



ISSN: 0976-3031

Available Online at <http://www.recentscientific.com>

CODEN: IJRSFP (USA)

*International Journal of Recent Scientific Research*  
Vol. 8, Issue, 5, pp. 16945-16948, May, 2017

**International Journal of  
Recent Scientific  
Research**

DOI: 10.24327/IJRSR

## Research Article

### THE INFLUENCE OF SOME IATROGENIC FACTORS ON A CYTOMORPHOMETRIC ANALYSIS OF GINGIVAL CELLS AND PERIODONTAL STATUS AMONG DIABETIC PATIENTS

Mohammed MA. Abdullah Al-Abdaly<sup>1</sup>, Mohammed Salman J. Almalki<sup>2</sup>  
and Kamel A. Saleh<sup>3</sup>

<sup>1</sup>Periodontics, College of Dentistry, King Khalid University, Abha, Saudi Arabia

<sup>2</sup>Dental Doctor, College of Dentistry, King Khalid University, Abha, Saudi Arabia

<sup>3</sup>Biology, College of Sciences- King Khalid University

DOI: <http://dx.doi.org/10.24327/ijrsr.2017.0805.0250>

#### ARTICLE INFO

##### Article History:

Received 10<sup>th</sup> February, 2017

Received in revised form 14<sup>th</sup>

March, 2017

Accepted 08<sup>th</sup> April, 2017

Published online 28<sup>th</sup> May, 2017

##### Key Words:

A cytomorphometric analysis/  
gingival cells / iatrogenic factors.

#### ABSTRACT

**Background:** The relation between dental restorations and periodontal health are more important so the healthy periodontal status requires correction of dental restorations that is mandatory for the protection of periodontal tissues.

**Aims of the study:** The present study was designed to evaluate the effect of some iatrogenic factors on a cytomorphometric analysis of gingival cells and periodontal status among diabetic patients.

**Subjects and Methods:** The sample of the present study consisted of three hundred diabetic patients and they were divided into three equal groups 100 patients per group control group, composite fillings group and metal ceramic crown group. They were selected from the clinics of the college of dentistry King Khalid University. Plaque index (PI), gingival index (GI), and clinical attachment loss (CAL) were recorded. The cytomorphometric changes were evaluated by the Cytological investigation (MN Apoptotic and Necrotic Test). The results were collected and analyzed by ANOVA test.

**Results:** In the present study there were significant differences were detected in all clinical parameters in the samples of the study groups where there was an increase in (PI), (GI), and (CAL) of study groups compared to control group. In a cytomorphometric analysis of gingival cells, there was no significant increase of apoptosis and necrosis compared to control group.

**Conclusions:** The results of a current study revealed that there were increased gingival inflammation and attachment loss in both study groups compared to the control group, but there was a significance induction of micronuclei formation in all groups.

Copyright © Mohammed MA. Abdullah Al-Abdaly *et al*, 2017, this is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution and reproduction in any medium, provided the original work is properly cited.

#### INTRODUCTION

Biocompatibility of dental materials is an imperative for the patient, clinician, research facility expert, and the maker. Preferably, a dental material that will be utilized as a part of the oral cavity ought to be innocuous to every single oral tissue, gingiva, mucosa, pulp, and bone. Moreover, it ought to contain no dangerous, leachable or diffusible substances that can be assimilated into the bloodstream, bringing on systemic reactions; including teratogenic or cancer-causing impacts and their contact with tissues should be biologically acceptable and non destructive<sup>1</sup>.

Moreover, the materials ought to likewise be free of operators that could evoke vulnerability or a hypersensitive reaction in a sensitive patient where dental filling materials may bring about various responses in the oral mucosa, for example, gingiva. It is

not clear today the amount of the in vivo watched cytotoxicity is brought on, either by the dental filling materials or by microbial dental plaque that collects on teeth and dental fillings<sup>2</sup>, but the filling materials display some cytotoxicity that will be diminished considerably in time as a result of the buffering and protein-restricting impacts of salivation to relieve against the cytotoxic impacts<sup>3</sup>.

Composite resin materials are at first exceptionally cytotoxic in vitro trial of direct contact with fibroblasts<sup>4</sup>. The chemical interaction of a toxic substance with biologically relevant molecules is mandatory for local toxicity, whereas the compatibility of tissue may also based on the causes of other material toxicity<sup>5</sup>. On the other hand, in other vitro study that was carried out on newly and developed restorative materials which revealed decreased water solubility and absorption and displayed high characteristics of biocompatibility<sup>6&7</sup>.

\*Corresponding author: Mohammed MA. Abdullah Al-Abdaly

Periodontics, College of Dentistry, King Khalid University, Abha, Saudi Arabia

Dental ceramics are used when the esthetics is required such as crowns, inlays, onlays and veneers on anterior teeth<sup>8</sup>, where the first used of ceramic abutment Ceramic Core was in 1993<sup>9&10</sup>. Furthermore, the metal ceramic systems are durability therapy characterized by long success rate, but there are disadvantages of these systems such as biocompatibility and aesthetics, through the last years there were many new ceramic systems that developed with the ability to anterior and posterior teeth restoration<sup>11</sup>.

The surface characteristics of the materials influence on the biofilm growth, i.e. surface roughness and surface-free energy where the surface roughness is more effective on biofilm retention more than surface-free energy<sup>12</sup>. Diabetes mellitus is considered a metabolic disease characterized by pancreas dysfunction in beta cells of the islets of Langerhans that is causing increased in the level of blood sugar and display of sugar in the urine secretion<sup>13</sup>, it influences on 2-10% of the population<sup>14</sup> so it is the most common metabolic disorders in the world<sup>15</sup> on the other hand periodontal disease is considered as a complication of diabetes mellitus but there is no direct relation between initiation of periodontal disease and diabetes mellitus occurrence<sup>16</sup>

So the present study was designed to evaluate the influence of some iatrogenic factors on a cytomorphometric analysis of gingival cells and periodontal status among diabetic patients.

### Subjects and Methods

#### Patients samples

Three hundred diabetic patients under dental restorative therapy by composite fillings and metal ceramic crowns were selected from outpatient clinics, college of dentistry, King Khalid University. They have not received any periodontal treatment since at least six months and they were divided according to dental reconstruction into three equal groups. Group I patients without dental reconstruction were considered as a control group and two group study included group II patients were received composite fillings for restoration of class II dental caries and group III patients were received metal ceramic crowns for reconstruction of destroyed crowns.

#### Clinical examination

Periodontal examination was carried out by an assessment of plaque index (PLI)<sup>17</sup> and (GI)<sup>18</sup> and clinical attachment loss (CAL) for clinical evaluation of gingival inflammation intensity and assessed also the relation existing between



Fig 1 Clinical Examination

the gingiva and the margins of dental reconstructions in the current study also to evaluate the attachment level and maintaining of normal insertion or migration of the epithelial junction, furthermore if there are recession or periodontal pockets due to iatrogenic factors (Figure 1).

#### Cytological study

Cytological examination was done for evaluating the cytotoxic effects of composite fillings and metallic crowns on gingival cells and periodontal status. The Cytological smears, then sent into the biology lab, College of Sciences, King Khalid University (Figure 2 A&B).

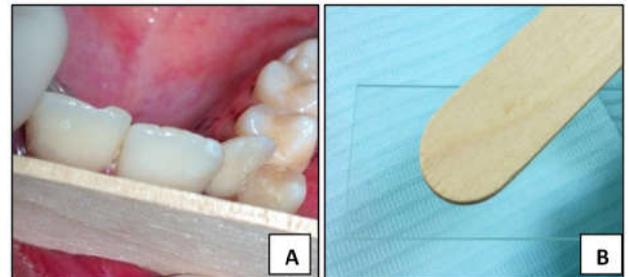


Fig 2 A& B Cytological gingival sample

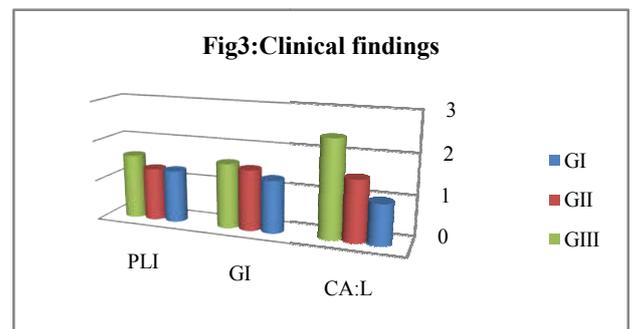
**Statistical analysis:** The data were collected and analyzed by ANOVA test.

### RESULTS

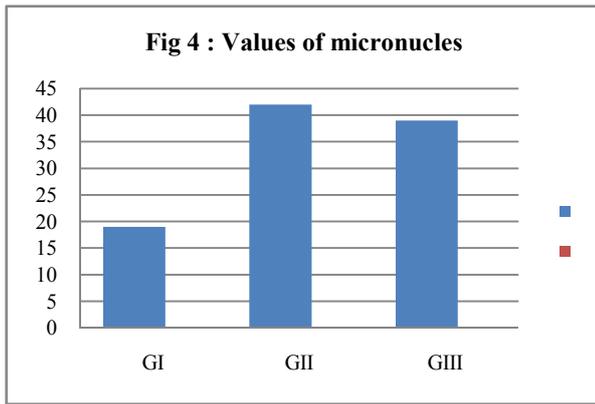
The table and figure 3 of clinical periodontal examination revealed significant differences between group I, II and III ( $p < 0.05$ ) where there were harmful effects of metallic ceramic crowns on periodontal tissues compared to control group (group I) whereas the destructive effects on periodontal tissue was more in group II compared to group I and group III in the present study.

Table of clinical Findings

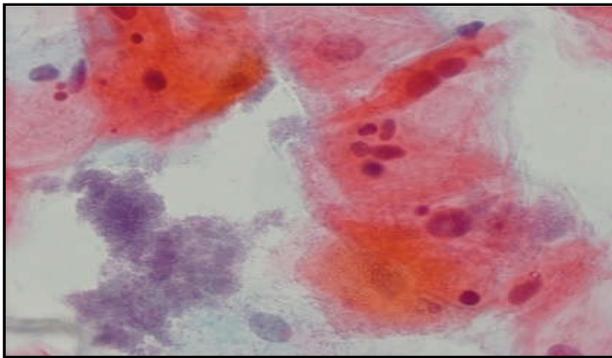
Group study	PLI	GI	CAL
GI	1.3±0,6	1.3±0,6	1.0±0,2
GII	1,3±0,6	1.5±0,5	1.5±0,1
GIII	1.6±0,6	1.6±0,6	2.4±0,2



The microscopic investigation displayed that there was no significant increase of apoptosis and necrosis in the study groups compared to control group, but, the results of the current study showed that (Composite filling) group had the highest value of micronucleus (MN) compared to metallic ceramic crowns group and the control group (Figure4). The microscopic investigation of Micronuclei also showed a variation in their shapes and number per cell.



The micronucleus type (M1) was found in all groups, while, type (M2) in metal ceramic crowns and (M3) micronucleus were not found in any group (Figure 5). Where the deformed nucleus (irregular in shape and has loops) and necrotic cells were not detected in any group.



**Fig 5** Micronuclei also showed a variation in their shapes and number per cell

## DISCUSSION

Iatrogenic factors in dentistry are the inappropriate diagnostic or dental therapy procedures by practitioners that resulted accidentally reverse effects on periodontal tissues, thus, the current study here is presented to evaluate the influence of some iatrogenic factors on a cytomorphometric analysis of gingival cells and periodontal status among diabetic patients. In general, dental restorations impact on periodontal tissues due to increased dental plaque accumulations on gingival margins and changes of the composition of the subgingival microbiota with an increase in periodontal pathogens<sup>19</sup> this in agreement with the cross sectional study of by van Dijken *et al.* where they found that the microbiological alterations occurring in the subgingival flora with different materials used in dentistry<sup>20</sup>

In the study that was done by Larato on composite resin restorations adjacent to subgingival region and also in the similar study of Hammer & Hotz where there were gingival inflammation in gingival tissues adjacent composite resin restorations more than metal restorations<sup>21,22</sup> furthermore, in another study that carried out by Willershausen *et al* there were increased gingival bleeding and probing depth of resin-based restorations, as compared with other restorative materials<sup>23</sup> these findings in agreement of our study where there were increases in GI and CAL in group II compared to control group (group I) but there were no significance differences in PLI between GI and GII.

The inappropriate prosthodontic treatment assists the appearance of arrogance pathological changes in periodontal tissues based on the roughness, surface status of the crown margins that participate in plaque accumulation and gingival tissue inflammation<sup>24</sup>. On the other hand, in the experimental study on patients were followed up 5 years after part of fixed prosthesis there were increases in pocket depth more than the control group<sup>25</sup> and clinical attachment loss was reported in other longitudinal studies that extended from 1 to 15 years where the meaning of attachment loss was from 0.1 to 1.3 mm during this study<sup>25,26,27</sup> these findings similar the results of the current study where there were increases in PLI, GI, and CAL in GIII compared GI. On the other hand the GI of GII and GIII more than GI but it was more in GIII than GII furthermore, there were significance differences in CAL in our study where it was more in GIII and GII than GI but it was more in GIII than GII.

In the comparison with the cytological samples of metal ceramic crowns of the current study there were percentage of apoptotic cells caused by composite filling and this is in agreement with Samuelsen *et al*<sup>28</sup> and Schweikl *et al*<sup>29</sup>. This effect may be due to an increase the duration of composite resin fillings (more than 1 week) that induced an elevation of the monomer transformation rate based on the chemical nature of these materials and also there was an increase in the dimethacrylates cytotoxicity after more than (72hr) where the treated cell cultures could not able to modify their form that were destroyed cells<sup>29</sup>. Furthermore, there were a toxic product generation like methacrylic acid (MA) due to the effect of enzymatic hydrolysis (by esterases) on composite resins<sup>30,31</sup>. which may be oxidized to form formaldehyde as a secondary product<sup>32</sup> consequently the biodegradation of resin may play important role in changes of the oral environment<sup>31,33</sup> On the other hand and in the current study there were no significant differences in increase of apoptosis and necrosis in the study groups compared to control group, but there were increase in value of micronucleus (MN) in group II compared to group I and group III, furthermore, the microscopic investigation of Micronuclei revealed differences in shapes and number of cells in group II compared to group I and group III where the micronucleus type (M1) was found in all groups, while, type (M2) in metal ceramic crowns and (M3) micronucleus were not found in any group. Finally, we did not find any nucleus abnormal and necrotic cells in all study groups.

## CONCLUSION

The results of a current study revealed that there were significant induction of micronucleus formulation in all groups compared to the control group, this induction and variation may due to the effect of composite resin filling materials, and it may need further studies to confirm these results.

## Acknowledgment

The authors would like to thank, the technicians in the lab of biology, College of sciences, King Khalid University, Abha, Kingdom of Saudi Arabia who investigated the cytological samples of this study.

## Reference

1. Bayne SC, Thompson GY (2006). Biomaterials. In: Roberson TM, Heyman HO, Swift EJ, editors. Sturdevant's art and science of operative dentistry. 5th ed. St. Louis: Mosby, 135-242.
2. Walker DM (2004). *Oral mucosal immunology: an overview*, Ann Acad Med Singapore, 33(4):27-30.
3. Schmid-Schwab M, Franz A, König F, Bristela M, Lucas T, Piehslinger E, Watts DC, Schedle A (2009), *Cytotoxicity of four categories of dental cements*, Dent Mater, 25(3):360-368.
4. Darmani H, Al-Hiyasat AS, Milhem MM (2007), *Cytotoxicity of dental composites and their leached components*, Quintessence Int,38(9):789-795.
5. Kanca J (1990). 3 rd. Pulpal studies: Biocompatibility or effectiveness of marginal seal? Quintessence Int21:775-9. Review
6. Palin WM, Fleming GJ, Burke FJ, Marquis PM, Randall RC (2005). The influence of short and medium-term water immersion on the hydrolytic stability of novel low-shrink dental composites. Dent Mater,21:852-63
7. Schweikl H, Schmalz G, Weinmann W (2004). The induction of gene mutations and micronuclei by oxiranes and siloranes in mammalian cells in vitro. J Dent Res, 83:17-21.
8. Rosenstiel SF, Land MF, Fugimoto J (2001). Contemporary fixed prosthodontics. 3rd ed. St. Louis: Mosby.
9. Prestipino V, Ingber A (1993). Esthetic high-strength implant abutments. Part I. J Esthet Dent, 5: 29-36.
10. Prestipino V, Ingber A (1993). Esthetic high-strength implant abutments. Part II. J Esthet Dent, 5: 63-68.
11. Barnfather KD, Brunton PA (2007). Restoration of upper dental arch using Lava all-ceramic crown and bridgework.Br. Dent J, 202(12):731-35.
12. Quirynen, M., Marechal, M., Busscher, H. J., Weerkamp, A.H., Darius, P. L. and Steenberghe, D. van. (1990). the influence of surface-free energy and surface roughness on early plaque formation. An in vivo study in man. J Clin Periodontol 17, 138-44.
13. Macleod J (1984). English language book society. 14th ed. Philadelphia (US): Churchill Livingstone; Davidson's principles and practice of medicine.
14. Firatli E, Yilmaz O, Onan V (1996). The relationship between clinical attachment loss and the duration of insulin dependent diabetes mellitus (IDDM) in children and adolescents. J Clin Periodontol, 23:362-6.
15. Shah SN, Tripathy BB (1994). API textbook of medicine. In: Sainani GS, Anand MP, Billimoria AR, Chugh KS, Joshi VR, Mehta PJ, editors. 15th ed. Bombay: Association of Physicians of India.
16. Grossi SG, Skrepanski FB, Caro TD, Zambon JJ, Cummins D, Genco RJ (1996). Response to periodontal therapy in diabetics and smokers. J periodontal, 67: 1094-102.
17. Silness J, Loe H (1964). Periodontal disease in pregnancy. II. Correlation between oral hygiene and periodontal condtion. Acta Odontol Scand, 22:121-35.
18. Loe H, Silness J (1963). Periodontal disease in pregnancy. I. Prevalence and severity. Acta Odontol Scand, 21:533-51.
19. Lang NP, Kiel RA (1963), Anderhalden K. Clinical and microbiological effects of subgingival restorations with overhanging or clinically perfect margins. J Clin Periodontol, 10: 563-78.
20. Van Dijken JW, Sjöström S (1987), Wing K. The effect of different types of composite resin fillings on marginal gingiva. J Clin Periodontol, 14: 185-9.
21. Larato DC (1972). Influence of a composite resin restoration on the gingiva. J Prosthet Dent, 28: 402-4.
22. Hammer B, Hotz P (1979). Nachkontrolle von 1 bis 5 jährigen amalgamkomposit- und gold full ungen. Schweiz Monatsschr Zahnheilkd, 89: 301-14.
23. Willershhausen B, Kottgen C, Ernst CP (2001), the influence of restorative materials on marginal gingival. Eur J Med R, 6(10): 433-9.
24. Tadumadze L (2005). Influence of the prosthodontic construction on the marginal gingiva nearby dental crowns and bridges. Georgian Med News, (126):31-3.
25. Valderhaugh J, Birkeland JM (1976). Periodontal conditions in patients 5 years following insertion offixed prostheses. Pocket depth & loss ofattachment. J Oral Rehabil, 3: 237-43.
26. Valderhaug J (1980). Periodontal conditions & carious lesions following the insertion of fixed prostheses: a 1 0-year followup study. Int Dent J, 30: 296-304.
27. Valderhaug J, Ellingsen JE, Jokstad A (1993). Oral hygiene, periodontal conditions & carious lesions in patients treated with dental bridges. A 1 5-year clinical & radiographic followup study. J Clin Periodontal, 20: 482-9.
28. Samuelsen JT, Holme JA, Becher R, Karlsson S, Morisbak E, Dahl JE (2008). HEMA reduces cell proliferation and induces apoptosis in vitro. Dent. Mater, 84(1): 134-40.
29. Schweikl H, Hiller KA, Bolay C, Kreissl M, Kreismann, Nusser A, Steinhauser S, wieczorek J, Vasold, Schmalz G (2005). Cytotoxic and mutagenic effects of dental composite materials.Biomaterials, 26: 1713-1719.
30. Munksgaard EC, Freund M (1990). Enzymatic hydrolysis of (di)methacrylates and their polymers. Scand. J. Dent. Res, 98: 261-7.
31. Yourtee DM, Smith RE, Russo KA, Burmaster S, Cannon JM, Eick JD, Kostoryz EL (2001). The stability of methacrylate biomaterials when enzyme challenged: kinetic and systematic evaluations. J. Biomed. Mater. Res, 57: 522-31.
32. Øysæd H, Ruyter IE, Sjøvik-Kleven IJ (1988). Release of formaldehyde from dental composites. J. Dent. Res, 67: 1289-94.
33. Geurtsen W, Leyhausen G (2001). Chemical-biological interactions of the resin monomer triethyleneglycol-dimethacrylat (TEGDMA). J. Dent. Res, 80: 2046-50.

\*\*\*\*\*