

Recent Advances in Pulp Capping Materials

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ABSTRACT

Pulp capping is a dental restoration procedure that protects the dental pulp from death of tissue after it has been exposed, or almost exposed during cavity preparation, from a severe injury, causing the pulp to die. Calcium hydroxide and Mineral Trioxide Aggregates (MTA) are the most often utilised pulp capping materials in dentistry, and they have had significant clinical success. In recent years various other materials like Bone morph genic protein, Bio dentin, Lasers are also introduced clinically. Therefore this review article will summarize recent pulp capping materials and their advantages.

Key words: Pulp capping, Pulp capping materials, Calcium hydroxide, MTA, Growth hormones, Enzymes

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INTRODUCTION

It is a dental restoration procedure that protects the dental pulp from death of tissue due to soft tissue injury [1]. There are two methods of pulp capping. Direct method is a treatment for exposed vital pulp that entails placing a dental substance over the exposed area to help form a preventive barrier [2-4]. As well as the preservation of vital pulp [5-6]. Indirect method is a treatment that involves placing a substance on a tiny portion of residual dentin that prevents essential exposure of pulp. Pulpotomy varies from pulp-capping in that it removes a part of the existing pulp prior to application of capping substance. Caries exposure occurs when pulp is exposed before caries is totally eradicated. Mechanical exposure occurs when pulp is exposed during cavity preparation with no caries. Mechanical exposures usually occur as a result of a failure during tooth preparation. A various dental substances used in direct pulp capping although they typically have poor effects because it shows toxic effect to the dental pulp [7]. Calcium hydroxide is the one of the best material materials, with a significant track record of clinical success. Mineral Trioxide Aggregates

(MTA), have gained a lot of attention as direct pulp capping materials in recent years. Apart from calcium hydroxide, MTA is also used as material [8]. Despite the fact that both these materials have an excellent track record in clinical trials, they both have some drawbacks, such as slow (or no) setting times, difficult handling, low physical strength, high solubility, and/or progressive resorption. They can be light cured, giving them a favourable condition it enhances the ability to withstand the stress of physical forces, it is easy to manipulate, and more accurate for placement [9].

Furthermore, the pulp-capping material should possess the desirable characteristics such as:

- Stimulate irritation dentin development.
- Maintain normal condition of pulp.
- Fluoride is released to inhibit the formation of a layer of dentin after the tooth's root has fully-developed.
- Bacteriostatic or bactericidal.
- Bind to the dentin
- Bind to restorative materials.
- Withstand the pressures when the restoration is placed.
- It is free from bacteria or microorganisms.
- Gives bacterial seal [10].

LITERATURE REVIEW

Following are the recent pulp capping materials

Calcium hydroxide: Hermann introduced in the year 1921 Calcium Hydroxide (Ca $(OH)_2$) as a best capping materials [11-13]. He also found that calcium hydroxide may be used to heal an exposure site in 1930 [14-15]. Since in the form of powder, paste, and cement has been utilised successfully in the clinic to aid in the creation of reparative dentin as well as the preservation of essential pulp, mineralization, and bacterial growth inhibition [16-17].

Advantages

- It is gold standard of direct pulp capping material.
- Initially bactericidal than bacteriostatic.
- Promote healing and repair.
- Balances low pH of acids.
- Stops destruction of dentinal tubules.
- Induction of mineralization.
- Poor cytotoxicity.
- Low cost and simple to use [18].

Disadvantages

- After one year, cavo surface disintegration may occur.
- Marginal failure with amalgam condensation
- Over time, it will disintegrate.
- The pulp chamber is completely obliterated due to extensive dentin development.
- Adhesion problems
- After acid etching, the material degrades [18].

Glass ionomer or resin modified glass ionomer

When utilised in close proximity but not direct contact with the pulp, glass ionomer provides an effective resistance against bacteria and it has good biocompatibility [18-19].

Advantages

- Amazing bacterial seal.
- Fluoride release, expansion coefficient, and elastic modulus are similar to dentin.
- Affiliate with both enamel and dentin.
- Excellent compatibility with pulp tissue [18].

Disadvantages

- Causes long-term inflammation.
- Formation of dentinal bridges is absent.
- It is cytotoxic when it comes into contact with cells.
- Physical characteristics are poor, solubility is high, and the setting rate is slow.
- Resin Modified Glass Ionomer Cement is more toxic than traditional GIC; administration to the pulp tissue should not be done directly [18].

Calcium phosphate: Because of its good compatibility with pulp tissue, it has better compressive strength, and ability to change into hydroxyapatite over time, calcium

phosphate cement has been considered as a possible replacement [18]. In contrast to calcium hydroxide, Yoshimine found that tetra calcium phosphate cement caused bridge development with no superficial death of tissue and no substantial pulp inflammation [20].

Advantages

- With no superficial tissue necrosis, it aids in the creation of bridges.
- Compared to Ca (OH)₂, there is a considerable lack of pulp inflammation.
- Good physical properties [18].

Disadvantages

Clinical trials are required to assess this item [18].

Lasers: Between 1985 and 1987, Melcer suggested applying a carbon dioxide (CO_2) (1W) laser for "direct pulp capping [21-24].

Advantages

- Following the use of a laser for pulpectomy in primary teeth, better clinical, radiographic, and histological results were obtained.
- In diode laser pulpotomies, the patient did not experience any pain or discomfort, and no painkiller was required.
- Secondary dentin formation.
- Targeted tissue sterilization.
- Bactericidal properties [18].

Disadvantages

- Techniques that is sensitive.
- In large doses, causes thermal damage to pulp.
- High price [18].

Mineral Trioxide Aggregates (MTA): Torabinejad in year 1990s discovered Mineral Trioxide Aggregates (MTA). Has become available as various types of dental materials. During the setting process initial PH 10.2 which increases up to 12.5 during first few hours [25].

Advantages

- Cell adhere and growth.
- Nontoxic and non-mutagenic.
- Promote formation of original tissue when it gets contact with cementum, alveolar bone, and periodontal ligament.
- It has the capacity to trigger cytokine release from bone cells, which aids in the creation of hard tissue.
- There is less pulpal irritation.
- In compared to calcium hydroxide, the creation of a hard tissue barrier is more predictable.
- Property of antibacterial.
- Radio sensitivity.
- Bioactive dentin matrix proteins are released [18].

Disadvantages

• It is a technique sensitive material, hard to manipulate

- Long time to set
- Tooth discolouration is caused by grey MTA.
- A two-step approach is followed.
- Solubility is high [18].

Zinc Oxide Eugenol (ZOE) Cement: ZOE cement is more effective for irritated and exposed pulp. When it comes in contact with soft tissues it causes inflammation [26]. Eugenol has anti-inflammatory response which reduces sensitivity in the dental pulp at low doses [18].

Advantages

- Zinc oxide eugenol cement destroys or inactivates pathogens.
- Kills bacteria present in carious lesions, so arrest its process this gives pulp chance for healing and regeneration.
- Excellent initial seal [18].

Disadvantages

- Formation of calcific bridges is not possible due to a lack of calcium.
- Releases a high concentration of eugenol, which is toxic to living cells or tissue.
- Demonstrate the presence of interfacial leakage [18].

Corticosteroids and antibiotics: For pulp capping, corticosteroids such as Cleocin, penicillin, neomycin, and Keflin were used with calcium hydroxide in the hopes of minimising or protecting pulp inflammation [18]. Gardner discovered that a combination of vancomycin and calcium hydroxide was somewhat more effective in inducing a more regular reparative dentin bridge than calcium hydroxide. According to Watts and Paterson, anti-inflammatory medicines should not be given to anyone who is at risk of bacteraemia [27-28].

Advantages

- Reduces the inflammation of the pulp
- Promoteda more regular reparative dentin with vancomycin+Ca (OH)₂ [18].

Disadvantages

• Should not be used in patients who are susceptible to bacteraemia [18].

Inert materials: Isobutyl cyanoacrylate and tri calcium phosphate ceramic are the Inert materials. Although there was less pulpal inflammation and unanticipated space between one or more teeth, none of these materials has been recommended to the dentistry as a viable approach [29-30]. Cyanoacrylate is an adhesive that result from chemical reaction between formaldehyde and esters of cyanoacrylate [18].

Advantages

- It is an excellent agent which shortens the clotting of blood and prevents the growth of bacteria.
- It triggers the formation of morphologically abnormal dentin [18].

Disadvantages

- It does not produce continues barrier of morphologically irregular dentin following application of not covered pulp tissue.
- None of these materials have been promoted in dentistry as a viable technique [18].

MTYA1-Ca filler: Atsuko Niinuma developed resin based agent containing calcium hydroxide. Dentine bridge construction occurred in MTYA₁-Ca and necrotic layer was not formed, despite the fact that it was not histopathological inferior to Dycal. Hence it promises to be a good material [31].

Advantages

- Aids in the production of morphologically irregular dentin formed in response to an irritant or other substance that reseals exposed tooth pulp tissue without forming a necrotic layer.
- Shear bond strength is comparable to RMGIC and higher than ordinary GIC.
- Dentin bridge formation occurs in MTYA₁-Ca without a reduction in pulp space.
- Dentin bonding is improved [18].

Disadvantages

• The presence of 10% Ca (OH)₂ prevents the material from fully curing, and leftover monomers cause toxicity in pulp tissue [18].

Bonding agents: Dentin bonding agents, according to miyakoshi. Give greater adherence to peripheral hard tissues. However, because of the cytotoxic effect and the lack of calcific bridge development, they have a poor prognosis [32].

Advantages

- Effective barrier against germs, oral fluids, ions, and chemicals getting into the contact between the teeth and filling material.
- The tendency of disparate objects or surfaces to adhere to each other due to physical forces in hard tissues [18].

Disadvantages

- Toxic to living tissue.
- The production of calcific bridges is not present.
- In vivo investigations have shown that placing an adhesive resin directly on the point of exposure of pulp or dentin thickness of less than 0.5 mm promotes blood vessel dilation and congestion, as well as a chronic inflammatory pulp response [18].

Growth factors: Growth factors promote wound healing and tissue regeneration as well as regulating growth and development.

Bone morphogenic protein: Bone Morphogenic proteins are the group of molecules work by producing adult stem cells to replicate into bone forming cells lines that form new bone. They are involved in many physiological and pathological processes such as-

• Inflammatory response.

- Bone forming and resorption.
- Growth signalling pathways.
- Oncogenesis and immune response.
- They are also referred to as osteogenic proteins.

Advantages

- It play important role in cell growth, apoptosis in variety of cell development including osteoblast and chondrocytes.
- Rapidly formed tertiary dentin and tubular dentin formation.
- Dentin formation that is more homogenous.
- In terms of mineralization inducing properties, Ca (OH)₂ is superior.
- After 28 days, the dentin bridge development was equal to dycal [18].

Disadvantages

- Increased infection rates
- Increased postoperative swellings
- The expense of their clinical application can be an impediment.
- In the case of an inflammatory pulp, failure to activate reparative dentin
- The half-life is shorter.
- A high level of concentration is needed.
- Immunological issues may arise as a result of the frequent implantation of active substances [18].

Bio Dentin: Bio dentine is novel bioactive cement that has mechanical qualities similar to dentin and can be used to replace dentin. It enhances the production of tertiary dentin and has a beneficial effect on vital pulp cells [33].

Advantages

- Biocompatible.
- Antimicrobial activity is good.
- Enhance the development of tertiary dentin.
- It is mechanically stronger, less soluble, and creates tighter seals than calcium hydroxide.
- Compared to MTA, it takes less time to set up and has better handling capabilities [18].

Disadvantages

• For a clear assessment of Bio dentine, more long-term clinical investigations are required [18].

Bone Sialoprotein: According to Goldberg M bone sialoprotein was the most beneficial bioactive molecule in producing uniform and well mineralized secondary dentin (BSP). In terms of mineralization inducing characteristics, both BSP and BMP-7 outperformed calcium hydroxide [34].

Advantages

- Induced reparative dentin that is homogeneous and well mineralized
- In terms of mineralization inducing properties, Ca (OH)₂ is superior [18].

Disadvantages

• Further clinical studies are needed [18].

Enzymes

Heme-Oxygenase-1: Hypoxic stress and nitric oxidemediated cytotoxicity are both protected by HO-1 induction. Furthermore, HO-1 expression in dental pulp cells produced by Bismuth oxide-containing Portland Cement (BPC) protects against BPC's cytotoxic effects [35].

Advantages

- They act as a barrier against cytokines and nitric oxide in human pulp cells.
- Prevent toxicity in pulp cells caused by H₂O₂ [18].

Disadvantages

• Damage to pulp tissue occurs at high concentrations.

Thera Cal: It is a light-cured, resin-modified liner for both pulp capping methods, as well as a protective base/ liner beneath composites, amalgams, and other base materials. It acts as a dental pulpal complex insulator, barrier, and protectant.

Advantages

- Protect the tooth pulpal complex by acting as a protectant. Dentin that is deep and wet
- Physical properties are strong, there is no solubility, and the radiopacity is high.
- Pro Root MTA and Dycal both have stronger calcium releasing ability but lesser solubility than Thera Cal [18].

Disadvantages

• It's opaque and "whitish" in colour, and must be thin to avoid showing through translucent composite materials and influencing final restoration shading [18].

Other enzymes

- Oil Bean (COB) Cement.
- Novel Endodontic Cement (NEC).
- Emdogain (EMD).
- Propolis (Russian penicillin) [36-40].
- Stem Cells.
- Simvastatin.

CONCLUSION

Mineral trioxide aggregate, calcium hydroxide, zinc oxide eugenol, zinc phosphate etc. are the materials used in direct pulp capping as well as indirect pulp capping. Materials are biocompatible to the patients and hence widely used in dental field. Pulp capping materials are easy to manipulate for dentist and clinicians. Various growth factors such as bio dentin, bone morphogenic protein and enzymes like heme-oxygenase-1 are also introduced clinically which enhance the growth and development of pulp capping.

REFERENCES

- 1. Komabayashi T, Zhu Q, Eberhart R, et al. Current status of direct pulp-capping materials for permanent teeth. Dent Mater J 2016; 35:1-2.
- Bergenholtz G, Mjor IA, Cotton WR, et al. The biology of dentin and pulp. Consensus report. J Dent Res 1985; 64:631-633.
- 3. Couve E. Ultrastructural changes during the life cycle of human odontoblasts. Arch Oral Biol 1986; 31:643-651.
- 4. Pashley DH. Dynamics of the pulpo-dentin complex. Crit Rev Oral Biol Med 1996; 7:104-133.
- 5. Zander HA, Glass RL. The healing of phenolized pulp exposures. Oral Surg Oral Med Oral Pathol 1949; 2:803-810.
- 6. Bergenholtz G. Advances since the paper by Zander and Glass (1949) on the pursuit of healing methods for pulpal exposures: historical perspectives. Oral Surg Oral Med Oral Pathol Oral Radiol Endod 2005; 100:102-108.
- 7. Hilton TJ. Keys to clinical success with pulp capping: a review of the literature. Oper Dent 2009; 34:615-25.
- 8. Bogen G, Kim JS, Bakland LK. Direct pulp capping with mineral trioxide aggregate: an observational study. J Am Dent Assoc 2008; 139:305-315.
- 9. Chen L, Suh BI. Cytotoxicity and biocompatibility of resin-free and resin-modified direct pulp capping materials: A state-of-the-art review. Dent Mater J 2017; 36:1-7.
- 10. Cohen BD, Combe EC. Development of new adhesive pulp capping materials. Dent update 1994; 21:57-62.
- 11. Cox CF, Subay RK, Ostro E, et al. Tunnel defects in dentin bridges: their formation following direct pulp capping. Oper dent 1996; 21:4-11.
- 12. Schroder U. Effects of calcium hydroxidecontaining pulp-capping agents on pulp cell migration, proliferation, and differentiation. J dent res 1985; 64:541-548.
- 13. Cox CF, Suzuki S. Re-evaluating pulp protection: calcium hydroxide liners vs. cohesive hybridization. J Am Dent Assoc 1994; 125:823-831.
- 14. Hermann B. CalciumhydroxydalsMittelzumBehandel und Fullen von ZahnwurzelkanalenWürzburg, Germany: Faculty of Medicine, University of Wurzburg. 2019.
- 15. Hermann BW. Dentinobliteration der wurzelkanalenachbehandlungmit calcium. ZahnärztlRundsch. 1930; 39:888-899.
- 16. Stuart KG, Miller CH, Brown Jr CE, et al. The comparative antimicrobial effect of calcium hydroxide. Oral Surg Oral Med Oral Pathol 1991; 72:101-104.
- 17. Cavalcanti BN, Rode SM, Marques MM. Cytotoxicity of substances leached or dissolved from pulp capping materials. Int Endod J 2005; 38:505-509.

- Qureshi A, Soujanya E, Nandakumar P. Recent advances in pulp capping materials: an overview. J Clin Diagn Res 2014; 8:316.
- 19. Tarim B, Hafez AA, Cox CF. Pulpal response to a resin-modified glass-ionomer material on nonexposed and exposed monkey pulps. Quintessence Int 1998; 29.
- 20. Yoshimine Y, Maeda K. Histologic evaluation of tetracalcium phosphate-based cement as a direct pulp-capping agent. Oral Surg Oral Med Oral Pathol Oral Radiol Endod 1995; 79:351-358.
- 21. Melcer J, Chaumette MT, Melcer F, et al. Experimental research on the preparation of dentin-pulp tissue of teeth exposed to CO_2 laser beams in dogs and macaques (Macacamulatta and Macacafascicularis). C R Seances Soc Biol Fil 1985; 179:577-585.
- 22. Melcer J. Latest treatment in dentistry by means of the CO_2 laser beam. Lasers Surg Med 1986; 6:396-398.
- 23. Melcer J, Chaumette MT, Melcer F. Dental pulp exposed to the CO_2 laser beam. Lasers in surgery and medicine 1987; 7:347-352.
- 24. Yasuda Y, Ohtomo E, Tsukuba T, et al. Carbon dioxide laser irradiation stimulates mineralization in rat dental pulp cells. Int Endod J 2009; 42:940-946.
- 25. Bogen G, Kim JS, Bakland LK. Direct pulp capping with mineral trioxide aggregate: an observational study. J Am Dent Assoc 2008; 139:305-315.
- Dummett CO, Kopel HM. Pediatric Endodontics. In Ingle and Bakland. 5th ed. Endodontics: B.C. Decker Elsevier 2002; 861-902.
- 27. Gardner DE, Mitchell DF, Mcdonald RE. Treatment of pulps of monkeys with vancomycin and calcium hydroxide. J Dent Res 1971; 50:1273-1277.
- 28. Watts A, Paterson RC. Cellular responses in the dental pulp: a review. Int endod j 1981; 14:10-21.
- 29. Bhaskar SN, Beasley JD, Ward JP, et al. Human pulp capping with isobutyl cyanoacrylate. J Dent Res 1972; 51:58-61.
- Heys DR, Cox CF, Heys RJ, et al. Histological considerations of direct pulp capping agents. J Dent Res 1981; 60:1371-1379.
- Niinuma A. Newly developed resinous direct pulp capping agent containing calcium hydroxide (MTYA1-Ca). Int Endod J 1999; 32:475-483.
- 32. Miyokoshi S. Interfacial interactions of 4-META-MMA/TBB resin and pulp. JCDR 1993; 72:220.
- 33. Laurent P, Camps J, De Meo M, et al. Induction of specific cell responses to a Ca3SiO5-based posterior restorative material. Dent mater 2008; 24:1486-1494.
- Goldberg M, Six N, Decup F, et al. Application of bioactive molecules in pulp-capping situations. Adv Dent Res 2001; 15:91-95.

- 35. Min KS, Lee HJ, Kim SH, et al. Hydrogen peroxide induces heme oxygenase–1 and dentin sialophosphoprotein mRNA in human pulp cells. J endod 2008; 34:983-989.
- 36. Madan K, Baliga S, Deulkar P, et al. A Comparative Evaluation between Propolis and Mineral Trioxide Aggregate as Pulpotomy Medicaments in Primary Molars. J Evol Med Dent 2020; 9:1256– 1260.
- 37. Mahapatra J, Nikhade PP, Belsare A. Comparative evaluation of the efficacy of theracallc, mineral trioxide aggregate and bio dentine as direct pulp capping materials in patients with pulpal exposure in posterior teeth-an interventional study. Int J Pharm Res 2019; 11:1819–1824.
- Rathi S, Nikhade P, Jaiswal A, et al. Management of Deep Carious Lesion with Single Visit Indirect Pulp Capping: A Case Report. Indian J Forensic Med Toxicol 2020; 14.
- 39. Bhonde R, Ikhar A, Palsodkar P. Comparative clinical evaluation of post endodontic pain in retreatment cases using calcium hydroxide and triple anti biotic paste an interventional study. Int J Pharm Res 2019; 11:1428–1430.
- 40. Chandak MG, Modi RR, Rathi BJJ, et al. In vitro comparative assessment of diffusion of ion from calcium hydroxide with three different phytomedicine pastes through dentin. World J Dent 2018; 9:366–371.