodontitis, and thereby improve the overall quality of periodontal treatment rendered in Europe

and worldwide, (ii) reduce tooth loss associated with periodontitis, and ultimately (iii) improve overall systemic health and quality of life.

2.1 | Target users of the guideline

Dental professionals, together with stakeholders related to oral health care, including patients. In addition, this CPG aims to inform medical professions, health systems, policymakers, patients and the public.

2.2 | Targeted environments

Academic/hospital environments, community-based dental clinics and practices were the targeted environments for this guideline.

2.3 | Targeted patient population

Patients were selected based on the following criteria:

- I. People with stage IV periodontitis.
- II. People with stage IV periodontitis, following successful periodontal treatment.
- III. People with stage IV periodontitis, following successful periodontal and multidisciplinary treatment.

2.4 | Exceptions from the guideline

This guideline does not consider in detail the health/economic cost-benefit ratio of the proposed therapies, since (i) the target users and patient populations include people in different countries with diverse, not readily comparable healthcare systems, and (ii) there is a paucity of sound scientific data available addressing this issue.

This guideline does not consider the treatment of gingivitis (although management of gingivitis is included as an indirect goal in a number of proposed interventions), the treatment of stages I–III periodontitis (already covered in a previously published guideline; Sanz, Herrera, et al., 2020), necrotising periodontitis (Herrera et al., 2018; Papapanou et al., 2018), periodontitis as manifestation of systemic diseases and mucogingival conditions (Jepsen et al., 2018). However, we emphasize that (i) treatment of gingivitis is a primary prevention strategy for periodontitis (Chapple et al., 2015), and (ii) maintenance of stable periodontal tissues requires the control of gingival inflammation (Chapple et al., 2018).

3 | METHODOLOGY

3.1 | General framework

This guideline was developed following methodological guidance published by the Standing Guideline Commission of the Association of

Scientific Medical Societies in Germany (AWMF) (<https://www.awmf.org/leitlinien/awmf-regelwerk/awmf-guidance.html>) and the Grading of Recommendations Assessment, Development and Evaluation (GRADE) working group (<https://www.gradeworkinggroup.org/>).

The guideline was developed under the auspices of the European Federation of Periodontology (EFP) and overseen by the EFP Workshop Committee. This guideline development process was steered by an organizing committee and a methodology consultant designated by the EFP. All members of the organizing committee participated in the EFP workshop committee.

To ensure adequate stakeholder involvement, the EFP established a guideline panel involving dental professionals representing national periodontal societies within the EFP, together with experts in orthodontics, prosthodontics, implant dentistry, oral surgery and oral diagnosis (Table 1). These delegates were nominated and selected by the organizing committee and participated in the guideline development process with voting rights in the consensus conference. For the guideline development process, delegates were assigned to four working groups that were chaired by selected members of the organizing committee and guided by the methodology consultant. This panel was supported by key stakeholders from European scientific societies with a strong professional interest in periodontal care and from European organizations representing key groups within the dental profession (Table 2), and key experts from non-EFP member regions, such as North America and Australia.

In addition, the EFP engaged an independent guideline methodologist to advise the panel and facilitate the consensus process (Prof. Dr. med. Ina Kopp [I.K.]). The guideline methodologist had no voting rights.

The EFP and the guideline panel attempted to involve patient forums/organizations but were unable to identify any groups focused on periodontal diseases at a pan-European level. In a future update, efforts will be undertaken to include the perspectives of citizens/patients (Brocklehurst et al., 2018), and national societies will be encouraged to involve patient groups within individual countries, as key stakeholders for the *Adaptation, Adoption, De Novo Development*—“ADOLOPMENT” of this GPG (Schünemann et al., 2017).

3.2 | Evidence synthesis

3.2.1 | Systematic search and critical appraisal of guidelines

To assess and utilize existing guidelines during the development of the present guideline, we performed electronic searches in a range of well-established guideline registers and the websites of large periodontal societies:

- Guideline International Network (GIN)
- Guidelinecentral.com
- The National Institute for Health and Clinical Excellence (NICE)
- Canadian Health Technology Assessment (CADTH)

TABLE 1 Guideline panel

Scientific society/organization	Delegate(s)
European Federation of Periodontology (EFP) *Selected experts not belonging to EFP (experts in orthodontics, prosthodontics, implant dentistry, oral surgery, oral diagnosis)	Organizing Committee, Working Group Chairs (in alphabetic order): Tord Berglundh, Iain Chapple, David Herrera, Søren Jepsen, Moritz Kepschull, Panos Papapanou, Mariano Sanz, Anton Sculean, Maurizio Tonetti Methodologists: Ina Kopp Clinical Experts (in alphabetic order): Mario Aimetti Bilal Al-Nawas* Juan Blanco Philippe Bouchard Maria Clotilde Carra Tali Chackartchi Francesco D'Aiuto Bettina Dannewitz Monique Danser Jan Derks Thomas Dietrich Henrik Dommisch Nikos Donos Elena Figuero Moshe Goldstein Marjolaine Gosset Filippo Graziani Lisa Heitz-Mayfield Karin Jepsen Ronald Jung Dimitrios Kloukos* Bahar Eren Kuru France Lambert Luca Landi Natalie Leow Rodrigo López Phoebus Madianos Conchita Martín* Paula Matesanz Ana Molina Virginie Monnet Corti Eduardo Montero Ian Needleman Luigi Nibali Spyridon N. Papageorgiou* Guillermo Pradies* Marc Quirynen Christoph Ramseier Stefan Renvert Mario Rocuzzo Irena Sailer* Giovanni Salvi Nerea Sánchez Ignacio Sanz-Sánchez Frank Schwarz Falk Schwendicke* Lior Shapira Andreas Stavropoulos Jean Suvan Wim Teughels Cristiano Tomasi Leonardo Trombelli Katleen Vandamme* Gernot Wimmer Stefan Wolfart* Nicola Zitzmann

(Continues)

TABLE 1 (Continued)

Scientific society/organization	Delegate(s)
Scientific societies	
European Association for Osseointegration	Henning Schliephake
European Federation of Conservative Dentistry	Sebastian Paris
European Federation of Periodontology— <i>Executive Committee</i>	Xavier Struillou
European Federation of Periodontology— <i>Executive Committee</i>	Nicola West
European Prosthodontic Association	Marco Ferrari
European Society of Endodontology	Lise Lotte Kirkevang
Other organizations	
Council of European Chief Dental Officers	Kenneth Eaton
Platform for Better Oral Health in Europe	Kenneth Eaton
Council of European Dentists	Paulo Melo
European Dental Students' Association	Tin Crnić

TABLE 2 Key stakeholders contacted and participants

Institution/society	Acronym	Answer ^a	Representative
Association for Dental Education in Europe	ADEE	No proposal	None
Continental European Division of IADR	CED-IADR	No proposal	None
Council of European Chief Dental Officers	CECDO	Participant	Kenneth Eaton
Council of European Dentists	CED	Participant	Paulo Melo
European Association for Osseointegration	EAO	Participant	Henning Schliephake
European Association of Dental Public Health	EADPH	No answer	None
European Dental Hygienists Federation	EDHF	No proposal	None
European Dental Students' Association	EDSA	Participant	Tin Crnić
European Federation of Conservative Dentistry	EFCD	Participant	Sebastian Paris
European Orthodontic Society	EOS	No answer	None
European Prosthodontic Association	EPA	Participant	Marco Ferrari
European Society of Endodontology	ESE	Participant	Lise Lotte Kirkevang
Platform for Better Oral Health in Europe	PBOHE	Participant	Kenneth Eaton

^aMessages sent on 9 April 2020; reminder sent on June 2020.

- European Federation for Periodontology (EFP)
- American Academy of Periodontology (AAP)
- American Dental Association (ADA)

The last search was performed on 1 November 2021. Search terms used were: “periodont*” “Periodontal”, “Guidelines”, “Clinical Practice Guidelines”. In addition, content was screened by hand searches, see Table 3.

Only guidelines published in English and with full texts available were included. The methodological quality of these guideline texts was critically appraised using the AGREE II framework (<https://www.agreetrust.org/agree-ii/>).

We did not identify guidelines/documents directly relevant to the current guideline development process due to: (i) their publication time, which frequently predated the workshop that defined stage IV

periodontitis; (ii) their methodological approach; or (iii) their stated inclusion criteria. We have referenced the EFP S3-Level clinical practice guideline (Sanz, Herrera, et al., 2020), where applicable.

3.2.2 | Systematic search and critical appraisal of the literature

For this guideline, a total of 13 systematic reviews (SRs) were conducted to support the guideline development process (Carra et al., 2021; Dommisch et al., 2021; Donos et al., 2021; Gennai et al., 2021; Gotfredsen et al., 2021; Kloukos et al., 2021; Leow et al., 2021; Martín et al., 2021; Montero et al., 2021; Orlandi et al., 2021; Papageorgiou et al., 2021; Ramanauskaitė et al., 2021; Tomasi et al., 2021). The corresponding manuscripts are published

TABLE 3 Results of the guideline search

Database	Identified, potentially relevant guidelines	Critical appraisal
Guideline International Network (GIN) International Guidelines Library ^a	Comprehensive periodontal therapy: a statement by the American Academy of Periodontology. American Academy of Periodontology. NGC:008726 (2011)	8 years old, recommendations not based on systematic evaluation of evidence, <i>not applicable</i>
	HealthPartners Dental Group and Clinics guidelines for the diagnosis and treatment of periodontal diseases. HealthPartners Dental Group. NGC:008848 (2011)	8 years old, unclear methodology, <i>not applicable</i>
Guidelinecentral.com “Dentistry” category	Health Partners Dental Group and Clinics Caries Guideline	<i>Not applicable</i>
The National Institute for Health and Clinical Excellence (NICE) ^b	No thematically relevant hits	<i>Not applicable</i>
National Guideline Clearinghouse (Agency for Healthcare Research and Quality) ^c	No thematically relevant hits	<i>Not applicable</i>
Canadian Health Technology Assessment (CADTH) ^d	Periodontal Regenerative Procedures for Patients with Periodontal Disease: A Review of Clinical Effectiveness (2010)	9-year-old review article, <i>not applicable</i>
	Treatment of Periodontal Disease: Guidelines and Impact (2010)	9-year-old review article, <i>not applicable</i>
	Dental Scaling and Root Planing for Periodontal Health: A Review of the Clinical Effectiveness, Cost-effectiveness, and Guidelines (2016)	Unclear methodology (follow-up, outcome variables, recommendations, guideline group), <i>not applicable</i>
	Dental Cleaning and Polishing for Oral Health: A Review of the Clinical Effectiveness, Cost-effectiveness and Guidelines (2013)	Unclear methodology (follow-up, outcome variables, recommendations, guideline group), <i>not applicable</i>
European Federation of Periodontology (EFP) ^e	EFP S3-Level CPG for stage I–III	<i>Indirectly applicable, high quality</i>
American Academy of Periodontology (AAP) ^f	The American Journal of Cardiology and Journal of Periodontology Editors' Consensus: Periodontitis and Atherosclerotic Vascular Disease (2009)	Unclear methodology, 10-year-old consensus-based article, only limited clinically applicable recommendations, <i>not applicable</i>
	Comprehensive Periodontal Therapy: A Statement by the American Academy of Periodontology (2011)	Unclear methodology (follow-up, outcome variables, recommendations, guideline group), almost a decade old, <i>not applicable</i>
	Academy Statements on Gingival Curettage (2002), Local Delivery (2006), Risk Assessment (2008), Efficacy of Lasers (2011)	Unclear methodology, 10-year-old consensus-based article, only limited clinically applicable recommendations, <i>not applicable</i>
American Dental Association (ADA) ^g	Nonsurgical Treatment of Chronic Periodontitis Guideline (2015)	Outcome variable CAL (not PPD), no minimal follow-up— <i>not applicable</i>

^a<https://guidelines.ebmportal.com/>.^b<https://www.nice.org.uk/guidance/published?type=csg,mg,ph,sg,sc>.^c<https://www.ahrq.gov/gam/index.html>.^d<https://www.cadth.ca/>.^e<http://www.efp.org/publications/index.html>.^f<https://www.perio.org/publications>.^g<https://ebd.ada.org/en/evidence/guidelines>.

within this special issue of the Journal of Clinical Periodontology. The SRs were updated in July–September 2021, and the reports of the updates are presented as an Addendum to the present CPG. The Addendum is accessible online (CPGstage4-Addendum).

All SRs were conducted following the “Preferred Reporting Items for Systematic Reviews and Meta-Analyses” (PRISMA) framework (Moher et al., 2009), and were prospectively registered in PROSPERO.

3.2.3 | Focused questions

In all 13 SRs, focused questions in PICOS format (Centre for reviews and dissemination, 2008; Guyatt et al., 2011) were proposed by the authors in January 2019 to a panel comprising the working group chairs and the methodological consultant in order to review and approve them (Table 4a–d). The panel took great care to avoid

TABLE 4 PICOS questions addressed by each systematic review, listed according to working group: (a) treatment of pathological tooth migration in stage IV periodontitis patients; (b) treatment of tooth loss/masticatory dysfunction/bite collapse in stage IV periodontitis patients—partial edentulism amenable for partial rehabilitation; (c) treatment of tooth loss/masticatory dysfunction/bite collapse in stage IV periodontitis patients with terminal dentition only amenable for full-arch rehabilitation; (d) long-term outcomes and impact of treatment in stage IV periodontitis patients

Reference	Systematic review title	Final PICOS question (as written in manuscripts)
(a)		
Martín et al. (2021)	Effect of orthodontic therapy in periodontitis and non-periodontitis patients: a systematic review with meta-analysis.	<p>#1: In adult patients with malocclusion (population), what are the effects of OTM on clinical attachment level (CAL) changes (outcome) in treated periodontitis patients with a healthy but reduced periodontium (exposure) compared with non-periodontitis patients (comparator)?</p> <p>#2: In adult patients with malocclusion and healthy but reduced periodontium (population), what is the efficacy of skeletal anchorage devices (implants or TADs—micro-screws or mini-plates—) (intervention) compared with conventional anchorage systems (comparator), in terms of orthodontic treatment (outcomes)?</p>
Papageorgiou et al. (2021)	Effect of periodontal–orthodontic treatment of teeth with pathological tooth flaring, drifting and elongation in patients with severe periodontitis: a systematic review with meta-analysis.	What is the influence of periodontal–orthodontic treatment of pathologically migrated teeth in patients with severe periodontitis on the periodontal status?
Kloukos et al. (2021)	Effect of combined periodontal and orthodontic treatment of tilted molars and of teeth with intra-bony and furcation defects in stage IV periodontitis patients. A systematic review.	<p>#1. In periodontitis patients with treated tilted molars, what is the effect of orthodontic treatment, as compared with no treatment, in terms of changes in PPD and CAL?</p> <p>#2. In periodontitis patients with treated intra-bony defects, what is the effect of orthodontic treatment, as compared with no treatment, in terms of changes in PPD and CAL?</p> <p>#3. In periodontitis patients with treated furcation defects, what is the effect of orthodontic treatment, as compared with no treatment, in terms of changes in PPD and CAL?</p>
(b)		
Dommisch et al. (2021)	Efficacy of tooth splinting and occlusal equilibration in patients with periodontitis exhibiting masticatory dysfunction—A systematic review.	What is the benefit of (I) tooth splinting (TS) or occlusal equilibration (OE) of teeth with adaptive and progressive mobility during non-surgical and surgical periodontal therapy in (P) patients with periodontitis exhibiting masticatory dysfunction when compared with (C) non-splinted teeth with adaptive and progressive mobility or no-TS within the same periodontitis patient at diseased sites and no OE with respect to (O) tooth loss (primary outcome parameter), and change in PPD, CAL change, mobility, and patient-reported outcome measures (PROMs) (secondary outcome parameters) for a follow-up of ≥12 months evidenced as shown by randomized clinical controlled trials, clinical controlled trials, retrospective and prospective case-control studies, and case series?
Gotfredsen et al. (2021)	Efficacy and risks of removable prosthesis in periodontitis patients. A systematic review.	In partially edentulous periodontitis patients (P), are removable dental prostheses (I), in comparison with no prosthetic treatment or treatment to a shortened dental arch, or with fixed dental prosthesis and comparison between different RDP designs (C), more efficacious in terms of tooth loss, periodontal parameters, mastication/chewing efficiency, and patient-related outcome measures (O), as evidenced in randomized controlled clinical trials (RCTs) or prospective and retrospective cohort controlled studies and with a follow-up period of ≥1 year (S)?
Montero et al. (2021)	Efficacy and risks of tooth-supported prostheses in the treatment of partially edentulous patients with stage IV periodontitis. A systematic review and meta-analysis.	#1. In partially edentulous patients (population), what is the efficacy of multi-unit tooth-supported fixed prostheses in patients with stage IV periodontitis, as compared with non-periodontitis patients (intervention and comparison), in terms of survival rate of teeth used as abutments (primary outcome), in RCTs (study design) with at least 12 months of follow-up?

(Continues)

TABLE 4 (Continued)

Reference	Systematic review title	Final PICOS question (as written in manuscripts)
Carra et al. (2021)	Effectiveness of implant-supported fixed partial denture in patients with history of periodontitis: a systematic review and meta-analysis.	<p>#2. In partially edentulous patients diagnosed with stage IV periodontitis (population), what is the effectiveness of multi-unit tooth-supported fixed prostheses (intervention and comparison), in terms of survival rate of teeth used as abutments (primary outcome), in RCTs, CCTs, prospective/retrospective cohort studies or prospective/retrospective case series (CS) with a minimum follow-up time of 12 months?</p> <p>What is the effectiveness (i.e., survival) and risks (i.e., biological and mechanical complications) of IS-FPD in patients with a history of periodontitis compared with patients with no history of periodontitis at >=1 year from implant loading?</p>
(c)		
Donos et al. (2021)	Efficacy of tooth-supported compared with implant-supported full-arch removable prostheses in patients with terminal dentition. A systematic review.	<p>#1. In patients with terminal dentition and/or stage IV periodontitis, what is the efficacy of tooth-supported (TSRP) compared with implant-supported (ISRP) full-arch removable prostheses in terms of survival rate of the implants/teeth and survival of the prosthesis, as reported in studies with at least 1 year of follow-up post prosthesis delivery?</p> <p>#2. In patients with terminal dentition due to stage IV periodontitis, what is the estimated cumulative survival of teeth/implants and prostheses in case of tooth-supported (TSRP) and in case of implant-supported (ISRP) full-arch removable prostheses, as reported in studies with at least 1 year of follow-up post prosthesis delivery?</p>
Tomasi et al. (2021)	Efficacy of rehabilitation of stage IV periodontitis patients with full-arch fixed prostheses: tooth-supported versus implant-supported. A systematic review.	<p>#1. In patients with a periodontally compromised dentition (due to stage IV periodontitis or equivalent), what is the evidence from controlled studies with a minimum follow-up of 1 year that implant-supported full-arch fixed prostheses are more efficacious than tooth-supported full-arch fixed prostheses in terms of survival (of restorations and supportive units) and complications?</p> <p>#2. In patients with a periodontally compromised dentition (due to stage IV periodontitis or equivalent), what is the performance of tooth-supported full-arch fixed prostheses as reported in interventional or observational studies with a minimum follow-up of 1 year?</p> <p>#3. In patients with a periodontally compromised dentition (due to stage IV periodontitis or equivalent), what is the performance of implant-supported full-arch fixed prostheses following extraction of the remaining teeth as reported in interventional or observational studies with a minimum follow-up of 1 year?</p>
Ramanauskaite et al. (2021)	Efficacy of rehabilitation with different approaches of implant-supported full-arch prosthetic designs: A systematic review.	In patients with at least one edentulous jaw, with tooth loss mainly due to periodontitis (Population), what is the efficacy of different types of rehabilitation with fixed or removable full-arch implant-supported prosthesis designs (Intervention and Comparison), in terms of implant loss and success rates (Outcome), as reported in prospective and retrospective observational one-arm and case series, randomized and non-randomized controlled clinical trials (Study design)?
(d)		
Leow et al. (2021)	Recurrence and progression of periodontitis and methods of management in long-term care. A systematic review and meta-analysis.	<p>#1. In people treated for periodontitis and in supportive maintenance care for 5 years or more, compared with no supportive maintenance care (SPC), how common is recurrence of the condition?</p> <p>#2. In people experiencing recurrence of periodontitis, what is the effect of different methods of treatment on the recurrence as assessed by measures of health, quality of life, cost and accessibility of care and harms?</p>
Orlandi et al. (2021)	In patients with severe periodontitis what is the effect of periodontal treatment on	#1. In patients with severe periodontitis (stages III or IV or equivalent) who are otherwise healthy, what is the effect of periodontal treatment in comparison with no treatment or control treatment, in

(Continues)

TABLE 4 (Continued)

Reference	Systematic review title	Final PICOS question (as written in manuscripts)
	systemic disease risk and adverse pregnancy outcomes?	terms of systemic health and quality of life outcomes, as reported in 6 month (minimum follow-up) randomized controlled trials?
		#2. In patients with periodontitis (stages III or IV or equivalent) and a non-communicable disease, what is the effect of periodontal treatment in comparison with no treatment or control treatment, in terms of systemic health and quality of life outcomes, as reported in 6 months minimum follow-up randomized controlled trials?
		#3. In patients with periodontitis (stages III or IV or equivalent) and pregnancy, what is the effect of periodontal treatment in comparison with no treatment or control treatment, in terms of perinatal, maternal and quality of life outcomes, as reported in randomized controlled trials?
Gennai et al. (2021)	Impact of rehabilitation versus edentulism in systemic health and quality of life in patients affected by periodontitis. A systematic review and meta-analysis.	What is the effect of bridges or dentures versus no treatment in fully or partially edentulous patients affected by stage IV periodontitis in terms of quality of life (as measured through psychometric testing) and systemic health (as measured through general disease's incidence and surrogate markers), as reported in randomized and non-randomized controlled clinical trials, case series, cohort studies, cross-sectional studies and case-control studies?

overlaps between the SRs or significant thematic omissions, in order to ensure that they encompass the main interventions currently undertaken in the management of stage IV periodontitis. Since the criteria to define stage IV periodontitis were only available after 2018, in order to include, in the SRs, those studies published/conducted before 2018, the terminology “stage IV (or equivalent) periodontitis” has been used, when applicable.

3.2.4 | Relevance of outcomes

One important outcome of interest in a clinical guideline in Dentistry is tooth loss. However, a narrative review article (Loos & Needleman, 2020), commissioned during the development of the CPG for the treatment of stage I–III periodontitis (Sanz, Herrera, et al., 2020) reported that the reduction in periodontal probing pocket depth (PPD) was an important predictor of tooth loss, in the context of periodontal therapy. Therefore, PPD reduction was also used as a primary outcome in SRs not addressing periodontal regeneration, and in instances where tooth loss data were not available. When reviewing periodontal regenerative interventions, gains in clinical attachment were used as the primary outcome measure. Secondary periodontal outcomes included the proportion of residual/open pockets as critical endpoint of therapy, which have been shown to be associated with disease recurrence (Loos & Needleman, 2020).

For the purposes of the present guideline, in addition to tooth survival, tooth-supported prosthesis survival and implant and implant-supported prosthesis survival were also considered. Additional outcomes related to complexity factors were also addressed in the SRs (e.g., vertical dimension or evaluation of masticatory dysfunction). Finally, patient-reported outcome measures (PROMs), quality of life indicators and economic factors were studied whenever possible.

3.2.5 | Search strategy

All SRs utilized a comprehensive search strategy of at least two different databases, supplemented by a hand search of periodontology-focused journals and the reference lists of included studies. In all SRs, the electronic and manual search, as well as the data extraction, was undertaken in parallel by, at least, two different investigators.

3.2.6 | Quality assessment of included studies

In all SRs, the risk of bias of controlled clinical trials was assessed using the Cochrane risk-of bias tool (<https://methods.cochrane.org/bias/resources/rob-2-revised-cochrane-risk-bias-tool-randomized-trials>). For observational studies, the Newcastle–Ottawa scale was used (http://www.ohri.ca/programs/clinical_epidemiology/oxford.asp).

3.2.7 | Data synthesis

Where applicable, the available evidence was summarized by means of a meta-analysis.

3.3 | From evidence to recommendation: Structured consensus process

The structured consensus development conference was held during the XVII European Workshop in Periodontology in La Granja de San Ildefonso Segovia, Spain, on 7–9 November 2021. Using the 13 SRs as background information, evidence-based recommendations were

formally debated by the guideline panel using the format of a structured consensus development conference. This consisted of small group discussions and open plenary discussions, where the proposed recommendations were presented, voted upon and adopted by consensus (Murphy et al., 1998). Delegates declaring potential conflicts of interest abstained from voting and abstentions were recorded. Prior to the in-person meeting, up to four online meetings were organized (one at the plenary level, and three at the working group level) in July, September and October 2021, in order to advance the process of guideline development to a mature stage prior to the face-to-face consensus meeting.

In the small group phase, delegates convened in four working groups (WGs) directed by two chairpersons belonging to the EFP Workshop Committee, addressing the following subtopics:

- WG 1. Treatment of pathological tooth migration in stage IV periodontitis patients (chairs Søren Jepsen and Mariano Sanz).
- WG 2. Treatment of tooth loss/masticatory dysfunction/bite collapse in stage IV periodontitis patients—partial edentulism amenable for partial rehabilitation (chairs Moritz Kebschull and Anton Sculean).
- WG 3. Treatment of tooth loss/masticatory dysfunction/bite collapse in stage IV periodontitis patients with a terminal dentition only amenable for full-arch rehabilitation (chairs Tord Berglundh, Panos Papapanou and Maurizio Tonetti).
- WG 4. Long-term outcomes and impact of treatment in stage IV periodontitis patients (chairs Iain Chapple and David Herrera).

With the support of the methodology expert, recommendations and draft background texts were generated and subsequently presented, debated and subjected to a vote in the plenary sessions with all delegates present. During these plenary sessions, the guideline development process and discussions and votes were overseen and facilitated by the independent guideline methodologist (I.K.). The plenary votes were recorded using an electronic voting system, checked for accuracy and then introduced into the guideline text.

The consensus process was conducted as follows:

3.3.1 | Plenary session 1 (online session, July 2021)

Introduction to guideline methodology (presentation, discussion) by the independent guideline methodologist (I.K.) and the chair of the workshop (D.H).

3.3.2 | Working group phase 1 (three online sessions, from July to October 2021)

- Peer evaluation of declarations of interest and management of conflicts of interest.
- Presentation of the evidence (SR results) by group chairs and reviewers.

- Invitation of all members of the working group to reflect critically on the quality of available evidence by group chairs, considering the GRADE criteria.
- Structured group discussions:
 - development of draft recommendations and their grading, considering the GRADE criteria.
 - development of draft background texts, considering the GRADE criteria.
 - invitation to comment on draft recommendations and background text to suggest reasonable amendments by group chairs.
 - collection and merging of amendments by group chairs.

3.3.3 | Plenary session 2 (in-person meeting, November 2021)

- Presentation of working group results (draft recommendations and background text) by working group chairs.
- Invitation to formulate questions, statements and reasonable amendments of the plenum by the independent guideline methodologist/facilitator.
- Answering of questions by working group chairs.
- Collection and merging of amendments by an independent moderator.
- Preliminary vote on all suggestions provided by the WGs and all reasonable amendments.
- Assessment of the strength of consensus.
- Recording of abstentions made due to potential conflicts of interest.
- Opening debate, where no consensus was reached or reasonable need for discussion was identified.
- Formulation of tasks to be solved within the WGs.

3.3.4 | Working group phase 2 (in-person meeting, November 2021)

- Discussion of tasks and potential amendments raised by the plenum.
- Formulation of reasonable and justifiable amendments, considering the GRADE framework.
- Initial voting within the working group on recommendations and guideline text in preparation for the plenary session.

3.3.5 | Plenary session 3 (in-person meeting, November 2021)

- Presentation of working group results by working group chairpersons.
- Invitation to formulate questions, statements and reasonable amendments of the plenary by the independent moderator.

- Collection and merging of amendments by an independent moderator.
- Preliminary vote.
- Assessment of the strength of consensus.
- Opening debate, where no consensus was reached or reasonable need for discussion was identified.
- Formulation of reasonable alternatives.
- Final vote of each recommendation, recording the consensus and abstentions due to potential conflicts of interest.

3.3.6 | Plenary session 4 (online meeting, January 2022)

- Presentation of pending recommendations and suggestions received.
- Preliminary vote.
- Assessment of the strength of consensus.
- Opening debate, where no consensus was reached or reasonable need for discussion was identified.
- Formulation of reasonable alternatives.
- Final vote of each recommendation, recording abstentions due to potential conflicts of interest.

3.4 | Definitions: Rating the quality of evidence, grading the strength of recommendations and determining the strength of consensus

For all recommendations and statements, this guideline makes transparent:

- the underlying quality of evidence, reflecting the degree of certainty/uncertainty of the evidence and robustness of study results;
- the grade of the recommendation, reflecting the criteria considered to make the judgement; the strength of consensus, indicating the degree of agreement within the guideline panel; the number of abstentions due to potential conflicts of interest.

3.4.1 | Quality of evidence

The quality of evidence was assessed using a recommended rating scheme (Balslem et al., 2011; Schunemann, Zhang, Oxman, & Expert Evidence in Guidelines, 2019).

3.4.2 | Strength of recommendations

The grading of the recommendations used the grading scheme (Table 5) by the German Association of the Scientific Medical

TABLE 5 Strength of recommendations: Grading scheme (German Association of the Scientific Medical Societies (AWMF) & Standing Guidelines Commission, 2012)

Grade of recommendation grade ^a	Description	Syntax
A	Strong recommendation	We recommend (↑↑)/We recommend not to (↓↓)
B	Recommendation	We suggest to (↑)/We suggest not to (↓)
0	Open recommendation	May be considered (↔)

^aIf the group felt that evidence was not clear enough to support a recommendation, Statements were formulated, including the need (or not) of additional research.

Societies (AWMF) and Standing Guidelines Commission (2012), taking into account not only the quality of evidence, but also considering a judgement guided by the following criteria:

- relevance of outcomes and quality of evidence for each relevant outcome;
- consistency of study results;
- direct applicability of the evidence to the target population/PICOS specifics;
- precision of effect estimates using confidence intervals;
- magnitude of the effects;
- balance of benefit and harm;
- ethical, legal, economic considerations;
- patient preferences.

The grading of the quality of evidence and the strength of a recommendation may therefore differ, but where they do, the justification and context are clearly documented in the background narrative that follows each recommendation table.

3.4.3 | Strength of consensus

The consensus determination process followed the recommendations by the German Association of the Scientific Medical Societies (AWMF) and Standing Guidelines Commission (2012). Where consensus could not be reached, different points of view were documented in the guideline text (see Table 6).

3.5 | Editorial independence

3.5.1 | Funding of the guideline

The development of this guideline and its subsequent publication was financed entirely by internal funds of the EFP, without any support from industry or other organizations.

TABLE 6 Strength of consensus: Determination scheme (German Association of the Scientific Medical Societies (AWMF) & Standing Guidelines Commission, 2012)

Unanimous consensus	Agreement of 100% of participants
Strong consensus	Agreement of >95% of participants
Consensus	Agreement of 75%–95% of participants
Simple majority	Agreement of 50%–74% of participants
No consensus	Agreement of <50% of participants

3.5.2 | Declaration of interests and management of potential conflicts

All members of the guideline panel declared secondary interests using the standardized form provided by the International Committee of Medical Journal Editors (ICMJE) (International Committee of Medical Editors, 2013).

Management of conflicts of interests (Cols) was discussed in the WGs and the plenary sessions, following the principles provided by the Guidelines International Network (Schunemann et al., 2015). According to these principles, panel members with relevant, potential Cols abstained from voting on guideline statements and recommendations within the consensus process. Those abstentions were recorded in each recommendation table.

3.6 | Peer review

All 13 SRs underwent a multi-step peer review process. First, the draft documents were evaluated by members of the EFP Workshop Committee and the methodological consultants using a custom-made appraisal tool to assess: (i) the methodological quality of the SRs using the AMSTAR 2 checklist (Shea et al., 2017); and (ii) whether all PICOS questions were addressed as planned. Detailed feedback was then provided to the SR authors. Subsequently, all 13 SRs underwent the regular editorial peer review process defined by the *Journal of Clinical Periodontology*. In addition, the reports of the updates of the SRs, presented in the Addendum (accessible online *CPGstage4-Addendum*), were also peer-reviewed.

The guideline text was drafted by the chairs of the WGs, in close cooperation with the methodological consultant, and circulated among the members of the guideline group prior to the workshop. The methodological quality was formally assessed by an external consultant using the AGREE framework. The guideline was subsequently peer-reviewed for its publication in the *Journal of Clinical Periodontology* following the standard evaluation process of the journal.

3.7 | Implementation and dissemination plan

For this guideline, a multi-stage dissemination and implementation strategy will be established and implemented by the EFP, supported by a communication campaign.

This will include:

- Publication of the guideline and the underlying SRs as an Open Access special issue of the *Journal of Clinical Periodontology*.
- Commentary, Adoption, or Adaptation (Schünemann et al., 2017) by national societies.
- Generation of educational material for dental professionals and patients, and dissemination via the EFP member societies.
- Dissemination via educational programmes at dental conferences.
- Dissemination via the EFP through European stakeholders via National Society members of the EFP.
- Long-term evaluation of the successful implementation of the guideline by a survey of EFP members.

The timeline of the guideline development process is detailed in Table 7.

3.8 | Validity and update process

The guideline is valid until 2027. However, the EFP, represented by the members of the organizing committee, will continuously assess current developments in the field. Where there are major changes of circumstances, for example, new relevant evidence, this will trigger an update of the guideline to potentially amend the recommendations. It is planned to update the current guideline regularly on demand and consistent with the format of a living guideline.

4 | PERIODONTAL DIAGNOSIS AND TREATMENT SEQUENCE FOR THE MANAGEMENT OF STAGE IV PERIODONTITIS PATIENTS

4.1 | Periodontal diagnosis

Defining a case of periodontitis and establishing a diagnosis should be performed using the 2018 classification system, developed following the 2017 World Workshop on the Classification of Periodontal and Peri-Implant Diseases and Conditions (Caton et al., 2018; Chapple et al., 2018; Jepsen et al., 2018; Papapanou et al., 2018; Tonetti et al., 2018).

According to this classification, stage IV periodontitis is identified from the broader population of individuals with stage III periodontitis, defined by periodontal inflammation, and attachment loss reaching the middle third of the root and beyond, based on the need for complex rehabilitation due to the presence of one or more of the following factors:

- Secondary occlusal trauma/tooth hypermobility attributable to a reduced periodontal attachment that is attributed to periodontitis.
- Tooth migration, drifting and opening of diastema are associated with severe attachment loss at the affected teeth.
- Loss of five or more teeth due to periodontitis.

TABLE 7 Timeline of the guideline development process

Time point	Action
April 2018	Decision by European Federation of Periodontology (EFP) General Assembly to develop comprehensive treatment guidelines for periodontitis, including periodontitis in stage IV
May–September 2018	EFP Workshop Committee assesses merits and disadvantages of various established methodologies and their applicability to the field
November 2019	EFP Workshop Committee decides on (i) topics covered by proposed guideline, (ii) working groups and chairs, (iii) systematic reviewers, and (iv) outcomes measures
February 2020	EFP Workshop Committee decides the systematic reviewers
March 2020	Submission of PICO(S) questions by systematic reviewers to group chairs for internal alignment
April 2020	Decision on PICO(S) and information sent to reviewers
May 2020	Decision of postponement from November 2020 until July 2021
June 2020	Decision on consensus group, invitations sent to participants, invitations sent to stakeholders
June–November 2020	Submission of Systematic reviews by reviewers, initial quality assessment by workshop committee
July 2020–March 2021	Submission to <i>Journal of Clinical Periodontology</i> , peer review and revision process
April 2021	Decision of postponement from July 2021 until November 2021, and schedule of preparatory online meetings
June 2021	Submission of declarations of interest by all delegates
5 July 2021	Online plenary meeting and working group meetings
27 September 2021	Online working group meetings
18 October 2021	Online working group meetings
July–October 2021	Electronic circulation of reviews, guideline draft, etc.
7–9 November 2021	Workshop in La Granja with moderated formalized consensus process
November 2021–January 2022	Formal stakeholder consultation, finalization of guideline method report and background text
26 January 2022	Online plenary meeting
February 2022	Submission of guideline document to the <i>Journal of Clinical Periodontology</i>
March–April 2022	Publication of guideline and underlying Systematic Reviews in the <i>Journal of Clinical Periodontology</i>
April–September 2022	Processes of adaptation/adoption by National Societies

- Loss of posterior support and/or flaring of anterior teeth due to periodontitis.
- Loss of masticatory function (masticatory dysfunction) secondary to a combination of the above.

The above signs and symptoms of functional impairment (masticatory dysfunction) may also be present as sequelae of multiple tooth loss due to caries or severe malocclusion in people without significant periodontal breakdown or even in people with periodontal breakdown compatible with stages I–II periodontitis, who do not meet the criteria for stage IV. Hence, differential diagnosis is important.

Stage IV periodontitis not only jeopardizes the survival of individual teeth, but that of the entire dentition. In these patients, control of periodontitis (through standard periodontal therapy, i.e., steps I–III plus supportive periodontal care) is not enough to stabilize the mouth, resolve masticatory dysfunction and improve quality of life. An inter-disciplinary treatment plan that may include the management of secondary occlusal trauma, orthodontic tooth movement and/or restorative dental care following successful periodontal therapy must therefore be implemented to adequately treat these patients.

4.1.1 | Specific diagnostic pathways in patients with stage IV periodontitis

The clinical assessment of a stage IV periodontitis case comprises five critical dimensions:

i. Evaluation of the amount of periodontal breakdown, patient function and aesthetics

A full periodontal examination/charting, combined with appropriate imaging, is required to assess the severity of the periodontal attachment loss and, hence, the complexity of treatment required. In stage IV periodontitis, such examinations must be complemented with an in-depth assessment of the functional and aesthetic status of the individual teeth and the overall dentition, including assessment of tooth hypermobility, assessment of tooth vitality, presence of secondary occlusal trauma, presence of stable posterior vertical stops, fremitus in centric occlusion and excursive movements, subjective and objective assessments of chewing function, aesthetics and phonetics.

ii. Number of teeth that have been lost due to periodontitis

Attributing missing teeth to periodontitis is challenging as it requires a complex assessment based on history of tooth loss/extraction and the symptoms associated with it (Sanz, Papapanou, et al., 2020; Ravidà et al., 2021; Uy et al., 2021). Recall bias and availability of previous records may influence the findings. Nevertheless, the use of a clinical history for the determination of the probable cause of tooth loss adds useful information that can be applied for individual case diagnosis (Ravidà et al., 2021).

iii. Prognosis of individual teeth

Establishment of tooth prognosis in stage IV periodontitis patients, and especially the differentiation of compromised/questionable teeth from those of hopeless prognosis is complex and requires a multidisciplinary approach to identify the ability of treatment(s) that may change the assigned prognosis, which is usually linked to the experience and technical ability of the individual operator(s). Individual tooth prognosis is frequently complicated by the need to assess the possibility of a periodontally compromised tooth that is to be used as an abutment for a fixed or a removable restoration. Studies focusing on periodontal prognosis have shown that it is difficult to accurately predict tooth survival, something that even specialists tend to underestimate (McGuire & Nunn, 1996). Key for long-term prognosis is the ability to achieve the endpoints of periodontal therapy that were identified in the guideline for the treatment of periodontitis in stages I–III (Loos & Needleman, 2020; Sanz, Herrera, et al., 2020), and the implementation of an effective supportive periodontal care programme.

iv. Restorative factors

The extent of the edentulous spaces, as well as the number, distribution and restorability of teeth that can be retained must be assessed, while considering all restorative scenarios, either focusing on teeth alone and/or with the addition of dental implants. These scenarios must consider the technical complexities of the planned prosthesis, as well as the interventions required to place implants, depending on the availability of adequate ridge dimensions.

v. Prognosis of the overall case

The overall case prognosis must be established while considering the individual susceptibility of the patient, based on a thorough analysis of the patient's modifiable and non-modifiable risk factors, using the primary grade criteria as well as the grade modifiers defined earlier (Papapanou et al., 2018; Tonetti et al., 2018). The prognosis should also estimate the probability of disease recurrence/progression, which is a differentiator in the management of stage IV periodontitis patients from other complex restorative cases (Baumer et al., 2011; Lang et al., 2015; Lang & Tonetti, 1996).

The case analysis required to make a sound prognostic assessment and an appropriate treatment plan is complex and requires a detailed assessment of what is technically and biologically feasible, cost-effective and in line with the patient's preferences and expectations.

4.1.2 | Differential diagnosis

For the practical implementation of the case definition and treatment planning, an appropriate differential diagnosis should be established based on the identification of cases with:

- Opening of diastemata or tooth migration secondary to orthodontic relapse.

- Primary occlusal trauma in people with periodontitis.
- Masticatory dysfunction in people who have experienced multiple tooth loss that cannot be attributed to periodontitis and in the absence of periodontitis.
- Masticatory dysfunction in people who have experienced multiple tooth loss that cannot be attributed to periodontitis, but in the presence of stage I–II periodontitis or localized stage III periodontitis
- Generalized stage III periodontitis without tooth loss, or without the other criteria that define stage IV periodontitis. This differential diagnosis may be difficult in some borderline cases. A recent cross-sectional study, with limited external applicability (Uy et al., 2021), has shown that stage IV periodontitis cases, when compared with other periodontitis cases, were more likely to present more severe periodontal breakdown, tooth hypermobility and loss of posterior functional tooth units, as well as self-reporting changes in dietary habits due to their oral condition, impaired chewing ability and lower quality of life.

4.1.3 | Phenotypic variation and identification of clinical case types

Stage IV periodontitis cases may present with great phenotypic variation based on the individual patterns of their periodontal breakdown, number of missing teeth, inter-maxillary relationships and residual alveolar ridge, which will result in different degrees of functional and aesthetic compromise, as well as different treatment needs.

To provide a simplified workable guideline, four major stage IV periodontitis phenotypes were recognized by the organizing committee, leading to specific clinical case types:

- Case type 1: the patient with tooth hypermobility due to secondary occlusal trauma that can be corrected without tooth replacement. It is recognized that there is a continuum of severity and complexity of management between some stage III periodontitis patients, and case type 1 of periodontitis in stage IV.
- Case type 2: the patient with pathological tooth migration, characterized by tooth elongation, drifting and flaring, which is amenable to orthodontic correction.
- Case type 3: partially edentulous patients who can be prosthetically restored without full-arch rehabilitation.
- Case type 4: partially edentulous patients with a dentition that need full-arch rehabilitation, either tooth- or implant-supported/retained.

These phenotypes and associated clinical case types may overlap on occasion, as one arch may require treatment according to a specific scenario while the other might require a different approach. Furthermore, after orthodontic treatment, it is frequently necessary to retain the teeth in the new position with a fixed type of retainer/splint. SRs covering the specific scenarios have been

commissioned and have been used as the basis for development of this clinical guideline.

Several treatment tools are available for the rehabilitation of the different case types, besides the treatment of periodontitis according to the guideline for the treatment of periodontitis in stages I–III (Sanz, Herrera, et al., 2020):

- Temporary control of secondary occlusal trauma (e.g., extra-coronal splinting and/or relief of fremitus by limited selective occlusal adjustment).
- Prosthetic splinting with a fixed dental prosthesis.
- Orthodontic therapy.
- Tooth-supported/retained removable or fixed partial dental prostheses.
- Implant-supported/retained removable or fixed partial dental prostheses.
- Tooth-supported/retained cross-arch dental prostheses, conventional full dental prostheses, implant-retained or -supported full dental prostheses.

The following **key messages** should be highlighted:

- Most cases of stage IV periodontitis can be successfully treated, maintaining the natural dentition in a state of adequate health and function.
- Before treatment planning for people with stage IV periodontitis, it is recommended undertaking a full diagnosis and case study, including a tooth-by-tooth prognosis to identify the number, distribution, residual support, periodontal maintainability and restorability of the remaining natural dentition.

4.2 | Sequence for the treatment of stage IV periodontitis

The treatment plan for the management of stage IV periodontitis should include a successful outcome after completing the interventions in steps 1, 2 and 3, according to the EFP S3 Level clinical practice guideline for treatment of stage I–III periodontitis (Sanz, Herrera, et al., 2020). The sequence of the different steps, however, requires the introduction of specific additional treatment measures to meet the specific demands of stage IV periodontitis. In these cases, rehabilitation of function, restoration of masticatory comfort and treatment of secondary occlusal trauma and, sometimes, restoration of the vertical dimension of the occlusion are also necessary and need to be planned from the beginning, and even implemented simultaneously with steps 1–3.

As is the case for the treatment of periodontitis in stages I–III, an essential pre-requisite to therapy is to inform the patient of the diagnosis, including aetiology of the condition, risk factors, treatment alternatives and expected risks and benefits, including the option of no treatment. This discussion should be followed by agreement on a personalized care plan. The plan might need to be modified during the course of treatment, depending upon initial treatment outcomes,

patient preferences, clinical findings and changes to the patient's overall state of health. It must be recognized that in stage IV periodontitis, a “no treatment” option must be discouraged, given the expected high risk of loss of the dentition.

Key to the care of these patients is:

- The need to combine periodontal therapy, which is modelled in line with the recent guidelines for the treatment of stage I–III periodontitis (Sanz, Herrera, et al., 2020), with rehabilitation.
- Identification of the appropriate timing/sequence of implementation of the adjunctive orthodontic/restorative treatment and the periodontal treatment (Figure 1).

For details of recommended approaches to periodontal therapy, readers are referred to the guideline for periodontal therapy for stage I–III periodontitis (Sanz, Herrera, et al., 2020), since that guideline also applies to the periodontal treatment of stage IV periodontitis patients. Of particular importance are the frequent re-evaluations to assess adherence to oral hygiene instructions for supragingival biofilm control and adherence to risk factor control interventions, and the two periodontal re-evaluation assessments following step 2, and later after step 3 (“final re-evaluation”), since achieving the desired treatment outcomes is particularly important in stage IV cases to be suitable to proceed with the planned restorative or orthodontic therapy, as well as to justify the significant resources required to proceed with the case. In stage IV periodontitis patients, re-evaluation following steps 2 and 3 treatment requires additional planning over and above purely periodontal maintenance. Restorative factors need to be adequately assessed. For example, the ability of a tooth to act as an abutment for a restoration needs to be assessed in terms of periodontal maintainability, but also residual periodontal support and restorative parameters such as the presence of adequate tooth structure. Similarly, dental implants that have been placed to assist in the restoration should present with healthy, maintainable soft- and hard-tissue interfaces.

When evaluating periodontal outcomes at the end of active periodontal treatment (control of periodontal inflammation, achievement of shallow maintainable pockets, management of furcation lesions), it is important to consider whether a specific tooth will be incorporated as an abutment for a fixed or removable prosthesis. Specific criteria must be considered to assess the ability of the tooth to function as a restorative abutment. While it has been shown that teeth with a reduced but healthy periodontium can function well as prosthetic abutments, a minimal threshold of residual periodontal support (10%–20%) may be controversial and dependent upon the design of the restoration (e.g., fixed vs. removable), the number and distribution of the abutments and the stability of the definitive prosthesis (Nyman & Lang, 1994).

4.2.1 | Specific treatment pathways according to the different stage IV periodontitis case types

Common to all clinical case types is the need to perform a careful diagnosis and a case study that includes both periodontal and

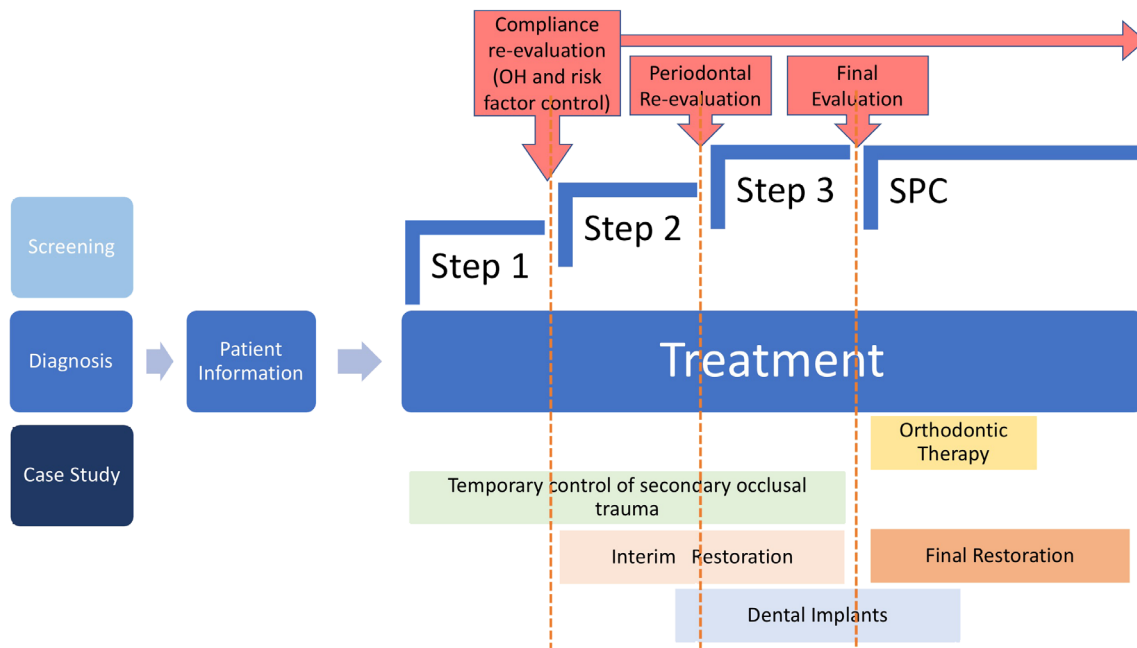


FIGURE 1 Visual description of timing/sequence of implementation of the adjunctive orthodontic/restorative treatment and the periodontal treatment (OH, oral hygiene)

rehabilitation phases (orthodontics and/or restorative dentistry, as appropriate). Furthermore, adequate self-performed oral hygiene, and risk factor control must be realized, alongside satisfactory initial treatment outcomes prior to progressing the case to subsequent periodontal/oral rehabilitation.

- **Case type 1:** the patient with tooth hypermobility due to secondary occlusal trauma that can be corrected without tooth replacement. Temporary tooth splinting and initial occlusal adjustment (mostly relief of fremitus in combination with splinting) can be implemented during step 1 of therapy to manage secondary occlusal trauma and the impact of tooth hypermobility on patient comfort. The need for and the implementation of longer-term splinting needs to be reassessed following completion of steps 2 and 3 of periodontal therapy.
- **Case type 2:** the patient with pathological tooth migration, characterized by tooth elongation, drifting and flaring, which is amenable to orthodontic correction. Orthodontic therapy can be planned during step 2 of care (subgingival instrumentation with or without adjunctives) and, in some cases, step 3 (subgingival re-instrumentation and periodontal surgery) of treatment, but should not be implemented before achieving the periodontal treatment objectives of shallow maintainable pockets and control of periodontal inflammation. Special considerations apply to the regenerative treatment of intra-bony defects (see Section 7).
- **Case type 3:** partially edentulous patients who can be prosthetically restored without full-arch rehabilitation. The timing of intermediate restorations, if required, should be carefully evaluated based on the individuality of the case and keeping in

mind patient wishes and aesthetic considerations. Ideally, interim tooth-retained restorations or dental implants should not be placed before completion of step 2 of treatment and, if possible, deferred until the periodontal treatment objectives have been achieved (after re-evaluation following steps 2 and 3 of periodontal treatment). Definitive restorative treatment or placement of dental implants must be performed after successful completion of periodontal therapy and any additional conservative treatment of the abutment teeth.

- **Case type 4:** partially edentulous patients who need to be restored by means of full-arch rehabilitation, either tooth- or implant-retained, and either fixed or removable.

The timing of treatment differs for cases with tooth-supported full-arch restorations and cases with implant-supported full-arch restorations.

In tooth-supported cases, an intermediate restoration is frequently placed following successful completion of step 1 of periodontal treatment. Step 2 of periodontal treatment, including scaling and root surface instrumentation of the abutment teeth, is performed with the intermediate restoration in place. Insertion of a definitive restoration (or long-term intermediate restoration) follows successful completion of periodontal therapy and achievement of shallow maintainable pockets and control of periodontal inflammation.

For implant-supported cases requiring extraction of the terminal dentition, in one or both arches, teeth are extracted, and implants placed after successful completion of step 1 of periodontal treatment if one arch still has natural teeth. The sequence of treatment and the insertion of intermediate fixed or removable prostheses aims to reconcile the biology of wound healing with the need to manage patient expectations and ensure an adequate level of comfort during the transition.

4.2.2 | Additional therapeutical interventions in stage IV periodontitis

The present guideline has specifically focused on the additional interventions for stage IV periodontitis patients, including:

- Temporary control of secondary occlusal trauma (e.g., extra-coronal splinting and/or relief of fremitus by limited selective occlusal adjustment) (Section 6).
- Orthodontic therapy (Section 7).
- Rehabilitation of one or multiple (small or large) tooth-delimited edentulous spaces (Section 9).
- Rehabilitation of unilateral or bilateral posterior free-end edentulism (Section 9).
- Tooth-supported full-arch fixed dental prostheses (Section 10).
- Tooth-supported full-arch removable dental prosthesis (Section 10).
- Implant-supported full-arch fixed dental prostheses (Section 10).
- Implant-supported removable dental prosthesis (Section 10).

Some of these interventions should be **performed simultaneously or within steps 1–3** of periodontal therapy, including relief of pain, symptoms of tooth hypermobility, functional impairment and conservative treatment of the dentition (including abutment teeth). Restorative treatment implemented during steps 1–3 of periodontal therapy usually consists of intermediate restorations that will only be replaced with definitive ones after final evaluation of the case and achievement of the periodontal and restorative objectives.

Conversely, certain other interventions will only be **performed in the last step (Step R, for rehabilitation)**, following successful completion of steps 1–3 periodontal therapy, once the patient has started supportive periodontal care, and after achievement of the periodontal and restorative treatment objectives. *Step R* treatment includes:

- Long-term periodontal splinting.
- Orthodontic tooth movement and retention.
- Definitive restorations designed to satisfy the functional and aesthetic demands of the patient and enable optimal self-performed oral hygiene and professional tooth cleaning during supportive periodontal care.

4.2.3 | Supportive periodontal care in stage IV periodontitis patients

As for the treatment of stage I–III periodontitis (Sanz, Herrera, et al., 2020), supportive periodontal care (Trombelli et al., 2015) is a crucial step to achieve periodontal stability and long-term tooth/implant retention. This guideline provides specific recommendations for supportive periodontal care in stage IV periodontitis patients (Section 11).

4.2.4 | Impact of therapy in stage IV periodontitis patients on systemic health and quality of life

It is well established that, in addition to positive effects on periodontal outcomes, periodontal therapy may also impact favourably upon systemic health (e.g., reduce systemic inflammation and lower the levels of markers of cardio-metabolic risk) and on quality of life. Since the treatment of stage IV periodontitis includes both periodontal therapy and rehabilitation, the impact of both interventions was considered within the present guideline and recommendations are presented for both the impact of periodontal therapy (Section 12.1) and the impact of rehabilitation (dental prostheses) in fully or partially edentulous patients (Section 12.2).

4.2.5 | Key aspects in the treatment of periodontitis in stage IV

We highlight the following **key messages**:

- To effectively manage stage IV periodontitis, it is recommended that patients are informed in detail about their condition, the various treatment options and associated risks, including the need for periodontal therapy, the design of the rehabilitation, and the sequence of interventions. In addition, patients should be aware that treatment planning may be modified depending upon several factors, including treatment outcomes at re-evaluations, and adherence/compliance with interventions, such as supragingival biofilm control, or risk factor control.
- The starting point for treatment of stage IV periodontitis initially attempts to preserve all periodontally compromised teeth that are deemed rational to treat. Early extraction of teeth with questionable (as opposed to hopeless) prognosis is strongly discouraged and is not supported by current evidence.
- Whenever tooth retention is possible, it is recommended that periodontal treatment of stage IV periodontitis patients should follow the guideline for the treatment of periodontitis in stages I–III. In these patients, the ability to successfully complete full periodontal therapy is a pre-requisite. In addition, management of these cases may also include orthodontics, tooth splinting, tooth-supported fixed and removable dental prostheses, and/or implant-supported fixed and removable dental prostheses.
- For stage IV periodontitis patients, it is recommended to frequently assess motivation and adherence to self-performed supragingival plaque control and risk factor control throughout the course of treatment and during supportive periodontal care, since this will greatly influence both the choice and the outcomes of therapy.
- In stage IV periodontitis patients, it is mandatory that restorations should be designed to achieve function and aesthetics while enabling effective self-performed oral hygiene and professional tooth cleaning.

The first part of this CPG document (Sections 1–4) was prepared by the steering group with the help of the methodology consultant, it was examined by the experts participating in the consensus and Section 4 was voted upon in a plenary session to form the basis for the specific recommendations.

Strength of consensus

Unanimous Consensus (0% of the group abstained due to potential Col).

5 | CLINICAL RECOMMENDATIONS: OVERALL STRATEGY FOR THE MANAGEMENT OF STAGE IV PERIODONTITIS PATIENTS

The pathognomonic features of stage IV periodontitis, as explained in Section 4, are the functional and aesthetic complications that arise following periodontal tissue breakdown and/or the resulting tooth loss. This process severely impacts the quality of life and puts the residual dentition at further risk of being lost if appropriate treatment is not rendered. As discussed, the disease spectrum covers a wide range of phenotypic variations characterized by a wide spectrum from subtle changes, that may be overlooked, to severe loss of function that raises the question of whether the dentition can be saved, and rehabilitation of function can translate in restoration of quality of life. The competences required for appropriate diagnosis and management of these cases are frequently complex and inter-disciplinary, while the evidence base supporting the different choices is frequently limited. In such situation of complexity and uncertainty, the experts and stakeholders participating in the workshop agreed on a series of expert-based recommendations that provide critical guidance in the management of these cases, in order to understand the general strategical principles for therapeutic management in patients with a compromised dentition due to stage IV periodontitis.

R5.1. Can stage IV periodontitis can be successfully managed?

Additional question raised by the WG

R5.1: Expert consensus-based statement

Stage IV periodontitis can be successfully managed with the combination of periodontal therapy, appropriate rehabilitation of function and improvement of aesthetics/quality of life, and supportive periodontal care.

Supporting literature Expert opinion

(Continues)

(Continued)

Additional question raised by the WG

R5.1: Expert consensus-based statement

Quality of evidence Not applicable

Grade of recommendation Statement

Strength of consensus Unanimous Consensus (0% of the group abstained due to potential Col)

Background

Experts wished to emphasize that stage IV periodontitis can be managed beyond the scope of palliative care and that management requires appropriate periodontal therapy delivered according to the S3 guideline for management of stage I–III periodontitis (Sanz, Herrera, et al., 2020) in combination with rehabilitation of lost function. Once active treatment is completed, supportive periodontal care is a critical component to enable long-term success. Further, due to the complexity, case instability and loss of function, option zero (no treatment) is rarely an option.

R5.2. How relevant is tooth retention?

Additional question raised by the WG

R5.2: Expert consensus-based recommendation

We recommend tooth retention to be the first line of treatment strategy in the rehabilitation of stage IV periodontitis patients.

Supporting literature (Montero et al., 2021; Tomasi et al., 2021)

Quality of evidence Low

Grade of recommendation Grade A–↑↑

Strength of consensus Strong Consensus (0% of the group abstained due to potential Col)

Background

In the long-term management of stage IV periodontitis, retention of the natural dentition with adequate therapy, whenever possible, provides a strategic advantage as it defers the time of implant-based options and shortens their required longevity. The option of tooth retention needs to be considered first and alternatives need to be justified for the specific case based on case and individual tooth prognosis, technical feasibility, patient preference and, ideally, cost–benefit considerations.

R5.3. How relevant is preservation of dental arch integrity

Additional question raised by the WG

R5.3: Expert consensus-based recommendation

In patients with stage IV periodontitis, we suggest preserving the integrity of the dental arch by avoiding extractions, if teeth can be retained.

Supporting literature Expert opinion

(Continues)

(Continued)

Additional question raised by the WG**R5.3: Expert consensus-based recommendation***Quality of evidence* Not applicable*Grade of recommendation* Grade B—↑*Strength of consensus* Consensus (0% of the group abstained due to potential Col)**Background**

In cases with an intact dental arch (no missing teeth apart from the molar area), the possibility to preserve all teeth, and thus avoid the need for tooth replacement, through periodontal and/or restorative treatment provides a strategic advantage. It may simplify treatment and reduce costs. A careful diagnosis of the possibility to preserve “hopeless” or severely compromised teeth by advanced periodontal therapy, combined with management of hypermobility and/or addressing patient concerns, should be performed before deciding to extract teeth in such cases.

R5.4. Can acceptability of tooth preservation be improved with simple measures?**Additional question raised by the WG****R5.4: Expert consensus-based recommendation**

In patients with stage IV periodontitis requesting improvement of aesthetics, phonetics, masticatory function and/or patient well-being, direct and indirect tooth restorations and/or epitheses may be considered.

Supporting literature Expert opinion*Quality of evidence* Not applicable*Grade of recommendation* Grade O—↔*Strength of consensus* Consensus (0% of the group abstained due to potential Col)**Background**

In many stage IV periodontitis cases, the aesthetic, phonetic and functional sequelae of periodontal breakdown severely affect patient well-being and quality of life. Once correctly identified, such concerns may raise questions on tooth-retention-based choices, unless costly approaches are considered, and may provide barriers to the acceptability of periodontal therapy and tooth retention. Simple approaches such as the use of gingival epitheses (custom-made removable silicone masks to replace missing gingiva that hide black triangles or correct the presence of uneven gingival margins) and/or direct/indirect adhesive restoration that re-shape teeth (and sometimes provide a splinting effect) may mitigate patient concerns and increase the acceptability of periodontal treatment in stage IV periodontitis patients.

6 | CLINICAL RECOMMENDATIONS: CASE TYPE 1**Case type 1: the patient with tooth hypermobility due to secondary occlusal trauma that can be corrected without tooth replacement.****6.1 | Intervention: Tooth splinting and occlusal adjustment in patients with periodontitis stage IV**

A systematic review (Dommisch et al., 2021) assessed the efficacy of tooth splinting (TS) and occlusal adjustment (OA), compared with no-TS or OA in patients with periodontitis exhibiting masticatory dysfunction. Both treatment modalities are especially relevant in fully dentate patients with stage IV periodontitis. In this context, TS reduces tooth mobility due to advanced attachment loss to facilitate biting and chewing abilities (patient's comfort), and OA addresses secondary occlusal trauma in the management of occlusal dysfunction. In the systematic review (Dommisch et al., 2021), no study specifically focused on patients with stage IV periodontitis, and thus, the efficacy of TS and OA was evaluated for patients with all stages of periodontitis.

The primary outcome criterion was tooth loss, and the secondary outcome parameters were change in PPD, change in clinical attachment level (CAL), tooth mobility (TM) and PROMs (Dommisch et al., 2021). For TS, two studies were included considering a follow-up time range between 3–15 and 2–32.4 years, respectively. Three studies addressed OA with a follow-up of 2–8 years.

As explained in Section 4, a degree of overlap between stage III periodontitis and case type 1 of periodontitis in stage IV exist.

R6.1. In fully dentate patients with periodontitis stage IV with increased tooth mobility due to advanced periodontal attachment loss, what is the effectiveness of tooth splinting and/or limited occlusal adjustment of hypermobile teeth?*(Common question for Recommendations R6.1, R6.2)***PICOS question addressed by a SR****R6.1: Evidence-based recommendation**

In patients with stage IV periodontitis, temporary splinting and/or limited selective occlusal adjustment of hypermobile teeth may be considered during all steps of periodontal therapy (but particularly during step 1 treatment) to increase patient comfort and enable/facilitate periodontal therapy.

Supporting literature (Dommisch et al., 2021; Sonnenschein, Ciardo, et al., 2021; Sonnenschein, Ziegler, et al., 2021)*Quality of evidence* Very low*Grade of recommendation* Grade O—↔

(Continues)

(Continued)

PICOS question addressed by a SR

R6.1: Evidence-based recommendation

Strength of consensus Unanimous Consensus (0% of the group abstained due to potential CoI)

Background

Intervention

TS and OA in fully dentate patients with periodontitis.

Available evidence

Number and design of included studies

For **TS**, the two included retrospective studies ($n = 72$ patients) analysed a time follow-up time period ranging between 2 and 32.4 years. After a minimum of 2 years following non-surgical periodontal therapy, synthesis of data revealed a weighted mean incidence of 8.4% of tooth loss for TS compared with 10.1% of tooth loss for no-TS. Outcome measures comprised, beside tooth loss, PPD, changes in CAL, changes in bone level (BL), and plaque scores. Biological complications, PROMs, health-economic parameters and adverse events were not consistently reported.

One additional randomized prospective study indicated a positive effect on the oral health-related quality of life 3 months after non-surgical periodontal therapy (Sonnenschein, Ziegler, et al., 2021). The time point of TS (prior or following step 2 of periodontal therapy) was addressed in a randomized trial, and the timing appeared not to influence the outcome of periodontal parameters (Sonnenschein, Ziegler, et al., 2021).

Throughout periodontal therapy, TS repairs were frequently required (Sonnenschein et al., 2017; Graetz et al., 2019). In patients with periodontitis stage IV, both TS and advanced attachment loss may widely impair the patient's aesthetics.

The main findings were that tooth TS does not affect tooth loss, and does not affect PPD, CAL and BL.

Finally, and in preparation for step 3 of periodontal therapy, since an increase of TM following flap mobilization is anticipated, TS may be beneficial when regenerative periodontal treatment is planned (Cortellini et al., 2001).

For **occlusal adjustment** (OA), three studies ($n = 205$ patients) were included, one randomized controlled clinical trial (RCT) (Burgett et al., 1992) and two prospective studies (randomization did not include OA procedures) (Fleszar et al., 1980; Kerry et al., 1982). All studies were of prospective design. While two studies included OA as part of the initial phase, one study allocated OA randomly facilitating a comparison of patients with and without OA and no-OA. The follow-up time ranged from 2 to 8 years. None of the studies reported on TL or PROMs.

The main findings were that the effects of OA on TL are unclear, that OA positively affects CAL, and that OA does not affect PPD and BL.

Risk of bias

For **TS**, study quality assessment using the Newcastle–Ottawa scale identified a low risk of bias for both studies included. For OA, study quality assessment using the Cochrane Collaboration's tool for assessing risk of bias identified unclear risk of bias, mainly related to potential centre bias.

Effect sizes and their clinical relevance

For **TS**, based on two retrospective case series ($n = 72$ patients), over at least 2 years mean TL was 8.4% for TS and 10.1% for no-TS. TS may be performed as an adjunctive intervention for teeth with a progressive increase in TM in order to improve the patient's chewing capabilities.

For **OA**, based on one study ($n = 50$ patients), CAL was monitored over a period of 2 years (Burgett et al., 1992). After 1 and 2 years, OA led to an improved CAL of approximately 0.4 mm (mean) following non-surgical periodontal therapy. Although this improvement may not be considered as clinically relevant, OA may remove jiggling forces, facilitate patients' chewing capabilities and resolve pain perception when premature contacts are removed.

Consistency

In the analysis of **TS**, only two articles exhibiting heterogenous data with a wide range of follow-up examinations (between 2 and 32.4 years) were in accordance with the inclusion criteria of the systematic review. The included studies differed regarding the definition of the control group as well as operators performing splinting and periodontal examination. In addition, one study reported a high drop-out rate (Sonnenschein et al., 2017).

In the analysis of **OA**, only three studies of heterogeneous design were in accordance with the inclusion criteria of the systematic review. Of these three studies, one study was designed as an RCT focusing on randomization for OA (Burgett et al., 1992), while the two remaining studies randomized for periodontal treatment modalities but not for OA (Fleszar et al., 1980; Kerry et al., 1982).

Balance of benefit and harm

For **TS**, the overall evidence suggests a very weak positive relationship between benefit and harm/risks. Although TS does not improve tooth survival, a positive effect on the individual patient comfort exists. However, if TS is performed, splinting technique, material and design should enable optimal self-performed and professional oral hygiene procedures.

Preliminary data exist on OHRQoL in periodontitis patients. Three months after non-surgical periodontal therapy, a positive impact on the OHIP-14 summary scores was documented, suggesting a trend towards improved patients' comfort (Sonnenschein, Ziegler, et al., 2021). There is a lack of information on PROMs and adverse events, and those need to be evaluated in future research.

For **OA**, the existing evidence suggests a weak positive relationship between benefit and harm/risks. There is no information on PROMs and adverse events, and those need to be evaluated in future research.

Overall certainty of the evidence

Very low for both TS and OA.

From evidence to recommendation—additional considerations

Acceptability

The certainty of evidence is graded as very low based on the lack of studies, study design and lack of generalizability of data.

Feasibility

Both TS and OA can be performed by general dentist as well as specialist as the minimum standard of care.

Ethical considerations

Based on the available evidence, no evaluation of ethical aspects could be performed. TS per se is a minimal-invasive, and in most cases, reversible therapeutic procedures that do not negatively influence tooth survival. OA is a minimal-invasive but not reversible therapeutic procedure that does not negatively influence periodontal outcome variables.

Economic considerations

Health-economic parameters were not evaluated in the identified studies.

Legal considerations

Not applicable.

R6.2. See question for R6.1

PICOS question addressed by a SR

R6.2: Evidence-based recommendation

In patients with stage IV periodontitis who do not require tooth replacement but show persistent hypermobility or increasing mobility after successful completion of periodontal therapy, long-term tooth splinting may be considered to improve patient comfort.

Supporting literature (Dommisch et al., 2021; Sonnenschein, Ciardo, et al., 2021; Sonnenschein, Ziegler, et al., 2021)

Quality of evidence Very low

Grade of recommendation Grade O—↔

Strength of consensus Unanimous Consensus (0% of the group abstained due to potential Col)

Background

See the background text for Recommendation R6.1.

7 | CLINICAL RECOMMENDATIONS: CASE TYPE 2

Case type 2: the patient with pathological tooth migration, characterized by tooth elongation, drifting and flaring, which is amenable to orthodontic correction.

7.1 | Introduction: Treatment of pathological tooth migration in stage IV periodontitis patients

As explained in Sections 1.1.1 and 4.1, stage IV periodontitis is characterized by similar severity and complexity as stage III periodontitis in terms of periodontal inflammation, attachment and bone loss, but these stage IV periodontitis patients have either lost five or more teeth due to periodontitis and/or are in need for complex rehabilitation due to one or more of the following criteria:

- Bite collapse, tooth migration, drifting, flaring and spacing are associated with severe attachment loss at the affected teeth.
- Loss of posterior support and/or flaring of anterior teeth consequent to periodontitis.
- Secondary occlusal trauma/tooth hypermobility, degree ≥ 2 , attributable to reduced periodontal attachment consequent to periodontitis.
- Less than 20 remaining teeth (10 opposing pairs)
- Loss of masticatory function (masticatory dysfunction) secondary to a combination of the above.

Common to these patients is that the lack of appropriate treatment, not only risks the loss of affected remaining teeth, but also loss of the whole remaining dentition. In stage IV periodontitis patients, the treatment of periodontitis (through standard periodontal therapy, i.e., steps I–III plus supportive periodontal care) is not enough to stabilize the case, resolve the masticatory dysfunction and improve their quality of life. An inter-disciplinary treatment plan, therefore, must be implemented, which may include orthodontic and/or prosthetic rehabilitation for the stabilization and/or restoration of the masticatory function, patient aesthetics and quality of life.

In stage IV periodontitis patients, there are specific phenotypic variations/clinical scenarios based on the individual patterns of the periodontal breakdown, which result in different degrees of functional and aesthetic compromise, as well as in different treatment needs. One of the most common phenotypic variations is the clinical case defining the patient with pathological tooth migration, characterized by tooth elongation, drifting and flaring, which is amenable to orthodontic correction. This consensus report presents the evidence-based and expert-based recommendations of the present clinical practice guideline for the treatment of the described case type (type 2) of stage IV periodontitis (Figure 2).

Therefore, recommendations refer to combined periodontal and orthodontic therapy in stage IV periodontitis patients where orthodontic therapy is indicated.

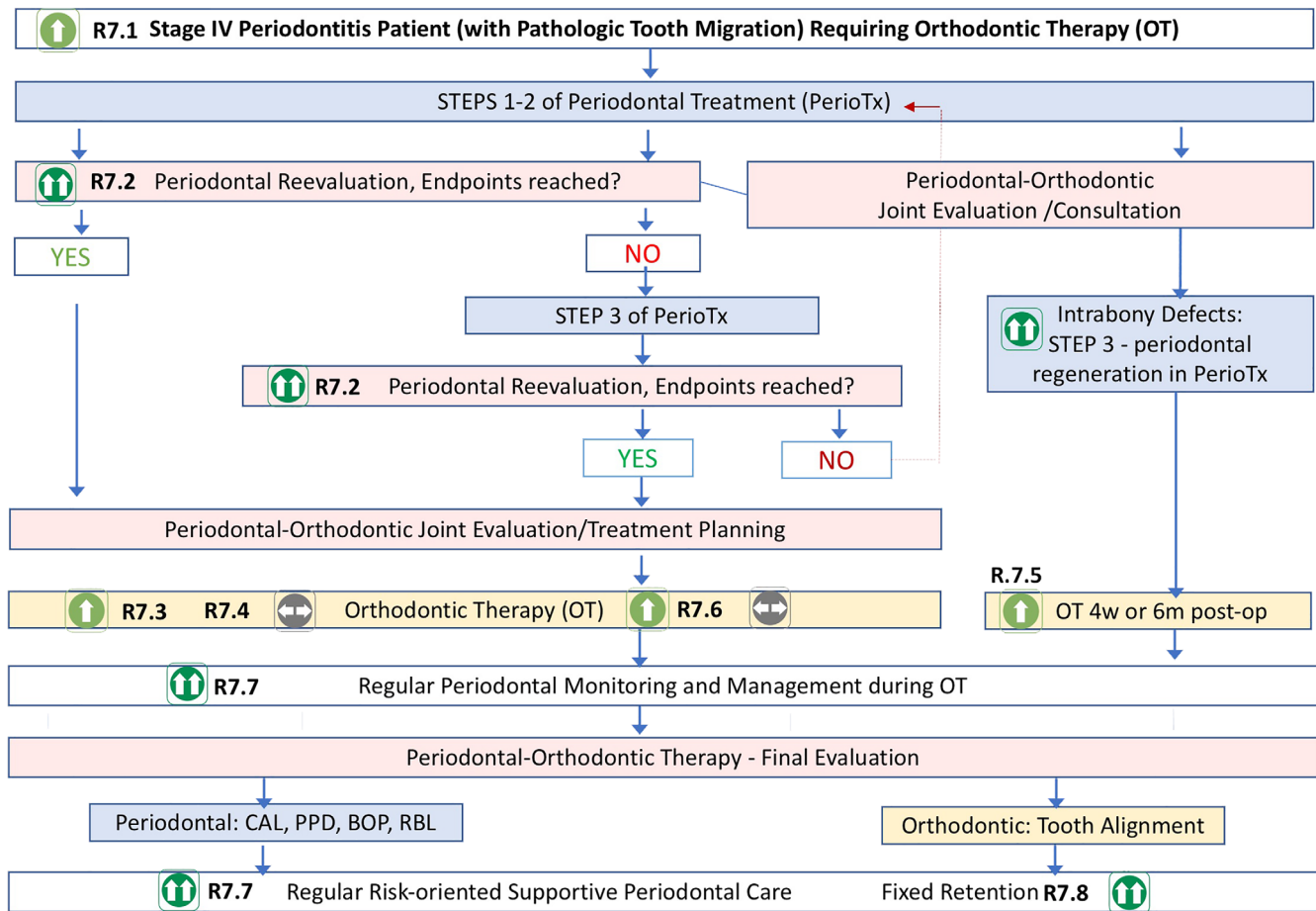


FIGURE 2 Flowchart illustrating how orthodontic therapy (OT) of stage IV periodontitis patients can be integrated in the overall periodontal treatment plan with reference to the recommendations R7.1–R7.8 of the S3-Level clinical practice guideline for the treatment of stage IV periodontitis. Steps of periodontal therapy were described in the S3-Level clinical practice guideline for the treatment of periodontitis stage I–III (Sanz, Herrera, et al., 2020). CAL, clinical attachment loss; PPD, probing pocket depth; BOP, bleeding on probing; RBL, radiographic bone loss; 4 weeks, 4 weeks; 6 months, 6 months; post-op, post-operative

7.2 | Clinical recommendations for case type 2: Orthodontic therapy in the treatment of stage IV periodontitis patients

R7.1. In stage IV periodontitis patients, when there is an indication for both periodontal and orthodontic therapy, what is the effect of orthodontic therapy (OT) on periodontal health and what are the possible adverse effects and complications?

PICOS question addressed by a SR

R7.1: Evidence-based recommendation

In successfully treated stage IV periodontitis patients in need of orthodontic therapy, we **suggest** undertaking OT based on evidence that:

- a. it does not significantly affect periodontal outcomes (probing pocket depth-PPD and clinical attachment levels-CAL);

(Continues)

(Continued)

PICOS question addressed by a SR

R7.1: Evidence-based recommendation

- b. it does not significantly affect gingival inflammation (bleeding on probing–BOP) and gingival recession;
- c. it does not lead to a significant increase in root resorption.

Supporting literature (Martín et al., 2021)

Quality of evidence Moderate

Grade of recommendation Grade B–↑

Strength of consensus Consensus (0% of the group abstained due to potential Col)

Background

Intervention

Treated stage IV periodontitis patients frequently require OT to restore the tooth position and functional occlusion, which during the

progression of periodontitis have pathologically migrated, as a consequence of periodontal attachment and bone loss. In these patients with a healthy but reduced periodontium, the results of OT may be different, compared with patients without attachment loss, or the orthodontic tooth movements may cause adverse effects on the affected teeth or on the remaining periodontium.

Available evidence

Number and design of included studies

The SR by Martin et al. (2021) included 35 studies. These studies were mainly case series with a pre–post design (15 prospective and 5 retrospective) or cohort studies (4 prospective and 2 retrospective) evaluating the outcomes of OT in periodontitis patients; or studies comparing the outcomes of OT between periodontitis versus non-periodontitis patients, either with RCTs (1 split mouth and 7 with parallel arms), or CCT (1 split mouth).

Risk of bias

The overall quality of the included studies was evaluated as “low”. From the eight RCTs assessed using RoB 2.0 tool, the overall bias assessment was rated as “high” for two RCTs; as “low risk of bias” for three RCTs and as “some concern” in the other three. Regarding the 20 pre–post intervention studies, only 1 was rated as “good” quality, 10 was rated as “fair” (moderate risk of bias) and 9 as “poor”. The cohort studies were rated with the Newcastle–Ottawa scale (NOS) scale with 4 being rated as “fair” quality, one as “poor” and one as “good”.

Effect sizes and clinical relevance

A. Periodontal outcomes (PPD and CAL)

In patients without periodontitis, changes in CAL (mm) (mean effect–ME = 0.248; 95% CI [–0.055; 0.551]; $p = .109$) were minimal and non-statistically significant across time ($p > .05$). For changes in PPD (mm), a statistically significant reduction was observed (ME = 0.325; 95% CI [0.123; 0.526]; $p < .001$).

In patients with treated periodontitis, no significant changes were observed in CAL (mm) (ME = 0.202; 95% CI [–0.018; 0.422]; $p = .072$). There was a slight reduction in PPD (mm) (ME = 0.129; 95% CI [0.058; 0.200]; $p < .001$), during OT.

B. Gingival inflammation and gingival recession

In patients without periodontitis, changes in BOP (%) (ME = 3.63; 95% CI [–2.11; 9.37]; $p = .215$) and in gingival recession (REC) (mm) (ns = 2, np = 26; ME = 0.23; 95% CI [–0.32; 0.77]; $p = .418$) were minimal and non-statistically significant ($p > .05$).

In patients with treated periodontitis, also changes in BOP (%) (ME = 0.96; 95% CI [–0.77; 2.69]; $p = .278$) or REC (mm) (ME = 0.08; 95% CI [–0.37; 0.53]; $p = .731$) were minimal during OT and not statistically significant.

C. Root resorption

Due to the scarcity of the data, no meta-analyses could be constructed for adverse effects. The advent of root resorption after

OT in periodontitis patients was reported in a few studies irrespective of the type of orthodontic movement or anchorage system used. In the conventional anchorage group, one study (Melsen et al., 1989) reported 1–3 mm of root resorption when the incisors were orthodontically intruded, while in another study (Corrente et al., 2003) no resorption using similar orthodontic movements was reported. In the skeletal anchorage group, only one study reported apical root resorption (ranging from 0.2 to 0.4 mm) in the periodontally treated group, while similar root resorption was also reported in periodontally healthy patients treated by OT (ranging from 1 to 1.5 mm).

Balance of benefit and harm

OT has no detrimental effects on periodontal conditions in periodontitis patients with a healthy but reduced periodontium, provided the results of periodontal therapy are maintained during the active OT. Overall, the benefits exceed the potential harms.

Consistency

All studies reported the same tendency.

Overall certainty of the evidence

The evidence was graded as moderate since a relevant number of studies reported consistent results, albeit with moderate to high risk of bias.

From evidence to recommendation—additional considerations

Acceptability

Considering the likely benefits (improved tooth position, functional occlusion, and aesthetics), this treatment is well accepted by the patients, although there is lack of evidence on patient-reported outcomes (PROMs) from the referred studies.

Feasibility, ethical and economic considerations

It should be considered that OT in patients with a reduced periodontium may be a complex treatment frequently requiring application of complicated devices and appliances, which requires that appropriate specialists or dentists with advanced training implement these treatments. Furthermore, these orthodontic treatments are usually long (up to 2 years) and require regular visits, which imposes the patient or the health system with considerable costs. There are, however, no available cost-effectiveness studies for these combined perio-ortho treatments. The considerable costs needed for these treatments are mainly financially self-supported by the affected individuals in many European countries, what may result in inequity when no healthcare funding is available. Furthermore, the availability for appropriate professionals to render these complex treatments may vary widely.

Legal considerations

There are usually no legal constraints to OT.

R7.2. In stage IV periodontitis patients, when should orthodontic therapy (OT) start?

Additional question addressed by the WG

R7.2: Expert consensus-based recommendation

In successfully treated stage IV periodontitis patients in need of orthodontic therapy, we **recommend** starting OT once the endpoints of periodontal therapy have been achieved [no sites with PPD = 5 mm and BOP and no sites with PPD ≥6 mm (Sanz, Herrera, et al., 2020)].

Supporting literature Expert opinion; data from pre-clinical studies

Quality of evidence Low (expert opinion)

Grade of recommendation Grade A—↑↑

Strength of consensus Strong consensus (0% of the group abstained due to potential Col)

Background

Intervention

OT is indicated for the functional and aesthetic rehabilitation of stage IV periodontitis patients with pathological tooth migration, characterized by tooth elongation, drifting and flaring. This OT must fulfil its objectives without jeopardizing the short- and long-term outcomes of periodontal therapy and hence it is important to stage it appropriately within the overall patient's treatment plan.

Available evidence

Number and design of included studies

The evidence for this recommendation is derived from expert opinion and from experimental in vivo investigations.

Risk of bias

Not applicable.

Effect sizes and their clinical relevance

The evidence derives from well-controlled pre-clinical in vivo investigations (Eliasson et al., 1982; Ericsson et al., 1977; Melsen, 1986; Wennstrom et al., 1993) and clearly indicates that when periodontitis is not fully treated (inflammation is not arrested) prior/during OT, these orthodontic (biomechanical) forces within the periodontal tissues with remaining inflammatory processes will re-initiate and/or accelerate the progression of periodontal destruction, leading to further loss of clinical attachment and supporting alveolar bone. In contrast, no detrimental effects of orthodontic tooth movements have been observed when these movements are exerted on

teeth with healthy (non-inflamed) reduced periodontal support.

Consistency

All the referred pre-clinical investigations report consistent results indicating that when periodontitis is not adequately controlled, OT will result in further tissue destruction, evidenced by loss of periodontal attachment and alveolar bone.

Balance of benefit and harm

These unwanted effects (further loss of periodontal attachment and bone) clearly outweigh the benefit of OT and hence orthodontic tooth movements should not be started until periodontal inflammation is arrested, when fulfilling the well-established end points of periodontal therapy (Sanz, Herrera, et al., 2020).

Overall certainty of the evidence

Low, expert opinion.

From evidence to recommendation—additional considerations

Acceptability

Patients usually accept and understand, once they have been explained the balance of the associated benefits versus harms, that OT can only be implemented once periodontal therapy has been completed and the end points of periodontal therapy have been reached.

Feasibility, ethical and economic considerations

The only perceived barrier is the length of overall treatment, but since the benefits clearly outweigh this inconvenience, this barrier is usually well accepted.

There are no ethical or economic considerations.

Legal considerations

There are no legal considerations.

R7.3. How should we manage stage IV periodontitis patients with pathological tooth migration (flaring, drifting and elongation)?

PICOS question addressed by a SR

R7.3: Evidence-based recommendation

In stage IV periodontitis patients with pathological tooth migration, we **suggest** undertaking orthodontic therapy once the endpoints of periodontal therapy have been reached, based on the evidence that this therapy:

- does not significantly affect periodontal outcomes [CAL, PPD, and radiographic bone levels (RBL)];
- seems to reduce gingival inflammation (BOP);
- does not significantly alter gingival margin levels;

(Continues)

(Continued)

PICOS question addressed by a SR**R7.3: Evidence-based recommendation**

- d. *seems to improve inter-dental papilla height;*
- e. *does not significantly affect root resorption and seems to reduce tooth mobility.*

Supporting literature (Papageorgiou et al., 2021)**Quality of evidence** Moderate**Grade of recommendation** Grade B-†**Strength of consensus** Consensus (0% of the group abstained due to potential Col)

Background

Intervention

Pathological tooth migration is a frequent sequela of periodontal attachment loss in stage IV periodontitis patients manifested by tooth drifting, flaring and elongation. Its correction requires OT after completion of periodontal therapy. The treatment of these sequelae usually involves intrusive, retrusive, and alignment tooth movements, which may potentially cause adverse effects (further periodontal attachment or bone loss, increased gingival inflammation, or increased root resorption) or secondary effects (undesired aesthetic outcomes such as gingival recession and loss of inter-dental papilla) to the affected teeth.

Available evidence

Number and design of included studies

Thirty-four studies (from 37 publications) were included, reporting data from 1090 participants with mean age of 43.7 years and a female/male ratio of 3/1. Included studies were either comparative (randomized or non-randomized) or single-group cohort studies with at least one group with pre-post data combining periodontal-orthodontic therapy in stage IV periodontitis patients with pathologic tooth migration.

Data were analysed either directly from comparative studies (randomized/non-randomized) or indirectly, by pooling all study arms from pre-post data studies, calculating the average effects for each outcome and identifying the associated factors through subgroup/meta-regression analyses. In this background text, the latter is referred to as *indirect meta-analyses* and their results presented as *pooled changes*; for *direct meta-analysis*, *mean differences* (MD) are presented. Data for periodontal outcomes were retrieved from two comparative studies for PPD and from one study for RBL. Indirect meta-analyses from single-group before-and-after studies assessed CAL (7 studies), PPD (7 studies) and RBL (7 studies).

Gingival inflammation was reported in one retrospective non-randomized comparative study, while gingival recession was reported

in three non-randomized single-group before-and-after studies. Inter-dental papilla height by indirect meta-analysis from two non-randomized single-group before-and-after studies and root resorption by indirect meta-analysis from three non-randomized single-group before-and-after studies.

Risk of bias

All studies were evaluated as high risk of bias based on the ROBINS-I or RoB 2.0 tools.

Effect sizes and their clinical relevance

There was no statistically significant difference in the overall PPD change between combined periodontal-orthodontic therapy compared with periodontal therapy alone, in a meta-analysis of two non-randomized multi-group comparative cohort studies with 92 patients (MD = -0.31 mm; 95% CI [-0.83; 0.22]; $p = .26$; $I^2 = 83\%$).

Indirect meta-analyses from single-group before-and-after studies on the effect of OT reported significant CAL gain with a pooled change of -0.24 mm (95% CI [-0.38; -0.10 mm]; $p < .001$; $I^2 = 79\%$ [7 studies]). Similarly, there was PPD reduction with a pooled change of -0.23 mm (95% CI [-0.49; 0.04]; $p = .09$; $I^2 = 95\%$ [7 studies]); and RBL gain with a pooled change of -0.36 mm (95% CI [-0.59; -0.13]; $p = .002$; $I^2 = 88\%$ [7 studies]).

There was no significant effect in terms of the percentage of RBL changes from the combined periodontal-orthodontic therapy (1 study; 20 patients; MD = -0.60%; 95% CI [-2.80; 1.60]; $p = .59$).

One study reported reduced *gingival inflammation* (bleeding index) with combined periodontal-orthodontic therapy compared with only periodontal treatment (1 study; 72 patients; MD = -13.89%; 95% CI [-16.06; -11.72]; $p < .001$).

Indirect meta-analyses reported no significant effect on average gingival recession either for the combined periodontal-orthodontic therapy with pooled change of -0.53 mm (95% CI [-2.07; 1.01]; $p = .50$; $I^2 = 98\%$ [3 studies]); or solely for OT subsequent to periodontal therapy, with pooled change of 0.09 mm (95% CI [-0.01; 0.20]; $p = .09$; $I^2 = 0\%$ [2 studies]).

Indirect meta-analysis reported increased inter-dental papilla height after OT with pooled change of -1.42 mm (95% CI [-1.98; -0.86]; $p < .001$; $I^2 = 94\%$ [2 studies]).

Indirect meta-analyses reported no significant increase in root resorption either for the combined periodontal-orthodontic therapy with pooled change of -0.42 mm (95% CI [-0.63; -0.22]; $p < .001$; $I^2 = 56\%$ [3 studies]); or solely for the subsequent OT with pooled change of -0.49 mm (95% CI [-1.04; 0.06]; $p = .08$; $I^2 = 71\%$ [2 studies]).

Consistency

The results on the periodontal outcomes and gingival inflammation indicated either minor non-relevant effects or benefits from the combined periodontal-orthodontic therapy. Consistency could not be formally assessed by direct meta-analysis due to the small number of included studies.

Balance of benefit and harm

In stage IV periodontitis patients in need of OT, the benefit after OT clearly outweighs the harm since the overall pooled effect resulted in a beneficial impact on CAL, PPD, RBL and gingival inflammation without being associated with significant adverse or secondary effects.

From evidence to recommendation—additional considerations

Acceptability

The addition of OT might impose an additional burden in patients, but the expected benefits resulting from the improved function and aesthetics are usually understood and well accepted by the affected patients, although there is lack of evidence on PROMs from the referred studies.

Feasibility, ethical and economic considerations

These combined treatments need the collective effort of many oral health providers (dental hygienists, periodontists, orthodontists or dentists with advanced training and skills) since beyond the management of the OT, patient's oral hygiene and periodontal status must be carefully monitored throughout the treatment. Availability of these needed human resources vary widely depending on the environment and may be influenced by the available public healthcare funding, since in many instances it must be self-funded by the affected patient, what may implicate inequalities in the access to these treatments. There is no evidence on the economic effects, since there are no cost-effectiveness studies, although these combined periodontal/orthodontic treatments are usually costly since they require complex interventions and require multiple care providers.

Legal considerations

There are no apparent legal constraints.

R7.4. How should we manage stage IV periodontitis patients with tilted molars?

PICOS question addressed by a SR

R7.4: Expert consensus-based statement

In stage IV periodontitis patients with tilted molars orthodontic therapy may be considered, although there is a lack of evidence on its possible effect on periodontal outcomes.

Supporting literature (Kloukos et al., 2021)

Quality of evidence Very low

(Continues)

(Continued)

PICOS question addressed by a SR

R7.4: Expert consensus-based statement

Grade of recommendation Grade O↔

Strength of consensus Strong consensus (0% of the group abstained due to potential CoI)

Background

Intervention

Tilted molars are a frequent sequela of tooth loss and periodontal attachment loss in stage IV periodontitis patients, often in combination with bite collapse and loss of the vertical dimension of the occlusion. The treatment of these sequelae usually involves OT using movements of tooth up-righting, which may cause adverse effects to the affected teeth (further attachment and/or bone loss).

Available evidence

Number and design of included studies

The SR by Kloukos et al. (2021) has only identified one study evaluating the effect of OT on tilted molars (Kraal et al., 1980). This split-mouth retrospective study reported a single cohort of patients, where, on one side, tilted molars were orthodontically treated, while, on the contra-lateral side, they were not.

Risk of bias

Using the ROBINS-I-tool, this study by Kraal et al. (1980) was rated as of critical risk of bias.

Effect sizes and their clinical relevance

There were no data available on PPD or CAL changes. Results only reported mean radiographic bone height changes. In 15 patients with treated and untreated sides, mean bone height change was -2.21 mm (SD = 4.53) in untreated, versus -1.07 mm (SD = 7.35) in treated teeth. The mean bone height change was -2.53 mm (SD = 4.53) in 15 untreated molars, versus -1.60 mm (SD = 7.01) in 21 uprighted molars. Due to the scarcity of available evidence, the clinical relevance of these results could not be considered.

Balance of benefit and harm

Adverse effects such as tooth loss, abscess or decay were not reported in this study. Similarly, success and duration of orthodontic tooth movement, as well as subjective/objective evaluation of masticatory function or PROMs were not reported. Due to the scarcity of available evidence, the balance of benefit versus harm of this type of combined perio-ortho therapy could not be assessed.

Overall certainty of the evidence

Considering the evidence derived from only one retrospective study, the overall certainty of the evidence was rated as very low.

From evidence to recommendation—additional considerations

Acceptability, feasibility, ethical, economic and legal considerations of this intervention could not be evaluated.

R7.5. In stage IV periodontitis patients presenting with intra-bony defects and in need of OT, what is the outcome of the combined periodontal and orthodontic therapy and what should be the time interval between the periodontal regenerative and orthodontic therapies?

PICOS question addressed by a SR

R7.5: Evidence-based recommendation

In stage IV periodontitis patients where intra-bony defects have been treated following the recommendations of the clinical practice guideline ([Sanz, Herrera, et al., 2020] using the appropriate regenerative interventions):

1. **We recommend** undertaking OT based on the evidence that the combined treatment significantly improves periodontal outcomes (increased CAL gain, PD reduction and RBL gain) and significantly reduces gingival inflammation (BOP).
2. **We suggest** not to wait for a prolonged healing period after the regenerative intervention, before initiating OT, since there is evidence that a short (1 month) and a prolonged (6 months) period between periodontal/regenerative and OT result in comparable outcomes.

Supporting literature (Kloukos et al., 2021; Martín et al., 2021; Papageorgiou et al., 2021)

Quality of evidence Moderate

Grade of recommendation Grade A—↑↑ (1); Grade B—↑ (2)

Strength of consensus Consensus (10.6% of the group abstained due to potential Col)

Background

Intervention

Teeth with intra-bony defects are frequently present in stage IV periodontitis patients. These defects are not only a complexity factor for the periodontal therapy, but may also affect the outcomes of OT since tooth movements may occur through the regenerated tissues. As reported in the previously published S3-Level CPG of periodontal treatment for stage I–III periodontitis patients (Sanz, Herrera, et al., 2020), these intra-bony defects should be treated during the step 3 of periodontal therapy by surgical periodontal regenerative interventions. After this therapy, it is relevant to understand whether orthodontic

tooth movements can be implemented safely in these affected teeth, and what is the appropriate time interval between the surgical intervention and the start of active orthodontic tooth movements.

Available evidence

Number and design of included studies

Data from the three SRs (Kloukos et al., 2021; Martín et al., 2021; Papageorgiou et al., 2021) were retrieved from 16 studies including a total number of 683 patients: being three RCTs, 10 prospective and three retrospective case series studies.

Risk of bias

Using the Risk of Bias 2.0. tool, two RCTs were rated high risk of bias. The RCT comparing early versus delayed OT (Jepsen et al., 2021) was rated low risk of bias. The remaining 13 studies, using the ROBINS-I-tool, were rated as critical risk of bias in eight studies, as serious risk of bias in four, and as fair risk in one study.

Effect sizes and their clinical relevance

Twelve studies reported on mean PPD changes in mm, while in one study the mean PPD changes were reported in percentages. Outcomes from two RCTs indicated a PPD reduction of 2.17 mm (SD = 0.20) and 4.21 mm (SD = 1.35), respectively, while the results from non-RCTs indicated mean PPD reduction ranging from 0.07 mm (SD = 0.75) to 5.5 mm (SD = 1.50). A retrospective cohort study with 48 patients (Tietmann et al., 2021) reported a mean PPD reduction of 2.5 mm and a mean RBL gain of 4.7 mm after 12 months. Pocket closure (PPD ≤ 4 mm) was attained in 87% of defects.

Mean CAL changes were reported in eight studies. In two RCTs, reported gain in CAL was 3.09 mm (SD = 0.47) and 3.67 mm (SD = 0.76), respectively, while in non-RCTs CAL gain ranged from 0.29 mm (SD = 0.17) to 5.93 mm (SD = 1.41). The results from the RCT comparing OT starting early versus late, after the periodontal regenerative intervention, demonstrated a potential beneficial effect of the early OT, since no statistically significant differences were observed in terms of CAL gain (5.4 mm [SD = 2.1] for early; 4.5 mm [SD = 1.7]) for late OT, or PPD reduction [4.2 mm (SD = 1.9) in the early group versus 3.9 mm (SD = 1.5) in the late group ($p > .05$)]. Similarly, pocket closure (PPD ≤ 4 mm) occurred in 91% of the early OT treated teeth versus 85% in late OT (Jepsen et al., 2021).

Consistency

There was general agreement among included studies that in patients with severe periodontitis (stage IV or equivalent) and presence of intra-bony lesions, the combination of periodontal regenerative treatment with OT positively influenced the periodontal outcomes.

Balance of benefit and harm

Five articles reported on harms/adverse effects; no significant adverse effects were reported. Furthermore, the benefits of early OT (similar

outcomes with significantly reduced overall treatment time) seem to outweigh the potential risks.

From evidence to recommendation—additional considerations

Acceptability

Although there is no evidence from PROMs, the combination of periodontal and orthodontic therapies was well accepted by the patients.

Feasibility

The combination of complex periodontal and orthodontic therapies requires the coordinated efforts of different oral care providers (specialists or dentists with advanced training and skills in periodontal and orthodontic therapies) and these special settings may not be encountered in every oral health care environment.

Ethical considerations

There is no evidence for ethical considerations.

Economic considerations

There is no evidence from cost-effectiveness studies, although the combined periodontal–orthodontic therapy of these patients is complex, requires multiple care providers and it is usually costly.

Legal considerations

There are no specific legal considerations.

R7.6. How should we implement OT in stage IV periodontitis patients to maintain/improve periodontal outcomes?

PICOS question addressed by a SR

R7.6: Evidence-based recommendation

In patients with severe periodontitis (stage IV or equivalent) with indication of OT to maintain/improve periodontal stability:

1. **we suggest** using fixed rather than removable appliances;
2. use of fiberotomy as an adjunct to orthodontic tooth movement **may be considered** to improve periodontal outcomes;
3. use of skeletal anchorage devices (implants or temporary anchorage devices—micro-screws or mini-plates) **may be considered** to enhance orthodontic tooth movement.

Supporting literature (Martín et al., 2021; Papageorgiou et al., 2021)

Quality of evidence Low

Grade of recommendation Grade B—↑ (1), Grade O—↔ (2, 3)

Strength of consensus Consensus (1.7% of the group abstained due to potential CoI)

Background

Intervention

1. Orthodontic tooth movements can be carried out either by fixed (braces) or removable (able to be inserted/removed by the patients, such as removable plates, thermoplastic aligners, etc.) orthodontic appliances.
2. Circumferential fiberotomy of the supracrestal periodontal fibres has been suggested as an adjunct surgical procedure to improve the post-treatment stability after correction of severely rotated teeth or as an intervention aimed to improve attachment levels during orthodontic tooth intrusion.
3. In stage IV periodontitis patients with a healthy but reduced periodontium, the use of skeletal anchorage devices, compared with conventional anchorage systems, may improve the efficacy of OT and its effect on periodontal outcomes.

Available evidence

Number and design of included studies

1. One non-randomized retrospective comparative study comparing fixed appliances (braces) with aligners, reporting results from before periodontal to after OT, in terms of CAL/RBL/PPD/root length outcomes.
2. The use of fiberotomy as an adjunct to combined periodontal–orthodontic therapy was assessed in two within-individual RCTs.
3. Anchorage systems:
 - In 4 studies, skeletal anchorage (temporary anchorage devices, TADs, or implants) was used in all patients.
 - In 12 studies, different types of conventional anchorage were used.
 - 2 studies presented results of patients treated with TADs mixed with patients without TADs.

Risk of bias

1. Type of orthodontic appliance
The only included study was judged as serious risk of bias using the ROBINS-I tool.
2. Use of fiberotomy
Both randomized trials were judged as high risk of bias using the RoB 2.0 tool for at least two domains each.
3. Anchorage systems
The overall quality of the included studies was judged as “low”.
 - Considering the 12 pre–post intervention studies assessed, 2 were rated as “good”, 4 of them were rated as “fair” (moderate risk of bias) and 6 were rated as “poor”.
 - The study rated according to ROBINS-I tool received an overall assessment of “serious” risk of bias.
 - Considering the five RCTs assessed using RoB 2.0 tool, the overall bias assessment was “high” for two RCTs; one RCTs was rated as “low risk of bias” and another two were rated with “some concerns”.

Effect sizes and their clinical relevance

1. The included study reported that fixed appliances have benefits over aligners for PPD (1 study; MD = -1.64 mm; 95% CI [-2.50 to -0.78]; $p < .001$), but not for RBL (1 study; MD = 0.01; 95% CI [-1.01 to 1.03]; $p = .99$).
2. Adjunctive use of fiberotomy was associated with benefits in CAL (1 study; MD = -0.63 mm; 95% CI [-1.10 to -0.16]; $p = .009$) and RBL (2 studies; MD = -0.98 mm; 95% CI [-1.87 to -0.10]; $p = .03$), but not PPD (1 study; MD = -0.03 mm; 95% CI [-0.48 to 0.42]; $p = .90$).
3. In terms of different anchorage systems, there is no evidence of added benefits on periodontal parameters from using a specific type of anchorage system during OT. In terms of orthodontic outcomes, the use of osseointegrated implants as anchorage (one study), TADs (three studies) and conventional anchorage resulted in different tooth movements (such as intrusion) ranging between 1 and 7 mm, without a clear advantage among the different systems.

Root resorption after OT in periodontitis patients was reported in studies using different anchorage systems, with similar levels of reported resorption (between 0.2 and 3.0 mm) with either conventional anchorage or skeletal anchorage. Root resorption was also reported in similar ranges in studies with OT in healthy periodontal patients.

Consistency

In relation to the type of orthodontic appliance, since only one study was available, the consistency could not be assessed. For the adjunctive use of fiberotomy, the results were not consistent for the periodontal outcomes. The results using different anchorage systems reported the same tendency.

Balance of benefit and harm

There are no considerable harms for the periodontal tissues and other dental tissues and structures, with the use of conventional fixed appliances (braces), other than the increased microbial burden, related to the difficulties in oral hygiene practices, which can however be managed appropriately. Similarly, there is no evidence of significant harms associated with the different anchorage systems.

Although fiberotomy is a surgical procedure, the possible side effects have not been adequately reported.

Overall certainty of the evidence

Low.

From evidence to recommendation—additional considerations

Acceptability

1. Some patients might prefer aligners over fixed appliances, due to their enhanced aesthetics and easiness for oral hygiene. However, braces have been historically well accepted by adults and there are

- currently more aesthetic alternatives in fixed appliances (ceramic/lingual appliances).
2. Fiberotomy is a surgical intervention, albeit minimally invasive, what might be less acceptable by some patients.
3. Some patients might experience discomfort when receiving TADs, although this fact has not been evaluated in the reported studies.

Feasibility

1. The use of fixed appliances requires specific knowledge of applied biomechanics, what is usually rendered by orthodontic specialists, while aligners treatment is mostly facilitated through third parties (companies or external laboratories) producing the aligners and enabling both specialists and general dentists to provide this kind of treatment. However, extensive knowledge of tooth biology and applied biomechanics is still a pre-requisite for treatment planning and monitoring.
2. Fiberotomy requires specific surgical experience from the operator.
3. In cases of using skeletal anchorage systems, knowledge of surgical anatomy and surgical skills are needed.

Ethical and economic considerations

Fiberotomy, as an additional surgical insult to the periodontal tissues, must be further evaluated in terms of its long-term effectiveness and since it is an additional surgical intervention, it might be associated with further costs.

Legal considerations

There are no specific legal considerations.

R7.7. How should we manage stage IV periodontitis patients during and after the completion of OT to prevent recurrence of periodontitis?

Additional question addressed by the WG

R7.7: Expert consensus-based recommendation

We recommend that during OT the patient's periodontal status is closely monitored and managed, ideally at each orthodontic appointment. If signs of periodontitis recurrence are detected, active OT should be interrupted, and the affected teeth should be maintained passively, while rendering proper periodontal treatment and oral hygiene reinforcement. Once periodontal health/stability has been re-established, OT can be re-instituted.

We recommend that after completion of OT, life-long supportive periodontal care and life-long orthodontic retention are provided tailored to the individual needs/risk profile of the patient.

Supporting literature (Arn et al., 2020; Jiang et al., 2021) and Expert opinion

Quality of evidence Low (expert opinion)

Grade of recommendation Grade A-↑↑

Strength of consensus Unanimous consensus (0% of the group abstained due to potential CoI)

Background

Intervention

The use of orthodontic appliances is associated with increased microbial colonization and increased plaque retention; hence, OT is often associated with gingival inflammation and a transient shift of the subgingival microbiota. It is, therefore, important to implement an appropriate oral hygiene and periodontal management protocol throughout OT to ensure periodontal health and avoid adverse effects like enamel demineralization, tooth discolorations, and further loss of periodontal support. Although, in some patients, an acceptable oral hygiene level can be attained, professional plaque control and other supportive oral and periodontal care must be implemented following the patient's risk profile.

Available evidence

Number and design of included studies

The evidence for this recommendation is derived from expert opinion and from a randomized trial with 48 patients, comparing three different periodic periodontal scaling protocols (every month, every 3 months or every 6 months) for the maintenance of periodontal health in adolescents with fixed orthodontic appliances (Jiang et al., 2021). A systematic review (Arn et al., 2020) assessed the available evidence in the literature for the effects of fixed orthodontic retainers on periodontal health. It included 11 RCTs, 4 prospective cohort studies, 1 retrospective cohort study and 13 cross-sectional studies.

Risk of bias

Not applicable.

Effect sizes and their clinical relevance

One RCT reported that periodontal scaling conducted monthly or once every 3 months during OT was more effective (in terms of PI, GI, PPD and crevicular inflammation biomarkers) than scaling administered once every 6 months. The systematic review concluded that the use of fixed orthodontic retainers was compatible with periodontal health, or at least not associated with detrimental effects on the periodontium.

Even though the RCT and the systematic review studied periodontally healthy patients and not adults with stage IV periodontitis, it is expected that these findings can be reasonably extrapolated.

Consistency, balance of benefit and harm

Not applicable.

Overall certainty of the evidence

Low, expert opinion.

From evidence to recommendation—additional considerations

Acceptability

Patients usually accept and understand the need for long-term supportive care.

Feasibility, ethical, economic and legal considerations

There are no specific considerations.

R7.8. How should we attain stability of the orthodontically treated dentition in stage IV periodontitis patients?

Additional question addressed by the WG

R7.8: Expert consensus-based recommendation

We **recommend** that an appropriately designed permanent fixed passive retention (with or without additional removable retention) is used after the completion of orthodontic therapy.

We also **recommend** that patients enter a life-long supporting protocol to identify early retainer failures and/or undesired tooth movements.

Supporting literature (Han et al., 2020) and Expert opinion

Quality of evidence Low (expert opinion)

Grade of recommendation Grade A—↑↑

Strength of consensus Consensus (0% of the group abstained due to potential Col)

Background

Intervention

Post-orthodontic relapse of the moved teeth towards their pre-treatment positions is often observed and can have aesthetic and functional consequences that compromise treatment outcomes and patient satisfaction. Therefore, some type of retention, either with removable or fixed appliances, is usually implemented depending on the pre-existing malocclusion, the type of tooth movements carried out, and patient preferences. Fixed retention is usually considered superior to removable appliance in terms of reducing post-treatment relapse, especially in anterior crowding (Littlewood et al., 2016). However, at the same time, it may be more prone to retention failure in the short or the long term, might lead to more plaque accumulation or gingival inflammation, and might even lead to inadvertent tooth movements due to the distortion of the bonded wire. It is, therefore, advisable that fixed retention (with additional removable appliances, if needed) is used after OT in patients with stage IV periodontitis, but a long-term supportive care protocol should be implemented to assess not only periodontal health, but the integrity of the retention appliances and the stability of the treatment outcomes.

Available evidence

Number and design of included studies

The evidence for this recommendation is derived from expert opinion and from a retrospective cohort study with 52 patients followed at least 2 years after OT (Han et al., 2020).

Risk of bias

Not applicable.

Effect sizes and their clinical relevance

Han et al. (2020) reported that orthodontic fixed retainers fail more often in stage III periodontitis patients compared with stage I periodontitis patients. This study did not include stage IV periodontitis patients, but these findings indicate that increased periodontitis severity was associated with higher retainer failure, and hence, in stage IV periodontitis patients, the need for frequent recalls to evaluate the retainer's integrity should be highlighted.

Consistency, balance of benefit and harm

Not applicable.

Overall certainty of the evidence

Low.

From evidence to recommendation—additional considerations

Acceptability

Patients usually accept and understand the need for long-term fixed passive retention and regular monitoring.

Feasibility, ethical, economic and legal considerations

There are no specific considerations.

8 | CLINICAL RECOMMENDATIONS: OVERALL STRATEGY FOR THE MANAGEMENT OF CASE TYPES 3 AND 4

In stage IV periodontitis patients, prosthetic rehabilitation requires adherence to both periodontal and reconstructive treatment principles. Such principles need to be tailored to the specific needs of this patient group. Several key issues need to be addressed, including:

1. Identification of the restorative needs with an emphasis on limiting the scope of the prosthetic treatment while ensuring patient comfort and stability.
2. Identification of the need for interim dental prostheses.
3. Timing of delivery of interim and definitive dental prostheses.
4. Need and timing of dental implant placement.

This section aims to provide recommendations for the general principles of case management in patients with a compromised dentition due to stage IV periodontitis requiring rehabilitation.

R8.1. How important is it to identify the restorative needs of the individual case in stage IV periodontitis patients?

Additional question raised by the WG

R8.1: Expert consensus-based recommendation

We *recommend* identifying the restorative needs of partially edentulous stage IV periodontitis patients based on the pattern of tooth loss, individual functional and aesthetic needs, patient comfort and prognostic factors. The level of function and design of the rehabilitation should be compatible with case stability over time.

Supporting literature Expert opinion

Quality of evidence Not applicable

Grade of recommendation Grade A—↑↑

Strength of consensus Strong consensus (0% of the group abstained due to potential Col)

Background

Stage IV periodontitis cases are characterized by high levels of phenotypic variation. While restorative needs to achieve case stability and address patient comfort are only one dimension of what needs to be considered in treatment planning, precise identification of these needs is critical. Treatment goals need to be based on the individual patient, its unique pattern of tooth loss, tooth prognosis both in terms of periodontal maintainability and restorative factors. Furthermore, the treatment plan should be able to stop or mitigate greatly the occlusal and functional aspects that contribute to the instability of the case and loss of patient comfort.

R8.2. Is there a need, and what is the timing of interim restorations in stage IV periodontitis patients?

Additional question raised by the WG

R8.2: Expert consensus-based recommendation

We *recommend* placing an interim dental prosthesis, if required, early during periodontal therapy, but only after establishing adequate oral hygiene.

Supporting literature (Donos et al., 2021; Montero et al., 2021; Ramanauskaitė et al., 2021; Tomasi et al., 2021) and Expert opinion

Quality of evidence Very low

Grade of recommendation Grade A—↑↑

Strength of consensus Strong consensus (1.9% of the group abstained due to potential Col)

Background

Once the restorative needs of the case have been identified, interim restorations may be required to provide stable posterior occlusal stops, relieve secondary occlusal trauma, replace functionally important missing teeth, or improve patient comfort. Ideally, the decision to place an interim restoration should be based on the individual response of periodontal therapy as assessed in the periodontal re-evaluation after completion of step 2 treatment. Frequently, the benefits to the treatment plan from interim restorations require their insertion as early as possible. Flexibility to move forward the interim restoration in the treatment plan, however, should not compromise the need to achieve adequate supragingival biofilm control in the context of step 1 treatment.

R8.3. Which are the general principles for the design and delivery of dental prostheses in stage IV periodontitis patients?

Additional question raised by the WG

R8.3: Expert consensus-based recommendation

We **recommend** designing the dental prostheses to allow optimal self-performed oral hygiene measures and professional mechanical plaque removal.

We **recommend** delivery of the definitive prosthesis after a final evaluation of the maintainability and prognosis of abutment teeth/implants.

Supporting literature (Donos et al., 2021; Montero et al., 2021; Ramanauskaitė et al., 2021; Tomasi et al., 2021) and Expert opinion

Quality of evidence Very low

Grade of recommendation Grade A–↑↑

Strength of consensus Unanimous consensus (0% of the group abstained due to potential Col)

Background

Oral biofilm control is important for the longevity of the dentition and restorations. This is critically important in subjects with stage IV periodontitis who have shown high susceptibility to periodontitis. Restorations may render access and effectiveness of self-performed biofilm control and professional mechanical plaque removal (PMPR) more challenging. While restorations are primarily designed to restore function, they should enable optimal biofilm control at the critical interface between the restoration, the tooth/root surface, the implant on one side and the soft tissues (gingiva, peri-implant or alveolar ridge mucosa) on the other. Planning and execution of these restorations should preserve the necessary access for oral hygiene aids including inter-dental brushes or dental floss, as well as instruments for PMPR (both supra- and sub-gingivally). This should be prioritized compared with avoidance of food impaction. The design should be finalized with the interim dental

prosthesis and achievement of tissue health needs to be verified along with the maintainability of the abutments before insertion of the final prosthesis. Insertion of the final dental prosthesis needs to follow the achievement of the treatment goals for natural teeth and/or implants at the completion of active periodontal therapy.

R8.4. Which are the general considerations when incorporating dental implants in stage IV periodontitis patients?

Additional question raised by the WG

R8.4: Expert consensus-based recommendation

When dental implants are considered in the rehabilitation of stage IV periodontitis patients, we **recommend** verifying (i) absence of contraindications to surgery, (ii) hard and soft tissue dimensions and (iii) the potential need for soft-/hard-tissue augmentation.

Supporting literature Expert opinion

Quality of evidence Not applicable

Grade of recommendation Grade A–↑↑

Strength of consensus Consensus (3.5% of the group abstained due to potential Col)

Background

Dental implants are frequently considered in the rehabilitation of stage IV periodontitis patients to replace missing teeth and restore function. The use of a dental implant, however, may present different levels of complexity and range from a simple procedure in a subject with no medical contraindications and adequate dimensions of soft and hard tissues, to a complex procedure requiring specific medical assessments and/or challenging local hard and/or soft tissue augmentation. As the complexity of the dental implant placement increases, its clinical performance may decrease, affecting the cost-benefit ratio and its attractiveness with respect to alternatives. Such considerations are part of the strategic assessment that concur in the determination of the treatment plan for the individual case.

R8.5. Which are the specific considerations when incorporating dental implants in stage IV periodontitis patients?

PICOS question addressed by a SR

R8.5: Expert consensus-based recommendation

When dental implants are considered in the rehabilitation of stage IV periodontitis patients, we **recommend** that information on the increased risk for peri-implantitis and implant loss should be provided.

Supporting literature (Lindhe, Meyle, & Group, 2008; Carra et al., 2021) and Expert opinion

Quality of evidence Low

Grade of recommendation Grade A–↑↑

Strength of consensus Consensus (23.8% of the group abstained due to potential Col)

Background

Substantial evidence indicates that subjects with advanced/rapidly progressing forms of periodontitis have a higher risk of implant loss and peri-implantitis, when compared with the general population, or with individuals without a history of periodontitis (Schwarz et al., 2018; Carra et al., 2021). While some of the risk seems to be associated with placement of dental implants before achievement of full control of periodontitis and, perhaps, with subjects with nicotine addiction who cannot quit cigarette smoking, the available evidence does not allow excluding that a portion of the increased risk associated with periodontitis may persist even after appropriate periodontal treatment. The assumption that the health and function of dental implants in adequately treated and well-maintained stage IV periodontitis patients parallels the longevity observed for dental implants in the general population is probably optimistic.

The systematic review by Carra et al. (2021) was based on 7 prospective and 10 retrospective studies and accounted for 1718 implants placed in patients with a history of periodontitis and 2879 implants placed in patients with no history of periodontitis to restore. Eight out of the 17 studies (47%) included both non-smokers and smokers, the latter group representing 1.7%–28.8% of the patient sample. One study considered non-smokers only, and the remaining seven studies did not provide any detail about smoking habits. The type of supportive periodontal/implant care was reported in only 11 of 17 studies (64.7%), but the patient's compliance to the recall intervals was rarely defined. Although the overall mean implant survival rate, at a follow-up of at least 5 years, was high also in patients with a history of periodontitis (94.7%, 95% CI [92.3; 97.1]), pooled data analysis demonstrated that implant-supported fixed dental prostheses have a greater risk of failure (risk ratio—RR: 1.9, 95% CI [1.31; 2.79]) and peri-implantitis (RR: 3.3, 95% CI [1.31; 8.3]) in patients with a history of periodontitis, compared with patients with no history of periodontitis. The magnitude of the risk was considered clinically relevant, particularly after 5 years of follow-up from implant loading (hazard ratio—HR: 2.11, 95% CI [1.18; 3.79]).

R8.6. Which are the specific considerations for the design of dental implants restorations in stage IV periodontitis patients?

PICOS question addressed by a SR

R8.6: Expert consensus-based recommendation

Due to the risk of tooth loss and prosthesis failure, we **suggest** avoiding combined tooth/implant-supported fixed partial dental prostheses if alternatives are feasible.

Supporting literature (Montero et al., 2021) and Expert opinion

Quality of evidence Low

Grade of recommendation Grade B—↑

Strength of consensus Unanimous Consensus (7.8% of the group abstained due to potential Col)

Background

Long-term studies have shown that dental prostheses retained by a combination of teeth and dental implants show higher rates of failure and tooth loss compared with dental prostheses supported by teeth or implants alone. In treatment planning, whenever alternatives exist, such designs should be avoided.

9 | CLINICAL RECOMMENDATIONS: CASE TYPE 3

Case type 3: partially edentulous patients who can be prosthetically restored without full-arch rehabilitation.

9.1 | Clinical recommendations for case type 3 with tooth-delimited gaps

R9.1. What is the effectiveness of prosthetic rehabilitation in patients with periodontitis stage IV where tooth preservation is feasible, with one or multiple tooth-delimited gaps, and adequate residual periodontal support and maintainability of the remaining teeth?

(Common question for Recommendations R9.1, R9.2, R9.3, R9.4)

PICOS question addressed by a SR

R9.1: Evidence-based recommendation

In partially edentulous stage IV periodontitis patients with tooth-delimited edentulous spaces, different options (namely tooth-supported fixed dental prostheses, implant-supported fixed dental prostheses, removable dental prostheses, or no prosthetic rehabilitation) **may be considered**.

Supporting literature (Carra et al., 2021; Gotfredsen et al., 2021; Montero et al., 2021)

Quality of evidence Low

Grade of recommendation Grade O—↔

Strength of consensus Consensus (1.4% of the group abstained due to potential Col)

Background

Intervention

Interventions include tooth-supported fixed dental prostheses, implant-supported fixed dental prostheses, removable dental prostheses (RDPS), or no prosthetic rehabilitation, in periodontitis stage IV patients with one or multiple tooth-delimited gaps, and adequate residual periodontal support and maintainability of the remaining teeth.

Available evidence

Number and design of included studies

For **implant-supported fixed dental prostheses (iFDPs)** (Carra et al., 2021), evidence is based on 7 prospective and 10 retrospective studies, reporting on 1718 implants placed in patients with a history of periodontitis and 2879 implants placed in patients with no history of periodontitis.

For **tooth-supported fixed dental prostheses (tFDPs)** (Montero et al., 2021), evidence is based on one RCT, one CCT, four prospective case series and 14 retrospective case series, including data from 1029 subjects at baseline, with a total of 1037 tFDPs, including 3186 abutment teeth. The follow-up ranged between 24 and 425 months. At the end of the follow-up period, 933 subjects with 915 tFDPs on 2989 abutment teeth were analysed.

For **removable dental prostheses (RDPs)** (Gotfredsen et al., 2021), evidence is based on three prospective (one RCT/two non-randomized) and one retrospective study that specifically reported having also included Kennedy class III or IV RDPs (including minimum 175 to maximum 234, in total), and reporting on 1–5 years results, in either exclusively periodontitis patients or in mixed periodontitis/non-periodontitis study populations. Several studies did not specify the type of Kennedy class.

Risk of bias

For **iFDPs**, 10 of 17 studies (58.8%) were considered at low risk of bias (Newcastle–Ottawa scale). Funding sources and conflict of interest declaration were not reported in the majority of studies.

For **tFDPs**, all investigations ($n = 20$) presented an unclear or high risk of bias, using the RoB 2.0 tool, the ROBINS-I tool or the Newcastle–Ottawa scale.

For **RDPs**, all four publications that specifically reported having also included Kennedy class III or IV RDPs were judged being of moderate risk of bias (the RCT with the RoB 2.0 tool and the three non-randomized studies with the Newcastle–Ottawa scale). Funding sources and conflict of interest declaration were inconsistently reported.

Effect sizes and their clinical relevance

For **iFDPs**, although the risk of peri-implantitis and implant loss is greater in patients with a history of periodontitis compared with subjects with no history of periodontitis, the overall mean implant survival rate at a follow-up of at least 5 years is favourable (94.7% [92.3%–97.1%]) and acceptable to counterbalance the risks.

For **tFDPs**, the incidence of abutment tooth loss after a follow-up period from 2 to 35.4 years was low ($n = 17$ studies; weighted mean incidence (WMI) = 4.8%; 95% CI [3.2; 6.5]). The corresponding figure for prosthesis failure was WMI = 6.9% ($n = 18$ studies; 95% CI [4.1; 9.7]). Therefore, tFDPs seemed to be a valid alternative to restore function in patients with stage IV periodontitis.

For **RDPs**, in one publication (a prospective cohort study), out of the four publications that specifically reported having also included Kennedy class III or IV RDPs, an abutment tooth failure rate (defined as abutment tooth restoration or loss) ranging from 16% to 48% was

reported; however, it was not possible to attribute specific rates to a particular Kennedy class or type of RDP. In another RCT, specifically reporting on Kennedy class III or IV RDPs, an abutment failure rate of 9%–15% was reported. However, it was not possible to attribute specific rates to a particular Kennedy class; the risk for failure was higher at non-vital compared with vital abutment teeth (HR = 2.29).

Consistency

For **iFDPs**, sensitivity analysis indicated consistency.

For **tFDPs**, a significant publication bias was observed for the main outcome measure (i.e., tooth loss).

For **RDPs**, no analysis of the consistency was possible based on the available evidence.

Balance of benefit and harm

Specific analyses on benefit/risk ratios, comparing different options for the rehabilitation of unilateral or bilateral posterior free-end edentulism, were not the focus of the SRs.

For **iFDPs**, potential harms related to surgery (e.g., intra- and post-operative complications) and peri-implant health maintenance (e.g., higher risk of peri-implantitis and implant loss over time) should be considered.

For **tFDPs**, technical complications (e.g., loss of retention, framework fracture, porcelain fracture, etc.) presented a WMI = 13.6% ($n = 10$ studies; 95% CI [8.3; 18.9]) and seemed to be more frequent than biologic complications (e.g., caries, endodontic failure, root fracture, etc.) that presented a WMI = 5.1% ($n = 7$ studies, 95% CI [2.5; 7.8]).

RDPs do not necessarily cause further periodontal breakdown or tooth loss. There is no information in the studies (having also included Kennedy class III or IV RDPs) on whether masticatory efficiency, including nutritional status, or OHRQoL were improved.

Overall certainty of the evidence

For **iFDPs**, the certainty of evidence is moderate.

For **tFDPs**, the certainty of evidence is low.

For **RDPs**, the certainty of evidence is low.

From evidence to recommendation—additional considerations

Acceptability

iFDPs are acceptable, and a widespread option for rehabilitation of partial edentulism in patients with stage IV periodontitis. Little is known about patients' preferences and satisfaction.

Patient satisfaction was considered in seven studies evaluating **tFDPs**, using questionnaires related to chewing function, aesthetics, phonetics, comfort and/or general satisfaction. In general, more than 85% of the patients were satisfied with the treatment provided or reported positive results for the different outcomes evaluated.

RDPs are acceptable and a widespread option for rehabilitation of partial edentulism in patients with stage IV periodontitis. Little is known about patients' preferences and satisfaction.

Feasibility

iFDPs are demanding in terms of professional competency and resources.

tFDPs are demanding in terms of professional competency and resources.

RDs are often less demanded compared with **tFDPs** or **iFDPs**.

Ethical considerations

Not applicable.

Economic considerations

iFDPs come with additional costs that are not covered by healthcare systems in most countries and may be a source of inequality.

The economic considerations related to **tFDPs** have not been properly evaluated. In any case, they may imply additional costs that are not covered by healthcare systems in most countries and may be a source of inequality.

RDs are more economical compared with **tFDPs** or **iFDPs** and often covered (at least partially) by most healthcare systems.

Legal considerations

Not applicable.

R9.2. For question, see R9.1

PICOS question addressed by a SR

R9.2: Evidence-based recommendation

We **suggest** using tooth-supported fixed dental prostheses in patients with stage IV periodontitis when abutment teeth are periodontally maintainable and restorable.

Resin-bonded fixed dental prostheses **may be considered** in certain circumstances (e.g., small tooth-delimited edentulous spaces).

We **suggest not to use** resin-bonded fixed dental prostheses for large tooth-delimited edentulous spaces.

Supporting literature (Montero et al., 2021)

Quality of evidence Low

Grade of recommendation Grade B-↑/Grade O-↔/Grade B-↑

Strength of consensus Consensus (4.9% of the group abstained due to potential Col)

Background

Intervention

For tooth-supported fixed dental prostheses (**tFDPs**), see the background text of Recommendation R9.1.

In certain cases, for example, in small, tooth-delimited edentulous spaces, minimally- or non-invasive resin-bonded fixed dental prostheses (**FDPs**) can be considered.

Available evidence

Number and design of included studies

Two case series (one prospective and one retrospective) evaluating 80 patients and 99 resin-bonded **FDPs** in mandibular anterior teeth.

Risk of bias

Both studies presented a high risk of bias according to the Newcastle–Ottawa scale.

Effect sizes and their clinical relevance

No tooth loss was reported, but a high incidence of prostheses failure (WMI = 27.4%, 95% CI [-6.7; 61.4]) was observed. Therefore, this treatment option may be considered only in specific clinical scenarios (e.g., missing single anterior teeth).

Consistency

No analysis of the consistency was possible based on the available evidence.

Balance of benefit and harm

Even if prosthetic complications may be expected in resin-bonded **FDPs**, most of the cases were solved with minor reparations. Moreover, it must be highlighted that teeth used as abutment were minimally prepared.

Overall certainty of the evidence

Low.

From evidence to recommendation—additional considerations

Acceptability

General satisfaction and aesthetics were evaluated in both studies with high rates for both outcomes (over 8.5 of 10 in visual analogue scales).

Feasibility

Resin-bonded **FDPs** are demanding in terms of professional competency and resources.

Ethical considerations

Not applicable.

Economic considerations

Resin-bonded **FDPs** may be a more affordable alternative to other tooth-supported **FDPs**.

Legal considerations

Not applicable.

R9.3. For question, see R9.1**PICOS question addressed by a SR****R9.3: Evidence-based recommendation**

We **suggest** using implant-supported fixed dental prostheses when abutment teeth are not periodontally maintainable and restorable.

Supporting literature (Carra et al., 2021)

Quality of evidence Low

Grade of recommendation Grade B—↑

Strength of consensus Strong consensus (2.1% of the group abstained due to potential Col)

Background**Intervention**

Rehabilitation with iFDP, in periodontitis stage IV partially edentulous patients, with one or multiple tooth-delimited gaps, and abutment teeth that are not periodontally maintainable and restorable.

For further detail, see the sections dealing with implant-supported fixed dental prostheses in the background text for Recommendation R9.1.

R9.4. For question, see R9.1**PICOS question addressed by a SR****R9.4: Evidence-based recommendation**

Metal-based frame removable dental prostheses may be considered as transitional or definitive treatment options when a fixed solution is not a consideration.

Supporting literature (Gotfredsen et al., 2021)

Quality of evidence Low

Grade of recommendation Grade O—↔

Strength of consensus Consensus (0% of the group abstained due to potential Col)

Background**Intervention**

Patients diagnosed with stage IV periodontitis frequently require an interim RDP to ensure chewing function by increasing the number of occlusal units (antagonistic pairs in pre-molar and molar region) and/or to replace anterior teeth for aesthetic reasons.

Steps 1 and 2 of periodontal therapy are accompanied by a reduction in inflammation and swelling, which becomes frequently visible as gingival recession, while tooth extractions lead to remodelling of the alveolar bone associated with morphological tissue changes. In the light of this background, any interim RDP has to be designed with an occlusal rest on the residual dentition to prevent the prosthesis from sinking into the mucosal tissue and subsequent trauma to periodontal tissues and teeth. Before taking the initial impression or intra-oral scan, the inter-maxillary situation needs to be evaluated and teeth with a good prognosis are considered as

abutment teeth. Depending on the occlusal situation and the available space, abutment teeth should be provided with a shallow occlusal rest preparation. On the resulting situation model, which is used to fabricate the interim RDP, teeth to be extracted are removed in plaster, and clasps are manually bent. These clasps with occlusal rests ensure sufficient stability during the phase of tissue remodelling, after which relining of the prosthesis basis or fabrication of a definitive RDP is frequently indicated.

After successful periodontal therapy and when post-extraction tissue changes have occurred, the interim prosthesis can be replaced by a definitive RDP. For retention of the RDP, 2–4 abutment teeth with good prognosis and a wide spread (distributed anterior and posterior) are used. The retention element is selected depending on the condition of the coronal tooth substance, aesthetic demands and financial conditions:

- An intact, caries-free tooth is best provided with a clasp or an adhesive attachment (Zitzmann et al., 2009). While a clasp abutment tooth can have fillings of small extent, a tooth intended for an adhesive element with an extra-coronal attachment should have sound enamel surfaces to ensure adhesive cementation. These resin-bonded attachments for precision-retained RPD are compatible with periodontal health, ensure occlusal rests, require minimal tooth preparation and provide very good aesthetics results. To ensure periodontal health, clasps are designed with vertical support by an occlusal rest (ideally located on sound tooth structure), and a stiff reciprocal part neutralizing lateral forces during prosthesis insertion and removal.
- Decayed teeth and those with extended fillings are better restored with telescopes or crowns with clasps (Zitzmann et al., 2009).
- For abutment teeth with a destroyed clinical crown, root canal treatment is frequently required and the root cap provided with a post and a retentive element is the most appropriate solution to retain the RDP (overdenture).

For further detail, see the sections dealing with RDP in the background text for Recommendation R9.1.

9.2 | Clinical recommendations for case type 3 with unilateral or bilateral posterior free-end edentulism**R9.5. In patients with a periodontally compromised dentition due to stage IV periodontitis or equivalent, what is the efficacy of various prosthetic restorative options for the rehabilitation of unilateral or bilateral posterior free-end edentulism?**

(Common question for Recommendations R9.5, R9.6, R9.7)

PICOS question addressed by a SR**R9.5: Evidence-based recommendation**

For the rehabilitation of partially edentulous stage IV periodontitis patients with free-end situations, different options (namely shortened

(Continues)

(Continued)

PICOS question addressed by a SR**R9.5: Evidence-based recommendation**

dental arch, implant-supported restorations or removable dental prostheses) may be considered.

Supporting literature (Carra et al., 2021; Gotfredsen et al., 2021; Montero et al., 2021)

Quality of evidence Low

Grade of recommendation Grade O—↔

Strength of consensus Strong consensus (0% of the group abstained due to potential CoI)

Background**Intervention**

Interventions include planning towards a shortened dental arch, implant-supported fixed dental prostheses (iFDPs) or RDPs, in periodontitis stage IV patients with unilateral or bilateral posterior free-end edentulism. See also the background text of Recommendation R9.1.

Available evidence*Number and design of included studies*

For **iFDPs**, evidence is based on 7 prospective and 10 retrospective studies, including accounting for 1718 implants placed in patients with a history of periodontitis and 2879 implants placed in patients with no history of periodontitis.

For the **shortened dental arch**, the evidence is based on seven RCTs and five non-randomized studies (four prospective and one retrospective), including from 10 to 79 patients per group at final examination, either having shortened dental arch or restored to shortened dental arch; however, most of the publications included <50 patients/RDPs per group. Publications report mostly on 1-year data (one study presents 10-year results), in either periodontitis patients or mixed periodontitis/non-periodontitis study populations.

For **RDPs**, evidence is based on 12 RCTs including from 19 to 79 patients/RDP per group at the final evaluation, six prospective cohort studies including from 10 to 703 patients/RDP per group at the final evaluation, and three retrospective studies, including from 15 to 25 patients/RDP per treatment group; however, most of the publications included <50 patients/RDP per group. Most studies reported on Kennedy class I and II. The studies reported on 1 to 10-year results, with several reporting on 5-year data, in either periodontitis patients or periodontitis/non-periodontitis study populations.

Risk of bias

For **iFDPs**, 10/17 studies (58.8%) were considered at low risk of bias (Newcastle–Ottawa scale). Funding sources and conflict of interest declaration were not reported in the majority of studies.

For the **shortened dental arch**, five out of seven RCTs present some concerns regarding risk of bias (RoB 2.0 tool), while among the five non-randomized studies, one was judged of high risk, four were judged of moderate risk, and only one was judged of low risk (Newcastle–Ottawa scale).

For **RDPs**, in 10 out of 12 RCTs, some concerns regarding risk of bias (RoB 2.0 tool) were found, while in the nine non-randomized studies, two were judged of high risk, six were judged of moderate risk, and only one was judged of low risk (Newcastle–Ottawa scale). Funding sources and conflict of interest declaration were inconsistently reported.

Effect sizes and their clinical relevance

For **iFDPs**, although the risk of peri-implantitis and implant loss is greater in patients with a history of periodontitis, compared with patients with no history of periodontitis, the overall mean implant survival rate at a follow-up of at least 5 years is favourable (94.7%, 95% CI [92.3–97.1]) and acceptable to counterbalance the risks.

For the **shortened dental arch**, two studies show a lower risk for tooth loss compared with patients with an RDP (odds ratio, OR = 1.92; HR = 1.24), but it was not consistently significant. In a few studies, the shortened dental arch is associated with significantly lower plaque values and/or gingival inflammation compared with RDPs, and less gingival recession, comparing to abutment teeth of RDPs. A shortened dental arch does not necessarily imply lower masticatory efficiency, including nutritional status, or worse OHRQoL compared with patients with RDPs.

For **RDPs**, abutment tooth failure rates of 9%–48% have been reported in two studies comparing different types of RDPs, but these were not possible to attribute to specific Kennedy class and the differences were not significant. Two studies show a higher risk for tooth loss in patients with an RDP compared with no treatment (OR = 1.92; HR = 1.24), but it was not consistently significant. In a few studies, RDPs are associated with significantly more plaque values and/or gingival inflammation, and more recession at abutment teeth with clasps, but not necessarily with more significant other periodontal problems. RDPs do not necessarily increase masticatory efficiency, including nutritional status, or improved OHRQoL compared with a short dental arch.

Consistency

For **iFDPs**, sensitivity analysis indicated consistency.

For the **shortened dental arch**, no analysis was possible.

For **RDPs**, no analysis was possible.

Balance of benefit and harm

Specific analyses on the benefit/risk ratios comparing different options for the rehabilitation of unilateral or bilateral posterior free-end edentulism were not the focus of the SRs.

For **iFDPs**, potential harms related to surgery (e.g., intra- and post-operative complications) and peri-implant health maintenance

(e.g., higher risk of peri-implantitis and implant loss over time) should be considered.

Preserving- or restoring to a **shortened dental arch**, does not necessarily cause further periodontal breakdown or tooth loss, but is also not necessarily associated with reduced masticatory efficiency or nutritional status, or reduced OHRQoL.

RDPs do not necessarily cause further periodontal breakdown or tooth loss but are also not necessarily associated with increased masticatory efficiency, including nutritional status, or improve OHRQoL.

Overall certainty of the evidence

For **iFDP**, the certainty of evidence is moderate.

For the **shortened dental arch**, the certainty of evidence is low.

For **RDPs**, the certainty of evidence is low.

From evidence to recommendation—additional considerations

Acceptability

iFDPs are acceptable, and a widespread option for rehabilitation of partial edentulism in patients with periodontitis. Little is known about patients' preferences and satisfaction.

No restorative treatment or restoring to a **shortened dental arch** intervention is an acceptable treatment option associated with little risks. Little is known about patients' preferences and satisfaction.

RDPs are acceptable and a widespread option for rehabilitation of partial edentulism in patients with periodontitis. Little is known about patients' preferences and satisfaction.

Feasibility

iFDPs are demanding in terms of professional competency and resources.

Restoring to a **shortened dental arch** may not be a very demanding treatment option.

RDPs are less demanding comparing to tooth- or implant-supported fixed dental prostheses.

Ethical considerations

Not applicable.

Economic considerations

iFDPs imply additional costs that are not covered by healthcare systems in most countries and may be source of inequality.

No restorative treatment or restoring to a **shortened dental arch** may be the most economical solution.

RDPs are more economical comparing to tooth- or implant-supported fixed dental prostheses and often covered (at least partially) from most healthcare systems.

Legal considerations

Not applicable.

R9.6. For question, see R9.5

PICOS question addressed by a SR

R9.6: Evidence-based recommendation

*In stage IV periodontitis patients with a shortened dental arch with sufficient occluding/masticatory units (e.g., from second pre-molar to second pre-molar, no evident risk for flaring or tooth elongation and adequate patient comfort) no tooth replacement **may be considered** in the free-end situation.*

Supporting literature (Kayser, 1981; Walter et al., 2018; Walter et al., 2020; Gotfredsen et al., 2021)

Quality of evidence Low

Grade of recommendation Grade O—↔

Strength of consensus Simple majority (0% of the group abstained due to potential Col)

Background

Intervention

No restorative treatment in situations with sufficient occluding/masticatory units. See the background text for Recommendation R9.5.

R9.7. For question, see R9.5

PICOS question addressed by a SR

R9.7: Evidence-based recommendation

*In stage IV periodontitis patients with free-end situations who require additional occluding units, we **suggest** implant-supported fixed dental prostheses.*

*When implants are not an option, we **suggest** removable dental prostheses with a metal-based framework.*

Supporting literature (Carra et al., 2021; Gotfredsen et al., 2021)

Quality of evidence Low

Grade of recommendation Grade B—↑/Grade B—↑

Strength of consensus Strong consensus (1.9% of the group abstained due to potential Col)

Background

Intervention

Interventions include implant-supported fixed dental prostheses or RDPs with a metal-based framework, to restore free-end partial edentulism in periodontitis stage IV patients. See the background text of Recommendation R9.1 for information on implant-supported fixed dental prostheses, and the background text of Recommendation R9.5 for information on RDPs, respectively.

10 | CLINICAL RECOMMENDATIONS: CASE TYPE 4

Case type 4: partially edentulous patients who need to be restored with full-arch rehabilitation, either tooth- or implant-retained.

R10.1. In patients with a compromised dentition due to stage IV periodontitis with a sufficient number of adequately distributed teeth, what is the performance of tooth-supported full-arch fixed prostheses?

PICOS question addressed by a SR

R10.1: Evidence-based recommendation

In patients with periodontitis stage IV with a sufficient number (≥ 4 abutment teeth) of periodontally maintainable, bilaterally distributed and restorable teeth in the maxilla and/or mandible, we suggest a tooth-supported full-arch fixed dental prosthesis.

Supporting literature (Montero et al., 2021; Tomasi et al., 2021)

Quality of evidence Low

Grade of recommendation Grade B— \uparrow

Strength of consensus Consensus (1.6% of the group abstained due to potential Col)

Background

Intervention

Tooth-supported full-arch fixed prostheses were used to restore function and aesthetics. The treatment strategy commonly included the use of a fixed interim restoration during the early steps of periodontal therapy.

Available evidence

Number and design of included studies

Seven studies ($n = 522$ patients), with a weighted mean follow-up period of 8.6 years, were included. All studies were of observational and retrospective design and outcome measures included tooth loss, loss of restoration and occurrence of technical complications. Biological complications, PROMs, health-economic parameters and adverse events were not consistently reported.

Risk of bias

All included studies were judged to have a high risk of bias, mainly due to confounding and outcome assessment.

Effect sizes and their clinical relevance

Based on three studies ($n = 165$ patients), tooth loss over a period of 12.7 years was estimated at 4.9% (95% CI [2; 14]). Based on four studies ($n = 415$ patients), loss of restoration over an observation period of 9.7 years was estimated at 4.6% (95% CI [2; 14]). During an observation period of 7.2 years in three studies ($n = 365$ patients), the overall occurrence of technical complications at the restoration level was 8.0% (95% CI [6; 11]).

Consistency

Results were consistent across studies. The evidence considered has been generated by a small group of investigators and external applicability is unclear.

Balance of benefit and harm

High survival rates and low incidence of complications were observed. PROMs were not considered in the available studies. A tooth-supported fixed restoration may not address some functional and/or aesthetic needs in specific patients (e.g., when in need for facial tissue and lip support, or in cases of soft tissue deficiencies or long abutment teeth).

Overall certainty of the evidence

The certainty of evidence is graded as low based on the observational, retrospective design of studies and the high risk of bias.

From evidence to recommendation—additional considerations

Acceptability

PROMs were not reported in the identified studies (Tomasi et al., 2021). Data summarized by Montero et al. (2021) on tooth-supported fixed partial and full-arch prostheses suggest a high rate of patient satisfaction.

Feasibility

Related procedures are clinically and technically demanding.

Ethical considerations

Not applicable.

Economic considerations

Health-economic parameters were not evaluated in the identified studies. Different restorative materials and technical approaches may increase affordability. Due to the complexity of execution and cost of tooth-supported full-arch fixed prostheses, some patients may elect to proceed with palliative care consisting of a transitional denture retained by few residual teeth or a full denture.

Legal considerations

Not applicable.

R10.2. In patients with a compromised dentition due to stage IV periodontitis, what is the performance of tooth-supported full-arch removable prostheses?

PICOS question addressed by a SR

R10.2: Evidence-based recommendation

In periodontitis stage IV patients with an insufficient number/distribution of periodontally maintainable teeth to support a tooth-supported full-arch fixed dental prosthesis, a tooth-supported full-arch removable dental prosthesis (overdenture) may be considered.

Supporting literature (Donos et al., 2021)

Quality of evidence Very low

Grade of recommendation Grade O↔

Strength of consensus Strong consensus (0% of the group abstained due to potential Col)

Background

Intervention

In patients with an insufficient number and/or distribution of periodontally maintainable teeth to support a full-arch fixed prosthesis in the maxilla/mandible, clinicians may consider a tooth-supported removable full-arch prosthesis (TSRP). TSRP can be performed through different retention/attachment systems (e.g., ball-cap, magnetic) and the abutment teeth may or may not be splinted.

Available evidence

Number and design of included studies

Twenty-two studies (10 prospective and 12 retrospective studies) were included reporting on 4579 abutment teeth and 1660 TSRPs. The prospective studies had a weighted mean follow-up of 36.9 months for tooth survival and 86.5 months for prosthesis survival, while the retrospective studies had a weighted mean follow-up of 83.3 months for tooth survival and 75.2 months for prosthesis survival. Outcome measures included tooth loss and loss of restoration. Occurrence of technical and biological complications as well as PROMs and health-economic parameters and adverse events were not consistently reported.

Risk of bias

All included studies were judged to present with a high risk of bias.

Effect sizes and their clinical relevance

The prospective studies suggested a tooth survival rate (tooth level) ranging from 86% to 100% over a weighted mean follow-up of 36.9 months. The retrospective studies showed a range of survival from 34% to 94% at tooth level (weighted mean follow-up

83.3 months) and from 38% to 100% at prosthesis level (weighted mean follow-up 75.2 months).

Consistency

Treatment strategies varied across studies and sensitivity analyses were not feasible. The evidence considered has been generated by a small group of investigators and external validity is unclear.

Balance of benefit and harm

The evidence suggests heterogeneous outcomes in terms of prosthesis and tooth survival. The lack of information on complications should be considered.

Overall certainty of the evidence

The certainty of evidence is graded as very low based on study design, heterogeneous outcomes and high risk of bias.

From evidence to recommendation—additional considerations

Acceptability

PROMs were reported in nine of the included studies. The tools used to assess PROMs were not validated and inconsistent across studies.

Feasibility

Related procedures are clinically and technically demanding.

Ethical considerations

The preservation of abutment teeth might offer a psychological benefit for patients who are transitioning to edentulism.

Economic considerations

Health-economic parameters were not evaluated in the identified studies. TSRPs entail costs related to periodontal and prosthetic treatment.

Legal considerations

Not applicable.

R10.3. In patients with a compromised dentition due to stage IV periodontitis in whom tooth preservation was deemed impossible, what is the performance of implant-supported full-arch fixed prostheses?

PICOS question addressed by a SR

R10.3: Evidence-based recommendation

In periodontitis stage IV patients in whom tooth preservation was deemed impossible, and a sufficient number (≥ 4) of bilaterally distributed and adequately sized dental implants are planned in the maxilla and/or mandible, we suggest an implant-supported full-arch fixed dental prosthesis.

(Continues)

(Continued)

PICOS question addressed by a SR**R10.3: Evidence-based recommendation****Supporting literature** (Ramanauskaite et al., 2021; Tomasi et al., 2021)**Quality of evidence** Low**Grade of recommendation** Grade B-†**Strength of consensus** Consensus (33.3% of the group abstained due to potential Col)**Background****Intervention**

Implant-supported full-arch fixed prostheses were used to restore function and aesthetics on ≥ 4 implants in the same jaw. Prosthetic restorations were predominantly screw-retained and installation protocols, that is, immediate versus delayed implant installation, varied across studies.

Available evidence*Number and design of included studies*

Nineteen studies ($n = 1189$ patients) with a weighted mean follow-up period of 3.6 years were included. The majority of studies were observational and designed as prospective case series. Consistently reported outcome measures included loss of implants, loss of restorations and occurrence of technical and biological complications. Data on PROMs and health-economic parameters were not consistently reported.

Risk of bias

All included studies were judged to present with a high risk of bias, mainly due to confounding and outcome assessment.

Effect sizes and their clinical relevance

Based on 15 studies ($n = 670$ patients), implant loss over an observation period of 3.9 years was estimated at 3.5% (95% CI [2; 7]). Loss of restorations, as reported in nine studies ($n = 766$ patients), was estimated to be 4.6% (95% CI [1; 18]) during an observation period of 3.2 years. Over a follow-up period of 2.6 years (9 studies, 723 patients), technical complications affected 41.7% (95% CI [25; 68]) of all restorations. Biological complications were evaluated in 12 studies ($n = 984$ patients), covering a time period of 3.1 years and 8.5% (95% CI [5; 13]) of all implants developed at least one biological complication.

Consistency

Results were consistent across studies. The evidence considered has been generated by a small group of investigators and external applicability is unclear.

Balance of benefit and harm

High survival rates of restorations and implants were observed. The high rate of technical complications over short observation periods should be considered, as should be the limited information on PROMs. An implant-supported fixed restoration may not address some functional and/or aesthetic needs in specific patients (e.g., when in need for facial tissue and lip support, or in cases of soft tissue deficiencies).

Overall certainty of the evidence

The certainty of evidence is graded as low based on study design and limited follow-up periods.

From evidence to recommendation—additional considerations*Acceptability*

PROMs were reported in two studies ($n = 22$ patients), indicating a high level of patient satisfaction.

Feasibility

Related procedures are demanding in terms of professional competence and resources.

Ethical considerations

Not applicable.

Economic considerations

Health-economic parameters were evaluated in one ($n = 56$ patients) of the identified studies.

Legal considerations

Not applicable.

R10.4. In patients with a compromised dentition due to stage IV periodontitis in whom tooth preservation was deemed impossible, what is the performance of implant-supported full-arch removable prostheses?**PICOS question addressed by a SR****R10.4: Evidence-based recommendation**

In periodontitis stage IV patients in whom tooth preservation was deemed impossible and adequately sized dental implants can be used, albeit not in sufficient number and/or adequate position to support a full-arch fixed dental prosthesis, an implant-supported full-arch removable dental prosthesis (overdenture) may be considered.

Supporting literature (Donos et al., 2021; Ramanauskaite et al., 2021)**Quality of evidence** Very low**Grade of recommendation** Grade O-↔**Strength of consensus** Strong consensus (37.0% of the group abstained due to potential Col)

Background

Intervention

In edentulous patients in whom the insertion of a sufficient number (≥ 4) of bilaterally distributed and adequately sized (in terms of length and width, i.e., implant sizes providing adequate bone to implant contact to support the functional load) dental implants is not feasible, clinicians may consider an implant-supported removable prosthesis (ISRP). ISRPs can be supported/retained by a number of different attachment systems. The most commonly used systems include stud, bar, magnetic and telescopic attachments. Implants may or may not be splinted.

Available evidence

Number and design of included studies

Five studies ($n = 136$ patients) with a weighted mean follow-up period of 3.3 years were included. Four studies were of prospective and one of retrospective design. All studies were observational and used conventional loading protocols. The consistently reported outcome measures included survival of ISRPs and implants. Data on technical and biological complications, PROMs and health-economic parameters were not (consistently) reported.

Risk of bias

All studies had a high or critical risk of bias, thus raising concerns on the possibility of drawing robust conclusions.

Effect sizes and their clinical relevance

Prosthesis survival during the weighted mean follow-up period of 3.3 years was 100% and implant survival ranged from 96% to 100%.

Consistency

Due to the limited number of studies, sensitivity analyses were not feasible. Treatment strategies varied considerably across studies.

Balance of benefit and harm

The evidence suggests favourable outcomes in terms of prostheses and implant survival. The lack of information on complications and PROMs as well as the short follow-up should be considered.

Overall certainty of the evidence

The certainty of evidence is graded as very low based on study design and high risk of bias.

From evidence to recommendation—additional considerations

Acceptability

PROMs were reported in two of the included studies. The tools used to assess PROMs were not validated and inconsistent across studies.

Feasibility

ISRP-related procedures are clinically and technically demanding.

Ethical considerations

Not applicable.

Economic considerations

Health-economic parameters were not evaluated in the identified studies. ISRPs entail costs related to surgical and prosthetic treatment.

Legal considerations

Not applicable.

11 | LONG-TERM OUTCOMES OF TREATMENT IN STAGE IV PERIODONTITIS PATIENTS

11.1 | Intervention: professionally administered supportive periodontal care

The systematic review (Leow et al., 2021) assessed the role of professionally administered supportive periodontal care (SPC) in preventing tooth loss, periodontitis progression (clinical attachment loss, CAL) and associated health outcomes in patients with periodontitis who had completed active periodontal treatment.

The role of SPC is highly relevant to patients with all stages of periodontitis. In the systematic review (Leow et al., 2021), it was not possible to identify studies focussed on stage IV periodontitis and the findings are therefore relevant to all stages of disease.

Five components or combinations thereof contribute to SPC interventions:

1. **Interview:** periodontal health symptoms, medical and social history, risk factors including tobacco use, stress, diabetes and reported plaque control regime;
2. **Assessment:** plaque and calculus deposits, periodontal health status, including inflammation, PPDs and bleeding pockets;
3. **Evaluation:** intervention needs including risk factor management, oral hygiene and re-treatment;
4. **Practical Intervention:** oral hygiene coaching, instrumentation of supra- and sub-gingival plaque and calculus, treatment of sites with recurrence (finding of periodontitis at a previously healthy/stable site) or residual periodontitis (a deep periodontal pocket remains despite active therapy);
5. **Planning:** interval before next SPC visit. The control was no or irregular SPC, defined as greater than a frequency of 3-monthly.

Specialist and non-specialist settings were considered, and protocols included adjunctive therapies.

The primary outcome was tooth loss. Secondary outcomes included one site with CAL loss ≥ 2 mm, number of sites with PPDs

≥5 mm with BOP; number of sites requiring re-treatment; OHRQoL questionnaires; care cost analysis; other patient-reported outcomes.

Studies of 5–10 years follow-up ($n = 17$), over 10-years ($n = 7$) and 20 years ($n = 2$) were included. Nine studies addressed 12-month follow-up of treatment of recurrence, but high heterogeneity prevented conclusions on different treatment methods.

11.2 | Is regular, professionally administered, SPC effective in preventing tooth loss or disease recurrence in the longer term?

R11.1. Does regular SPC reduce tooth loss?

PICOS question addressed by a SR

R11.1: Evidence-based recommendation (1), Expert consensus-based recommendation (2)

1. **We recommend** provision of and adherence to regular professionally administered supportive periodontal care (SPC) to reduce tooth loss in the long term (≥5 years).
2. **We recommend** that SPC should initially be provided at 3-monthly intervals. Medium to long-term frequency should be personalized to each individual patient, taking into account clinical and behavioural circumstances.

Supporting literature (Leow et al., 2021)

Quality of evidence Moderate

For tooth loss: Seventeen prospective cohorts judged as “low” risk of bias except one study with a “moderate” risk using the Newcastle–Ottawa scale.

For CAL loss ≥2 mm: Seven prospective cohorts whereby all were judged as “low” risk of bias.

Grade of recommendation Grade A–↑↑ (1); Grade A–↑↑ (2)

Strength of consensus Consensus (0% of the group abstained due to potential Col)

Background

Intervention

SPC includes preventative and/or therapeutic interventions for periodontitis patients who have been successfully treated for periodontitis. The goals of SPC are to maintain periodontal stability, by preventing disease recurrence or progression and ultimately to prevent tooth loss. Adherence to professionally administered SPC at regular time intervals allows ongoing monitoring of periodontal status, practical interventions (as required) and formulation of individually tailored SPC time intervals based on patient risk.

Available evidence

Number and design of included studies

Twenty-four prospective cohort studies. The primary outcome of the systematic review (Leow et al., 2021) was tooth loss. The secondary outcome to inform on disease recurrence/progression was CAL loss ≥2 mm.

Tooth loss: 17 studies reported tooth loss and eight of these studies were included in the meta-analysis, contributing to tooth loss at a patient level (192 participants). The remaining nine studies were reported qualitatively.

Clinical attachment level loss ≥2 mm: seven studies reported on CAL loss ≥2 mm. Three studies contributed data to the meta-analysis for estimating the number of patients experiencing CAL loss ≥2 mm (86 participants), while the remaining four studies were reported qualitatively.

Risk of bias

Study quality assessment using the Newcastle–Ottawa scale, identified low risk of bias for all but two studies, which had a moderate risk of bias.

Effect sizes and their clinical relevance

“Regular” SPC was defined as 3-monthly, while a lack of adherence to or no SPC was described as “irregular”.

Tooth loss: the mean proportion of patients who experienced tooth loss was 9.6% (95% CI [5; 14]) with low heterogeneity $I^2 = 28%$ ($p = .161$); 8% (95% CI [2; 14]) of regular attendees experienced tooth loss compared with 11.9% (95% CI [5; 19]) of irregular attendees. The difference between sub-groups was, however, not statistically significant ($p = .161$). A greater length of follow-up (≥10 years) was associated with an increased experience of tooth loss of 12.7% (95% CI [4; 22]) compared with 8.2% (95% CI [3; 13]) for 5–10 years follow-up.

Clinical attachment level loss ≥2 mm: the mean proportion of patients who experienced overall CAL loss ≥2 mm was 24.8% (95% CI [11; 38]) with substantial heterogeneity $I^2 = 63%$ ($p = .013$); 30.2% (95% CI [–2; –63]) of patients who attended regular SPC appointments experienced CAL loss ≥2 mm, compared with 21.4% (95% CI [10; 33]) of those who attended irregularly. The difference between subgroups was not statistically significant ($p = .332$). The unexpected result indicating that regular SPC may lead to a greater experience of CAL loss ≥2 mm is imprecise (large confidence interval). A greater length of follow-up (≥10 years) was associated with a slight increase in CAL loss ≥2 mm of 26.3% (95% CI [8; 45]) compared with 22.1% (95% CI [5; 39]) for 5–10 years follow-up.

Consistency

Heterogeneity across studies was identified for both tooth loss [$I^2 = 28%$ ($p = .161$)] and CAL loss ≥2 mm [$I^2 = 63%$ ($p = .013$)]. This may be explained by the limited number of studies fulfilling the inclusion criteria of the systematic review and/or the type of therapy carried out in the active periodontal treatment phase (regenerative or non-regenerative). Studies differed in terms of a large variety of factors/steps related to an SPC appointment and frequently did not report the operator(s) performing each visit.

Balance of benefit and harm

Few studies reported adverse events. An overall consideration of the benefit versus harm of regular SPC supports the strength of the recommendation.

Overall certainty of the evidence

Moderate.

From evidence to recommendation—additional considerations

Acceptability

Little is known about patient preferences in relation to SPC. However, SPC has been recommended for oral healthcare for several decades.

Feasibility

Little is known about implementation. SPC is a routinely provided intervention in a number of healthcare systems, although the regularity of visits (3–4 times per year) may be a barrier for some patients (financial and logistical).

Ethical considerations

No evaluation of equity or access to SPC has been conducted. However, utilization of dental services is unequally distributed, and therefore, it is reasonable to assume the same would be the case for SPC. Identifying barriers to and facilitators of SPC, and using this information to enhance access would seem to be a priority.

Economic considerations

Limited evidence exists on the cost-effectiveness of SPC. SPC delivered in specialist care when compared with general dental practice may result in less CAL loss and higher tooth survival rates, but at a greater financial cost (Gaunt et al., 2008). Economic modelling indicates SPC is cost-effective in developed economies (Pennington et al., 2011).

Legal considerations

Not applicable.

R11.2. Does residual probing pocket depth, following active periodontal treatment, affect disease recurrence during SPC?

PICOS question addressed by a SR

R11.2: Evidence-based statement

In patients with periodontitis, the presence of residual probing pocket depths (≥ 5 mm) following active therapy increases the risk of disease recurrence/progression, even though the patient is enrolled in SPC.

Supporting literature (Leow et al., 2021)

Quality of evidence Moderate

Two studies, which were a prospective cohort and a controlled clinical trial, with risk of bias designated as “low” and “serious”, respectively.

Grade of recommendation Statement: unclear, additional research needed

Strength of consensus Unanimous consensus (0% of the group abstained due to potential CoI)

Background

Intervention

The goals of SPC are to maintain periodontal stability (BOP in $<10\%$ of sites; shallow PPDs of 4 mm or less, and no 4 mm sites with BOP) (Chapple et al., 2018), by preventing disease recurrence or progression and ultimately to prevent tooth loss. Ideally, patients enter SPC with periodontal stability; however, in some situations, the strict definition in the 2018 Classification of stability/health following periodontal treatment is not always achieved. Residual periodontal probing depths (PPDs) ≥ 4 mm with BOP are likely to be unstable and pose a risk for disease progression.

Available evidence

Number and design of included studies

Two studies addressing disease recurrence (CAL loss ≥ 2 mm) during SPC were included, one controlled clinical trial and one prospective cohort.

Risk of bias

Study quality assessment using the Robins-I tool showed a “serious” risk of bias and “low” risk using the Newcastle–Ottawa scale.

Effect sizes and their clinical relevance

Based on one controlled clinical trial (Jenkins et al., 2000), patients with residual PPD ≥ 6 mm after active periodontal therapy demonstrated disease recurrence (CAL loss ≥ 2 mm) in the range of 20.5%–28.6% of sites over 12 months follow-up. In contrast, shallower residual PPD (4.0–5.9 mm) showed disease recurrence in the range of 11.6%–11.8%. No statistically significant difference between groups was found with regard to the modality of treatment during SPC (coronal scaling vs. subgingival instrumentation).

The prospective cohort study (Cortellini et al., 2017) reported the highest rate of disease recurrence at sites PPD ≥ 5 mm (compared with PPD <5 mm) over a 20-year period. Regression analysis showed that residual PPD significantly correlated with disease recurrence ($p = .0024$, $R^2 = 0.31$, root mean square error = 0.75).

Consistency

Studies were heterogeneous in design and reporting of outcomes. All studies describe increased disease recurrence with increasing residual PPD after active periodontal therapy.

Balance of benefit and harm

One study reported adverse events following subgingival instrumentation of residual pockets. These were, however, not deemed to be serious. An overall consideration of the benefit versus harm of achieving periodontal stability (no residual deepened PPDs), following active periodontal therapy, in order to reduce the risk of disease recurrence supports the strength of the statement.

Overall certainty of the evidence

Moderate.

From evidence to recommendation—additional considerations

Acceptability

Little is known about patient preferences in relation to periodontal stability at the completion of active periodontal therapy. Dental professionals have a strong preference for PPD \leq 4 mm without BOP after completion of periodontal therapy.

Feasibility

Little is known about implementation. Direct and indirect costs of reaching periodontal stability at completion of SPC may be a barrier for some patients.

Ethical and legal considerations

Not applicable.

Economic considerations

Limited evidence exists on direct/indirect costs of the treatment of disease recurrence during SPC. Clearly, disparities exist between countries, healthcare systems and the modality of treatment chosen to treat the recurrence.

R11.3. Should recall intervals for SPC be guided by patients' risk status?

Additional question raised by the WG

R11.3: Expert consensus-based recommendation

We **recommend** that recall intervals for supportive periodontal care (SPC) should be guided by patients' risk profile as determined by individual risk factors (e.g., smoking, hyperglycaemia) and disease-associated clinical measures (such as pocket depths and bleeding on probing).

Supporting literature (Rosling et al., 2001; Matuliene et al., 2008; Lang et al., 2015; Trombelli et al., 2015; Trombelli et al., 2017; Trombelli et al., 2020) and Expert opinion.

Quality of evidence Not applicable

Grade of recommendation Grade A- $\uparrow\uparrow$

Strength of consensus Unanimous consensus (0% of the group abstained due to potential CoI)

Background

Intervention

A patient's previous experience of periodontitis is a strong predictor of future disease activity in the absence or presence of periodontal

treatment (Machtei et al., 1993; Martin et al., 2009). Regular SPC is associated with reductions in tooth loss relative to irregular SPC (Saminsky et al., 2015). Risk factor control is an important element of the first step of therapy in the treatment of periodontitis (Ramseier et al., 2020; Sanz, Herrera, et al., 2020). It is important in an era of personalized dental medicine to determine whether frequency of SPC should be tailored according to a patient's risk profile. Patient risk profile can be estimated through different validated patient risk assessment tools, such as the Periodontal Risk Assessment tool (PRA) (Lang & Tonetti, 2003), the Periodontal Risk Calculator (PRC) and DenPlan/PreViser Patient Assessment tool (Page et al., 2003), Peri-oRisk (Trombelli et al., 2017). Patient risk profile can also be expressed through the "Grade" system (Tonetti et al., 2018).

Available evidence

Number and design of included studies

In the systematic review (Leow et al., 2021), none of the studies that were included addressed risk factor control in SPC. No detail of factors that influenced the frequency of the recall interval was given. One 12-year prospective cohort study (Rosling et al., 2001) evaluated disease progression in normal and high susceptibility patients during 12 years of SPC, where SPC frequency was determined by risk/susceptibility categorization at the end of active periodontal therapy.

Risk of bias

Not applicable.

Effect sizes and their clinical relevance

No included studies documented details on how risk factor control might have influenced recall intervals in SPC. Only one study (Rosling et al., 2001) included in the systematic review (Leow et al., 2021) investigated risk profiles in association with tooth loss, and suggested that tailoring frequency of SPC to risk profiles may prevent tooth loss in cohorts with different susceptibilities to periodontitis. Substantial evidence for the validity of different risk assessment tools based on individual risk factors (e.g., smoking, hyperglycaemia) and disease-associated clinical markers (such as pocket depths and BOP) to predict tooth loss has been presented (Lang & Tonetti, 2003; Page et al., 2003; Matuliene et al., 2008; Martin et al., 2009; Martin et al., 2010; Trombelli et al., 2017; Ramseier et al., 2019). Such tools may be used to inform the frequency of SPC recalls (Lang et al., 2015; Trombelli et al., 2020). Initial evidence suggests that "Grading" (Tonetti et al., 2018) may be able to predict tooth loss (Ravida et al., 2020; Al-Harathi et al., 2021; Saleh et al., 2021) and may be used to set SPC frequency recalls, although more evidence is needed.

Consistency

Not applicable.

Balance of benefit and harm

Evidence suggests that the frequency of recall intervals for SPC may be set according to a patient's risk status, with high-risk individuals benefiting from 3-monthly SPC and lower-risk patients remaining largely stable with a frequency of 6–12 months. An overall consideration of the benefit versus harm of risk factor control to influence recall intervals of SPC supports the strength of the statement.

Overall certainty of the evidence

Not applicable.

From evidence to recommendation—additional considerations

Acceptability

Little is known about patients' preferences in relation to frequency of recall visits. Oral healthcare professionals are increasingly collaborating with other healthcare providers and health authorities to raise awareness of and manage risk factors for the treatment of periodontitis.

Feasibility

Little is known about the implementation of risk-driven recall intervals for SPC; however, experience of oral healthcare professionals suggests that 3-monthly SPC intervals are feasible and acceptable to patients at high risk of recurrence of periodontitis and associated tooth loss.

Ethical considerations

It would seem appropriate to recommend more regular (3–4 monthly) SPC visits for those with a higher grade/risk of periodontitis than those with a lower grade/risk.

Economic considerations

Some barriers may exist for patients at high risk due to the need for more frequent recall visits and the associated costs. This is likely to increase health inequalities in countries with no or limited public healthcare funding for periodontitis.

Legal considerations

Not applicable.

R11.4. What are the important components to consider when designing a successful SPC programme?

Additional question raised by the WG

R11.4: Expert consensus-based recommendation

We **recommend** a number of important components when designing an SPC programme, including:

- a. Specific interventions include interview, assessment, evaluation, practical intervention and planning (see introduction).

(Continues)

(Continued)

Additional question raised by the WG

R11.4: Expert consensus-based recommendation

- b. Delivery by a variety of oral healthcare professionals, under the supervision of a suitably trained general dentist or a specialist, as appropriate to case complexity.
- c. Clear two-way communication between the oral healthcare team and the patient, and between healthcare professionals (medical or dental).

Supporting literature Expert Opinion.

Quality of evidence Not applicable

Grade of recommendation Grade A–↑↑

Strength of consensus Unanimous consensus (0% of the group abstained due to potential Col)

Background

Intervention

SPC is the fourth step of therapy (Sanz, Herrera, et al., 2020). It is a complex intervention that elicits key questions relevant to both oral healthcare providers and patients and is crucial to the long-term stability of the periodontium. Important aspects of SPC are documented below:

- (a) Specific interventions to include in an SPC programme.

In designing an SPC programme, it is important to regularly consider a number of components which include:

- **Interview:** elicit information on periodontal health symptoms, medical and social history, risk factors including tobacco use, stress, diabetes and reported plaque control regime, patient motivation towards continuous risk factor control and PMPR/subgingival instrumentation;
- **Assessment:** plaque and calculus deposits, periodontal health including inflammation, PPDs, and bleeding pockets;
- **Evaluation:** of intervention needs, including risk factor management, oral hygiene and re-treatment;
- **Communicating:** findings to patients to enhance their ownership of periodontal health and agreement on required interventions
- **Practical Intervention:** oral hygiene coaching, instrumentation of supra- and sub-gingival plaque and calculus, treatment of sites with recurrence or residual periodontitis.
- **Planning:** interval before next SPC visit.

(b) The specific oral healthcare professional that undertakes/oversees the SPC programme.

A variety of qualified and trained oral healthcare professionals can carry out the components of an SPC programme. Nine prospective cohort studies included in the systematic review (Leow et al., 2021) utilized undergraduate dental students, dental hygienists and periodontists.

The SPC programme should be overseen by a suitably qualified and trained general dentist or specialist. Effective

communication is essential should the operator carrying out the components of the SPC programme differ from the person overseeing the SPC programme.

(c) Key communication steps to undertake for long-term patient benefit?

- Oral healthcare professionals and the patient: communication between oral healthcare professionals delivering the SPC programme and the patient is crucial at each visit. The patient should have a clear understanding of his/her periodontal status, treatment needs (if any) and the recommended home-care regime.
- Communication between healthcare professionals: the person overseeing the SPC programme may be different to that delivering the components of an SPC programme. Clear and transparent communication between these oral healthcare professionals should occur at each recall visit. Importantly, a number of healthcare professionals may be involved with a patient's care, particularly with regard to medical health and/or risk factor management. In order to work synergistically for long-term patient benefit, the person overseeing the SPC programme should initiate and maintain communications.
- Oral healthcare professional overseeing the SPC programme and the patient: the person overseeing the SPC programme should communicate with the patient on a regular basis. Patients should be actively engaged in the decision-making process regarding long-term care.

Available evidence

Number and design of included studies

None of the studies included addressed specific components that should be included in an SPC programme, who should undertake or oversee the SPC programme nor which key communications steps should be undertaken for long-term patient benefit.

Risk of bias, effect sizes and their clinical relevance, consistency, balance of benefit and harm

Not applicable.

Overall certainty of the evidence

Not applicable.

From evidence to recommendation—additional considerations

Acceptability

Little is known about patient preferences with regard to design and delivery of SPC programmes. Clear and transparent communication between healthcare professionals and with the patient is essential for long-term patient benefit.

Feasibility, ethical, economic, legal considerations

Not applicable.

R11.5. What is the best approach, when treating the recurrence of periodontitis during SPC to reduce tooth loss and/or influence other outcomes (e.g., measures of periodontal health, quality of life, cost and accessibility of care and harms)?

PICOS question addressed by a SR

R11.5: Evidence-based recommendation

We **suggest not** to use adjunctive approaches to subgingival instrumentation when treating recurrence of periodontitis during supportive periodontal care.

Supporting literature (Leow et al., 2021)

Quality of evidence Moderate

Nine studies (seven randomized controlled trials, one controlled clinical trial and one prospective cohort) with ≥ 12 months follow-up. Risk of bias—6 “of some concern”, 1 “high”, 2 “serious”.

Grade of recommendation Grade B—↓

Strength of consensus Consensus (0% of the group abstained due to potential Col)

Background

Intervention

Periodontitis may become unstable during long-term SPC and may present as disease recurrence, occurrence or progression.

- Recurrence refers to a finding of periodontitis at a site that was rendered periodontally healthy/stable through active treatment;
- Occurrence refers to a diseased site arising within a periodontitis patient that did not previously exhibit signs of disease;
- Progression is characterized by deterioration (e.g., clinical attachment loss) at a site that exhibited residual disease despite active treatment.

It may be challenging to distinguish between these, particularly if the SPC programme has not been continuously delivered in a single dental practice/office setting. Sites with PPDs ≥ 4 mm and BOP require further treatment in order to reduce the risk of further deterioration and/or tooth loss.

Available evidence

Number and design of included studies

Nine studies, with ≥ 12 months follow-up, were included. Seven studies were RCTs, one study was a controlled clinical trial and one study was a prospective cohort.

Tooth loss: two studies (one RCT and one prospective cohort) reported on tooth loss. These studies were reported qualitatively.

CAL loss ≥ 2 mm: two studies reported on CAL loss ≥ 2 mm (one RCT and one controlled clinical trial) and were included in the qualitative analysis.

Pockets of ≥ 5 mm with BOP: no studies reported on the number of PPD ≥ 5 mm with bleeding on probing during SPC.

Sites that need/experience re-treatment: one study (split-mouth RCT) reported on breakdown sites that required re-treatment.

PROMs: one RCT reported on oral health-related quality of life utilizing the Italian translation of the Oral Health Impact Profile (OHIP) questionnaire –14.

Health-economic outcomes: two RCTs provided information on total cumulative costs for operative interventions.

Other PROMs: a number of studies reported on other outcomes/adverse events. These included periodontal abscess (one RCT), masticatory function and aesthetics (one RCT) and adverse events (one controlled clinical trial and an RCT).

Risk of bias

Nine studies were included. Study quality assessment using the Cochrane Risk of Bias tool 2.0 showed six studies were of, “some concern”, and one study was classified as, “high”. Two studies were judged as, “serious” using the Robins-I tool.

Effect sizes and their clinical relevance

Tooth loss: one RCT (Bogren et al., 2008) on 128 patients with 3-year outcomes, compared subgingival instrumentation (SRP) alone (control) with SRP and locally delivered 8.8% doxycycline gel applications (test) in PPD ≥ 5 mm. SPC was conducted every 6 months. Twenty-five test sites were lost due to tooth extractions (3.6% of initially included sites) in the test group, while 45 sites were lost due to extractions (4.9% of those initially included) in the control group. The difference between groups was not statistically significant ($p > .05$). A prospective cohort (Costa et al., 2015), which included 212 patients, assessed surgical therapy versus non-surgical therapy in both compliant and non-compliant patients, over 5 years of follow-up. While mean tooth loss for compliant patients was significantly less (0.3 for non-surgical and 0.8 for surgical therapy) than for non-compliant patients (2.2 and 2.8, respectively), no statistically significant differences were noted overall between ST and NST.

CAL loss ≥ 2 mm: one controlled clinical trial (Jenkins et al., 2000) compared coronal scaling in 17 patients (146 sites) with subgingival scaling in 14 patients (130 sites) over 12 months, and 21 sites exhibited CAL loss ≥ 2 mm from each group over the course of SPC, with no statistically significant differences between groups. A 12-month multi-centre RCT (Tonetti et al., 2012) on 202 subjects compared SRP (control) with SRP and 14% doxycycline gel application (test) on PPD ≥ 5 mm with BOP: 8 test and 7 control patients experienced CAL loss ≥ 2 mm with no statistically significant difference between the groups.

Sites that need/experience re-treatment: one split-mouth RCT (Kaldahl et al., 1996b) reported 685 “breakdown” sites during the course of SPC which required re-treatment. About 5%–12% of these breakdown sites experienced CAL loss ≥ 3 mm and were re-treated and subsequently experienced further loss of attachment.

PROMs: the Italian translation of the OHIP-14 questionnaire was used in one RCT (Cortellini et al., 2020) at baseline, 1, 5 and 10 years after regenerative treatment (other treatment groups were not relevant to this review). The mean OHIP-14 score 1 year after the regenerative procedure was 6.6 (SD = 2.4), however no data were presented at 10 years.

Health-economic outcomes: one RCT (Cortellini et al., 2017) reported cumulative costs over a 20-year period (with 3-monthly SPC) using an average of fees from private practices in Italy. The cumulative costs ranged from a mean of €3090.98 (± 210.66) to €3382 (± 88.95) and were dependent on the treatment carried out during the active phase of therapy. Another study (Cortellini et al., 2020) reported cumulative costs for a regenerative procedure over 10 years (excluding SPC).

Other PROMs: 27 periodontal abscesses were reported in one RCT (Kaldahl et al., 1996a) with 84 months of follow-up, with most (85%) occurring in the group originally treated by coronal scaling alone and a large proportion (63%) occurring in sites with PPD ≥ 7 mm at the initial examination. Cortellini et al. (2020) assessed masticatory function and aesthetics using a 5-point Likert scale with a 10-year follow-up, and the proportion of people reporting “some concern” for both masticatory function and aesthetics appeared to increase over the 9 years of SPC (graphical information available only).

Adverse events were largely not described, but when judged in two studies, it was mainly judged as “not serious” (Jenkins et al., 2000; Tonetti et al., 2012). Both studies reported no difference between study groups in regard to adverse events.

Consistency

Not applicable.

Balance of benefit and harm

The best approach to treatment of recurrence of periodontitis in SPC is currently unknown. However, as demonstrated in those studies included, some clinical benefit can be achieved regardless of treatment modality. Additionally, no difference between treatment approaches could be determined with respect to adverse events.

Overall certainty of the evidence

Low.

From evidence to recommendation—additional considerations

Acceptability

Little is known about patient preferences in regard to treatment of recurrence in SPC. Determining the best approach to treatment of recurrence in conjunction with patients' views on these should undoubtedly be a research priority.

Feasibility, ethical, legal considerations

Not applicable.

Economic considerations

The costs and cost-effectiveness of re-treatment of recurrence are currently unknown.

11.3 | Is regular, professionally administered, SPC associated with positive and/or negative outcomes, aside from tooth loss and/or disease recurrence?

R11.6. Are there disadvantages to regular long-term SPC (e.g., more gingival recession/clinical attachment loss)?

PICOS question addressed by a SR

R11.6: Evidence-based statement

There is no evidence of clinical disadvantages to regular long-term SPC, such as gingival recession/clinical attachment loss; however, the possibility of these side effects cannot be excluded based on the evidence reviewed. Patients should be advised of this as part of their informed consent.

Supporting literature (Leow et al., 2021)

Quality of evidence Three prospective cohorts with a “low” risk of bias.

Grade of recommendation Grade O→ Statement: unclear, additional research needed

Strength of consensus Strong consensus (0% of the group abstained due to potential Col)

Background

Intervention

Periodontitis patients in long-term SPC should attend regular recall appointments in order to reduce their risk of tooth loss. At times, periodontitis may become unstable and require re-treatment, which may take the form of a non-surgical and/or surgical approach to resolve PPDs ≥ 4 mm with BOP. One common consequence of re-treatment may be CAL in the form of gingival recession, the magnitude of which is difficult to predict.

Available evidence

Number and design of included studies

Three prospective cohort studies contributed data to the meta-analysis for estimating the number of patients experiencing CAL loss ≥ 2 mm (86 participants) during SPC.

Risk of bias

All three studies were classified as having, “low” risk of bias using the Newcastle–Ottawa scale.

Effect sizes and their clinical relevance

The primary outcome of the systematic review (Leow et al., 2021) was tooth loss. The secondary outcome, to inform on disease recurrence/

progression, was CAL loss ≥ 2 mm. “Regular” SPC was defined as 3-monthly, while a lack of adherence to or no SPC was described as “irregular”. The mean proportion of patients who experienced overall CAL loss ≥ 2 mm was 24.8% (95% CI [11; 38]) with substantial heterogeneity $I^2 = 63%$ ($p = .013$). 30.2% (95% CI [-2; 63]) of patients who attended regular SPC appointments experienced CAL loss ≥ 2 mm, compared with 21.4% (95% CI [10; 33]) of those who attended irregularly. The difference between subgroups was not statistically significant ($p = .332$).

Although it appears that regular SPC does not lead to significantly greater experience of CAL loss ≥ 2 mm, this should be interpreted with caution. Only a small number of studies ($n = 3$) contributed to the meta-analysis and the resulting data for the regular SPC group appear imprecise (large confidence interval). Furthermore, the disparity may be explained by a single outlier where participants in that group presented with an increased number of residual PPD at the start of SPC.

A greater length of follow-up (≥ 10 years) was associated with a slight increase in CAL loss ≥ 2 mm of 26.3% (95% CI [8; 45]) compared with 22.1% (95% CI [5; 39]) for 5–10 years follow-up.

Consistency

Heterogeneity for studies included to inform on CAL loss ≥ 2 mm was substantial ($I^2 = 63%$; $p = .013$).

Balance of benefit and harm

The desirable effects of long-term regular SPC (reduced prevalence of tooth loss) would undoubtedly outweigh possible undesirable effects regarding CAL. An overall consideration of the benefit versus harm of regular SPC supports regular SPC.

Overall certainty of the evidence

Low.

From evidence to recommendation—additional considerations

Acceptability, feasibility, ethical, economic, legal considerations

See recommendation R11.1.

Legal considerations

Not applicable.

R11.7. Is long-term SPC cost-effective when considering direct and indirect costs?

Additional question raised by the WG

R11.7: Expert consensus-based statement

We suggest that regular long-term SPC in specialist practice may result in greater periodontal stability and tooth survival when compared with SPC in general practice.

(Continues)

(Continued)

Additional question raised by the WG**R11.7: Expert consensus-based statement**

We do not know if long-term SPC is cost-effective when considering direct and indirect costs.

Supporting literature (Gaunt et al., 2008) and Expert opinion

Quality of evidence Not applicable

Grade of recommendation Grade O—↔ Statement: unclear, additional research needed

Strength of consensus Unanimous consensus (0% of the group abstained due to potential Col)

Background**Intervention**

Costs incurred by SPC are complex and include both direct and indirect costs. Direct costs include elements such as the recall visits themselves and/or costs of re-treatment, while the consequences of periodontitis (e.g., cost of oral rehabilitation following tooth extraction or root caries) and time commitments from the patient (i.e., absenteeism from work) are considered indirect costs. There are also “intangible” costs such as impact on quality of life and number of healthy life years.

Available evidence*Number and design of included studies*

No studies reported on direct, indirect and intangible costs in SPC.

Risk of bias

Not applicable.

Effect sizes and their clinical relevance

One systematic review (Gaunt et al., 2008) concluded that SPC delivered in a specialist environment led to greater periodontal stability (clinical attachment) and higher tooth survival rates when compared with general practice, but was more expensive to deliver (direct costs) in the specialist environment. The systematic review conducted an analysis on cost-effectiveness based on data from one study (Axelsson & Lindhe, 1981) being extrapolated over a 30-year period. The analysis was from the perspective of a single patient with the primary patient-based outcome of tooth loss and secondary outcome of CAL. The authors calculated “tooth years lost”, which considered time as a factor. Thus, one tooth lost after the first year of SPC, equated to 30 tooth years lost over the course of the evaluation. The comparison was taken for those patients who received SPC in specialist care (charges based on one specialist practice in North East England) versus in a general dental practice (charges based on state health service charges in Scotland). Importantly, this model assumed that only SPC would be undertaken (no periodontal re-

treatment) and highlights the uncertainty in the values used in the analysis. An incremental cost-effectiveness ratio (ICER) was created by considering the increase in cost of a particular programme (i.e., specialist or general dentist in either setting) by the increase in benefit (outcome). The results of the analysis showed that SPC when provided in specialist practice was more effective than that delivered by a general practitioner, in terms of tooth loss and clinical attachment. SPC in specialist practice, however, costs €4466 more than a general dentist in private practice, and €5938 more than a general dentist in a state health service system. The ICER for SPC delivered in specialist care (general dentist in private practice as baseline) was €217 for one extra tooth year or an extra €1130 for 1 mm less attachment loss. When using the general dentist in a state-supported health service as a baseline, specialist care equated to €288 for one extra tooth year or an extra €1503 per 1 mm loss of attachment. Undoubtedly, compliance with recall appointments and efficiency and appropriateness of care is crucial to our understanding of cost-effectiveness. No data on compliance were given for the cohort of patients who received SPC in general practice.

Consistency, balance of benefit and harm

Not applicable.

Overall certainty of the evidence

Very low.

From evidence to recommendation—additional considerations*Acceptability*

It is unknown whether patients prefer SPC appointments in the specialist environment or in general practice. Often, a shared responsibility exists (between specialist and general dental practices), which may reduce overall costs, possibly at the expense of tooth or CAL.

Feasibility

Not applicable.

Ethical considerations

Limited evidence exists that regular SPC is beneficial to the patient (reducing tooth loss and disease progression/recurrence). Although SPC delivered in specialist practice appears to increase tooth survival and reduce CAL when compared with general practice, this comes at a higher cost, which may be a barrier for some patients.

Economic considerations

Limited evidence of the cost-effectiveness of SPC regarding direct and indirect costs exists and should be a clear future research priority.

Legal considerations

Not applicable.

R11.8. Does long-term SPC impact upon patient-reported outcome measures (PROMs) (OHRQoL, masticatory function, aesthetics)?

PICO question addressed by a SR

R11.8: Evidence-based statement

We **do not know** if long-term SPC impacts upon patient-reported outcomes.

Supporting literature (Leow et al., 2021)

Quality of evidence Three prospective cohorts

Grade of recommendation Grade O→ Statement: unclear, additional research needed

Strength of consensus Unanimous consensus (0% of the group abstained due to potential Col)

Background

Intervention

The impact of a disease and its treatment on a patient's quality of life may be captured by PROMs. A number of tools have been utilized to assess PROMs, often in the form of questionnaires or scales. Periodontitis has been demonstrated to have a negative impact on a patient's OHRQoL, while non-surgical therapy and surgical therapy may improve this (Shanbhag et al., 2012), using validated PROMs.

Available evidence

Number and design of included studies

Three prospective cohorts reporting on PROMs were included.

Risk of bias

All three studies were judged as, "low" risk of bias using the Newcastle–Ottawa scale.

Effect sizes and their clinical relevance

Patient-reported outcomes: the Italian translation of the OHIP-14 questionnaire was used in one RCT (Cortellini et al., 2020) at baseline, 1, 5 and 10 years after regenerative treatment (other treatment groups were not relevant to this review). The mean OHIP-14 score 1 year after the regenerative procedure was 6.6 (±2.4). However, no data were presented at 10 years.

Other PROMs: 27 periodontal abscesses were reported in one RCT (Kaldahl et al., 1996a), with 84 months of follow-up, with most (85%) occurring in the group originally treated by coronal scaling alone and a large proportion (63%) occurring in sites with PPD ≥7 mm at the initial examination.

Cortellini et al. (2020) assessed masticatory function and aesthetics using a 5-point Likert scale with a 10-year follow-up. The proportion of people reporting "some concern" for both masticatory

function and aesthetics appears to increase over the 9 years of SPC (graphical information available only).

Consistency

Not applicable.

Balance of benefit and harm

Regular SPC benefits the patient to reduce the risk of tooth loss and CAL, while limited evidence exists on PROMs.

Overall certainty of the evidence

Low.

From evidence to recommendation—additional considerations

Acceptability, feasibility, ethical, economic, legal considerations

Not applicable.

12 | IMPACT OF PERIODONTAL TREATMENT ON SYSTEMIC HEALTH AND QUALITY OF LIFE

12.1 | Periodontal treatment in severe periodontitis (stages III or IV or equivalent): Impact on systemic health and quality of life

A systematic review (Orlandi et al., 2021) addressed the impact of periodontal interventions, including non-surgical (steps 1 and 2), surgical (step 3) and adjunctive therapies, on systemic health and quality of life in patients with severe (stages III or IV or equivalent) periodontitis who:

1. were systemically healthy by anamnesis (medical history);
2. exhibited one or more co-morbid systemic non-communicable diseases (NCDs);
3. were pregnant.

NCDs assessed were: cardiovascular diseases, arrhythmias, hypertension, rheumatic, neurological, respiratory, metabolic, kidney, liver and inflammatory gastrointestinal diseases, malignancy, osteoporosis and mental health conditions (G.B.D. Diseases and Injuries Collaborators, 2020).

The included intervention studies did not specifically provide treatment to patients to a defined endpoint/outcome of success; therefore, there may be an underestimation of the effect size upon the systemic outcomes defined by the PICOS. Furthermore, treatment of periodontitis is instigated for the purpose of improving oral health, and systemic health benefits in patients without a co-morbid NCD are therefore a secondary consideration because an absence

of any beneficial systemic effect would not contraindicate the periodontal therapy. While the most relevant analysis refers to the treatment of periodontitis in patients with co-morbid NCDs ($n = 29$ RCTs), data were also combined with that from systemically healthy individuals ($n = 3$ RCTs). This approach was not defined a priori and it represents a methodological limitation of the review, but it was deemed to be justified on the basis that there would likely be patients within the systemically healthy group who had undiagnosed NCDs.

Sixteen RCTs addressed adverse pregnancy outcomes (pre-term birth <37, <35, and <32 weeks, low birth weight <2500 g and <1500 g, pre-term low birth weight, pre-eclampsia, gestational age at delivery, C-reactive protein [CRP], stillbirth, birthweight, and perinatal loss).

Following an update of the systematic search, five new studies were identified (Buwembo et al., 2020; Montero et al., 2020; Nguyen et al., 2021; Qureshi et al., 2021; Rapone et al., 2021) with data from four of them included in meta-analyses (Montero et al., 2020; Nguyen et al., 2021; Qureshi et al., 2021; Rapone et al., 2021). The updated meta-analyses did not change the interpretation of the available evidence apart from the increase in diastolic blood pressure previously observed for PICOS #2, which became non-statistically significant (0.15 mmHg 95% CI [-0.14; 0.44]; $p = .311$). Furthermore, a statistically significant reduction in tumour necrosis factor (TNF)- α [-0.27 pg/ml 95% CI [-0.53; 0.014]; $p = .039$) was observed for PICOS #1 and PICOS #2 populations combined.

12.1.1 | Impact of periodontal treatment on systemic inflammation and cardio-metabolic risk in people with no reported systemic co-morbidity (presumed systemically healthy)

R12.1. Does periodontal treatment have an impact on systemic inflammation and cardio-metabolic risk in people with no reported systemic co-morbidity (presumed systemically healthy)?

PICOS question addressed by a SR

R12.1: Evidence-based statement

Treatment of periodontitis *may* improve levels of biomarkers of systemic inflammation and cardio-metabolic risk in people with no reported systemic co-morbidity.

Supporting literature (Orlandi et al., 2021)

Quality of evidence Low—Three RCTs at low risk of bias

Grade of recommendation Grade O \leftrightarrow Statement: unclear, additional research needed

Strength of consensus Strong consensus (2.1% of the group abstained due to potential Col)

Background

Intervention

Treatment of periodontitis consisted of step 1 (behaviour change, oral hygiene coaching and supra-gingival PMPR), step 2 (subgingival instrumentation, including removal of dental biofilm and calcified deposits and use of adjunctive therapies), step 3 (surgical periodontal therapy) and step 4 (supportive periodontal care) in order to reduce gingival inflammation (Sanz, Herrera, et al., 2020). Observational evidence has linked periodontitis to increased risk of NCDs (Sanz et al., 2018; Sanz, Marco Del Castillo, et al., 2020). Multiple mechanisms have been proposed to explain the biological plausibility of a systemic effect of periodontitis (Hajishengallis & Chavakis, 2021). Furthermore, evidence from interventional data suggests that the treatment of periodontitis may have an impact on systemic health outcomes (D'Aiuto et al., 2013; Hajishengallis & Chavakis, 2021; Simpson et al., 2015).

Available evidence

Number and design of included studies

Three randomized controlled trials (Fu et al., 2016; Tonetti et al., 2007; Zhou et al., 2017) provided data on the effect of the treatment of periodontitis in comparison with no/control treatment on systemic health in systemically healthy participants with periodontitis. Systemic outcomes reported in these trials included high sensitivity C-reactive protein (hs-CRP), fasting plasma glucose, TNF-alpha, interleukin 6 (IL-6), total cholesterol (TC), high density lipoprotein (HDL) cholesterol, low density lipoprotein (LDL) cholesterol, triglycerides (TG), flow-mediated dilatation (FMD), systolic blood pressure (SBP), diastolic blood pressure (DBP) and body mass index (BMI), at 6 months follow-up. Meta-analyses were performed by systemic outcome for hs-CRP, IL-6, TC, HDL cholesterol, LDL cholesterol, TG, SBP, and DBP, at 6 months.

Risk of bias

The three trials included in the analysis were considered at low risk of bias according to the RoB 2.0 tool.

Effect sizes and their clinical relevance

Although reductions in systemic biomarkers of inflammation and metabolic control were reported, no statistically significant systemic effect of periodontal therapy was observed after 6-months of follow-up.

Consistency

Meta-analysis of three studies showed a reduction in eight biomarkers (hs-CRP, IL-6, TC, HDL, LDL, TG, SBP and DBP) with high levels of heterogeneity in six meta-analyses and wide confidence intervals.

Balance of benefit and harm

No evidence of harm was reported in any of the clinical trials.

Overall certainty of the evidence

Low.

From evidence to recommendation—additional considerations

Acceptability

Little is known about patient preferences in relation to the treatment of periodontitis, due to a paucity of patient-reported outcome data.

Feasibility

Little is known about the implementation of periodontal therapy that is targeted at reducing cardiovascular biomarkers or those targeted at lowering metabolic risk, predominantly because the primary purpose of periodontal treatment is to improve periodontal rather than systemic health outcomes. Treatment of periodontitis is routinely performed in many healthcare systems across the world, and therefore it is in itself feasible for those who can afford care or those able to gain access to care. There are, however, groups in society who struggle to access periodontal care for various reasons, some being cost (The Economist Intelligence Unit, 2021).

Ethical considerations

Evaluation of the efficacy of the treatment of periodontitis on systemic health outcomes is ethically challenging as it would entail comparison with no treatment.

Economic considerations

Cost-effectiveness has not been evaluated in these studies.

Legal considerations

Not applicable.

12.1.2 | Impact of periodontal treatment on systemic inflammation and cardio-metabolic risk in people with co-morbid systemic non-communicable disease (NCD)

R12.2. Does treatment of periodontitis impact upon “hard” outcomes or complications of systemic NCDs, in periodontitis patients with a co-morbid NCD?

PICOS question addressed by a SR

R12.2: Expert consensus-based statement

It is currently unclear if the treatment of periodontitis improves “hard” outcomes or complications of systemic NCDs in patients with periodontitis with a co-morbid NCD.

(Continues)

(Continued)

PICOS question addressed by a SR

R12.2: Expert consensus-based statement

Supporting literature (Orlandi et al., 2021)

Quality of evidence Very Low—One feasibility RCT at high risk of bias

Grade of recommendation Grade O↔ Statement: unclear, additional research needed

Strength of consensus Consensus (2.4% of the group abstained due to potential Col)

Background

Intervention

Treatment of periodontitis was exactly as described under R12.1.

Available evidence

Number and design of included studies

One randomized controlled study (feasibility trial) reported data on the effect of the treatment of periodontitis in comparison with no/control treatment on “hard” outcomes or complications of systemic diseases, in patients with a non-communicable disease (Beck et al., 2008). “Hard” outcomes have been defined as “patient-important endpoints that are definitive with respect to the disease process, and reflect how a patient feels, functions or survives” (Institute of Medicine, 2010).

Risk of bias

The trial was considered at high risk of bias according to the RoB 2.0 tool.

Effect sizes and their clinical relevance

No statistically significant overall effect of periodontal therapy on cardiovascular events was observed.

Consistency

Not applicable.

Balance of benefit and harm

No evidence of harm was reported in this study.

Overall certainty of the evidence

Low.

From evidence to recommendation—additional considerations

Acceptability

Little is known about patient preferences in relation to treatment of periodontitis in patients with an NCD.

Feasibility

Little is known about the implementation of periodontal treatment in patients with NCDs, or where periodontal therapy is prioritized by those patients relative to other co-morbidities within their overall healthcare portfolio. Treatment of periodontitis is routinely performed in many healthcare systems across the world, and therefore it is in itself feasible, accepting the limitations outlined in R12.1.

Ethical considerations

Evaluation of the efficacy of the treatment of periodontitis on systemic health outcomes is ethically challenging as it entails comparison with no treatment or a delayed treatment group, where adverse outcomes may arise in periodontal health, irrespective of systemic health outcomes.

Economic considerations

Insufficient evidence exists on the cost-effectiveness of the treatment of periodontitis when evaluating hard outcomes of systemic disease and/or complications of systemic conditions.

Legal considerations

Not applicable.

R12.3. Does periodontal treatment impact on systemic inflammation, metabolic control and cardiovascular risk in periodontitis patients with a co-morbid NCD?

PICOS question addressed by a SR

R12.3: Evidence-based recommendation

We **suggest** that treatment of periodontitis is **performed** to reduce systemic inflammation, to reduce cardiovascular risk profile and to improve metabolic control in patients with co-morbid NCD; however, treatment protocols should include careful consideration of the general health status of the patient (e.g., quadrant vs. full-mouth approach).

Supporting literature (Orlandi et al., 2021)

Quality of evidence Moderate—Six RCTs were considered at high risk of bias, nine at moderate risk and 18 at low risk of bias

Grade of recommendation Grade B—†

Strength of consensus Consensus (0% of the group abstained due to potential CoI)

Background

Intervention

Treatment of periodontitis was exactly as described under R12.1.

Available evidence

Number and design of included studies

Thirty-three randomized controlled trials reported data on the effect of the treatment of periodontitis in comparison with

no/control treatment on systemic health in patients with a non-communicable disease. Populations with type 2 diabetes, cardiovascular diseases, polycystic ovary syndrome, end-stage renal disease, multiple co-morbidities, rheumatoid arthritis, and chronic kidney disease were reported. Systemic outcomes evaluated included hs-CRP, TNF-alpha, IL-6, erythrocyte sedimentation rate (ESR), HbA1c, fasting plasma glucose (FPG), TC, HDL cholesterol, LDL cholesterol, TG, very low density lipoproteins (VLDL), FMD, BMI, SBP, DBP, pulse rate, serum creatinine (sCR) and albumin, at 6 months follow-up. Meta-analyses were conducted by systemic outcome for hs-CRP, FPG, TNF-alpha, IL-6, TC, HDL cholesterol, LDL cholesterol, TG, FMD, SBP, DBP and BMI, at 6-month follow-up, and for hs-CRP, IL-6, HbA1c, TC, HDL cholesterol, estimated glomerular filtration rate (eGFR) and asymmetric dimethylarginine (ADMA), at 12-month follow-up.

Risk of bias

Of 33 trials, 6 were considered at high risk, 9 at moderate risk and 18 at low risk of bias according to the RoB 2.0 tool.

Effect sizes and their clinical relevance

Treatment of periodontitis demonstrated a statistically significant reduction in hs-CRP (0.47 mg/L, 95% CI [0.20; 0.74]), reduction in FPG (1.07 mmol/L, 95% CI [0.25; 1.89]) and increase in FMD (0.31%, 95% CI [0.07; 0.55]), at 6-month follow-up.

Consistency

Fifteen of 16 clinical trials reported a reduction in serum hs-CRP but with high heterogeneity. Six out of eight studies reported a reduction in FPG but with high heterogeneity. Both of the studies investigating FMD as a primary outcome measure, reported an increase in FMD with low heterogeneity. No evidence of publication bias was observed. Meta-analyses on IL-6 (6 studies), TNF- α (4 studies) and HbA1c (14 studies) reported statistically non-significant reductions following treatment of periodontitis.

Balance of benefit and harm

No evidence of harm was reported in any studies.

Overall certainty of the evidence

Moderate.

From evidence to recommendation—additional considerations

Acceptability

Little is known about patient preferences in relation to treatment of periodontitis in people with a co-morbid systemic condition, due to the lack of studies addressing PROMs.

Feasibility

Little is known about implementation of protocols targeted at reducing systemic biomarkers of cardio-metabolic risk. Treatment of periodontitis

is routinely performed in many healthcare systems across the world, and therefore is in itself feasible, accepting the limitations outlined in R12.1.

Ethical considerations

The evaluation of the efficacy of the treatment of periodontitis on systemic health outcomes is ethically challenging for the reasons stated in R12.2.

Economic considerations

Insufficient evidence exists on the cost-effectiveness of the treatment of periodontitis when evaluating biomarker or other surrogate outcomes of systemic disease and/or complications of systemic conditions.

Legal considerations

Not applicable.

12.1.3 | Does periodontal treatment reduce the risk of adverse pregnancy outcomes?

R12.4. Does periodontal treatment during pregnancy reduce adverse pregnancy outcomes?

PICOS question addressed by a SR

R12.4: Evidence-based statement

It is unclear whether treatment of periodontitis during pregnancy may reduce pre-term births (<37 weeks) or reduces other adverse pregnancy outcomes.

Supporting literature (Orlandi et al., 2021)

Quality of evidence Moderate—3 RCTs were of high risk of bias, 10 studies were graded as moderate and the remaining 3 as low risk of bias.

Grade of recommendation Grade O—↔ Statement: unclear, additional research needed, specifically focused on high-risk individuals.

Strength of consensus Strong consensus (0% of the group abstained due to potential Col)

Background

Intervention

Treatment of periodontitis was as described under R12.1. Epidemiological studies have shown that periodontal diseases increase the odds of several pregnancy complications, with the strength of the associations varying depending upon the study population (Offenbacher et al., 1996; Jeffcoat et al., 2001; Ide & Papapanou, 2013). Prevention strategies including oral hygiene instructions and supra/subgingival instrumentation of the dentition are encouraged during pregnancy to maintain/restore oral health. However, it is still unclear whether the treatment of periodontal diseases has an effect on preventing adverse pregnancy outcomes such as pre-term birth or low birth weight.

Subgingival instrumentation has traditionally been delivered during multiple sessions (e.g., quadrant-wise). As an alternative, full-mouth protocols have been suggested. Full-mouth protocols included single-stage and two-stage therapy within 24 hours; however, protocols including adjunctive antiseptics (full-mouth disinfection) were not included in this analysis.

Available evidence

Number and design of included studies

Sixteen RCTs reported on the effect of treatment of periodontitis on pregnancy outcomes. The pregnancy outcomes assessed in the meta-analysis comprised pre-term birth <37, <35, and <32 weeks, low birth weight <2500 g and less than <1500 g, pre-term low birth weight, pre-eclampsia, gestational age at delivery, CRP, stillbirth, birthweight and perinatal loss.

Risk of bias

Of the 16 studies, 3 were at high risk of bias, 10 studies were graded as moderate and the remaining 3 as low risk of bias based on the ROB 2 tool.

Effect sizes and their clinical relevance

Based on 14 RCTs, the treatment of periodontitis resulted in a statistically significant reduction of pre-term births (<37 weeks) (risk ratio = 0.77, 95% CI [0.60; 0.98]). No statistically significant impact was observed for other pregnancy outcomes.

Consistency

Nine of 14 RCTs resulted in a reduction of pre-term births (<37 weeks).

Balance of benefit and harm

Of the studies included that reported adverse effects, none reported maternal mortality following non-surgical periodontal interventions. Clinicians should be aware that there is evidence of systemic consequences (e.g., acute-phase systemic inflammatory response and vascular dysfunction, i.e., FMD) with full-mouth protocols. While the clinical relevance of such changes remains to be determined, such an approach should always include careful consideration of the general health status of the pregnant patient.

Overall certainty of the evidence

Moderate.

From evidence to recommendation—additional considerations

Acceptability

Little is known about patient preferences regarding the treatment of periodontitis during pregnancy, due to the lack of studies addressing relevant PROMs.

Feasibility

Little is known about implementation protocols targeted at reducing adverse pregnancy outcomes. Treatment of periodontitis is routinely performed in many healthcare systems across the world, and therefore is in itself feasible, accepting the limitations outlined in R12.1.

Ethical considerations

Evaluation of the efficacy of the treatment of periodontitis on adverse pregnancy outcomes is ethically challenging as it would entail comparison with no treatment or treatment delayed until post-partum. There is another potential ethical dilemma in that patient preference may conflict with the clinician's recommendation in terms of mode of treatment delivery or timing of treatment. Patient autonomy should be respected.

Economic considerations

Insufficient evidence exists on the cost-effectiveness of the treatment of periodontitis when evaluating adverse pregnancy outcomes.

Legal considerations

Not applicable.

12.2 | Prosthetic rehabilitation in fully or partially edentulous patients: Impact on systemic health and quality of life

A systematic review (Gennai et al., 2021) explored the impact of treating partial or complete edentulism in patients with periodontitis and those with tooth loss from any cause, on OHRQoL and general health.

The review considered the impact of rehabilitation of edentulous spaces of 5 or more teeth, in order to identify patients with stage IV periodontitis. However, only 13 studies recorded data from periodontitis patients, and even in these studies, the reasons for the tooth loss were not documented. Forty-three studies were available that met the criteria of analysing oral health quality of life outcomes and specific measures of general health (cognitive impairment, nutritional status, frailty, systemic serum markers) in patients with edentulous spaces of 5 or more teeth.

The recommendations made by the workshop focussed on the available data from 13 studies of periodontitis patients. However, as periodontitis is a major cause of tooth loss in adults (overall prevalence 45%–50% and severe periodontitis on 7%–11%), data were also considered from 43 studies where the presence of periodontitis was not specified, on the assumption that it is likely to have contributed to tooth loss in a number of cases studied. This decision reflected the limitations of the available literature, but the guideline group endorsed this pragmatic approach.

OHRQoL was analysed using various validated tools including: OHIP-14, OHIP-20, OHIP-49, OHIP-54, OHIP-EDENT, OHIP-EDENT-21, OHQoL-UK, OIDP, DIDL and GOHAI, and VAS, Likert

scales or questionnaires for patient satisfaction (12 different methods). Quality of life instruments used to analyse general health status included EORTC QLQ-C30, EORTC QLQ-H&N35, GHQ, SIQ, SF-36, WHOQoL-BREF, GSS and EQ-5D ($n = 8$). For explanations of the abbreviations of questionnaires/tools, the reader should read the referred systematic review (Gennai et al., 2021).

12.2.1 | In people with a minimum of five teeth missing for any reason (including stage IV periodontitis patients), does prosthetic rehabilitation of edentulous spaces improve quality of life?

R12.5. Does prosthetic rehabilitation of partial edentulism improve quality of life in people with tooth loss (for any reason, including periodontitis)?

PICOS question addressed by a SR

R12.5: Evidence-based recommendation (1), Evidence-based statements (2, 3)

1. **We recommend** rehabilitation of people with partial edentulism of at least 5 teeth (including those affected by periodontitis) to improve quality of life.
2. Rehabilitation of partial edentulism with tooth-supported, fixed or removable prostheses improves quality of life.
3. Rehabilitation of partial edentulism with implant-supported prostheses improves quality of life.

Supporting literature (Gennai et al., 2021)

Quality of evidence Moderate

Grade of recommendation Grade A—↑↑ (1); Statements (2, 3)

Strength of consensus Strong consensus (1.9% of the group abstained due to potential CoI)

Background

Intervention

Partially edentulous spaces may be left un-restored or may be restored using a variety of approaches, including fixed or removable prostheses, which may or may not be supported by teeth or dental implants. Such restorations provide numerous benefits that are reported to provide positive impacts beyond the oral cavity including but not limited to function and aesthetics. There is also a body of evidence suggesting improvements in quality of life following the restoration of partially edentulous spaces.

OHRQoL has been measured with psychometric questionnaires. Restoration of edentulous spaces appears to incur significant changes in OHRQoL and such changes appear tangible to those patients treated by such reconstructions.

Available evidence

Number and design of included studies

The evidence underpinning this recommendation includes five RCTs (465 patients), one cross-sectional study, a case-control trial (14 patients) and one prospective case series (248 patients), with all studies consistently highlighting that the restoration of partial edentulism led to an improvement in quality of life.

Risk of bias

Risk of bias was evaluated via the Cochrane reviewers' Handbook for interventional studies identifying three studies at high risk of bias and three with moderate risk of bias. One case-control study was at low risk of bias (5 stars out of 6) on the Newcastle-Ottawa scale.

Effect sizes and their clinical relevance

Improvements were consistently statistically significant and the magnitude of improvement was superior to the "minimally important difference" when available, suggesting clinically meaningful effect sizes.

There were no randomized studies evaluating treatment, versus no treatment. Thus, pre-operative to post-operative differences were evaluated in order to assess the impact of the treatment protocol studied.

Consistency

Overall, studies reported that restoration of partial edentulism with tooth-supported restorations led to an improvement in quality of life.

Balance of benefit and harm

Benefits and harms were not reported, although it may be speculated that tooth-supported fixed restorations, may be associated with some potential harm due to complications associated with the endodontic and prosthodontic components of the treatment. In the case of implant-supported restorations, potential harms associated with implant placement (potential intra- and post-operative complications) and maintenance (especially in people affected by periodontitis showing higher risk of peri-implantitis and implant loss) should be carefully considered.

However, overall, the perceived benefits in terms of masticatory function and improvements in quality of life appear to exceed the potential harms.

Overall certainty of the evidence

High (consistency and effect size) to Moderate (quality of the included studies).

From evidence to recommendation—additional considerations

Acceptability

Restoration of edentulism is broadly accepted by both patients and institutions and widely requested by patients.

Feasibility

Approaches to restoring edentulous spaces vary according to the knowledge and skills of individual dental practitioners, dental practice/office protocols, and healthcare funding systems which vary widely from country to country. There seem to be challenges, however, for frail populations and the elderly living in care homes. Moreover, surgical options as pre-treatment for the restorations (e.g., for implant placement) are not universally available, and may reflect healthcare funding and the financial circumstances of individuals, thereby also reflecting health inequalities.

Ethical considerations

Many national health systems cover/reimburse the costs for treatment of edentulism. However, in some countries, fees for treatment are borne entirely by the individual patient.

Economic considerations

The costs associated with the restoration of edentulous spaces vary widely according to the type of restoration provided, with removable options (removable dental prostheses) generally costing significantly less than fixed restorations (tooth/implant-supported fixed partial dental prostheses).

Legal considerations

Not applicable.

R12.6. Does provision of conventional complete removable prostheses, in one or in both dental arches, improve quality of life in fully edentulous patients when compared with no rehabilitation?

PICOS question addressed by a SR

R12.6: Evidence-based recommendation

We recommend providing complete conventional removable prostheses to treat fully edentulous patients (including those who have lost teeth due to periodontitis), in one or both arches, to improve quality of life.

Supporting literature (Gennai et al., 2021)

Quality of evidence Moderate

Grade of recommendation Grade A-↑↑

Strength of consensus Strong consensus (0% of the group abstained due to potential CoI)

Background

Intervention

The traditional approach to treating complete edentulism involves rehabilitation with complete conventional removable prostheses. Overall, this approach to treatment is technically less demanding,

universally accessible and associated with lower costs than implant-supported overdentures.

Available evidence

Number and design of included studies

The evidence supporting this recommendation is based on a systematic review and meta-analysis (Gennai et al., 2021), 44 studies, of which eight were interventional and 36 observational.

Risk of bias

Risk of bias was evaluated through the Cochrane reviewers' Handbook for interventional studies and the Newcastle–Ottawa scale for observational studies. Overall, studies were judged to be of moderate to high risk of bias and none of the manuscripts included were found to be at low risk of bias. Interventional studies were judged to be of moderate to high risk of bias as these studies showed on average 45% of items that were considered adequate. The observational studies showed 57% of items judged positively, thereby indicating a moderate overall risk of bias.

Effect sizes and their clinical relevance

The effect measured through psychometric questionnaires was both statistically significant and clinically meaningful as significant changes of GOHAI, OHIP-14 and OHIP-49 were noticed after rehabilitation.

Based on the analysis, the magnitude of the effect is considered as relevant for the patient, since it was superior, on average, to the value of the “minimally important difference”, as explained before (Tsakos et al., 2012).

Consistency

There is a high level of consistency throughout the selected studies. None of the studies reported an absence of effect. Indeed, all the studies reported significant improvements in the various psychometric questionnaires used.

Balance of benefit and harm

Harms associated with treatment of edentulism with complete conventional removable prostheses are negligible. Patients may have temporary difficulties in adapting to new prostheses. Moreover, poorly fitting complete conventional removable prostheses may be a cause of discomfort.

Overall certainty of the evidence

High.

From evidence to recommendation—additional considerations

Acceptability

Complete conventional RDPs are traditionally the most commonly used method of rehabilitation for fully edentulous patients. In general,

they are safe and effective and the positive changes in quality of life reported improve their acceptability. In some patients, however, the lack of retention associated with the prosthesis may constitute an important psychological issue.

Feasibility

Complete conventional RDPs are routinely provided by dentists worldwide. The design and provision of these prostheses are practical, time-effective and safe. Importantly, the relatively modest costs associated with this type of rehabilitation significantly allow its widespread feasibility.

Ethical considerations

Not applicable.

Economic considerations

Complete conventional RDPs are the least expensive form of treatment to rehabilitate fully edentulous patients. Many national health systems cover/reimburse the costs of treating edentulism. However, in some countries, treatment fees may have to be borne entirely by the patient.

Legal considerations

Not applicable.

R12.7. Are implant-supported full-arch removable dental prosthesis (overdenture) superior to conventional full-arch removable prostheses in terms of improvement of quality of life?

Additional question raised by the WG

R12.7: Expert consensus-based recommendation

We suggest treating fully edentulous people (including those affected by periodontitis) with implant-supported full-arch removable dental prosthesis (overdenture), rather than conventional full-arch removable prostheses, to improve quality of life.

Supporting literature (Awad et al., 2000; Heydecke et al., 2005; Allen et al., 2006; Harris et al., 2013; Muller et al., 2013), included in the systematic review (Gennai et al., 2021)

Quality of evidence Moderate

Grade of recommendation Grade B—↑

Strength of consensus Unanimous consensus (14.6% of the group abstained due to potential Col)

Background

Intervention

Implant-supported full-arch RDPs (“overdentures”) are frequently advocated to overcome complications associated with complete conventional removable prostheses, such as fit, aesthetics and masticatory function.

Available evidence

Number and design of included studies

The evidence for this recommendation is based on four RCTs (3 cohorts, 342 patients) evaluating implant-supported full-arch overdentures versus conventional removable prostheses in fully edentulous patients. Quality of life was evaluated through OHIP-49 and social impact questionnaires.

Risk of bias

Risk of bias was evaluated through the Cochrane reviewers' Handbook for interventional studies identifying two studies at high risk of bias and two with moderate risk of bias.

Effect sizes and their clinical relevance

Differences in outcome appear socially meaningful and statistically relevant.

Consistency

Three studies reported significantly greater improvements in quality of life for patients treated with implant-supported overdentures.

Balance of benefit and harm

Benefits and harms were not reported, although it is plausible that implant placement may be associated with some potential harms. Implant placement procedures may be associated with potential intra- and post-operative complications. Moreover, implants, especially in people previously affected by periodontitis, show higher risk of developing peri-implantitis and implant loss.

Overall certainty of the evidence

Moderate.

From evidence to recommendation—additional considerations

Acceptability

Restoration of edentulism is widely advocated and accepted. However, some patients may not be willing to undergo a surgical procedure for implant placement.

Feasibility

The restoration of edentulous spaces using implant-supported restorations is widely undertaken. However, in certain situations, access to surgical treatment may be limited as surgical facilities and skills may not be present in every dental setting. Furthermore, there are a high number of national healthcare systems that do not provide implant treatments. Moreover, implants require supportive maintenance care in all patients, but especially in those previously affected by periodontitis, which may not always be accessible.

Ethical considerations

The fact that dental implant-supported overdentures are more expensive than complete conventional removable prostheses, may raise some ethical concerns as there may be a gradient of improvement of OHRQoL, for implant-supported overdentures, while for some individuals there are problems with the affordability of the procedure, thereby impacting adversely upon social inequalities.

Economic considerations

Restoration of edentulous spaces with implant-retained prostheses is funded in many countries out-of-pocket with costly implications that may limit uptake.

Legal considerations

Not applicable.

12.2.2 | In people with a minimum of five teeth missing for any reason (including stage IV periodontitis patients), does restoration of edentulous spaces improve systemic health?

R12.8. Is rehabilitation of partial/full edentulism associated with better systemic health (teeth lost for any reason, including stage IV periodontitis)?

PICOS question addressed by a SR

R12.8: Evidence-based recommendation (1), Expert consensus-based recommendation (2)

1. **We recommend** prosthetically treating fully edentulous people in order to improve nutritional status.
2. **We do not know** if treating full edentulism is associated with improvement of frailty, cognitive function or other systemic health benefits.

Supporting literature (Gennai et al., 2021)

Quality of evidence Low

Grade of recommendation Grade A—↑↑ (1); Grade O—↔ Unclear, additional research needed (2)

Strength of consensus Unanimous consensus (0% of the group abstained due to potential CoI)

Background

Intervention

People with untreated edentulism exhibit a higher prevalence of malnutrition, frailty and cognitive impairment compared with those who have been treated for complete edentulism. Restoration of complete edentulism significantly reduces the number of patients at risk of malnutrition.

Available evidence

Number and design of included studies

Evidence for malnutrition derives from one RCT (34 patients), one prospective cohort study (51 patients), and one cross-sectional study (343 patients) utilizing the Mini Nutritional Assessment (MNA), a validated instrument covering numerous aspects of the patient's general health, nutritional state and habits. Evidence for frailty derives from a cross-sectional study (1026 patients) assessing frailty with the Groening Frailty Indicator. Cognitive Impairment was assessed in 240 patients with Mini-Mental Status Examination in a cross-sectional study.

Risk of bias

Risk of bias was evaluated through the Cochrane and Newcastle–Ottawa scale according to study design. A moderate to high risk of bias was detected. Intervention studies showed 50% of items as adequate, generating a moderate risk of bias. Observational studies were judged as moderate to high risk of bias as they reported a range of adequate items varying from 4 to 7.

Effect sizes and their clinical relevance

Treatment reduces the proportion of people at risk of malnutrition to a significant extent.

Consistency

Studies are limited in number but consistent in detecting enhancement of nutritional status after rehabilitation. Levels of frailty, cognitive impairment and malnutrition are higher in untreated edentulous subjects.

Balance of benefit and harm

The potential harms of any restoration are outweighed by the benefits of reducing malnutrition.

Overall certainty of the evidence

Low.

From evidence to recommendation—additional considerations

Acceptability

Restoration of edentulism is widely accepted and requested by patients to enhance masticatory function and improve nutritional status.

Feasibility

The restoration of edentulous spaces/edentulism is widely undertaken and techniques and methods vary significantly among different countries, dental settings and healthcare systems. Overall, the vast majority of the population in many countries benefit from restoration of their edentulous spaces/edentulism.

Ethical considerations

Restoration of edentulous spaces should be advocated, especially in elderly populations, to improve masticatory function, improve nutritional status and reduce the likelihood of frailty.

Economic considerations

Restoration of edentulous spaces/edentulism incurs a wide range of costs dependent upon the type of restoration provided.

Legal considerations

Not applicable.

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CONFLICT OF INTEREST

Individual potential conflict of interest forms were completed by all participants and are available on file at the European Federation of Periodontology and extracted in the Supporting Information, available online (CPGstage4- *Potential conflict of interest*). Potential conflicts of interest, in the previous 36 months, reported by the chairs of the workshop (in alphabetic order) are listed here:

Tord Berglundh (Chair) reports—Grants or contracts from any entity: Dentsply Implants IH AB (Institution grants); Consulting fees: Dentsply Implants IH AB (Personal); Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: Dentsply Implants IH AB, Straumann (Lecture, personal); Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: National Board of Health and Welfare, Sweden, National guidelines in dentistry—periodontal and peri-implant diseases (Institution grant).

Iain Chapple (Chair) reports—Grants or contracts from any entity: GSK (Three grants for clinical research); Unilever (One grant for clinical research); Royalties or licences: Quintessence Book (Royalties); Consulting fees: J&J, GSK, Unilever (Consultancy fees); Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: P&G, Philips Oral Healthcare, Unilever (Sponsored Lecture Fees); Patents planned, issued or pending: Patents (Filed with Philips Oral Healthcare); Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: British Society of Periodontology, EFP, UK Oral and Dental Research Trust; Other financial or non-financial interests: Non-financial (my wife runs Oral Health Innovations that has the licence for PreViser and DEPPA risk assessment software in the UK).

David Herrera (Chair) reports—Grants or contracts from any entity: DentaId, Klockner Implants (Research contract with University, establishing a Chair with unrestricted yearly funds for research); Sunstar, Bonyf AG, Lacer, Isdin, ZIZ Dental Care, GSK, Straumann (Research contract with university); Consulting fees: Unilever (Consulting fees, for continuing education); Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: Colgate, Procter and Gamble-OralB, Klockner Implants, ITI, Johnson & Johnson, Lacer (Lecture fees); Support for attending meetings and/or travel: Procter and Gamble-OralB, Johnson & Johnson, ITI, Klockner Implants (Support for travel when delivering lectures); Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: Sociedad Española de Periodoncia Foundation (Patron), World Dental Federation (Committee member of Global Periodontal Health Project), European Federation of Periodontology (Chair of the Workshop Committee, Scientific chair of EuroPerio10).

Søren Jepsen (Chair) reports—Grants or contracts from any entity: Osteology Foundation (Research contract with University); Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: Straumann, Geistlich Pharma (Lecture Fees); Participation on a Data Safety Monitoring Board or Advisory Board: Colgate, Procter & Gamble (Participation in Advisory Board); Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: German Society of Dentistry and Oral Medicine (DGZMK), Advisory Council on continuing dental education.

Moritz Kepschull (Chair) reports—Grants or contracts from any entity: Genolytic GmbH; Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: Dexcel, Geitslich, GSK, Hu-Friedy, Procter; Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: EFP EC, BSP Council, ESE Guideline Group Co-Chair, DG PARO Guideline Office.

Panos N. Papapanou (Chair) reports—Grants or contracts from any entity: Nobel Biocare, National Institutes of Health (Research grants); Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: Straumann (Personal honorarium), GSK (consultancy fees); Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: Editor-in-Chief, *Journal of Clinical Periodontology*.

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Anton Sculean (Chair) reports—Grants or contracts from any entity: Oral Reconstruction Foundation, Camlog, Geistlich, Straumann, ITI, Regedent, EMS (Research contract with university); Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: Geistlich, Osteology Foundation, Straumann, Camlog, EMS, Regedent (Lecture fees); Support for attending meetings and/or travel: Geistlich, Osteology Foundation.

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SUPPORTING INFORMATION

Additional supporting information may be found in the online version of the article at the publisher's website.

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APPENDIX

Workshop Participants: Mario Aimetti, Bilal Al-Nawas, Juan Blanco, Philippe Bouchard, Maria Clotilde Carra, Tali Chackartchi, Tin Crnić, Francesco D'Aiuto, Bettina Dannewitz, Monique Danser, Jan Derks, Thomas Dietrich, Henrik Dommisch, Nikos Donos, Kenneth Eaton, Marco Ferrari, Elena Figuero, Moshe Goldstein, Marjolaine Gosset, Filippo Graziani, Lisa Heitz-Mayfield, Karin Jepsen, Ronald Jung, Lise Lotte Kirkevang, Dimitrios Kloukos, Bahar Eren Kuru, France Lambert, Luca Landi, Natalie Leow, Rodrigo López, Phoebus Madianos, Conchita Martín, Paula Matesanz, Paulo Melo, Ana Molina, Virginie Monnet Corti, Eduardo Montero, Ian Needleman, Luigi Nibali, Spyridon N. Papageorgiou, Sebastian Paris, Guillermo Pradies, Marc Quirynen, Christoph Ramseier, Stefan Renvert, Mario Rocuzzo, Irena Sailer, Giovanni Salvi, Nerea Sánchez, Ignacio Sanz-Sánchez, Henning Schliephake, Frank Schwarz, Falk Schwendicke, Lior Shapira, Andreas Stavropoulos, Xavier Struillou, Jean Suvan, Wim Teughels, Cristiano Tomasi, Leonardo Trombelli, Katleen Vandamme, Nicola West, Gernot Wimmer, Stefan Wolfart, Nicola Zitzmann.

Methodological Consultant: Ina Kopp.

Workshop Organization: European Federation of Periodontology.

Scientific societies involved in the guideline development process: European Association for Osseointegration; European Federation of Conservative Dentistry; European Prosthodontic Association; European Society for Endodontology.

Other organizations involved in the guideline development process: Council of European Chief Dental Officers; Council of European Dentists; European Dental Students' Association; Platform for Better Oral Health in Europe.