



ISSN: 0975-833X

Available online at <http://www.ijournalcra.com>

International Journal of Current Research
Vol. 12, Issue, 05, pp.11369-11372, May, 2020

DOI: <https://doi.org/10.24941/ijcr.38636.05.2020>

INTERNATIONAL JOURNAL
OF CURRENT RESEARCH

RESEARCH ARTICLE

BIOCERAMICS- BOON IN ENDODONTICS

¹Shivani Parmar and ^{2,*}Ankita Sharma

¹MDS Prosthodontics and Crown & Bridge

²Junior Resident, Department of Pathology, IGMC, Shimla

ARTICLE INFO

Article History:

Received 08th February, 2020

Received in revised form

24th March, 2020

Accepted 18th April, 2020

Published online 30th May, 2020

Key Words:

Bioceramics, Bioactive glass, calcium phosphate, calcium silicate, hydroxyapatite.

Copyright © 2020, Shivani Parmar and Ankita Sharma. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Citation: Shivani Parmar and Ankita Sharma. 2020. "Bioceramics- Boon in Endodontics", *International Journal of Current Research*, 12, (5), 11369-11372.

ABSTRACT

Bioceramics are materials which include alumina, zirconia, bioactive glass, glass ceramics, hydroxyapatite, resorbable calcium phosphates. They have been used in dentistry as root repair materials, bony defects repair, apical filling material, perforation repair and help in tissue regeneration. They have certain advantages like biocompatibility, non toxicity, dimensional stability and bio-inertness. They have a structural similarity to Hydroxyapatite, an intrinsic osteo conductive activity and have an ability to induce regenerative responses in the human body. This review focuses on an overview of Bioceramics, classification and their advantages and also gave a detailed insight into individual bioceramic materials currently used in the fields of endodontics.

INTRODUCTION

With the introduction of new techniques and technology the field of endodontics is widely changing. Various advancements in endodontic materials has led to significant growth in endodontics. As with the introduction of Bio-ceramics the prognosis and treatment outcome of certain procedures have totally revolutionised in endodontics. Bioceramics are chemically processed biocompatible ceramic compounds which can be obtained both in situ and in vivo. Bioceramics are quite similar to biological hydroxyapatite thus exhibit excellent biocompatibility properties. Different bioceramic products are obtained during the hydration process e.g. hydroxyapatites, they have the ability to induce a regenerative response in the human body. Mineral hydroxyapatite has an osteoconductive effect, which leads to the bone formation at the interface. Bioceramics have an intrinsic osteoinductive capacity, because of documented ability to absorb osteoinductive substance.¹ Bioceramics have antibacterial properties, as a result of precipitation they form porous powders containing nanocrystals with the diameter of 1–3 nm, which prevent bacterial adhesion.² In addition, bioceramics can be combined with synthetic hydroxyapatite, and might constitute of fluoride ions incorporated in apatite crystals resulting in antibacterial properties.³ The aim of this paper is to perform a literature review on bioceramic materials currently used in endodontics and on their various characteristics.

Classification of Bioceramics:

Bioinert: These are non-interactive with the biological systems (Alumina, zirconia).

Bioactive: These are durable tissues that can undergo interfacial interactions with surrounding tissue (bioactive glasses, bioactive glass ceramics, hydroxyapatite, calcium silicates).

Biodegradable: These are soluble or resorbable materials, those eventually can be replaced or incorporated into the tissue (Tricalcium phosphate, Bioactive glasses).

Advantages of Bioceramics:

- Due to their similarity with biological hydroxyapatite crystals they have excellent biocompatibility.
- Bioceramics have intrinsic osteoinductive capacity because of their ability to absorb osteoinductive substance if there is a bone healing process nearby.
- These provide a framework of resorbable lattices that act as a regenerative scaffold that is eventually dissolved as the body rebuilds tissue.
- They have ability to achieve excellent hemetic seal, form a chemical bond with the tooth structure and have good radiopacity^{4,5}.
- Bioceramics have antibacterial properties these form porous powders containing nanocrystals with

*Corresponding author: Ankita Sharma,
Junior Resident, Department of Pathology, IGMC, Shimla.

diameters of 1-3 nm as a result of precipitation, which prevents bacterial sequestration and adhesion. Sometimes, fluoride ions are constituents of apatite crystals, which further lead to antibacterial properties⁶.

Bioceramics In Operative Dentistry And Endodontics

Mineral Trioxide Aggregate MTA was introduced by Dr Torabian in 1993. This material has osseointegrative, osseo inductive, and biocompatible properties. It has been marketed as Pro Root since its approval by FDA in 1998. It is used primarily to seal lateral root perforations and as a root-end filling material and also used for pulp capping, pulpotomy, apexogenesis, repair of root perforations, and as a root canal filling material⁷. MTA powder contains fine hydrophilic particles that set in the presence of moisture. It is currently available in grey and white forms that differ in their chemical composition. Grey form composed of dicalcium and tricalcium silicate and bismuth oxide whereas white form consists of tricalcium silicate and bismuth oxide⁸. Bismuth oxide provides radioopacity which is present in both hydrated and nonhydrated MTA. When mineral trioxide powder is mixed with water, initially calcium hydroxide and calcium silicate hydrate are formed resulting in high alkalinity of MTA which later precipitates into a poorly crystallized and porous solid gel.

Properties of MTA

- Long setting time when compared to other materials, which is their major drawback.
- Mineral trioxide aggregate has less compressive strength when compared to other materials after 24 hours.¹⁰ The compressive strength and bond strength reach their maximum after several days mixing because the hydration rate of dicalcium silicate is slower than tricalcium silicate¹¹.
- The pH value of mineral trioxide aggregate is 10.2 after mixing and it rises to 12.5 at 3 hours.

Certain limitations of MTA

- It does not come in premixed form, so difficult to use as retrofills and they have large particle size that cannot be extruded through a small syringe.

Advantages: MTA has excellent tissue biocompatibility so proposed as a potential medicament for pulp capping with reversible pulpitis^{12,13,14}. It has superior tissue reaction, amount and type of dentin bridge formation as compared to calcium hydroxide. With MTA, dentin bridge formation after pulp capping was seen at about 1 week which progressively increased in length and thickness in duration of 3 months whereas following pulp capping with calcium hydroxide, the dentin bridge was less consistent with numerous tunnel defects.¹⁴ In a histological study by Jabbarifar et al¹⁵ MTA was found to be a better choice as a pulpotomy material. Several materials have been used as root-end filling agents but the main disadvantage is micro leakage and the lack of biocompatibility. MTA was found to be material of choice for pulpotomy as compared to formocresol and hydroxyapatite treated teeth as, MTA exhibited significant reduction in inflammation, more cementum formation and regeneration of periradicular tissues¹⁶.

Apexification is a process to obtain an apical barrier in immature non vital permanent tooth so as to prevent the extrusion of the obturating material¹⁷. An MTA plug of 4mm thickness is placed at the apical region to form a barrier, sealing the canal from the periapical area^{18,19}. Mineral Trioxide Aggregate can be used to obturate the root canal of retained primary tooth where the succedaneous permanent tooth is absent but not recommended for obturation of primary teeth that are expected to exfoliate as it is observed that MTA would be absorbed slowly. Lee and associates²⁰ found that MTA had significantly less micro leakage and less tendency for overfilling or under filling, when compared with amalgam. Torabinejad and Chivian have suggested the use of MTA for sealing vertical root fractures. MTA needs to be more explored by clinicians so that its complete beneficial properties can be obtained.

Biodentine: Biodentine was developed and introduced by Septodont research group as a new class of dental material that has high mechanical properties, excellent biocompatibility and bioactive behaviour. Chemical composition of already known endodontic repair cements are based on the Ca₃SiO₅-water chemistry which made them highly biocompatible, Septodont increased the physicochemical properties which makes Biodentine²¹. Clinically Biodentine is easy to handle and biocompatible for the restorative procedures and for classical endodontic procedures. According to all the ISO standards, as well as different preclinical and clinical collaborations, Biodentine turns out to be one of the most biocompatible materials of all the biomaterials as demonstrated in dentistry.

Setting reaction of Biodentine: The chemical reaction involves hydration of tricalcium silicate which produces hydrated calcium silicate gel and calcium hydroxide. The hydrated calcium silicate gel and calcium hydroxide tends to precipitate at the surface of the particle. The CSH (calcium silicate hydration) gel formation is due to the permanent hydration of the tricalcium silicate, which gradually fills the spaces between the tricalcium grains.

Properties of Biodentine: The working time of Biodentine is up to 6 minutes with a final set at around 10-12 minutes. When tested according to ISO standard with Gilmore needles, the working time is over 1 minute and setting time is between 9-12 minutes. Biodentine has a consistency after mixing which can be manipulated with a spatula, with an amalgam carrier which is used for endodontic cements in retrograde fillings. Biodentine has superior mechanical properties as determined by the lower water content in the mixing stage. After the initial setting of Biodentine, decrease in porosity has been observed leading to improvement in the internal structure. There is a sharp increase in the compressive strength of the material in the first hour and reach up to 200Mpa at 24 hours which is more than glass ionomers. The bending resistance of Biodentine is superior to GIC but much lower than the composite resins²². It has the surface hardness in the same range as natural dentine. Biodentine is suitable for endodontic treatment as it contains zirconium dioxide for radio opacity indications. Biodentine is used as a dentine substitute under a composite restoration, as a direct pulp capping material and as an endodontic repair material.

Bioaggregate: BioAggregate is composed of nano particle sized tricalcium silicate, tantalum oxide, calcium phosphate

and silicon dioxide presents improved performance as compared with MTA. Tricalcium silicate is the main component phase, tantalum oxide is added as a radio pacifier and it is free of aluminium²³

Setting reaction: On hydration, the tricalcium silicate produces calcium silicate hydrate and calcium hydroxide. The former is deposited around the cement grains, while the latter reacts with the silicon dioxide to form additional calcium silicate hydrate. This results in reduction of calcium hydroxide in the aged cement. MTA reacts in a similar fashion; however, since it contained no additives, the calcium hydroxide was still present in the aged cement²⁴.

Biocompatibility-Bioactivity was demonstrated by deposition of hydroxyapatite. The tantalum oxide was inert as compared to bismuth oxide, and tantalum was not leached in solution.²⁵

Differences between MTA and Bioaggregate

- Bio Aggregate does not contain aluminium and contains additives such as calcium phosphate and silicon dioxide as opposed to MTA.
- MTA exhibited the presence of aluminium, while Bio Aggregate had phosphorus.
- Bio Aggregate exhibits high calcium ion release early, which is maintained over the 28-day period as compared to MTA.
- Reactivity of Bioaggregate was slower when compared to MTA²⁶.
- BioAggregate is more biocompatible, has better sealing ability, higher fracture and acidic resistance than MTA²⁷.
- Bio Aggregate has a greater potential to induce odontoblastic differentiation and mineralization than that of MTA in pulp capping²⁸.

Calcium Phosphate Cement: In vivo and in vitro studies shown calcium phosphate cement as a promising material for grafting applications. It is an bioactive and biodegradable grafting material in the form of powder and liquid. Calcium phosphate cement can be used as a canal obturation material. Good ell et al: recommend CPC as a substitute for calcium hydroxide in apexification cases.

E. Calcium Hydroxide Based Cement: Rhoner in 1940 first clinically used calcium hydroxide as a root canal filling material²⁹. A ‘miracle’ material’ Biocal was developed by French researchers, and it was believed to make radical changes to endodontic instrumentation methods³⁰. Biocal/Endocal is a root canal filler that uses calcium oxide in ethyl glycol, when calcium oxide combines with water in the tooth it becomes calcium hydroxide which has well known long term use as an excellent root canal material. The calcium hydroxide based pulpcapping agent, Dycal became popular as a sealer among some clinicians in late 1970s, later on many root canal sealers based on calcium hydroxide became popular³¹. The rationale for the addition of calcium hydroxide to root canal sealers is from its antibacterial and tissue repair abilities as observed in calcium hydroxide based base and lining materials. When compared with zinc oxide eugenol (ZOE), AH 26 and Ketac-Endo sealers in dye leakage studies, Sealapex, CRCS, and Apexit showed no significant difference in leakage at 30 days to 32 weeks^{32,33}. Sealapex exhibited slower bacterial penetration than AH 26 and AH Plus. In an animal study, Seal apex in tissue contact will dissolve and was

partially replaced by ingrowths of connective tissue³⁴. Soares et al reported the presence of disintegrated Seal apex sealer particles in macrophage away from the root-filling material in the periapical regions of dog teeth.

Conclusion

Bioceramics have evolved to become an integral part of our modern dental health care systems specially in field of endodontics. The advantages of bioceramics are its biocompatibility and antimicrobial properties. Bioceramics offer treatment options for better prognosis in various operative and endodontic procedures. They have transcend many traditionally used materials such as calcium hydroxide. Still various research is still going on to improve the properties of bioceramics so that they can become more widely used.

REFERENCES

- 1.Cheng L, Ye F, Yang R, Lu X, Shi Y, Li L. et al., 2010. Osteoinduction of hydroxyapatite/ beta-tricalcium phosphate bioceramics in mice with a fractured fibula. *Acta Biomater.* 6(4):1569–1574
2. Hermansson L. 2014. Nanostructural bioceramics: Advances in Chemically Bonded Bioceramics. CRC Press.
- 3.Tomoaia G, Tomoaia-Cotisel M, Pop LB, Mocanu A, Pop A. 2013. Nanopowders of hydroxyapatite and its substituted derivatives with medical applications and their fabrication procedure. A20100523. Romanian Patent. Apr 10;
- 4.Prati C, Gandolfi MG. 2015. Calcium silicate bioactive cements: biological perspectives and clinical applications. *Dent Mater.* April;31(4):351–70.
- 5.Utneja S, Nawal RR, Talwar S, Verma M. 2015. Current perspectives of bio-ceramic technology in endodontics: calcium enriched mixture cement - review of its composition, properties and applications. *Restor Dent Endod.* February;40(1):1–13.
- 6.Jitaru S, Hodisan I, Timis L, Lucian A, Bud M. 2016. The use of bioceramics in endodontics - literature review. *Clujul Med.*, 89(4):470–3.
- 7.Torabinejad M, Chivian N. 1999. Clinical applications of mineral trioxide aggregate. *J Endod.*, 25:197–205.
- 8.Camilleri J, Pitt Ford TR. Mineral trioxide aggregate: a review of the constituents.
- 9.Torabinejad M, Hong CU, McDonald F, Pitt Ford TR. 1995. Physical and chemical properties of a new root-end filling material. *J Endod.* 21:349–53.
- 10.Roberts HW, Toth JM, Berzins DW, Charlton DG. 2008. Mineral trioxide aggregate material use in endodontic treatment: a review of the literature. *Dent Mater*, 24: 149–64.
- 11.Myers K, Kaminski E, Miller. 1996. The effects of mineral trioxide Aggregate on the Dog Pulp. *J Endod.*, 22: 198.
- 12.Schwartz R S, Mauger M, Clement D J, Walker WA. 1999. Mineral Trioxide Aggregate: A new material for endodontics. *J Am Dent Assoc.*, 30: 967–975.
- 13.Bogen G, Kim JS, Bakland LK. 2008. Direct pulp capping with mineral trioxide aggregate. An observational study. *JADA*, 139: 305–315.
- 14.Chacko V, Kurikose S. 2006. Human pulpal response to Mineral Trioxide Aggregate (MTA): A histologic study. *J Clin Pediatr Dent*, 30: 203–209.
- 15.Ebrahim J, Mohammad RS, Neda A. 2007. Histopathologic Responses of Dog’s Dental Pulp to Mineral Trioxide Aggregate, Bio Active Glass, Formocresol, Hydroxyapatite. *Dental Research Journal*, 4: 83–87.

16. Torabinejad M, Hong CU, Lee SF, Monsef M, Pitt Ford TR. 1995. Investigation of mineral trioxide aggregate for root-end-filling in dogs. *J Endod.*, 21: 603–608.
17. Frank AL. 1966. Therapy for the divergent pulpless tooth by continued apical formation. *J Am Dent Assoc.*, 72: 87–93.
18. Lawley GR, Schindler WG, Walker WA, Kolodrubetz D. 2004. Evaluation of ultrasonically placed MTA and fracture resistance with intracanal composite resin in a model of apexification. *J Endo.*, 30: 167–172.
19. Matt GD., Thorpe JR, Strother JM, McClanahan SB. 2004. Comparative study of white and grey mineral trioxide aggregate (MTA) simulating a one or two step apical barrier technique. *J Endod.*, 30: 876–879.
20. Lee SJ., Monsef M., Torabinejad M. 1993. Sealing ability of a mineral trioxide aggregate for repair of lateral root perforations. *J Endod.*, 19: 541–544.
21. Malkondu Ö, Karapinar Kazandag M, Kazazoglu E. 2014. A review on biodentine, a contemporary dentine replacement and repair material. *Bio Med Res Int.*
22. Ranjan M. 2014. Review on biodentine—a bioactive dentin substitute. *J Dent Med Sci.*
23. Pariookh M., Torabinejad M. 2014. Calcium silicate-based cements in mineral trioxide aggregate: Properties and clinical applications. Hoboken, NJ, USA: John Wiley & Sons.
24. Camilleri J, Sorrentino F, Damidot D. 2015. Characterization of un-hydrated and hydrated Bio Aggregate™ and MTA Angelus™. *Clin Oral Investig.* April; 19(3):689–98.
25. Skrtic D, Antonucci JM, Eanes ED, Eichmiller FC, Schumacher GE. 2000. Physicochemical evaluation of bioactive polymeric composites based on hybrid amorphous calcium phosphates. *J Biomed Mater Res.*, 53(4):381–91.
26. Tuloglu N, Bayrak S. 2016. Comparative evaluation of mineral trioxide aggregate and bioaggregate as apical barrier material in traumatized nonvital, immature teeth: A clinical pilot study. *Niger J Clin Pract.*, Jan-Feb; 19(1):52–7.
27. Zhang S, Yang X, Fan M. 2013. Bio Aggregate and iRoot BP Plus optimize the proliferation and mineralization ability of human dental pulp cells. *Int Endod J.*, October; 46(10):923–9.
28. Leonardo M., Leal J., Filho A. 1980. Pulpectomy: immediate root canal filling with calcium hydroxide. *Oral Surg Oral Med Oral Pathol.*, 49:441–50.
29. Hendra L. 1970. Biocalx—a new approach to endodontia dependent upon biological principles and chemical action only. *J Br Endod Soc.*, Autumn: 37–41.
30. Goldberg F, Gurfinkel J. 1979. Analysis of the use of Dycal with guttapercha points as an endodontic filling technique. *Oral Surg Oral Med Oral Pathol.*, 47:78–82.
31. Hovland E, Dumsha T. 1985. Leakage evaluation in vitro of the root canal sealer cement Sealapex. *Int Endod J.*, 18:179–82
32. Jacobsen E., BeGole E., Vitkus D. et al, 1987. An evaluation of two newly formulated calcium hydroxide cements: a leakage study. *J Endod.*, 12:164–9.
33. Yucel A., Guler E., Guler A. et al., 2006. Bacterial penetration after obturation with four different root canal sealers. *J Endod.*, 32:890–3.
34. Soares I, Goldberg F, Massone E, et al. 1990. Periapical tissue response to two calcium hydroxide containing endodontic sealers. *J Endod.*, 16:166–9.
