

"Comparative Evaluation of Concentrated Growth Factor versus Bone Graft in Crestal Sinus Lift with Simultaneous Implant Placement in the Posterior Maxilla: A Prospective Case–Control Study"

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Abstract

Background: Implant placement in the posterior maxilla is often limited by reduced residual bone height due to sinus pneumatization and ridge resorption. Crestal sinus lift with simultaneous implant placement is a minimally invasive option, but the ideal grafting material remains debated.

Objective: To compare clinical and radiographic outcomes of concentrated growth factor (CGF) versus conventional bone graft in crestal sinus lift with simultaneous implant placement in the posterior maxilla.

Materials and Methods: In this prospective case–control study, 30 ASA I–II patients requiring implant placement in the posterior maxilla with residual bone height ≤ 5 mm were randomly allocated to two groups (n=15 each): Group A (bone graft) and Group B (CGF). All patients underwent osseodensification-assisted crestal sinus lift with simultaneous implant placement. Outcomes included implant stability (ISQ), insertion torque, radiographic bone gain and density (IOPA and CBCT), marginal bone loss, postoperative pain (VAS), soft-tissue healing (Landry index), and complications over 6 months.

Results: Primary implant stability was comparable between groups. At 4 months, the bone graft group showed significantly higher ISQ (78.2 ± 2.8 vs 75.1 ± 3.3 ; $p=0.008$). Bone gain at 3 months was greater with bone graft (3.4 ± 0.6 mm vs 2.9 ± 0.5 mm; $p=0.021$), while CGF showed significantly higher early bone density at 1 month (110.5 ± 13.6 vs 95.2 ± 12.8 grayscale units; $p=0.004$). At 6 months, implant stability, bone gain, bone density, marginal bone loss, implant survival, and prosthetic success were comparable. Postoperative pain and morbidity trends favored CGF, though not significantly.

Conclusion: Both CGF and bone graft are effective for crestal sinus lift with simultaneous implant placement. Bone graft provides better early mechanical stability and bone gain, whereas CGF offers advantages in early bone density and biological healing. CGF can be considered a viable autologous alternative in appropriately selected cases.

Keywords: Crestal sinus lift; Concentrated growth factor; Bone graft; Posterior maxilla; Dental implants; Osseodensification.

How to cite this article: Yadav M, Bhutani H, Kumar S, Kashyap K, Mittal A., Comparative Evaluation of Concentrated Growth Factor versus Bone Graft in Crestal Sinus Lift with Simultaneous Implant Placement in the Posterior Maxilla: A Prospective Case–Control Study. *Int J Drug Deliv Technol.* 2026;16(78s): 105; DOI: 10.25258/ijddt.16.78s.12

Introduction

Implant-supported rehabilitation has become a cornerstone of modern dentistry, offering predictable restoration of function, aesthetics, and phonetics. Long-term success depends on stable osseointegration and healthy peri-implant hard and soft tissues, which in turn require adequate bone volume and quality at the recipient site. The posterior maxilla remains one of the most challenging regions due to progressive alveolar ridge resorption after tooth extraction and inferior pneumatization of the maxillary sinus, often leaving insufficient vertical bone height for standard implant placement.

To overcome these anatomical limitations, maxillary sinus floor elevation (MSFE) techniques have been widely adopted. Among these, the crestal (transcrestal) approach is less invasive than the lateral window technique and is particularly suitable when only limited vertical augmentation is required and simultaneous implant placement is planned. However, maintaining the space created beneath the elevated Schneiderian membrane remains critical to ensure predictable bone formation and implant stability.

Traditionally, particulate bone grafts (xenografts, allografts, alloplasts, or autografts) are used to stabilize the membrane, maintain volume, and act as an osteoconductive scaffold. Despite their success, graft

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materials have drawbacks, including cost, variable resorption, potential donor-site morbidity (for autografts), and risk of complications such as sinusitis or graft migration, especially in cases of membrane perforation.

Concentrated Growth Factor (CGF), an autologous platelet concentrate obtained through differential centrifugation of venous blood, forms a dense fibrin matrix enriched with growth factors such as PDGF, TGF- β , and VEGF. CGF is proposed to enhance angiogenesis, cell migration, and early wound stability, potentially improving healing and bone quality in sinus augmentation. Unlike particulate grafts, CGF lacks rigid volumetric structure, raising questions about its ability to maintain space under the membrane in cases with severely reduced residual bone height.

Although systematic reviews suggest that autologous platelet concentrates may improve early healing and reduce postoperative discomfort when combined with grafts, high-quality comparative trials of CGF versus conventional bone grafts in crestal sinus lift with simultaneous implants are limited. This study aims to compare the clinical and radiographic outcomes of CGF versus bone graft in crestal sinus lift with simultaneous implant placement in the posterior maxilla.

Materials and Methods

Study Design and Setting

This prospective case–control study was conducted at the Department of Oral and Maxillofacial Surgery, I.T.S. Dental College, Hospital and Research Centre, Greater Noida, India, between 2023 and 2026. The study protocol was approved by the Institutional Ethical Committee (IEC), and all participants provided written informed consent.

Sample Size and Power Calculation

A total of 30 patients were recruited based on a priori sample size calculation targeting a clinically meaningful difference in implant stability (ISQ) and radiographic bone gain between groups, with $\alpha=0.05$ and power=80%.

Inclusion and Exclusion Criteria

Inclusion criteria:

- Adults aged 18–65 years
- ASA physical status I or II
- Partially edentulous posterior maxilla requiring implant placement
- Residual alveolar bone height ≤ 5 mm and adequate bone width for implant placement
- Good general and oral health

Exclusion criteria:

- Uncontrolled systemic diseases (e.g., uncontrolled diabetes, bleeding disorders)
- History of radiation therapy to the head and neck
- Active sinus pathology or severe sinusitis
- Heavy smoking (>10 cigarettes/day) or substance abuse
- Poor oral hygiene or untreated periodontal disease
- Pregnancy or lactation

Randomization and Group Allocation

Eligible patients were randomly allocated into two groups using a computer-generated randomization sequence:

Group A (Bone Graft): Crestal sinus lift with particulate bone graft and simultaneous implant placement

Group B (CGF): Crestal sinus lift with CGF and simultaneous implant placement

Each group comprised 15 patients.

Surgical Protocol

All surgeries were performed under local anesthesia by the same surgeon using a standardized protocol.

Preoperative Preparation:

Clinical examination, OPG, and CBCT to assess residual bone height, width, sinus anatomy, and pathology.

Oral prophylaxis and preoperative instructions.

Crestal Sinus Lift (Osseodensification-Assisted):

A crestal incision with or without releasing incisions was made.

Full-thickness mucoperiosteal flap reflection exposed the alveolar crest.

Osteotomy was performed using Densah burs (osseodensification) in a counter-clockwise direction to compact bone laterally and elevate the sinus floor hydraulically.

The Schneiderian membrane was carefully elevated using specialized instruments.

Graft Placement:

Group A: Particulate bone graft material was placed into the created sub-antral space to maintain elevation and support new bone formation.

Group B: CGF was prepared by drawing 9 mL of venous blood from each patient into special tubes without anticoagulant and centrifuging according to the manufacturer's protocol. The middle CGF layer (fibrin-rich) was harvested and placed into the sub-antral space.

Implant Placement:

Implants were placed immediately after graft/CGF placement.

Primary stability was assessed via insertion torque (Ncm) and Implant Stability Quotient (ISQ) using resonance frequency analysis.

Closure:

Flaps were repositioned and sutured with 3-0 or 4-0 non-resorbable/resorbable sutures to achieve primary closure.

Postoperative Care and Follow-up

All patients received standard postoperative instructions, analgesics, and antibiotics. Sutures were removed at 7–10 days. Clinical and radiographic evaluations were performed at 0, 1, 3, 4, and 6 months. Second-stage surgery and prosthetic loading were performed at approximately 4–6 months, depending on clinical and radiographic evidence of osseointegration.

Outcome Measures

Primary outcomes:

Implant stability (ISQ) at insertion, 4 months, and 6 months

Radiographic bone gain (mm) at 1, 3, and 6 months via IOPA

Secondary outcomes:

Insertion torque (Ncm)

Bone density (grayscale units via IOPA/Image J; HU via CBCT subset)

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Marginal bone loss at 6 months

Implant survival and prosthetic success at 6 months

Postoperative pain (VAS scores at days 1, 3, 7, and suture removal)

Soft-tissue healing (Landry index at 1 week)

Complications (sinus perforation, sinusitis, wound dehiscence, graft/CGF exposure, persistent pain)

Operator-assessed handling properties of graft material (ease of placement, containment, adhesion, overall satisfaction)

Radiographic Assessment

IOPA: Standardized intra-oral periapical radiographs were taken at 0, 1, 3, 4, and 6 months. Bone gain was measured from the implant apex to the highest point of new bone formation. Bone density was assessed using Image J software (grayscale analysis).

CBCT: Performed in a subset of 5 patients per group at baseline and 6 months to evaluate 3D bone gain, bone density (HU), and dome formation.

Statistical Analysis

Data were analyzed using SPSS. Normality was assessed using Shapiro–Wilk test. Continuous variables were compared using unpaired t-test, and categorical variables using Chi-square or Fisher's exact test. A p-value <0.05 was considered statistically significant.

Results

Baseline Characteristics

A total of 30 patients (15 per group) were included. Mean age was 48.7±6.2 years in Group A and 50.1±5.8 years in Group B (p=0.524). Gender distribution, ASA status, residual bone height, and residual bone width were comparable between groups (all p>0.05), confirming baseline homogeneity.

Implant Stability (ISQ)

At insertion: ISQ was 68.4±3.1 (Group A) vs 67.9±3.4 (Group B); p=0.682.

At 4 months: ISQ increased to 78.2±2.8 (Group A) vs 75.1±3.3 (Group B); p=0.008 (significant).

At 6 months: ISQ was 82.6±1.9 (Group A) vs 81.8±2.1 (Group B); p=0.285.

Primary stability was comparable, but bone graft showed significantly higher secondary stability at 4 months.

Insertion Torque and Prosthetic Outcomes

Mean insertion torque was 33.5±2.1 Ncm (Group A) vs 32.8±2.4 Ncm (Group B); p=0.401.

Implant survival at 6 months was 100% (15/15) in Group A and 93.3% (14/15) in Group B (p=0.500). Prosthetic success at 1 month post-loading was 100% in Group A and 93.3% in Group B (p=0.500). Time to prosthesis loading was 126.3±4.5 days (Group A) vs 124.8±5.2 days (Group B); p=0.391.

Radiographic Bone Gain (IOPA)

Baseline: 0.0 mm in both groups.

1 month: 2.1±0.5 mm (Group A) vs 1.8±0.4 mm (Group B); p=0.092.

3 months: 3.4±0.6 mm (Group A) vs 2.9±0.5 mm (Group B); p=0.021 (significant).

6 months: 3.8±0.5 mm (Group A) vs 3.5±0.6 mm (Group B); p=0.143.

Bone graft showed significantly greater bone gain at 3 months, but differences were not significant at 6 months.

Bone Density (Grayscale via IOPA/Image J)

Baseline (adjacent bone): 142.3±15.1 (Group A) vs 138.7±14.3 (Group B); p=0.521.

0 month: 95.2±12.8 (Group A) vs 110.5±13.6 (Group B); p=0.004 (significant).

4 months: 125.7±11.4 (Group A) vs 130.8±12.1 (Group B); p=0.253.

6 months: 140.1±10.2 (Group A) vs 142.9±9.8 (Group B); p=0.445.

CGF demonstrated significantly higher early bone density at 0 month, but values converged by 6 months.

CBCT Analysis (Subset: n=5 per group)

Bone gain at 6 months: 4.1±0.7 mm (Group A) vs 3.7±0.6 mm (Group B); p=0.328.

Bone density at baseline: 762.4±125.3 HU (Group A) vs 745.8±118.7 HU (Group B); p=0.823.

Bone density at 6 months: 1125.6±156.4 HU (Group A) vs 1210.3±143.9 HU (Group B); p=0.394.

Dome formation at 6 months: 100% in both groups.

CBCT confirmed comparable 3D bone gain and density at 6 months.

Postoperative Healing and Complications

Intra-op sinus perforation: 1 case (6.7%) in Group A; none in Group B (p=1.000).

Post-op sinusitis: 1 case (6.7%) in Group A; none in Group B (p=1.000).

Wound dehiscence: 2 cases (13.3%) in Group A; 1 case (6.7%) in Group B (p=1.000).

Persistent pain (>1 week): 3 cases (20.0%) in Group A; 1 case (6.7%) in Group B (p=0.598).

Graft/CGF exposure: 1 case (6.7%) in Group A; none in Group B (p=1.000).

No statistically significant differences were observed, though complication trends favored CGF.

Postoperative Pain (VAS)

Day 1: 6.5±1.3 (Group A) vs 5.8±1.4 (Group B); p=0.167.

Day 3: 4.2±1.1 (Group A) vs 3.5±1.0 (Group B); p=0.083.

Day 7: 1.3±0.8 (Group A) vs 1.0±0.7 (Group B); p=0.291.

At suture removal: 0.5±0.6 (Group A) vs 0.3±0.5 (Group B); p=0.337.

Pain scores were consistently lower in the CGF group but not significantly.

Soft-Tissue Healing (Landry Index at 1 Week)

Very poor/poor: 1 case (6.7%) in Group A; none in Group B.

Fair: 3 cases (20.0%) in Group A; 1 case (6.7%) in Group B.

Good: 7 cases (46.7%) in Group A; 9 cases (60.0%) in Group B.

Excellent: 4 cases (26.7%) in Group A; 5 cases (33.3%) in Group B.

No significant differences were observed (p>0.05).

Operator-Assessed Handling Properties

Ease of placement: 4.1±0.7 (Group A) vs 3.5±0.9 (Group B); p=0.048 (significant).

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Containment in site: 4.4 ± 0.6 (Group A) vs 3.0 ± 0.8 (Group B); $p=0.001$ (significant).
Adhesion to implant: 2.2 ± 0.8 (Group A) vs 4.6 ± 0.5 (Group B); $p=0.001$ (significant).

Overall satisfaction: 3.9 ± 0.7 (Group A) vs 3.7 ± 0.8 (Group B); $p=0.482$.
Bone graft was easier to place and contain, while CGF showed superior adhesion to the implant surface.

Armamentarium



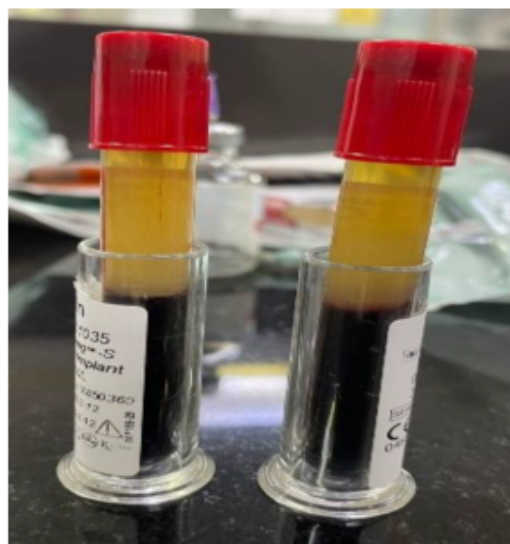
Minor oral surgical instruments



densah bur kit

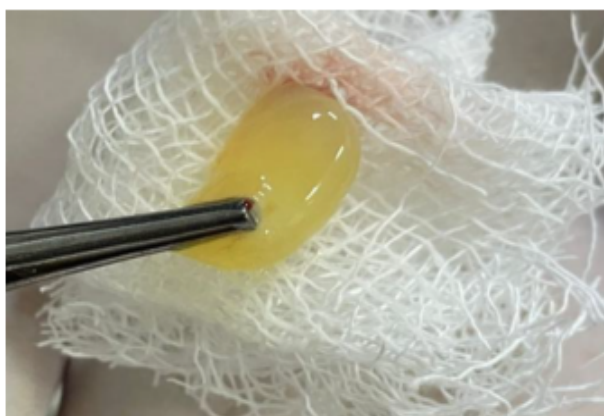


bone graft



CGF

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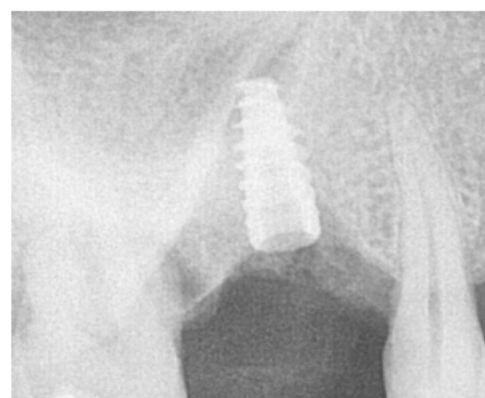


CGF CLOT

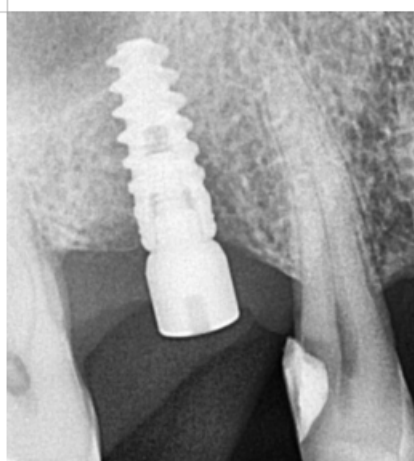


MEMBRANE

**Group A- Crestal Sinus Lift And Implant With Bone Graft
Intraoral Periapical Radiograph Of Group A**



Pre Operative 0 Month



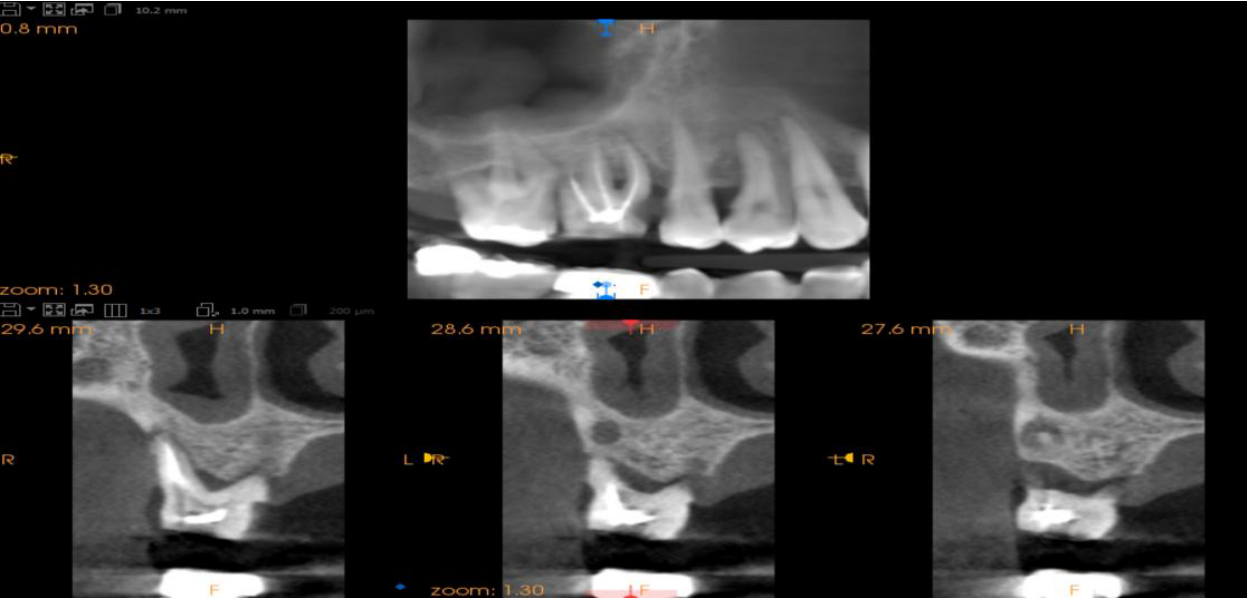
4 MONTH



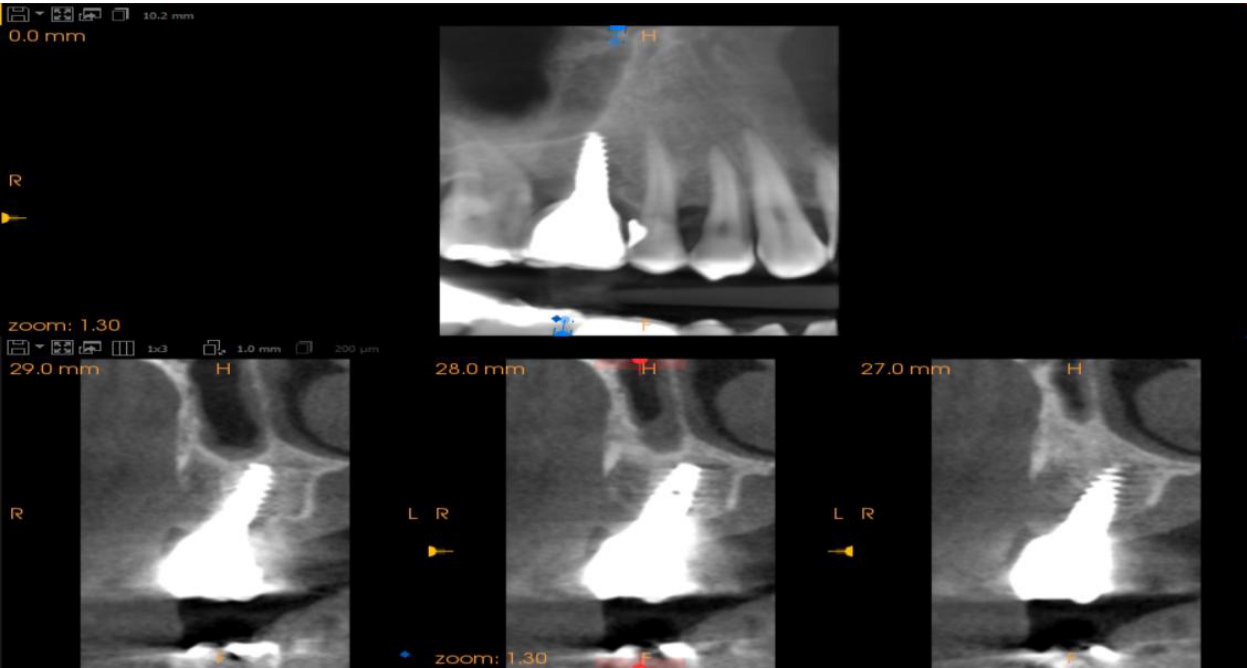
6 MONTH

CBCT

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PRE OPERATIVE



6 MONTHS POST OPERATIVE

Group B - Indirect Sinus Lift With Cgf
Intraoral Periapical Radiograph Group B

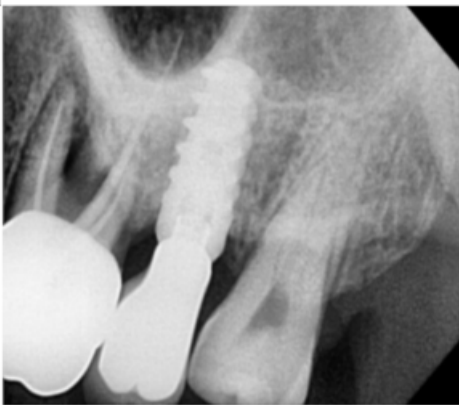
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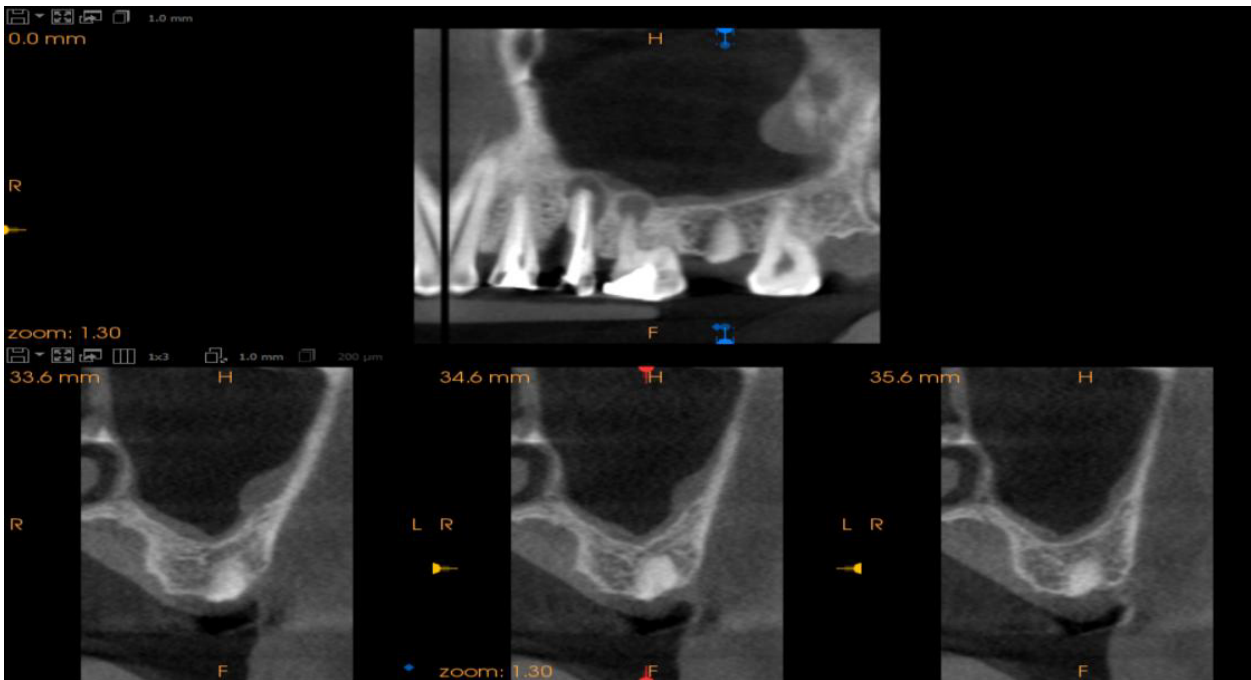
0 MONTH



4 MONTH



6 MONTH
CBCT



PRE OPERATIVE

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POST OPERATIVE

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