Influence of Depression and Anxiety on Non-Surgical Periodontal Treatment Outcomes: A 6-Month Prospective Study

Catherine Petit 1,2,3, Victor Anadon-Rosinach 1,2, Nicolas Tuzin 4, Jean-Luc Davideau 1,2 and Olivier Huck 1,2,3, * 

Abstract: Periodontal treatment could be worsened by risk factors. Depression and anxiety have been suggested as potentially influencing periodontal treatment outcomes. The aim of this study was to determine their association with non-surgical periodontal treatment outcomes in patients with generalized severe periodontitis (stage III/IV generalized periodontitis) at 6 months. A total of 68 patients diagnosed with generalized severe periodontitis were treated with scaling and root planing (SRP) and were followed at 3 and 6 months. The data of the 54 patients that followed the entire protocol were considered for analysis. Depression and anxiety levels were determined at baseline by the Beck Depression Inventory (BDI) and State-Trait Inventory (STAI) questionnaires. The association between psychological scores and periodontal parameters was evaluated by multivariate analysis. At 3 and 6 months, SRP induced an improvement for all periodontal parameters (plaque index (PI), bleeding on probing (BOP), periodontal probing depth (PPD) and clinical attachment loss (CAL)). BDI and STAI scores were associated with the evolution of PI, BOP, mean PPD and number of sites with PPD > 3 mm and with CAL > 3 mm. Depression and anxiety should be considered as risk factors for SRP and the identification of at-risk patients should be performed using well-established tools.

1. Introduction

Periodontitis is a chronic inflammatory disease induced by oral dysbiosis and characterized by clinical symptoms, including gingival swelling, gingival bleeding, clinical attachment loss (CAL) and tooth mobility [1]. This highly prevalent disease impacts the oral-health-related quality of life and its long-term management is at a high cost [2,3]. The development of periodontitis involves the disruption of the balance between the host inflammatory process will modulate the bone homeostasis through the promotion of osteoclastogenesis, contributing to the destruction of the alveolar bone [7]. Interestingly, several inflammatory mediators (IL-1b, IL-6 and IL-8) implicated in the development of periodontal diseases have correlated with the level of stress, emphasizing a link between periodontal status and psychological status [8,10,12,13].
Currently, the treatment of periodontitis is mainly non-surgical (scaling and root planning (SRP)) and aims to remove oral biofilms and to restore homeostasis between bacterial commensals and the host response [14]. Several factors can also negatively influence SRP outcomes. While smoking and chronic diseases, such as diabetes, are the most described factors [15–17], the psychological status has also been suggested as worsening SRP outcomes [10,18–22]. Indeed, it was observed that patients with increased stress, anxiety and depression scores, as well as those exhibiting negative coping strategies, demonstrate worsened SRP outcomes [18–20]. Depression represents a major health concern, as it was shown that depressed patients are more affected by chronic inflammatory diseases, such as diabetes [23,24]. In the context of periodontal diseases, it was observed that the incidence of depression was higher in periodontitis patients, highlighting a potential relationship between both conditions [20,25,26]. The same type of association was depicted regarding the prevalence of anxiety in patients affected by chronic inflammatory diseases, including periodontitis [27,28]. However, some contradictory results could be found within the scientific literature [29]. Such discrepancies could be explained by the specific type of population investigated and the use of different methods of psychological evaluation. Indeed, several questionnaires are commonly used to determine depression or anxiety, such as the Depression Anxiety Stress Scale (DASS-42), State-Trait Anxiety Inventory (STAI) or Beck Depression Inventory (BDI). DASS-42 is a self-reported questionnaire that evaluates the negative emotional states of depression, anxiety and stress. Each of the three DASS-scales contain 14 items. Indeed, the depression scale assesses dysphoria, hopelessness, the devaluation of life, self-deprecation, lack of interest/involvement, anhedonia and inertia. The essential function of the DASS is to assess the severity of the core symptoms of depression, anxiety and stress [30]. The STAI is a commonly used measure of trait and state anxiety used in clinical settings to diagnose anxiety and to distinguish it from depressive syndromes [31]. It consists of 20 items assessing trait anxiety and 20 items assessing state anxiety. BDI is a 21-item questionnaire that measures characteristic attitudes and symptoms of depression [32].

As the tools used to characterize the association between psychological status and periodontal treatment outcomes are diverse, the aim of this study was to determine the influence of depression and anxiety assessed by self-administered questionnaires i.e., STAI and BDI, on SRP outcomes at 6 months in a French population suffering from severe chronic periodontitis (stage III–IV generalized periodontitis).

2. Materials and Methods

2.1. Study’s Protocol

This is an additional analysis of a study conducted in accordance with the Declaration of Helsinki and approved by the local ethical committee (Comité de protection des personnes 3/30; clinical trials: NCT02568163) [11]. All participants were enrolled at the Department of periodontology, University hospital, Strasbourg, France and informed about the protocol and aim of the study, and gave written consent prior to entering the study.

2.2. Inclusion Criteria

As described previously, adult patients (>18 years old) diagnosed with generalized severe chronic periodontitis [33] (stage III-IV generalized periodontitis) characterized by alveolar bone loss assessed on radiographs, CAL > 4 mm, more than 15 teeth with at least 5% of the sites with PPD > 5 mm and radiographic bone loss were included [11]. Patients with systemic diseases (diabetes, auto-immune diseases, chronic inflammatory diseases, etc.) and/or treated with medications that could affect periodontium, such as antibiotics and anti-inflammatory or psychotropic drugs, in the last 6 months, were excluded. Patients that were treated with specialized periodontal treatment in the last 6 months, as well as pregnant women and patients wearing orthodontic appliances, were also excluded. Patients that did not attend recall, having used antibiotics or anti-inflammatory drugs, were excluded from the study.
2.3. Psychological Measurements

Psychological status was evaluated using self-administered questionnaires. The French versions of BDI and STAI-YA/STAI-YB questionnaires were filled by patients at baseline. BDI questionnaire is a 21-item questionnaire that requires 10 min to be filled. This questionnaire allows for the quantitative evaluation of the intensity of the depressive feelings. STAI has 20 items for assessing trait anxiety and 20 for state anxiety, each item being scored on a 4-point scale, and requires 10 min to be filled. Questionnaire’s scores were blinded to both patients and investigators in charge of the periodontal examination and SRP (Figure 1).

Figure 1. Flow chart of the study.

2.4. Non-Surgical Periodontal Treatment and Follow-Up

Periodontal indexes (PPD, CAL, PI and BOP) were collected prior to the treatment using a PCPUNC 15 periodontal probe (Hu-Friedy, Chicago, IL, USA) at baseline, 3 and 6 months. Patients were treated as follows: oral hygiene instructions, SRP at sites with PPD > 3 mm using ultrasonic devices (Suprasson Newton, Satelec, France) and manual curets (Deppeler, Switzerland) under local anesthesia, in 2 sessions within 7 days. Following each session, patients were instructed to rinse with chlorhexidine (0.12%) mouthwash (Eludril, Pierre Fabre, Cahors, France) for 15 days. At 3 months, residual sites with PPD > 3 mm were re-instrumented. All periodontal examinations, as well as SRP, have been performed by the same operator blinded to psychological measurements results.

2.5. Statistical Analysis

The assessment of examiner reliability showed that more than 90% of repeated measurements were within +/−1 mm of the original, indicating good intra-examiner reliability and agreement (kappa score > 0.9). For an alpha risk of 5% and a power of 80%, the number of subjects required to obtain a correlation coefficient of 0.4 is 47. By adding 15% in case of possible drop out, the minimal number of subjects required is equal to 54 patients [20]. All interactions were tested by the Wald procedure in generalized linear models. Influencing parameters with a p-value < 0.2 were considered in a multivariable model. All statistical tests were two-tailed. A p-value < 0.05 was considered statistically significant. All analyses were performed using R software under its version 3.0 (R Core Team (2014). R: A language and environment for statistical computing. Vienna, Austria).

3. Results

3.1. Study’s Sample and Demographics

Among the 68 patients fulfilling the inclusion criteria that were included in this study, 54 completed the 6 months of follow-up. The main reasons for exclusion were the non-respect of the study’s protocol, especially any cancelled or delayed follow-up sessions. The mean age of the sample population was 51.2 years old (SD = 10.1) and 56% of the sample population was female. Nevertheless, 39% of the patients were smokers, with 17 patients consuming more than 10 cigarettes/day and 26% being an ex-smoker. Seventeen percent were consuming alcohol more than 1 time/day. Seventy percent of the patients were married and 72% were employed. Regarding the psychological status of the patients, the mean score of BDI was 9.6 (6.7), 37.9 (10.8) for STAI-YA and 40.8 (10.4) for STAI-YB.
3.2. Non-Surgical Periodontal Treatment Outcomes

After 6 months, SRP was effective in significantly reducing PI (−0.7 (0.11)), BOP (−35% (11)), mean PPD (−0.7 mm (0.1)) and the number of shallow (PPD > 3 mm) (−27.4 (17.7)) and deep sites (PPD > 5 mm) (−12.2 (12)) (p < 0.05) (Figure 2). Interestingly, age, gender, alcohol consumption and smoking habits negatively influenced SRP outcomes (p < 0.05) and, therefore, such parameters were considered in our analysis in order to determine the association between the BDI and STAI scores and SRP outcomes (Table 1). Interestingly, it was estimated that the BDI score was associated with the evolution of PI (p < 0.001), PPD > 3 mm (p < 0.001) and ∆CAL > 3 mm (p = 0.03). Additionally, STAI-YA was associated with ∆BOP (p < 0.001) and ∆PPD (mean) (p < 0.05), whereas STAI-YB was associated with ∆BOP (p < 0.001) only.

**Figure 2.** Evolution of periodontal parameters following SRP. PI, mean % of sites with BOP, mean PPD (mm), % of sites with PPD > 3 mm, >5 mm, 7 mm, mean CAL (mm), % sites with CAL > 3 mm, >5 mm, 7 mm were measured at baseline, 3 and 6 months. *: p < 0.05 vs. baseline.

**Table 1.** Multivariate analysis of covariance between periodontal treatment outcomes and psychological scores. Associations were analyzed by linear and logistic mixed regression models considering psychological parameters at baseline and the evolution of periodontal parameters between baseline and 6 months. All potential influencing variables were considered (age, smoking, gender) in the mathematical model. Significant associations are in bold (p < 0.05). ∆PI, ∆BOP, ∆PPD and ∆CAL represent the difference for mean PI, BOP, PPD and CAL between baseline and 6 months, respectively; ∆PPD > 3 mm, ∆PPD > 5 mm and ∆PPD > 7 mm represent the difference for number of sites with PPD > 3 mm, >5 mm and 7 mm between baseline and 6 months. ∆CAL > 3 mm, ∆CAL > 5 mm and ∆CAL > 7 mm represent the difference for number of sites with PPD > 3 mm, >5 mm and 7 mm between baseline and 6 months.

| Variable | Score | Estimate (SD) | OR (CI) | p-Value |
|----------|-------|---------------|---------|---------|
| ΔPI      | BDI   | −0.01 (0.003) | <0.001  |
|          | STAI-YA | −0.00 (0.002) | 0.96    |
|          | STAI-YB | 0.01 (0.002) | <0.001  |
| ΔBOP     | BDI   | 0.00 (0.003) | 0.91    |
|          | STAI-YA | 0.019 (0.002) | <0.0001 |
|          | STAI-YB | −0.017 (0.002) | <0.0001 |
### Table 1. Cont.

| Variable       | Score     | Estimate (SD)  | OR (CI)     | p-Value |
|----------------|-----------|----------------|-------------|---------|
| ΔPPD (mean)    | BDI       | −0.010 (0.006) | 0.070       |         |
|                | STAI-YA   | 0.010 (0.004)  | 0.014       |         |
|                | STAI-YB   | −0.006 (0.004) | 0.202       |         |
| ΔPPD > 3 mm    | BDI       | 0.97 (0.95–0.99) | 0.004       |         |
|                | STAI-YA   | 1.01 (0.99–1.02) | 0.113       |         |
|                | STAI-YB   | 0.999 (0.98–1.01) | 0.874      |         |
| ΔPPD > 5 mm    | BDI       | 1.011 (0.98–1.04) | 0.435       |         |
|                | STAI-YA   | 1.01 (0.99–1.03)  | 0.290       |         |
|                | STAI-YB   | 0.99 (0.97–1.01)  | 0.584       |         |
| ΔPPD > 7 mm    | BDI       | 1.03 (0.97–1.10)  | 0.321       |         |
|                | STAI-YA   | 1.01 (0.97–1.05)  | 0.523       |         |
|                | STAI-YB   | 0.98 (0.93–1.02)  | 0.282       |         |
| ΔCAL (mean)    | BDI       | −0.014 (0.008)  | 0.075       |         |
|                | STAI-YA   | 0.003 (0.005)  | 0.637       |         |
|                | STAI-YB   | 0.007 (0.006)  | 0.272       |         |
| ΔCAL > 3 mm    | BDI       | 0.980 (0.96–1.00) | 0.03        |         |
|                | STAI-YA   | 1.007 (0.99–1.02) | 0.453       |         |
|                | STAI-YB   | 1.007 (0.99–1.02) | 0.332       |         |
| ΔCAL > 5 mm    | BDI       | 0.99 (0.97–1.00)  | 0.344       |         |
|                | STAI-YA   | 1.002 (0.98–1.02) | 0.787       |         |
|                | STAI-YB   | 1.008 (0.99–1.02) | 0.268       |         |
| ΔCAL > 7 mm    | BDI       | 0.99 (0.96–1.01)  | 0.486       |         |
|                | STAI-YA   | 1.003 (0.98–1.02) | 0.785       |         |
|                | STAI-YB   | 1.006 (0.98–1.02) | 0.552       |         |

### 4. Discussion

Psychological status has been demonstrated as a risk factor for worsened periodontal treatment outcomes. In this study, a specific focus was made on depression and anxiety, which were evaluated using BDI and STAI self-administered questionnaires, and associations between the score and SRP outcomes were observed. This confirms that several questionnaires can be used by dentists to detect at-risk patients prior to SRP.

Psychological factors have been suggested as being associated with the increased prevalence of periodontitis due to the inflammatory involvement of the diseases, but also due to the impact on the patients’ behaviors [34,35]. However, this hypothesis remains under discussion, as, in some studies, no differences were observed between the healthy population and patients diagnosed with psychological disorders, such as severe depression [36]. A recent meta-analysis pointed out that subjects with periodontal disease had a higher depression scale score and anxiety score; however, the heterogeneity between studies highlighted the necessity of additional ones [37]. Indeed, the characterization of depression could be difficult, as several questionnaires are commonly used.

In this study, we aimed to evaluate the impact of depression and anxiety assessed with STAI and BDI questionnaires on periodontal treatment outcomes. The French versions of STAI and BDI were selected due to their strong validation, reliability, internal consistency and extensive use in order to evaluate the association between psychological status and chronic diseases [38–41]. In a previous analysis where depression level was measured using the depression score of the DASS-42 self-administered questionnaire, an association between depression and worsened SRP outcomes was observed, especially regarding the evolution of BOP and reduction in deep sites with PPD > 5 mm and >7 mm [20]. When the patients’ depression status was evaluated using BDI, a negative association with the SRP outcomes for PI, PPD > 3 mm and CAL > 3 mm was measured. It was previously observed that the depression component of DASS was correlated with the
BDI scores [42]. Here, it should be mentioned that different periodontal parameters were influenced by each questionnaire’s score. This may be explained by the distribution of the patient’s scores on each questionnaire’s scale, and the limited number of patients with high scores of depression. Regarding anxiety, the STAI questionnaire was used to determine the state and trait of anxiety of the patients. It was observed in several clinical studies that anxiety was related to periodontitis prevalence and severity. It is worth noting that anxiety can lead to behavioral changes that might worsen treatment outcomes, or, in a worst case scenario, lead to treatment interruption [43–46]. In a murine model of ligature-induced periodontitis, the presence of anxiety-like behaviors was observed in the acute phase of periodontitis, highlighting a potential role for such a psychological trait or state in the pathological process [47]. Here, the association between STAI scores and periodontal parameters evolution was observed. However, it would have been of interest to precisely determine the influence of the anxiety level on the oral hygiene habits of the patients in order to assess the potential modification on the patient’s behavior and its impact on SRP outcomes. For instance, psychological diseases will be associated with the adoption of health-damaging behaviors, such as tobacco smoking or increased alcohol consumption, inadequate sleep quality, poor diet choices, insufficient hygiene practice or a poor compliance with appointments/postoperative treatment suggestions [48]. Such an observation was observed for an instance in a population of stressed police recruits, where patients with a high level of stress presented more plaque accumulation [49].

Several mechanisms have been suggested to link depression and anxiety to periodontitis. Indeed, neuroinflammation sustained by the chronic inflammation associated with periodontitis, increased systemic lipopolysaccharide and bacterial translocation from periodontal sites through a more permeable blood–brain barrier has been investigated [50,51]. The development of depression-like behavior with brain modifications was observed in mice with experimental periodontitis induced by oral gavage of Porphyromonas gingivalis, such as an increased number of activated astrocytes and decreased levels of mature brain-derived neurotrophic factor and astrocytic p75NTR in the hippocampus [52]. Nevertheless, during periodontitis, cytokines, such as IL-1β, IL-6 and TNF-α, are secreted in response to bacterial insult and could sustain the neuroinflammation process after passage through the blood–brain barrier [34,50]. Such increased levels of cytokines could be measured in the gingival crevicular fluid of depressed patients, as observed in a cohort of depressed women on long-term sick leave [53] and women with stress-related depression [54]. However, these changes vary according to the severity of both periodontitis and psychological disease [55]. A depression state is also associated with a decreased immune response, characterized by decreased leukocyte mobilization, decreased innate and adaptive responses, increased Th1 and Th2 response and decreased immunoprotection, altogether contributing to an increased susceptibility to infection and delayed wound healing [48].

The present study presents some limitations. Firstly, as only patients with severe forms of periodontitis were enrolled, the magnitude of the influence of risk factors, such as depression and anxiety, on SRP outcomes may have been blunted by local factors. In addition, only a reduced number of patients presented high levels of depression or anxiety, and large studies with more patients with different levels of depression or anxiety should be conducted in order to determine clear thresholds for an increased risk. We also decided to include smokers, even if smoking is a well-described risk factor for worsened periodontal treatment outcomes. This strategy was selected in order to avoid excluding patients that potentially exhibit a high level of anxiety or depression. This could be considered as a limitation, and future studies may focus exclusively on non-smoker or smoker populations.

5. Conclusions

In conclusion, it can be mentioned that BDI and STAI questionnaires could be proposed as interesting tools to evaluate the risk of worsened SRP outcomes associated with psychological status. These quick and reliable questionnaires could be used by dentists. This strategy may help to detect patients in need of specialized psychological consulta-
tion and to adapt periodontal treatment strategy. For patients exhibiting psychological diseases, short periodontal maintenance recalls will be of interest in order to maintain periodontal health through constant motivation of the patients and the reinforcement of oral hygiene procedures. During the active phase of the treatment, the use of adjunctive therapies may be of interest (antibiotics, antiseptics), especially in immune-compromised patients [48]. However, the impact of such modified treatment procedures should be evaluated in future studies.

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References

1. Curtis, M.A.; Diaz, P.I.; Van Dyke, T.E. The Role of the Microbiota in Periodontal Disease. *Periodontol. 2000* 2020, 83, 14–25. [CrossRef]

2. GBD 2017 Oral Disorders Collaborators; Bernabe, E.; Marcenes, W.; Hernandez, C.R.; Bailey, J.; Abreu, L.G.; Alipour, V.; Amini, S.; Arabloo, J.; Arefi, Z.; et al. Global, Regional, and National Levels and Trends in Burden of Oral Conditions from 1990 to 2017: A Systematic Analysis for the Global Burden of Disease 2017 Study. *J. Dent. Res. 2020*, 99, 362–373. [CrossRef]

3. Schwendicke, F.; Krois, J.; Engel, A.S.; Seidel, M.; Graetz, C. Long-Term Periodontitis Treatment Costs According to the 2018 Classification of Periodontal Diseases. *J. Dent. 2020*, 99, 103417. [CrossRef] [PubMed]

4. Hajishengallis, G.; Lamont, R.J. Polymicrobial Communities in Periodontal Disease: Their Quasi-Organismal Nature and Dialogue with the Host. *Periodontol. 2000* 2020, 86, 210–230. [CrossRef] [PubMed]

5. Bugueno, I.M.; Batool, F.; Keller, L.; Kuchler-Bopp, S.; Benkirane-Jessel, N.; Huck, O. Porphyromonas Gingivalis Bypasses Epithelial Barrier and Modulates Fibroblastic Inflammatory Response in an in Vitro 3D Spheroid Model. *Sci. Rep. 2018*, 8, 14914. [CrossRef] [PubMed]

6. Cekici, A.; Kantarci, A.; Hasturk, H.; Van Dyke, T.E. Inflammatory and Immune Pathways in the Pathogenesis of Periodontal Disease. *Periodontol. 2000* 2014, 64, 57–80. [CrossRef] [PubMed]

7. Graves, D.T.; Li, J.; Cochran, D.L. Inflammation and Uncoupling as Mechanisms of Periodontal Bone Loss. *J. Dent. Res. 2011*, 90, 143–153. [CrossRef] [PubMed]

8. Decker, A.; Askar, H.; Tatman, M.; Taichman, R.; Wang, H.-L. The Assessment of Stress, Depression, and Inflammation as a Collective Risk Factor for Periodontal Diseases: A Systematic Review. *Clin. Oral Investig. 2020*, 24, 1–12. [CrossRef]

9. Omer-Chiang, R.; Baser, U.; Kucukgurgen, C.; Aykol-Sahin, G.; Huck, O.; Yalcin, F. Impact of Smoking on Neutrophil Enzyme Levels in Gingivitis: A Case-Control Study. *Int. J. Environ. Res. Public Health 2021*, 18, 8075. [CrossRef] [PubMed]

10. Akcali, A.; Huck, O.; Tenenbaum, H.; Davideau, J.L.; Buduneli, N. Periodontal Diseases and Stress: A Brief Review. *J. Oral Rehabil. 2013*, 40, 60–68. [CrossRef]

11. Caton, J.G.; Armitage, G.; Berglundh, T.; Chapple, I.L.C.; Jepsen, S.; Kornman, K.S.; Mealey, B.L.; Papapanou, P.N.; Sanz, M.; Tonetti, M.S. A New Classification Scheme for Periodontal and Peri-Implant Diseases and Conditions—Introduction and Key Changes from the 1999 Classification. *J. Periodontol. 2018*, 89 (Suppl. 1), S1–S8. [CrossRef]

12. Giannopoulou, C.; Kamma, J.J.; Mombelli, A. Effect of Inflammation, Smoking and Stress on Gingival Crevicular Fluid Cytokine Level. *J. Clin. Periodontol. 2003*, 30, 145–153. [CrossRef]

13. Kamma, J.J.; Giannopoulou, C.; Vasdeksis, V.G.S.; Mombelli, A. Cytokine Profile in Gingival Crevicular Fluid of Aggressive Periodontitis: Influence of Smoking and Stress. *J. Clin. Periodontol. 2004*, 31, 894–902. [CrossRef]

14. Van Dyke, T.E.; Sima, C. Understanding Resolution of Inflammation in Periodontal Diseases: Is Chronic Inflammatory Periodontitis a Failure to Resolve? *Periodontol. 2000* 2020, 82, 205–213. [CrossRef] [PubMed]
15. Baeza, M.; Morales, A.; Cisterna, C.; Cavalla, F.; Jara, G.; Isamitt, Y.; Pino, P.; Gamonal, J. Effect of Periodontal Treatment in Patients with Periodontitis and Diabetes: Systematic Review and Meta-Analysis. J. Appl. Oral Sci. Rev. FOB. 2020, 28, e20190248. [CrossRef] [PubMed]

16. Kannaz, B.; Lappin, D.F.; Niel, C.J.; Buduneli, N. Effects of Smoking on Non-Surgical Periodontal Therapy in Patients with Periodontitis Stage III or IV, and Grade C. J. Periodontol. 2020, 91, 442–453. [CrossRef]

17. Boulaamaim, T.; Tenenbaum, H.; Davideau, J.-L.; Huck, O. Selective Influence of Smoking on Periodontal Treatment Outcomes after 3 Years of Follow-Up. Oral Health Prev. Dent. 2020, 18, 823–831. [CrossRef] [PubMed]

18. Wimmer, G.; Janda, M.; Wieselmann-Penker, K.; Jakse, N.; Polansky, R.; Peth, C. Coping with Stress: Its Influence on Periodontal Disease. J. Periodontol. 2002, 73, 1343–1511. [CrossRef] [PubMed]

19. Vettore, M.; Quintaniilha, R.S.; Monteiro da Silva, A.M.; Lamarca, G.A.; Leão, A.T.T. The Influence of Stress and Anxiety on the Response of Non-Surgical Periodontal Treatment. J. Clin. Periodontol. 2005, 32, 1226–1235. [CrossRef] [PubMed]

20. Petit, C.; Anadon-Rosinach, V.; Rettig, L.; Schmidt-Mutter, C.; Tuzin, N.; Davideau, J.-L.; Huck, O. Influence of Psychological Stress on Non-Surgical Periodontal Treatment Outcomes in Patients with Severe Chronic Periodontitis. J. Periodontol. 2020, 92, 186–195. [CrossRef] [PubMed]

21. Rawlinson, A.; Vettore, M.V.; Baker, S.R.; Robinson, P.G. Periodontal Treatment, Psychological Factors and Oral Health-Related Quality of Life. J. Clin. Periodontol. 2021, 48, 226–236. [CrossRef]

22. Rawlinson, A.; Vettore, M.V.; Baker, S.R.; Robinson, P.G. Do Psychological Factors Predict Changes in Oral Health-Related Quality of Life and Clinical Status after Periodontal Treatment? J. Clin. Periodontol. 2021, 48, 795–804. [CrossRef]

23. Lustman, P.J.; Clouse, R.E. Depression in Diabetic Patients: The Relationship between Mood and Glycemic Control. J. Diabetes Complicat. 2005, 19, 113–122. [CrossRef]

24. Lwin, M.N.; Serhal, L.; Holroyd, C.; Edwards, C.J. Rheumatoid Arthritis: The Impact of Mental Health on Disease: A Narrative Review. Rheumatol. Ther. 2020, 7, 457–471. [CrossRef]

25. Aldosari, M.; Helmi, M.; Kennedy, E.N.; Badania, R.; Odani, S.; Agaku, I.; Vardavas, C. Depression, Periodontitis, Caries and Missing Teeth in the USA, NHANES 2009–2014. Fam. Med. Community Health 2020, 8, e00583. [CrossRef] [PubMed]

26. Hsu, C.-C.; Hsu, Y.-C.; Chen, H.-J.; Lin, C.-C.; Chang, K.-H.; Lee, C.-Y.; Chong, L.-W.; Kao, C.-H. Association of Periodontitis and Subsequent Depression: A Nationwide Population-Based Study. Medicine 2015, 94, e2347. [CrossRef]

27. Buchberger, B.; Hupertz, H.; Krabbe, L.; Lux, B.; Mattivi, F.; Siafarikas, A. Symptoms of Depression and Anxiety in Youth with Type 1 Diabetes: A Systematic Review and Meta-Analysis. Psychoneuroendocrinology 2016, 70, 70–84. [CrossRef] [PubMed]

28. Laforgia, A.; Corsalini, M.; Stefanachi, G.; Pettini, F.; Di Venere, D. Assessment of Psychopatologic Traits in a Group of Patients with Adult Chronic Periodontitis: Study on 108 Cases and Analysis of Compliance during and after Periodontal Treatment. Int. J. Med. Sci. 2015, 12, 832–839. [CrossRef] [PubMed]

29. Kim, S.-R.; Nam, S.-H. Comparison of Diagnosed Depression and Self-Reported Depression Symptom as a Risk Factor of Periodontitis: Analysis of 2016–2018 Korean National Health and Nutrition Examination Survey Data. Int. J. Environ. Res. Public Health 2021, 18, 871. [CrossRef]

30. Overview of the DASS and Its Uses. Available online: http://www2.psy.unsw.edu.au/dass/over.htm (accessed on 27 August 2021).

31. Spielberger, C.; Gorsuch, R.; Lushene, R.; Vagg, P.; Jacobs, G. Manual for the State-Trait Anxiety Inventory (Form Y1–Y2); Consulting Psychologists Press: Palo Alto, CA, USA, 1983; Volume IV.

32. Beck, A.T.; Ward, C.H.; Mendelson, M.; Mock, J.; Erbaugh, J. An Inventory for Measuring Depression. Arch. Gen. Psychiatry 1961, 4, 561–571. [CrossRef]

33. Armitage, G.C. Development of a Classification System for Periodontal Diseases and Conditions. Ann. Periodontol. 1999, 4, 1–6.

34. Hashioka, S.; Inoue, K.; Miyaoka, T.; Hayashida, M.; Wake, R.; Oh-Nishi, A.; Inagaki, M. The Possible Causal Link of Periodontitis to Neuropsychiatric Disorders: More Than Psychosocial Mechanisms. Int. J. Mol. Sci. 2019, 20, 3723. [CrossRef]

35. Eger, T.; Wörner, F.; Simon, U.; Konrad, S.; Wolowski, A. Dental Anxiety and Higher Sensory Processing Sensitivity in a Sample of German Soldiers with Inflammatory Periodontal Disease. Int. J. Environ. Res. Public. Health 2021, 18, 1584. [CrossRef]

36. Solis, A.C.O.; Marques, A.H.; Pannuti, C.M.; Lotufo, R.F.M.; Lotufo-Neto, F. Evaluation of Periodontitis in Hospital Outpatients with Major Depressive Disorder. J. Periodontol. 2014, 49, 77–84. [CrossRef]

37. Zheng, D.-X.; Kang, X.-N.; Wang, Y.-X.; Huang, Y.-N.; Pang, C.-F.; Chen, Y.-X.; Kuang, Z.-L.; Peng, Y. Periodontal Disease and Emotional Disorders: A Meta-Analysis. J. Clin. Periodontol. 2021, 48, 180–204. [CrossRef] [PubMed]

38. Bredart, A.; Kop, J.-L.; Fall, M.; Pelissier, S.; Simondi, C.; Dolbeault, S.; Livartowski, A.; Tardivon, A. Magnetic Resonance Imaging study group (STIC IRM 2005) Anxiety and Specific Distress in Women at Intermediate and High Risk of Breast Cancer before and after Surveillance by Magnetic Resonance Imaging and Mammography versus Standard Mammography. Psychooncology 2012, 21, 1185–1194. [CrossRef] [PubMed]

39. Lahlou-Laforêt, K.; Ledru, F.; Niarra, R.; Consoli, S.M. PANIC Investigators Validity of Beck Depression Inventory for the Assessment of Depression in Chronic Heart Failure Patients. J. Affect. Disord. 2015, 184, 256–260. [CrossRef] [PubMed]

40. Cakmak, O.; Alkan, B.A.; Satici, E.; Tasdemir, Z. The Effect of Nonsurgical Periodontal Treatment on Gingival Crevicular Fluid Stress Hormone Levels: A Prospective Study. Oral Dis. 2019, 25, 250–257. [CrossRef] [PubMed]
41. Vergne-Salle, P.; Pouplin, S.; Trouvin, A.P.; Bera-Louville, A.; Soubrier, M.; Richez, C.; Javier, R.M.; Perrot, S.; Bertin, P. The Burden of Pain in Rheumatoid Arthritis: Impact of Disease Activity and Psychological Factors. *Eur. J. Pain* 2020, 24, 1979–1989. [CrossRef] [PubMed]

42. Hekimoglu, L.; Altun, Z.O.; Kaya, E.Z.; Bayram, N.; Bilgel, N. Psychometric Properties of the Turkish Version of the 42 Item Depression Anxiety Stress Scale (DASS-42) in a Clinical Sample. *Int. J. Psychiatry Med.* 2012, 44, 183–198. [CrossRef]

43. Kesim, S.; Unalan, D.; Esen, C.; Ozturk, A. The Relationship between Periodontal Disease Severity and State-Trait Anxiety Level. *J. Pak. Med. Assoc.* 2012, 62, 1304–1308. [PubMed]

44. Khambaty, T.; Stewart, J.C. Associations of Depressive and Anxiety Disorders with Periodontal Disease Prevalence in Young Adults: Analysis of 1999-2004 National Health and Nutrition Examination Survey (NHANES) Data. *Ann. Behav. Med.* 2013, 45, 393–397. [CrossRef]

45. Delgado-Angulo, E.K.; Sabbah, W.; Suominen, A.L.; Vehkalahti, M.M.; Knuuttila, M.; Partonen, T.; Nordblad, A.; Sheiham, A.; Watt, R.G.; Tsakos, G. The Association of Depression and Anxiety with Dental Caries and Periodontal Disease among Finnish Adults. *Community Dent. Oral Epidemiol.* 2015, 43, 540–549. [CrossRef]

46. Kolte, A.P.; Kolte, R.A.; Lathiya, V.N. Association between Anxiety, Obesity and Periodontal Disease in Smokers and Non-Smokers: A Cross-Sectional Study. *J. Dent. Res. Dent. Clin. Dent. Prospect.* 2016, 10, 234–240. [CrossRef] [PubMed]

47. Varotto, B.L.R.; Martinez, R.C.R.; Gouveia, F.V.; Antunes, G.F.; de Campos Fabri, G.M.; Ballester, G.; Antequera, R.; de Siqueira, S.R.D.T.; Fonoff, E.T.; Teixeira, M.J.; et al. Increased Anxiety-Like Behavior in the Acute Phase of a Preclinical Model of Periodontal Disease. *Front. Neurol.* 2020, 11, 598851. [CrossRef] [PubMed]

48. Decker, A.M.; Kapila, Y.L.; Wang, H.-L. The Psychobiological Links between Chronic Stress-Related Diseases, Periodontal/Peri-Implant Diseases, and Wound Healing. *Periodontol.* 2000 2021, 87, 94–106. [CrossRef]

49. Hashioka, S.; Inoue, K.; Hayashida, M.; Wake, R.; Oh-Nishi, A.; Miyaoka, T. Implications of Systemic Inflammation and Periodontitis for Major Depression. *Front. Neurosci.* 2018, 12, 483. [CrossRef]

50. Martínez, M.; Martín-Hernández, D.; Virto, L.; MacDowell, K.S.; Montero, E.; González-Bris, Á.; Marín, M.J.; Ambrosio, N.; Herrera, D.; Leza, J.C.; et al. Periodontal Diseases and Depression: A Pre-Clinical in Vivo Study. *J. Clin. Periodontol.* 2021, 48, 503–527. [CrossRef] [PubMed]

51. Wang, Y.-X.; Kang, X.-N.; Cao, Y.; Zheng, D.-X.; Lu, Y.-M.; Pang, C.-F.; Wang, Z.; Cheng, B.; Peng, Y. Porphyromonas Gingivalis Induces Depression via Downregulating P75NTR-Mediated BDNF Maturation in Astrocytes. *Brain Behav. Immun.* 2019, 81, 523–534. [CrossRef] [PubMed]

52. Johannsen, A.; Rylander, G.; Söder, B.; Asberg, M. Gingival Inflammation, Increased Periodontal Pocket Depth and Elevated Interleukin-6 in Gingival Crevicular Fluid of Depressed Women on Long-Term Sick Leave. *J. Periodontal Res.* 2007, 42, 546–552. [CrossRef] [PubMed]

53. Johannsen, A.; Rylander, G.; Söder, B.; Asberg, M. Dental Plaque, Gingival Inflammation, and Elevated Levels of Interleukin-6 and Cortisol in Gingival Crevicular Fluid from Women with Stress-Related Depression and Exhaustion. *J. Periodontol.* 2006, 77, 1403–1409. [CrossRef] [PubMed]

54. de Solis, A.C.O.; Marques, A.H.; Dominguez, W.V.; de Prado, E.B.A.; Pannuti, C.M.; Lotufo, R.F.M.; Lotufo-Neto, F. Evaluation of Periodontitis in Hospital Outpatients with Major Depressive Disorder. A Focus on Gingival and Circulating Cytokines. *Brain Behav. Immun.* 2016, 53, 49–53. [CrossRef] [PubMed]