

Sundell et al. BMC Oral Health (2010) 11:85
DOI 10.1186/1472-6556-11-85

RESEARCH ARTICLE Open Access

Comparing caries risk profiles between 5- and 10-year-old children with cleft lip and/or palate and non-cleft controls

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Abstract
Background: Previous studies have suggested that children with clefts may have higher caries prevalence as compared to non-cleft controls, but the relative importance of the potential risk factors is not clear. The aim of this study was to compare the caries risk profiles in a group of cleft lip and/or palate (CLP) children with non-cleft controls in terms of age at diagnosis and a complicated caries risk assessment model.

Methods: A cross-sectional study was conducted among 5- and 10-year-old children with cleft lip and/or palate (CLP) aged 5 years and 164 aged 10 years and 252 non-cleft controls (113 aged 5 years and 164 aged 10 years). A questionnaire was used to collect data concerning the child's oral hygiene routines, dietary habits and fluoride exposure. Oral hygiene was assessed using Görtz Oral Hygiene Index (GOHI) and dental caries were assessed using the World Health Organization (WHO) Caries Detection and Assessment System. Whole saliva samples were analyzed for mutans streptococci, lactobacilli, buffering capacity and lactate levels. The risk factors and risk profiles were calculated by comparing the groups with and without caries experience and categorized as "high" or "low".

Results: Children with CLP (the entire study group) had significantly higher caries rates (salivary lactobacilli > 6.02) and displayed less good oral hygiene ($\text{GOHI} < 0.5$). More 10-year-old children in the CLP group had low secretion rate than non-cleft controls. There were no significant differences in the odds of being categorized with high caries risk in the CLP group (OR 1.25–2.86). In both groups, children in the high risk category had a higher caries experience than those with low risk.

Conclusion: Children with CLP displayed increased odds of being categorized as high caries risk with impaired oral hygiene and low salivary lactobacilli. According to the results, the results suggest that a caries risk assessment model should be applied in the routine CLP care as a basis for the clinical decision-making and implementation of primary and secondary caries prevention.

Keywords: Cleft lip, Cleft palate, Cleft lip and/or palate, Caries Risk, Children

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Cleft Lip and Palate An Evidence-Based Review

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KEY WORDS

- Cleft lip • Cleft palate • Evidence-based medicine • Outcomes

KEY POINTS

- The repair of unilateral cleft lip is performed using a rotation advancement, genovate, straight-line, or hybrid technique.
- For bilateral cleft lip repair, most surgeons use either the Millard or Mulliken techniques, and their variations.
- Most cleft centers perform cleft lip repair at the age of 3 to 6 months.
- Pre surgical infant orthopedics, which can include nasoalveolar molding, is used before definitive surgery.
- For cleft palate repair, the 2-flap palatoplasty and Furlow double-opposing Z-plasty are most commonly used.

INTRODUCTION

An estimated prevalence of 16.66 cases per 10,000 live births, bilateral cleft palate, as well as cleft lip with or without cleft palate, is the most common congenital orofacial malformation in the United States. Worldwide, cleft lip and palate may require a multitude of physical and developmental challenges. There also may be psychosocial and emotional concerns for the patients and their families. As a consequence, cleft lip and palate repair with cleft lip or palate repair requires an interdisciplinary team. The guidelines for team care outlined in this article are intended to assist the cleft lip and palate surgeon in endeavors that may include anesthesia, otolaryngology, genetics, neurosurgery, nursing, ophthalmology, oral maxillofacial surgery, orthodontics, otolaryngology-head and neck surgery,

pediatrics, pediatric dentistry, physical anthroplogy, plastic surgery, prosthodontics, psychiatry, psychology, social work, and speech-language pathology.¹ Although every specialist may not be represented on the team, it is important to have a strong collaborative division and coordination of care.

Broadly speaking, orofacial cleft anomalies may be congenital or acquired. As such, the literature is both. Although there have been considerable publications on this topic, most are single-surgeon/career experience papers or are limited to case reports.^{2–4} As such, the literature on cleft palate literature regarding the clinical and surgical decision points lacks consensus. This review article will provide an evidence-based approach to management of cleft lip and palate. It will present plans, describe the various viewpoints, and suggest recommendations based on the levels of evidence (Table 1) on the management of cleft lip.

Funding sources: none.

Competing interests: none.

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Original Paper

Caries Res 2013;47:406–415
DOI: 10.1007/s00155-013-0090-y
Published online May 3, 2013

Caries Prevalence in Non-Syndromic Patients with Cleft Lip and/or Palate: A Meta-Analysis

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Key Words: Caries prevalence – Cleft lip and/or palate – Meta-analysis

Abstract: **Aim:** To evaluate caries prevalence in non-syndromic patients with cleft lip and/or palate (CLP) in comparison with non-CLP individuals. **Method:** A systematic search of the literature was conducted in order to identify articles reporting on the prevalence of caries in CLP versus non-CLP individuals. The outcome measures were caries prevalence and mean difference between the groups. Unpublished manuscripts and abstracts were used to expand the search. Only studies with a suitable matched control group were included. From each study, the mean difference and standard deviation were extracted, as were results. The meta-analysis was done to score for caries prevalence in each study, using a well-defined index. The heterogeneity was tested using the Q statistic and a meta-analysis was performed using the random effects model.

Results: From the 502 articles initially identified, 7 were chosen for inclusion. The remaining 495 articles were excluded. All the studies were cross-sectional in nature, and used the decayed, missing, and filled (DMF) index as the final outcome measure. The caries prevalence in CLP patients aged 1–29 years was 20.7% compared to 14.5% in non-CLP patients aged 1–29 years. When looking at permanent teeth, data from 3 studies suggest that CLP patients have a higher number of DMF teeth than the controls (mean difference 1.39 ± 0.60), for deciduous teeth, data from 4 studies suggest

that CLP patients have a higher number of dental fissures than the controls (mean difference 1.33, $p = 0.03$). **Conclusion:** Non-syndromic patients with CLP tend to have higher caries prevalence, both in the permanent and the deciduous dentition, in comparison with non-CLP individuals.

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MEDICAL SCIENCE MONITOR

Received: 2013-01-10
Accepted: 2013-03-10
Published: 2013-05-23

Author Contributions: Agnieszka Gornowicz – Conceptualization, Data curation, Formal analysis, Methodology, Project administration, Resources, Software, Supervision, Validation, Writing – original draft preparation, Writing – review and editing; Grzegorz Tokajuk – Conceptualization, Data curation, Formal analysis, Methodology, Project administration, Resources, Software, Supervision, Validation, Writing – original draft preparation, Writing – review and editing; Małgorzata Maksymczak – Conceptualization, Data curation, Formal analysis, Methodology, Project administration, Resources, Software, Supervision, Validation, Writing – original draft preparation, Writing – review and editing; Elżbieta Majkowska – Conceptualization, Data curation, Formal analysis, Methodology, Project administration, Resources, Software, Supervision, Validation, Writing – original draft preparation, Writing – review and editing; Robert Jabłoski – Conceptualization, Data curation, Formal analysis, Methodology, Project administration, Resources, Software, Supervision, Validation, Writing – original draft preparation, Writing – review and editing; Krzysztof Bielański – Conceptualization, Data curation, Formal analysis, Methodology, Project administration, Resources, Software, Supervision, Validation, Writing – original draft preparation, Writing – review and editing.

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Source of Support: This publication was supported by Research grant 2014-03-06-00-07 from Medical University of Białystok, Poland.

Background: Saliva contains a number of protective factors such as enzymes, immunoglobulins (e.g. IgA, IgG, and IgM), and enzymes (e.g. lysozyme and lactoperoxidase) that play an important role in the maintenance of oral health. The aim of this study was to compare levels of sIgA, histatin-5, and lactoperoxidase in saliva of adolescents with dental caries.

Material/Methods: 14 healthy adolescents (age 14 years) from high school were examined. Eight subjects with DMF > 1 (Group I) and 24 adolescents with DMF ≤ 1 (Group II) were enrolled for this study. Clinical evaluation procedures comprised oral examination (including teeth, periodontal and oral mucosal status) and collection of saliva samples. Saliva samples were analyzed for total lactoperoxidase (LPO) assay (ELISA) and was used for determination of sIgA, histatin-5, and lactoperoxidase levels.

Results: Our results showed that adolescents with very high intensity of dental caries (DMF > 1) had increased levels of sIgA, histatin-5, and lactoperoxidase compared to adolescents with lower intensity of caries. The increase was statistically significant ($p < 0.05$). We suggest that high levels of caries are associated with increased levels of some salivary components – sIgA, histatin-5, and lactoperoxidase – that possess strong antimicrobial or bactericidal effects, resulting in aggregation of oral bacteria and their clearance from the oral cavity.

Conclusions: Dental Caries Prevalence – Histatin-5 + Lactoperoxidase + Saliva

MeSH Keywords: Dental Caries Prevalence • Histatin-5 • Lactoperoxidase • Saliva

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UNIVERSITAS ISLAM SULTAN AYYUBI جامعة سلطان اوجي الإسلامية

Association between Maternal Folate Intake and Polymorphism MTHFR A1298C as Risk Factor of Non-Syndromic Cleft Lips

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ABSTRACT

Background: Methylentetrahydrofolate reductase (MTHFR) is often associated with the incidence of orofacial clefts. Folic acid deficiency has gained considerable attention because of its promising role in reducing diverse clinical condition such as cleft. The objective of the study is to describe the association of MTHFR A1298C polymorphism and maternal folic acid intake with an orofacial cleft in Sultak Population.

Method: This study used control case design, the number of the subjects were 148 who were divided into case groups and their mother (70 women) and control groups and their mother (78 items). The detection of Polymorphism MTHFR A1298C used PCR-RFLP and sequencing for confirmation. The information on the dietary pattern and folic acid intake used PQ (Food Frequency Questionnaire).

Results: MTHFR A1298C polymorphism was associated with maternal folic acid intake in Sultak ($p = 0.00$, $\chi^2 = 14.7$ CI 95% (2.49–85.53) for cases and 0.041, OR = 4.4 CI 95% (3.19–19.16) for control group. Maternal folic acid intake was associated with cleft ($p = 0.007$) OR = 2.7 CI 95% (1.06–6.94) in Sultak Population.

Conclusion: Maternal folic acid was as the risk factor cleft lip/palate in Sultak population and association with MTHFR A1298C Polymorphism.

Keywords: Polymorphism MTHFR A1298C, folic acid, orofacial cleft

A1298C is MTHFR genotype variants that are thought to contribute to folate deficiency.^{1–3} MTHFR is a genetic enzyme that converts 5-(deoxyribose)folic acid into 5-methyltetrahydrofolate in folate cycle. The endogenous folate cycle is a predominant methyl donor to methylethyl homocysteine (MeHCy) into methionine.⁴

Pregnant women with MTHFR polymorphism have a higher risk to get folate deficiency.⁵ The food sources that rich in folic acid are liver, fish, and meat, mushrooms, green leafy vegetables such as spinach, beans leaves, nuts, and yeast. Food processing can decrease 50–90% of folate content. Heating, cooking, and boiling can destroy folate.⁶ Folic acid and folate are conjugated by Gamma-glutamyl-folic acid into megaloblastic acid by Zick.⁷ Folate deficiency and abnormal metabolism of folic acid and MeHCy play a significant role in the incidence

KARGER
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 ISSN: 0939-6068 • Volume 56 • Number 1 • DOI: 10.1159/000356068
 Available online at www.karger.com
Review Article
Saliva as a diagnostic tool for oral and systemic diseases

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ARTICLE INFO

ABSTRACT

Early disease detection is not only vital to reduce disease severity and prevent complications but also to reduce the cost of treatment. Saliva is a potential biomarker to detect the disease due to its ease and non-invasive sampling along with its abundance of biomarkers, such as genetic material and proteins. Saliva is a non-invasive diagnostic sample that can be used to detect various diseases (cancer, systemic diseases, diabetes, hypertension, heart diseases, oral diseases, and periodontal diseases). Considering data accuracy, efficacy, and convenience, saliva is considered as a promising diagnostic tool. The aim of this study is to evaluate the usefulness of saliva as a diagnostic tool for oral and systemic diseases. The saliva can be used as a wide range of oral and systemic diseases as the early warning and check-up tool.

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1. Introduction

Early diagnosis of diseases is crucial to prevent complications that could have a negative impact on a patient's quality of life.

For instance, elevation rates of the 50% most common cancers and most common diseases in the world are increasing rapidly at stage 0 in comparison to 90% if diagnosed at stage I. Similarly, type 2 diabetes, 90% of the adult population in Britain, can be detected by molecular diagnostics, and antibodies in specific serols and proteins present in the saliva can be used to detect oral and systemic diseases (Table 1). Likewise, if the diagnosis is made earlier, for instance, despite the early warning and check-up diseases can

be detected and treated at an early stage, the cost of treatment will be reduced.⁸

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ISSN 0939-6068 • Volume 56 • Number 1 • DOI: 10.1159/000356068

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RESEARCH ARTICLE

Psychosocial Stress Increases Salivary Alpha-Amylase Activity Independently from Plasma Noradrenaline Levels

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Abstract

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Citation: Potruska L, Doering BK, Vitz S, Engler H, Haef M, Schmidova M, Grigoleit JS (2013) Psychosocial Stress Increases Salivary Alpha-Amylase Activity Independently from Plasma Noradrenaline Levels. *PLoS ONE* 8(8): e73881. doi:10.1371/journal.pone.0073881

Editor: Ulrich Schmidt, Max Planck Institute of Psychiatry, GERMANY

Received: February 18, 2013

Accepted: July 12, 2013

Published: August 6, 2013

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Competing interests: No competing interests declared.

Data Availability Statement: All relevant data are within the paper.

Funding: This study was funded by grants from Deutsche Forschungsgemeinschaft, Dr. Rennkamp-Stiftung, and the Max-Planck-Institut für Psychiatrie. The funders had no role in study design or in the collection, analysis, interpretation of data, or writing of this manuscript.

Introduction

Salivary alpha-amylase (sAA) is one of the major enzymes in the oral cavity. Beyond its primary function, the hydrolysis of starch and glycogen, it is involved in defense against bacteria with low sAA activity being related to a higher risk of oral infections [1]. In addition, increased salivary sAA activity suggests sAA as surrogate marker of sympathetic activation [2].

Since salivary cortisol became the standard indicator of hypothalamic-pituitary-adrenal (HPA) axis activity, a comparably easy-to-use salivary measure for activity of the sympathetic-adrenomedullary system (SAS) has highly been desired. First evidence that stress-inducible changes in sAA activity in humans may be dependent on beta-adrenergic transmission came from a study in which an increase in sAA activity in response to a cold water stressor was perceived

Research article

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The relationship between the level of salivary alpha-amylase activity and pain severity in patients with symptomatic irreversible pulpitis

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Abstract

Background: Assessment of dental pain severity is very challenging in dentistry. Previous studies have suggested that elevated salivary alpha-amylase may contribute to localized dental distress. There is a lack of evidence concerning salivary alpha-amylase activity in patients with symptomatic irreversible pulpitis. The aim of this study was to evaluate the relationship between pain severity and salivary-alpha-amylase activity in patients with symptomatic irreversible pulpitis. **Methods:** Thirty healthy volunteers were selected. The visual analog scale (VAS) score was used to assess the pain intensity in each patient. Unstimulated whole saliva was collected, and salivary alpha-amylase activity was measured by spectrophotometry at 405 nm. Statistical analysis was performed using SPSS 13. **Results:** The level of salivary alpha-amylase activity significantly correlated with VAS pain score in association with pulpitis induced by carious lesions and/or traumatic fractures. **Conclusion:** There was a significant correlation between the VAS pain scale and salivary alpha-amylase levels, which indicates this biomarker may be a good index for the subjective assessment of pain severity. **Keywords:** Alpha Amylase, Pain, Pulpitis, Saliva

Introduction

Pain is an unpleasant feeling, a generic term for the sensory response to the body's internal or external environment. It can also be induced by psychological causes or relieved from other sites. Therefore, difficulty sometimes arises in finding an aetiological cause. Pain expression, which may happen after dental caries or dental trauma, is a complex process involving both peripheral and central nervous systems. Significant temporal changes could follow long and severe pulpal pain, which may even be permanent after the removal of the stimulus. The pain will be more severe and persistent in comparison to the peripheral afferents and the surrounding tissue.

Saliva has many functions, and 50–60% of salivary proteins consist of salivary amylases.¹ Salivary amylase is a single-chain protein with a molecular weight of approximately 60 kDa and it provides the highest enzyme activity of all the salivary enzymes.² Although salivary amylase is not a physiologically strong anti-bacterial agent,³ it cannot be considered a circumscribed indicator of the caries-associated levels.⁴ Salivary amylase found a significant relationship between pain and the level of salivary alpha-amylase.⁵

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RESEARCH ARTICLE

Open Access

Comparing caries risk profiles between 5- and 10-year-old children with cleft lip and/or palate and non-cleft controls

Anna Lena Sundell¹, Christine Ulter², Agneta Marcusson¹ and Svante Twetman^{1*}

Abstract

Background: Previous studies have suggested that children with oral clefts may have higher caries prevalence in comparison with non-cleft controls, but the relative importance of the potential risk factors is not clear. The aim of this study was to compare caries risk profiles between children with cleft lip and/or palate (CLP), children with non-clift controls in the same age using a comprehensive caries risk assessment model.

Methods: The study group consisted of 133 children with CLP (77 subjects aged 5 years and 56 aged 10 years) and 130 non-clift controls (70 subjects aged 5 years and 60 aged 10 years). The caries risk was assessed by means of a questionnaire concerning the child's oral hygiene routines, dietary habits and fluoride exposure. Oral hygiene was assessed using Quigley-Hein plaque index and the caries prevalence and frequency was scored according to the International Caries Detection and Assessment System (ICDAS).

Results: Children with CLP in the study group had significantly higher counts of salivary lactobacilli ($p = 0.02$) and displayed less good oral hygiene ($p < 0.05$). More 10-year-old children in the CLP group had low secretion rate for saliva ($p = 0.03$) and more children in the CLP group had a high caries risk ($p = 0.01$). However, there were no significant differences between the groups. The odds of being categorized with high caries risk in the CLP group was significantly reduced ($OR = 0.89$, 95% CI = 0.75–1.06). In both groups, children in the high risk category had a significantly higher caries risk ($p < 0.001$).

Conclusion: Children with CLP displayed increased odds of being categorized at high caries risk with impaired oral hygiene and elevated salivary lactobacilli counts as most influential factors. The results suggest that a caries risk assessment model can be used to identify children at high caries risk as a basis for the clinical decision-making and implementation of primary and secondary caries prevention.

Keywords: Cleft lip, Cleft palate, Oral health and/or caries, Caries risk, Program, Children

Background

Cleft lip and/or palate (CLP) is the most common congenital craniofacial deformity, affecting nearly two in every 1000 live births worldwide. The relationship between CLP and dental caries in children is not fully clear but a number of studies indicate a higher caries prevalence in children with CLP compared to non-clift controls with non-clift controls (2–4). Several factors can contribute to this higher susceptibility such as impaired hygiene (2, 5, 7, 8), enamel hypoplasia (9, 10) and diet (11). Furthermore, parents to children with CLP tend to overestimate the children and offer them more protection than non-clift children due to their medical condition (12, 13). The prolonged oral clearance time of the caries-associated bacteria in a cleft environment (14), the role and relative importance of the potential risk factors are however not

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