

Do conventional glass ionomer cements release more fluoride than resin-modified glass ionomer cements?

Maria Fernanda Costa Cabral¹, Roberto Luiz de Menezes Martinho², Manoel Valcácio Guedes-Neto¹, Maria Augusta Bessa Rebelo², Danielson Guedes Pontes^{2,3}, Flávia Cohen-Carneiro^{2*}

¹School of Dentistry, Federal University of Amazonas, Manaus, Brazil

²Postgraduate Program in Dentistry, School of Dentistry, Federal University of Amazonas, Manaus, Brazil

³School of Health Sciences, State University of Amazonas, Manaus, Brazil

Objectives: The aim of this study was to evaluate the fluoride release of conventional glass ionomer cements (GICs) and resin-modified GICs. **Materials and Methods:** The cements were grouped as follows: G1 (Vidrion R, SS White), G2 (Vitro Fil, DFL), G3 (Vitro Molar, DFL), G4 (Bioglass R, Biodinâmica), and G5 (Ketac Fil, 3M ESPE), as conventional GICs, and G6 (Vitremar, 3M ESPE), G7 (Vitro Fil LC, DFL), and G8 (Resiglass, Biodinâmica) as resin-modified GICs. Six specimens (8.60 mm in diameter; 1.65 mm in thickness) of each material were prepared using a stainless steel mold. The specimens were immersed in a demineralizing solution (pH 4.3) for 6 hr and a remineralizing solution (pH 7.0) for 18 hr a day. The fluoride ions were measured for 15 days. Analysis of variance (ANOVA) and Tukey's test with 5% significance were applied. **Results:** The highest amounts of fluoride release were found during the first 24 hr for all cements, decreasing abruptly on day 2, and reaching gradually decreasing levels on day 7. Based on these results, the decreasing scale of fluoride release was as follows: G2 > G3 > G8 = G4 = G7 > G6 = G1 > G5 ($p < 0.05$). **Conclusions:** There were wide variations among the materials in terms of the cumulative amount of fluoride ion released, and the amount of fluoride release could not be attributed to the category of cement, that is, conventional GICs or resin-modified GICs. (*Restor Dent Endod* 2015;40(3):209-215)

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¹Cabral MFC; Guedes-Neto MV, School of Dentistry, Federal University of Amazonas, Manaus, Brazil

²Martinho RLM; Rebelo MAB; Pontes DG; Cohen-Carneiro F, Postgraduate Program in Dentistry, School of Dentistry, Federal University of Amazonas, Manaus, Brazil

³Pontes DG, School of Health Sciences, State University of Amazonas, Manaus, Brazil

***Correspondence to**

Flávia Cohen-Carneiro, DDS, PhD. Adjunct Professor, Postgraduate Program in Dentistry, School of Dentistry, Federal University of Amazonas, Rua Rio Mar, n. 1203/901, Nossa Senhora das Graças, Manaus, AM, Brazil 69053-120
 TEL, +55-92-98855-1101; FAX, +55-92-3305-4905; E-mail, flaviacohen Carneiro@gmail.com

Introduction

Dental caries is one of the most prevalent chronic diseases affecting humanity. Its progression or control depends on the balance between pathological and protective factors, and the best strategy for caries management is focused on methods of improving the remineralizing process.^{1,2} When a restorative intervention is necessary, the use of materials with minimal aggression to the tooth structure and cariostatic, adhesive, and biocompatible properties must be prioritized.³ Since their introduction by Wilson and Kent and clinical development by McLean and Wilson, glass ionomer cements (GICs) have been largely used in restorative dentistry because of their ability to reduce the incidence of caries affecting unrestored tooth surfaces.⁴⁻¹¹ One of the main characteristics of GICs is their continuous fluoride release, allowing reduction of the mineral dental structure solubility and inhibition of microbial metabolism and favoring re-establishment of the balance of the oral environment.^{5,12-15}

Fluoride ions are essential elements in the prevention and treatment of dental caries.^{3,15-18} Thus, many new studies are interested in finding ways to maintain fluoride

ions in the oral environment, particularly at the interface of the tooth with a bacterial biofilm.^{15,19} The measurement of fluoride ions released within a determined period of time is one of the applicable methodologies to evaluate the cariostatic properties of GICs for *in vitro* studies. This evaluation, associated with laboratory tests such as adhesive and physical resistance and manipulation characteristics, guides the choice of materials for clinical practice.²⁰

Laboratory tests demonstrated that fluoride release varies in accordance with the cement category - conventional or resin modified - and a variation among cements in the same category can also be seen. This affirmation can be inferred from the different results obtained by several researchers for different tested materials.^{12,13,17,21-27} However, there is no consensus on the principal factor responsible for the variation of fluoride release from ionomer cements, that is, whether the difference is due to the cement category or the specific composition of some materials, independent of their category. Therefore, the present study

aims to compare the fluoride release of five new and three established brands of conventional and resin-modified GICs. The null hypothesis of the study is that there is no difference between conventional and resin-modified GICs with respect to the fluoride release.

Material and Methods

The fluoride release from the GICs tested was analyzed in an experimentally designed *in vitro* study. The materials used are listed in Table 1.

Immersion media

Three different solutions were used during the fluoride release test. 1) Total ionic strength adjustment buffer II (TISAB II), a buffer solution containing deionized water, acetic acid, sodium chloride, and cyclohexane-diamino-tetra-acid (CDTA), which was used to provide constant background ion strength, decomplex F to make

Table 1. Materials used in the study

Group	Material	Type	Composition	Manufacturer
G1	Vidrion R	GIC	Powder: sodium, calcium, aluminum fluorosilicate, barium sulfate, polyacrylic acid, and pigments Liquid: tartaric acid, distilled water	S.S. WHITE Artigos Dentários Ltda, Rio de Janeiro, Brazil
G2	Vitro Fil	GIC	Powder: strontium, aluminum silicate, dehydrated polyacrylic acid, and iron oxide Liquid: polyacrylic acid, tartaric acid, and distilled water	DFL Indústria e Comércio S.A., Rio de Janeiro, Brazil
G3	Vitro Molar	GIC	Powder: barium, aluminum silicate, dehydrated polyacrylic acid, and iron oxide Liquid: polyacrylic acid, tartaric acid, and distilled water	DFL Indústria e Comércio S.A., Rio de Janeiro, Brazil
G4	Bioglass R	GIC	Powder: calcium, barium, aluminum fluorosilicate, polyacrylic acid, and inorganic fillers Liquid: polyacrylic acid, tartaric acid, and deionized water	BIODINÂMICA Química e Farmacêutica Ltda, Paraná, Brazil
G5	Ketac Fil	GIC	Powder: lanthanum, aluminum, strontium fluorosilicate glass, and pigments Liquid: tartaric acid and water	3M ESPE Dental Products, Minnesota, USA
G6	Vitremer	RMGIC	Powder/liquid: methacrylate polyacids, water, aluminum fluorosilicate glass, methacrylate monomers, and initiators	3M ESPE Dental Products, Minnesota, USA
G7	Vitro Fil LC	RMGIC	Powder: strontium, aluminum silicate, excipients, activators, and iron oxide Liquid: 2-hydroxyethyl methacrylate polyacids, stabilizer, catalyzer, and ethyl alcohol	DFL Indústria e Comércio S.A., Rio de Janeiro, Brazil
G8	Resiglass R Restore	RMGIC	Powder: calcium, barium, aluminum fluorosilicate, polyacrylic acid, and inorganic fillers Liquid: dimethacrylate groups, deionized water, and catalyst	BIODINÂMICA Química e Farmacêutica Ltda, Paraná, Brazil

GIC, glass ionomer cement; RMGIC, resin-modified glass ionomer cement.

it available for determination, and adjust the solution pH. 2) Demineralization solution containing deionized water, calcium chloride dehydrate, sodium phosphate monobasic, acetate buffer, sodium hydroxide, and thymol and having a pH of 4.3, in which the specimens were immersed for 6 hours a day. 3) Remineralization solution containing deionized water, calcium chloride dehydrate, sodium phosphate monobasic, potassium chloride, tris(hydroxymethyl)aminomethane buffer (TRIS buffer), sodium hydroxide, and thymol and having a pH of 7.0, in which the specimens were immersed for 18 hours a day. Solutions 2 and 3 compose the pH cycling system used in this study. During the immersion period, the specimens were agitated using a shaker equipment at room temperature, and the storage solutions were changed daily as described above (after 6 hours in the demineralization solution, the specimens were immersed for 18 hours in the remineralizing solution).

Preparation of the specimens

Six disc specimens (8.60 mm in diameter; 1.65 mm in thickness) of each material were prepared using a standard stainless steel mold, two glass plates, and a transparent matrix in between. The tested materials were mixed in accordance with the manufacturer's recommendations at room temperature ($25 \pm 1^\circ\text{C}$). A piece of nylon thread was incorporated into the cement during setting to suspend the samples in the test medium. After 7 minutes, they were removed from the mold and placed in a humidifier recipient at 37°C , 100% humidity for 24 hours to simulate the oral environment.

Fluoride release measurement

The fluoride ion concentration was measured by the potentiometric method using a fluoride-specific electrode (Orion model 96-09, Orion Research, Cambridge, MA, USA) coupled with an ion analyzer (Orion SA-720, Orion Research) according to Carvalho and Cury (1999).²³ The instrument was calibrated with a series of standard fluoride solutions (1.25 to 15.00 and 5.00 to 25.00 $\mu\text{gF}/\text{mL}$ prepared in TISAB) according to the fluoride concentration of the samples. The specimens of each material were placed in plastic test tubes containing 2 mL of the remineralization or demineralization solution according to the pH cycling system, and after the immersion period, the same volume of TISAB was mixed for the fluoride analysis.

The fluoride release was measured for 15 days. To calculate the total daily fluoride release, the fluoride ion concentration found in the remineralization and demineralization solutions were added. The fluoride concentration was given in millivolts (mV) by the electrode and was converted into the quantity of fluoride per material area ($\mu\text{gF}/\text{cm}^2$). Analysis of variance (ANOVA) and Tukey's test were used for the statistical analysis ($\alpha = 0.05$).

Results

The daily mean fluoride release from GICs in the demineralization-remineralization (de-re) solutions over 15 days is displayed in Figure 1. Table 2 shows the cumulative fluoride release of each material over 15 days. The daily mean fluoride release during the test period for each material is presented in Table 3. The highest amount of

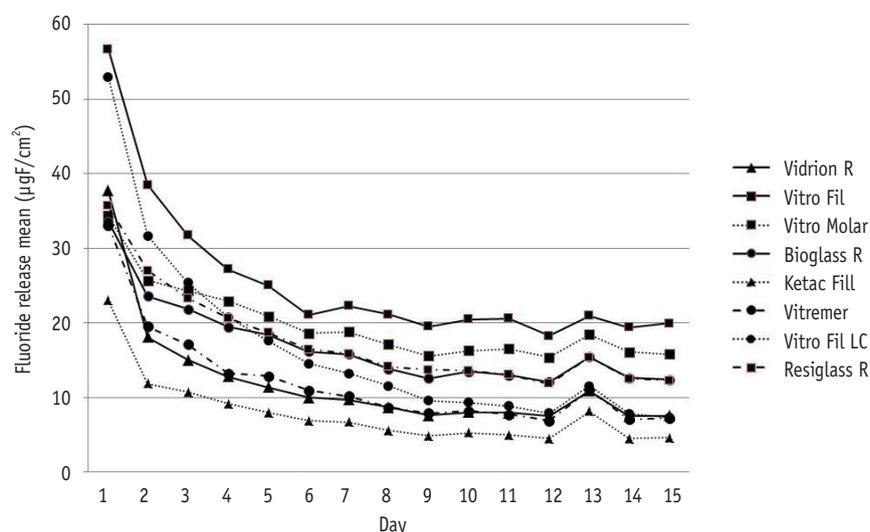


Figure 1. Daily mean fluoride release of conventional and resin-modified glass ionomer cements (GICs) in demineralization-remineralization solutions over 15 days. Vidrion R, Group 1; Vitro Fil, Group 2; Vitro Molar, Group 3; Bioglass R, Group 4; Ketac Fil, Group 5; Vitremer, Group 6; Vitro Fil LC, Group 7; Resiglass, Group 8.

Table 2. Total amount of fluoride released by each tested group ($\mu\text{gF}/\text{cm}^2$) during the 15 day study period

Group/cement	Fluoride released				Cumulative fluoride
	Day 1	Day 2	Day 7	Day 15	
G1: Vidrion R	37.75 \pm 1.28	18.08 \pm 1.25	9.69 \pm 0.38	7.54 \pm 0.36	180.45 \pm 7.76
G2: Vitro Fil	56.72 \pm 0.73	38.49 \pm 1.12	22.28 \pm 0.49	19.98 \pm 1.57	382.77 \pm 10.21
G3: Vitro Molar	34.39 \pm 2.13	25.69 \pm 2.16	18.81 \pm 1.07	15.78 \pm 0.84	296.97 \pm 5.22
G4: Bioglass R	33.58 \pm 3.52	23.58 \pm 4.39	15.81 \pm 3.24	12.38 \pm 2.07	254.26 \pm 5.84
G5: Ketac Fil	22.98 \pm 1.69	11.84 \pm 1.21	6.68 \pm 0.86	4.59 \pm 0.59	118.56 \pm 4.78
G6: Vitremer	33.09 \pm 4.35	19.53 \pm 5.37	10.14 \pm 2.29	7.23 \pm 1.30	181.73 \pm 6.91
G7: Vitro Fil LC	52.98 \pm 3.80	31.69 \pm 6.32	13.23 \pm 3.08	7.29 \pm 1.78	250.22 \pm 12.29
G8: Resiglass	35.71 \pm 0.81	26.99 \pm 2.53	15.84 \pm 1.72	12.91 \pm 1.47	264.15 \pm 6.65

Table 3. Daily mean of fluoride released ($\mu\text{gF}/\text{cm}^2$) during the study period for each tested material and the corresponding significance levels

Group	Material	Daily fluoride release*
G2	Vitro Fil	25.52 \pm 10.21 ^a
G3	Vitro Molar	19.80 \pm 5.22 ^b
G8	Resiglass	17.61 \pm 6.65 ^c
G4	Bioglass R	16.95 \pm 5.84 ^c
G7	Vitro Fil Lc	16.68 \pm 12.29 ^c
G6	Vitremer	12.12 \pm 6.91 ^d
G1	Vidrion R	12.03 \pm 7.76 ^d
G5	Ketac Fil	7.90 \pm 4.78 ^e

*Different letters represent statistically significant differences ($p < 0.05$) according to analysis of variance (ANOVA) and Tukey's test.

fluoride was released by Vitro Fil (DFL), a conventional GIC, differing statistically from all other tested materials ($p < 0.05$). There were no statistically significant differences among Resiglass (Biodinâmica), Bioglass R (Biodinâmica), and Vitro Fil LC (DFL), or between Vitremer (3M ESPE) and Vidrion R (SS WHITE) ($p > 0.05$). The least amount of fluoride was released by Ketac Fil (3M ESPE), differing statistically from all other tested cements ($p < 0.05$).

A pattern of fluoride release was observed in all tested materials. The highest mean values of fluoride release were found in the first 24 hours, decreasing abruptly on day 2 and reaching gradually decreasing levels on day 7.

Discussion

Several studies tested the *in vitro* fluoride release from conventional and resin-modified GICs.^{17,25,26,28-31} However, the use of different methodologies and materials leads

to a considerable variation of research results. Some of the factors related to *in vitro* fluoride release from restorative materials are the fluoride concentration in the set materials, size and composition of the inorganic filler, powder-liquid ratio of two-phase systems, mixing procedure, curing time, inner-material porosity, surface treatment and amount of exposed area of the specimen, and the type, temperature, and pH of the immersion media used.^{6,12,15} Depending on the powder composition, the material is capable of releasing different amounts of fluoride. Hattab and Amin found a strong positive correlation between the released fluoride and the fluoride concentration in the set materials.¹² Further, it is common knowledge that the inorganic filler composition of ionomer cements, compomers, and composite resins interferes with fluoride release.^{25,26} Fluoroaluminosilicate glass is the major component filler of GICs. As it is more soluble than the barium and strontium present in most compomers and

composites, it is able to release more fluoride.³² Particle size also has a significant influence on fluoride release. Reducing the filler particle size can increase fluoride release because smaller particles have larger surface areas.^{1,26}

Reduction of the inorganic filler size in compomers and GICs is a way for manufacturers to increase fluoride release.^{5,26} Another item related to the release and recharge on fluoride ions is the material's porosity. Xu and Burgess found that higher porosity allows deeper diffusion of the recharge agent into the sample and results in more fluoride storage and release.²⁶ They also reported that materials with less resin content, such as glass ionomers and resin-modified glass ionomers, have higher porosity and exhibit higher fluoride recharge capabilities than compomers or composites.

Regarding immersion media, the most cited ones in fluoride release methodologies are deionized water, saliva, artificial saliva, saline solution, and pH cycling system (de-re solutions).^{6,8,12,23,27,33} An analysis of these findings reveals that pH and ionic saturation from an immersion medium can influence the quantity of fluoride released. Indeed, for conventional and resin-modified GICs, the highest fluoride release is found in acidic and de-re solutions.^{15,30}

The increasing amount of fluoride in acidic media can be explained by the fact that a decrease in pH increases the dissolution of the material, leading to a higher fluoride level in the acidic immersion.¹⁵ In the case of neutral solutions, Hattab and Amin observed that GICs released significantly less fluoride in artificial saliva than in deionized water.¹² Besides the pH, the type of acid found in immersion solutions has an important effect on cement degradation.³³ McKenzie *et al.* observed that solutions containing carboxylic acids, capable of chelating calcium ions present in the cement and forming complexes of reasonable solubility in water (such as in orange and apple juices), have shown a greater degradation potential over ionomer cements than solutions containing phosphoric acid (such as Coca Cola), which although capable of chelating with calcium, form essentially insoluble complexes.³³ It is also important to explain that the degradation of ionomer cements is directly related to fluoride release.³⁰

Carvalho and Cury, evaluating the fluoride released from different dental materials in deionized water, artificial saliva, and de-re solutions, concluded that de-re solutions were a better immersion medium for the specimens because of the better cariogenic challenge simulation, which could not have been conducted in deionized water or artificial saliva, whereas other elements such as pH and proteins potentially influence dissolution and erosion, as reported by Shiozawa.^{6,23} In the present study, the de-re solutions were chosen to represent oral conditions.

The use of a surface coating agent (SCA) over GICs can be controversial. Although manufacturers recommended protecting the ionomer cement against degradation in

its first setting phase, the use of SCA in *in vitro* studies interferes with microleakage and significantly reduces ionomer cement fluoride release.¹² Thus, covering freshly set glass ionomers with varnish or resin significantly reduces the rate and amount of fluoride released in both deionized water and artificial saliva.¹² Further, unlike the *in vivo* physiological abrasion caused by SCA due to chewing, occlusal grinding, and tooth brushing, there is no abrasion during *in vitro* tests and a thin layer of fluid resin remains over the material.¹² In such cases, an SCA was not used as it could not be removed because the finishing and polishing of the specimen would decrease its surface area, modifying the amount of fluoride release.

Williams *et al.*, while analyzing the influence of the surface area and the volume of specimens during *in vitro* fluoride release tests, concluded that the quantity of released ions is directly related to the surface area of the specimen and that the volume does not have any influence on the quantity of ions released.²⁴ Thus, finishing and polishing could change the specimen dimensions, resulting in possible alterations to the surface area. Otherwise, the specimens' superficial smoothness was obtained by a transparent matrix, which is a good substitute for finishing and polishing.

In the present study, the highest or lowest amounts of fluoride release of the tested GICs could not be justified by the material category, that is, conventional or resin-modified GICs. While the cement that released the highest amount of fluoride was a conventional GIC (Vitro Fil, DFL), the one that released the least amount belonged to the conventional category as well (Ketac Fil, 3M ESPE). The different methodologies applied to the *in vitro* fluoride release research and the absence of published research about the five new materials tested in the present study make it even more difficult to compare these results to other *in vitro* fluoride release results. Carvalho and Cury, using a similar methodology, found the daily means of fluoride release for Vitremer ($12.27 \pm 1.16 \mu\text{gF}/\text{cm}^2$) and for Chelon Fil ($8.35 \pm 0.51 \mu\text{gF}/\text{cm}^2$), agreeing with the results found with Vitremer and Ketac Fil in the present study.²³ Other researchers have demonstrated the greatest fluoride release from Vitremer compared with Ketac Fil, Ketac Molar, and Fuji IX, all conventional GICs with regular viscosity (Ketac Fil) or high viscosity (Ketac Molar and Fuji IX).^{25,27,28,34} The explanation for the higher fluoride release by resin-modified GICs is as follows: Acid-based reactions are slowed down by the resin component, which makes the ionic matrix less mature and capable of releasing more fluoride, if compared with a conventional material of the same age; larger pore size and porosity of resin-modified GICs; low solubility and high powder-liquid proportion of GICs with high viscosity.^{25,28,34}

However, when a great variety of conventional and resin-modified GICs were compared, the variation of the amount

of fluoride release could not be attributed to the type of material or GIC category; this is in agreement with the results of the present study.^{26,35} Xu and Burgess found a negative linear correlation between the compressive strength and fluoride release and that restorative materials with high fluoride release had lower mechanical properties.²⁶ This may be in accordance with other studies that attribute the rates of fluoride release to the solubility and erosion of GICs.^{3,30} Additionally, Xu and Burgess found a considerably large discrepancy in the amount of fluoride release within the resin-modified GIC category, from 162 $\mu\text{gF}/\text{cm}^2$ (Vitremar, 3M ESPE) to 375 $\mu\text{gF}/\text{cm}^2$ (Photac Fil, 3M ESPE), cumulated in 21 days.²⁶

The fluoride release of conventional and resin-modified GICs follows the same pattern. Most of the reviewed studies mentioned that the tested materials present a significantly higher release during the first 24 hours, declining from day 2 and tending to stabilize towards day 7.^{13,17,29} This may be caused by the initial superficial rinsing effect, while the constant fluoride release during the following days occurs because of the fluoride's ability to diffuse through cement pores and fractures.³² An initial high release from glass ionomers over the first 24 hours is likely due to the burst of fluoride released from the glass particles when reacting with the polyalkenoate acid during the setting reaction.¹⁵ In the present study, the higher means were observed on day 1, decaying from day 2, and then tenuously diminishing the fluoride release towards day 7.

Fluoride-releasing restorative materials, particularly GICs, present antibacterial properties and can affect *in vitro* conditions of the microbial metabolism during caries process simulation. However, prospective clinical studies did not verify whether the secondary caries incidence can be significantly reduced by these materials. The cariogenicity and frequency of the patient's diet, presence of saliva, and bacteria-producing acid challenges are important clinical variables to be considered.^{5,36} Dijkman *et al.* showed that a monthly cumulative fluoride release of 200 - 300 mg/cm^2 is sufficient to completely inhibit enamel demineralization *in vivo*.³⁷ In the present study, for a 15 day period, the materials that showed a fluoride release of 200 $\mu\text{g}/\text{cm}^2$ or more were Vitro Fil, Vitro Molar, and Bioglass R among the conventional GICs, and Vitro Fil LC and Resiglass among the resin-modified GICs.

Conclusions

This study showed a wide variation among the materials in terms of the cumulative amounts of fluoride ions released. This variation was not related to the type of material. Therefore, the amount of fluoride released could not be attributed to the category of GIC, that is, conventional and resin-modified GICs, and this confirms the null hypothesis of the present study.

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Orcid number:

Roberto Luiz de Menezes Martinho, 0000-0002-0854-5190

Danielson Guedes Pontes, 0000-0003-2417-0672

Flávia Cohen-Carneiro, 0000-0002-5497-6514

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References

1. Mungara J, Philip J, Joseph E, Rajendran S, Elangovan A, Selvaraju G. Comparative evaluation of fluoride release and recharge of pre-reacted glass ionomer composite and nano-ionomeric glass ionomer with daily fluoride exposure: an *in vitro* study. *J Indian Soc Pedod Prev Dent* 2013;31:234-239.
2. Upadhyay S, Rao A, Shenoy R. Comparison of the amount of fluoride release from nanofilled resin modified glass ionomer, conventional and resin modified glass ionomer cements. *J Dent (Tehran)* 2013;10:134-140.
3. Markovic DL, Petrovic BB, Peric TO. Fluoride content and recharge ability of five glass ionomer dental materials. *BMC Oral Health* 2008;8:21.
4. Wilson AD, Kent BE. A new translucent cement for dentistry. The glass ionomer cement. *Br Dent J* 1972;132:133-135.
5. Neelakantan P, John S, Anand S, Sureshbabu N, Subbarao C. Fluoride release from a new glass-ionomer cement. *Oper Dent* 2011;36:80-85.
6. Shiozawa M, Takahashi H, Iwasaki N. Fluoride release and mechanical properties after 1-year water storage of recent restorative glass ionomer cements. *Clin Oral Investig* 2014;18:1053-1060.
7. McLean JW, Wilson AD. The clinical development of glass-ionomer cements III. The erosion lesion. *Aust Dent J* 1977;22:190-195.
8. Levallois B, Fovet Y, Lapeyre L, Gal JY. *In vitro* fluoride release from restorative materials in water versus artificial saliva medium (SAGF). *Dent Mater* 1998;14:441-447.
9. Tiwari S, Nandlal B. Comparative evaluation of fluoride release from hydroxyapatite incorporated and conventional glass ionomer cement: an *in vitro* study. *J Indian Soc Pedod Prev Dent* 2012;30:284-287.
10. Qvist V, Manscher E, Teglers PT. Resin-modified and conventional glass ionomer restorations in primary teeth: 8-year results. *J Dent* 2004;32:285-294.
11. Qvist V, Poulsen A, Teglers PT, Mjör IA. Fluorides

- leaching from restorative materials and the effect on adjacent teeth. *Inter Dent J* 2010;60:156-160.
12. Hattab FN, Amin WM. Fluoride release from glass ionomer restorative materials and the effects of surface coating. *Biomaterials* 2001;22:1449-1458.
 13. Diaz-Arnold AM, Holmes DC, Wistrom DW, Swift EJ Jr. Short-term fluoride release/uptake of glass ionomer restoratives. *Dent Mater* 1995;11:96-101.
 14. Dionysopoulos D, Koliniotou-Koumpia E, Helvatzoglou-Antoniades M, Kotsanos N. Fluoride release and recharge abilities of contemporary fluoride-containing restorative materials and dental adhesives. *Dent Mater J* 2013;32:296-304.
 15. Wiegand A, Buchalla W, Attin T. Review on fluoride-releasing restorative materials--Fluoride release and uptake characteristics, antibacterial activity and influence on caries formation. *Dent Mater* 2007;23:343-362.
 16. Billington RW, Williams JA, Dorban A, Pearson GJ. Glass ionomer cement: evidence pointing to fluorine release in the form of monofluorophosphate in addition to fluoride ion. *Biomaterials* 2004;25:3399-3402.
 17. Can-Karabulut DC, Batmaz I, Solak H, Taştekin M. Linear regression modeling to compare fluoride release profiles of various restorative materials. *Dent Mater* 2007;23:1057-1065.
 18. Bahadure RN, Pandey RK, Kumar R, Gopal K, Singh RK. An estimation of fluoride release from various dental restorative materials at different pH: *in vitro* study. *J Indian Soc Pedod Prev Dent* 2012;30:122-126.
 19. Moreau JL, Xu HH. Fluoride releasing restorative materials: effects of pH on mechanical properties and ion release. *Dent Mater* 2010;26:e227-e235.
 20. Jeong YN, Yang SY, Park BJ, Park YJ, Hwang YC, Hwang IN, Oh WM. Physical and chemical properties of experimental mixture of mineral trioxide aggregate and glass ionomer cement. *J Korean Acad Conserv Dent* 2010;35:344-352.
 21. Seppä L, Forss H, Ogaard B. The effect of fluoride application on fluoride release and the antibacterial action of glass ionomers. *J Dent Res* 1993;72:1310-1314.
 22. Grobler SR, Rossouw RJ, Van Wyk Kotze TJ. A comparison of fluoride release from various dental materials. *J Dent* 1998;26:259-265.
 23. Carvalho AS, Cury JA. Fluoride release from some dental materials in different solutions. *Oper Dent* 1999;24:14-19.
 24. Williams JA, Billington RW, Pearson GJ. The influence of sample dimensions on fluoride ion release from a glass ionomer restorative cement. *Biomaterials* 1999;20:1327-1337.
 25. Dionysopoulos P, Kotsanos N, Pataridou A. Fluoride release and uptake by four new fluoride releasing materials. *J Oral Rehabil* 2003;30:866-872.
 26. Xu X, Burgess JO. Compressive strength, fluoride release and recharge of fluoride-releasing materials. *Biomaterials* 2003;24:2451-2461.
 27. Hayacibara MF, Ambrosano GM, Cury JA. Simultaneous release of fluoride and aluminum from dental materials in various immersion media. *Oper Dent* 2004;29:16-22.
 28. Hsu HM, Huang GF, Chang HH, Wang YL, Guo MK. A continuous flow system for assessing fluoride release/uptake of fluoride-containing restorative materials. *Dent Mater* 2004;20:740-749.
 29. Chan WD, Yang L, Wan W, Rizkalla AS. Fluoride release from dental cements and composites: a mechanistic study. *Dent Mater* 2006;22:366-373.
 30. Gandolfi MG, Chersoni S, Acquaviva GL, Piana G, Prati C, Mongiorgi R. Fluoride release and absorption at different pH from glass-ionomer cements. *Dent Mater* 2006;22:441-449.
 31. Selimović-Dragaš M, Hasić-Branković L, Korać F, Đapo N, Huseinbegović A, Kobašlija S, Lekić M, Hatibović-Kofman Š. *In vitro* fluoride release from a different kind of conventional and resin modified glass-ionomer cements. *Bosn J Basic Med Sci* 2013;13:197-202.
 32. Mousavinasab SM, Meyers I. Fluoride release by glass ionomer cements, compomer and giomer. *Dent Res J (Isfahan)* 2009;6:75-81.
 33. McKenzie MA, Linden RW, Nicholson JW. The physical properties of conventional and resin-modified glass-ionomer dental cements stored in saliva, proprietary acidic beverages, saline and water. *Biomaterials* 2003;24:4063-4069.
 34. Rothwell M, Anstice HM, Pearson GJ. The uptake and release of fluoride by ion-leaching cements after exposure to toothpaste. *J Dent* 1998;26:591-597.
 35. Gao W, Smales RJ. Fluoride release/uptake of conventional and resin-modified glass ionomers, and compomers. *J Dent* 2001;29:301-306.
 36. Rodrigues JA, Marchi GM, Serra MC, Hara AT. Visual evaluation of *in vitro* cariostatic effect of restorative materials associated with dentifrices. *Braz Dent J* 2005;16:112-118.
 37. Dijkman GE, de Vries J, Lodding A, Arends J. Long-term fluoride release of visible light-activated composites *in vitro*: a correlation with *in situ* demineralization data. *Caries Res* 1993;27:117-123.