
Pattern of tooth mobility in smokers and non-smokers with chronic periodontitis

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Abstract: Periodontitis is an infectious inflammatory disease that is caused by the bacteria of dental plaque resulting in the progressive destruction of the tissue supporting the teeth, that is gingival, periodontal ligament, cementum and the alveolar bone. The progression of the disease is accelerated by tobacco consumption. The aim of this study is to find the pattern of tooth mobility in smokers and non smokers. This retrospective study was conducted in a Private dental college, chennai. The sample size was 100, with 50 smokers and 50 non smokers. The data was collected from the hospital digital database. It was observed that tooth mobility in smokers was prevalent in the posterior region (16%) ($p < 0.05$) and in the maxilla and both the jaws (11% & 23% respectively). Tooth mobility in smokers was commonly seen in the posterior region of maxilla (10%) and anterior region of mandible (20%) for which the P value was found to be statistically significant (< 0.05). For non-smokers, there was an increased tooth mobility observed in the anterior region of the mandible (34%) for which the P value was found to be statistically significant (< 0.05). Within the limits of the study, it can be concluded that smoking causes increased periodontal destruction to the surrounding tissues, with tooth mobility and loss commonly observed in the posterior regions of maxilla and posterior region in both the jaws. Thus, Progression of disease can be prevented by early diagnosis and by tobacco cessation programmes.

Keywords: smoking, periodontitis, tooth mobility, recession, furcation

INTRODUCTION

Periodontitis is an infectious inflammatory disease that is caused by the bacteria of dental plaque resulting in the progressive destruction of the tissue supporting the teeth, that is gingival, periodontal ligament, cementum and the alveolar bone (Newman, 2007),(Pihlstrom, 2001).It is an infective condition and is caused by pathogens namely, *aggregatibacter actinomycetemcomitans*, *porphyromonas gingivalis*, *prevotella intermedia*, *fusobacterium nucleatum* and few other microorganisms (Rai, Kaur and Kharb, 2009).

The first sign of periodontal destruction is from the displacement of junctional epithelium which results in pocket formation between the tooth and the gingiva (Löe *et al.*, 1978). The other signs of disease include root exposure due to recession, mobility furcation environment which can, in later stages, lead to tooth loss. The risk factors involved in periodontal destruction are ageing, tobacco consumption, alcohol consumption, stress and few systemic conditions like diabetes mellitus (Arigbede, Babatope and Bamidele, 2012).

Tooth loss remains a major concern, especially in young adults. Physiological tooth mobility is defined as the slight displacement of the clinical crown of a tooth, that is allowed by the resilience of an intact and healthy periodontium, under the application of a moderate force (Mühlemann, 1954), (Glargia and Lindhe, 1997). The degree of tooth mobility may be influenced by a wide variety of factors, such as: the root surface area with connective tissue attachment and therefore tooth type and morphology (e.g. crown-to-root ratio, number, shape and length of roots etc.) and the structural, biophysical (e.g. viscoelasticity and resilience) and metabolic properties of the periodontal ligament and the supporting alveolar bone.

These properties may be affected by functional (e.g. intensity and direction of occlusal forces), local (e.g. severity of periodontal inflammation) or systemic conditions (e.g. diabetes); such conditions could be either

physiological (e.g. pregnancy) or pathological (e.g. periodontal or periapical abscess) (Mühlemann, Savdir and Rateitschak, 1965).

There are both local and systemic factors that could result in tooth loss. Poor oral hygiene could cause caries and periodontal disease leading to the loss of teeth (Axelsson and Lindhe, 1978). Additionally, there are a number of systemic diseases such as diabetes, hypophosphatasia, leukemia, hyperthyroidism, etc whose effects on the oral cavity could make the teeth susceptible to exfoliation (Fure and Zickert, 1997).

According to studies, tobacco consumption is one of the main risk factors associated with chronic periodontitis (BERGSTROM and J, 2006). Cigarette smoking or other forms of tobacco usage has shown five times increased risk of developing periodontal diseases (Papapanou, 1996). Smokers have a greater risk of tooth loss than a non-smoker (Ahlqvist *et al.*, 1989) and also the furcation involvement in the molars are more frequent in smokers (Mullally and Linden, 1996). Though the mechanism behind smoking and progression of the disease remains unclear, few studies suggest that the local effect as in Vasco construction caused by nicotine along with the decreased oxygen tension can create a favourable environment for colonisation by anaerobic bacteria (Salvi *et al.*, 1997). Smoking also influences factors like chemotaxis, phagocytosis, antibody production which in turn causes progression of the disease (Palmer *et al.*, 2005). The CSTK levels are elevated in smokers with chronic periodontitis when compared to non-smokers (Gajendran, Parthasarathy and Tadeballi, 2018).

In chronic periodontitis, there is destruction of gingival, periodontal ligament, cementum, and the bone. There is a need to regenerate the lost tissue which according to previous studies have shown that stem cells are effective in self renewal and in differentiation to produce specialised tissues (Avinash and Malaippan, 2017). Growth factors such as platelet rich fibrin (PRF) are considered vital mediators in inducing differentiation, proliferation and migration of periodontal progenitor cells (Panda, Jayakumar and Sankari, 2014). The PRGF along with GTR has shown good results in improving clinical and radiographic parameters in patients with chronic periodontitis (Ravi *et al.*, 2017).

Previously our team had conducted numerous clinical trials (Kavarthapu and Thamaraiselvan, 2018), (Ramesh, Ravi and Kaarthikeyan, 2017), (Ramesh, Vellayappan and Ravi, 2019), (Priyanka, Kaarthikeyan and Nadathur, 2017), (Ramesh, Varghese and Doraiswamy, 2016) over the past 5 years. Now we are focusing on retrospective studies. The idea for the current study stemmed from current interest in our community. Our team has rich experience in research and we have collaborated with numerous authors over various topics in the past decade (Deogade, Gupta and Ariga, 2018; Ezhilarasan, 2018; Ezhilarasan, Sokal and Najimi, 2018; Jeevanandan and Govindaraju, 2018; J *et al.*, 2018; Menon *et al.*, 2018; Prabakar *et al.*, 2018; Rajeshkumar *et al.*, 2018, 2019; Vishnu Prasad *et al.*, 2018; Wahab *et al.*, 2018; Dua *et al.*, 2019; Duraisamy *et al.*, 2019; Ezhilarasan, Apoorva and Ashok Vardhan, 2019; Gheena and Ezhilarasan, 2019; Malli Sureshbabu *et al.*, 2019; Mehta *et al.*, 2019; Panchal, Jeevanandan and Subramanian, 2019; Rajendran *et al.*, 2019; Ramakrishnan, Dhanalakshmi and Subramanian, 2019; Sharma *et al.*, 2019; Varghese, Ramesh and Veeraiyan, 2019; Gomathi *et al.*, 2020; Samuel, Acharya and Rao, 2020)

Thus, the current study was conducted to find the pattern of tooth mobility in smokers and non-smokers.

MATERIALS AND METHODS

The study was conducted in a Private dental College, Chennai which is a University set up. The population chosen for the study included patients with chronic periodontitis and tooth mobility with two groups; 1. Patients having a habit of smoking and 2. Patients who don't have the habit of smoking. The data was collected from the hospital digital database. Two examiners were included in the study

The study is a retrospective study. The data was collected over a period of nine months- from June 2019 to March 2020. The sample size was 100, of which 50 were smokers and 50 were non-smokers. The collected data was cross- verified with photographs. The inclusion criteria was all patients with chronic periodontitis and tooth mobility. The exclusion criteria was insufficient or unavailable data on habits and periodontal status.

Data collection

The collected data was based on patients having periodontitis with tooth mobility. The population was divided into 2 groups namely; (i) Smokers and (ii) Non- smokers. The tooth loss pattern was studied under the sub headings; SITE-(i) anterior, (ii) posterior and (iii) both, JAWS- (i) maxilla, (ii) mandible, and (iii) both.

Statistical analysis

The collected data was entered in an excel sheet and tabulated statistically using SPSS software (version 23: IBM Corporation NY USA). Pearson Chi square test was performed and the level of significance was set at 0.05.

Ethical approval

The ethical approval for the retrospective study was obtained from the institutional ethical committee.

RESULTS AND DISCUSSION

Comparing smokers and non-smokers with sites of tooth mobility, it was found that in smokers, tooth mobility was prevalent in the posterior region (16%) for which the P value was found to be statistically significant (0.011) (Figure 1). Association between smokers and non-smokers with jaws was done, it was found that tooth mobility was prevalent in the maxilla and both the jaws (11% & 23% respectively), however the P value was found to be not statistically significant (Figure 2). Comparing the tooth mobility with site and jaws in smokers, it was found that the tooth mobility in the posterior region was more prevalent in maxilla (10%), anterior region was common in mandible (20%) and prevalence for both the sites in both the jaws (26%) (Figure 3). The p value was found to be 0.004 which was statistically significant (<0.05). Smokers in the age group 53-62 years had more prevalence of tooth mobility in the maxilla and mandible (12% & 16% respectively) and 40-52 years age group had prevalence in both the jaws (20%) (Figure 4). The P value was found to be statistically not significant (>0.05). The age group 53-62 years had more prevalence of tooth mobility in anterior, posterior and both the regions (12%,12% and 20% respectively). The P value was found to be statistically not significant (>0.05). (Figure 5). Comparing the tooth mobility with site and jaws in non-smokers, it was observed that, tooth mobility in the posterior region was prevalent in maxilla (6%), anterior region was prevalent in mandible (34%). The P value was found to be statistically significant (0.000) which is <0.05 (Figure 6). Non- smokers in the age group 41-52 years had prevalence for maxilla and both the regions (6% & 22% respectively) and the age group 25-40years had prevalence for mandible region (14%). The P value was found to be statistically not significant (>0.05) (Figure 7). The age group 41-52 years had prevalence for both the sites (28%) and the age group 25-40 years had prevalence for anterior and posterior region (18% & 4% respectively). The P value was found to be statistically not significant (>0.05) (Figure 8).

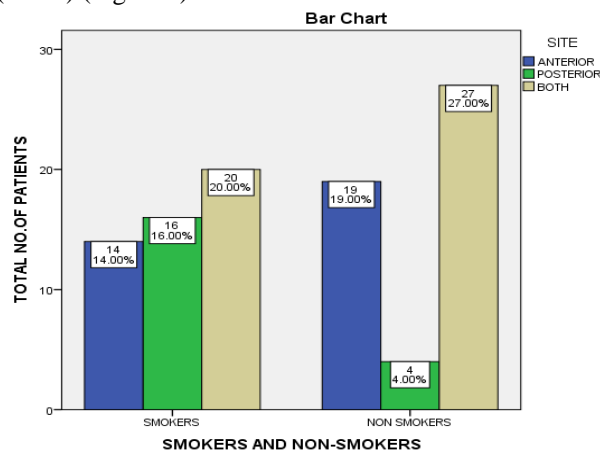


Fig.1: Bar graph showing association of tooth mobility between smokers and non smokers with site. The X axis represents tooth mobility in smokers and non smokers with site and the Y axis represents total number of patients with tooth mobility. It is observed that, in smokers, tooth mobility was more common in the posterior region (green) than the anterior region (blue). In non-smokers, tooth mobility was more common in the anterior region (blue) and both the sites (grey). Chi- square test was done. Pearson Chi-square value-9.000, df-2, p value-0.011, which means there is significant association of tooth mobility between smokers and non smokers with site

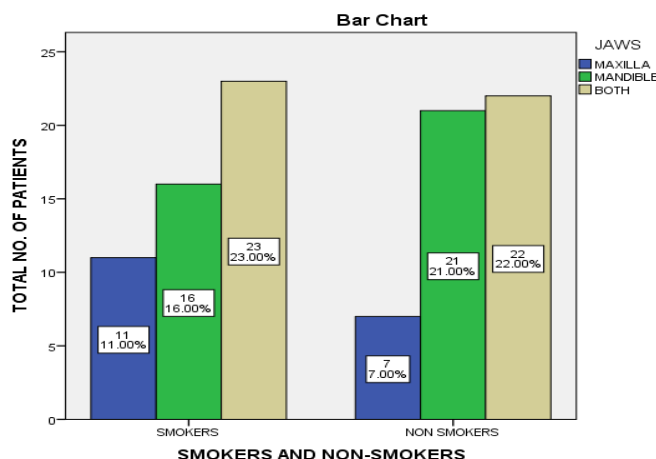


Fig.2: Bar graph showing association of tooth mobility between smokers and non smokers with jaws. The X axis represents smokers and non smokers and the Y axis represents total number of patients with tooth mobility. It is observed that, in smokers, tooth mobility was more prevalent in the maxilla (blue) and both the jaws (grey) than in mandible (green) which was prevalent in non smokers. However, the association is statistically not significant (Pearson Chi square value-1.587, df-2, p value-0.452).

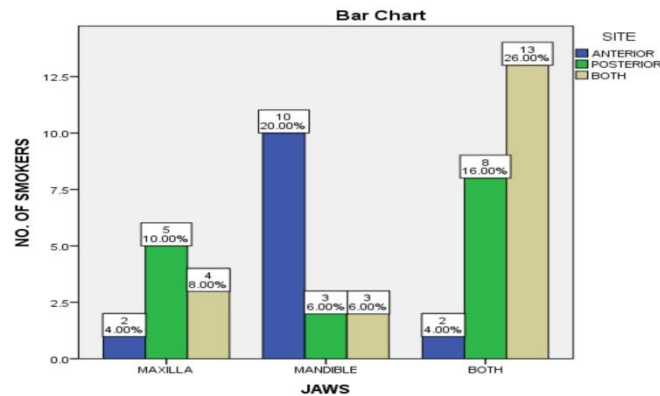


Fig.3: Bar graph showing association of tooth mobility between the site and jaws in smokers. X axis represents jaws and Y axis represents the number of smokers with tooth mobility in anterior, posterior and both the regions. Tooth mobility was seen in both maxilla and mandible with both the sites prevalence. Chi square test was done. There is statistically significant association between the site and jaws in smokers (Pearson Chi-square value-15.209, df-4, p value-0.004).

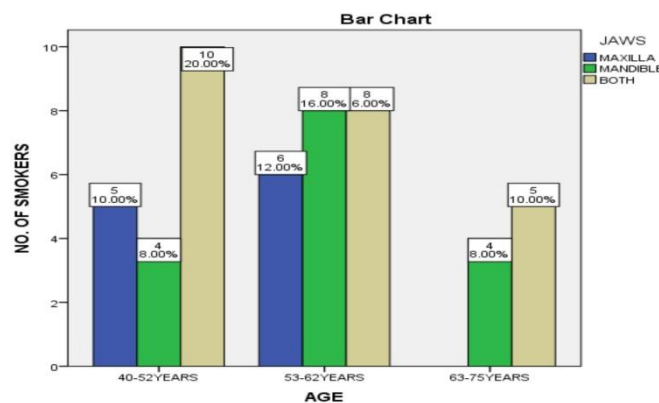


Fig.4: Bar graph showing association of tooth mobility between age and jaws in smokers. X axis represents age and the Y axis represents the number of smokers. Patients in the age group of 53-62 years had prevalence of tooth mobility in maxilla (blue) and mandible (green). Chi square test was done. Pearson Chi square value-4.501, df-4, p value-0.342. However, there is no statistically significant association of tooth mobility between age and jaws in smokers.

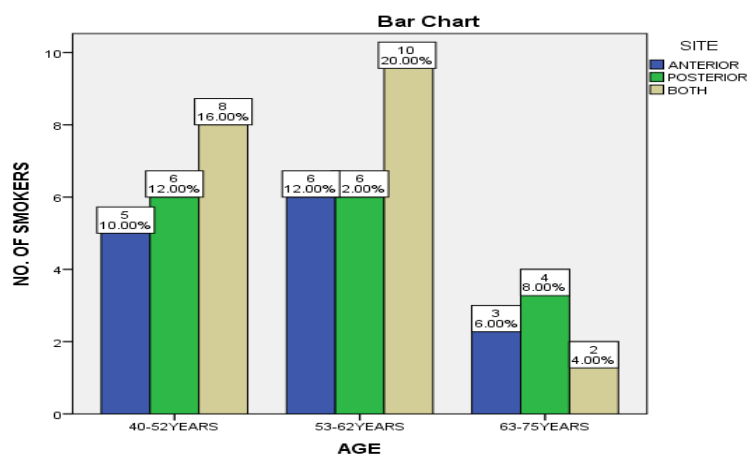


Fig.5: Bar graph showing association of tooth mobility between age and site in smokers. X axis represents age Y axis represents the number of smokers. Patients in the age group of 53-62 years had prevalence of tooth mobility in all three sites observed; anterior, posterior and both the sites (blue, green and grey respectively). Chi square test was done. Pearson Chi square value-1.601, df-4 p value-0.809. However, there is no statistically significant association of tooth mobility between age and site in smokers.

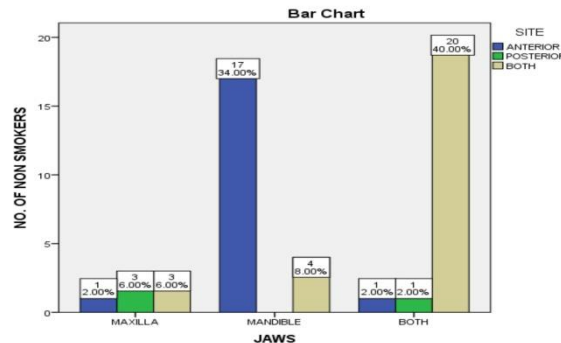


Fig.6: Bar graph showing association of tooth mobility between the site and jaws in non- smokers. X axis represents jaws and Y axis represents the number of non-smokers with tooth mobility in anterior, posterior and both the regions. Tooth mobility was more prevalent in the anterior region of the mandible (blue). Chi square test was done. Pearson Chi-square value-40.813, df-4, p value-0.000. There is statistically significant association between the site and jaws in smokers.

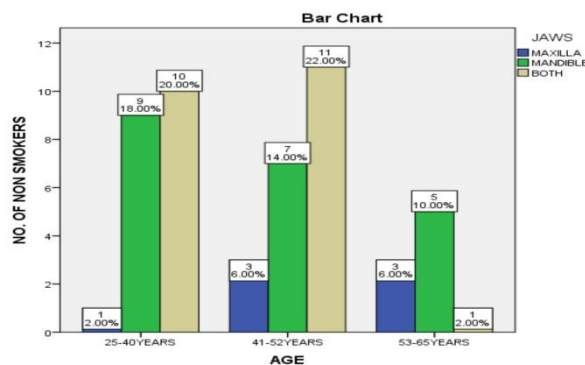


Fig.7: Bar graph showing association of tooth mobility between age and jaws in non- smokers. X axis represents age and the Y axis represents the number of non-smokers. Patients in the age group of 41-52 years had prevalence of tooth mobility in maxilla (blue) and both the jaws (grey) observed. Chi square test was done. Pearson Chi square value-7.085, df-4, p value-0.131. However, there is no statistically significant association of tooth mobility between age and jaws in non smokers.

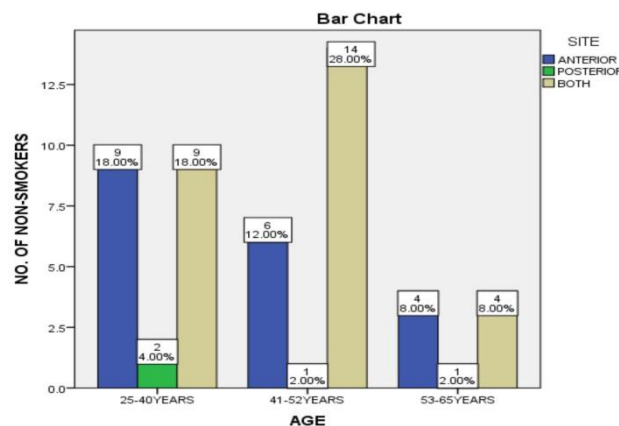


Fig.8: Bar graph showing association of tooth mobility between age and site in non smokers. The X axis represents the age Y axis represents the number of non-smokers. Patients in the age group of

41-52 years had prevalence of tooth mobility in both the sites (grey) observed. Chi square test was done. Pearson Chi square value-2.408, df-4 p value-0.661. However, there is no statistically significant association of tooth mobility between age and site in non smokers.

From the study it is observed that, in smokers, tooth mobility was prevalent in the posterior region (16%) ($p < 0.05$) and was prevalent in the maxilla and both the jaws (11% & 23% respectively). In smokers, the tooth mobility in the posterior region was more prevalent in maxilla (10%), anterior region was common in mandible (20%) and both the sites had prevalence of 26% and non-smokers, it was observed that, tooth mobility in the posterior region was prevalent in maxilla (6%), anterior region was prevalent in mandible (34%) for which the P values were found to be statistically significant (< 0.05).

From Maddipati S et al study, it was observed that the smokers had more teeth with mobility (29%) when compared to non smokers (16%) (Sreedevi, Ramesh and Dwarakanath, 2012). This study is in accordance with the present study. In Ankola A et al study, smokers in the age group 55-60 years had 12.7 % tooth mobility and non-smokers in the age group 55-60 had 11.1% tooth mobility (Pankaj *et al.*, 2007). The current study is in agreement with their study.

According to Smith S et al study, there was a male prevalence among the population, which is in accordance to our study, and also found that the vertical bone loss was more in smokers than in non- smokers (Smith *et al.*, 2019). According to Bergstrom J et al, smokers have significantly greater probing depth, clinical attachment loss and tooth mobility than non smokers (Bergström, Eliasson and Preber, 1991). From Lucinara I et al study, it is observed that there was a tendency for upper tooth loss in smokers and lower tooth loss in non smokers (Luzzi *et al.*, 2007) which is in agreement with the current study where there is 34% of tooth mobility in mandible among non smokers and 10 % tooth mobility in maxilla among smokers. The study also showed a male predominance which is in accordance with our study.

From M Razali et al study, it is observed that smokers had more prevalence for tooth loss and the age group 45 years and above was the most commonly affected group (Razali *et al.*, 2005), which is in accordance with the present study. Surekha V et al study states that smoking increases the periodontal destruction, commonly in the maxillary anterior and premolar region (Velidandla *et al.*, 2019) to which our study shows a similar association.

From Murugan T et al study, it is observed that the coronally advanced flap was found to be a predictable treatment for isolated Miller, class I and II recession defects (Thangakumaran, Gadagi and Arthie, 2015). Few studies show that the salivary TNF-alpha levels are significantly higher in patients with chronic periodontitis (Varghese, Thomas and Jayakumar, 2015). Archana M et al, observed that there is an elevated serum IL-21 levels in patients with chronic periodontitis (Mootha *et al.*, 2016). According to Waleed K et al, serum ET-1 is increased in chronic periodontitis (Khalid, Varghese and Sankari, 2017), (Khalid, Vargheese and Lakshmanan, 2016). Study conducted by Asha R et al, discussed the role of neutrophils in periodontitis which shows hyper/hypo activity to bacterial stimuli (Ramesh *et al.*, 2016).

Thus it is observed that the present study is in accordance with the previous literature and was found to be statistically significant. It can be used as a reference in clinical practice. Though the study was found to be statistically significant, the limitations faced in the study was smaller sample size. Thus, future studies should be done with a larger sample size and equal distribution of study parameters for a better view on the point of study. Our institution is passionate about high quality evidence based research and has excelled in various fields ((Pc, Marimuthu and Devadoss, 2018; Ramesh *et al.*, 2018; Ezhilarasan, Apoorva and Ashok Vardhan, 2019; Ramadurai *et al.*, 2019; Sridharan *et al.*, 2019; Vijayashree Priyadharsini, 2019; Mathew *et al.*, 2020)

CONCLUSION

Within the limits of the study, it can be concluded that smoking causes increased periodontal destruction to the surrounding tissues, with tooth mobility and tooth loss observed in the posterior region of maxilla and posterior region in both the jaws. Thus, Progression of disease can be prevented by early diagnosis and by tobacco cessation programmes.

AUTHOR CONTRIBUTION

Preetha Parthasarathy carried out the retrospective study, planning the study design, collection and analysis of data and drafted the manuscript. Dr. Jeevitha and Dr. Sree Devi aided in conception of the topic, supervision and appraisal of the manuscript.

ACKNOWLEDGEMENT

The study was supported by the Institution who provided the insights and expertise that greatly assisted the study. We would also like to thank the reviewers of the articles for these insights.

Conflict of interest

Authors have no conflict of interest.

REFERENCES

1. Ahlqwist, M. *et al.* (1989) 'Smoking habits and tooth loss in Swedish women', *Community dentistry and oral epidemiology*, 17(3), pp. 144–147.
2. Arigbede, A., Babatope, B. and Bamidele, M. (2012) 'Periodontitis and systemic diseases: A literature review', *Journal of Indian Society of Periodontology*, p. 487. doi: 10.4103/0972-124x.106878.
3. Avinash, K. and Malaippan, S. (2017) 'Methods of isolation and characterization of stem cells from different regions of oral cavity using markers: a systematic review', *International journal of*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5488772/>.
4. Axelsson, P. and Lindhe, J. (1978) 'Effect of controlled oral hygiene procedures on caries and periodontal disease in adults', *Journal of clinical periodontology*, 5(2), pp. 133–151.
5. BERGSTROM and J (2006) 'Periodontitis and smoking : an evidence-based appraisal', *The journal of evidence-based dental practice*, 6, pp. 33–41.
6. Bergström, J., Eliasson, S. and Preber, H. (1991) 'Cigarette smoking and periodontal bone loss', *Journal of periodontology*. Available at: <https://aap.onlinelibrary.wiley.com/doi/abs/10.1902/jop.1991.62.4.242>.
7. Deogade, S., Gupta, P. and Ariga, P. (2018) 'Effect of monopoly-coating agent on the surface roughness of a tissue conditioner subjected to cleansing and disinfection: A Contact Profilometric In vitro study', *Contemporary Clinical Dentistry*, p. 122. doi: 10.4103/ccd.ccd_112_18.
8. Dua, K. *et al.* (2019) 'The potential of siRNA based drug delivery in respiratory disorders: Recent advances and progress', *Drug development research*, 80(6), pp. 714–730.
9. Duraisamy, R. *et al.* (2019) 'Compatibility of Nonoriginal Abutments With Implants: Evaluation of Microgap at the Implant-Abutment Interface, With Original and Nonoriginal Abutments', *Implant dentistry*, 28(3), pp. 289–295.
10. Ezhilarasan, D. (2018) 'Oxidative stress is bane in chronic liver diseases: Clinical and experimental perspective', *Arab journal of gastroenterology: the official publication of the Pan-Arab Association of Gastroenterology*, 19(2), pp. 56–64.
11. Ezhilarasan, D., Apoorva, V. S. and Ashok Vardhan, N. (2019) 'Syzygium cumini extract induced reactive oxygen species-mediated apoptosis in human oral squamous carcinoma cells', *Journal of oral pathology & medicine: official publication of the International Association of Oral Pathologists and the American Academy of Oral Pathology*, 48(2), pp. 115–121.
12. Ezhilarasan, D., Sokal, E. and Najimi, M. (2018) 'Hepatic fibrosis: It is time to go with hepatic stellate cell-specific therapeutic targets', *Hepatobiliary & pancreatic diseases international: HBPD INT*, 17(3), pp. 192–197.
13. Fure, S. and Zickert, I. (1997) 'Incidence of tooth loss and dental caries in 60-, 70-and 80-year-old Swedish individuals', *Community dentistry and oral epidemiology*. Available at: <https://onlinelibrary.wiley.com/doi/abs/10.1111/j.1600-0528.1997.tb00911.x>.
14. Gajendran, P., Parthasarathy, H. and Tadepalli, A. (2018) 'Comparative evaluation of cathepsin K levels in gingival crevicular fluid among smoking and nonsmoking patients with chronic periodontitis', *Indian Journal of Dental Research*, p. 588. doi: 10.4103/ijdr.ijdr_95_17.
15. Gheena, S. and Ezhilarasan, D. (2019) 'Synergic acid triggers reactive oxygen species-mediated cytotoxicity in HepG2 cells', *Human & experimental toxicology*, 38(6), pp. 694–702.
16. Glargia, M. and Lindhe, J. (1997) 'Tooth mobility and periodontal Disease', *Journal of Clinical Periodontology*, pp. 785–795. doi: 10.1111/j.1600-051x.1997.tb01190.x.
17. Gomathi, A. C. *et al.* (2020) 'Anticancer activity of silver nanoparticles synthesized using aqueous fruit shell extract of Tamarindus indica on MCF-7 human breast cancer cell line', *Journal of Drug Delivery Science and Technology*, p. 101376. doi: 10.1016/j.jddst.2019.101376.
18. Jeevanandan, G. and Govindaraju, L. (2018) 'Clinical comparison of Kedo-S paediatric rotary files vs manual instrumentation for root canal preparation in primary molars: a double blinded randomised clinical trial', *European Archives of Paediatric Dentistry*, pp. 273–278. doi: 10.1007/s40368-018-0356-6.
19. J, P. C. *et al.* (2018) 'Prevalence and measurement of anterior loop of the mandibular canal using CBCT: A cross sectional study', *Clinical implant dentistry and related research*, 20(4), pp. 531–534.
20. Kavarthapu, A. and Thamaraiselvan, M. (2018) 'Assessing the variation in course and position of inferior alveolar nerve among south Indian population: A cone beam computed tomographic study', *Indian journal of dental research: official publication of Indian Society for Dental Research*, 29(4), p. 405.
21. Khalid, W., Vargheese, S. S. and Lakshmanan, R. (2016) 'Role of endothelin-1 in periodontal diseases: A structured review', *Indian journal of dental research: official publication of Indian Society for Dental Research*. Available at: <http://www.ijdr.in/tocd.asp?2016/27/3/323/186247/3>.
22. Khalid, W., Vargheese, S. S. and Sankari, M. (2017) 'Comparison of serum levels of endothelin-1 in chronic periodontitis patients before and after treatment', *Journal of clinical and*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5449924/>.
23. Loe, H. *et al.* (1978) 'The Natural History of Periodontal Disease in Man: The Rate of Periodontal

- Destruction Before 40 Years of Age', *Journal of periodontology*, 49(12), pp. 607–620.
24. Luzzi, L. I. T. *et al.* (2007) 'Evaluation of clinical periodontal conditions in smokers and non-smokers', *Journal of applied oral science: revista FOB*, 15(6), pp. 512–517.
 25. Malli Sureshababu, N. *et al.* (2019) 'Concentrated Growth Factors as an Ingenious Biomaterial in Regeneration of Bony Defects after Periapical Surgery: A Report of Two Cases', *Case reports in dentistry*, 2019, p. 7046203.
 26. Mathew, M. G. *et al.* (2020) 'Evaluation of adhesion of Streptococcus mutans, plaque accumulation on zirconia and stainless steel crowns, and surrounding gingival inflammation in primary ...', *Clinical oral investigations*. Available at: <https://link.springer.com/article/10.1007/s00784-020-03204-9>.
 27. Mehta, M. *et al.* (2019) 'Oligonucleotide therapy: An emerging focus area for drug delivery in chronic inflammatory respiratory diseases', *Chemico-biological interactions*, 308, pp. 206–215.
 28. Menon, S. *et al.* (2018) 'Selenium nanoparticles: A potent chemotherapeutic agent and an elucidation of its mechanism', *Colloids and Surfaces B: Biointerfaces*, pp. 280–292. doi: 10.1016/j.colsurfb.2018.06.006.
 29. Mootha, A. *et al.* (2016) 'The Effect of Periodontitis on Expression of Interleukin-21: A Systematic Review', *International journal of inflammation*, 2016. doi: 10.1155/2016/3507503.
 30. Mühlemann, H. R. (1954) 'Tooth Mobility: The Measuring Method. Initial and Secondary Tooth Mobility', *Journal of periodontology*, 25(1), pp. 22–29.
 31. Mühlemann, H. R., Savdir, S. and Rateitschak, K. H. (1965) 'Tooth Mobility — Its Causes and Significance', *Journal of Periodontology*, pp. 148–153. doi: 10.1902/jop.1965.36.2.148.
 32. Mullally, B. H. and Linden, G. J. (1996) 'Molar furcation involvement associated with cigarette smoking in periodontal referrals', *Journal of clinical periodontology*, 23(7), pp. 658–661.
 33. Newman, M. G. (2007) 'Classification and epidemiology of periodontal diseases. Carraza's Clinical Periodontology'. Philadelphia: WB Saunders Company.
 34. Palmer, R. M. *et al.* (2005) 'Mechanisms of action of environmental factors--tobacco smoking', *Journal of clinical periodontology*, 32, pp. 180–195.
 35. Panchal, V., Jeevanandan, G. and Subramanian, E. M. G. (2019) 'Comparison of post-operative pain after root canal instrumentation with hand K-files, H-files and rotary Kedo-S files in primary teeth: a randomised clinical trial', *European archives of paediatric dentistry: official journal of the European Academy of Paediatric Dentistry*, 20(5), pp. 467–472.
 36. Panda, S., Jayakumar, N. D. and Sankari, M. (2014) 'Platelet rich fibrin and xenograft in treatment of intrabony defect', *Contemporary clinical*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4229771/>.
 37. Pankaj *et al.* (2007) 'Assessment of periodontal status and loss of teeth among smokers and non-smokers in Belgaum city', *Indian Journal of Community Medicine*, p. 75. doi: 10.4103/0970-0218.53413.
 38. Papapanou, P. N. (1996) 'Periodontal Diseases: Epidemiology', *Annals of Periodontology*, pp. 1–36. doi: 10.1902/annals.1996.1.1.1.
 39. Pc, J., Marimuthu, T. and Devadoss, P. (2018) 'Prevalence and measurement of anterior loop of the mandibular canal using CBCT: A cross sectional study', *Clinical implant dentistry and related research*. Available at: <https://europepmc.org/article/med/29624863>.
 40. Pihlstrom, B. L. (2001) 'Periodontal risk assessment, diagnosis and treatment planning', *Periodontology 2000*, pp. 37–58. doi: 10.1034/j.1600-0757.2001.22250104.x.
 41. Prabakar, J. *et al.* (2018) 'Comparative Evaluation of Retention, Cariostatic Effect and Discoloration of Conventional and Hydrophilic Sealants - A Single Blinded Randomized Split Mouth Clinical Trial', *Contemporary clinical dentistry*, 9(Suppl 2), pp. S233–S239.
 42. Priyanka, S., Kaarthikeyan, G. and Nadathur, J. D. (2017) 'Detection of cytomegalovirus, Epstein-Barr virus, and Torque Teno virus in subgingival and atheromatous plaques of cardiac patients with chronic ...', *Journal of Indian*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5846241/>.
 43. Rai, B., Kaur, J. and Kharb, S. (2009) 'Pregnancy gingivitis and periodontitis and its systemic effect', *The Internet Journal of Dental Science*, 6(2).
 44. Rajendran, R. *et al.* (2019) 'Comparative Evaluation of Remineralizing Potential of a Paste Containing Bioactive Glass and a Topical Cream Containing Casein Phosphopeptide-Amorphous Calcium Phosphate: An in Vitro Study', *Pesquisa Brasileira em Odontopediatria e Clínica Integrada*, pp. 1–10. doi: 10.4034/pboci.2019.191.61.
 45. Rajeshkumar, S. *et al.* (2018) 'Biosynthesis of zinc oxide nanoparticles using Mangifera indica leaves and evaluation of their antioxidant and cytotoxic properties in lung cancer (A549) cells', *Enzyme and microbial technology*, 117, pp. 91–95.
 46. Rajeshkumar, S. *et al.* (2019) 'Antibacterial and antioxidant potential of biosynthesized copper nanoparticles mediated through Cissus arnotiana plant extract', *Journal of photochemistry and photobiology. B, Biology*, 197, p. 111531.
 47. Ramadurai, N. *et al.* (2019) 'Effectiveness of 2% Articaine as an anesthetic agent in children: randomized

- controlled trial', *Clinical oral investigations*, 23(9), pp. 3543–3550.
48. Ramakrishnan, M., Dhanalakshmi, R. and Subramanian, E. M. G. (2019) 'Survival rate of different fixed posterior space maintainers used in Paediatric Dentistry - A systematic review', *The Saudi dental journal*, 31(2), pp. 165–172.
 49. Ramesh, A. et al. (2016) 'Chronic obstructive pulmonary disease and periodontitis--unwinding their linking mechanisms', *Journal of oral biosciences / JAOB, Japanese Association for Oral Biology*, 58(1), pp. 23–26.
 50. Ramesh, A. et al. (2018) 'Comparative estimation of sulfiredoxin levels between chronic periodontitis and healthy patients - A case-control study', *Journal of periodontology*, 89(10), pp. 1241–1248.
 51. Ramesh, A., Ravi, S. and Kaarthikeyan, G. (2017) 'Comprehensive rehabilitation using dental implants in generalized aggressive periodontitis', *Journal of Indian Society of*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5771115/>.
 52. Ramesh, A., Varghese, S. S. and Doraiswamy, J. N. (2016) 'Herbs as an antioxidant arsenal for periodontal diseases', *Journal of*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4805154/>.
 53. Ramesh, A., Vellayappan, R. and Ravi, S. (2019) 'Esthetic lip repositioning: A cosmetic approach for correction of gummy smile--A case series', *Journal of Indian*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6519099/>.
 54. Ravi, S. et al. (2017) 'Additive Effect of Plasma Rich in Growth Factors With Guided Tissue Regeneration in Treatment of Intrabony Defects in Patients With Chronic Periodontitis: A Split-Mouth Randomized Controlled Clinical Trial', *Journal of Periodontology*, pp. 839–845. doi: 10.1902/jop.2017.160824.
 55. Razali, M. et al. (2005) 'A retrospective study of periodontal disease severity in smokers and non-smokers', *British dental journal*, 198(8), pp. 495–498.
 56. Salvi, G. E. et al. (1997) 'Influence of risk factors on the pathogenesis of periodontitis', *Periodontology 2000*, pp. 173–201. doi: 10.1111/j.1600-0757.1997.tb00197.x.
 57. Samuel, S. R., Acharya, S. and Rao, J. C. (2020) 'School Interventions-based Prevention of Early-Childhood Caries among 3-5-year-old children from very low socioeconomic status: Two-year randomized trial', *Journal of public health dentistry*, 80(1), pp. 51–60.
 58. Sharma, P. et al. (2019) 'Emerging trends in the novel drug delivery approaches for the treatment of lung cancer', *Chemico-biological interactions*, 309, p. 108720.
 59. Smith, S. et al. (2019) 'VERTICAL RADIOGRAPHIC ALVEOLAR BONE LOSS IN SMOKERS AND NON-SMOKERS WITH PERIODONTITIS: A PILOT STUDY', *Pakistan Oral & Dental Journal*, 39(4), pp. 317–324.
 60. Sreedevi, M., Ramesh, A. and Dwarakanath, C. (2012) 'Periodontal Status in Smokers and Nonsmokers: A Clinical, Microbiological, and Histopathological Study', *International journal of dentistry*, 2012. doi: 10.1155/2012/571590.
 61. Sridharan, G. et al. (2019) 'Evaluation of salivary metabolomics in oral leukoplakia and oral squamous cell carcinoma', *Journal of oral pathology & medicine: official publication of the International Association of Oral Pathologists and the American Academy of Oral Pathology*, 48(4), pp. 299–306.
 62. Thangakumaran, S., Gadagi, J. S. and Arthie, T. (2015) 'Comparative clinical evaluation of coronally advanced flap with or without platelet rich fibrin membrane in the treatment of isolated gingival recession', *Journal of Indian*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4365161/>.
 63. Varghese, S. S., Ramesh, A. and Veeraiyan, D. N. (2019) 'Blended Module-Based Teaching in Biostatistics and Research Methodology: A Retrospective Study with Postgraduate Dental Students', *Journal of dental education*, 83(4), pp. 445–450.
 64. Varghese, S. S., Thomas, H. and Jayakumar, N. D. (2015) 'Estimation of salivary tumor necrosis factor-alpha in chronic and aggressive periodontitis patients', *Contemporary clinical*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4632215/>.
 65. Velidandla, S. et al. (2019) 'Distribution of Periodontal Pockets Among Smokers and Nonsmokers in Patients with Chronic Periodontitis: A Cross-sectional Study', *Cureus*. Available at: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6820887/>.
 66. Vijayashree Priyadharsini, J. (2019) 'In silico validation of the non-antibiotic drugs acetaminophen and ibuprofen as antibacterial agents against red complex pathogens', *Journal of periodontology*, 90(12), pp. 1441–1448.
 67. Vishnu Prasad, S. et al. (2018) 'Report on oral health status and treatment needs of 5-15 years old children with sensory deficits in Chennai, India', *Special care in dentistry: official publication of the American Association of Hospital Dentists, the Academy of Dentistry for the Handicapped, and the American Society for Geriatric Dentistry*, 38(1), pp. 58–59.
 68. Wahab, P. U. A. et al. (2018) 'Scalpel Versus Diathermy in Wound Healing After Mucosal Incisions: A Split-Mouth Study', *Journal of oral and maxillofacial surgery: official journal of the American Association of Oral and Maxillofacial Surgeons*, 76(6), pp. 1160–1164.