Overview of Advances of Dental Composite Materials

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Abstract: The current review aims to review the advancement/development approaches that have been taken to improve dental composite material performance, and the changes that occurs since last two decades to these composite properties and how it improvement effects in dental health. Comprehensive review of evincive discussing the dental composite material advances, was performed through biomedical databases as: PubMed, and Embase. All studies search was limited to English language with human subject published studies up to September, 2017. Composite dental repairs represent a unique class of biomaterials with serious restrictions on biocompatibility, healing actions, esthetics, and also best material buildings. These products are presently limited by contraction as well as polymerization-induced shrinkage tension, limited durability, the existence of unreacted monomer that remains following the polymerization, as well as several other factors. These products are currently restricted by shrinking and polymerization generated contraction tension, minimal sturdiness, the presence of unreacted monomer that continues to be adhering to the polymerization, as well as a number of various other aspects. Luckily, various advancements take place in enhancing remediation efficiency by changing the initiation system, monomers, as well as fillers, and also by creating unique polymerization approaches.

Keywords: dental health, composite material, biocompatibility, healing actions, esthetics.

1. INTRODUCTION

Composite materials (also called as composition materials or composites) are products composed of two or even more materials to create a new valuable product. Representing one of the many successes of contemporary biomaterials research, given that they replace organic tissue in both appearance as well as function. At the very least fifty percent of posterior direct repair placements currently count on composite products [1]. Dental composites are complex combined products which typically consist of an organic polymerizable matrix, strengthening fillers, which are generally inorganic and also a silane-coupling representative [2]. The polymerizable matrix includes several monomers, co-onomers and different ingredients, like an initiator (camphorquinone), coinitiator, a prevention of polymerization, and also a photostabilizer (e.g., benzophenone), numerous not natural products are used as fillers: quartz, borosilicate, lithium aluminum silicate glasses, as well as amorphous silica. In order to attain radiopacity, oxide glasses with barium, strontium, zinc or other steels are added to fillers of modern material composites [3].

Picking an appropriate resin-based composite (RBC) for a remediation in modern-day dental care needs balancing a great deal of requirements. This needs functional residential properties, consisting of enhanced long life of the restoratives by outstanding mechanical properties such as high stamina, fracture sturdiness, surface solidity, maximized modulus of elasticity, low wear, low tide sorption as well as solubility, low polymerization contraction, reduced exhaustion and also degradation, high radiopacity and a much better discovery of the product throughout removal of a composite restoration. Several scientific adverse effects in resin-based composite repairs, such as low inconsistencies [4,5]. minimal staining, white lines around the repair, cusp cracks, microleakage, debonding, secondary cavities, postoperative sensitivity or pain are still observed in repairs with modern RBCs [6,7]. These are regularly attached to polymerization shrinkage anxiety. Even if these assertions are nowadays generally accepted, just minimal scientific proof exists to sustain a clear relationship in between these adverse end results and also polymerization shrinking anxiety [5,6,7].

Vol. 5, Issue 2, pp: (182-187), Month: October 2017 - March 2018, Available at: www.researchpublish.com

There are some records about leaching substances from dental composite materials and also problems on their biocompatibility which could influence growth and immune responsivity of gingival fibroblasts [8,9]. In long-term teeth, dental material composites are the most essential tooth tinted filling up materials, in the primary dentition crack sealers, conventional as well as resin-modified glass ionomer concretes, and compomers additionally play a crucial duty [3,8,9]. Inevitably, these advancements decrease a repair's lifetime and also stand for the driving pressure for enhancement in dental composites. Clinical evaluations as well as laboratory-based researches focused on composite toughness likewise remain to highlight this need for brand-new products [9].

The current review aims to review the advancement/development approaches that have been taken to improve dental composite material performance, and the changes that occurs since last two decades to these composite properties and how it improvement effects in dental health.

2. MATERIALS AND METHODS

Comprehensive review of evincive discussing the dental composite material advances, was performed through biomedical databases as: PubMed, and Embase. all studies search was limited to English language with human subject published studies up to September, 2017. moreover, articles were recruited from searching references lists of included articles.

3. DISCUSSION

• Composites with lighter weight than before and effect of DMAEMA on mechanical properties of composites:

In alternate to magnesium, aluminum and also steel alloys have been made use of for in automobiles for their lightweight. Emerging patterns in polymers and also fiber-based crossbreed composite has caused the growth of lightweight cars. The other course of lightweight materials in automobile market are all-natural fiber enhanced composites. Numerous auto industries are making use of the all-natural fiber reinforced composites (density 1.5 g/cc) as an alternative to glass fiber (2.5 g/cc). In industries producing premium items, carbon fiber is being made use of as an alternative to glass fiber. Crossbreed design of carbon as well as glass fiber have actually caused maximum performance. Recycling of carbon fibers as well as including nano fibers in the carbon fibers improves the efficiency of cars. College of Toronto, Canada, in association with vehicle companions have developed a lightweight crossbreed prototypes which is presently under study. Current advances in use of cellulose and also carbon fiber- enabled composite frameworks have actually raised the ergonomics style of automobiles [10].

In one important research [11] compression tests were carried out for both bought (wet colloidal crystal in this situation) and non-ordered composite samplings (**Table 1**). All composite samples made use of as-received (i.e. no silane finish) silica grains as filler (60 wt%) as well as consisted of CQ (0.5 wt%). The only distinction was that the non-ordered example contained 0.5 wt% DMAEMA, while there was no dimethylaminoethyl methacrylate (DMAEMA) in the bought example. For comparison objectives, cool resin examples of TEGDMA with or without DMAEMA were additionally prepared. The ASTM criterion does not supply a straight method to measure toughness from compression tests, we approximated the strength as the incorporated location under the stress-strain curve [12].

	TEGDMA (with 0.5 wt% DMAEMA)	TEGDMA (no DMAEMA)	Non-ordered Composite (with 0.5 wt% DMAEMA)	Ordered Composite (no DMAEMA)
Modulus (GPa)	1.91 (0.11)	1.88 (0.24)	4.92 (0.30)	4.49 (0.48)
Failure Strain (%)	29 (2.9)	29 (4.7)	8.3 (1.2)	14.2 (2.5)
Compressive Strength (MPa)	198.9 (26.7)	220.7 (56.3)	165.4 (21.2)	192.1 (17.2)
Toughness [*] (J/m ³ × 10 ⁷)	3.12 (0.65)	3.05 (0.86)	0.77 (0.31)	1.81 (0.48)

Table 1: Mechanical properties (in compression) of neat resins and monodisperse, non-silanized silica-filled composites

• Improvement and modeling of Composite Materials:

The improvement of composite products specifically glass fiber reinforced plastic (GFRP) in pulstration production market was accomplished by procedures like Re-engineering, re-cycling and re-designing. This enhancement was gone for eco-efficiency performance. The term 'eco-efficiency' indicates doing a lot more with less resources. It was observed that eco efficiency renovation can be attained by minimizing the intake of material, power and waste. In the Re-engineering method, the temperature account of the die was verified experimentally by Thermography strategy and also

Vol. 5, Issue 2, pp: (182-187), Month: October 2017 - March 2018, Available at: www.researchpublish.com

evaluated by FEM, which showed the decrease of energy intake by 17 % and also warm up time by 50 %' In re-cycling method, the non-conformities, byproducts of polycarbonate products and carbon fiber enhanced plastic (CFRP) were taken into observation as well as located to be reliable by 15 %' replacement [13].

As a result of innovations in newer products, the composite materials call for modeling and also simulation. FEM is the mathematical technique of structural evaluation in which the data for the needed for structure analysis such as anxieties, contortions, variations can be gotten. FEM indicates splitting the structures right into small dimensions and forms and then creating a Mesh and also analysing it. Catia, pro-E, strong works, 'CFD are several of the software applications available for modeling as well as analyzing the warm transfer, liquid moves in numerous structures [14].

• Dental Composites with Nano-fillers:

Considerable attention has actually been dedicated to nano filled materials, consisting of improvements recognized by the consolidation of nanofillers right into industrial composite materials and study targeted at the growth of new nanofillers. A recent review focused on nano filled dental composite products [15], and a different report fixated just how nanofillers impact composite mechanical homes as well as behave distinctly differently compared with mini- or macro-scale fillers [16]. Nano-sized fillers can be classified as either separated discrete fragments, with dimensions of approximately 5 to 100 nm, or merged accumulations of primary nanoparticles, where the cluster size might dramatically surpass 100 nm. The massive increase in filler surface and the corresponding thickening result on composite paste consistency connected with reducing filler dimension limit the content of discrete nanoparticles to fairly low loading levels, whereas high materials of nanoparticle collections are convenient with suitable surface area therapy. A spatially solved nanoindentation study examined Filtek Supreme XT (A3 Dentin) as a nanofilled composite as well as demonstrated considerable differences in the dynamic complex modulus as a function of placing within the matrix, within a filler collection, or at the matrix-filler user interface [17]. A research on the influence of mono-, bi-, and also tri-modal circulations of fillers on the wear residential properties of composites showed that filler shapes and size dramatically influence wear resistance, with the inclusion of nano-sized filler a crucial function, frequently bring about enhanced residential or commercial properties [18]. A similar dependence of toothbrush abrasion resistance on the presence of nanoparticles in commercial dental composites has actually been revealed [19].

• Camphorquinone/Amine system improvement:

The camphorquinone/amine initiation system remains to be the topic of research in initiatives to figure out maximum initiator as well as co-initiator focus for kinetics and also polymer mechanical residential or commercial properties [20]. In this visible-light-activated initiator system, camphorquinone (CQ) takes in a photon to produce a short-term excited-state types that complicated with the tertiary amine to advertise a sequential electron and also proton transfer that creates the energetic α-aminoalkyl-initiating radical. Added studies have reviewed alternate co-initiators to the generally utilized ethyl-4-dimethylaminobenzoate (EDMAB), consisting of N,N-dimethylaminobenzyl alcohol, 4-(N,N-dimethylamino) phenethyl alcohol (DMPOH), and also N,N-3,5- tetramethylaniline (TMA) as well as a range of aliphatic as well as aromatic amines [21,22]. With the goal of minimizing or getting rid of the amount of possibly toxic amine co-initiator, cyclic acetals and also the normally taking place 1,3-benzodioxole and also its by-products were examined as potential substitutes for traditional amine co-initiators as well as were discovered to be reliable co-initiators, resulting in kinetics as well as polymer residential or commercial properties just like those of equal systems launched by CQ/EDMAB [23,24]. PPD (1-phenyl-1,2- propanedione) is a non-yellowing photosensitizer that has actually been proposed as a choice to CQ, though it does not appear to be as reliable as CQ for visible light initiation [20,25].

One fascinating technique for attending to amine co-initiator toxicity has actually been to develop polymerizable monomers that likewise function as co-initiators. Therein, the amine part is polymerized right into the network through the methacrylate, which substantially restricts the extractable amine. Several methacrylated amine co-initiators, such as ethylene glycol 3-diethylamino-propionate methacrylate, have actually been synthesized and also demonstrated to exhibit polymerization kinetics as well as residential or commercial properties almost equal to those of typical BisGMA/TEGDMA systems with a CQ/EDMAB initiation system [26,27].

• Acidic Monomers improvement:

Recent growths in methacrylate materials have explored the incorporation of acidic practical groups right into the monomer framework. Integrating acidic monomers in fairly tiny mole portions into methacrylate materials might make it possible for a different sticky layer to be gotten rid of and result in boosted overall efficiency. Current acidic materials do

Vol. 5, Issue 2, pp: (182-187), Month: October 2017 - March 2018, Available at: www.researchpublish.com

not exhibit the essential mechanical residential or commercial properties to function as resin-based composites, as well as thus study has actually focused on developing acidic monomers with improved mechanical residential properties. Acidic monomers have been manufactured from o-hydroxyaryl phosphonates that exhibited fast polymerization kinetics [15]. Bisphenol-A by-products have been generated both with carboxylic acid and also phosphonic acid practical teams without degradable esters and also with carboxylic, amide, and also hydroxy functional teams to improve adhesion [23,28]. Other by-products of BisGMA, including carboxylic acid functionalized monomers, show equivalent mechanical buildings but enhanced water sorption [29]. The enhanced moisture absorption related to the acid is a usual issue, given that the existence of the acidic team raises the hydrophilicity of the material through raised polarity as well as charge thickness. Acidic monomer-containing composites with various fragrant core frameworks were reviewed for mechanical properties as well as found to exhibit homes just like those of BisGMA/TEGDMA [30]. The use of acidic monomers requires consideration of hydrolytic stability. Acrylamides show boosted hydrolytic security compared to esters, as well as especially when used under acidic problems, acrylamides offer an encouraging option to methacrylates. When contrasted with comparable dimethacrylate resins [bis-acrylamides showed comparable reactivity, flexural toughness, as well as flexural modulus [31]. The use of inner amide useful teams have likewise been taken into consideration. These monomers are derivatives of bisphenol-An and also displayed considerable decreases in volume shrinkage, yet likewise led to boosted thickness and also, because of reduced solubility, have the ability to be incorporated just at levels up to 5 mol% in BisGMA/TEGDMA materials [32].

• Composite advances in Ring-opening Polymerization:

The primary reason that ring-opening polymerization has actually gotten focus is underpinned by the one-of-a-kind shrinking actions observed in these polymerizations. Whereas methacrylate (and also thiol-ene) photopolymerizations entail the conversion of a carbon-carbon double bond right into solitary bonds, the ring-opening reaction relies on the opening of a cyclic framework to assist in intermonomer bonding as well as crosslinking. When rings are opened [integral to the cyclic structures is that substantially much less volume contraction takes place [33].

A recent interesting growth in ring-opening polymerization is the business release of the cationically photopolymerizable silorane product (Filtek LS) by 3M/ESPE [34]. The material chemistry counts on the ring-opening polymerization of a combination of proprietary as well as conveniently available cycloaliphatic monomers. The silorane terminology originates from the unique monomer made up of a cyclic siloxane core added with 4 oxirane responsive teams. The cyclohexene oxide-type oxirane rings are dramatically more responsive compared with a simple epoxy analog, and the very hydrophobic siloxane framework efficiently stabilizes the prospective hydrophilicity of the polyether backbone generated by the ring-opening polymerization. The cationic reaction is turned on by a visible-light photoinitiator system with camphorquinone as photosensitizer, a tertiary fragrant amine as a photoreductant, as well as an iodonium salt as an electron contributor that produces the active cationic varieties. The filler selection, the filler surface area treatment, and the adhesive required for bonding the composite to dentin and enamel are also created to suit the cationic curing procedure. A successful, cationically started dental corrective material definitely stands for a substantial difficulty, as implied by the reality that Bowen at first tried and also abandoned cationic epoxide products before creating BisGMA in his critical work on dental corrective materials [33]. The offered literature on silorane physical as well as mechanical buildings has recently been reviewed [35]. A mechanical building characterization targeting numerous various size scales has been conducted with the silorane composite product together with a number of business composites based on traditional dimethacrylate material chemistry [17].

4. CONCLUSION

Composite dental repairs represent a unique class of biomaterials with serious restrictions on biocompatibility, healing actions, esthetics, and also best material buildings. These products are presently limited by contraction as well as polymerization-induced shrinkage tension, limited durability, the existence of unreacted monomer that remains following the polymerization, as well as several other factors. These products are currently restricted by shrinking and polymerization generated contraction tension, minimal sturdiness, the presence of unreacted monomer that continues to be adhering to the polymerization, as well as a number of various other aspects. Luckily, various advancements take place in enhancing remediation efficiency by changing the initiation system, monomers, as well as fillers, and also by creating unique polymerization approaches.

Vol. 5, Issue 2, pp: (182-187), Month: October 2017 - March 2018, Available at: www.researchpublish.com

REFERENCES

- Sadowsky SJ. (2006). An overview of treatment considerations for esthetic restorations: a review of the literature. J Prosthet Dent 96:433-442
- [2] Craig RG. Chap 9. 11th ed. United States: Mosby Inc; 2002. Restorative dental materials.
- [3] Stein PS, Sullivan J, Haubenreich JE, Osborne PB. Composite resin in medicine and dentistry. J Long Term Eff Med Implants. 2005;15:641–54.
- [4] Irie M, Suzuki K, Watts DC. Marginal gap formation of light-activated restorative materials: effects of immediate setting shrinkage and bond strength. *Dent Mater* 2002;18:203–210.
- [5] Sakaguchi RL, Peters MC, Nelson SR, Douglas WH, Poort HW. Effects of polymerization contraction in composite restorations. *J Dent* 1992;20:178–182.
- [6] Ferracane JL. Buonocore Lecture. Placing dental composites-a stressful experience. Oper Dent2008;33:247-257.
- [7] Ferracane JL, Mitchem JC. Relationship between composite contraction stress and leakage in Class V cavities. *Am J Dent* 2003;16:239–243.
- [8] Willershausen B, Schäfer D, Pistorius A, Schulze R, Mann W. Influence of resin-based restoration materials on cytotoxicity in gingival fibroblasts. Eur J Med Res. 1999;4:149–55.
- [9] Lapp CA, Schuster GS. Effects of DMAEMA and 4-methoxyphenol on gingival fibroblast growth, metabolism, and response to interleukin-1. J Biomed Mater Res. 2002;60:30–5.
- [10] Tanaka M, Matsunaga K, Kadoma Y, et al. Use of fluoride-containing sealant on proximal surfaces. J Med Dent Sci 2000;47(1):49–54.
- [11] Mousavinasab SM. Biocompatibility of composite resins. Dental Research Journal. 2011;8(Suppl1):S21-S29.
- [12] Pu Z, Mark JE, Jethmalani JM, Ford WT. Effects of dispersion and aggregation of silica in the reinforcement of poly(methyl acrylate) elastomers. Chemistry of Materials. 1997;9:2442–2447.
- [13] Hebling J, Feigal R. Use of one-bottle adhesive as an intermediate bonding layer to reduce sealant microleakage on saliva-contaminated enamel. Am J Dent 2000; 13(4):187.
- [14] Kemp-Scholte C, Davidson C. Complete marginal seal of Class V resin composite restorations effected by increased flexibility. J Dent Res 1990;69(6):1240.
- [15] Soh MS, Sellinger A, Yap AU. (2006). Dental nanocomposites. Curr Nanosci 2:373-381
- [16] Grosby AJ, Lee J-Y. (2007). Polymer nanocomposites: the 'nano' effect on mechanical properties. Polym Rev 47:217-229
- [17] Ilie N, Hickel R, Watts DC. (2009). Spatial and cure-time distribution of dynamic-mechanical properties of a dimethacrylate nano-composite. Dent Mater 25:411-418
- [18] Turssi CP, Ferracane JL, Vogel K. (2005). Filler features and their effects on wear and degree of conversion of particulate dental resin composites. Biomaterials 26:4932-4937
- [19] Cavalcante LM, Masouras K, Watts DC, Pimenta LA, Silikas N. (2009). Effect of nanofillers' size on surface properties after toothbrush abrasion. Am J Dent 22:60-64
- [20] Vljanen EK, Lassila LV, Skrifvars M, Vallittu PK. (2005). Degree of conversion and flexural properties of a dendrimer/methyl methacrylate copolymer: design of experiments and statistical screening. Dent Mater 21:172-177
- [21] Schroeder WF, Cook WD, Vallo CI. (2008). Photopolymerization of N,N-dimethylaminobenzyl alcohol as amine co-initiator for light-cured dental resins. Dent Mater 24:686-693
- [22] Kim O. (2005). Effect of photo-accelerator on the polymerization behavior of light-activated polymeric dental restorative composites. J Ind Eng Chem 11:287-292
- [23] Shi S, Nie J. (2007). A natural component as coinitiator for unfilled dental resin composites. J Biomed Mater Res Part B: Appl Biomater 82:44-50

Vol. 5, Issue 2, pp: (182-187), Month: October 2017 - March 2018, Available at: www.researchpublish.com

- [24] Liu SJ, Shi SQ, Hou GY, Nie J. (2007). Benzodioxole derivative as coinitiator for dental resin. Acta Odontol Scand 65:313-318
- [25] Schneider LF, Cavalcante LM, Consani S, Ferracane JL. (2009a). Effect of co-initiator ratio on the polymer properties of experimental resin composites formulated with camphorquinone and phenyl-propanedione. Dent Mater 25:369-375.
- [26] Wu G, Nie J. (2007). Synthesis and evaluation of ethylene glycol 3-diethylamino-propionate methacrylate as a polymerizable amine coinitiator for dental application. Dent Mater 23:623-629
- [27] Nie J, Bowman CN. (2002). Synthesis and photopolymerization of N,N'-dimethyl,-N,N'-di(methacryloxy ethyl)-1,6-hexanediamine as a polymerizable amine coinitiator for dental restorations. Biomaterials23:1221-1226
- [28] Sahin G, Albayrak AZ, Bilgici ZS, Avci D. (2009). Synthesis and evaluation of new dental monomers with both phosphonic and carboxylic acid functional groups. J Polym Sci Part A: Polym Chem 47:1953-1965
- [29] Atai M, Nekoomanesh M, Hashemi SA, Amani S. (2004). Physical and mechanical properties of an experimental dental composite based on a new monomer. Dent Mater 20:663-668.
- [30] López-Suevos F, Dickens SH. (2008). Degree of cure and fracture properties of experimental acid-resin modified composites under wet and dry conditions. Dent Mater 24:778-785
- [31] Moszner N, Fischer UK, Angermann J, Rheinberger V. (2006b). Bis-(acrylamide)s as new cross-linkers for resinbased composite restoratives. Dent Mater 22:1157-1162.
- [32] Xia Y, Zhang F, Xie H, Gu N. (2008). Nanoparticle-reinforced resin-based dental composites. J Dent 36:450-455.
- [33] Bowman CN, Kloxin CJ. (2008). Toward an enhanced understanding and implementation of photopolymerization reactions. AIChE J 54:2775-2795
- [34] Weinmann W, Thalacker C, Guggenberger R. (2005). Siloranes in dental composites. Dent Mater 21:68-74
- [35] Duarte S, Botta AC, Phark JH, Sadan A. (2009). Selected mechanical and physical properties and clinical application of a new low-shrinkage composite restoration. Quintessence Int 40:631-638