

CASE REPORT

Rehabilitation of Lower Anteriors with Implant Therapy in Localized Aggressive Periodontitis

Varshal J Barot

ABSTRACT

Periodontitis is the most common, destructive, inflammatory disease of supporting tissues of the teeth in humans, with profound effects on general health. Aggressive periodontitis (AgP) comprises a rare group of population, a rapidly progressing form of periodontitis, characterized by severe destruction of periodontal attachment apparatus and tooth loss at an early age. Considering the psychological problems that these patients have faced during the early stages of their life, the main aim of treatment is to achieve functional, esthetic, and phonetic rehabilitation. Dental implant is a widely used treatment option that provides functional and esthetic resolution. Implant placement in patients with a history of AgP might be considered a viable option to restore oral function; however, the risk for implant failure is significantly higher.

This report presents rehabilitation of lower anteriors with implant therapy in a 24-year-old systemically healthy male with history of localized AgP, who had previously received complete periodontal therapy for periodontal stabilization.

Keywords: Aggressive periodontitis, Dental implants, Periodontal diseases.

How to cite this article: Barot VJ. Rehabilitation of Lower Anteriors with Implant Therapy in Localized Aggressive Periodontitis. *Int J Clin Dent Res* 2017;1(1):40-44.

Source of support: Nil

Conflict of interest: None

INTRODUCTION

Every periodontal entity has a distinct progressive pattern associated with different bacteria. Thus, it becomes an utmost necessity to treat and control periodontal disease, regardless of its progression pattern and subtype before implant therapy is initiated to improve implant longevity, in the pursuit of excellence in implant dentistry. A higher incidence of peri-implantitis and a lower implant survival rate are being reported in patients susceptible to periodontitis;^{1,2} it is important to mention that the pre-existing ecologic conditions of the oral cavity influence

biofilm formation on implants. Success in implant dentistry relies on the initial osseointegration and long-term stability.³ Patient systemic factors and susceptibility to periodontal diseases, implant macrodesign and microdesign, and periodontal pathogenic bacteria, among others, have all been shown to play a role in achieving long-term implant stability.⁴

CASE REPORT

A 24-year-old healthy male was a known case of localized aggressive periodontitis (AgP) (Figs 1 and 2) who had received periodontal therapy and extractions with severely drifted lower central incisors before 4 months. He received scaling, root planing, and soft tissue curettage in



Fig. 1: Preoperative (before periodontal therapy for treatment of AgP) clinical, front view



Fig. 2: Preoperative (before periodontal therapy for treatment of AgP) clinical, lingual view

Periodontist and Implantologist

Satyam Multispeciality Dental Clinic, Vadodara, Gujarat, India

Corresponding Author: Varshal J Barot, Periodontist and Implantologist, Satyam Multispeciality Dental Clinic, Vadodara Gujarat, India, Phone: +919558805347, e-mail: varshalb@gmail.com



Fig. 3: Postoperative (after periodontal therapy for treatment of AgP) clinical, front view

conjunction with systemic antibiotics. His medical history was not significant, and he had no adverse habits.

Intraoral examination revealed anterior open bite along with pathologic migration of maxillary and mandibular anteriors (Fig. 3). Maxillary arch showed largely spaced, flared anterior teeth, whereas in mandibular arch a four-unit bridge spacing in anterior region was present with history of extracted both central incisors. Radiographic examination using cone beam computed tomography (CBCT) of mandible was done to assess the existing bone morphology (Fig. 4). After informed consent, prosthetic rehabilitation was planned with a four-unit bridge on two implants in the lower anteriors.

The surgery was performed under local anesthesia – bilateral mental nerve block. Patient was given an antibiotic prophylaxis for 48 hours (amoxicillin 500 mg given orally 1 hour preoperatively and every 6 hours postoperatively). Following midcrestal incision, full-thickness flaps were reflected and implant sites were prepared bilaterally (Fig. 5). Two titanium plasma spray-coated implants, 13 mm in length and 3.3 mm in diameter, were placed in the anterior region of the mandible (Figs 6A and B). Flaps were approximated using 4-0 absorbable



Fig. 5: After full-thickness flaps reflection at proposed implant sites in the anterior region of the mandible

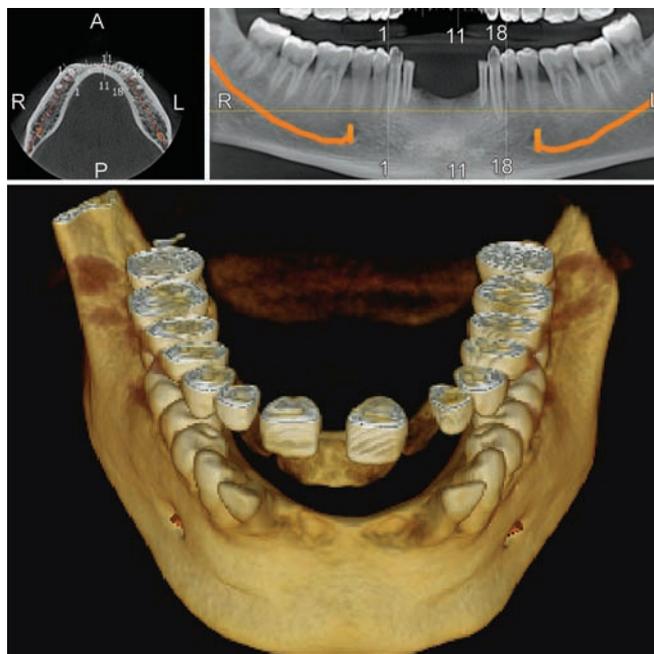
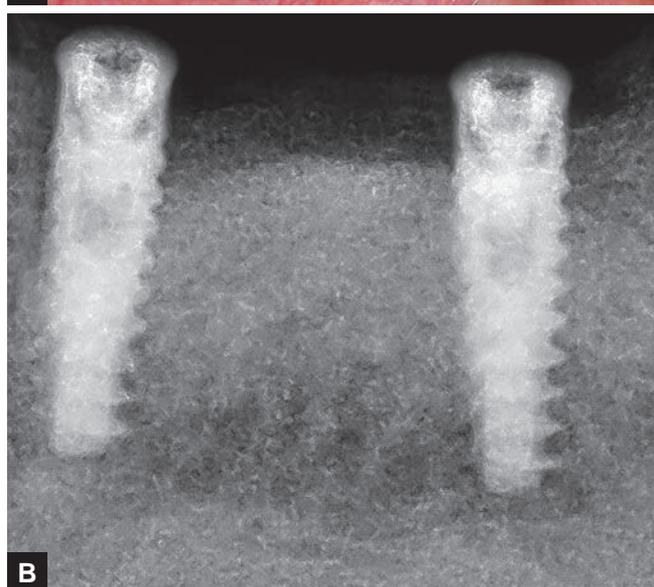
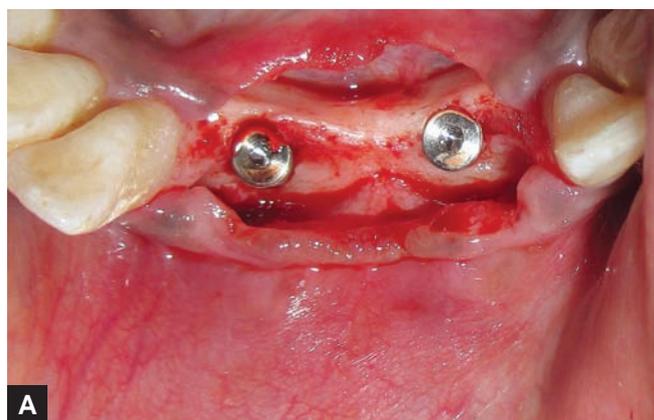


Fig. 4: Radiographic examination using CBCT mandible to assess existing bone morphology



Figs 6A and B: (A) Two implants, 13 mm in length and 3.3 mm in diameter, placed in the anterior region of the mandible; and (B) The IOPA radiograph showing well-aligned implants in the anterior region of the mandible, immediate postinsertion



Fig. 7: Flaps were approximated using 4-0 absorbable (vicryl) interrupted sutures

(vicryl) interrupted sutures (Fig. 7). Analgesics (tablet ibuprofen, 400 mg every 4–6 hours) along with antimicrobial rinse (0.2% chlorhexidine gluconate twice-a-day for 4 weeks) were prescribed, and the patient was recalled after 7 days for follow-up.

At 3 months postsurgery, the implant sites were reopened with crestal incisions and gingival former was screwed. Healing was uneventful with both the implants well osseointegrated, and phase 2 surgical uncovering was accomplished. The gingival former was replaced with abutments 3 weeks later (Fig. 8) and a four-unit porcelain-fused-to-metal (PFM) prosthesis was inserted (Figs 9A and B). Patient was kept on regular maintenance and follow-up every 4 months (Fig. 10). Both implants were functioning and immobile when tested clinically, with no evidence of peri-implant radiolucency in intraoral periapical (IOPA) radiograph and CBCT at 1-year follow-up postloading (Figs 11 and 12).

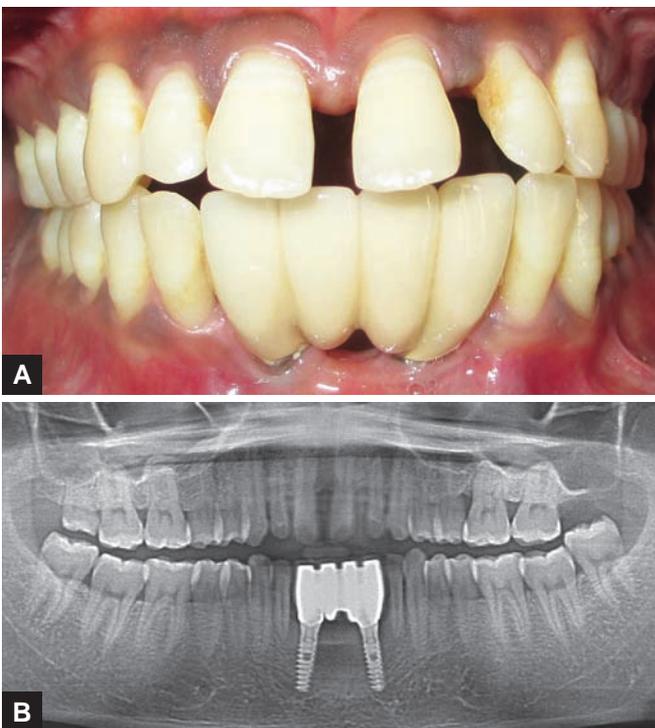


Fig. 8: Two abutments placed 3 weeks later of gingival former

DISCUSSION

Baer⁵ defined AgP as “a disease of the periodontium occurring in an otherwise healthy adolescent, which is characterized by a rapid loss of alveolar bone around more than one tooth of the permanent dentition.” According to the American Academy of Periodontology⁶ classification, the term “AgP” was adopted as a new name for this unique disease classification, replacing the term “early-onset periodontitis” and the disease being classified into localized and generalized forms⁷:

- Localized aggressive periodontitis:
 - Circumpubertal onset.
 - Localized first molar/incisor presentation with interproximal attachment loss (AL) on at least two



Figs 9A and B: (A) Four-unit PFM prosthesis inserted; and (B) The OPG showing well-connected prosthesis



Fig. 10: Clinical view showing patient's normal smile before and after prosthetic insertion in the anterior region of the mandible

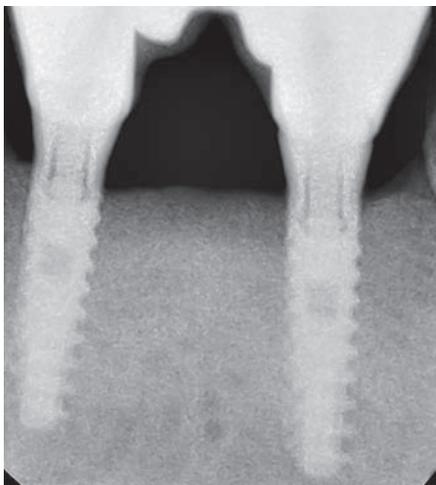


Fig. 11: Intraoral periapical radiograph at 1-year follow-up postloading

permanent teeth (one of which is a first molar) and involving no more than two teeth other than first molars and incisors.

- Robust serum antibody response to infecting agents.
- Generalized aggressive periodontitis:
 - Usually affecting persons under 30 years of age, but patients may be older.
 - Generalized interproximal AL affecting at least three permanent teeth other than first molars and incisors.
 - Pronounced episodic nature of the destruction of attachment and alveolar bone.
 - Poor serum antibody response to infecting agents.

Three major characteristics were used to define the aggressive disease^{7,8}:

1. Clinically healthy with the exception of periodontitis;
2. Rapid AL and bone breakdown; and
3. Familial aggregation.

Other characteristics can also be used in the diagnosis of the disease⁷⁻⁹:

- Amounts of microbial deposits inconsistent with the severity of periodontal tissue breakdown;
- Elevated proportions of *Aggregatibacter actinomycetem-comitans* and *Porphyromonas gingivalis*;
- Phagocyte abnormalities;
- Hyperresponsive macrophage phenotype, including elevated levels of prostaglandin E2 and interleukin (IL)-1b; and
- Self-arresting progression of AL and bone loss.

Polymorphisms in genes regulating the expression of IL-1, IL-6, IL-10, tumor necrosis factor, E-selectins, Fc-g receptor, cluster of differentiation 14, toll-like receptors, caspase recruitment domain 15, vitamin D receptor, lactoferrin, caldesmon, heat shock protein 70, and Stac protein 23 and major histocompatibility complexes A9 and B1524 were associated with AgP. As a consequence of these polymorphisms, the inflammatory profile is altered, including, but not limited to, polymorphonuclear neutrophil (PMN) transendothelial migration and signaling functions,¹⁰ reduced chemotactic response, and depression in neutrophil phagocytosis and superoxide production.¹¹

Apse et al¹² stated that the peri-implant sulcus behaves similar to the periodontal sulcus, and, therefore, an inflammatory process similar to periodontitis occurs around implants, i.e., peri-implantitis. Mombelli et al¹³ identified the same pathogens in peri-implant lesions as the ones that were present 6 months before in natural dentition. Implant placement in patients with a history

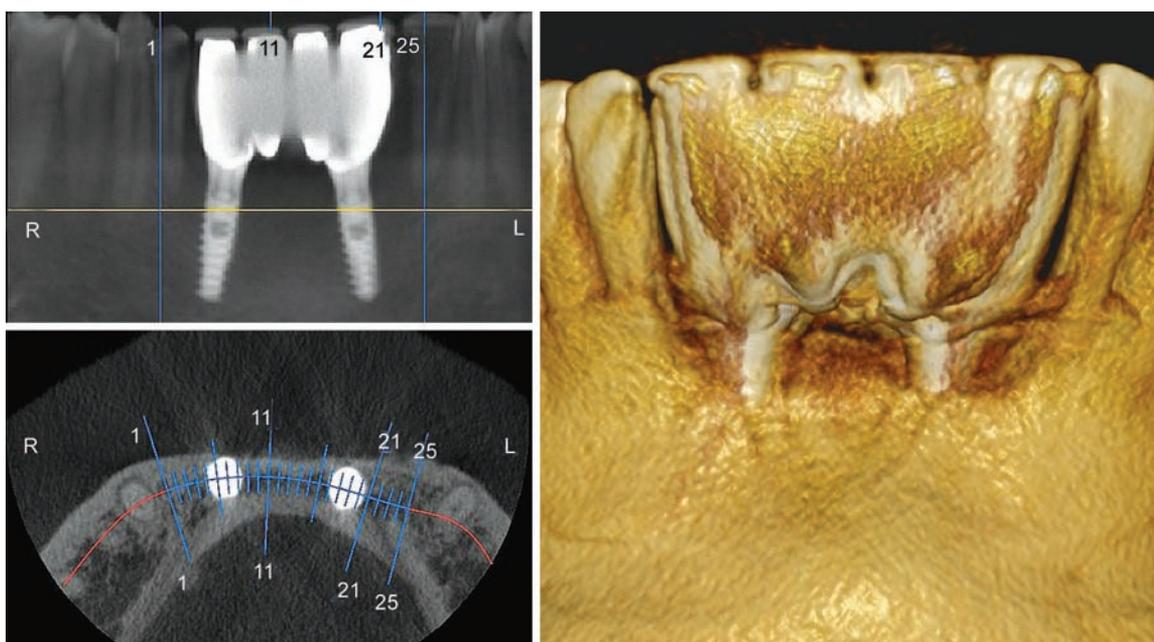


Fig. 12: Radiographic examination using CBCT at 1-year follow-up postloading

of AgP might be considered a viable option to restore oral function, with survival outcomes similar to those found in both patients with chronic periodontitis and healthy patients. However, the risk ratio for implant failure in patients with AgP is significantly higher when compared with healthy patients and those with chronic periodontitis.¹⁴

In AgP patients, often unmodifiable factors would potentially play a role in implant success. These factors include^{10,11,15}

- Genetic polymorphisms;
- Alterations of the immune system (phagocyte abnormalities and hyper-responsive macrophage phenotype, altered PMN transendothelial migration and signaling functions, reduced chemotactic response, and depression in phagocytosis and superoxide production)
- Depression, stress, and loneliness;
- Oral hygiene; and
- Tobacco consumption.

For an implant to be deemed successful, the first year mean bone loss (MBL) of about 0.9 to 1.6 mm has been reported to be acceptable. During the subsequent years, the MBL has been reported to decrease to 0.05–0.13 mm annually.^{16,17} Mengel et al¹⁸ showed that patients with generalized AgP exhibited MBL of 2.07 mm during the first year after implant placement.

CONCLUSION

Both in AgP and peri-implantitis processes, a comprehensive implant maintenance program to identify peri-implant bone loss early is highly encouraged, specifically in patients with a history of aggressive periodontal disease due to unmodifiable conditions that might play a dominant role. A good case selection, meticulous treatment plan, and good patient compliance with long-term regular follow-up confer successful placement of implants in patients with AgP.

REFERENCES

1. Al-Zahrani MS. Implant therapy in aggressive periodontitis patients: a systematic review and clinical implications. *Quintessence Int* 2008 Mar;39(3):211-215.
2. Donos N, Laurell L, Mardas N. Hierarchical decisions on teeth vs. implants in the periodontitis-susceptible patient: the modern dilemma. *Periodontol* 2000 2012 Jun;59(1):89-110.
3. Linkow LI, Rinaldi AW, Weiss WW Jr, Smith GH. Factors influencing long-term implant success. *J Prosthet Dent* 1990 Jan;63(1):64-73.
4. Quirynen M, De Soete M, van Steenberghe D. Infectious risks for oral implants: a review of the literature. *Clin Oral Implants Res* 2002 Feb;13(1):1-19.
5. Baer PN. The case for periodontosis as a clinical entity. *J Periodontol* 1971 Aug;42(8):516-520.
6. Armitage GC. Development of a classification system for periodontal diseases and conditions. *Ann Periodontol* 1999 Dec;4(1):1-6.
7. Lang N, Bartold PM, Cullinan M, Jeffcoat M, Mombelli A, Murakami S, Page R, Papapanou P, Tonetti M, Van Dyke T. Consensus report – aggressive periodontitis. *Ann Periodontol* 1999;4:53.
8. American Academy of Periodontology. Parameter on aggressive periodontitis. *J Periodontol* 2000 May;71(Suppl 5):867-869.
9. Demmer RT, Papapanou PN. Epidemiologic patterns of chronic and aggressive periodontitis. *Periodontol* 2000 2010 Jun;53:28-44.
10. Gronert K, Kantarci A, Levy BD, Clish CB, Odparlik S, Hasturk H, Badwey JA, Colgan SP, Van Dyke TE, Serhan CN. A molecular defect in intracellular lipid signaling in human neutrophils in localized aggressive periodontal tissue damage. *J Immunol* 2004 Feb 1;172(3):1856-1861.
11. Nishimura F, Nagai A, Kurimoto K, Isoshima O, Takashiba S, Kobayashi M, Akutsu I, Kurihara H, Nomura Y, Murayama Y, et al. A family study of a mother and daughter with increased susceptibility to early-onset periodontitis: microbiological, immunological, host defensive, and genetic analyses. *J Periodontol* 1990 Dec;61(12):755-762.
12. Apse P, Ellen RP, Overall CM, Zarb GA. Microbiota and crevicular fluid collagenase activity in the osseointegrated dental implant sulcus: a comparison of sites in edentulous and partially edentulous patients. *J Periodontal Res* 1989 Mar;24(2):96-105.
13. Mombelli A, Nyman S, Bragger U, Wennstrom J, Lang NP. Clinical and microbiological changes associated with an altered subgingival environment induced by periodontal pocket reduction. *J Clin Periodontol* 1995 Oct;22(10):780-787.
14. Monje A, Alcoforado G, Padiar-Molina M, Suarez F, Lin GH, Wang HL. Generalized aggressive periodontitis as a risk factor for dental implant failure: a systematic review and meta-analysis. *J Periodontol* 2014 Oct;85(10):1398-1407.
15. Monteiro da Silva AM, Oakley DA, Newman HN, Nohl FS, Lloyd HM. Psychosocial factors and adult onset rapidly progressive periodontitis. *J Clin Periodontol* 1996 Aug;23(8):789-794.
16. Adell R, Lekholm U, Rockler B, Brånemark PI. A 15-year study of osseointegrated implants in the treatment of the edentulous jaw. *Int J Oral Surg* 1981 Dec;10(6):387-416.
17. Albrektsson T, Zarb G, Worthington P, Eriksson AR. The long-term efficacy of currently used dental implants: a review and prognosis criteria for success. *Int J Oral Maxillofac Implants* 1986 Summer;1(1):11-25.
18. Mengel R, Behle M, Flores-de-Jacoby L. Osseointegrated implants in subjects treated for generalized aggressive periodontitis: 10-year results of a prospective, long-term cohort study. *J Periodontol* 2007 Dec;78(12): 2229-2237.