# An Exploratory Study Comparing Salivary CAVI Levels in Children Undergoing Fixed and Interceptive Orthodontic Treatments Versus Caries-Matched Controls

# Abstract

Aim: The aim of this study is to determine the influence of salivary CAVI on the dental caries status of children receiving fixed and interceptive orthodontic treatment and compare them to children who were not receiving orthodontic treatment.

**Methods:** The study participants comprised 60 patients aged 9–14 years allocated to one of three groups Control patients (with no fixed or interceptive treatment carried out), patients with interceptive orthodontic appliance and patients with fixed orthodontic appliance. All the participants were examined for DMFT/dmft, OHI and PH. Saliva was collected by sterile pasture pipette or by passive drool method. CAVI was estimated by using a commercially available Elisa kit.

**Results:** CAVI levels in the fixed orthodontic group exhibit the highest levels of the enzyme whereas in the control groups had the lowest levels. When the CAVI levels among groups were subjected to the Scheffe's post hoc test it was observed that while a significant difference existed between the CAVI levels and the other two groups (p<0.05) no significant differences were observed between the control group and the interceptive orthodontic group.

**Conclusion:** Children undergoing fixed orthodontic treatment have significantly higher CAVI expression than those undergoing interceptive orthodontic treatment or controls.

There is no significant difference in the CAVI levels of children undergoing interceptive orthodontic treatment and controls who were matched for oral hygiene and DMFT.

Reem AlSakr<sup>1\*</sup>, Sharat Pani<sup>1</sup> and Deema AlShammery<sup>2</sup>

<sup>1</sup>Department of Preventive Dental Sciences, Riyadh Elm University, Riyadh, Saudi Arabia

<sup>2</sup>Department of Orthodontic, College of Dentistry, Riyadh Elm University, Riyadh, Saudi Arabia

\*Corresponding author: Reem Alsakr, Department of Preventive Dental Sciences, Postgraduate Student in Pediatric Dentistry, Riyadh Elm University, Riyadh, Saudi Arabia, Email: reemsakr0555@gmail.com

Received: Feb 4 2024, Accepted: Mar 10 2024; Published: Mar 15, 2024, DOI: 10.59462/jodt.1.1.102

**Copyright:** © 2024 Alsakr R, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited

Citation: AlSakr R, Pani S, AlShammery D (2024) An Exploratory Study Comparing Salivary CAVI Levels in Children Undergoing Fixed and Interceptive Orthodontic Treatments Versus Caries-Matched Controls. Journal of Oral Diseases and Treatment, 1(1):102.

**Keywords:** Carbonic Anhydrase 6; Interceptive Orthodontic Treatment; Fixed orthodontic

# Introduction

One of the most common dental disorders is malocclusion that may lead to increasing the risk of periodontal disease and dental caries orthodontic treatment olen resolve malocclusion, or at least prevent further progression [1]. However, it may a9ect oral hygiene by influencing several factors including the saliva properties and microbial count [2]. Uese changes in saliva include decrease in pH, flow rate and bu9ering capacity. Uis may contribute to demineralization of enamel and increase the susceptibility to dental caries [3].

Another important factor in the dental caries dynamic process is the bu9ering capacity of saliva, which also plays an important role in maintaining the oral tissues homeostasis. Among its multiple functions, the clearance promoted by the salivary flow and the pH stability at acceptable levels stand out, mainly due to carbonate and phosphate bu9ers [4]. Among the defense systems of saliva, salivary carbonic anhydrase isoenzyme VI (CAVI) is the only known secreted isoenzyme of the carbonic anhydrase family, which has been detected in the saliva secreted by the serous acinar cells of mammalian parotid and submandibular glands. It catalyzes the reversible reaction of carbon dioxide in a reaction of CO<sub>2</sub>+H<sub>2</sub>O H+HCO<sub>3</sub>. By catalyzing this reaction, CAVI is believed to provide a greater bu9ering capacity to saliva by penetrating dental biofilm and facilitating acid neutralization by salivary bicarbonate [5].

Given that orthodontic appliances create biofilm stagnation areas and complicate the oral hygiene, patients are susceptible to a higher biofilm accumulation, and mostly subjected to important biochemical and microbiological changes in saliva and biofilm. Uere is also debate in literature suggesting that the role played by fixed orthodontic appliances on the oral hygiene of children may di9er from the role played by interceptive or removable appliances [3].

Uus, deliberating the behavior of CAVI in saliva of di9erent types of orthodontic patient would be at great importance to investigate [6].

# Methodology

Ue proposal was registered with the research center of Riyadh Elm University (FPGRP/43735003/228) and ethical approval was obtained from the Institutional Review Board (IRB) of Riyadh Elm University (RC/IRB/2018/1053).

A total of 60 patients, age 9 to 14 years, mixed dentition period, good general health attending the orthodontic clinics of Riyadh Elm University hospitals, were participated in the study and allocated to one of three groups; Group A, Group B, Group C. Represent respectively. Control patients (with no fixed or interceptive treatment carried out), patients with interceptive orthodontic appliance and patients with fixed orthodontic appliance.

#### Examination of oral health

Subjects were examined for caries using WHO (DMFT) Index Decay, Missing, and Filled Tooth. Oral hygiene level was assessed using the Oral Hygiene Index (Simplified) OHI(S) using a mirror and a WHO CPTIN type E probe.

OHI(S) was calculated as the mean score for the examined surfaces and was used as a measure for the average level of plaque accumulation. Ue salivary pH was calculated using test strips.

#### Saliva collection

Saliva of about 1 ml was collected using a sterile Pasteur pipette and transferred to a 1.5 ml Eppendorf tube from the participants. Uese sterile plastic collection tubes were transferred to the deep freezer of a  $-600^{\circ}$ C until they can be analyzed.

## Assessment of CAVI activity

Ue assessment of CAVI activity was performed by the zymography method. It was performed on saliva since this isoenzyme can adhere to the acquired pellicle and promote the neutralization of excess acid by catalyzing the reaction of  $H^++HCO_3^- \leftrightarrow CO_2+H_2O$ , which constitutes the most important bu9er in the oral environment [7]. Salivary CAVI was measured using the commercially available CAVI ELISA kit.

Uis kit is based on sandwich enzyme-like immuno-sorbent assay technology. An antibody specific to CAVI is pre-coated onto a 96-well plate. Ue standards and samples are added to wells and incubated. Biotin conjugated anti-CAVI antibody. Next, Avidin conjugated to HRP is added to each microplate well and incubated. AIer TMB substrate solution is added only wells that contain CAVI, biotinconjugated antibody and enzyme-conjugated Avidin will produce a blue color product that changes into yellow aIer adding acidic stop solution. Ue intensity of the color yellow is proportional to the CAVI amount bound on the plate. AIer that, CAVI O.D. absorbance is measured spectrophotometrically at 450 nm in a microplate reader, and then the concentration of CAVI can be calculated.

#### Result

## Descriptive statistics

Ue sample comprised of a total of 60 subjects (20 Fixed orthodontic, 20 interceptive orthodontic and 20 control) aged between 9 and 14 years. Ue mean age of the sample was 10.38 years (SD  $\pm$  2.1 years). Uere were no significant di9erences between genders (Table 1), Patients receiving fixed orthodontic treatment were significantly older than those receiving interceptive orthodontic treatment.

		Gender				
		Male age		Fem		
		Mean	Standard deviation	Mean	Standard deviation	
group	Controlab	10.56	1.72	10.62	1.5	0.346
	Interceptive Orthodonticsa	8.5	1.88	9.64	2.01	0.564
	Fixed orthodonticsb	11.32	1.2	12.31	2.1	0.455

\*calculated using the independent t test

a,b : Differences in superscript indicate significant difference in age (calculated using One-Way ANOVA and Scheffe's Post hoc test).

Table 1: Age distribution of the sample.

		Mean	Std. deviatio n	Skewness		Kurtosis	
Group		Statisti c	Statistic	Statisti c	Std. error	Statisti c	Std. error
Control	CAVI	26.245	5.17631	0.716	0.512	-0.065	0.992
Interceptive orthodontics	CAVI	17.675	1.61404	0.689	0.512	1.977	0.992
Fixed orthodontics	CAVI	13.005	2.46907	0.042	0.512	-0.993	0.992

Table 2: Descriptive statistics of the CAVI profile of the sample.

Ue descriptive statistics of the sample revealed no significant skew suggesting that the sample was not normally distributed and that parametric statistics should be used. Ue sample also revealed no significant Kurtosis (<3) suggesting the use of two-tailed tests (Table 2).

### Oral hygiene of the di9erent groups

		Mean	Std. deviation	F*	Sig		
	Control	6.55	0.51042		0.657		
Saliva	Interceptive orthodontics	6.7	0.80131	0.427			
pH	Fixed orthodontics	6.75	0.7864				
	Control	2.54	0.63279		0.138		
	Interceptive orthodontics	2.325	0.43392	2.068			
оні	Fixed orthodontics	2.66	0.50513				
* differences tested using the One-Way ANOVA							

Table 3: Comparison mean pH and OHI among groups.

Ue control group had the lowest salivary pH. While the fixed orthodontics group had the worst oral hygiene (Table 3). However, the One-Way ANOVA showed that the di9erences were not statistically significant.

Dental caries experience of the population

Ue dental caries in primary teeth was not recorded as the protocol of the university called for the extraction of all carious primary teeth prior to orthodontic treatment. When the Decayed (D), Missing (M) and Filled (F) permanent teeth were compared it was observed that there were no significant di9erences in the overall DMFT of any of the three groups (Table 4).

	Mean	Std. deviation	F*	Sig			
Control	4.05	1.79991	0.673	0.123			
Interceptive orthodontics	4.45	1.87715					
Fixed orthodontics	4.05	2.18789					
*differences tested using the One Way ANOVA							

\*differences tested using the One-Way ANOVA

#### Table 4: Comparison of Mean DMFT across groups.

#### CAVI activity and orthodontic treatment

When the CAVI levels were compared in the saliva across groups it was observed that the control group had the lowest levels and the fixed orthodontics groups had the highest level. Ue di9erences among groups were statistically significant (Table 5).

	N	Mean	Std. deviation	F*	Sig		
Fixed orthodontics	20	19.245	3.17631	46.221	<0.001**		
Interceptive orthodontics	20	17.675	1.61404				
Control	20	13.005	2.46907				
* Calculated using the One-Way ANOVA							
** Differences significant at p<0.05							

Table 5: Comparison of CAVI levels among groups.

When the CAVI levels among groups were subjected to the Sche9e's post hoc test it was observed that while a significant di9erence existed between the CAVI levels and the other two groups (p<0.05) no significant di9erences were observed between the control group and the interceptive orthodontic group (Table 6).

### Means for groups in homogeneous subsets are displayed.

When a regression model was formulated with CAVI levels as the dependent variable it was shown that the presence of orthodontic treatment and caries levels had a significant association with CAVI

levels. Ue salivary pH and OHI scores had no significant association with the CAVI levels (Table 7).

CAVI						
Scheffea	Subset for alpha=0.05					
Group	N	1	2			
Fixed orthodontics	20	19.245				
Interceptive orthodontics	20	17.675				
Control	20		13.005			
Sig.		0.401	0.171			
Means for groups in homogeneous subsets are displayed.						
a. Uses Harmonic Mean Sample Size=20.000.						

Table 6: Post hoc comparisons among di9erent groups.

Model	Unstandardized coefficients		Stand coefficie	Sig.			
	В	Std. error	Beta	t			
(Constant)	35.267	5.392		6.541	0		
Orthodontic treatment	-6.815	0.557	-0.866	-12.244	.000*		
DMFT	-0.397	0.196	-0.152	-2.028	.047*		
Saliva pH	0.296	0.658	0.032	0.45	0.655		
OHI	-0.553	0.87	-0.046	-0.635	0.528		
*Association statistically significant							

Table 7: Regression model showing factors associated with CAVI levels. Dependent Variable: CAVI. Calculated using linear regression modelling.

# Discussion

Dental caries is one of the most serious challenges of orthodontic treatment [8,9]. Dental caries is an irreversible microbial disease of the calcified tissues of teeth characterized by demineralization of the inorganic portion and destruction of the organic substance of the tooth which oIen leads to cavitation [10]. Caries is a complex and dynamic process where a multitude of factors influence and initiate the progression of the disease. It is well known that orthodontic treatment has the potential to cause damage to the hard and soI tissues. Ue presence of archwires complicates cleaning and makes access to plaque retaining areas difficult, especially when multiple loops, auxiliary archwires and di9erent types of elastics are used [11]. Ue significantly higher expression of CAVI in the fixed orthodontic group confirms the hypothesis that the presence of arch wires is probably the most likely cause of dental caries in children undergoing orthodontic treatment.

Boersma et al. [3] showed positive correlation with caries prevalence was found for the bleeding and oral hygiene index scores of children undergoing orthodontic treatment (similar observations have been reported by various authors) [12,13]. Ue current study controlled for the overall oral hygiene among the di9erent groups studied. Ue presence of an increased inflammatory reaction in children undergoing orthodontic treatment has been previously documented [14]. Ue results of this study show that in fixed orthodontic treatment there is the greatest CAVI activity, a fact that is confirmatory of an increased response to inflammation in patients undergoing orthodontic treatment. Given that the di9erence was only significant in the fixed orthodontic group, it can be assumed that the continuous presence of wires in the patient's mouth presents a significantly higher inflammatory challenge, even when oral hygiene and dental caries are controlled for.

Salivary pH is an indicator of the bu9ering capacity of the saliva. Human saliva not only lubricates the oral tissues, making oral functions such as speaking, eating, and swallowing possible, but also protects teeth and oro-mucosal surfaces in di9erent ways [15].

Ue result of our study showed that there was no significant di9erence in the pH among the di9erent groups suggests the hypothesis that in cases of increased inflammatory challenges the salivary bu9er systems are able to maintain the oral homeostasis [16].

Dental enamel is the hardest tissue in the human body, and the main challenge to it comes from acidic conditions in the oral cavity, which can cause dissolving of the mineral, i.e. dental caries or erosion [17-21]. However as shown by several studies, the bu9ering systems of the oral cavity are able to overcome the challenges of mild inflammation [22,23]. Ue bu9ering capacity (is the power to resist changes of pH when acid or alkali are added) of a complex solution like saliva will vary at di9erent pH values because di9erent systems of bu9ers are e9ective over di9erent levels of the pH range [24].

Ue three major bu9er systems responsible for bu9er capacity of both unstimulated and stimulated saliva are the phosphate, the carbonic acid and the protien bu9ers [1,24]. Carbonic anhydrase is genetically expressed indicator for the functionality of the carbonic acid bu9er.

Carbonic anhydrases are evolutionally old enzymes [25]. Uey are expressed in most tissues of the human body, participating in pH regulation, carbon dioxide and bicarbonate [26,27]. transport, as well as in the maintenance of water and electrolyte balance [28,29]. To date, seven isoenzymes have been identified in mammals, and all of them are expressed in the alimentary tract [30-34].

Of the isoenzymes expressed in the salivary glands, Cytoplasmic CAII is a high-activity isoenzyme which has been proposed to catalyze the production of salivary bicarbonate [35]. However, more recent research has shown that CAVI which is secreted into the saliva by the serous acinar cells of the parotid and submandibular glands is perhaps a better expressor of salivary bu9ering capacity [33,36,37]. Ue results of our study showed that the CAVI levels of children undergoing fixed orthodontics were elevated significantly higher than those of the interceptive orthodontic and the control group. However there were no significant di9erences in pH. Uis, combined with the fact that the study design called for the matching of oral hygeine across groups lends credence to the argument that in cases where oral hygiene is maintained, the natural bu9ereing capacity of the saliva will help children compensate the inflammatory e9ects of arch wires [12,13].

Ue secretion of salivary CAVI is characterized [37,38] by a circadian periodicity, the concentration in saliva being very low during sleep and rapidly rising to the daytime levels aler awakening [39-58]. Uis could help explain why fixed orthodontic patients, who had the presence of arch wires continuously, had significantly higher [59-94].

CAVI activity than those who had interceptive orthodontic appliances that were removed at night. Uis also explains the finding that there was no significant di9erence in the CAVI levels between control patients and those with interceptive orthodontic appliances [95-124].

# Conclusion

Children undergoing fixed orthodontic treatment have significantly higher CAVI expression than those undergoing interceptive orthodontic treatment or controls.

Uere is no significant di9erence in the CAVI levels of children undergoing interceptive orthodontic treatment and controls who were matched for oral hygiene and DMFT.

#### References

- Bollen AM, Cunha-Cruz J, Bakko DW, Huang GJ, Hujoel PP et al (2008) Ue e9ects of orthodontic therapy on periodontal health: A systematic review of controlled evidence. J Am Dent Assoc 139: 413-422.
- Smiech-slomkowska G, Jablonska-zrobek J (2007) Ue e9ect of oral health education on dental plaque development and the level of cariesrelated Streptococcus mutans and Lactobacillus spp. Eur J Orthod 29: 157-160.
- Boersma JG, Van der Veen MH, Lagerweij MD, Bokhout B, Prahl-Andersen B et al. (2005) Caries prevalence measured with QLF aler treatment with fixed orthodontic appliances: Influencing factors. Caries Res 39: 41-47.
- Bardow A, Moe D, Nyvad B, Nauntole B (2000) Ue bu9er capacity and bu9er systems of human whole saliva measured without loss of CO<sub>2</sub>. Arch of Oral Biol 45: 1-12.
- 5. Kimoto M, Kishino M, Yura Y, Ogawa Y (2006) A role of salivary carbonic anhydrase VI in dental plaque. Arch Oral Biol 51: 117-122.
- Bardow A, Hofer E, Nyvad B, Ten Cate JM, Kirkeby S et al. (2005) NauntoIe B. E9ect of saliva composition on experimental root caries. Caries Res 39: 71-77.
- Leinonen J, Kivelä J, Parkkila S, Parkkila AK, Rajaniemi H (1999) Salivary carbonic anhydrase isoenzyme VI is located in the human enamel pellicle. Caries Res 33: 185-190.
- 8. Newbrun E (1989) Cariology. Quintessence Books.
- Dean JA, Avery DR, Mc Donald RE (2005) McDonald and Avery Dentistry for the Child and Adolescent, Elsevier Health Sciences.
- 10. Shafer WG, Hine MK, Levy BM. Dental caries. In: A Textbook of Oral Pathology.
- Årtun J, Brobakken BO (1986) Prevalence of carious white spots aler orthodontic treatment with multibonded appliances. Eur J Orthod 8: 229-234.
- Mattousch TJ, Van Der Veen MH, Zentner A (2007) Caries lesions aler orthodontic treatment followed by quantitative light-induced fluorescence: A 2-year follow-up. Eur J Orthod 29: 294-298.
- 13. Wisth PJ, Nord A (1977) Caries experience in orthodontically treated individuals. Angle Orthod 47: 59-64.
- 14. Mitchell L (1992) Decalcification during orthodontic treatment with fixed appliances: An overview. Br J Orthod 19: 199-205.
- Lenander-Lumikari M, Loimaranta V (2000) Saliva and dental caries. Adv Dent Res 14: 40-47.
- 16. Stookey GK (2008) Ue e9ect of saliva on dental caries. J Am Dent Assoc 139: 11S-17S.
- 17. Clarke JK (1924) On the bacterial factor in the aetiology of dental caries. Br J Exp Pathol 5: 141-147.
- Stephan RM (1944) Intra-oral hydrogen-ion concentrations associated with dental caries activity. J Dent Res 23: 257-266.
- Muntz JA (1943) Production of acids from glucose by dental plaque material. J Boil Chem 148: 225-236.

- 20. Geddes DA (1975) Acids produced by human dental plaque metabolism in situ. Caries Res 9: 98-109.
- 21. Ferguson DB (1981) Environment of the teeth. Karger 3: 162-173.
- Rudney JD (1995) Does variability in salivary protein concentrations influence oral microbial ecology and oral health? Crit Rev Oral Biol Med 6: 343-367.
- 23. Peeters FP, Devries MW, Vissink A (1998) Risks for oral health with the use of antidepressants. Gen Hosp Psychiatry 20: 150-154.
- 24. Dowd FJ (1999) Saliva and dental caries. Dent Clin North Am 43: 579-597.
- 25. Tashian RE, Hewett-emmet D, Venta PJ (1991) Diversity and evolution in the carbonic anhydrase gene family, Weinheim, VCH VerlagsgesellschaI 8: 151-161.
- 26. Al Nimri K, Richardson A (2000) Interceptive orthodontics in the real world of community dentistry. Int J Paediatr Dent 10: 99-108.
- 27. Cardoso AA, Lopes LM, Rodrigues LP, Teixeira JJ, Steiner-Oliveira C et al. (2017) Influence of salivary parameters in the caries development in orthodontic patients: An observational clinical study. Int J Paediatr Dent 27: 540-550.
- 28. Tashian RE (1989) Ue carbonic anhydrases: Widening perspectives on their evolution, expression and function. Bioessays 10: 186-192.
- Tashian RE (1992) Genetics of the mammalian carbonic anhydrases. Adv Genet 30: 321-356.
- Lönnerholm G, Selking Ö, Wistrand PJ (1985) Amount and distribution of carbonic anhydrases CA I and CA II in the gastrointestinal tract. Gastroenterology 88:1151-1161.
- Fleming RE, Parkkila S, Parkkila AK, Rajaniemi H, Waheed A et al. (1995) Carbonic anhydrase IV expression in rat and human gastrointestinal tract regional, cellular, and subcellular localization. J Clin Invest 96: 2907-2913.
- 32. Parkkila S, Parkkila AK (1996) Carbonic anhydrase in the alimentary tract. Roles of the di9erent isozymes and salivary factors in the maintenance of optimal conditions in the gastrointestinal canal. Scand J Gastroenterol 31: 305-317.
- **33**. Parkkila S, Parkkila AK, Juvonen T, Rajaniemi H (1994) Distribution of the carbonic anhydrase isoenzymes I, II, and VI in the human alimentary tract Gut 35: 646-650.
- 34. Pastorekova S, Parkkila S, Parkkila AK, Opavsky R, Zelnik V, et al. (1997) Carbonic anhydrase IX, MN/CA IX: Analysis of stomach complementary DNA sequence and expression in human and rat alimentary tracts. Gastroenterology 112: 398-408.
- Parkkila S, Kaunisto K, Rajaniemi L, Kumpulainen T, Jokinen K, et al. (1990) Immunohistochemical localization of carbonic anhydrase isoenzymes VI, II, and I in human parotid and submandibular glands. J Histochem Cytochem 38: 941-947.
- Fernley RT, Darling P, Aldred P, Wright RD, Coghlan JP et al. (1989) Tissue and species distribution of the secreted carbonic anhydrase isoenzyme. Biochem J 259: 91-96.
- Parkkila S, Parkkila AK, Vierjoki T, Stahlberg T, Rajaniemi, H (1993) Competitive time-resolved immunofluorometric assay for quantifying carbonic anhydrase VI in saliva. Clin Chem 39: 2154-2157.
- Kivelä J, Parkkila S, Waheed A, Parkkila AK, Sly WS et al. (1997) Secretory carbonic anhydrase isoenzyme (CA VI) in human serum. Clin Chem 43:2318-2322.
- Alhaija ES, Al-Khateeb SN, Al-Nimri KS (2005) Prevalence of malocclusion in 13-15 year-old North Jordanian school children. Community Dent Health 22: 266-271.
- Abu-Saad HH (2000) Challenge of pain in the cognitively impaired. Lancet 9245: 1867-1868.
- **41**. Abu-Tahun I, Rabah'ah A, Khraisat A (2012) A review of the questions and needs in endodontic diagnosis. Odontostomatol Trop 35: 11-20.
- Ackerman JL, Proffit WR (1980) Preventive and interceptive orthodontics: A strong theory proves weak in practice. Angle Orthod 50: 75-86.

- **43.** Afify AR, Zawawi KH (2012) Ue prevalence of dental anomalies in the Western region of Saudi Arabia. ISRN Dent 19: 837270.
- Aggarwal BB, Vijayalekshmi RV, Sung B (2009) Targeting inflammatory pathways for prevention and therapy of cancer: short-term friend, long- term foe. Clin Cancer Res 15: 425-430.
- Al-Emran S, Wisth PJ, Böe OE (1990) Prevalence of malocclusion and need for orthodontic treatment in Saudi Arabia. Community Dent Oral Epidemiol 18: 253-255.
- Al-Hashimi I, Levine MJ (1989) Characterization of in vivo salivaryderived enamel pellicle. Arch Oral Biol 34: 289-295.
- Bäßler-Zeltmann S, Kretschmer I, Göz G (1998) Malocclusion and the need for orthodontic treatment in 9-year-old children. J Orofacial Orthopedics/Fortschritte der Kieferorthopädie 59: 193-201.
- Belstrøm D, Damgaard C, Könönen E, Gürsoy M, Holmstrup P et al. (2017) Salivary cytokine levels in early gingival inflammation. J oral microbiol 9: 1364101.
- **49**. Birkhed D (1989) Salivary secretion rate, bu9er capacity, and pH, in Human Saliva. Clin Chemistry Microbiol 1: 50-52.
- 50. Borghi GN, Rodrigues LP, Lopes LM, Parisotto TM, Steiner-Oliveira C et al. (2017) Relationship among α amylase and carbonic anhydrase VI in saliva, visible biofilm, and early childhood caries: A longitudinal study. Int J Paediatr Dent 27: 174-182.
- Borzabadi-Farahani A, Borzabadi-Farahani A, Eslamipour F (2009) Malocclusion and occlusal traits in an urban Iranian population. An epidemiological study of 11-to 14-year-old children. Eur J Orthod 31: 477-484.
- 52. Burden DJ, Holmes A (1994) Ue need for orthodontic treatment in the child population of the United Kingdom. Eur J Orthod 16: 395-399.
- Chattopadhyay S, Davis RM, Menezes DD, Singh G, Acharya RU (2012) Application of Bayesian classifier for the diagnosis of dental pain. J Med Syst 36: 1425-1439.
- Alonso Chevitarese AB, Valle DD, Moreira TC (2003) Prevalence of malocclusion in 4-6 year old Brazilian children. J Clin Pediatr Dent 27: 81-85.
- 55. Ciancio DG (1985) A comparison of plaque accumulation in bonded versus banded teeth. J Dent Res 64: 325.
- Coetzee CE (1999) Development of an index for preventive and interceptive orthodontic needs (IPION) Doctoral dissertation. University of Pretoria 1-16.
- 57. Dawes C (2008) Salivary flow patterns and the health of hard and soI oral tissues. J Am Dent Assoc 139: 18S-24S.
- Edgar WM (1992) Saliva: its secretion, composition and functions. Br Dent J 172: 305-312.
- 59. Edgar WM, Higham SM, Manning RH (1994) Saliva stimulation and caries prevention. Adv Dent Res 8: 239-245.
- ERICSON D, BRATTHALL D (1989) Simplified method to estimate salivary bu9er capacity. Scand J Dent Res 97: 405-407.
- **61.** Ericsson Y (1959) Clinical investigations of the salivary bu9ering action. Acta Odontologica Scandinavica 17: 131-165.
- Estioko LJ, Wright FA, Morgan MV (1994) Orthodontic treatment need of secondary schoolchildren in Heidelberg, Victoria: an epidemiologic study using the Dental Aesthetic Index. Community dent Health 11: 147-151.
- FARSI NM, SALAMA FS (1996) Characteristics of primary dentition occlusion in a group of Saudi children. Int J Paediatr Dent 6: 253-259.
- Featherstone JD, Behrman JM, Bell JE (1993) E9ect of whole saliva components on enamel demineralization in vitro. Crit Rev Oral Biol Med 4: 357-362.
- Fenoll-Palomares C, Muñoz-Montagud JV, Sanchiz V, Herreros B, Hernández V et al. (2004) Unstimulated salivary flow rate, pH and bu9er capacity of saliva in healthy volunteers. Rev Esp Enferm Dig 96: 773-783.
- Fernley RT, Farthing J, Cooper EJ (1995) Radioimmunoassay for salivary carbonic anhydrase in human parotid saliva. Arch Oral Biol 40: 567-569.
- Finoti LS, Nepomuceno R, Pigossi SC, Corbi SC, Secolin R et al.(2017) Association between interleukin-8 levels and chronic periodontal disease:

A PRISMA-compliant systematic review and meta-analysis. Medicine Baltimore 96: e6932.

- 68. Frasseto F, Parisotto TM, Peres RC, Marques MR, Line SR et al. (2012) Relationship among salivary carbonic anhydrase VI activity and flow rate, biofilm pH and caries in primary dentition. Caries Res 46: 194-200.
- Geiger AM, Gorelick L, Gwinnett AJ, Griswold PG (1998) Ue e9ect of a fluoride program on white spot formation during orthodontic treatment. Am J Orthod Dentofacial Orthop 93: 29-37.
- Gopinath VK, Arzreanne AR (2006) Saliva as a diagnostic tool for assessment of dental caries. Archives of orofacial sciences 1: 57-59.
- Gorelick L, Geiger AM, Gwinnett AJ (1982) Incidence of white spot formation aler bonding and banding. Am J Orthod 81: 93-98.
- Gornowicz A, Bielawska A, Bielawski K, Grabowska SZ, Wójcicka A, et al. (2012) Pro-inflammatory cytokines in saliva of adolescents with dental caries disease. Ann Agric Environ Med 19: 711-716.
- 73. Heintze U, Birkhed D, Björn H (1983) Secretion rate and bu9er e9ect of resting and stimulated whole saliva as a function of age and sex. Swed Dent J 7: 227-238.
- Hill PA (1992) Ue prevalence and severity of malocclusion and the need for orthodontic treatment in 9-, 12-, and 15-year-old Glasgow schoolchildren. Br J Orthod 19: 87-96.
- 75. Holmes A (1992) Ue subjective need and demand for orthodontic treatment. British Journal of Orthodontics 19: 287-297.
- Ingervall B, Seeman L, Uilander B (1972) Frequency of malocclusion and need of orthodontic treatment in 10-year old children in Gothenburg. Sven Tandlak Tidskr 65: 7-21.
- 77. Jolley CJ, Huang GJ, Greenlee GM, Spiekerman C, Kiyak HA et al. (2010) Dental e9ects of interceptive orthodontic treatment in a Medicaid population: Interim results from a randomized clinical trial. Am J Orthod Dentofacial Orthop 137: 324-333.
- Karaiskos N, Wiltshire WA, Odlum O, Brothwell D, Hassard TH et al. (2005) Preventive and interceptive orthodontic treatment needs of an inner-city group of 6-and 9-year-old Canadian children. J Can Dent Assoc 71: 649.
- Kerosuo H, Väkiparta M, Nyström M, Heikinheimo K (2008) Ue seven- year outcome of an early orthodontic treatment strategy. J Dent Res 87: 584-548.
- Baelum V, Fejerskov O, Nyvad B (2008) Dental caries: Ue disease and its clinical management. Blackwell Munksgaard.
- Kiliçoğlu H, Yildirim M, Polater H (1997) Comparison of the e9ectiveness of two types of toothbrushes on the oral hygiene of patients undergoing orthodontic treatment with fixed appliances. Am J Orthod Dentofacial Orthop 111: 591-594.
- King GJ, Brudvik P(2010) E9ectiveness of interceptive orthodontic treatment in reducing malocclusions. Am J Orthod Dentofacial Orthop 137: 18-25.
- Kirstilä V (1997) Clinical significance of oral defense systems for oral health. In: ANNALES-UNIVERSITATIS TURKUENSIS SERIES D. Turun Yliopisto 259.
- Kivelä J, Parkkila S, Parkkila AK, Rajaniemi H (1999) A low concentration of carbonic anhydrase isoenzyme VI in whole saliva is associated with caries prevalence. Caries Res 33: 178-184.
- Kousvelari EE, Baratz RS, Burke B, Oppenheim FG (1980) Basic Biological Sciences: Immunochemical Identification and Determination of Proline-rich Proteins in Salivary Secretions, Enamel Pellicle, and Glandular Tissue Specimens. J Den Res 59: 1430-1438.
- Lagerlöf F(1994) Caries-protective factors in saliva. Adv Den Res 8: 229-238.
- Lamkin MS, Oppenheim FG (1993) Structural features of salivary function. Critl Rev Oral Biol Med 4: 251-259.
- Lundmark A, Johannsen G, Eriksson K, Kats A, Jansson L et al.(2017) Mucin 4 and matrix metalloproteinase 7 as novel salivary biomarkers for periodontitis. J Clin Periodontol 44: 247-254.

- Lundström FR, Hamp SE(1980) E9ect of oral hygiene education on children with and without subsequent orthodontic treatment. Eur J Oral Sci 88: 53-59.
- **90.** Lundström F, Krasse BO (1987) Caries incidence in orthodontic patients with high levels of Streptococcus mutans. Eur J Orthod 9: 117-121.
- **91.** Maren TH (1967) Carbonic anhydrase: Chemistry, physiology, and inhibition. Physiol Rev 47: 595-781.
- Meurman JH, Frank RM (1991) Scanning electron microscopic study of the e9ect of salivary pellicle on enamel erosion. Caries Res 25: 1-6.
- **93.** MizrahI E (1982) Enamel demineralization following orthodontic treatment. Am J Orthod 82: 62-67.
- **94.** MizrahI E (1983) Surface distribution of enamel opacities following orthodontic treatment. Am J Orthod 84: 323-331.
- Navazesh M, Christensen CM (1982) A comparison of whole mouth resting and stimulated salivary measurement procedures. J Dent Res 61: 1158-1162.
- Ogaard B (1989) Incidence of filled surfaces from 10-18 years of age in an orthodontically treated and untreated group in Norway. Eur J Orthod 11: 116-119.
- Onyeaso CO (2004) Prevalence of malocclusion among adolescents in Ibadan, Nigeria. Am J Orthod Dentofacial Orthop 126: 604-607.
- 98. Ozturk LK, Furuncuoglu H, Atala MH, Ulukoylu O, Akyuz S, et al. (2008) Association between dental-oral health in young adults and salivary glutathione, lipid peroxidation and sialic acid levels and carbonic anhydrase activity. Braz J Med Biol Res 41: 956-959.
- Parkkila S, Parkkila AK, Lehtola J, Reinila A, Sodervik HJ, et al. (1997) Salivary carbonic anhydrase protects gastroesophageal mucosa from acid injury. Dig Dis Sci 42: 1013-1019.
- Parkkila S, Parkkila AK, Rajaniemi H (1995) Circadian periodicity in salivary carbonic anhydrase VI concentration. Acta Physiol Scand 154: 205-211.
- Peros K, Mestrovic S, anic-milosevic S, Slaj M (2011) Salivary microbial and nonmicrobial parameters in children with fixed orthodontic appliances. Angle Orthod 81: 901-906.
- 102. Popovich F, Uomson GW (1975) Evaluation of preventive and interceptive orthodontic treatment between three and eighteen years of age. Uird International Orthodontic Congress 20: 260-281.
- 103. Rangbulla V, Nirola A, Gupta M, Batra P, Gupta M et al. (2017) Salivary IgA, Interleukin-1beta and MMP-8 as Salivary Biomarkers in Chronic Periodontitis Patients. Chin J Dent Res 20: 43-51.
- 104. Rechenberg DK, Galicia JC, Peters OA (2016) Biological markers for pulpal inflammation: A systematic review PLoS One 11: e0167289.
- 105. Russell JI, Macfarlane TW, Aitchison TC, Stephen KW, Burchell CK <sup>et al.</sup> (1990) Caries prevalence and microbiological and salivary caries activity tests in Scottish adolescents. Community Dent Oral Epidemiol 18: 120-125.
- Saloum FS, Sondhi A (1987) Preventing enamel decalcification aler orthodontic treatment. J Am Dent Assoc 115: 257-261.
- 107. Sharma V, Gupta N, Srivastava N, Rana V, Chandna P, et al. (2017) Diagnostic potential of inflammatory biomarkers in early childhood caries: A case control study. Clin Chim Acta 471: 158-163.
- 108. Shaw WC, Richmond S, O'Brien KD (1995) Ue use of occlusal indices: A European perspective. Am J Orthod Dentofacial Orthop 107: 1-10.
- 109. Stratemann MW, Shannon IL (1974) Control of decalcification in orthodontic patients by daily self-administered application of a water-free 0.4 per cent stannous fluoride gel. Am J Orthod 66: 273-279.
- 110. Symons FJ, Elghazi I, Reilly BG, Barney CC, Hanson L, et al. (2015) Can biomarkers di9erentiate pain and no pain subgroups of nonverbal children with cerebral palsy? A preliminary investigation based on noninvasive saliva sampling. Pain Med 16: 249-256.
- Syndergaard B, Al-sabbagh M, Kryscio RJ, Xi J, Ding X, et al. (2014) Salivary biomarkers associated with gingivitis and response to therapy. J Periodontol 85: e295-303.
- **112**. Szabo I (1974) Carbonic anhydrase activity in the saliva of children and its relation to caries activity. Caries Res 8: 187-191.

- Tenovuo J (1997) Salivary parameters of relevance for assessing caries activity in individuals and populations. Community Dent Oral Epidemiol 25: 82-86.
- 114. Uilander B, Myrberg N (1973) Ue prevalence of malocclusion in Swedish schoolchildren. Scand J Dent Res 81: 12-20.
- 115. Uilander B, Pena L, Infante C, Parada SS, DE Mayorga C (2001) Prevalence of malocclusion and orthodontic treatment need in children and adolescents in Bogota, Colombia. An epidemiological study related to di9erent stages of dental development. Eur J Orthod 23: 153-167.
- 116. Uilander B, Wahlund S, Lennartsson B (1984) Ue e9ect of early interceptive treatment in children with posterior cross-bite. Eur J Orthod 6: 25-34.
- 117. Tulloch JF, Proffit WR, Phillips C (2004) Outcomes in a 2-phase randomized clinical trial of early Class II treatment. Am J Orthod Dentofacial Orthop 125: 657-667.
- **118.** Westbom L, Rimstedt A, Nordmark E (2017) Assessments of pain in children and adolescents with cerebral palsy: A retrospective population-based registry study. Dev Med Child Neurol 59: 858-863.

- **119.** Wheeler TT, Mcgorray SP, Yurkiewicz L, Keeling SD, King GJ et al. (1994) Orthodontic treatment demand and need in third and fourth grade schoolchildren. Am J Orthod Dentofacial Orthop 106: 22-33.
- Zachrisson BJ (1977) A posttreatment evaluation of direct bonding in orthodontics. Am J Orthod 71: 173-189.
- Zachrisson BU, Zachrisson S (1971) Caries incidence and oral hygiene during orthodontic treatment. Scand J Dent Res 79: 394-401.
- 122. Zahradnik RT, Moreno EC, Burke EJ (1976) E9ect of salivary pellicle on enamel subsurface demineralization in vitro. J Dent Res 55: 664-670.
- 123. Zahradnik RT, Propas D, Moreno EC (1977) In vitro enamel demineralization by Streptococcus mutans in the presence of salivary pellicles. J Dent Res 56: 1107-1110.
- 124. Zahradnik RT, Propas D, Moreno EC (1978) E9ect of salivary pellicle formation time on in vitro attachment and demineralization by Streptococcus mutans. J Dent Res 57: 1036-1042.