





Bone Graft Cements Features and Clinical Benefits



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SECTION 1



Biphasic Calcium Sulfate Overview

Calcium Sulfate

More than 100 years of documented clinical success in bone augmentation

Calcium sulfate (CS) features a unique position among all regenerative bone graft substitutes. With more than 100 years of documented clinical success, (Dreesmann 1892, Thomas and Puleo 2008) it has a longer history of clinical use than most currently available biomaterials.

Researchers and clinicians have continued to explore the effectiveness of the material in various applications. The long clinical history as an augmentation material was reported and published in thousands of scientific articles during 120 years of research, not only in dental and maxillofacial applications (Coetzee, 1980), furthermore in orthopaedic, plastic surgery, oncology, revision arthroplasty and spinal arthrodesis (Bucholz, 2002).

CS is widely recognized as a well-tolerated bone regeneration material. It undergoes virtually complete resorption *in vivo* and has consistently been considered as highly biocompatible, osteoconductive, and easy to use (Boden, 1999 or more recently by Pietrzak and Ronk, 2000; Ricci et al., 2000; Tay et al., 1999; Thomas et al., 2005; Thomas and Puleo, 2008).

The raw material, medical grade calcium sulfate is clinically used in 2 different forms differing in their individual (crystal) water content and their physicochemical behavior. (Anusavice 2003).



Biphasic Calcium Sulfate Chemical & Physical Structure

In an in vivo scenario, proteins and other biological macromolecules may further retard the setting time to up to 200 min. This is impeding the application procedure as well as the clinical performance for a typical dental application significantly (Ricci 2000).

Although Orsini and co-workers have shown that there are no differences in the bone healing pattern between preset CS dihydrate granules and moldable CShemihydrate cement. (Orsini 2004), by using preset CS dihydrate the surgeon loses advantageous handling properties of the moldable cement derivative.



By making use of the Biphasic Calcium Sulfate, the previous setting issues can be solved: The CS hemihydrate component is controlling the cement consistency and moldability characteristics. The dihydrate component is regulating the setting properties.

Using this formulation, the setting process can be reduced from about 20 minutes to 3 minutes, also under in vivo conditions in presence of blood and saliva. The key performance criteria of CS-derived graft materials that was shown in numerous studies is the fast and effective vital bone formation as a result of the rapid dissolution in vivo. (Thomas and Puleo 2008, Toloue, Dahlin) These characteristics provoke advantageous clinical results in small and well-contained bony defects like extraction socket filling. (Toloue, Guarnieri).



Bond Apatite®

Bond Apatite[®] is composed of biphasic calcium sulfate and synthetic hydroxyapatite granules. With this formulation, the concept of a graft cement can be successfully applied for a wider range of indications.

The BCS affects an injectable graft cement and features fast resorption and new bone formation. The hydroxyapatite granules provide a slow-resorbing osteoconductive scaffold to preserve the augmented volume also over a longer time period.

The BCS matrix is supposed to feature complete resorption. The slow-resorbing hydroxyapatite granule matrix serves as longer range space maintainer for optimized volume control.

Bond Apatite[®] is delivered in a specially designed ready-to-use syringe containing the granulated powder and physiological saline. Mixing the powder component with the liquid in the driver results in a viscous composite that is suitable for direct injection into the graft site.





Histological Evidence Complete Regeneration 8 Months Post-Op

At 12 weeks after placing the Bond Apatite[®], nearly 90% is replaced by the patient's own bone.

The HA particles occupying 1/3 of the composite graft are intended to slow down the overall resorption rate of the graft. The HA particles do not integrate with the bone, they are first encapsulated by connective tissue, slowly resorb, and later on, the remaining connective tissue under goes ossification.

8 Months Post-Op Complete Regeneration

3 Months Post-Op Nearly 90% Bone Regeneration



Properties and Clinical Applications of Biphasic Calcium Sulfate, Yahav et al, April 2020.



Properties and Clinical Applications of Biphasic Calcium Sulfate, Yahav et al, April 2020.

After eight months of healing time, the Bond Apatite[®] graft is almost entirely resorbed with no encapsulated materials remaining due to the complete transformation of the graft into the new vital bone.

The small and medium-sized HA particles resorb in 3-4 months. In contrast, the larger size particles, which comprise less than 10% of Bond Apatite[®], will remain for a prolonged period, almost fully resorbed after eight months, and continues its resorption process up to a complete regeneration.



SECTION 2



Features and Clinical Benefits



Biphasic Calcium Sulfate Clinical Benefits

Feature	Clinical Benefit
Osteoconductive and Bioactive	Complete bone regeneration
Prevention of infiltration of epithelio-connective cells	No membrane is required
Bacteriostatic	Can be left partially exposed
Biocompatible	Better healing and bone growth
Minimal inflammation	Following the surgical protocols leads to minimal swelling and post- operative pain to the patient
Encourages cells proliferation & angiogenesis	Promotes osteogenesis

FEATURE

Osteoconductive & Bioactive: Calcium Sulfate is entirely resorbed between 4-10 weeks and is replaced by the patient's own bone.

Calcium Sulfate is a short-term space maintainer which resorbs within up to 10 weeks. As it breaks down, Calcium Sulfate leaves behind key ingredients for stimulating bone growth. Due to its rapid absorption, Biphasic Calcium Sulfate is ideal for use in small bone defects such as socket grafting with four boney walls.

For larger defects Hydroxyapatite is mixed with Biphasic Calcium Sulfate to act as scaffolding as the new bone forms. With Bond Apatite[®], complete degradation of the Biphasic Calcium Sulfate is quickly replaced by the patient's own bone, while the volume is maintained by the different particle sizes of Hydroxyapatite that eventually resorb and replaced by the new formed bone.

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Fig. 1



#44 (28) is to be extracted.

Fig. 2



CBCT of #44 (28). We decided to graft and then to place an implant in a second step, while trying to preserve the buccal bone plate.

Fig. 3



Verifying the presence of 4 bony walls using a probe.



Application of the Biphasic Calcium Sulfate.



Fig. 5



Coverage of the exposed Biphasic Calcium Sulfate by using a collagen sponge.

Fig. 6



Periapical X-ray from the day of the operation.

Fig. 7



CBCT 4 months post-op displays bone formation and the process of its calcification.



Reentry - 4 months post-op, revealing 100% of bone regeneration.



Fig. 9



Placement of the implant in young, vascularized bone.

Fig. 10



Periapical X-ray, 5.5 months postop.

CONCLUSION

Biphasic Calcium Sulfate by itself is the bone filling material of choice for small bone defects with 3-4 bony walls.

In addition, Biphasic Calcium Sulfate forms a barrier which prevents soft tissue infiltration into the augmented site.



FEATURE

Biphasic Calcium Sulfate forms a barrier which prevents soft tissue infiltration into the material.

In traditional bone grafting, membranes are obligatory for the purpose of preventing soft tissue invagination. However, with Biphasic Calcium Sulfate the material itself acts as a graft and a barrier. Rather than penetrating the material, epithelial and connective tissue cells proliferate on the surface of the cement.

Due to the barrier effect, no membrane, or any other kind of additional barrier should be used for coverage of the cement. This has the added benefits of shorter and less invasive surgery.

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Fig. 1



#32 (23) and #34 (21) are to be extracted, #32 (23) has significant inflammatory signs.

Fig. 2



Panoramic X-ray represents the lesion in #32 (23).



Very significant buccal bone deficiency in #32 (23).

Fig. 4



Placement of a Bond Apatite[®] without membrane.



Fig. 5



The flap is stretched & sutured under tension (not tension free).

Fig. 6



CBCT 4 months post-op Shows entire bone regeneration with complete reconstruction of the buccal plate.

Fig. 7



Placement of 2 implants in a young, native bone that is well vascularized.

CONCLUSION

Bond Apatite[®] forms a barrier preventing the invagination of epithelial and connective cells. For obtaining good results, ensure the cement is well compacted. No membrane or other forms of barriers are required.



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FEATURE

Biphasic Calcium Sulfate is bacteriostatic.

Biphasic Calcium Sulfate is a salt which is known for its bacteriostatic nature. This feature enables the cement to be left partially exposed to the oral cavity.

Consequently, primary closure is not obligatory, and the cement is less susceptible for infection due to exposure.

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Fig. 1



#46 (30) needed to be extracted due to a large lesion between the roots.

Fig. 2



Extraction of #46 (30) and cleaning of the granulation tissue.

Fig. 3



The buccal plate is missing. Following protocol #2, the flap has been opened to expose the entire defect. Fig. 4



Placement of Bond Apatite[®].



Fig. 5



The flap was stretched under tension and sutured (No periosteal releasing incision or any soft tissue manipulation to gain tension free were required).

Fig. 6



Periapical X-ray with Bond Apatite[®] in place.



CBCT 3 months post-op represents a complete bone regeneration.

Fig. 8



Reentry 3 months post-op – Placement of the implant in a vascularized bone.

Fig. 9



Periapical X-ray – 3 months post implant placement.

CONCLUSION

The bacteriostatic properties of Bond Apatite[®] allow the cement to be partially exposed during flap closure with minimal risk of contamination.



FEATURE

Biphasic Calcium Sulfate is biocompatible.

Biocompatibility is a dynamic term that largely refers to the ability of a material or product to be accepted by, and function with, the human body. Calcium Sulfate is not only well tolerated by the body, but in fact works together with the body in many ways to promote optimal bone and soft tissue growth and healing.

Biphasic Calcium Sulfate stimulates the activity of osteoblasts needed for bone formation. It is completely resorbed, dissolves into components that are found in the human body and do not provoke an immune response.

Biphasic Calcium Sulfate is used without risk of a negative reaction from the body, and as a biocompatible material it aids in the healing and growth process.

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A large lesion of bone in mesial #36 (19).

Fig. 2



#36 (19) is to be extracted.



A small flap has been opened in order to visualize the bone defect.

Fig. 4



Bond Apatite® in place serves as both graft and a barrier.



Fig. 5



Experience has shown us that flap sutured under tension will not open.

Fig. 6



CBCT 8 months post-op showing very good bone regeneration with cortical bone formation.

Fig. 7



Reentry 8 months post-op shows the old defect has fully regenerated.



Bone sampling was harvested with a trephine for histological analysis.



Fig. 9



8 months post-op – Placement of the implant.



Radiographic appearance:1 year post implant placement.



Complete regeneration 8 months post-op

Fig. 11



CONCLUSION

Uneventful healing and true bone regeneration is achieved.



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FEATURE

Biphasic Calcium Sulfate provokes minimal inflammation.

Traditional GBR methods require tension free and primary closure and mostly require extensive flap reflections. Such procedures are intrusive to the patient and are largely associated with post-operative pain, swelling and hematomas.

The use of bone cement's surgical protocols require minimally invasive surgery that less traumatic to the patient with negligible post-operative complications, if any.

READ MORE

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Post-op surgery using traditional bone grafting materials.



Examples of swelling and hematoma reactions, due to invasive surgeries during traditional augmentation procedures with traditional bone grafting materials.



Bone cement surgical protocols resulting in minimal inflammatory reaction.

Fig. 1



Flap for lateral ridge augmentation is minimally reflected to expose the defect. No periosteal releasing incision or soft-tissue manipulation is used to gain tension-free. Fig. 2



Bond Apatite[®] in place.

Fig. 3



1 day post-op, no signs of hematoma or swelling.

No swelling reaction or hematoma, and less pain.

Fig. 1



Four extractions and two implants in place.

Fig. 2



Filling gaps with Bond Apatite[®].

Fig. 3



1 day post-op, no signs of hematoma or swelling.



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No traumatic experience for the patient.

Fig. 1



Tooth extraction of #36 (19) and immediate implant placement.

Fig. 2



Augmentation of the site with Bond Apatite[®].

Fig. 3



1 day post-op, no signs of hematoma or swelling.

CONCLUSION

The patient benefits from minimally invasive surgical protocols resulting in less pain, swelling and hematomas.



FEATURE

Biphasic Calcium Sulfate promotes angiogenesis.

Clinical studies done on Biphasic Calcium Sulfate feature a micro (0- 10μ m) and macro (50-500 μ m) porous structure, and an overall porosity of more than 40%. This allows for a high rate of angiogenesis, a key process in bone regeneration. Multiple studies have shown that Calcium Sulfate promotes vascularization and brings osteoblasts and growth factors into the site/grafted area.

The high rate of angiogenesis that occurs with Biphasic Calcium Sulfate is a key factor in healing process and the quick development of true, new bone.

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Fig. 1



Fig. 2



#36 (19) presents a fractured root and a large bony lesion that is needed to be extracted.



The blood clot should be preserved.

Do not aspirate the blood clot.

Fig. 5

Bond Apatite[®] in place.

Fig. 6

Flap closure with tension.

CBCT – 3 months post-op.

Fig. 8

Reentry – 3 months post-op.

Fig. 9

Periapical X-ray – 1 year post implant placement.

CONCLUSION

Calcium Sulfate chemical structure and its intra-structure porosity of more than 40% encourage growth factor infiltration, angiogenesis formation, and cell proliferation, all of which contribute to fast bone formation.

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SECTION 3

Radiographic Appearance

Radiographic Appearance with Bond Apatite[®]

The radiographic appearance after using Bond Apatite[®] is quite different from that obtained with other types of grafts. Bond Apatite[®] does not behave as traditional grafts since it does not integrate with the bone, instead it reabsorbs and transforms into the patient own bone.

This progressive transformation over time is reflected in the radiographic appearance, therefore the radiographic appearance will appear radiopaque on day 1.

As the cement resorbs it is replaced by young bone osteoid matrix, which will appear radiolucent.

As the osteoid gets calcified, the bone tissue becomes dense and then it appears radiopaque again.

With time, trabeculation and cortical plate formation will take place. The bone defect will disappear and instead we will see continuity between the patient's bone and the newly regenerated bone.

After 3-4 months, CBCT might underestimate the actual dimensions of bone formation as a new young bone is formed. This is because the bone is in the process of maturation.

Regarding the augmentation of the sinus floor by lateral approach, it is common to see clear radiological gaps after 3 months. These gaps represent immature bone that is still young and not yet calcified, however, after 5 months the bone will appear more radiopaque.

SOCKET GRAFTING

Fig. 1



Pre-op

Fig. 2



Day of surgery. Immediate post-op.

Fig. 3



1 week post-op.

Fig. 4



One month post-op.



SOCKET GRAFTING



3 months post-op.

Fig. 6



X-ray 3 months post-op – implant in place.

Fig. 7



6 months post-op.



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LATERAL AUGMENTATION

Fig. 1



CBCT – 3 months post lateral augmentation.

Note that the bone volume appears less than in the reentry (Fig. 2).

Fig. 2



Reentry – 3 months post lateral augmentation and implant placement.

Fig. 3

LATERAL AUGMENTATION



CBCT – 3 months post-op.

Fig. 4



Reentry – 3 months post-op.



- Notice the differences between the CBCT and the clinical appearance upon reopening of the site.
- In lateral augmentation, it is recommended to view the CBCT 5 months post-op where we see higher levels of bone maturation and calcification.

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OPEN SINUS LIFT



CBCT 3 months post-op.

Fig. 2



CBCT panoramic view. 3 months post-op.

Fig. 3



CBCT 6 months post-op.



- The radiolucency appearing in *Fig. 1-2*, 3 months post-op is misleading.
- In fact, the bone is in the process of calcification which can be seen upon reentry.
- Radiography after 6-months illustrates the calcification process as seen in *Fig. 3*.



OPEN SINUS LIFT



CBCT – 3 months post-op.

CBCT 3 months post-op – Panoramic view.

Fig. 6



Periapical radiographic view – 6 months post-op.





GAP FILLING

X-ray, 3 weeks after placing Bond Apatite[®] around the implant.

Fig. 1



Fig. 2



X-ray, 3 months after placing Bond Apatite[®] around the implant.

GAP FILLING

Fig. 1



Immediate implant placement in a socket.

Notice the gap between the buccal plate and the implant.

Fig. 3



X-ray, 3 weeks after placing Bond Apatite[®] around the implant.

Fig. 2



Bond Apatite® in place.

Fig. 4



X-ray, 3 months after placing Bond Apatite[®] around the implant.



SECTION 4



Surgical Bone Cement Protocols



Bone graft cement content Organized by your clinical preferences

Created for dental practitioners



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SURGICAL PROTOCOL

Socket Preservation with 4 Bony Walls

Coverage with a collagen sponge or wound dressing





Watch the protocol with a collagen sponge



Watch the protocol with a wound dressing

After extraction and complete debridement, activate and place the cement into the socket.

Place a dry, unfolded gauze above the cement and press firmly for 3 seconds, first with a finger and then with a periosteal elevator.

> Protect the exposed cement with a suitable collagen sponge or a wound dressing.



Suture and secure the collagen sponge or wound dressing in place.

EXPERT TIPS

- Press firmly for 3 seconds with a dry gauze, first with a finger and then with a periosteal elevator.
- A suitable collagen sponge a sponge that can last at least 7 days.
- The cement should be well compacted in the cervical zone, though filling the apex zone is not crucial.
- Secure the sponge in place to the surrounding soft tissue by an initial suture and thereafter with a cross-stitch.



CASE #1 – Coverage with a collagen sponge

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS





#16 (3) is to be extracted.

Fig. 2



X-ray of #16 (3)'s large roots.



After extraction, the height of the keratinized gingiva is important.

Fig. 4



Bond Apatite® in place.



CASE #1 – Coverage with a collagen sponge

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 5



Collagen sponge sutured and secured in place, over Bond Apatite[®].

Fig. 6



2 weeks post-op.



4 weeks post-op – soft tissue appearance.

Fig. 8



4 months post-op – Implant placement.



CASE #2 – Coverage with a collagen sponge

CLINICAL EVIDENCE	SOCKET GRAFTING WITH 4 BONY WALLS
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Pre-op radiography.

Fig. 2



Pre-op.



Extract the tooth and prepare the socket for grafting.

Fig. 4



Bond Apatite® in place.



CASE #2 – Coverage with a collagen sponge

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 5



Protect the cement by covering it with a collagen plug or sponge.

Fig. 6



Secure the sponge in place to the surrounding soft tissue by an initial suture, thereafter with a cross suturing above.

Fig. 7



Due to exposure of more than 3mm, the cement is covered with a collagen sponge.



3 months post-op – soft tissue appearance.



CASE #2 – Coverage with a collagen sponge

CLINICAL EVIDENCE SOCKET GRA

SOCKET GRAFTING WITH 4 BONY WALLS



X-ray – 3 months post-op.



Reentry – 3 months post-op.



CASE #3 – Coverage with a collagen sponge

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS



Pre-op radiography.

Fig. 2



Pre-op.

Fig. 3



Extract the teeth and prepare the site for grafting.

Fig. 4



Bond Apatite® in place.



CASE #3 – Coverage with a collagen sponge

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 5



Collagen sponge secured in place.

Fig. 6



CBCT 4 months post-op.





CBCT 4 months post-op.



Reentry 4 months post-op.



CASE #3 – Coverage with a collagen sponge

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 9



Implants in place.

Fig. 10



Implants in place.



CASE #4 – Coverage with a wound dressing

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 1



#36 (19) needed to be extracted.

Fig. 2



Radiographic appearance before extraction.

Fig. 3



After extraction, we verified the presence of the 4 bony walls.



Bond Apatite® in place.



CASE #4 – Coverage with a wound dressing

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 5



The Bond Apatite[®] is covered with the wound dressing.

Fig. 6



Sutures over the wound dressing to hold it in place.



1 week post-op. Notice the rapid proliferation of soft tissue.

Fig. 8



2 weeks post-op – soft tissue appearance.



CASE #4 – Coverage with a wound dressing

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 9



CBCT 3 months post-op showing a true bone regeneration.

Fig. 10



Reentry - 3 months post op. Implant placement in a young, vascularized bone.

Fig. 11



Immediate X-ray post implant placement.



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CASE #5 – Coverage with a wound dressing

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS



#16 (3) with fractured root and large lesion is needed to be extracted.

Fig. 2



The height of keratinized gingiva is a reason for not advancing the flap, leaving the Bond Apatite[®] exposed and protected by the wound dressing until soft tissue proliferation takes place.

Fig. 3

Fig. 1



Socket after extraction.



Bond Apatite[®] in place, before removing excess.



CASE #5 – Coverage with a wound dressing

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS





The wound dressing sticks above the soft tissue.

Fig. 6



Sutures over the wound dressing.

Fig. 7



1 week post-op.

Fig. 8



2 weeks post-op.



CASE #5 – Coverage with a wound dressing

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 9



3 months post-op – soft tissue appearance.

Fig. 10



CBCT 3 months post-op – Bone regeneration can be seen.

Fig. 11



Reentry 3 months post-op. Well vascularized new bone can be seen with a few particles of HA on the surface.



X-ray 3 months post implant placement.



CASE #6 – Coverage with a wound dressing

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS



#21 (9), #22 (10), #23 (11), #24 (12), and #25 (13) are to be extracted.

Fig. 2



All extractions will take place at the same time.

Fig. 3

Fig. 1



Extractions performed.



Implants in place. Primary closure could not be achieved.



CASE #6 – Coverage with a wound dressing

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 5



Bond Apatite[®] in place.

Fig. 6



Some of the Bond Apatite[®] was not covered.

Fig. 7



Placing of the wound dressing after cleaning the mucosa with a dry gauze followed by a moist gauze.



Sutures of the wound dressing in an X to stabilize it in place.



CASE #6 – Coverage with a wound dressing

CLINICAL EVIDENCE

SOCKET GRAFTING WITH 4 BONY WALLS

Fig. 9



Fast healing after one week.

Fig. 10



3 months post-op – showing perfect healing.



More than 3mm of exposed cement should be protected and secured by a collagen sponge or a wound dressing.



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LEARN MORE

SOCKET GRAFTING WITH 4 BONY WALLS





SURGICAL PROTOCOL

Socket Preservation without 4 Bony Walls

STEPS



Watch the video

Perform mesial oblique vertical incision (up to 2-3 mm beyond the MGJ into the mobile mucosa).

Place the cement.

2 Raise full thickness flap, minimally as needed to expose the entire defect.

Clean the socket.

- Press firmly for 3 seconds, with a finger above a dry gauze.
- Then exert additional pressure for few seconds on the gauze with a periosteal elevator.
- The material should be well compacted from all directions, buccally and occlusally.

No soft tissue manipulation should be performed to gain a tension free flap. The flap is stretched for closure. Therefore, no periosteal releasing incision

Suturing: Start from mesial, then distal, then in the middle, after that sutures can be placed in between.

Reposition the flap for
maximal closure with tension.

Suture.



CASE #1 CLINICAL EVIDENCE

SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 1



Radiographic image of #46 (30).

Fig. 2



#46 (30) is to be extracted.

Fig. 3

Pre-op



According to the protocol, after minimal flap reflection a missing buccal plate can be noticed.

Fig. 4



Bond Apatite[®] in placed, after pressing with the dry gauze.



CASE #1 CLINICAL EVIDENCE

SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 5



Bond Apatite[®] begins to absorb the surrounding blood. Reapply pressure with a sterile gauze.

Fig. 6



First suture the mesial corner of the flap, the flap is sutured with tension by stretching.

Fig. 7



Then sutures according to the sequence, mesial, distal, middle, in between, and finally the vertical incision is sutured.



CBCT 3 months post-op. Complete bone formation.



CASE #1 CLINICAL EVIDENCE

SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 9



3 months post-op.Soft tissue healing appearance.

Fig. 10



3 months post-op.

Placement of the implant in young, vascularized bone.

Fig. 11



Periapical X-ray, 6 months from the day of augmentation.



Keratinized gingiva around the implant.



CASE #2 CLINICAL EVIDENCE

SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 1



#45 (29) and #47 (31) are to be extracted.

Fig. 2



Panoramic X-ray showing the bone deficiency in #47 (31).

Fig. 3



The buccal plates of #45 (29) and #47 (31) are partially missing.

Fig. 4



Bond Apatite[®] in place after complete debridement in #47 (31).



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CASE #2 CLINICAL EVIDENCE

SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 5



Flap stretched for closure with tension, according to the surgical protocol.

Fig. 6



CBCT 4 months post-op - #45 (29), showing total bone regeneration.

Fig. 7



CBCT 4 months post-op - #47 (31).

Fig. 8



4 months post-op – Placement of 3 implants in a new regenerated bone.



CASE #3 CLINICAL EVIDENCE

SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 1



Tooth #25 (13) will be extracted.

Fig. 2



8mm bone deficiency can be seen in the palatine wall.

Fig. 3



Bond Apatite[®] in place.



Soft tissue sutures with tension.


SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 5



6 months post-op. Total bone regeneration with palatal cortex.

Fig. 6



6 months post-op. Placement of the implant.

Fig. 7



3 years post-op. Periapical X-ray.



In this clinical case we traditionally would have had to make a palatal flap and use a membrane, but instead, the simple application of Bond Apatite[®] was highly effective to regenerate the bone, with the formation of a palatal cortex.



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SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 1



#35 (20) and 36 (19) are to be extracted.

Fig. 2



X-ray shows a bony lesion in both #35 (20) and #36 (19).

Fig. 3



After extracting we found that parts of the buccal bone walls are missing.



Oblique mesial vertical incision to expose the defect.



SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 5



Reflection of a small flap of full thickness revealed the extent of the bone deficiency.

Fig. 6



Bond Apatite® in place.



Flap under tension is sutured with Rapid PGA.

Fig. 8







SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 9



Reentry 3 months post-op. Perfect regeneration of the bone defect can be seen.

Fig. 10



Placement of 3 implants.



Since a tension free flap is not indicated, in cases where multiple teeth are being extracted a single vertical releasing incision is sufficient to allow the flap to cover the grafted sites by stretching.



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SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 1



#37 (18) to be extracted.

Fig. 2



Probing after extraction to assess the extent of the bone defect.

Fig. 3



Small flap is reflected to reveal the extent of the buccal bone deficiency.



Bond Apatite® in place.



SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 5



Sutures, flap closure with tension.

Fig. 6



CBCT showing the perfect regeneration of the buccal plate.

Fig. 7



3 months post-op. At this stage, some HA granules can be seen and are in the process of resorption.

Placement of the implant.



- The vertical releasing incision should be in a distance of at least 3-5mm from the bone defect and should allow visualization of the entire missing buccal plate.
- The use of Bond Apatite[®] allowed bone regeneration, with cortical formation, in a defect where the buccal plate was absent.
- Following the protocols ensures high predictability and clinical success.



SOCKET GRAFTING WITHOUT 4 BONY WALLS



#37(18) – needed to be extracted.

Fig. 2



Large caries under the crown of #37 (18).



After socket debridement, do not aspirate the blood clot before grafting.

Fig. 4



A minimal vertical incision and no membrane is required.



SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 5



Maximal closure (3mm of graft exposure is acceptable).

Fig. 6



X-ray – Day of surgery.

Fig. 7



CBCT – 3 months post-op.

Fig. 8



Reentry – 3 months post-op.

A true vital bone with small crystals of hydroxyapatite that are in the process of resorption.



SOCKET GRAFTING WITHOUT 4 BONY WALLS

Fig. 9



Placement of the implant 3 months post-op.

Fig. 10



18 months post-op.

Very stable and satisfactory results.



- The possibility of suturing the flap under tension enables us to respect the stability of the material and to limit inflammatory reactions.
- When a buccal wall is missing, a small incision and a full thickness flap sutured with tension will give very good results when using Bond Apatite[®].



LEARN MORE

SOCKET GRAFTING WITHOUT 4 BONY WALLS

INTRUCTIONAL VIDEO



ANIMATED PROTOCOL



ON-DEMAND WEBINAR





SURGICAL PROTOCOL

Lateral Ridge Augmentation

STEPS



Watch the video

Minimally reflect the flap, enough to expose the entire grafted site.

(After placing the cement) Press with a dry gauze, first with a finger and then with a periosteal elevator.

B Reposition the flap by stretching it directly above the cement for maximal closure.



- Perform a short vertical incision at a distance from the grafted site. That vertical incision should not pass the mucogingival junction by more than 2-3 mm. As well during undermining with the periosteal elevator in order to reflect the flap, it should not go deeper than 2-3 mm into the mobile mucosa.
- It is recommended to do decortication before placement of the cement.
- Suture first the mesial corner, then distal, then in the middle. After these first steps, sutures can be placed in between and finally for the vertical releasing incision.



LATERAL RIDGE AUGMENTATION

Fig. 1



Thin ridge, #24 (12) - 2mm thick.

Fig. 2



#24 (12) was extracted few years earlier without preserving the socket. Therefore, the bone is very narrow.

Fig. 3



During a full thickness flap reflection, It is necessary to visualize the entire area.



Placement of Bond Apatite[®], augmenting the area of #22 (10).



LATERAL RIDGE AUGMENTATION

Fig. 5



Closure of the flap with tension.

Fig. 6



1 day post-op - No sign of inflammation.



CBCT 3 months post-op.

Fig. 8



X-ray 3 months post implant placement.

LATERAL RIDGE AUGMENTATION

Fig. 9



Notice the thickness of the soft tissue.



To further improve the result, it is preferable to do decortication before grafting.



LATERAL RIDGE AUGMENTATION

Fig. 1



Old and unstable bridge from #13 (6) to #23 (11) needed to be replaced.

Fig. 2



It was decided to keep #13 (6), #22 (10) and #23 (11).

Fig. 3



The treatment plan was to place implants in #12 (7) and #21 (9).

Fig. 4



After flap reflection, a narrow ridge can be seen near #12 (7). Decortication was performed before augmentation.



LATERAL RIDGE AUGMENTATION

Fig. 5



Bond Apatite® in place.

Fig. 6



Sutures and collagen sponge covering the Bond Apatite[®] in #11 (8) and #21 (9).



Temporary bridge placed on the day of the operation.

Fig. 8



2 weeks post-op - Soft tissue appearance.



LATERAL RIDGE AUGMENTATION

Fig. 9



CBCT - 4 months post-op.



Fixed prosthetics is essential to prevent graft movement.



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LATERAL RIDGE AUGMENTATION

Fig. 1



Buccal collapse in #14 (5) which was extracted without socket grafting.

Fig. 2



CBCT confirming a narrow ridge.



Placement of a 3/16 implant and visualization of the area to be grafted.

Fig. 4



Bond Apatite[®] in place, forming a compact block.



LATERAL RIDGE AUGMENTATION

Fig. 5



Sutures.

Fig. 6



4 months post-op – sufficient bone width is achieved.

Fig. 7



4 months post-op.



In cases of simultaneous implant placement with augmentation procedure, it is necessary to ensure the primary stability of the implant.



LATERAL RIDGE AUGMENTATION



CBCT - Pre-op of #14 (5) and #15 (4).

Fig. 2



Decortication of the buccal wall.



Bond Apatite® in place.

Fig. 4



4 weeks post-op – soft tissue healing appearance.



LATERAL RIDGE AUGMENTATION

Fig. 5



Reentry 4 months post-op.

Fig. 6



CBCT 4 months post-op.



- Graft stability during the healing phase is crucial.
- Do not use removable temporary prostheses.
- Instruct the patient to apply a soft diet for avoiding pressure in the area.



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LATERAL RIDGE AUGMENTATION



CBCT pre-op - #22 - #25 (33-41)

Fig. 2



CBCT - #22 - #25 (33-41) represents a narrow crest.

Fig. 3



Clinical appearance after flap reflection.



Bond Apatite® in place.



LATERAL RIDGE AUGMENTATION

Fig. 5



Sutures

Fig. 6



Reopening - 4 months-post-op and osteotomy.

Fig. 7



CBCT – 4 months post-op.



- According to the protocol, graft stabilization is achieved by minimal flap reflection.
- Perform the *predictability test* by placing your finger in the vestibule and vibrating it to ascertain that the sutures don't move at all.
- Following the protocol prevents flap movement due to the traction of the muscles.



LEARN MORE

INTRUCTIONAL VIDEO



ANIMATED PROTOCOL



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SURGICAL PROTOCOL

Closed Sinus Lift/Intra-Crestal Approach



- Reload the particles into a Bond Apatite[®] syringe, or any other bone carrier syringe.
- Introduce the cement, then push with an osteotome.



CLOSED SINUS LIFT



#25 (13) and #27 (15) will be extracted due to a significant infection resulting in increased bone resorption.

Fig. 2



Identical situation in #27 (15). The goal is to obtain 3mm-4mm of bone.

Fig. 3

Fig. 1



Placement of Bond Apatite[®] after extraction of #25 (13), #27 (15).



3 months post-op – soft tissue appearance.



CLOSED SINUS LIFT



CBCT 3 months post-op of #27 (15), we obtained our 4mm of bone.

Fig. 6



CBCT of #25 (13). We can consider a Closed Sinus Lift.

Fig. 7



Reentry 3 months post-op.



Osteotomies were performed to reach up to the sinus floor.



CLOSED SINUS LIFT

Fig. 9



Placement of Bond Apatite[®].

Fig. 10



Pushing of material using the osteotome.

Fig. 11



Placement of 2 implants, 5 by 10mm.









CASE #1



- We went from a situation where we only had 1mm of bone and we ended up with 4mm below the sinus floor. This allowed us to perform a closed sinus lift procedure.
- The characteristics of Bond Apatite[®] reduce the risk of perforating the membrane due to the soft texture of the cement.
- Moreover, Bond Apatite[®] enabled us to achieve true bone regeneration in the sinus and around the implants.



CLOSED SINUS LIFT

Fig. 1



Pre-op – augmentation is required in teeth #14 (5) - #16 (3).

Fig. 2



CBCT indicates we have 4mm of bone which is required to perform an intra-crestal sinus lift (Summers' technique).

Fig. 3



#15 (4) - #16 (3) - two implant osteotomies reach up to the sinus floor.



Bond Apatite[®] is used to augment the sinus.



CLOSED SINUS LIFT

Fig. 5



Implant in place after performing the Summers' technique in #15 (4) - #16 (3).

Fig. 6



Periapical X-ray – immediate postop.

Fig. 7



Soft tissue closures & sutures.

Fig. 8



3 months post-op – soft tissue appearance.



CLOSED SINUS LIFT

Fig. 9



preserving the keratinized gingiva during placement of the healing abutments.

Fig. 10



X-ray – 1 year post-op.



- Radiographic appearance at the time of graft placement appear radiopaque.
- 3 weeks post-op the appearance is radiolucent.
- 3-4 months post-op the appearance will again become radiopaque.
- 1-year post-op X-ray demonstrates the bone gained in the augmentation procedure.



CLOSED SINUS LIFT



Pre-op CBCT of the area #16 (3) - #17 (2).

Fig. 2



Pre-op.



Pushing the Schneiderian membrane with an osteotome.

Fig. 4



Applying Bond Apatite[®] in a sterile dish and waiting for 3 minutes.



CLOSED SINUS LIFT

Fig. 5



Beginning Summers' technique sinus lift.

Fig. 6



Filling the osteotomy with Bond Apatite[®].

Fig. 7



2 implants in place.









CLOSED SINUS LIFT

Fig. 9



X-ray – 1 year post-op.



While filling the osteotomy:

- Inject the Bond Apatite[®] particles.
- Push the particles with an osteotome.
- Add additional layer of Bond Apatite[®] as needed and push again with an osteotome to elevate the membrane.
- Insert the implant into the osteotomy.


LEARN MORE

Closed Sinus Lift/Intra-Crestal Approach

INTRUCTIONAL VIDEO



ON-DEMAND WEBINAR

LIVE SURGERY





SURGICAL PROTOCOL

Open Sinus Lift/Lateral Window Approach

STEPS



Watch the video

Prepare your access window and elevate the sinus membrane.

Activate the syringe and wait one minute.

EXPERT TIPS

During ejection, the cement should be ejected through the crest. Then, compact it through the crest with a periosteal elevator above a dry gauze.



Repeat the process until 2/3 of the sinus is filled.

Eject the cement into the sinus cavity.

- Repeat this operation in the mesial, distal and the middle area until 2/3 of the sinus is filled.
- The final syringe should be injected directly after activation into the sinus cavity.
- Immediately after, press firmly with a dry sterile gauze and close the access window.
- No membrane is required.

Activate a new syringe to fill the rest of the sinus.

Suture.



CASE #1 CLINICAL EVIDENCE

OPEN SINUS LIFT



Pre-op

CBCT indicates the level of the left sinus floor.

Fig. 2



Opening the lateral access window.

Fig. 3



One minute post activating the syringe, applying Bond Apatite[®] in the sinus.

Fig. 4



Beginning to fill up the sinus floor.



CASE #1 CLINICAL EVIDENCE

OPEN SINUS LIFT

Fig. 5



After filling of the sinus according the protocol, we use Bond Apatite[®] to close the window.

Fig. 6



Soft tissue closure & sutures.

Fig. 7



CBCT – 4 months post-op.

Fig. 8



CBCT – 4 months post-op.



CASE #1 CLINICAL EVIDENCE

OPEN SINUS LIFT

Fig. 9



X-ray – 3 months post placing implants.



- For closing the sinus window, the last syringe is injected immediately after its activation.
- Notice the bone volume differences between CBCT 4 months post-op and CBCT 3 months post placing the implants (the bone is in a continuous process of maturation).



CASE #2 CLINICAL EVIDENCE

OPEN SINUS LIFT



Pre-op.

Panoramic X-ray shows the left sinus floor.

Fig. 2



Pre-op.

Fig. 3



Opening the lateral access window.

Fig. 4



Filling up the sinus floor with Bond Apatite[®] according to the protocol.



Fig. 1

CASE #2 CLINICAL EVIDENCE

OPEN SINUS LIFT

Fig. 5



Closing the lateral window floor with Bond Apatite $\ensuremath{^\mathbb{R}}$.

Fig. 6



Healing – 2 weeks post-op.

Fig. 7



Reentry – 4 months post-op. The window is closed with the patient's own bone.

Fig. 8



Placing 3 implants.



CASE #2 CLINICAL EVIDENCE

OPEN SINUS LIFT

Fig. 9



CBCT – 3 months post-op.



When drilling in the site 3-4 months post-op, you may feel the bone is soft. This is because the bone continues its calcification process.

Therefore, it is recommended to place implants 5 months post augmentation.



LEARN MORE

Open Sinus Lift/Lateral Window Approach

INTRUCTIONAL VIDEO



ON-DEMAND WEBINAR

ANIMATED PROTOCOL



LIVE SURGERY





Clinical cases by David Baranes, DMD

Meet the Expert



David Baranes, DMD

- Graduated from the Doctorate program in Dental Surgery at the University of Paris 7 in 1987.
- Completed a C.E.S biomaterials applied to dentistry.
- C.E.S in Periodontology at the University of Paris 7.
- Completed a post-graduate course in implantology (A.U.I) at the PARIS 7 Faculty.
- Since 1991 working exclusively in periodontal and basal implantology in Jerusalem and Netanya.
- Fellowship and DIPLOMAT status in 2015 with the INTERNATIONAL CONGRESS of ORAL IMPLANTOLOGIST.
- International speaker with focus on clinical use of cements for bone regeneration.
- Published internationally in professional journals.





References

Dreesmann, H. über Knochenplombierung. Klinische Chirurgie, 1892;9:804-810.

Thomas, M.V., Puleo, D.A., Calcium sulfate: A review. J. Long Term Eff. Med. Implants,

2005;15(69):599-607. b) Thomas, M.V., Puleo, D.A., Calcium sulfate: Properties and clinical applications. J. Biomed. Mater. Res. B Appl. Biomater., 2008 Nov 24. [Epub ahead of print]

Peltier, L.F., The use of plaster of Paris to fill defects in bone. Clin. Orthop. Res., 1961;21:1-29.

Peltier, L.F., Speer, D.P., Calcium Sulfate. In: Habal M. B., Reddi A.H. (eds) Bone grafts and bone substitutes, Saunders Co., Philadelphia, 1981:243-246.

Coetzee, A.S., Regeneration of bone in the presence of calcium sulfate. Arch. Otolaryngol., 1980;106:405-409.

Bucholz, RW. Nonallograft osteoconductive bone graft substitutes. Clin Orthop Relat Res., 2002;395:44-52.

Boden SD, Stevenson S. Bone grafting and bone graft substitutes. Philadelphia: Saunders, 1999.

Pietrzak, W.S., Ronk, R., Calcium sulfate bone void filler: A review and a look ahead. J.Craniofac. Surg., 2000;11(4):327-333.

Ricci JL, Alexander H, Nadkarni P, Hawkins M, Turner J, Rosenblum SF, Brezenoff L, De Leonardis D, Pecora G. Biological mechanisms of calcium- sulfate replacement by bone. In:

Davies JE , editor. Bone Engineering. Toronto: Em Squared Inc., 2000:332-344. b)Ricci, J.L.,

Report on the use of calcium sulfate cement with dental cement. In: Use of Calcium Sulfate

Cement for Dental and maxillofacial Bone Repair, edited by Pecora, G. and De Leonardis, D., Elite Publishing, Rome, 7 pages, 2001.

Tay, B.K.B., Patel V.V., Bradford, D.S., Calcium sulfate- and calcium phosphate-based bone substitutes: Mimicry of the mineral phase of bone. Orthop. Clin. North Am., 1999;30(4):615-623.

Anusavice KJ. Gypsum products. In: Anusavice KJ, editor. Phillips' Science of Dental Materials.

St. Louis, MO: Saunders, 2003:255-281.

Strocchi, R., Orsini, G., Iezzi, G., Scarano, A., Rubini, C., Pecora, G. & Piatelli, A. Bone regeneration with calcium sulphate: evidence for increased angiogenesis in rabbits. The Journal of Oral Implantology, 2002;28:273–278.

Turri A, Dahlin C. Comparative maxillary bone-defect healing by calcium-sulphate or deproteinized bovine bone particles and extra cellular matrix membranes in a guided bone regeneration setting: an experimental study in rabbits. Clin. Oral Impl. Res. 00, 2014, 1–6 doi:10.1111/clr.12425

Toloue SM, Chesnoiu-Matei I, Blanchard SB. A clinical and histomorphometric study of calcium sulfate compared with freeze-dried bone allograft for alveolar ridge preservation. J Periodontol, 2012;83:847–855.

Collins JR, Jiménez E, Martínez C, Polanco RT, Hirata R, Mousa R, Coelho PG, Bonfante EA,

Tovar N.. Clinical and Histological Evaluation of Socket Grafting Using Different Types of Bone Substitute in Adult Patients. Implant Dent, 2014;23:489–495.

Payne JM, Cobb CM, Rapley JW, Killoy WJ, Spencer P. Migration of human gingival fibroblasts over guided tissue regeneration barrier materials. J Periodontol, 1996;67:236-244.



References

Machtei EE, Mayer Y, Horwitz J, Zigdon-Giladi H. Prospective randomized controlled clinical trial to compare hard tissue changes following socket preservation using alloplasts, xenografts vs no grafting: Clinical and histological findings. Clin Implant Dent Relat Res. 2019;21:14–20. https://doi.org/10.1111/cid.12707

Walsh, W. R. PhD*; Morberg, P. MD, PhD*; Yu, Y. PhD*; Yang, J. L. PhD*; Haggard, W. PhD**; Sheath, P. C. MBBS*; Svehla, M. PhD*; Bruce, W. J. M. MBBS† Response of a Calcium Sulfate Bone Graft Substitute in a Confined Cancellous Defect, Clinical Orthopaedics and Related Research: January 2003 - Volume 406 - Issue 1 - p 228-236

Yahav A, Kurtzman GM, Katzap M, Dudek D, Baranes D. Bone Regeneration: Properties and Clinical Applications of Biphasic Calcium Sulfate. Dent Clin North Am. 2020 Apr;64(2):453-472. doi: 10.1016/j.cden.2019.12.006. Epub 2020 Jan 18. PMID: 32111280.

Mukherji A, Rath SK. Calcium sulfate in periodontics: A time tested versatile alloplast. J Sci Soc [serial online] 2016 [cited 2021 Feb 14];43:18-23. Available from: <u>https://www.jscisociety.com/text.asp?2016/43/1/18/175447</u>

Li Shue, Zhang Yufeng & Ullas Mony (2012) Biomaterials for periodontal regeneration, Biomatter, 2:4, 271-277, DOI: 10.4161/biom.22948

Lombardo G, Corrocher G, Rovera A, Pighi J, Marincola M, Lehrberg J, Nocini PF, "Decontamination Using a Desiccant with Air Powder Abrasion Followed by Biphasic Calcium Sulfate Grafting: A New Treatment for Peri-Implantitis", Case Reports in Dentistry, vol. 2015, Article ID 474839, 7 pages, 2015. https://doi.org/10.1155/2015/474839

Lombardo G, Marincola M, Cicconetti A, Simancas-Pallares MA, Pighi J, Lehrberg J, Signoriello A, Corrocher G, Serpa-Romero X, Vila Sierra LA, Arevalo-Tovar L, Nocini PF. Successful Management of Peri-Implantitis around Short and Ultrashort Single-Crown Implants: A Case Series with a 3-Year Follow-Up. Int J Dent. 2019 Sep 15;2019:5302752. doi: 10.1155/2019/5302752. PMID: 31636671; PMCID: PMC6766094.

Baranes D, Kurtzman GM; Biphasic Calcium Sulfate as an Alternative Grafting Material in Various Dental Applications. J Oral Implantol 1 June 2019; 45 (3): 247–255. doi: <u>https://doi.org/10.1563/aaid-joi-D-18-00306</u>

Sinjab YH, Sinjab KH, Navarrete-Bedoya C, Gutmann JL. Calcium sulfate applications in dentistry: A literature review. Endodontology [serial online] 2020 [cited 2021 Feb 14];32:167-74. Available from: <u>https://www.endodontologyonweb.org/text.asp?2020/32/4/167/307310</u>

Thomas MV, Puleo DA, Al-Sabbagh M. Calcium sulfate: a review. J Long Term Eff Med Implants. 2005;15(6):599-607. doi: 10.1615/jlongtermeffmedimplants.v15.i6.30. PMID: 16393128.

Thomas MV, Puleo DA. Calcium sulfate: Properties and clinical applications. J Biomed Mater Res B Appl Biomater. 2009 Feb;88(2):597-610. doi: 10.1002/jbm.b.31269. PMID: 19025981.

Nilsson M, Zheng MH, Tägil M. The composite of hydroxyapatite and calcium sulphate: a review of preclinical evaluation and clinical applications. Expert Rev Med Devices. 2013 Sep;10(5):675-84. doi: 10.1586/17434440.2013.827529. Erratum in: Expert Rev Med Devices. 2013 Nov;10(6):857. PMID: 24053255.



References

Damian D, Edyta RW, Gregori KM, Lanka M. The use of biphasic calcium sulfate (Bond Apatite®) for surgical treatment of osseous defects resulting from radicular cysts – Clinical study of 6 months follow-up. J Int Clin Dent Res Organ [serial online] 2020 [cited 2021 Feb 14];12:55-61. Available from: https://www.jicdro.org/text.asp?2020/12/1/55/291106

Dasmah A, Sennerby L, Rasmusson L, Hallman M. Intramembraneous bone tissue responses to calcium sulfate: an experimental study in the rabbit maxilla. Clin Oral Implants Res. 2011 Dec;22(12):1404-8. doi: 10.1111/j.1600-0501.2010.02129.x. Epub 2011 Mar 21. PMID: 21435007.

Machtei EE, Mayer Y, Horwitz J, Zigdon-Giladi H. Prospective randomized controlled clinical trial to compare hard tissue changes following socket preservation using alloplasts, xenografts vs no grafting: Clinical and histological findings. Clin Implant Dent Relat Res. 2019 Feb;21(1):14-20. doi: 10.1111/cid.12707. Epub 2018 Dec 28. PMID: 30592368.

Hughes E, Yanni T, Jamshidi P, & Grover L. M. (2015) Inorganic cements for biomedical application: calcium phosphate, calcium sulphate and calcium silicate, Advances in Applied Ceramics, 114:2, 6 https://doi.org/10.1179/1743676114Y.00000002195-76, DOI: 10.1179/1743676114Y.0000000219

Walsh WR, Morberg P, Yu Y, Yang JL, Haggard W, Sheath PC, Svehla M, Bruce WJ. Response of a calcium sulfate bone graft substitute in a confined cancellous defect. Clin Orthop., 2003:228-236.

Stubbs D, Deakin M, Chapman-Sheath P, Bruce W, Debes J, Gillies RM, Walsh WR, In vivo

evaluation of resorbable bone graft substitutes in a rabbit tibial defect mode, Biomaterials, 2004; 25(20): 5037-44.

Urban, R.M., Turner, T.M., Hall, D.J., Inoue, N., Gitelis, S., Increased bone formation using calcium sulfate-calcium phosphate composite graft. Clin. Orthop. Relat. Res., 2007;459:110-117.

Orsini G,Ricci J, Scarano A, Pecora G, Petrone G, Lezzi G, Piattelli A. Bone-defect healing with calcium-sulfate particles and cement: An experimental study in rabbit. J Biomed Mater Res B

Appl Biomater, 2004;68:199-208. a) Guarnieri R, Grassi R, Ripari M, Pecora G. Maxillary sinus augmentation using granular calcium sulfate (surgiplaster sinus): Radiographic and histologic study at 2 years. Int J Periodontics

Restorative Dent, 2006;26:79-85. b) Guarnieri R,Bovi M. Maxillary sinus augmentation using prehardened calcium sulfate: A case report. Int J Periodontics Restorative Dent, 2002;22:503-508



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