A COMPARATIVE EVALUATION OF EFFICACY OF AUTOGENOUS BONE GRAFT AND BIOACTIVE GLASS ALLOPLAST IN THE TREATMENT OF INTRAOSSEOUS PERIODONTAL DEFECTS- A CLINICO-RADIOLOGICAL STUDY

Dr. Pratibha Chaudhary\* Dr. Amandeep Dhillon\*, Dr. Sujata Masamatti\* Dr. Vidya Dodwad\*\*, Dr. Subhra Vaish\*\*\*, Dr. Nikhil\*\*\*\*

## ABSTRACT

**Aim :** The aim of this study was to compare the efficacy of autogenous bone graft and bioactive glass in the treatment of infrabony periodontal defects.

**Material and methods:** The present study is a randomized controlled trial of 6 months duration with intrabony defects of three and two wall defects which were equally divided into test and control groups.

**Results :** The mean Pocket depth decreased significantly (p < 0.05) from baseline to 6 months in the control group and test groups.

**Conclusion:** Periodontal osseous surgery tends to improve the clinical condition of gingiva of the patient leading to enhanced clinical results of periodontal therapy in periodontally affected sites.

#### **KEY WORDS**

Periodontal, autogenous bone graft, alloplast, perioglas etc

#### **ABOUT THE AUTHORS**

\* MDS, \*\*Principal at Bhartiya Vidyapeeth, Pune \*\*\*Professor, \*\*\*\*HOD, Department Of Periodontolgy And Oral Implanolgy

#### **CORRESPONDING AUTHOR**

Dr. Pratibha Chaudhary MDS e- Mail Id : pratibha.chaudhary17@gmail.com

## INTRODUCTION

Periodontal disease is among the most prevalent disease worldwide and is the major cause of tooth morbidity. The disease is characterized by presence of gingival inflammation, periodontal pocket formation, loss of connective tissue attachment and alveolar bone around the affected teeth. The primary goal of periodontal therapy is to arrest the progression of periodontal disease and maintain the natural dentition in health and comfortable function.<sup>1</sup>

Different types of bone deformities can result from periodontal disease. Vertical and angular defects are those that occur in an oblique direction, leaving a hollowed-out trough in the bone alongside the root, the base of the defect is located apical to the surrounding bone.<sup>2</sup>

Bone grafting procedures with autogenous bone grafts, allografts, xenografts, and alloplasts are used to promote periodontal regeneration.<sup>3</sup> Among the different graft materials available, autogenous bone remains the gold standard for osseous regeneration<sup>4</sup> because autograft bone have osteogenic property, less chances of immunological reactions, minimal inflammatory reactions, rapid revascularization graft particle and potential release of growth factors(Marx1994)<sup>5</sup>

Clinical measurements usually involve measuring pocket depths and clinical attachment loss. Both of these measurements rely on the use of a periodontal probe. Probing measurements can often be inaccurate due to a variety of operator, armamentarium and patient dependent factors. These factors include probe angulation, force, diameter of the tine, amount of inflammation, anxiety and discomfort during clinical evaluation.<sup>5</sup>Van der Zee has found that probe tiny diameter and calibration may have an effect on the measuring pocket depth.<sup>6</sup> Theil found that probe readings were not a precise measurement of attachment loss, especially in areas of increasing destruction and on multi-rooted teeth.<sup>7</sup>

Three-dimensional imaging in dentistry has become popular and gives the clinician an ability to visualize and measure bone level without structures being superimposed and enhance periodontal diagnosis compared to regular radiographs.<sup>8</sup>

Due to limited radiation exposure, ability to take a small focused field, and ability to avoid having distortion independent of the location of the tooth, CBCT has great potential for evaluation of depth and architecture of infrabony defects. The measurements from CBCT is as accurate as using periodontal probe and superior to PA's in diagnosing buccal and lingual defects. In difficult treatment planning cases, CBCTs have tremendous potential for making the initial exam more informative and the process of diagnosis and treatment planning more precise.<sup>9</sup>

Hence this prospective, randomized controlled trial is planned to evaluate the efficacy of autogenous bone graft in treatment of periodontal infrabony defects using CBCT as a diagnostic tool for appropriate diagnosis and treatment.

## **MATERIALS AND METHODS**

The subjects for this study were selected from the Outpatient Department of Periodontology, I.T.S-

CDSR, Muradnagar, Ghaziabad; Uttar Pradesh. The sample size of this study was 20 sites with three and two wall defects ,with angular defect. Patients will be assigned to the following two groups randomly and was followed for 6 months

GROUP 1- Defect sites were treated with application autogenous bone graft (TEST GROUP)

GROUP 2- Defect sites were treated with application of perioglas (CONTROL GROUP)

#### Inclusion criteria:

- 1. Systemically healthy subjects.
- 2. Subjects with age between 18-50 years.

3. Patients suffering from chronic periodontitis and each subject having minimum number of 20 teeth present.

4. Patients having periodontal pockets with probing depth≥5mm.

5. Patients showing radiographic evidence of infrabony osseous defects.



**Figure 1:** Armamentatrium and Surgical Procedure (test site) **A**. Contrangle Hand Piece with Trephine Bur and Bone Mill **B**. Pocket probing depth and RCAL at baseline **C**. Debridement of the defect **D**. Bone graft procured from extracted site **E**. Placement of the bone graft **F**. Sutures placed **G**. Pocket probing depth and relative clinical attachment level at 6 months **H**. CBCT at Baseline **I**. CBCT at 6 Months.



Figure 2. Surgical procedure (control site) A. Debridement of the defect B. Placement of the graft (perioglas) at the defect C. Sutures placed D. CBCT at baseline E. CBCT at 6months

6. Subjects who were willing to comply with all the study related procedure were allowed sign the inform consent form.

#### **Exclusion criteria:**

1. Patients who were medically compromised.

2. Patients with any history of allergy to the material used in this study

3. Patients who were on antibiotic or antimicrobial therapy in previous 6 months.

4. Patients on any drug therapy which was known to influence the periodontium.

5. Patients who were pregnant or lactating.

6. Patients who had undergone any type of periodontal flap surgical procedure or regenerative therapy 6 months prior to the initial examination.

- 7. Patients who were tobacco users.
- 8. Subjects unable to provide informed consent.

### SURGICAL PROCEDURE

All the periodontal surgical procedure were performed on patients suffering from chronic periodontitis<sup>1</sup> under aseptic conditions. Standardized surgical procedure for the test sites was performed as follows:

Surgical area was anesthetized using local anesthesia (2% lignocaine with adrenaline 1:80,000) After intracrevicular incision, full thickness mucoperiosteal flap was elevated without any vertical incisions. Fully expose the infrabony defects followed by meticulous debridement and thoroughlt irrigated wit sterile saline. After the surgical site was prepared an adequate amount of particulate cortical bone harvested with the trephine bur from the buccal cortical plate 10 adjacent to the defect or any extracted site or parasymphysis region and put it into the bone crusher to squeeze out the bone and placed to the defect site. The flaps were repositioned to the presurgical level and sutured with 3-0 silk. Post operative medications included a single standard regimen of oral dministration of amoxicillin 500mg

thrice a day for 5 days, Ibuprofen 400 mg+ Paracetamol 325mg thrice daily for 5 days and 0.2% chlorhexidiene gluconate twice daily for a period of 2 weeks. sutures were removed after 7 days.

## STATISTICAL ANALYSIS

Statistical analysis was performed using a commercially available software program SPSS (Statistical Package for Social Sciences) version 21.0 and Epi-info version 3.0.Mann-Whitney U test is used for comparison of mean value between 2 groups when the data does not follow the normal distribution. Wilcoxon sign rank pair test was used for comparison of 2 mean values obtained from a same group or a pair of values obtained from the same sample when the data does not follow the normal distribution. Friedman's test was used for comparison of more than 2 mean values obtained from the same sample when the data does not follow the normal distribution.

## RESULT

The purpose of the present clinical trial was to evaluate the efficacy of bioactive glass particles (PerioGlas®) and autogenous bone graft in the treatment of periodontal intrabony defects. For this study 20 patients in the age of 18-60 years fulfilling the inclusion and exclusion criteria contributing to a total of 20 intrabony defects, were recruited. The test group with 10 intrabony defects was treated with autogenous bone as bone graft material and while the control group 10 intrabony defects were treated with bioactive glass particle (PerioGlas®) alone.

All the surgical sites healed uneventfully. Neither allergic reactions nor suppuration or abscess formation was observed at any surgical site. No teeth were extracted during the course of the study.

#### Measurement of GI

The mean GI 11at baseline for the control group and the test group was  $1.20\pm0.42$  and  $1.00\pm0.00$ respectively. The mean GI at 6 months was found to be  $0.70\pm0.42$  and  $0.20\pm0.42$  for the control and the test group respectively (TABLE1, GRAPH 1). The mean Gingival Index decreased significantly (p<0.05) from baseline to 6 months for test group (TABLE 3) as compared to the control group.

#### Measurement of PI

The mean Plaque Index12 at baseline for the control group and the test group was  $1.20\pm0.42$  and  $1.20\pm0.42$  respectively. The mean Plaque Index for the control and the test group at 6 months was  $0.90\pm0.32$  and  $0.30\pm0.48$  at 6 months respectively (TABLE 1, GRAPH 1). The mean Plaque Index decreased significantly from baseline to 6 months in test group (TABLE 3).But there was no significant change in the mean Plaque Index for the inter-interval comparison for control group (TABLE 3).

		Test Group		Contr	ol Group			
		Mean	Std. Deviation	Mean	Std. Deviation	Mean Difference	t-test value	p-value
Plaque index	Baseline	1.20	0.42	1.20	0.42	0.00	0.000	1.000
	3 months	0.80	0.42	1.00	0.00	-0.20	-1.500	0.151
	6 months	0.30	0.48	0.90	0.32	-0.60	-3.286	0.004*
Gingival index	Baseline	1.00	0.00	1.20	0.42	-0.20	-1.500	0.151
	3 months	0.70	0.48	1.00	0.00	-0.30	-1.964	0.065
	6 months	0.20	0.42	0.70	0.48	-0.50	-2.466	0.024*
	Baseline	6.50	1.43	6.20	1.14	0.30	0.519	0.610
Pocket depth	3 months	3.10	0.57	4.10	1.10	-1.00	-2.554	0.020*
	6 months	2.40	0.52	3.70	1.06	-1.30	-3.488	0.003*
RCAL	Baseline	9.70	1.95	9.70	1.16	0.00	0.000	1.000
	3 months	6.60	1.51	8.00	0.94	-1.40	-2.492	0.023*
	6 months	5.10	1.45	7.10	0.88	-2.00	-3.735	0.002*
defect depth	Baseline	8.34	2.07	8.48	1.87	-0.14	-0.159	0.876
	6 months	3.69	1.79	6.43	2.15	-2.74	-3.100	0.006*

 TABLE 1: INTER GROUP COMPARISON OF CLINICAL PARAMETERS AT VARIOUS TIME INTERVALS.

 \*SIGNIFICANT DIFFERENCE



**GRAPH 1: MEAN PLAQUE INDEX, GINGIVAL INDEX, PROBING DEPTH, RCAL AND RADIOGRAPHIC DEPTH AT VARIOUS TIME INTERVALS IN TEST AND CONTROL GROUPS** 

		Test Group		Control Group				
		Mean	SD	Mean	SD	Mean Difference	t-test value	p-value
Difference in PI	from baseline to 3 months	0.40	0.52	0.20	0.42	0.20	0.949	0.355
	from baseline to 6 months	0.90	0.32	0.30	0.48	0.60	3.286	0.004*
	from 3 to 6 months	0.50	0.53	0.10	0.32	0.40	2.058	0.044*
Difference in GI	from baseline to 3 months	0.30	0.48	0.20	0.42	0.10	0.493	0.628
	from baseline to 6 months	0.80	0.42	0.50	0.53	0.30	3.406	0.047*
	from 3 to 6 months	0.50	0.53	0.30	0.48	0.20	2.885	0.038*
Difference in PD	from baseline to 3 months	3.40	1.43	2.10	0.74	1.30	2.555	0.020*
	from baseline to 6 months	4.10	1.45	2.50	0.71	1.60	3.138	0.006*
	from 3 to 6 months	0.70	0.48	0.40	0.52	0.30	1.342	0.196
Difference in RCAL	from baseline to 3 months	3.10	1.45	1.70	0.67	1.40	2.769	0.013*
	from baseline to 6 months	4.60	1.35	2.60	0.70	2.00	4.160	0.001*
	from 3 to 6 months	1.50	0.85	0.90	0.57	0.60	1.857	0.080
Difference in Defect Depth from baseline to 6 months		4.65	1.07	2.05	0.74	2.60	6.325	0.001*

# TABLE 2: INTER GROUP COMPARISION OF MEAN DIFFERENCE IN CLINICALPARAMETERS AT VARIOUS TIME INTERVALS.

\* SIGNIFICANT DIFFERENCE

			Test C	Broup	Control Group		
			Mean Difference	p-value	Mean Difference	p-value	
Plaque index	Baseline	3 months	0.40	0.046*	0.20	0.157	
	Baseline	6 months	0.90	0.003*	0.30	0.083	
	3 months	6 months	0.50	0.025*	0.10	0.317	
Gingiyal	Baseline	3 months	0.30	0.043*	0.20	0.110	
index	Baseline	6 months	0.80	0.014*	0.50	0.243	
	3 months	6 months	0.50	0.043*	0.30	1.000	
	Baseline	3 months	3.40	< 0.001*	2.10	< 0.0 01 *	
Pocket depth	Baseline	6 months	4.10	< 0.001*	2.50	< 0.0 01 *	
	3 months	6 months	0.70	0.015*	0.40	0.110	
RCAL	Baseline	3 months	3.10	< 0.001*	1.70	< 0.001 *	
	Baseline	6 months	4.60	< 0.001*	2.60	< 0.0 01 *	
	3 months	6 months	1.50	0.003*	0.90	0.002*	
Defect depth	Baseline	6 months	4.65	< 0.001*	2.05	< 0.001 *	

TABLE 3: INTRA GROUP COMPARISION OF MEAN DIFFERENCE IN CLINICAL PARAMETERS AT VARIOUS TIME INTERVALS IN CONTROL GROUP. \* SIGNIFICANT DIFFERENCE

	Test Group		Contro	l Group			
Radiographic defect depth	Mean	Std. Deviation	Mean	Std. Deviation	Mean Difference	t-test value	p-value
Baseline	8.34	2.07	8.48	1.87	-0.14	-0.159	0.876
6 months	3.69	1.79	6.43	2.15	-2.74	-3.100	0.006*
Difference from baseline to 6 months	4.65	1.07	2.05	0.74	2.60	6.325	0.001*

#### TABLE 4: THE COMPARISON OF MEAN DEFECT DEPTH AT BASELINE AND 6 MONTHS

## Measurement of POCKET PROBING DEPTH (PD).

The mean PD at baseline for the control and the test group was  $6.20\pm1.14$  mm and  $6.50\pm1.43$ mm respectively. At 3 months the mean PD reduced to  $4.10\pm1.10$  mm and  $3.10\pm0.57$  mm for the control and the test group respectively. The mean PD for the control and the test group was  $3.70\pm1.06$  mm and  $2.40\pm0.52$ mm at 6 months respectively (TABLE 1, GRAPH 1). The mean difference in Pocket depth

from baseline to 3 months and from baseline to 6 months was significantly more (p>0.05) in the Test group in comparison to Control group.(TABLE 3).

## Measurement of RELATIVE CLINICAL ATTACHMENTLEVEL (RCAL)

The mean RCAL at baseline for the control and the test group was  $9.70\pm1.16$  mm and  $9.70\pm1.95$  mm respectively. At 6 months RCAL for the control





GRAPH 3 :MEAN DIFFERENCE IN DEFECT DEPTH BETWEEN TEST AND CONTROL GROUPS FROM BASELINE TO 6 MONTHS

7.10 $\pm$ 0.88 and for the test group was 5.10 $\pm$ 1.45 mm (TABLE 1, GRAPH 1). The reduction of RCAL in test group was significantly more from baseline to 6 months (p>0.05) as compared to Control group. (TABLE 2).

## Measurement of RADIOGRAPAHIC DEFECT DEPTH

The mean radiographic defect depth at baseline for the control and the test group was  $8.48\pm 1.87$  mm and  $8.34\pm 2.07$  mm respectively. At 6 months, the mean defect depth reduced to  $6.43\pm 2.15$ mm and  $3.69\pm 1.79$ mm for the control and the test groups respectively. (TABLE 4, GRAPH3). There was significant difference in mean difference in defect depth from baseline to 6 months between Control and Test groups. (TABLE 4, GRAPH2).The mean Defect depth decreased significantly from baseline to 6 months for control group and test group. (TABLE 3)

## DISCUSSION

The goal of periodontal surgery is access for definitive calculus removal and surgical management of bony irregularities which have resulted from the disease process to reduce pockets as much as possible and improving the clinical attachment level. The mean **Relative Clinical Attachment Level** at baseline for the control and the test group was  $9.70\pm1.16$  mm and  $9.70\pm1.95$  mm respectively. At 3 months the mean RCAL reduced to  $8.00\pm0.94$  mm and  $6.60\pm1.51$  mm for the control and the test group respectively. The mean RCAL for the control and the

test group was  $7.10\pm0.88$  mm and  $5.10\pm1.45$  mm at 6 months respectively. The results of our study are consistent with the study done by keles et al<sup>14</sup> who found the significant CAL gain of 4.50±0.80mm in the sites treated with Autogenous bone graft in a period of 6months which was in accordance with our study. **Orsini et al**<sup>15</sup> showed the CAL gain of  $5.25\pm$ .75 in the group treated with autogenous bone graft at a period of 6 months. Nevins et al<sup>16</sup> in series of cases also recorded a mean CAL gain of 2.2 mm after 6 month following (Perioglas). The findings of our study are consistent with study by **Park et al**<sup>13</sup> who found a significant CAL gain in sites treated with bioactive glass compared to OFD at 6 months. Similar findings were observed in the study done by **Froum et al**<sup>17</sup> who reported significant improvement in clinical attachment level gain in the bioactive glass sites compared to the control sites. The gain in the CAL might have resulted from periodontal regeneration via new attachment or healing characterized by the formation of long junctional epithelium between the new regenerated tissues and the root surface.95

To substantiate the improvement in clinical parameters, radiographic assessment were done for alveolar bone changes. CBCT scans were done for radiographic evaluation of the test and control sites showed changes in the appearance of the graft material from the time of placement to 6 months. The mean **Radiographic Defect Depth** at baseline for the control and the test group was  $8.48\pm 1.87$  mm and  $8.34\pm 2.07$  mm respectively. At 6 months, the mean defect depth was reduced to  $6.43\pm 2.15$ mm and  $3.69\pm 1.79$ mm respectively. The mean defect depth at 6 months was significantly more (p>0.05) among test

group in comparison to control group. These findings are in accordance with the study done by **Keles et al**<sup>14</sup> who showed the bone fill of  $5.92\pm1.83$  mm in the ACB graft-treated group. **Crea et al**<sup>18</sup> who reported statistically significant gain in defect depth of 3.03 mm in sites treated with (IMP + OFD) while the sites treated with OFD alone showed a linear bone growth of 1.69 mm. **Frorm et al**.<sup>17</sup> also found that defect depth reduction was significantly greater in the bioactive glass sites (4.36 mm) compared to the OFD sites (3.15 mm).

The success of periodontal therapy depends on many factors, one of the most important factors is an accurate image of the morphology of periodontal bone destruction for the differential therapeutic treatment plan. The clinical diagnosis of periodontal osseous lesions, such as vertical bone defects or furcation involvement, presents a challenge for the practitioner<sup>20</sup>. Therefore, it is widely recognized that the treatment of patients with advanced periodontal diseases requires not only extensive clinical recording but also radiological examination<sup>21</sup>. Radiographs are unavoidable to determine the extent and severity of the periodontal lesions<sup>22</sup>. In general, the accurate detection of bony defects is only possible after direct intraoperative/surgical control. Threedimensional information is represented by twodimensional plane, thus losing essential diagnostic value<sup>23</sup>. For this reason, the spatial representation of the alveolar bone has a significant role in periodontology, as therapy decisions and long-term estimates of tooth-related prognosis is based on it. Cone beam computed tomography (CBCT) images of periodontal bone lesions offer a highly informative value.

## CONCLUSION

Within the limitations of this study, both Autogenous bone graft and alloplastic grafting led to similar improvements in clinical and radiographic parameters 6 months after the treatment of intraosseous periodontal defects. Autogenous bone grafts, a rich source of bone and marrow cells, have been accepted as the gold standard for bone grafting procedures. However, harvesting of intraoral bone is restricted to donor sites that yield comparatively limited graft volume. For the present study, more number of randomized clinical controlled trials with larger sample size and a longer follow up period along with histological examinations are required to further explore the benefits of Autogenous bone graft in the management of intrabony defects.

## REFERENCES

1. Paul SR, MarkAR, Gerald MB. The treatment of intrabony defects with bone grafts. Periodontol 2000;22:88-103.

2. Carranza, FA. Bone Loss and Patterns of Bone Destruction. In Newman, MG; Takei, HH; Carranza, FA; editors: Carranza's Clinical Periodontology, 9th Edition. Philadelphia: W.B. Saunders company, 2002;363

3. Kim CS, Choi SH, Cho KS, Chai JK, Wikesjo UM, Kim CK. Periodontal healing in one wall intrabony defects in dogs following implantation of autogenous bone or a coral-derived biomaterial. J Clin Periodontol 2005;32:583-9.

4. Marx RE. Clinical application of bone biology to mandibular and maxillary reconstruction. Clin Plast surgery 1994;21:377-92.

5. Van der Zee E, Davies EH, Newman HN. Marking width, calibration from tip and tine diameter of periodontal probes. J Clin Periodontol 1991;18:516-20.

6. Theil EM, Heaney TG. The validity of periodontal probing as a method of measuring loss of attachement. J Clin Periodontol 1991;18:648-53.

7. Noujeim M, Prihoda T, Langlais R, Nummikoski P. Evaluation of high-resolution cone beam computed tomography in the detection of simulated interradicular bone lesions. Dento Maxillofacial radiology 2009;38:156-62.

8. Mohan R, Singh A, Gundappa M. Three dimesnsional imaging in periodontal diagnosis – Utilization of cone beam computed tomography. J Ind Soc Periodontol 2011;15:11-7.

9. Rost A. Using CBCT as a diagnostic tool for evaluation of infrabony defects in vivo. Virginia commonwealth university 2014;21:4-7.

10. Mellonig JT. Autogenous and allogeneic bone grafts in periodontal therapy. Crit Rev Oral Biol Med. 1992;3:333-52.

11. Loe H, Silness J.Periodontal disease in pregnancy. I. Prevalence and severity. Acta Odontol Scand 1964;22:121-35.

12. Turesky S, Gilmore ND, Glickman I. Reduced plaque formation by the chloromethyl analogue of victamin C. J Periodontol 1970;41:41-3.

13. Park JS, Suh JJ, Choi SH, Moon IS, Cho KS, Kim CK et al. Effects of pretreatment clinical parameters on bioactive glass implantation in intrabony periodontal defects. J Periodontol. 2001;72:730-740.

14. Keles GC, Sumer M, Cetinkaya BO, Tutkun F, Simsek SB. Effect of autogenous cortical bone grafting in conjunction with guided tissue regeneration in the treatment of intraosseous periodontal defects. Eur J Dent. 2010;4:403–11

15. Orsini G, Benlloch D. Comparison of Calcium sulfate and Autogenous bone graft in intrabony defects: A split mouth technique. J Periodontol 2001;22:296-309.

16. Nevins M, Giannobile WV, McGuire MK, Kao RT, Mellonig JT, Hinrich JE, McAllister BS, Murphy

KS, McClain PK, Nevins ML, Paquette DW, Han TJ, Reddy M, Lavin PT, Genco RJ, Lynch SE. Platelet derived growth factor stimulates bone fill and rate of attachment level gain: Results of a large multicenter randomized controlled trial. J Periodontol 2005;76: 2205-15.

17. Froum SJ, Weinberg MA, Tarnow D. Comparison of bioactive glass synthetic bone graft particles and open debridement in the treatment of human periodontal defects. A clinical study. J Periodontol. 1998;69:698-709.

18. Crea A, Deli G, Littarru C, Lajaolo C, Orgeas GV, Tatakis DN. Intrabony Defects, open flap debridement, and decortications: a randomized clinical trial. J Periodontol. 2014;85:34-42.

19. Listgarten MA. Periodontal probing what does it mean. J Clin Periodontol 1980;7:165–76.

20. Tyndall D, Rathore S. Cone-beam CT diagnostic applications: caries, periodontal bone assessment, and endodontic applications. Dent Clin North Am 2008;52:825–41.

21. Tugnait A, Clerehugh V, Hirschmann PN. The usefulness of radiographs in diagnosis and management of periodontal diseases: a review. J Dent 2000;28:219–26.

22. Jeffcoat MK, Wang IC, Reddy MS. Radiographic diagnosisin periodontics. Periodontology 2000;7:54–68.

23. Pihlstrom BL. Periodontal risk assessment, diagnosis and treatment planning. Periodontology 200025:37–58.