Universidade de Lisboa Faculdade de Medicina Dentária



Evaluation of marginal discrepancy on Allceramic crowns manufactured by CAD/CAM versus Conventional methods

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Dissertação

Mestrado Integrado em Medicina Dentária

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Dissertação orientada pelo Prof. Doutor Tiago Mourão

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Abstract:

Introduction: The marginal integrity of a dental prosthesis can determine longevity and predictability. This gap is important because the amount of space will determine the amount of possible cement dissolution. Margin inaccuracy can lead to the accumulation of plaque and bacteria, the dissolution of luting material, and the introduction of unfavorable inflammation or the periodontal tissues. With the introduction of new technologies to fabricate dental ceramic crowns (CAD/CAM), marginal fit is a valuable way to determine the prognosis of the restoration in comparison to the more conventional methods.

Objectives: The objective of this work was to review the current literature in regards of the marginal gap/fit of all-ceramic crowns manufactured by conventional methods (Heat-Pressing and Slip-Casting) versus digital methods (CAD/CAM).

Materials and Methods: A research on PubMed electronic database was conducted for articles with the following combination of key words: (discrepancy or fit or gaps or adaptation) and (disilicate or ceramic) and (copings or crowns). The studies considered for this research were in English from peer-reviewed publications that focused on the evaluation of the marginal fit in ceramic single crowns.

Results: An overall review of the data retrieved for marginal gap showed that 86.8% of the values measured were less than or equal to 120 μ m described by McLean and Von Fraunhofer. The widest marginal gap measured was 180 μ m, and the smallest was 17 μ m. CAD/CAM ceramic crowns showed, an overall, better marginal fit than conventional crowns.

Conclusion: Based on the results obtained, the digital method seems to be a legitimate alternative to the traditional methods. Analysis of the results of this study suggested that the digital method exceeds the standards of clinical acceptability and can sometimes surpass the vertical marginal fit of conventionally fabricated crowns.

Keywords: marginal fit; marginal discrepancy; marginal gap; ceramics; CAD/CAM; coping; disilicate

Resumo:

Introdução: A utilização de qualquer prótese cerâmica em tratamentos restauradores tornou-se popular e muitas destas restaurações podem ser fabricadas por ambos os métodos laboratoriais tradicionais e de CAD/CAM. Os métodos tradicionais de fabricação de cerâmica têm sido descritos como sendo especialmente exigentes em termos de tempo de laboratório bem como tempo clínico, técnica sensível e imprevisível, devido a diversas variáveis. Assim os sistemas de CAD/CAM podem ser uma boa alternativa tanto para os dentistas bem como para os laboratórios. A tecnologia CAD/CAM pode também reduzir o tempo de fabricação de cerâmica de alta resistência, tais como InCeram (Vita Zahnfabrik, Bad Sackingen, Alemanha), até 90%. Além disso, os blocos fabricados industrialmente apresentam-se mais homogéneos traduzindo-se em menor número de defeitos intrínsecos. Podemos, assim dizer, que os avanços na tecnologia CAD/CAM são fundamentais para a pesquisa e desenvolvimento de cerâmicas de alta resistência, tais como o dióxido de zircônio estabilizado que não poderiam ser praticamente processado por métodos laboratoriais tradicionais. Estes materiais tornaram possível a utilização de coroas cerâmicas bem como pontes em espaços posteriores com elevadas cargas oclusais.

A integridade marginal de uma prótese dentária pode determinar a longevidade e a sua previsibilidade a longo prazo, logo a sua mensuração requer avaliação precisa e quantificação dos parâmetros marginais. Holmes *et al.* definiram geometricamente a relação da linha cavo-superficial da preparação dentária com a margem da prótese em termos de oito variáveis: diferença interna, "gap marginal", discrepância marginal vertical e horizontal, margem sobre-extendidas, margem sob-extendida, discrepância marginal absoluta e discrepância de adaptação. A adaptação marginal, vertical ou na horizontal. Esta diferença é importante porque a quantidade de espaço irá determinar a quantidade de dissolução possível de cimento. Imprecisão ao nível da margem da restauração pode levar à acumulação de placa bacteriana, dissolução do material de cimentação, e aparecimento de inflamação nos tecidos periodontais. McLean e Von Fraunhofer descreveram discrepâncias marginais clinicamente aceitáveis de $\leq 120 \mu m$. No entanto, de acordo com a American Dental Association (ADA) Especificação N ° 8, o espaço marginal de restaurações cimentadas deve estar entre os 25-40 µm para permitir a espessura adequada para a cimentação, no entanto este intervalo raramente é realizável.

Objectivos: O objectivo deste trabalho foi realizar uma revisão da literatura sobre a adaptação marginal de coroas cerâmicas totais fabricadas por métodos convencionais, em comparação com coroas em cerâmica total fabricadas por métodos digitais (CAD/CAM).

Materiais e Métodos: Uma pesquisa na base de dados electrónica PubMed foi realizada com a seguinte combinação de palavras-chave: (discrepancy or fit or gaps or adaptation) and (disilicate or ceramic) and (copings or crowns). A última pesquisa foi realizada a 25 de abril de 2015. Os estudos considerados para esta pesquisa foram em Inglês a partir de publicações revistas cientificamente, que abordavam a avaliação da adaptação marginal ou coroas individuais totalmente em cerâmica. Estudos "in vivo" e "in vitro" foram incluídos. Artigos que incidiam sobre a adaptação marginal de restaurações não-cerâmicas não foram considerados para esta revisão. Foram ainda excluídos estudos que incluíssem próteses parciais fixas, facetas, "inlays", "onlays", coroas parciais, restaurações diretas, coroas metalo-cerâmicas e restaurações implantosuportadas. Estudos que mediram a adaptação marginal de coroas cerâmicas fabricadas por sistemas menos populares ou desatualizados foram excluídos. Após a leitura e análise do resumo de uma artigo possível para ser incluído, o texto integral do artigo foi revisto e sujeito aos critérios de inclusão e exclusão. A pesquisa eletrônica também foi complementada por uma pesquisa manual através das referências dos artigos selecionados. Os seguintes dados foram extraídas de cada artigo: tipo de sistema estudado, estágio de conclusão da restauração, tamanho da amostra, tipo de limite da preparação dentária, a ocorrência de cimentação, método de exame da adaptação marginal e os valores da discrepância marginal absoluta ou "gap" marginal.

Resultados: Entre 525 relatórios identificados através da pesquisa eletrônica, 15 foram selecionados, todos publicados entre 2005 e Abril de 2015. Todos os estudos foram realizados "in vitro". Uma análise global dos dados obtidos para a adaptação marginal mostrou que 86,8% dos valores medidos foram inferiores ou iguais a 120 μ m como descrito por McLean e Von Fraunhofer. A maior diferença marginal medida foi de 180 μ m, e a menor foi de 17 μ m. Quatro estudos afirmam que o método

convencional demonstra melhores resultados de adaptação marginal, dois estudos mostraram existir nenhuma diferença significativa entre os métodos convencionais e de CAD/CAM e nove estudos mostraram melhores resultados para os grupos de CAD/CAM.

Discussão: O ajuste marginal ao nível das coroas cerâmicas é fundamental para o sucesso da restauração; coroas com ajuste deficiente estão propensas a falhas devido a micro-infiltração, dissolução do cimento e cárie dentária. Neste trabalho, a adaptação das coroas foi avaliada com base na medição da discrepância marginal vertical, que foi escolhido como o fator mais crítico de fenda marginal, no entanto será o fator menos suscetível à manipulação de pós-fabricação, conforme indicado por Holmes *et al.* Discrepâncias horizontais, como as saliências da coroa, podem ser modificadas até certo ponto intra-oralmente, no entanto, a discrepância marginal vertical, apenas pode ser fechado com cimento de cimentação, o que é propenso a dissolution. Por esta razão, este tipo de adaptação vertical tem a maior relevância clínica e deve ser considerado como o factor mais importante na avaliação da margem da coroa.

Restaurações em CAD/CAM estão atualmente a ser utilizadas por um grande número de dentistas em todo o mundo; no entanto, a precisão desses sistemas ainda é questionável, apresentando bons resultados em alguns estudos. No entanto, a precisão da aquisição de dados varia de acordo com várias tecnologias impressão óptica do sistema e do fabricante. A precisão do software de design bem como a tecnologia de fresagem também sofrem de diferenças entre os sistemas. Além disso, dentro do mesmo sistema, podem haver diferenças substanciais entre os valores de medição que podem ser explicadas por diferentes protocolos experimentais utilizados em cada estudo. A utilização do método convencional de fabricação de coroas tem sido utilizado durante décadas com resultados comprovados de longo prazo, tanto para a sobrevivência e longevidade. A seleção cuidadosa de materiais e procedimentos de fabricação meticulosos são necessários para compensar as expansões e contração dos diferentes materiais envolvidos na criação de uma coroa. No entanto, a impossibilidade de controlar todas as variáveis, combinada com a propensão para o erro humano, pode resultar em má adaptação marginal ou mesmo desajuste. O uso de um método digital parece diminuir a margem de erro. A intervenção humana no fabrico da coroa pode desempenhar um papel relativo à perícia do técnico de laboratório dentário sendo uma variável difícil de controlar. O número de passos envolvidos no processo é um outro elemento importante porque a probabilidade de erro aumenta consideravelmente com cada passo adicional necessário. Por exemplo, os sistemas não-CAD / CAM não apresentam a necessidade de aplicação de um espaçador por parte de um técnico, e o tradicional sistema In-Ceram foi descrito como técnica singularmente sensível.

Conclusão: Com base nos resultados obtidos, o método digital parece ser uma alternativa legítima para os métodos tradicionais. A análise dos resultados deste estudo sugeriram que o método digital excede as normas de aceitação clínica e, por vezes, pode superar a adaptação marginal vertical de coroas convencionalmente fabricadas.

Palavras-chave: adaptação marginal; discrepância marginal; cerâmicas; CAD/CAM; sistemas digitais; disilicato Evaluation of marginal discrepancy on All-ceramic crowns manufactured by CAD/CAM versus Conventional methods

Chapter 1: Introduction

Dental crowns have been used for many years to restore compromised or heavily restored dentition, and for esthetic changes and improvements. New CAD/CAM materials and systems have been developed and evolved in the last decade for fabrication of all-ceramic restorations. Dental CAD/CAM technology is gaining popularity because of its benefits in terms of manufacturing time, material savings, standardization of the fabrication process, predictability of the restorations and economic value. When the CAD/CAM manufacturing process is employed, the number of steps required for the fabrication of a restoration is less compared to traditional methods, which can bring fewer errors to the process. Another benefit of CAD/CAM dentistry includes the use of contemporary materials and data acquisition instruments; which represents a non-destructive method of saving impressions, restorations and information that are saved on a computer and constitutes an extraordinary communication tool for evaluation. Cooper (2011) stated that: "CAD/CAM technology is an efficient and effective point for critical evaluation of the proposed restorations prior to its fabrication". The incorporation of dental technology has not only brought a new range of manufacturing methods and material options but also some concerns about the processes involving restorations fit, quality, accuracy, short and long-term prognosis (MIYAZAKI et al. 2009).

1. Ceramics used for the fabrication of permanent dental crowns

1.1. Glass Ceramics

1.1.1. Feldspathic Glass ceramics

Feldspathic glass ceramics are silica-based ceramics with low to moderate crystalline leucite filler ($K_2O.AL_2O_3.4SiO_2$) with around 5-25% of volume, which is created by firing feldspar at 1150°C (DENRY *et al.* 1996; GIORDANO and MCLAREN 2010). This high glass content in the feldspathic ceramics results in excellent aesthetic properties resembling the natural tooth substance (PJETURSSON *et al.* 2007). Leucite particles are used to provide high translucency and alter the coefficient of thermal expansion, as well as to improve the material strength by inhibiting crack propagation. However, the original feldsphatic ceramics have a random distribution and large size of leucite particles, which contributes to the material's low fracture strength (FISCHER *et al.* 2008), so they are commonly used as a veneering

ceramic for metal-ceramic restorations (GIORDANO and MACLADEN 2010). In 1985, Vita Mark I blocks (VITA Zahnfabrik, Bad Säckingen, Germany) were developed, becoming the first glass ceramics for the Sirona CAD/CAM system (Sirona Dental System, Bensheim, Germany). This material had a flexural strength of around 120MPa (GIORDANO *et al.* 1995) and was intended to be used for fabrication of inlays, onlays and veneers. New generations with around 30% by volume fine grain (10µm to 20µm) and very evenly distributed particles we developed (TINSCHERT *et al.* 2000, GIORDANO and MCLAREN 2010) in 1991 as Vita Mark II block (VITA Zahnfabrik, Bad Säckingen, Germany). The fine crystal microstructure and the CAD/CAM processing technique produce the enamel-like abrasion characteristic of this material (KREJCI *et al.* 1994). According to the manufacturer data, this material is suitable for the fabrication of inlays, onlays and monolithic anterior crowns and veneers (POSSELT and KERSCHBAUM 2003, FRADEANI *et al.* 2005). This material can also be etched with hydrofluoric acid to create micromechanical retention for adhesive cementation (FASBINDER 2002, OTTO 2004).

1.1.2. Leucite-reinforced glass ceramics

The glass matrix in leucite-reinforced ceramic is based on an alumino-silicate glass. A high proportion of leucite crystal ranging from 35% to 45% by volume (DEANY 1996), is used to reinforce the glass ceramic and improve its biomechanical properties. Adding more leucite can increase the flexural strength of glass ceramic up to 105-120 MPa (SEGHI et al. 1990). Leucite reinforced are highly translucent (HEFFERNAN et al. 2002). The type of ceramic was first introduced as VITA VMK 68 ceramic (VITA Zahnfabrik, Bad Säckingen, Germany) in 1968 in powder/liquid form as metal-ceramic veneering material (GUESS el at. 2011). To improve this powder/liquid ceramics in terms of micro-porosity and shrinkage, the IPS Empress ceramic (Ivoclar Vivadent, Schaan, Liechtenstein) was introduced in 1990 and must be the most widely used leucite-reinforced pressable ceramic (GIORDANO and MCLAREN 2010). The ceramic ingots, supplied by the manufacturer in a variety of shades, can be pressed under heat (1050-1080°C) and pressure (0.3-0.4 MPa) (GONZAGA et al. 2008). The produced ceramic microstructure consists of uniformly distributed leucite crystals in a glassy matrix with a size range between 1-5µm (ET 2008). Fine leucite crystals and heat-pressing techniques have contributed to the increased material flexural strength of 160-180MPa (GROTEN and PROBSTER 1997). This ceramic material is indicated for inlays, onlays, veneers or crown restorations in anterior teeth (FRADEANI and REDEMAGNI 2002). IPS Empress CAD (Ivoclar Vivadent, Schaan, Liechtenstein) is the CAD/CAM machinable version. It was introduced in 2006 with flexural strength of around 160 MPa and designed to be used either with chairside or in lab-side CEREC systems to fabricate veneers, inlays, onlays and anterior crowns (GIORDANO and MCLAREN 2010).

1.1.3. Lithium disilicate glass ceramics

In order to construct anterior three-unit all-ceramic bridge restorations, a glass ceramic based on lithium disilicate (Li₂Si₂O₅) was developed. Arranged in a dense way lithium disilicate crystals had a concentration of 70% by volume (GUAZZATO et al. 2004) with a length of $4\mu m$ and a diameter of $0.5\mu m$, and are uniformly distributed in a glass matrix. This interlocking structure prevents crack propagation and elevates the flexural strength of lithium disilicate ceramic to 300-400 MPA, which is more than twice the strength of leucite-reinforced glass ceramic (DENRY and HOLLOWAY 2010). Ivoclar Vivadent introduced the first lithium disilicate ceramic (IPS Empress II, Ivoclar Vivadent, Schaan, Lichtenstein) in 1998 as an ingot form to be used with the press technique at approximately 920°C. Further improvement in physical properties and translucency of lithium disilicate glass ceramics was provided with the introduction of IPS e.max Press (Ivoclar Vivadent, Schaan, Liechtenstein) (STAPPERT et al. 2006). Pressable ingots are available in a variety of opacities, from high opacity to high translucency. This material is recommended in the fabrication of monolithic inlays, onlays and posterior crowns, or as a core for crowns and anterior 3-unit fixed dental prostheses (FDPs) (HOLLAND et al. 2000). As CAD/CAM production of dental restorations has become more common, a new innovation in lithium disilicate glass ceramics was developed in 2005 as IPS e.max CAD (Ivoclar Vivadent, Schaan, Liechtenstein) for milling techniques. The IPS e.max CAD block is a partially crystallized block consisting of 40% lithium meta-silicate crystals, allowing the material to be easily milled. After processing the block, a recrystallization process takes place at 850°C for 10 minutes, through which the lithium meta-silicate is transformed into lithium disilicate crystals. This transformation provides the restoration with its final mechanical and aesthetic properties. According to the manufacturer's data, the flexural strength of a fully crystallized IPS e.max CAD is about 360 MPA. This material is indicated for the fabrication of monolithic inlays, onlays, single crowns, and anterior FDPs, but also for short posterior FDPs (HOLLAND *et al.* 2008), with either conventional or adhesive cementation (BINDL *et al.* 2006).

1.2. Yttrium-tetragonal zirconia polycrystal ceramics (Y-TZP)

Zirconia in a pure state is polymorphic and exhibits three crystallographic phases at different temperatures: a cubic phase (c), stable from 2680°C to 2370°C, a tetragonal phase (t) stable from 2370°C to 1170°C and a monocyclic phase (m), stable from 1170°C to room temperature (DENRY and KELLY 2008). This transformation is associated with substantial volume increase (4%), and causes high internal stress, which can induce severe cracking (GUAZZATO et al. 2005). Addition of minor components such as magnesium oxide (MgO), calcium oxide (CaO), or yttrium oxide (Y_2O_3) to pure zirconia provides formation of multiphase materials known as partially stabilized zirconia (PSZ) at room temperature (CATTANI-LORENTE et al. 2011). Advances in CAD/CAM technology enable the use of zirconia in dentistry. Two CAD/CAM processing techniques are available, hard processing and soft processing of zirconia blanks (VAGKOPOULOU et al. 2009). The first method is involves milling fully sintered zirconia blanks to the desired framework shape and diminution. Unfortunately, fully sintered zirconia requires special milling equipment and long processing times (GIORDANO and MCLAREN 2010). The second method is based on milling partiallysintered zirconia blanks. Enlarged frameworks are designed and fabricated using CAD/CAM technology to compensate for about 20-25% material shrinkage after final sinter firing at 1300-1500°C for around 2-6 hours (SUTTOR et al. 2001). The most frequently used CAD/CAM systems for the processing of pre-sintered zirconia include CERCON (Dentsply Friadent, Mannheim, Germany), CEREC (Sirona, Bensheim, Germany), LAVA (3M ESPE, Seefeld, Germany), and Procera (Nobel Biocare, Gothenburg, Sweden). Recently, monolithic, fully anatomic zirconia ceramic restorations have been introduced to serve in high stress-loading posterior teeth, to avoid chipping failure as with veneering glass-ceramic. Lava all-Zirconia (3M ESPE, Seefeld, Germany), Zircon Zahn (ZIRCONZAHN GMBH, Bruneck, Italy), and BruxZir Solid Zirconia (Gildewell laboratories, California, USA) have been introduced to the market as all-zirconia monolithic restorations. According to the manufacturers, allzirconia monolithic restorations are indicated for fabrication of posterior single crowns restorations in patients with parafunctional habits or limited occlusal space. However, as zirconia is a high value with opaque material, staining the restoration prior to sintering

develops the desired tooth shade. Due to the inferior aesthetic properties of the monolithic zirconia ceramic, its application is restricted to the less aesthetically demanding posterior area (HOLT and BOKSMAN 2012, GRIFFIN 2013).

1.3. Bi-layered materials

Despite the superior aesthetic appearance of ceramic restorations, brittleness and susceptibility to fracture in high-stress areas are the common disadvantages of ceramic materials. In order to overcome this problems high-strength core materials were needed. These high strength ceramics tend to be opaque and therefore require veneering with glass ceramic to achieve a natural aesthetic look.

1.3.1. Aluminium oxide ceramics

The first application of aluminium oxide (Al₂O₃) in all-ceramic dental restorations was with the development of In-ceram Alumina (VITA Zahnfabrik, Badsäckingen, Germany) in 1989 (HASELTON et al. 2000). A infrastructure is produced by sintering a slurry of densely packed aluminium oxide (70-80%) at 1120°C for 10 hours, followed in a second stage by infiltration of a lanthanum silicate glass at 1100°C for 4 hours (XIAO-PING et al. 2002). Either traditional slip casting or CAD/CAM processing of pre-sintered blocks with CEREC (Sirona dental system, Charlotte, NC) can be used for fabrication of the ceramic core (BINDL and MORMANN 2002; GIORDANO and MCLAREN 2010). The aesthetic appearance of the restoration is achieved by veneering the core with feldspathic porcelain (HASELTON et al. 2000). The produced material has a high flexural strength of around 450 MPa and moderate translucency, making it suitable for fabricating anterior and posterior single crowns (GIORDANO and MCLAREN 2010). Increased attention to improve the ceramic core materials led to the development of Procera In-Ceram Alumina crowns (Nobel Biocare, Gothenburg, Sweden) in 1993. Densely sintered pure alumina consisting of 99.9% alumina oxide of 5 µm grain size is formed by compacting alumina powder into an enlarged die under high pressure, and then sintering at about 1600°C (ANDERSSON and ODEN 1993). This technique compensates for about 20% shrinkage of the alumina core, which is the veneered with feldspathic porcelain (ANDERSSON and ODEN 1993). Flexural strength of approximately 490-700 MPa were reported (RAIGRODSKI 2004). In-Ceram Spinell (VITA Zahnfabrik, Badsäckingen, Germany), is an oxide ceramic based on a magnesium-aluminum mixed

oxide, and was first introduced in 1993 with an improved dentin-like translucency (WASSERMANN *et al.* 2006). In-Ceram Zirconia (VITA Zahnfabrik, Badsäckingen, Germany), is an alumina oxide ceramic reinforced with 35% partially stabilized zirconium oxide. It was introduced in 1999 with a flexural strength of 420-800 MPa (GUAZZATO *et al.* 2002). Due to its high strength the material is suitable for fabrication of posterior single crown restorations and 3-unit FDPs (WASSERMANN *et al.* 2006). In-Ceram zirconia frameworks can be made either by using conventional slip casting or CAD/CAM processing techniques (TINSCHERT *et al.* 2000).

1.3.2. Zirconia Ceramics

Zirconium dioxide (ZrO₂), is a glass-free ceramic material formed by the addition of oxygen to the pure, elemental zirconium metal (PICONI and MACCAURO 1999). Zirconia in a polycrystalline form is a white opaque material with high flexural strength ranging from 900 to 1200 MPa and a high fracture toughness (MANICONE et al. 2007). The absence of a glassy-phase in zirconia impairs the effectiveness of the traditional hydrofluoric acid etching to aid adhesion (DERAND et al. 2005). Therefore, several surface treatments, especially airborne abrasion and selective infiltration etching, have been reported to facilitate the bond strength between resin cement and zirconia ceramics (ABOUSHELIB et al. 2007). A weak point in bi-layered zirconia CAD/CAM fabricated dental restorations is their need to be veneered with low strength glass ceramics. Chipping of the veneer ceramic layer is the most widely reported failure mode with this system (BEUER et al. 2009; GUESS et al. 2010). By using soft or hard CAD/CAM machining, dental restoration frameworks can be fabricated. Examples of (YTZP) blocks include Lava Frame (3M ESPE), Everest ZS and ZH (KaVo), In-Ceram YZ (VITA), Zerion (Straumann), and Cercon Smart Ceramics (DeguDent) (BEUER et al. 2008).

2. CAD/CAM technology

2.1. History

The acronym CAD/CAM is the abbreviation for computer aided design/computer aided manufacturing. The technology of computer aided design (CAD) applies the use of computer systems to assist in the creation, modification, analysis or optimization of a design (ET 2008), and computer aided manufacturing (CAM) applies

the use of the computer systems that plan, manage and control the manufacturing operations (PICHLER *et al.* 2000). This technology was first developed in the 1960s and used in aircraft industries. A decade later, Dr. Francois Duret was the first to develop a dental CAD/CAM device known as the Sopha System (Sopha Bioconcept, Inc.Los Angeles USA) (DURET and PRESTON 1991). However, due to high cost and complexity of use, the Sopha System was unsuccessful in the dental market. In the early 1980s, Dr. Mörmann and his team succeeded in developing the first dental chairside CAD/CAM system known as the CEREC system (MÖRMANN 2004). Digital impression of an inlay prepared cavity was performed using optical intra-oral camera, and digitized data was used to design and fabricate the first single visit chairside CAD/CAM inlay restoration. In 1983, a CAD/CAM technology for fabricating composite veneered restorations was introduced. This systems was later known as the Procera System (ANDERSSON and ODEN 1993). Since then many different systems have been introduced to the dental market.

2.2. CAD/CAM Components

Every developed dental CAD/CAM systems are composed of three basic components (BEUER et al. 2008);

2.2.1. Scanner

The scanner is one of the most critical components of any dental CAD/CAM system, since the accuracy of the design is limited by the accuracy of the captured and imported data (FASBINDER 2010). A digital scanner collects 3-dimensional data of the prepared teeth, neighboring structures and opposing teeth either intra-orally or extraorally from cast models. Following image acquisition, the final data is either used for chairside fabrication of restorations or digitally transmitted to a laboratory. Today, there are many different scanning devices. The most widely used is the optical scanner, in which a laser or white light source is used based on a triangulation procedure to capture several static or video images of the prepared tooth surfaces (CEREC, Lava Scan, Everest Scan) (BEUER *et al.* 2008). To enhance the intra-oral scanning quality, application of a high reflective oxide powder to the scanned tooth surfaces is required in some optical scanner systems (RONALD *et al.* 2011). The first intra-oral digital scanner (CEREC) was introduced in the early 1980s by Dr. Mörmann and Brandstinian, and has since been upgraded (MORMANN 2006). Whereas the earlier versions of CEREC were powered by and infrared camera, advances in the performance of short-wavelength blue light (BlueCam, Sirona dental system, Bensheim, Germany) have surpassed the quality of longer-wavelength infrared cameras. According to manufacturer data, the shorterwavelength intense blue light allows for higher precision in the captured optical image. However, it does continue to require an optical powder to properly image the desired area. Recently the CEREC Omnicam scanner (Sirona dental system, Bensheim, Germany) was introduced in the market. This system provided the advantages of the unrivalled handling and powder-free scanning with precise 3D images in natural colors.

2.2.2. Design Software

Design software can be defined as a computer unit equipped with software programs for visualization of the scanned data, planning and designing 3D dental restorations (ET 2008). A variety of dental restorations can be designed, including inlays, onlays, single crowns, copings and fixed dental prostheses. Software engineer Alain Ferru in cooperation with Mörmann designed the first dental software. Using the anatomy of natural tooth and the collected intra-oral preparation data, the CEREC 1 software was able to design the first chairside CAD/CAM inlay restoration (MORMANN 2006). The design was displayed two-dimensionally. With subsequent development of CEREC 2 software in 1994, the dentist was able to design and fabricate full crown restorations and copings. However, the design was still displayed in a 2D format. Partially due to recent improvements in computer speed and memory, the CEREC 3, with 3D capability, was introduced in 2005 (FASBINDER 2010). The software has become much simpler and enables automatic virtual occlusal adjustment (MORMANN 2006). This generated 3D data can be transformed into various data formats. Standard transformation language (STL) is used with open systems and allows free choice among different CAM processing systems (WITKOWSKI 2005). However, many closed systems are linked through the specific data format of the user (BEUER et al. 2008).

2.2.3. Processing devices

Virtual restorations provided by CAD software systems are converted to dental restorations using computer-controlled milling devices. A variety of prefabricated material blocks, such as ceramics, composites and metals can be machined in different axes to produce the desired restoration (RONALD *et al.* 2011). Final manual correction,

polishing and staining must be carried out by the dental technician. Two processing CAD/CAM systems can be defined.

2.2.3.1. Chairside system

With this system, all the components of the CAD/CAM system are located in the dental office, which offers the dentist the ability to fabricate a tooth-colored restoration in one appointment (BEUER *et al.* 2008). Different dental ceramic material blocks for chairside milling are available. Currently there are two chairside CAD/CAM systems, CEREC (Sirona dental system, Bensheim, Germany) and E4D (D4D Technologies, Texas, USA) in the market. The CEREC system is the most widely used and it is found to be well documented. CEREC AC (Sirona dental system, Bensheim, Germany) is the newest version of CEREC, which provide the ability to fabricate chairside dental restorations in one visit. Through the Sirona digital network (CEREC Connect) the optical impression can be also send by email to the dental laboratory. Recently, CEREC AC, powered by Omnicam, was introduced in the market. This systems provides the advantage of powder-free scanning and wide indication spectrum to fabricate chairside inlays, onlays, veneers, single crowns, bridges and surgical guides.

2.2.3.2. Lab-side system

All the component and production steps of a CAD/CAM system are located in the laboratory (BEUER *et al.* 2008). To generate 3D data of the preparation, a conventional dental impression is used to produce a master cast, which is later digitally scanned, or chairside digitally scanned data can be sent or mailed from the dental office to a laboratory. Many of the lab-side systems, such as Lava (3M ESPE, St.Paul, USA), Everest (KaVo, Biberach, Riss) or Cerec InLab (Sirona, Bensheim, Germany), produce monolithic restorations and copings and frameworks, which later require veneering with either manual or CAD/CAM techniques.

2.2.4. CAD/CAM technique

There are two types of processing techniques capable of generating a desired geometry of a dental restoration.

2.2.4.1. Substractive technique

The traditional CAD/CAM manufacturing of dental restorations is based on the subtractive technique in which sharp machine tools such as diamond drills are used to cut sintered or pre-sintered material blocks to the desired geometry. Computer programs are used to control all steps of the manufacturing (VAN NOORT 2012). Although this technique allows manufacturers to fabricate a more sophisticated dental restoration, considerable waste in the raw material drives manufacturers to save costs by using the additive technique (EBERT *et al.* 2009).

2.2.4.2. Additive technique

Instead of machining sintered or pre-sintered ceramic blocks, a 3D component can be built up layer-by-layer using a computerized numerical control (CNC) machine (SILVA *et al.* 2011b). Five additive manufacturing processes are present (EBERT *et al.* 2009): 1. Stereolithgraphy; 2. 3D printing; 3. Selective laser sintering; 4.Selective laser melting; 5. Direct inject printing. Additive manufacturing can produce complex shapes with little or no waste of materials, however it should be remembered that the materials currently being used are not suitable for medical industries (VAN NOORT 2012).

3. Conventional Methods

3.1. Slip Casting

Slip-cast ceramics for dental restorations were introduced in mid 1990s. A porous infrastructure is produced by slip-casting, sintered, and later infiltrated with a lanthanum-based glass, producing two interpenetrating continuous networks, one composed of the glassy phase and the other being the crystalline infrastructure. Three crystalline phases are available, namely alumina (Al2O3), spinel (MgAl2O4) and zirconia-alumina (12 Ce-TZP-Al2O3). Alumina-based slip-cast ceramics contain 68 vol % alumina, 27 vol % glass and 5 vol % porosity (GUAZZATO *et al.* 2004). The microstructure consists of blocky alumina grains of various sizes and shapes. Evidence of grain pull-out, bridging and crack deflection was reported with this type of ceramic (GUAZZATO *et al.* 2004), indicative of efficient crystalline reinforcement, and accounting for mechanical properties in the range of heat-pressed lithium disilicate glass-ceramics. It has also been suggested that the coefficient of thermal expansion mismatch between the alumina crystals and the infiltration glass could contribute to

strengthening due thermal residual stresses. The presence of large alumina crystals with a high refractive index, and a non-negligible amount of porosity, account for some degree of opacity in this all-ceramic system. Spinel-based slip-cast ceramics offer better translucency (HEFFERNAN *et al.* 2002), similar to that of lithium disilicate heatpressed ceramics, at the expense of mechanical properties (JUNG *et al.* 1999)

3.2. Heat-Pressing

The popularity of heat-pressed ceramics relies on the ability to use the lost-wax technique to produce dental ceramic restorations. Dental technicians are usually familiar with this technique, commonly used to cast dental alloys. In addition, the equipment needed to heat-press dental ceramics is relatively inexpensive. The first generation of heat-pressed dental ceramics contains leucite as reinforcing crystalline phase. The second generation is lithium disilicate-based. First generation heat-pressed ceramics contain between 35 and 45 vol % leucite as crystalline phase (DENRY et al. 1995). Flexural strength and fracture toughness values that are about two times higher than those of feldspathic porcelains (SEGHI et al. 1995). This increase in strength and toughness was explained by dispersion of fine leucite crystals from the heat pressing process (GUAZZATO et al. 2004). It should be noted, however, that coalescence of micro cracks can also cause decoupling of the crystals from the matrix and lead to degradation in strength and fracture toughness (MACKERT et al. 1996). The presence of about 9% porosity should also be considered, when analyzing the mechanical properties of this system (GUAZZATO et al. 2004). Further work revealed that the flexural strength of these ceramics was significantly improved after additional firings, due to additional leucite crystallization (DONG et al. 1992).

Second generation heat-pressed ceramics contain about 65 vol % lithium disilicate as the main crystalline phase, with about 1% porosity (GUAZZATO *et al.* 2004). Lithium disilicate glass-ceramics have been extensively studied (HÖLAND *et al.* 2006). All studies seem to agree that the mechanisms leading to the crystallization of lithium disilicate in these systems are somewhat complex, due to the presence of nanosized crystal phases (BOROM *et al.* 1975). High temperature X-ray diffraction studies revealed that both lithium metasilicate (Li2SiO3) and cristobalite (SiO2) form during the crystallization process, prior to the growth of lithium disilicate (Li2Si2O5) crystals (HÖLAND *et al.* 2006). The final microstructure consists of highly interlocked lithium disilicate crystals, 5 μ m in length, 0,8 μ m in diameter. The interlocked microstructure and layered crystals are also likely to contribute to strengthening. Crack propagation is easy along the cleavage planes, but more difficult across the planes, leading to multiple crack deflections due to an array of crystal orientations.

4. Marginal Fit

The marginal integrity of a dental prosthesis can determine longevity and predictability, and its measurement requires accurate assessment and quantification of marginal parameters so as to differentiate fir from misfit. Holmes et al. defined geometrically the relation of the cavosurface finish line to the prosthesis margin and defined fir in terms of eight variables: internal gap, marginal gap, vertical marginal discrepancy and horizontal marginal discrepancy, overextended margin, under extended margin, absolute marginal discrepancy and seating discrepancy (HOLMES et al. 1989). Fit has been described in both in vitro and in vivo studies as the marginal discrepancy, either vertically or horizontally (GARDNER et al. 1982). This gap is important because the amount of space will determine the amount of possible cement dissolution. Margin inaccuracy could lead to the accumulation of plaque and bacteria (GRASSO et al. 1985), the dissolution of luting material (JACOBS et al. 1991), and the introduction of unfavorable inflammation or the periodontal tissues (JANENKO et al. 1979). McLean and Von Fraunhofer described clinically acceptable marginal gaps of $\leq 120 \ \mu m$ (MCLEAN and VON FRAUNHOFER 1971). However, according to the American Dental Association (ADA) specification No. 8, the marginal fir of cemented restorations should be in the range of 25-40 µm to allow for luting cement thickness, however this range is rarely achievable (MAY et al. 1998).

Evaluation of marginal discrepancy on All-ceramic crowns manufactured by CAD/CAM versus Conventional methods

Chapter 2: Objective / Materials and Methods

1. Objective

The objective of this work was to review the current literature in regards of the marginal gap/fit of all-ceramic crowns manufactured by conventional methods (Heat-Pressing and Slip-Casting) versus digital methods (CAD/CAM).

2. Materials and Methods

A research on PubMed electronic database was conducted for articles with the following combination of key words: (discrepancy or fit or gaps or adaptation) and (disilicate or ceramic) and (copings or crowns). The last search was conducted on April 25th of 2015. The studies considered for this research were in English from peerreviewed publications that focused on the evaluation of the marginal fit or ceramic single crowns. Both in vivo and in vitro studies were included. Articles that focused on the marginal fit of restorations other than ceramic restorations were not considered for this review. This excluded studies of partial fixed dental prostheses, veneers, inlays, onlays, partial crowns, direct restorations, cast crowns, metal ceramic crowns, studies that focused only on the marginal adaptation of conventionally manufactured ceramic crowns/copings implant-supported restorations and studies that focused only on the marginal adaptation of ceramics manufactured by digital systems. Studies that measured the marginal fit of ceramic crowns manufactured by less popular or outdated systems were excluded. After the identification of an abstract for possible inclusion, the full text of the article was reviewed, and subject to the inclusion and exclusion criteria. The electronic search was also supplemented by manual searching through the references of selected articles.

The following data was extracted from each article: type of system studied, stage of completion of the restoration, sample size, type of finish line, occurrence of cementation, examination method and value of the absolute marginal discrepancy or marginal gap measured.

Inclusion Criteria	Exclusion Criteria
 English language 	 Partial fixed dental prostheses
 Single crowns 	 Veneers
 All-Ceramic 	 Inlays / Onlays
 In-vivo 	 Partial crowns
 In-vitro 	 Direct restorations
 Coping 	 Cast crowns
 Crown 	 Metal ceramic crowns
• Studies that focused	 Implant-supported restorations
simultaneously on the marginal adaptation of all-ceramic crowns/copings using conventional and digital methods	 Studies about marginal adaptation on all-ceramic crowns/copings made by conventional methods
	 Studies about marginal adaptation on all-ceramic crowns/copings made by digital systems

Table 1: Inclusion and Exclusion Criteria for the selection of the studies

Evaluation of marginal discrepancy on All-ceramic crowns manufactured by CAD/CAM versus Conventional methods

Chapter 3: Results

Among 525 reports identified through the electronic search, 15 were selected, all published between 2005 and April 2015. All the studies were conducted in vitro. An overall review of the data retrieved for marginal gap showed that 86.8% of the values measured were less than or equal to 120 µm described by McLean and Von Fraunhofer. The widest marginal gap measured was 180 µm, and the smallest was 17 µm. The selected articles displayed a significant heterogeneity in terms of experimental protocols, which led to different discrepancies being measured, sometimes even for the same system. Four reports stated that the conventional method showed better MG values (PELEKANOS *et al.* 2009; MOUSLY *et al.* 2014; SULAIMAN *et al.* 1997; BESCHIT and STRUB 1999). Two studies reported no significant difference between the conventional methods and CAD/CAM (RINKE *et al.* 1995, ANADIOTI *et al.* 2003; QUINTAS *et al.* 2004; BINDL and MORMANN 2005; KORKUT *et al.* 2011; YUKSEL and ZAIMOGLU 2011; ASAPVAPANUMAS *et al.* 2013; NG *et al.* 2014; DEMIR *et al.* 2014; PIMENTA *et al.* 2015).

Seven studies measured the MG on crowns (RINKE *et al.* 1995; BESCHIT and STRUB 1999; YEO *et al.* 2003; ANADIOTI *et al.* 2013; NG *et al.* 2014; MOUSLY *et al.* 2014; DEMIR *et al.* 2014), one did not report where they made the measurements (SULAIMAN *et al.* 1997) and seven measured on copings (QUINTAS *et al.* 2004; BINDL and MORMANN 2005; PELEKANOS *et al.* 2009; KORKUT *et al.* 2011; YUKSEL and ZAIMOGLU 2011; ASAPVAPANUMAS *et al.* 2013; PIMENTA *et al.* 2015). Five studies used a finish line in shoulder (RINKE *et al.* 1995; BESCHIT and STRUB 1999; YEO *et al.* 2003; ASAPVAPANUMAS *et al.* 2013; DEMIR *et al.* 2014), seven used a chamfer (BINDL and MORMANN 2005; PELEKANOS *et al.* 2013; DEMIR *et al.* 2014), seven used a chamfer (BINDL and MORMANN 2005; PELEKANOS *et al.* 2014; MOUSLY *et al.* 2014; PIMENTA *et al.* 2015) and 3 didn't report the kind of finish line in their preparations (SULAIMAN *et al.* 1997; QUINTAS *et al.* 2004; ANADIOTI *et al.* 2013).

Nine studies presented their results without cementing the crowns/copings (RINKE *et al.* 1995; SULAIMAN *et al.* 1997; YEO *et al.* 2003; PELEKANOS *et al.* 2009; ASAPVAPANUMAS *et al.* 2013; ANADIOTI *et al.* 2013; NG *et al.* 2014; MOUSLY *et al.* 2014; PIMENTA *et al.* 2015), three studies presented their results after cementation (BINDL and MORMANN 2005; KORKUT *et al.* 2011; YUKSEL and ZAIMOGLU 2011) and three studies reported values of MG before and after cementation (BESCHIT and STRUB 1999; QUINTAS *et al.* 2004; DEMIR *et al.* 2014).

Four studies used direct examination with optical microscope (RINKE *et al.* 1995; SULAIMAN *et al.* 1997; YEO *et al.* 2003; QUINTAS *et al.* 2004), four studies used scanning with micro-XCT technology (PELEKANOS *et al.* 2009; MOUSLY *et al.* 2014; DEMIR *et al.* 2014; PIMENTA *et al.* 2015), two studies used cross-sectioning of the copings/crowns prior to the use of optical microscopes (KORKUT *et al.* 2011; YUKSEL and ZAIMOGLU 2011), two studies used scanning with stereomicroscope and photographs (ASAPVAPANUMAS *et al.* 2013; NG *et al.* 2014), one study used direct examination with SEM (BINDL and MORMANN 2005), one study used an epoxy replica of the marginal area and then measured the values using an optical microscope (BESCHIT and STRUB 1999) and finally, one study did not state the type of method used to make the measurement (ANADIOTI *et al.* 2013).

							Marginal	
		Sample		Finish		Examination	Gap	
Article	System Manufacturer	Size	State	Line	Cemented	Method	(µm)	Conclusion
Rinke <i>et al</i> .	In-Ceram Alumina	10 Pm	Crown	S	No	In vitro	45	No significant
1995	(split-casting)	10 Inc				Direct examination with optical	33.5	difference
	Celay	10 Pm				microscope	45	
	(block In-Ceram Alumina)	10 Inc					38	
Sulaiman <i>et al.</i> 1997	Procera AllCeram	30	NA	NA	No	In vitro Direct examination	82	Significant difference
	IPS Empress I	30				with optical	62	between the 3
	(veneering technique)					microscope		methods
	In-Ceram Alumina (slip-casting)	30					160.66	

Beschnidt and	In-Ceram Alumina	10	Crown	S	No	In vitro	60	Significant
Strub	(split-casting)				Yes		82	difference
1999						Direct examination		between
	Celay	10			No	on an epoxy resin	99	CAD/CAM
	(block Vita Celay)				Yes	replica of the	117	and
						marginal area with		conventional
	Celay	10			No	optical microscope	78	method
	(block In-Ceram Alumina)				Yes		91	
Yeo et al.	In-Ceram Alumina	30	Crown	S	No	In vitro	112	In-Ceram has
2003	(slip-casting)							worse MG
						Direct examination		than the Celay
	Celay	29				with optical	83	
	(block In-Ceram Alumina)					microscope		
Quintas <i>et al</i> .	IPS Empress II	60	Coping	NA	No	In vitro	68	Procera
2004	-				Yes		110	copings
	In-Ceram Alumina	60			No	Direct examination	57	presented
	(slip-casting)				Yes	with optical	117	better MG
						microscope		
	Procera AllCeram	60			No	-	25	
					Yes		44	

Bindl and Mormann 2005	In-Ceram Zirconia (Slip-casting)	12	Coping	С	Yes	In vitro Direct examination	25	Procera copings presented
	IPS Empress II	12				with SEM	44	better MG
	Cerec InLab	12					43	All crowns
	(block In-Ceram Zirconia)							were within clinical
	DCS Précident	12					33	acceptable values
	Decim	12					23	
	Procera AllCeram	12					17	
Pelekanos <i>et al.</i> 2009	WolCeram (in-Ceram Alumina)	4	Coping	С	No	In vitro	34.86	Conventional methods
	In-Ceram Alumina (slip-casting)	4				Micro-XCT	21.08	presented the best results
	Cerec inLab	4					55.09	
	(block In-Ceram Alumina) Celay							Cerec inLab presented
	(block In-Ceram Alumina)	4					139.27	clinically acceptable values

Korkut <i>et al</i> .	Cercon	10	Coping	С	Yes	In vitro	43.02	Best MG in
2011	IPS Empress II	10				and optical microscope	47.51	group
	Procera Zirconia	10				examination	50.29	Differences were statistically significant
Yuksel and Zaimoglu	Lava	12	Coping	С	Yes	In vitro Cross-sectioning	82.7	Better MG results on the
2011	IPS e.max Press	12				and optical microscope examination	92.6	CAD/CAM group
Asavapanumas <i>et al.</i> 2013	IPS e.max Press Cercon	12	Coping	S	No	In vitro Stereomicroscope and photographed	132.61 128.35	The LAVA group demonstrated the worst MG
	Lava						162.23	The Cercon group presented the best results for MG

Anadioti <i>et al.</i> 2013	IPS e.max Press	30	Crown	NA	No	In vitro	75	There was no statistical
	LAVA COS (block IPS						74	between the
	e.max CAD)							two groups
								two groups
Ng et al.	IPS e.max Press	15	Crown	С	No	In vitro	74	CAD/CAM
2014						Stereomicroscope		group showed
	DMG (block IPS e.max	15				and photographed	48	better marginal
	CAD)							fit than the
	,							mathad
								method
Mously et al.	IPS e.max Press	30	Crown	С	No	In vitro	30.8	The heat-press
2014						Scanning by micro-		group showed
	E4D (block IPS e.max Cad					XCT	49.35	the best
								crown
								adaptation
								1
Demir <i>et al</i> .	Cerec InLab (block	10	Crown	S	Yes	In Vitro	60	Cerec inLab
2014	Vitabloc Mark II)	10			No		20	with Vitablocs
						Micro-XCT		Mark II
	Cerec InLab (block In-	10			Yes	examination	180	showed the
	Ceram 2000 AL)	10			No		80	best marginal
								adaptation
	IPS e max Press	10			Ves		160	and much
	11 5 C.1110A 1 1055	10			No		70	
		10			INO		/0	

Table 2. Summary of in	cluded studies
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7:rkonZohn							
ZIIKOIIZaIIII	5	Coping	С	No	In vitro	35.5	CAD/CAM
					Scanning by micro-		showed the
IPS e.max Press	5				XCT	76.19	best results for
							MG
							All the groups
							were within
							clinically
							acceptable
							values
	IPS e.max Press	IPS e.max Press 5	IPS e.max Press 5 XCT	IPS e.max Press 5 XCT 76.19			

Chapter 4: Discussion and Conclusions

Crown marginal fit is critical for success of the restoration; crowns with poor fit (marginal gap) are prone to failure due to micro-leakage, cement dissolution, and dental caries. In this paper, the fit of crowns was assessed based on the vertical gap measurement which was selected as the most critical factor of marginal gap while being the least susceptible to manipulation post-fabrication, as indicated by Holmes *et al.* in 1989. Horizontal discrepancies, such as crown overhangs, can be adjusted to some degree intra-orally, however, vertical MG can only be closed with luting cement, which is prone to dissolution (NG *et al.* 2014). For this reason, the vertical MG has the most clinical relevance and should be regarded as the most critical in crown margin evaluation.

Chairside CAD/CAM restorations are currently being used by a large number of dentists all-over the world; however, the accuracy of these systems is still questionable. The coping, and sometimes even the crown, could be completed without the use of a die through intraoral impression. However, the accuracy of the data acquisition varies according to the system's various optical impression technologies and manufacturer's (CONTREPOIS *et al.* 2013). Software technology and milling accuracy also suffers from differences between systems. In addition even for the same system, there can be substantial differences and variations among the measured values which can be explained by different experimental protocols used in each study as can be showed in this results.

The use of the conventional method of crown fabrication has been used for decades with proven long-term results for both longevity and survival. Careful selection of materials and meticulous fabrication procedures are necessary to compensate for expansions and contraction of the different materials involved in creating an accurately fitting crown (NG *et al.* 2014). However, the impossibility of controlling all the variables, combined with the propensity for human error, can result in poor marginal adaptation or even misfit. The use of a digital method seems to decrease the margin of error. Human intervention in the manufacturing of the crown could play a role according to the skill of the dental laboratory technician and the relative importance of his contribution (PELEKANOS *et al.* 2009).

The number of steps involved in the process is another important element because the probability of error increased with each additional step required For example, non-CAD/CAM systems required the use of a die spacer applied by a technician, and the traditional In-Ceram slip-casting system was described as singularly technique sensitive. (BINDL and MORMANN 2005).

Different measurement methods were used among the various studies, and this could have impacted results significantly. The first and most widely used method involved direct microscopic examination of the marginal area. Unfortunately, this method has two great disadvantages. First, identifying reference points to measure may prove difficult. Second, it may lead to projection errors (GROTEN et al. 2000). In the second method, cemented specimens were cross-sectioned, and the marginal area was then examined under a microscope. However, only a limited number of sections could be cut on any one specimen. These two techniques were also sometimes used to measure an epoxy resin replica of the marginal area instead of the area itself. This technique does not provide accurate results (CONTREPOIS et al. 2013). A third method involved creating a light bodied silicone replica of the gap between the crown and the tooth. This replica was then sectioned, and the zone that corresponded to the marginal area was observed by microscopy. This provided only a limited number of marginal gap measurements (BESCHIT and STRUB 1999). The last technique used was x-ray microtomography. This innovative and nondestructive technique, which delivers 2-dimensional and 3-dimensional imaging of the space between the reconstitution and the die, and it can provide very close sections of the marginal area, which allows for a great number of measurement sites and for easy recognition of the critical distances. (PELEKANOS et al. 2009). This method has several advantages over other technologies including the 3-dimensional evaluation of the marginal and internal gaps. Furthermore, it is easy to perform, nondestructive, and more time efficient and accurate than other methods. The main disadvantages are radiation artifacts, which are caused by the differences in the coefficient of radiation absorption among the different materials and the difficulty in using luting agents because they have some radiopacity, which might affect the evaluation of the marginal gap. (MOUSLY et al. 2014).

Another important factor is that, a better approximation of clinical conditions may be reached by conducting measurements upon completion of the crown (CONTREPOIS *et al.* 2013). In addition, measuring fit at the crown stage is necessary to compare single-layer crowns and multi-layer crowns.

Some studies also measured the marginal gap before and after cementation of the crown or coping. Measurements made solely after cementation do not allow for the determination of the relative impact on the marginal fit of cementation and of a system's intrinsic precision (GROTEN *et al.* 2000) It is also more convenient to conduct measurements without cementing the crown as most studies did. Further evaluation of the capacity of all-ceramic conventional methods and CAD/CAM systems marginal adaptation after cementation, should be studied with different objectives and not mixed with non-cementing studies.

The type of finish line used in the studies varied from shoulder or chamfer. This variable also could have been responsible for the different results obtained between the conventional and digital methods or even inside the same system. Finish lines made in in vitro studies should be prepared in accordance to the most realistic clinical conditions. So with this in mind, the use of models that bear no relation to an actual tooth anatomy should be discontinued. Furthermore, finish lines that present some degree of curvature should be preferred since they can better simulate the presence of a gingival margin (CONTREPOIS *et al.* 2013).

Analysis of the results of this study suggests that more studies support the idea that digitally made crowns/copings can have better marginal adaptation values. However, most results seem to be well within clinical acceptable values ($\leq 120 \mu m$), which means digital or conventional made crowns are still two well supported options for fixed rehabilitation. Better protocols should be implemented to study the adaptation or CAD/CAM ceramic crowns versus conventional ceramic crowns. Although there are many studies made all over the years regarding marginal discrepancy on all-ceramic crowns, little has been done to clearly compare the digital method and the conventional method. This fact could be appointed to the fact that CAD/CAM is a relatively new technology that is slowly making its way to medical office because of its high costs, making the conventional method still a very low cost/benefit method, still preferred my most dentists in the world.

Conclusions:

Based on the results obtained, the digital method seems to be a legitimate alternative to the traditional methods. Analysis of the results of this study suggested that the digital method exceeds the standards of clinical acceptability and can sometimes surpass the vertical marginal fit of conventionally fabricated crowns.

Further studies are encouraged using standardized protocols as well as systems and techniques, in order to better evaluate the capabilities of this digital systems.

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