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Correlation between Radiomorphometric Indexes and Low Bone Quality in the Success of Osseointegration in oral Rehabilitation

By Gabriela dos Santos Gonçães Lima, Angelinna Zanesco, DDS, PhD,
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Abstract- Introduction: Oral rehabilitation through implants has grown significantly in recent years, and with them also come the problems related to implant dentistry, the most important of which is the impossibility of bone tissue to establish osseointegration.

Objective: The aim of this study was to correlate radiomorphometric indices and poor bone quality in osseointegration failures in oral rehabilitation.

Material and Method: 104 missing implants were evaluated in 74 individuals, verifying in the panoramic radiographs the radiomorphometric indices Mental (IM) and the Mandibular Cortical (ICM).

Keywords: low bone quality, radiomorphometric indexes, implants.

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Correlation between Radiomorphometric Indexes and Low Bone Quality in the Success of Osseointegration in oral Rehabilitation

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Objective: The aim of this study was to correlate radiomorphometric indices and poor bone quality in osseointegration failures in oral rehabilitation.

Material and Method: 104 missing implants were evaluated in 74 individuals, verifying in the panoramic radiographs the radiomorphometric indices Mental (IM) and the Mandibular Cortical (ICM).

Results: It was possible to evaluate the correlation (-0.721) for $p < 0.001$ inverse correlation between the MI and the MCI. The lower the value found in the MI, the worse the bone quality evaluated in the MCI. It was also possible to verify the relationship (0.275) to $p < 0.001$ between MI and the arch where it was evaluated that bone quality in the upper arch showed worse quality when compared to the lower arch. It there was correlation to $p < 0.001$ the length of the lost implant and the region (-0.339), smaller length implants were lost in the posterior. Age was correlated (0.198) with the MCI to $p < 0.05$ where the older the age the worse the bone quality evaluated. The MCI correlated with the arch (-0.235) for $p < 0.05$ a worse bone quality when correlated with the implants lost in the upper arch.

Conclusion: Radiomorphometric indices can be used in the preoperative evaluation to assist in the detection of patients with poor bone quality.

Keywords: low bone quality, radiomorphometric indexes, implants.

1. INTRODUCTION

Oral rehabilitation through implants has been growing significantly in recent years and reaching high rates of clinical success. Currently, bone-implant contact is considered predictable, safe, and lasting¹. The long-term success of

dental implants depends on osseointegration². With the advancement of success also comes the problems related to implant dentistry³.

Failures can be classified as biological, mechanical or iatrogenic or due to insufficient patient adaptation⁴.

Among these faults, the most dangerous is the biological fault, which can be defined as the impossibility of bone tissue to establish osseointegration^{4,5}. This lack of osseointegration that requires implant removal is considered a biological failure⁶.

Biological faults are classified as primary and secondary. If the failure occurs during the osseointegration process, it is considered a primary failure; if it occurs after charging it is a minor fault^{3,7}. Primary failures result from lack of bone repair, where the connection between the implant surface and the bone does not occur. Instead, fibrous tissue forms between the implant and bone, causing the implant to lose its stability^{5,7,8}.

Failures can be prevented by appropriate patient selection; proper treatment planning is critical to successful implant dentistry⁹.

Some information has been reported on factors that influence implant osseointegration, such as: biocompatibility, implant design, surface conditions, surgical site, a surgical technique for implant installation, and loads applied to them¹⁰.

Studies relate early loss in short-length implants in posterior regions where space and volume are insufficient¹¹.

To acquire adequate healing conditions, the implant, after insertion, must exhibit good primary stability, which may correspond to the clinical manifestation of osseointegration. Primary stability is primarily determined by factors related to bone biomechanical properties, implant design, and surgical technique; while secondary stability is also determined by bone tissue response to surgical trauma and implant surface¹². Primary stability is achieved when the implant locks into the apical or marginal portion of the site due

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a sufficient amount of compact bone or cancellous bone¹³.

The world population has been aging over the years and inevitably, the increase in individuals with low bone density¹⁴. Several studies have shown a positive correlation between low bone density and implant loss¹⁵⁻¹⁸. The primary stability of the implant, as well as its survival, is affected by low bone density^{15,16}. The high rate of implant loss is related to bone type IV when compared to bone type I, II, III¹⁹.

The aim of this study was to correlate radiomorphometric indices and poor bone quality in osseointegration failure in oral rehabilitation.

II. MATERIAL AND METHOD

The project was submitted to the Research Ethics Committee (CEP) of the Metropolitan University of Santos and approved. It was used the database of the postgraduate course in implantology of the Metropolitan University of Santos, the data of 104 lost implants of various diameters in 74 individuals aged 33 - 84 years of both genders with an average of 59 years, in addition to their images and clinical records. Individuals with a

history of hormone replacement therapy or calcium therapy under the age of six months, and those who did not have all the tests necessary for this study were excluded.

The implants lost in the upper and lower arch in the anterior and posterior regions were separated, also grouped by diameter and length.

All digital panoramic radiographs images used in this study are from a partner institute of the Metropolitan University of Santos and performed by the ORTHOPHOS XG 3D PAN/TELE/TOMO device (Sirona Dental Systems GmbH, Bensheim, Germany) following the same protocol of acquisition: The Panoramic Radiography 69 kV, 15 mA, and exposure time 14.1 s.

In panoramic radiographs, radiomorphometric indices were evaluated. Among, them the mandibular cortical index (MCI)²⁰ a bilateral evaluation with results established in: C1 - clear and sharp posterior mandibular cortical, C2 - the endosteal surface presents semilunar defects (lacunar resorption), or the surface presents cortical residues, C3 - a cortical layer is extremely porous (Figure 1).

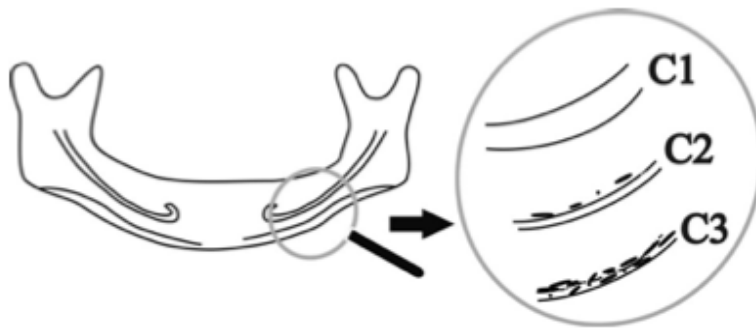


Figure 1: Illustration of the Mandibular Cortical Index

Mental index (MI)²¹ bilateral assessment determined by the width of the mandibular cortex, measured on the line perpendicular to the base of the

mandible, at the height of the center of the mental foramen (Figure 2).

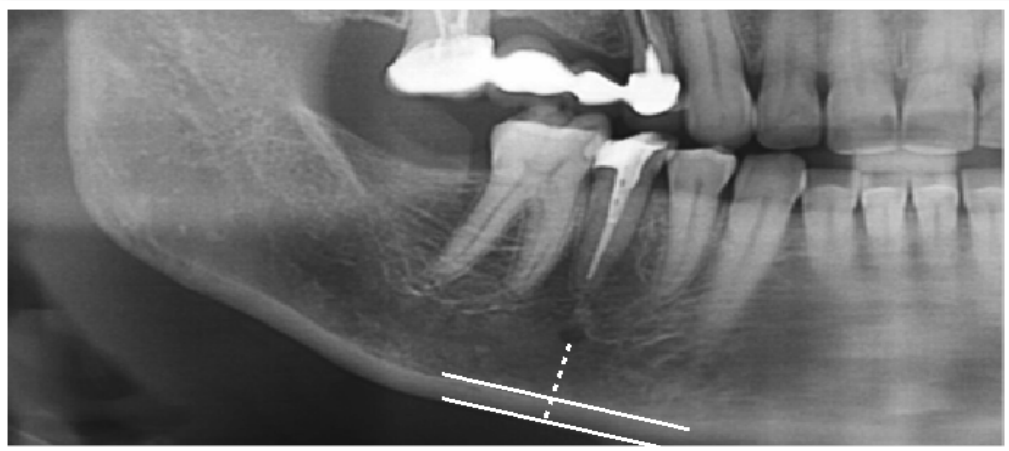


Figure 2: Mental Index Illustration

III. RESULTS

We evaluated data from 104 implants of various diameters in 74 individuals, 48 women and 26 men, aged 33 - 84 years with a mean of 59.4 years. As for the upper and lower arches, 46 implants lost in the lower

arch and 58 in the upper arch. We evaluated 76 implants in the posterior region and 28 in the anterior region. Spearman's correlation coefficient test was performed to assess the relationship between the variables of the lost implants (Table 1).

Table 1: Spearman correlation coefficient

	Age	Sex	MI	MCI	ARCADE	Region	Diameter	Lenght
Age								
Sex	R:0,209* P:0,034							
MI	R:-0,106 P:0,282	R:-0,038 P:0,701						
MCI	R:0,198* P:0,044	R:-0,021 P:0,833	R:-0,721** P:0,000					
ARCADE	R:0,214* P:0,029	R:0,052 P:0,597	R:0,275** P:0,005	R:-0,249* P:0,011				
Region	R:0,116 P:0,242	R:0,158 P:0,110	R:0,054 P:0,587	R:0,035 P:0,724	R:0,235* P:0,016			
Diameter	R:-0,028 P:0,779	R:-0,128 P:0,196	R:-0,05 P:0,616	R:0,044 P:0,655	R:0,038 P:0,700	R:0,072 P:0,467		
Lenght	R:-0,032 P:0,751	R:-0,040 P:0,889	R:-0,060 P:0,547	R:0,104 P:0,292	R:-0,241* P:0,014	R:-0,339** P:0,000	R:0,050 P:0,615	

* $p < 0,05$ and ** $p < 0,001$

According to the result of the correlation coefficient, it was possible to verify the correlation (-0.721) for $p < 0.001$ inverse relationship between the MI and the MCI. The ratio (0.275) to $p < 0.001$ between MI and the arch where bone quality was evaluated in the upper arch showed worse quality when compared to the lower arch.

It was also correlated (-0.339) to $p < 0.001$ the length of the lost implant and the region, smaller length implants were lost in the posterior region.

Age was correlated (0.198) with the MCI to $p < 0.05$ where the older, the age the worse, the bone quality evaluated.

The MCI correlated with the arch (-0.235) for $p < 0.05$ a worse bone quality when correlated with the implants lost in the upper arch.

IV. DISCUSSION

The use of radiomorphometric indices to assess bone quality is an effective and inexpensive instrument for this detection since the evaluation test is difficult to access and high for the population.

Several studies corroborate our results in the high correlation between radiomorphometric indices evaluated in panoramic radiographs, where these exams may be predictive in helping to assess bone quality²²⁻²⁴.

Some studies contradict these findings showing that there is inconsistency in the data obtained when in smaller groups, although the mandibular cortical index

is among the most reproducible radiomorphometric indices^{25,26}.

The influence of age with the loss of bone quality was evidenced in this study. Older individuals had worse bone quality when evaluated by the mandibular cortical index. This result was also found in other studies. Zlataric et al. 2002, verified the values of radiomorphometric indices in elderly individuals and showed that the values of these indices decreased in both sexes up to 78 years²⁷. Edgerton et al. 1999 evaluated British women and observed that radiomorphometric indices gradually decreased with increasing age²⁸.

When the arches were evaluated, the upper arch showed worse bone quality compared to the lower arch, implants installed in the posterior regions of the upper arch showed a higher loss rate when compared to the other regions.

It was evidenced in this study when the radiomorphometric indices IM and ICM were evaluated and correlated with the length of the implants and the region where the highest concentration of loss occurred in the posterior maxilla and implants of shorter lengths. These data were also evidenced by other studies where the incidence of loss of short posterior maxillary implants was higher when compared, for example, with the mandible²⁹⁻³¹.

Contrary to the results obtained in this study, implants installed in the posterior maxilla of short length and diameter presented excellent fixation results³².

V. CONCLUSION

According to the results of this study, radiomorphometric indices may be used the preoperative evaluation to assist in the detection of patients with poor bone quality.

BIBLIOGRAPHIC REFERENCES

1. Brånemark P I, Zarb G A, Albrektsson T. Tissue integrated prosthesis: osseointegration in clinical dentistry. Chicago: Quintessence; 1985.
2. Matos GRM. Tratamento de superfície de implantes dentários e osseointegração. Rev Dental Press Periodontia Implantol. 2008 out-dez ; 2(4): 40-50.
3. Givol N, Taicher S, Halamish-Shani T, Chaushu G. Risk management aspects of implant dentistry. Int J Oral Maxillofac Implants. 2002; 17(2): 258-62.
4. Buser D, Mericske-Stern R, Bernard JP, et al. Long-term evaluation of non-submerged ITI implants. Part 1: 8-year life table analysis of a prospective multicenter study with 2359 implants. Clin Oral Implants Res. 1997;8:161-172.
5. Esposito M, Hirsch J M, Lekholm U, et al. Biological factors contributing to failures of osseointegrated oral implants. (I). Success criteria and epidemiology. Eur J Oral Sci. 1998; 106: 527-551.
6. Misch C E, Perel M L, Wang H L, et al. Implant success, survival, and failure: The International Congress of Oral Implantologists (ICOI) Pisa Consensus Conference. Implant Dent. 2008; 17: 5-15.
7. Alsaadi G, Quirynen M, Michiles K, et al. Impact of local and systemic factors on the incidence of failures up to abutment connection with modified surface oral im- plants. J Clin Periodontol. 2008; 35: 51-57.
8. Esposito M, Thomsen P, Ericson L E, et al. Histopathologic observations on early oral implant failures. Int J Oral Max- illofac Implants. 1999; 14: 798-810.
9. Smith R S. Long-term complications of osseointegrated implants. In: Kaban B L, Progel M A, Perrot H D, editors. Complications in oral and maxillofacial surgery. Philadelphia: Saunders; 1997.
10. Albrektsson T, Zarb G, Worthington P, Eriksson A R. The longterm efficacy of currently used dental implants: a review and proposed criteria of success. Int J Oral Maxillofac Implants. 1986 summer; 1(1): 11-25.
11. Baqain Z H, Moqbel W Y, Sawair F A. Early dental implant failure: risk factors. Br J Oral Maxillofac Surg. 2012 Apr; 50(3): 239-43.
12. Sennerby L, Miyamoto I. Insertion torque and RFA analysis of Tiunite and SLA implants. A study in the rabbit. Appl Osseointegration Res. 2000; 1(1): 31-3.
13. Ashley E T, Covington L L, Bishop B G, Breault L G. Ailing and failing endosseous dental implants: a literature review. J Contemp Dent Pract. 2003 May 15; 4(2): 35-50.
14. Reginster J Y, Burlet N. Osteoporosis: a still increasing prevalence. Bone 2006; 38: S4-9.
15. Åstrand P, Billström C, Feldmann H, Fischer K, Henricsson V, Johansson B, et al. Tapered implants in jaws with soft bone quality: a clinical and radiographie 1-year study of the Brånemark System Mark IV fixture. Clin Implant Dent Relat Res 2003; 5: 213-8.
16. Busenlechner D, Fürhauser R, Haas R, Watzek G, Mailath G, Pommer B. Long-term implant success at the Academy for Oral Implantology: 8-year follow-up and risk factor analysis. J Periodontal Implant Sci 2014; 44: 102-8.
17. Niedermeier R, Stelzle F, Riemann M, Bolz W, Schuh P, Wachtel H. Implant- supported immediately loaded fixed full-arch dentures: evaluation of implant survival rates in a case cohort of up to 7 years. Clin Implant Dent Relat Res 2017; 19: 4-19.
18. Becker W, Hujoel P P, Becker B E, Willingham H. Osteoporosis and implant failure: an exploratory case control study. J Periodontol 2000; 71: 625-31
19. Lekholm U, Zarb G A. Patient selection and preparation. Tissue integrated prostheses: osseointegration in clinical dentistry. Chicago: Quintessence Pub Co; 1985. p. 199-209.
20. Klemetti E, Kolmakov S; Kröger H. Pantomography in assessment of the osteoporosis risk group. Scand. J. Dent. Res., 1994, Copenhagen, v.102, p. 68-72.
21. Taguchi A, Tanimoto K, Sueti Y, Wada T. Tooth loss and mandibular osteopenia. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 1995 Jan; 79(1): 127-32.
22. Miliuniene E, Alekna V, Peculiene V, Tamulaitiene M. Evaluation of bone mineral density in postmenopausal women with alterations of the mandible cortical bone. Stomatologija. 2016; 18(3): 86-91.
23. Dutra V, Yang J, Devlin H, Susin C. Radiomorphometric indices and their relation to gender, age, and dental status. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 2005; 99: 479-84.
24. Drozdowska B, Pluskiewicz W, Tarnawska B. Panoramic-based mandibular indices in relation to mandibular bone mineral density and skeletal status assessed by dual energy X-ray absorptiometry and quantitative ultrasound. Dentomaxillofac Radiol 2002; 31: 361-7.
25. Mansour S, AlGhamdi A S, Javed F, Marzouk H, Khan E A. Panoramic radiomorphometric indices as reliable parameters in predicting osteoporosis. Am J Med Sci. 2013 Dec; 346(6): 473-8.

26. Akay G, Akarslan Z, Karadag O, Gungor K. Does tooth loss in the mandibular posterior region have an effect on the mental index and panoramic mandibular index? *Eur Oral Res.* 2019 May; 53(2): 56-61.
27. Zlataric' D K, Celebic' A, Kobler P. Relationship between body mass index and local quality of mandibular bone structure in elderly individuals. *J Gerontol A Biol Sci Med Sci* 2002; 57: M588- 93.
28. Ledgerton D, Horner K, Devlin H, Worthington H. Radiomorphometric indices of the mandible in a British female population. *Dentomaxillofac Radiol* 1999; 28: 173-81.
29. Olate S, Lyrio M C, de Moraes M, Mazzonetto R, Moreira R W. Influence of diameter and length of implant on early dental implant failure. *J Oral Maxillofac Surg.* 2010 Feb; 68(2): 414-9.
30. Machtei E E, Mahler D, Oettinger-Barak O, Zuabi O, Horwitz J. Dental implants placed in previously failed sites: survival rate and factors affecting the outcome. *Clin Oral Implants Res.* 2008 Mar; 19(3): 259-64.
31. Alsaadi, G. Quirynen, M. & Van Steenberghe, D. (2006) The importance of implant surface characteristics in the replacement of failed implants. *The International Journal of Oral & Maxillofacial Implants* 21: 270–274.
32. Moriwaki H, Yamaguchi S, Nakano T, Yamanishi Y, Imazato S, Yatani H. Influence of Implant Length and Diameter, Bicortical Anchorage, and Sinus Augmentation on Bone Stress Distribution: Three-Dimensional Finite Element Analysis. *Int J Oral Maxillofac Implants.* 2016 Jul-Aug; 31(4): e84-91.



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Concentrated Growth Factor (CGF) in Alveolar Bone Grafting Procedures: Dilemma or a Reality? - A Detailed Review

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Concentrated Growth Factor (CGF) in Alveolar Bone Grafting Procedures: Dilemma or a Reality? - A Detailed Review

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& Dr. Madhav Naik P ^ω

Abstract- Alveolar bone grafting is considered to be one of the most challenging task in the management of cleft lip and palate patients as it is expected to bridge the gap between the alveolar segments and provide the solid base for the unerupted tooth/teeth to erupt through it, besides effectively healing the Oro-antral fistula if any passing through the cleft into the nasal cavity. Although the traditional approach is to use autogenous bone graft from iliac crest, which has its own side-effects like pain, morbidity, additional surgery, etc. However with the advent of Tissue engineering technique, it has become possible to overcome most of these challenges by using a tissue friendly yet innovative material by name CGF (Concentrated Growth Factor) which in reality is the latest version of platelet-rich plasma having in it abundant of growth factors thereby serving the purpose of a regenerative medicine, wherein three factors are important to stimulate the bone regenerative effect: 1) Scaffold 2) Growth factors and 3) Autologous cells. All these are found to be present in sufficient quantity in CGF which was first introduced by Sacco in 2006 followed by Corigliano in 2010.

Keywords: concentrated growth factors, platelet-rich plasma, autologous bone, alveolar bone grafting, osteogenic potential.

I. SEARCH STRATEGY

A secondary research based upon the library referencing from the year 2000 to 2018 through online search using key words as concentrated growth factors, platelets, wound healing, osteogenic potential, platelet-rich plasma, alveolar bone grafting, etc.

II. INTRODUCTION

Concentrated growth factor is considered to be an updated and latest version of platelet-rich plasma (PRP) found to be exhibiting remarkable wound healing potential in open surgical wound sites, where its application has proven to give both soft tissue and hard tissue healing potential by virtue of its wound healing

and osteogenic potential. Such desirable clinical and biotechnological potential of this group of platelets is found to be of immense use in different craniofacial surgical procedures including alveolar bone grafting in cleft lip and palate individuals wherein the golden standard of treatment is to use cortical bone from the iliac crest. Although the conventional method has its good bone induction and conduction property to bridge the gap in the cleft-alveolar segments, yet in most of the cases would result in several side-effects like surgical site morbidity and altered gait of the individuals post-surgery creating a new surgical site to treat the existing one.

III. HISTORY AND BACKGROUND

Bone defects involving the Oral and maxillofacial anatomical regions have been treated for long time by using variable platelet-concentrates, which differ from each other by their preparation methods to induce and accelerate the bone formation when mixed with bone grafts and barrier membranes. These concentrates are found to be abundantly rich in growth factors which are in reality the "PROTEINS" involved in regulating the complex process of wound healing. Since the growth factors are located in the blood plasma and platelets, these platelet-concentrates are commonly used to accelerate the tissue repair and regeneration of clefts of the alveolar region. Platelet's regenerative potentiality was introduced in the year 1974 by Rose et al who described the presence of growth factors in the blood-plasma.¹ It was also reported that based upon their methods of preparations and evolution of their clinical applications periodically, the platelet-concentrates have been grouped into various generations.

a) First-generation of platelets

Includes PRP (Platelet-rich plasma) and PRGF (Platelet-rich in growth factor) which were introduced by Marx and Anitua respectively. The method of preparation involved addition of calcium and bovine thrombin to achieve stability which in turn made it technique sensitive for all the clinical applications.

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b) *Second-generation of platelets*

Involved PRF (Platelet-rich fibrin) introduced by Choukroun. The method of preparation involved using venous blood from the patient's veins and centrifuging the same to produce a fibrin-rich gel with aggregated platelets obtained in the middle of the tube, just between the red corpuscles at the bottom and platelet poor plasma at the top.

c) *Preparation of CGF*

Its method of preparation is similar to PRF but the centrifugation speed differs. Its preparation uses 2400 rpm to 2700 rpm to separate all the cells in the venous blood. As a result a block is obtained which is rich in fibrin having a thicker, larger and denser amount of growth factors compared to PRF. This makes CGF better in bone regenerative capacity improving its versatility.³⁻⁵ After centrifugation the CGF was obtained in three parts, the upper white Part (PPP), the lower red Part (RBC) and the middle "buffy coat" Part (interface between white and red part). The results showed that platelets and leukocytes were localized in the buffy coat, whereas the erythrocytes were present only in the red part of CGF. The in vitro cumulative release of growth factors revealed the presence of PDGF-AB, TGF- β 1 and IGF-1 having a constant kinetic release, reaching the maximum accumulation at day 3rd and 6th respectively. VEGF and BMP-2 had a slow kinetic release reaching the maximum accumulation at day 8th. TNF α and BDNF had a faster kinetic release reaching the maximum accumulation at day 1st and 3rd respectively. These findings have largely supported the clinical use of CGF for its enhanced healing and osteogenic-effects.^{6,7} According to Professor Rodella⁸ at University of Brescia-Department of Biomedical Sciences and Biotechnologies, CGF shows higher tensile strength, more growth factors, higher viscosity and higher adhesive strength than PRF. Therefore it is recommended that surgeons can use the CGF as a barrier membrane to accelerate soft tissue healing or be mixed with bone graft to accelerate new bone formation. CGF doesn't require any chemical or allergenic additives such as bovine thrombin or anticoagulants into it. Therefore it is usually free from any viral disease transmission.⁹⁻¹¹

IV. MATERIALS AND METHODS

A secondary research based on hand – search and online search using the key terms was done for the articles from 2000 to March 2018 involving discussion about the platelets (CGF) and their osteogenic capabilities.

a) *Inclusion criteria*

All original researches, reviews, case reports on the POSITIVE outcome of the osteogenic properties of the CGF were included. Studies that dealt with craniofacial surgery for total or partial reconstruction of

mandibles/maxilla, cleft lip and palate surgeries, distraction osteogenesis and Osseo-integration were also added in the search strategies.

b) *Exclusion criteria*

Studies involving gene therapy, etc.

c) *Research Dilemma*

Therefore the need of the hour is to use this material which can help in efficient wound healing and also provide osteogenic impulse so that second surgical site can be avoided which would be beneficial to the patient in terms of pain, morbidity and cost.

d) *Advantages of CGF*

It is accepted that CGF is capable of better regenerative-capacity and higher versatility. The fibrin block is thick and dense with a very high concentration of fibrinogen, factor XIII and thrombin. The activated form of XIII (XIIIa) cross links the fibrin clot to increase the stability, strength, and protection against plasmin mediated degradation. Mixing this combination with autogenous or other fillers makes it user-friendly material to handle and fill even the larger bone defects improving the overall bone regenerative effect.¹² Sohn in 2009³ reported that CGF has multipurpose use associated to its higher levels of active proteins which can support healing, growth and cell morphogenesis. CGF provides sufficient stability to implant- root interface during healing process providing necessary bone formation around the implant which would be capable of absorbing the functional load during healing. With the help of Resonance Frequency-Analysis (RFA) which is considered to be very important tool for assessing the Osseo-integration process and also tracks the changes in the stability not only during the implant placement but also during the healing and later periods, the active role played by CGF and other growth factors containing products during Osseo-integration and bone healing was reported. In a sum up of all its advantages, CGF appear to show superior potential for tissue-regeneration in the clinical and biotechnological applications including sinus and alveolar ridge augmentation.¹³⁻¹⁹

e) *Possible mode of action by CGF*

The various research studies have shown that the improved regenerative effect of CGF when used to treat the bone defect could be attributed to its biologic constituents which are active in presence of chemotactic and mitogenic PDGF (Platelet Derived Growth Factor) which are basically involved in tissue regeneration process. Unlike PRP, CGF does not dissolve rapidly following its application on the surgical site. However it was found that the thick fibrin-matrix is remodeled slowly like a natural blood-clot. Thus prolonging the duration of growth factors activity which in turn is found to be effective for growth factor synergy, enhancing the cell proliferation and osteogenic differentiation.^{20,21}

V. DISCUSSION

CGF, therefore is considered as the latest progeny of the platelets containing the abundant number of growth factors, leucocytes and fibrin-matrix in it. The growth factors released are responsible for aiding in tissue regeneration (both hard and soft tissue) which is the reason that it can provide immediate wound healing and repair of the surgical site by outgrowing the number of growth factors.²²⁻²⁷ The greater advantage of this latest PRP is in it being an AUTOGENOUS preparation, which not only improves the tissue healing but also enhances the clinical outcome of the surgical procedure at a considerable short period of time.²⁸⁻³⁰ It is also reported that the presence of growth factors in CGF are considered to be a class of natural biological mediators that regulate the key cellular events in the tissue repair, including cell proliferation, differentiation and extracellular matrix synthesis. Release of biological factors responsible for tissue repair occurs through platelet activation and de-granulation. These factors are PDGF (Platelet-Derived Growth Factors), VEGF (vascular-endothelial growth factors), IGF (Insulin-Like Growth Factors), TGF (Tissue-Growth Factor), BDGF (Brain-Derived Growth Factor) and BMP (Bone-Morphogenetic Protein).³¹⁻³³ Recent studies have reported that when CGF is used alone or mixed with bone allograft increased the bone growth process by accelerating the healing of the soft tissue and facilitating periodontal ligament repair in both the animal and human studies. Furthermore, the findings suggested that CGF along with CD34 positive cells has played an effective role in vascular maintenance, neo-vascularization and angiogenesis when used at the surgical wound site. This was all attributed to the presence of the growth factors in CGF which in turn would enhance the lost or the damaged tissue regrowth at the local site of application.³⁴⁻³⁶ Although it is suggestive that the best bone grafts are loaded with the desirable qualities of Osteo-conduction, Osteo-induction and Osteo-genesis, it is the autogenous bone which is still considered the "gold standard" since it has all the three characteristics without any associated immune response. However it was found that CGF when used along with the autogenous bone could harness the osteogenic effect to achieve longer and better osteogenesis. This property is also found to be helpful in providing a complete root coverage for denuded roots in orthodontic cases as well.³⁷⁻³⁹ Additionally it is reported that use of CGF and other platelet- concentrates during bone grafting offers the advantages in terms of formation of a dense fibrin clot which primarily plays an important mechanical role, secondly it promotes cellular migration, particularly of endothelial cells which are necessary for neo-angiogenesis, vascularization and survival of the graft. Finally the presence of leukocytes and cytokines in the fibrin network play a significant role

in the self- regulation of inflammatory and infectious phenomenon within the grafted material.⁴⁰

VI. CONCLUSION

CGF with its well-known tissue-regeneration capability can be the next material of choice to be used for its beneficial wound healing and osteogenic property at the Cranio-maxillo-facial surgical wound site and therefore can be recommended to be used in augmenting the bone in alveolar bone grafting procedures alone or in combination with other graft materials for the best clinical outcome. Besides providing relief to the patient in terms of reduced wound healing time and an excellent opportunity to avoid another surgical procedure of harvesting bone from the iliac crest, its versatility and regeneration potentials should be given a fair chance to help the patients suffering from cleft lip and palate so that the burden of care giving is taken care by the clinicians. Although CGF has proven to be effective in Osseo-integration and bone healing further investigations and research at the clinical and histopathological levels should be encouraged to understand its role played at different levels for the benefit of mankind.

REFERENCES RÉFÉRENCES REFERENCIAS

1. Naik B, Karunakar P, Jayadev M, Marshal R V. Role of platelet rich fibrin in wound healing: A critical review. *Journal of Conservative Dentistry* 2013; 16(4): 284-93.
2. Sacco L. Lecture. International academy of Implant prosthesis and osteoconnection. 2006; 12: 4.
3. Sohn D S. The effect of concentrated growth factors on ridge augmentation. *Implant Journal*. 2009; 34-40.
4. Anitua E. The use of plasma-rich growth factors (PRGF) in oral surgery. *Pract Proced Aesthet Dent*. 2001; 13(6): 487-93.
5. J. Choukroun, F. Adda, C. Schoeffer, and A. Vervelle, "PRF: an opportunity in perio-implant tology," *Implantodontie* 2000; 42: 55-62.
6. Anitua et al. The potential impact of the preparation rich in growth factors (PRGF) in different medical fields. *Biomaterials*. 2007; 28: 4551.
7. Anitua et al. Autologous platelets as a source of proteins for healing and tissue regeneration. *Thromb Haemost*. 2004; 91: 4.
8. Rodella L F, Favero G, Boninsegna R et al. Growth factors, CD34 positive cells, and fibrin network analysis in concentrated growth factors fraction. *Microsc. Res. Tech*. 2011; 74(8): 772-777.
9. D. M. Dohan, J. Choukroun, A. Diss et al., "Platelet-rich fibrin (PRF): a second-generation platelet concentrate. Part II: platelet-related biologic features," *Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology and Endodontology*. 2006; 101(3): E45-E50,

10. J. Choukroun, A. Diss, A. Simonpieri et al., "Platelet-rich fibrin (PRF): a second-generation platelet concentrate. Part IV: clinical effects on tissue healing," *Oral Surgery, Oral Medicine, Oral Pathology, Oral Radiology and Endodontology*. 2006; 101(3): E56–E60, 2006.
11. Qi Li, Shuang Pan, Smit J. Dangaria, et al., "Platelet-Rich Fibrin Promotes Periodontal Regeneration and Enhances Alveolar Bone Augmentation," *BioMed Research International*, vol. 2013; Article ID 638043: 13 pages.
12. Upadhayayaa V, Arorab A, Goyal A. Bioactive Platelet Aggregates: Prp, Prgf, Prf, Cgf And Sticky Bone. *IOSR Journal of Dental and Medical Sciences*. 2017; 16(5) : PP 05-11.
13. Nurden A T, Nurden P, Sanchez M, Andia I, Anitua E. Platelets and wound healing. *Front Biosci*. 2008; 13: 3532–48.
14. Anitua E, Sanchez M, Zalduendo M M, de la Fuente M, Prado R, Orive G, et al. Fibroblastic response to treatment with different preparations rich in growth factors. *Cell Prolif*. 2009; 42(2): 162–70.
15. He L, Lin Y, Hu X, Zhang Y, Wu H. A comparative study of platelet-rich fibrin (PRF) and platelet-rich plasma (PRP) on the effect of proliferation and differentiation of rat osteoblasts in vitro. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod*. 2009; 108(5): 707–13.
16. Salvi G E, Lang N P. Diagnostic parameters for monitoring peri-implant conditions. *Int J Oral Maxillofac Implants*. 2004; 19(Suppl): 116–27.
17. Atsumi M, Park S H, Wang H L. Methods used to assess implant stability: current status. *Int J Oral Maxillofac Implants*. 2007; 22(5): 743–54.
18. Quesada-Garcia M P, Prados-Sanchez E, Olmedo-Gaya M V, Munoz-Soto E, Gonzalez-Rodriguez M P, Vallecillo-Capilla M. Measurement of dental implant stability by resonance frequency analysis: a review of the literature. *Med Oral Patol Oral Cir Bucal*. 2009; 14(10): e538–46.
19. Forabosco A, Gheno E, Spinato S, Garuti G, Forabosco E, Consolo U. Concentrated growth factors in maxillary sinus floor augmentation: A preliminary clinical comparative evaluation. *Int J Growth Factors Stem Cells Dent* 2018; 1: 2-7.
20. Gassling V L, Açil Y, Springer IN et al. Platelet-rich plasma and platelet-rich fibrin in human cell culture. *Oral Surg. Oral Med. Oral Pathol. Oral Radiol. Endod*. 2009; 108(1): 48–55.
21. Yu B, Wang Z. Effect of concentrated growth factors on beagle periodontal ligament stem cells in vitro. *Mol. Med. Rep*. 2014; 9(1): 235–242.
22. Anitua E, Sánchez M, Orive G, Andia. Delivering growth factors for therapeutics. *Trends Pharmacol Sci* 2008; 29: 37-41.
23. Dohan D M, Choukroun J, Diss A, Dohan S L, Dohan A J, et al. Platelet-rich fibrin (PRF): a second-generation platelet concentrate. Part I: technological concepts and evolution. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2006; 101: e37-e44.
24. Dohan D M, Choukroun J, Diss A, Dohan S L, Dohan A J, et al. Platelet-rich fibrin (PRF): a second-generation platelet concentrate. Part III: leucocyte activation: a new feature for platelet concentrates? *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2006; 101: e51-e55.
25. Choukron J. Advanced PRF and i-PRF: Platelet concentrates or blood concentrates? *J Periodont Med Clin Practice* 2014; 1: 3.
26. Anitua E, Alkhraisat M H, Orive G. Perspective and challenges in regenerative medicine using plasma rich in growth factors. *Journal of controlled release* 2012; 157: 29-38.
27. Sohn D S, Heo J U, Kwak D H, Kim D E, Kim J M, et al. Bone regeneration in the maxillary sinus using an autologous fibrin-rich block with concentrated growth factors alone. *Implant Dent* 2011; 20: 389-395.
28. Honda H, Tamai N, Naka N, Yoshikawa H, Myoui A. Bone tissue engineering with bone marrow-derived stromal cells integrated with concentrated growth factor in *Rattus norvegicus* calvaria defect model. *J Artif Organs* 2013; 16: 305-315.
29. Romano F, Paolino F M, Rizzo B A, et al. The use of growth factors, CD34+ cells and fibrin for the management of chronic venous ulcers. *Int Wound J* 2016; 13(5): 1011-3.
30. Eppley B L, Woodell J E, Higgins J. Platelet quantification and growth factor analysis from platelet-rich plasma: implications for wound healing. *Plast Reconstr Surg* 2004; 114: 1502-1508.
31. Kalen A, Wahlstrom O, Linder CH, Magnusson P. The content of bone morphogenetic proteins in platelets varies greatly between different platelet donors. *Biochem Biophys Res Commun*. 2008; 375: 261-264.
32. Kim T H, Kim S H, Sandor G K, et al. Comparison of the platelet rich plasma (PRP), platelet rich fibrin (PRF), and concentrated growth factor (CGF) in rabbit skull defect healing. *Arch Oral Biol* 2014; 59: 550-58.
33. Behnia H, Khojasteh A, Soleimani M, Tehrani A, Atashi A. Repair of alveolar cleft defect with mesenchymal stem cells and platelet derived growth factors: A preliminary report. *Journal of Cranio-Maxillofacial surgery* 2012; 40: 2-7.
34. Masuki H, Okedura T, Watanebe T, Suzuki M et al. Growth factor and pro-inflammatory cytokine contents in platelet rich plasma (PRP), Plasma rich in growth factors (PRGF), advanced platelet rich fibrin (A-PRF), and concentrated growth factors (CGF). *International journal of advanced dentistry* 2016; 2: 19.

35. Rosano et al. Immediate post extraction implant placement using plasma rich in growth factors technology in maxillary premolar region: a new strategy for soft tissue management. *J Oral Implantology* 2013; 39: 98.
36. Kikuchi-Taura A, Soma T, Matsuyama T et al. A new protocol for quantifying CD34 (+) cells in peripheral blood of patients with cardiovascular disease. *Tex. Heart Inst. J.* 33(4), 427–429 (2006).
37. Min L, Li Z, Yanmin W. (2013). Present research situation of the orthodontic related alveolar bone defects. *Int J Stomatol.* 40: 500-502.
38. Doğan Ş B, Dede F Ö, Ball U, et al. (2015) Concentrated growth factor in the treatment of adjacent multiple gingival recessions: a split-mouth randomized clinical trial. *J Clin Periodontol.* 42: 868-875.
39. García-Gareta E, Coathup M J, Blunn G W. Osteoinduction of bone grafting materials for bone repair and regeneration. *Bone.* 2015; 81: 112-121.
40. Agrawal A A. Evolution, current status and advances in application of platelet concentrate in periodontics and implantology. *World J Clin Cases.* 2017; 5(5): 159–171.



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Monostotic Fibrous Dysplasia of Maxilla - A Case Report and Review

By Dr. Y. Raghavendra Reddy, Dr. C. N. V. Akhila, Dr. A. Ravi Prakash
& Dr. M. Rajinikanth

Abstract- Fibrous dysplasia belongs to a group of fibro-osseous lesions in which cellular fibrous connective tissue stroma replaces bone. It is a developmental hamartomatous lesion with cases occurring below the age of puberty. Fibrous dysplasia can occur as the monostotic form in which single bone is affected and polyostotic where multiple bones are involved. Majority of the cases reported are the monostotic form with the common site of involvement being the craniofacial skeleton. Polyostotic form are often associated with McCune- Albright syndrome, Jaffe-Lichtenstein syndrome and, Mazabraud syndrome. The syndromic lesions manifest as triad of symptoms - fibrous dysplasias, endocrine abnormalities (endocrinopathies like precocious puberty and hypophosphatemia) and skin pigmentations (a cafe- au-lait spots). Fibrous dysplasias are expansile lesions and cause complications associated with the site of origin. The maxilla is the most common site of involvement in the craniofacial skeleton. In this case, a 13-year-old male patient presented who was having maxillary fibrous dysplasia.

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Monostotic Fibrous Dysplasia of Maxilla - A Case Report and Review

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Abstract- Fibrous dysplasia belongs to a group of fibro-osseous lesions in which cellular fibrous connective tissue stroma replaces bone. It is a developmental hamartomatous lesion with cases occurring below the age of puberty. Fibrous dysplasia can occur as the monostotic form in which single bone is affected and polyostotic where multiple bones are involved. Majority of the cases reported are the monostotic form with the common site of involvement being the craniofacial skeleton. Polyostotic form are often associated with McCune-Albright syndrome, Jaffe-Lichtenstein syndrome and, Mazabraud syndrome. The syndromic lesions manifest as triad of symptoms - fibrous dysplasias, endocrine abnormalities (endocrinopathies like precocious puberty and hypophosphatemia) and skin pigmentations (a cafe-au-lait spots). Fibrous dysplasias are expansile lesions and cause complications associated with the site of origin. The maxilla is the most common site of involvement in the craniofacial skeleton. In this case, a 13-year-old male patient presented who was having maxillary fibrous dysplasia.

I. INTRODUCTION

Fibro-osseous lesions comprise distinct clinical features with complications and co-morbid features. Fibrous dysplasia, cherubism, juvenile ossifying fibroma, osteoma, and aneurysmal bone cyst are the fibro-osseous lesions commonly encountered in the oral cavity. Fibrous dysplasia occurs as relatively rare neoplasm occurring during infancy or childhood. Monostotic fibrous dysplasia accounts for 80-85 % of all the cases with jaws being the prevalent site of involvement ¹.

The etiology is mainly genetic with postzygotic somatic mutations of the GNAS gene. Mutation of which leads to altered bone-forming cells which have fibroblastic phenotype and produce neoplasms at particular sites ². Thereby bone is replaced by fibrous connective tissue leading to irregular trabecular pattern and woven bone formation. Mostly the diagnosis of the lesion is made by clinical and radiographic features not

requiring biopsy. But some of them pose diagnostic difficulties in which biopsy is mandatory.

Treatment modalities differ based on the age and clinical behavior of the neoplasm. Surgical interventions may be difficult as they are more likely to be associated with significant anatomical structures. Follow up plays a major role if incomplete resection (remodeling) when done, lesions are more likely to recur over time. Bisphosphonate therapy can also be indicated in polyostotic fibrous dysplasia. Radiotherapy is contraindicated in these neoplasms as it increases the rate of malignant transformation, with the frequency of sarcoma occurrence ³.

In this article, we present a case of monostotic fibrous dysplasia of maxilla in a 13-year-old patient.

II. CASE REPORT

A 13-year-old male patient reported to G. Pullareddy dental college and hospital, oral and maxillofacial surgery department, with growth in the right maxillary region for one year. The patient complained that the lesion was insidious in onset with unusual growth pattern and attained the present size (Figure 1). The swelling was not associated with pain. No known family history is revealed by the patient. No associated trauma is admitted by patient. Laboratory investigations were normal. On extraoral examination, gross facial asymmetry was noticed on the right side of the face, with a deviation of the nose to left side (Figure 2). Anteriorly the lesion extended below the orbit region at the zygomatic process of the right maxilla without eye involvement. Posteriorly it extended till the malar area of the right side face. Upon inspection the growth was round circumscribed and well-demarcated, measuring 3 X 5 cm. On palpation, the lesion was bony hard, non-tender and was not associated with pain. Intraoral examination revealed expansile swelling extending 5cm anteroposteriorly from the mesial aspect of 14 to distal side of 17 with obliteration of vestibule and 4cm Buccopalatally with bi-cortical expansion, without crossing the mid-palatal line (Figure 3). Retained 54, 55 and clinically missing 13, 15 are observed. Blanching of alveolar mucosa at 55 and 54 is observed. OPG findings revealed radiopaque, ground glass appearance borders blending with adjacent bone (Figure 4). No resorption or displacement of teeth involved is found. CBCT axial view showed varied degrees of opacifications, and coronal view showed thinning of cortical plates, with

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pushing lateral wall and septum of nose medially (Figure 5 & Figure 6).

Provisionally a diagnosis of fibrous dysplasia was considered. Differential diagnosis of other fibro-osseous lesions such as, ossifying fibroma and osseous dysplasia are considered. Taking into consideration the age of the patient surgical recontouring and long term follow up was planned. Under general anesthesia the lesion was shaved, recontoured and reshaped. A vestibular approach was used exposing the anterior aspect till infraorbital region and the bone was shaved using osteotomes and rotary instruments and while periodically checking the amount of bone removal. After adequate removal of the swelling, the area was sutured. The bone bits are sent for histopathological examination (Figure 7).

Microscopically the H & E stained decalcified sections showed irregularly shaped trabeculae of woven bone in a cellular connective tissue stroma. The bony trabeculae are not rimmed by osteoblasts. Curvilinear shaped trabeculae were evident in focal areas. Lesional bone is adjacent to the normal bone. Artifactual peritrabecular clefting is also noticed. The fibrous stroma showed coarse, irregularly arranged collagen fibers with fibroblasts (Figure 8). Based on clinical, radiographic and histological features a final diagnosis of monostotic fibrous dysplasia of maxilla was given, and the patient was advised long term follow up.

III. DISCUSSION

Fibrous dysplasia is a benign congenital fibro-osseous lesion with physiological bone replaced by fibrous tissues. It was first described by Albright et al. in 1937 in a patient with syndromic symptoms of skeletal neoplasms, skin pigmentation, and endocrine abnormalities⁴. Pathogenesis of fibrous dysplasia mainly involves mutations of the *GNAS1* gene (Guanine nucleotide-binding protein) located at chromosome 20q13.2⁵. Early postzygotic mutations cause polyostotic fibrous dysplasia with multiple sites of involvement, and late postzygotic mutations cause monostotic fibrous dysplasia with a single area of occurrence. As these mutations are somatic familial history is not expected, as seen in the present case. These mutations cause hyperfunction of bone progenitor cells which acquire fibroblastic phenotype. The medullary and cortical bones is replaced by sheets of fibro-osseous tissues along with trabecular bone. The fibrous dysplasia occurs below the age of ten years, equally in both males and females.

Clinically the lesion exhibits severe asymmetry and associated symptoms like visual impairment, diplopia, proptosis, hearing loss, nasal obstruction, epistaxis, epiphora, pain, and paresthesia⁶. Laboratory investigations show increased levels of alkaline phosphatase, serum calcium, and serum phosphate

levels. But in monostotic forms, the laboratory investigations are not elevated like in the present case. Radiographic features are generally diagnostic with characteristic ground glass appearance, orange peel or cotton wool appearance of bony trabeculae. Displacement and root resorption of associated teeth are absent in fibrous dysplasia as seen in the present case. Sometimes the density of fibrous dysplasia mimics multilocular appearance often misleading the diagnosis as ameloblastoma. Differential diagnosis of ameloblastoma, ameloblastic fibroma, ameloblastic fibro-odontoma, central giant cell granuloma, odontogenic cyst, ossifying fibroma, chronic sclerosing osteomyelitis, and osteosarcoma are considered for fibrous dysplasia. Treatment may involve radical surgery for devastating forms or shaving and debridement as a part of conservative treatment. Sometimes the lesion is said to be regressed as the patient reaches adulthood, but this concept is not proven. Few cases reports also mentioned about deferring treatment till the patient is above ten years of age or as such with a close follow up.

IV. CONCLUSION

In our case report, we presented clinical, radiographic, histopathological features and, treatment plan for a case of monostotic fibrous dysplasia of maxilla. Proper clinical and radiographic features of the patient are mandatory for confirmative diagnosis. Based on the clinical behavior and age of the patient, appropriate treatment has to be planned with early intervention to avoid complications. Recurrences are frequently seen in fibrous dysplasia. Hence long term follow up should be done. Our case with its clinical and radiographic features represents an addition to the literature of monostotic fibrous dysplasia.

REFERENCES RÉFÉRENCES REFERENCIAS

1. Bansal, Puja & Vaid, Neha. Monostotic Fibrous Dysplasia of the Maxilla: Report of a Rare Case. *Journal of Oral Health & Research*. 2013; 4: 46-50.
2. Bhattacharyya N, Wiench M, Dumitrescu C, Connolly B M, Bugge T H, Patel H V, Gafni R I, Cherman N, Cho M, Hager G L, Collins M T. Mechanism of FGF23 processing in fibrous dysplasia. *Journal of bone and mineral research*. 2012 May; 27(5): 1132-41.
3. Sherman N H, Rao V M, Brennan R E, Edeiken J. Fibrous dysplasia of the facial bones and mandible. *Skeletal radiology*. 1982 May 1; 8(2): 141-3.
4. Warrick C K. Polyostotic Fibrous Dysplasia Albright's Syndrome. *The Journal of bone and joint surgery. British volume*. 1949 May; 31(2): 175-83.
5. Adetayo O A, Salcedo S E, Borad V, Richards S S, Workman A D, Ray A O. Fibrous dysplasia: an

overview of disease process, indications for surgical management, and a case report. *Eplasty*. 2015; 15.

6. Liakos G M, Walker C B, Carruth J A. Ocular complications in craniofacial fibrous dysplasia.

British Journal of Ophthalmology. 1979 Sep 1; 63(9): 611-6.



Figure 1: Extraoral photograph of the patient



Figure 2: Deviation of nose is noticed to left side

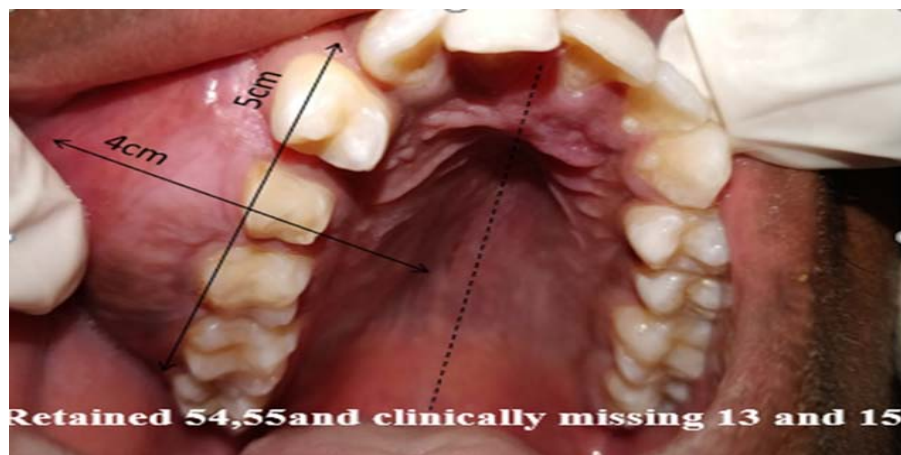


Figure 3: Intraoral Photograph of the patient



Figure 4: OPG



Figure 5: CBCT (axial view)



Figure 6: CBCT (coronal view)



Figure 7: Grossing picture of the lesion after biopsy

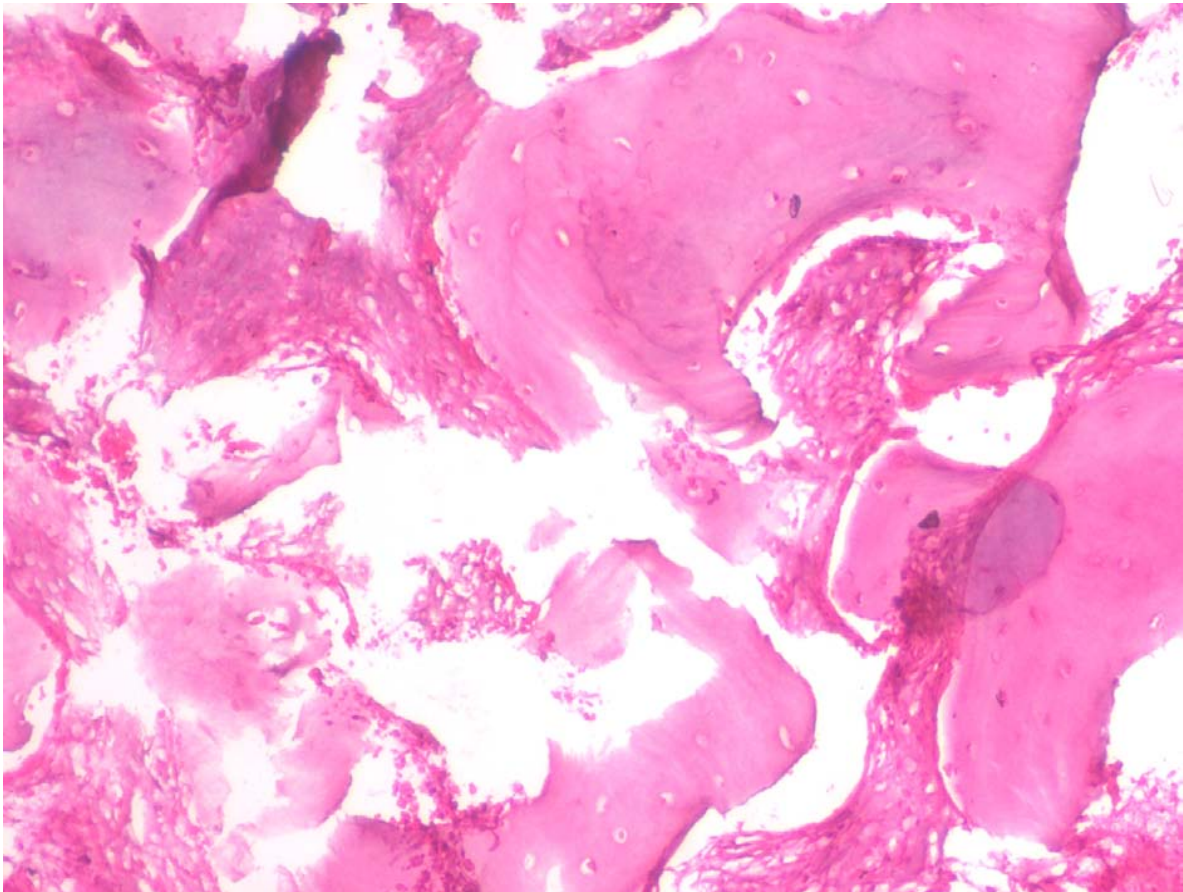


Figure 8: H & E picture of the lesion

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Pterygoid Implant: A Graftless Approach for Maxillary Rehabilitation using TTPHIL-ALL TILT[®] (Tall Tilted Pin Hole Placement with Immediate Loading) Technique

By Dr. P. Venkat Ratna Nag, Dr. P. Sarika & Dr. Tejashree Bhagwatkar

Abstract- Edentulism is a common dental problem in the elderly population. The maxillary posterior edentulous region offers a unique and challenging condition in implant dentistry. The maxillary posterior region presents with numerous anatomic complications in terms of bone quantity, quality, maxillary sinus pneumatization, and poor approachability. Most remarkable surgical approaches include sinus lift, bone grafting, pterygoid implant, considered as valid solutions to this problem. Pterygoid implants require lot of skill of the dentist and also is proven to be statistically superior. In this case series, the pterygoid implants placed and restored using TTPHIL (Tall Tilted Pin Hole placement with Immediate Loading)- ALL TILT[®] technique in the patients with atrophic posterior maxilla are discussed.

Keywords: pterygoid implants, graftless solutions, immediate loading, TTPHIL-ALL TILT[®].

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Dr. P. Venkat Ratna Nag ^α, Dr. P. Sarika ^σ & Dr. Tejashree Bhagwatkar ^ρ

Abstract- Edentulism is a common dental problem in the elderly population. The maxillary posterior edentulous region offers a unique and challenging condition in implant dentistry. The maxillary posterior region presents with numerous anatomic complications in terms of bone quantity, quality, maxillary sinus pneumatization, and poor approachability. Most remarkable surgical approaches include sinus lift, bone grafting, pterygoid implant, considered as valid solutions to this problem. Pterygoid implants require lot of skill of the dentist and also is proven to be statistically superior. In this case series, the pterygoid implants placed and restored using TTPHIL (Tall Tilted Pin Hole placement with Immediate Loading)- ALL TILT® technique in the patients with atrophic posterior maxilla are discussed.

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I. INTRODUCTION

Oral health plays important role to uphold proper mastication, digestion, phonation, aesthetic, and emotional well-being. The loss of permanent teeth can occur for a variety of reasons, ranging anywhere from hereditary factor, congenital absence, diseases of the dentition (e.g., caries or periodontal disease) and trauma.¹ An edentulous condition existing in the posterior maxilla poses a challenge to all the clinicians. The reasons behind this are the anatomic factors like bone quality (type III or IV) according to Lekholm and Zarb ², quantity, location of the maxillary sinus, and poor accessibility in the posterior region^{3,4}.

The pterygoid implants was introduced by Tulasne JF (1992). The advantage of a pterygoid implant is that it allows anchorage, eliminates posterior cantilever and improved axial loading in the posterior atrophic maxilla.⁵ It eradicates the need for

other techniques like sinus lifts or bone grafts^{6,7}, increased implant diameters or short implant of 8mm or lesser and implant placement in zygoma⁹. Placement of implants in the pterygoid region provides posterior bone support, bypass sinus augmentation, or bone grafts. Because of inadequate accessibility, placement of pterygoid implants is more technically demanding than placing short implants anterior to the maxillary antrum. However, there are no risks associated with implant placement in pterygoid area³.

The TTPHIL-ALL TILT® – (Tall Tilted Pin Hole Immediate Loading) concept has evolved from various ideologies in implantology: basal, pterygoid and angulated/tilted implants under immediate loading. To maximize the success of rehabilitation, the TTPHIL technique utilizes the use of long tilted bicortical implants by engaging the pterygoid region. Pterygoid implants were intended to pass through maxillary tuberosity, the pyramidal process of palatine bone and the pterygoid process of the sphenoid bone.^{3,10,11} In the literature, success rates of pterygoid implants for maxillary rehabilitation have been more than 94 to 99%. Thus, the TTPHIL-ALL TILT® technique is a novel approach for the placement of implants; it was derived from established schools of thought in implantology.¹²

In this article, following cases are examples of patients indicated for pterygoid implants using TTPHIL-ALL TILT® in partially edentulous and completely edentulous maxilla.

II. CASE REPORT

Case 1: Partially Edentulous Maxillary Arch (Immediate Implantation)

A 42-year-old male patient reported with a chief complaint of loose teeth in the upper left back region of the jaw. Clinical examination revealed Grade II mobility about tooth 27, gingival recession, missing 26, and poor oral hygiene status. (Fig: 1 a, b) The patient was willing to go ahead with a fixed prosthesis for the same region. TTPHIL- ALL TILT® technique was planned for the same; a thorough radiographic evaluation was suggested to the patient. These studies included panoramic films, CBCT, and stereolithography models. The mouth

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opening was also evaluated and found adequate for pterygoid implants. Routine blood investigations were performed. Radiographic picture (Fig: 1 c) and computed tomography suggested moderate bone loss present in the maxillary sinus region. Considering the amount of bone, atraumatic extraction of 27, followed by the placement of two implants in the left maxillary posteriori. e. 26, 28 (pterygoid implant) was planned and discussed with patient. Patient consent for the treatment was taken. Under aseptic precautions, local anesthesia was given at the surgical sites, and atraumatic extraction of 27 was performed. The socket was curetted carefully and irrigated with sterile saline solution, drilling was done through extraction socket at the planned length. The pilot drill of 1.2mm was then inserted through the mucosa into the alveolar bone of 26 for point of entry to a depth of 6mm. Single drill concept was used, i.e. long stepped drill with diameters of 1.4 or 2.2mm with enough coolant. The under-drilling concept was followed, wherein the dimension of the drills is less than the implant to be positioned for better anchorage. The bioline implant 3.75mm × 13mm and 3.75mm × 22mm mounted on the implant driver, and torque ratchet was used for final placement of 26, 28 (Fig: 2 a). Post-implant placement orthopantomogram was exposed (Fig: 2 b). Multiunit abutments were placed at correct angulations to compensate for the tilt of the implants. The surgical protocol was followed by the prosthetic phase of impression making with transfer copings, Jig trial, bite records, metal framework trial, bisque trial followed by placement of final prosthesis using CAD-CAM technology for design and fabrication was done (Fig: 2 c). The patient was rehabilitated with screw-retained metal-ceramic fixed prosthesis and orthopantomogram was exposed (Fig: 2 d). The patient was advised to adhere to regular oral hygiene maintenance.



a.



b.

Fig. 1 a and b: Intra-oral view showing gingival recession with 27, missing 26



c.

Fig. 1 c: Pre-operative OPG showing moderate bone loss with 27 and minimal bone in the region of maxillary sinus and adequate bone in the pterygoid region.



Fig. 2 a: Clinical photograph showing two implant placements (26, 28) in left maxillary region



Fig. 2 b: Post-operative OPG showing placement of 2 implants in 26, 28 region



Fig. 2 c: Final Metal-ceramic prosthesis was given



Fig. 2 d: Postoperative OPG after fixed implant prostheses

Case 2: Partially Edentulous Maxillary Arch (Delayed Implantation)

A 37-year-old female patient came with a chief complaint of missing teeth and wanted to get them replaced. On intraoral examination (Fig: 3 a), missing 15, 16, 24, 25, 27, root stumps about 17, 26 and badly decayed 11, 12, 13, 21 were observed. The patient was keen on having a fixed prosthesis. On detailed examination of the orthopantomogram (Fig: 3 b), there appeared to be adequate width of the available bone anterior to the maxillary sinus but the inadequate bone in the maxillary sinus region. Treatment plan for the patient was atraumatic extraction of root stumps and immediate placement of six implants i.e. 14, 15, 27 (pterygoid implant), 25, 26, 28 (pterygoid implant). Under aseptic precautions, local anesthesia was administered at the planned surgical sites and atraumatic extraction of 17, 26 was done. Extraction sockets were thoroughly debrided and inspected with the help of periodontal probe for any defect or possible perforation of the cortical plate. Six Bioline implants were placed in the regions of 14, 15, 17 (Pterygoid implant), 25, 26, 28 (Pterygoid implant) (Fig: 4 a). The dimensions of the implants were 3.75 mm × 20 mm (14, 15, 25, 26) and 3.75 mm × 22 mm (Pterygoid implant i.e 17, 28) respectively. The orthopantomogram of post-implant placement was exposed (Fig: 4 b). Multiunit abutments were placed at correct angulations to compensate for the tilt of the implants. Final prosthesis was designed and fabricated using CAD-CAM technology

(Fig: 4 c). Postoperative OPG was taken immediately after fixed implant prostheses. (Fig: 4 d) The patient was advised to adhere to strict soft diet regimen for a few days with regular oral hygiene maintenance and recalled for clinical and radiographic assessment.



Fig. 3 a: Intra-oral view shows missing posterior maxillary teeth and badly decayed upper front teeth

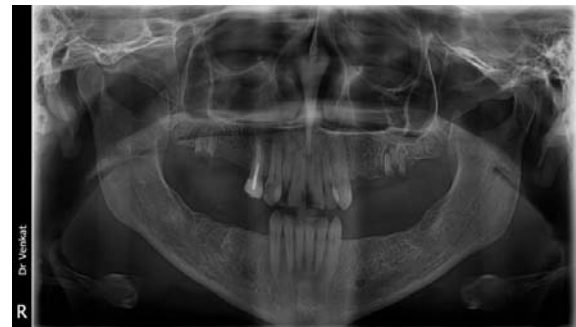


Fig. 3 b: Pre-operative OPG shows adequate width of the available bone anterior to maxillary sinus but inadequate bone in maxillary sinus region



Fig. 4 a: Clinical photograph showing implant placement



Fig. 4 b: Post-operative OPG showing placement of 6 implants



Fig. 4 c: Final Metal-ceramic prosthesis was given

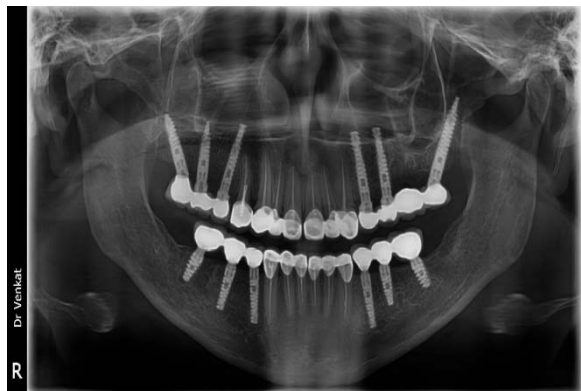


Fig 4 d: Postoperative OPG after fixed implant prostheses

Case 3: Completely Edentulous Maxillary Arch (Immediate Implantation)

A 61-year-old male patient complained of missing teeth in upper back region of jaw and desired replacement. Clinical examination (Fig: 5 a, b) revealed missing 16, 17, 26, 27, Grade II mobility about tooth 25, Grade I mobility about tooth 14, 15, 24 and splinting was present concerning 11, 12, 13. The patient was willing to receive a fixed prosthesis. Medical history revealed a heart problem and high blood pressure. On detailed examination of the orthopantomogram (Fig: 5 c), CT scan shows severe bone loss and inadequate bone in the maxillary sinus region. Extraction of all teeth, and immediate implant placement 13, 15, 17 (pterygoid implant), 23, 25, 27 (pterygoid implant) was planned and executed. Routine blood investigations were performed, and the physician's fitness for the surgical procedure was acquired. Patient consent was obtained before a surgical procedure. Surgical protocol emphasized complete asepsis and infection control. Under local anesthesia, atraumatic extractions of all remaining teeth were completed. Extraction sockets were thoroughly debrided and irrigated with saline solution. Six Bioline implants were placed in the regions of 13, 15, 17 (Pterygoid implant), 23, 25, 27 (Pterygoid implant) (Fig 6 a). Post-implant placement orthopantomogram was exposed (Fig: 6 b). The patient was rehabilitated with screw retained metal-ceramic fixed prosthesis (Fig: 6 c). Postoperative OPG was taken

immediately after fixed implant prostheses. (Fig: 6 d) The patient was followed up after few years for clinical and radiographic assessment.



a.



b.

Fig. 5 a, b: Intra-oral view showing missing posterior maxillary teeth and splinting present in anterior teeth



Fig. 5 c: Pre-operative OPG showing severe bone loss and inadequate bone in maxillary sinus region



Fig. 6 a: Clinical photograph showing implant placement



Fig. 6 b: Post-operative OPG showing placement of 6 implants



Fig. 6 c: Final Metal-ceramic prosthesis was given



Fig. 6 d: Postoperative OPG immediately after fixed implant prostheses

Case 4: Completely Edentulous Maxillary Arch (Delayed Implantation)

A 61-year-old female patient complained of missing upper teeth and wanted replacement of her denture. Intraoral examination revealed edentulous maxillary arch (Fig: 7 a). The patient was using denture for edentulous maxillary arch for two years and wanted a fixed prosthesis. Medical history revealed diabetes and insulin-dependent. On detailed examination of the OPG (Fig: 7 b), CT scan, there appeared severe bone loss and inadequate bone in the maxillary sinus region. Treatment plan was immediate implant placement 13, 15, 17 (Pterygoid implant), 23, 25, 27 (Pterygoid implant). Routine blood investigations were performed, and the physician's fitness was obtained for surgical procedure. Six Bioline implants were inserted in the

planned sites (Fig: 8 a). Post-implant placement orthopantomogram was exposed (Fig: 8 b). The prosthetic phase was carried out. The patient was rehabilitated with a screw-retained metal-ceramic fixed prosthesis (Fig: 8 c). Postoperative OPG was taken immediately after fixed implant prosthesis. (Fig: 8 d)



Fig. 7 a: Intra-oral view showing edentulous maxillary arch



Fig. 7 b: Pre-operative OPG showing severe bone loss and inadequate bone in maxillary sinus region



Fig. 8 a: Clinical photograph showing implant placement

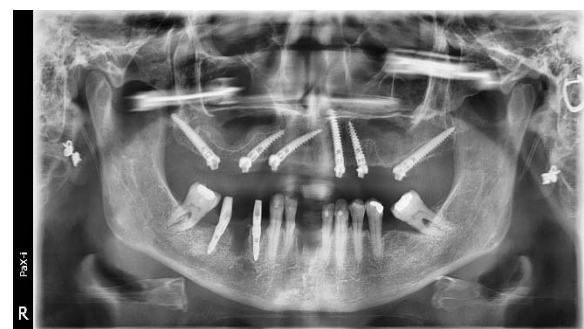


Fig. 8 b: Post-operative OPG showing placement of 6 implants



Fig. 8 c: Final Metal–ceramic prosthesis was given



Fig. 8 d: Postoperative OPG was taken after fixed implant prostheses

III. DISCUSSION

Treatment of edentulous patients has always been a challenging task for the dentists. In the modern era, implant dentistry gained popularity and considered as the best alternative, irrespective of resorbed bone, reduced bone density in the maxillary posterior regions and systemic conditions. Krump et al and Barnett et al reported higher success rates for immediate implant placement.^{13,14}

In the literature, several articles have assigned many labels for implant placement in the posterior maxillary region. Implants placed in the posterior maxilla have been discussed as pterygoid plate implants¹¹, tuberosity implants¹⁵⁻¹⁶, and pterygomaxillary implants¹⁷.

The structures that offer support for implant placement are the tuberosity of the maxillary bone, the pyramidal process of the palatine bone, and the pterygoid process of the sphenoid bone¹⁸.

Conventionally, because of the anatomic factors and biomechanical factors¹⁹, the success rates for implants placed in the atrophic posterior maxilla was lower than that for other locations.

In literature, few studies were carried out on pterygoid implants for the survival of implant and prosthesis. Bahat¹⁵ performed a study on 45 patients, 72 implants were engaged in the tuberosity region, and they noted 93% survival rate over an average loading time of 1.7 years. Balshi¹⁶ reported satisfactory 3-year results for 51 implants placed in the pterygoid area supported fixed prostheses in partially edentulous

patients. Khayat et al²⁰ reported on cases of implants placed in the pterygoid region with four years of follow up. Graves¹¹ described forty-three implants in the pterygoid plate area.

TTPHIL- ALL TILT® (Tall Tilted Pin Hole Immediate Loading) technique²¹⁻²⁵ was evolved after studying and analyzing all the advantages and shortcomings of other traditional and advanced techniques. Using this concept two tall tilted implants engaged the pterygoid region (junction of the palatine process of maxilla, the pyramidal process of palatine bone and pterygoid process of the sphenoid bone), thus eliminating distal cantilever, avoiding sinus encroachment and other augmentation procedures in the posterior maxilla. Tilting of implants avoids anatomical structures like the maxillary sinus. In literature, many studies were carried out on tilted implants, the success rates were found to be 95.7 to 100%²⁶, which have been improved by bicortical engagement. Tall implants with bicortical engagement offer primary stability which improves implant success.²⁷

The subcrestal implant placement decreases the stress in the crestal cortical bone around dental implants. Flapless guided implant placement helps in mucointegration, good aesthetics, reduces the time of treatment, and improves patient satisfaction. Nowadays, practice-driven implantology has been evolved the dimensions (lengths 15-25mm) of implants.²⁸ Multi-unit abutments, and their components help dentists to correct the angulations caused by the tilted implant.²⁹ This aids in immediate loading of the fixed prosthesis.³⁰

IV. CONCLUSION

Based on the outcomes of the case reports, Immediate/Delayed implant placement with immediate loading is considered as a feasible treatment option for the patients with the severely atrophic maxilla. The TTPHIL-ALL TILT® technique considered as a graftless solution which is characterized by Tall, tilted implant with bicortical engagement, minimal invasiveness (flapless approach), screw-retained prosthetic solutions with rigid cross arch stabilization with no cantileverage can be delivered in 2-5 days. However, this technique is highly sensitive and requires experts in implant dentistry for its execution. Careful selection of patients, thorough radiographic evaluation, proper treatment planning, and adequate follow-up of surgical and prosthetic protocols are the keys to success.

REFERENCES RÉFÉRENCES REFERENCIAS

1. Singh M, Kumar L, Anwar M, Chand P. Immediate dental implant placement with immediate loading following extraction of natural teeth. *Natl J Maxillofac Surg* 2015; 6: 252-5.
2. Lekholm U, Zarb G (1985) Patient selection and preparation. In: Brånemark PI, Zarb G, Albrektsson

- T (eds) Tissue-integrated prostheses Osseointegration in clinical dentistry. Quintessence, Chicago, pp 199–209.
3. Balshi T J, Wolfinger G J, Balshi S F II (1999) Analysis of 356 pterygomaxillary implants in edentulous arches for fixed prosthesis anchorage. *Int J Oral Maxillofac Implants* 14: 398–406.
4. Jaffin R A, Berman C L (1991) The excessive loss of Branemark fixtures in type IV bone: a 5-year analysis. *J Periodontol* 62: 2–4.
5. Ali S A, Karthigeyan S, Deivanai M, Kumar A. Implant Rehabilitation For Atrophic Maxilla: A Review. *J Indian Prosthodont Soc* 2014; 14(3): 196–207.
6. Van Steenberghe D, Lekholm U, Bolender C (1990) The applicability of osseointegrated oral implants in the rehabilitation of partial edentulism: a prospective multicenter study on 558 fixtures. *Int J Oral Maxillofac Implants* 5: 272–281.
7. Johansson B, Wannfors K, Ekenbaeck J, Smedberg J I, Hirsch J (1999) Implants and sinus-inlay bone grafts in a one-stage procedure on severely atrophied maxillae: surgical aspects of a 3-year follow-up study. *Int J Oral Maxillofac Implants* 14: 811–818.
8. Hallmann M A (2001) Prospective study of treatment of severely resorbed maxillae with narrow nonsubmerged implants: results after one year of loading. *Int Oral Maxillofac Implants* 16: 731–736.
9. Sorní M, Guarinos J, Penarrocha M (2005) Implants in anatomical buttresses of the upper jaw. *Med Oral Patol Oral Cir Bucal* 10:163–168
10. Bidra AS, Huynh BG. Implants in the Pterygoid Region: A Systematic Review of the Literature. *Int J Oral Maxillofac Surg* 2011; 40: 773–781.
11. Graves S L. The Pterygoid Plate Implant: A Solution for Restoring the Posterior Maxilla. *Int J Periodont Rest Dent* 1994; 14: 513–523
12. Nag PVR, Dhara V, et al. Treatment of the Complete Edentulous Atrophic Maxilla: The Tall Tilted Pin Hole Placement Immediate Loading (TTPHIL)-ALL TILT™ Implant Option. *J Contemp Dent Pract* 2019; 20(6): 754–763.
13. Krump J L, Barnett B G. The immediate implant: A treatment alternative. *Int J Oral Maxillofac Implants* 1991; 6: 19–23.
14. Singh M, Kumar L, Anwar M, Chand P. Immediate dental implant placement with immediate loading following extraction of natural teeth. *Natl J Maxillofac Surg* 2015; 6: 252–5.
15. Bahat O (1992) Osseointegrated implants in the maxillary tuberosity: report on 45 consecutive cases. *Int J Oral Maxillofac Implants* 7: 459–467.
16. Balshi T J (1992) Single tuberosity osseointegrated implant support for a tissue integrated prosthesis. *Int J Periodont Restor Dent* 12: 345–357.
17. Balshi T J, Hy L, Hernandez R E (1995) The use of pterygomaxillary implants in the partially edentulous patient. *Int J Oral Maxillofac Implants* 10: 89–99.
18. Grant JCB (1956) An atlas of anatomy. Williams & Wilkins, Baltimore, p 541
19. Anandakrishna G N, Rao G. Pterygomaxillary Implants: A Graftless Solution to Deficient Maxillary Bone. *J Indian Prosthodont Soc* 2012; 12(3): 182–186.
20. Khayat P, Nader N (1994) The use of osseointegrated implants in the maxillary tuberosity. *Pract Periodontics Aesthet Dent* 6: 53–61.
21. Venkat Ratna Nag P. Immediate Implant Placement and Loading with Tall and Tilted Pinhole Immediate Loading (TTPHIL) Technique. *Guident Sep* 2017: 26–27.
22. Venkat Nag P, Sarika P, Pavankumar, A. (2017). TTPHIL-ALL TILT™ Concept – An Innovative Technique in Immediate Functional Loading Implant Placement in Maxilla. *Scholars Journal of Dental Sciences*. 4(9), 397–399.
23. Venkat Nag P, Sarika P, Khan R, Bhagwatkar T. (2018). Tall and tilted pin hole immediately loaded implants (TTPHIL) technique for maxillary arch rehabilitation. *International Journal of Research and Review*, 5: 104–110.
24. Venkat Nag P, Sarika P, Khan R, Bhagwatkar T. (2018). Immediate implantation and loading in just two days with TTPHIL technique using CAD/CAM Prosthesis. *International Journal of Applied Dental Sciences*, 4(3): 209–213.
25. Venkat Ratna Nag, Sarika P, Bhagwatkar T, Dhara V. Tilted Implant Placement for Rehabilitation of Severe Atrophic Maxilla Supported by Finite Element Analysis. *The Journal of Implant & Advanced Clinical Dentistry* 2019; 11(1): 14–26.
26. Penarrocha O D, Candel M E, Ata A J, et al. Rehabilitation of the Atrophic Maxilla with Tilted Implants: Review of the Literature. *Journal of Oral Implantology* 2013; 38(5) 625–632.
27. Andrea H, Wook-Jin S, Ryan W. Comparison of initial implant stability of implants placed using bicortical fixation, Indirect sinus elevation, and unicortical fixation. *Int J of Oral Max Implants* 2016.31; 2: 459–468.
28. Bruno Salles S M, Camila de Andrade L, Plinio Mendes S. Biomechanical evaluation of subcrestal dental implants with different bone anchorages. *Braz Oral Res.*, (São Paulo) 2014; 28(1): 1–7.
29. Singh A V, Singh S. Tilted Implant Concept for Full Mouth Immediate Loading Restoration. *Int J Oral Implantol Clin Res* 2014; 5(1): 12–23.
30. Nag PVR, Sarika P, Bhagwatkar T, Dhara V. Pterygoid Implant: Option for Rehabilitation of the Atrophic Posterior Maxilla. *Int J Contemp Dent Med Rev* vol.2019, Article ID: 011218, 2019.

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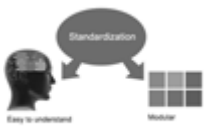
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Institutional Fellow of Open Association of Research Society (USA) - OARS (USA)

Global Journals Incorporation (USA) is accredited by Open Association of Research Society, U.S.A (OARS) and in turn, affiliates research institutions as “Institutional Fellow of Open Association of Research Society” (IFOARS).

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The IFOARS institution is entitled to form a Board comprised of one Chairperson and three to five board members preferably from different streams. The Board will be recognized as “Institutional Board of Open Association of Research Society”-(IBOARS).

The Institute will be entitled to following benefits:



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The author fees of such paper may be waived off up to 40%.

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We shall provide you intimation regarding launching of e-version of journal of your stream time to time. This may be utilized in your library for the enrichment of knowledge of your students as well as it can also be helpful for the concerned faculty members.



After nomination of your institution as “Institutional Fellow” and constantly functioning successfully for one year, we can consider giving recognition to your institute to function as Regional/Zonal office on our behalf.

The board can also take up the additional allied activities for betterment after our consultation.

The following entitlements are applicable to individual Fellows:

Open Association of Research Society, U.S.A (OARS) By-laws states that an individual Fellow may use the designations as applicable, or the corresponding initials. The Credentials of individual Fellow and Associate designations signify that the individual has gained knowledge of the fundamental concepts. One is magnanimous and proficient in an expertise course covering the professional code of conduct, and follows recognized standards of practice.



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- The professional accredited with Fellow honor, is entitled to various benefits viz. name, fame, honor, regular flow of income, secured bright future, social status etc.



- In addition to above, if one is single author, then entitled to 40% discount on publishing research paper and can get 10% discount if one is co-author or main author among group of authors.
- The Fellow can organize symposium/seminar/conference on behalf of Global Journals Incorporation (USA) and he/she can also attend the same organized by other institutes on behalf of Global Journals.
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- • This individual has learned the basic methods of applying those concepts and techniques to common challenging situations. This individual has further demonstrated an in-depth understanding of the application of suitable techniques to a particular area of research practice.

Note :

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- In future, if the board feels the necessity to change any board member, the same can be done with the consent of the chairperson along with anyone board member without our approval.
- In case, the chairperson needs to be replaced then consent of 2/3rd board members are required and they are also required to jointly pass the resolution copy of which should be sent to us. In such case, it will be compulsory to obtain our approval before replacement.
- In case of “Difference of Opinion [if any]” among the Board members, our decision will be final and binding to everyone.

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We accept the manuscript submissions in any standard (generic) format.

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Acknowledgments

Contributors to the research other than authors credited should be mentioned in Acknowledgments. The source of funding for the research can be included. Suppliers of resources may be mentioned along with their addresses.

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The following is the official style and template developed for publication of a research paper. Authors are not required to follow this style during the submission of the paper. It is just for reference purposes.



Manuscript Style Instruction (Optional)

- Microsoft Word Document Setting Instructions.
- Font type of all text should be Swis721 Lt BT.
- Page size: 8.27" x 11", left margin: 0.65, right margin: 0.65, bottom margin: 0.75.
- Paper title should be in one column of font size 24.
- Author name in font size of 11 in one column.
- Abstract: font size 9 with the word "Abstract" in bold italics.
- Main text: font size 10 with two justified columns.
- Two columns with equal column width of 3.38 and spacing of 0.2.
- First character must be three lines drop-capped.
- The paragraph before spacing of 1 pt and after of 0 pt.
- Line spacing of 1 pt.
- Large images must be in one column.
- The names of first main headings (Heading 1) must be in Roman font, capital letters, and font size of 10.
- The names of second main headings (Heading 2) must not include numbers and must be in italics with a font size of 10.

Structure and Format of Manuscript

The recommended size of an original research paper is under 15,000 words and review papers under 7,000 words. Research articles should be less than 10,000 words. Research papers are usually longer than review papers. Review papers are reports of significant research (typically less than 7,000 words, including tables, figures, and references)

A research paper must include:

- a) A title which should be relevant to the theme of the paper.
- b) A summary, known as an abstract (less than 150 words), containing the major results and conclusions.
- c) Up to 10 keywords that precisely identify the paper's subject, purpose, and focus.
- d) An introduction, giving fundamental background objectives.
- e) Resources and techniques with sufficient complete experimental details (wherever possible by reference) to permit repetition, sources of information must be given, and numerical methods must be specified by reference.
- f) Results which should be presented concisely by well-designed tables and figures.
- g) Suitable statistical data should also be given.
- h) All data must have been gathered with attention to numerical detail in the planning stage.

Design has been recognized to be essential to experiments for a considerable time, and the editor has decided that any paper that appears not to have adequate numerical treatments of the data will be returned unrefereed.

- i) Discussion should cover implications and consequences and not just recapitulate the results; conclusions should also be summarized.
- j) There should be brief acknowledgments.
- k) There ought to be references in the conventional format. Global Journals recommends APA format.

Authors should carefully consider the preparation of papers to ensure that they communicate effectively. Papers are much more likely to be accepted if they are carefully designed and laid out, contain few or no errors, are summarizing, and follow instructions. They will also be published with much fewer delays than those that require much technical and editorial correction.

The Editorial Board reserves the right to make literary corrections and suggestions to improve brevity.



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It is necessary that authors take care in submitting a manuscript that is written in simple language and adheres to published guidelines.

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The title page must carry an informative title that reflects the content, a running title (less than 45 characters together with spaces), names of the authors and co-authors, and the place(s) where the work was carried out.

Author details

The full postal address of any related author(s) must be specified.

Abstract

The abstract is the foundation of the research paper. It should be clear and concise and must contain the objective of the paper and inferences drawn. It is advised to not include big mathematical equations or complicated jargon.

Many researchers searching for information online will use search engines such as Google, Yahoo or others. By optimizing your paper for search engines, you will amplify the chance of someone finding it. In turn, this will make it more likely to be viewed and cited in further works. Global Journals has compiled these guidelines to facilitate you to maximize the web-friendliness of the most public part of your paper.

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A major lynchpin of research work for the writing of research papers is the keyword search, which one will employ to find both library and internet resources. Up to eleven keywords or very brief phrases have to be given to help data retrieval, mining, and indexing.

One must be persistent and creative in using keywords. An effective keyword search requires a strategy: planning of a list of possible keywords and phrases to try.

Choice of the main keywords is the first tool of writing a research paper. Research paper writing is an art. Keyword search should be as strategic as possible.

One should start brainstorming lists of potential keywords before even beginning searching. Think about the most important concepts related to research work. Ask, "What words would a source have to include to be truly valuable in a research paper?" Then consider synonyms for the important words.

It may take the discovery of only one important paper to steer in the right keyword direction because, in most databases, the keywords under which a research paper is abstracted are listed with the paper.

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Numerical methods used should be transparent and, where appropriate, supported by references.

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Authors must list all the abbreviations used in the paper at the end of the paper or in a separate table before using them.

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Authors are advised to submit any mathematical equation using either MathJax, KaTeX, or LaTeX, or in a very high-quality image.

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Tables: Tables should be cautiously designed, uncrowned, and include only essential data. Each must have an Arabic number, e.g., Table 4, a self-explanatory caption, and be on a separate sheet. Authors must submit tables in an editable format and not as images. References to these tables (if any) must be mentioned accurately.



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Figures are supposed to be submitted as separate files. Always include a citation in the text for each figure using Arabic numbers, e.g., Fig. 4. Artwork must be submitted online in vector electronic form or by emailing it.

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1. Choosing the topic: In most cases, the topic is selected by the interests of the author, but it can also be suggested by the guides. You can have several topics, and then judge which you are most comfortable with. This may be done by asking several questions of yourself, like "Will I be able to carry out a search in this area? Will I find all necessary resources to accomplish the search? Will I be able to find all information in this field area?" If the answer to this type of question is "yes," then you ought to choose that topic. In most cases, you may have to conduct surveys and visit several places. Also, you might have to do a lot of work to find all the rises and falls of the various data on that subject. Sometimes, detailed information plays a vital role, instead of short information. Evaluators are human: The first thing to remember is that evaluators are also human beings. They are not only meant for rejecting a paper. They are here to evaluate your paper. So present your best aspect.

2. Think like evaluators: If you are in confusion or getting demotivated because your paper may not be accepted by the evaluators, then think, and try to evaluate your paper like an evaluator. Try to understand what an evaluator wants in your research paper, and you will automatically have your answer. Make blueprints of paper: The outline is the plan or framework that will help you to arrange your thoughts. It will make your paper logical. But remember that all points of your outline must be related to the topic you have chosen.

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7. Revise what you wrote: When you write anything, always read it, summarize it, and then finalize it.

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11. Pick a good study spot: Always try to pick a spot for your research which is quiet. Not every spot is good for studying.

12. Know what you know: Always try to know what you know by making objectives, otherwise you will be confused and unable to achieve your target.

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Verbs have to be in agreement with their subjects. In a research paper, do not start sentences with conjunctions or finish them with prepositions. When writing formally, it is advisable to never split an infinitive because someone will (wrongly) complain. Avoid clichés like a disease. Always shun irritating alliteration. Use language which is simple and straightforward. Put together a neat summary.

14. Arrangement of information: Each section of the main body should start with an opening sentence, and there should be a changeover at the end of the section. Give only valid and powerful arguments for your topic. You may also maintain your arguments with records.

15. Never start at the last minute: Always allow enough time for research work. Leaving everything to the last minute will degrade your paper and spoil your work.

16. Multitasking in research is not good: Doing several things at the same time is a bad habit in the case of research activity. Research is an area where everything has a particular time slot. Divide your research work into parts, and do a particular part in a particular time slot.

17. Never copy others' work: Never copy others' work and give it your name because if the evaluator has seen it anywhere, you will be in trouble. Take proper rest and food: No matter how many hours you spend on your research activity, if you are not taking care of your health, then all your efforts will have been in vain. For quality research, take proper rest and food.

18. Go to seminars: Attend seminars if the topic is relevant to your research area. Utilize all your resources.

19. Refresh your mind after intervals: Try to give your mind a rest by listening to soft music or sleeping in intervals. This will also improve your memory. Acquire colleagues: Always try to acquire colleagues. No matter how sharp you are, if you acquire colleagues, they can give you ideas which will be helpful to your research.



20. Think technically: Always think technically. If anything happens, search for its reasons, benefits, and demerits. Think and then print: When you go to print your paper, check that tables are not split, headings are not detached from their descriptions, and page sequence is maintained.

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23. Upon conclusion: Once you have concluded your research, the next most important step is to present your findings. Presentation is extremely important as it is the definite medium through which your research is going to be in print for the rest of the crowd. Care should be taken to categorize your thoughts well and present them in a logical and neat manner. A good quality research paper format is essential because it serves to highlight your research paper and bring to light all necessary aspects of your research.

INFORMAL GUIDELINES OF RESEARCH PAPER WRITING

Key points to remember:

- Submit all work in its final form.
- Write your paper in the form which is presented in the guidelines using the template.
- Please note the criteria peer reviewers will use for grading the final paper.

Final points:

One purpose of organizing a research paper is to let people interpret your efforts selectively. The journal requires the following sections, submitted in the order listed, with each section starting on a new page:

The introduction: This will be compiled from reference matter and reflect the design processes or outline of basis that directed you to make a study. As you carry out the process of study, the method and process section will be constructed like that. The results segment will show related statistics in nearly sequential order and direct reviewers to similar intellectual paths throughout the data that you gathered to carry out your study.

The discussion section:

This will provide understanding of the data and projections as to the implications of the results. The use of good quality references throughout the paper will give the effort trustworthiness by representing an alertness to prior workings.

Writing a research paper is not an easy job, no matter how trouble-free the actual research or concept. Practice, excellent preparation, and controlled record-keeping are the only means to make straightforward progression.

General style:

Specific editorial column necessities for compliance of a manuscript will always take over from directions in these general guidelines.

To make a paper clear: Adhere to recommended page limits.



Mistakes to avoid:

- Insertion of a title at the foot of a page with subsequent text on the next page.
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- Submitting a manuscript with pages out of sequence.
- In every section of your document, use standard writing style, including articles ("a" and "the").
- Keep paying attention to the topic of the paper.
- Use paragraphs to split each significant point (excluding the abstract).
- Align the primary line of each section.
- Present your points in sound order.
- Use present tense to report well-accepted matters.
- Use past tense to describe specific results.
- Do not use familiar wording; don't address the reviewer directly. Don't use slang or superlatives.
- Avoid use of extra pictures—include only those figures essential to presenting results.

Title page:

Choose a revealing title. It should be short and include the name(s) and address(es) of all authors. It should not have acronyms or abbreviations or exceed two printed lines.

Abstract: This summary should be two hundred words or less. It should clearly and briefly explain the key findings reported in the manuscript and must have precise statistics. It should not have acronyms or abbreviations. It should be logical in itself. Do not cite references at this point.

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Approach:

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Approach:

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Approach:

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<i>References</i>	Complete and correct format, well organized	Beside the point, Incomplete	Wrong format and structuring



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