Estimation of Salivary Magnesium Levels in Patients with Oral Squamous Cell Carcinoma

Abstract

India holds the highest rank in terms of incidence of oral cancer. Early detection is essential to reduce the mortality rate results in late cancer diagnosis. Recent studies emphasize micronutrients or trace elements as harbingers in oral squamous cell carcinoma (OSCC). These trace elements play a dual role in cancer cell proliferation and act as anti-cancer agents. Magnesium (Mg) forms an essential component of the micronutrient family of the body. It has many functions, including regulating the cell cycle and proliferation, metabolic activities, and other physiological functions. Impaired Mg homeostasis is associated with various pathological conditions. It has gained much importance in the past decade due to its complex relationship with cancer, as alterations in serum and salivary Mg have been reported in cancer patients. This study aimed to estimate salivary Mg levels in patients with OSCC and compare them with those of healthy individuals for Mg to serve as a diagnostic tool. A case-control study was performed on 36 subjects with 18 per group of cases, with OSCC and control being healthy individuals. Unstimulated saliva was collected from each individual and subjected to analysis of salivary magnesium using the xylidyl blue method. The mean salivary Mg levels in patients with OSCC and healthy individuals did not exhibit a statistically significant difference using the ANOVA test. The comparison of salivary Mg in patients with OSCC and healthy individuals did not provide sufficient evidence of Mg as an alleged diagnostic tool or a potential biomarker for OSCC.

Keywords: Biomarkers, Diagnosis, Micronutrients, Trace elements

Introduction

Globally, India holds the highest rank in terms of incidence of oral cancer, with a higher prevalence in males. A delay in cancer diagnosis results in a high mortality rate. Hence early detection is essential to reduce the mortality rate. Consumption of alcohol and tobacco in smoking or non-smoking forms is the main contributor to oral cancer. Tobacco consists of many carcinogenic components, of which polycyclic hydrocarbons and nitrosamines are the key players. Consumption of alcohol is linked to a higher risk of oral cancer even in patients who are non-smokers. Recent studies emphasize micronutrients or trace elements as harbingers in oral squamous cell carcinoma (OSCC). These trace elements play a dual role in cancer cell proliferation and act as anti-cancer agents, thus proving their versatile nature. Many researchers have evaluated the correlation of trace elements like copper (Cu), zinc (Zn), calcium (Ca), iron (Fe), and magnesium (Mg) with cancer mortality.

In living cells, the most abundant cation present is Mg. It acts as a secondary messenger controlling intracellular signaling, cell growth, and cell survival. It performs various physiological functions, including cell-energy metabolism, activation of cell-cytoskeleton, maintaining cell membrane integrity, synthesis of proteins, antioxidation, and replication of DNA. Mg's physiological levels are required in DNA repair and maintenance of genomic stability concerning cancer. There is a complex relationship between magnesium levels in the body and cancer occurrence. It has been found that both excess and deficiency of magnesium can impact carcinogenesis. The potential role of Mg in cell energy metabolism is solely responsible for increasing cell proliferation. Thus, neoplastic cells show significant Mg influx, leading to cancer progression.

Although histopathological examination remains the gold standard for diagnosing OSCC, recent studies focus on parallel investigations. The emphasis is solely made on the examination of certain salivary variables to determine the occurrence of
OSCC. However, the results are sometimes the least likely.[13] Saliva is a substantial fluid that is important in the human body. It is required to maintain the integrity of oral structures and protect the oral mucosa from infection.[17] The oral cavity is continuously bathed by saliva. Therefore, saliva could represent the changes associated with the oral cavity at cellular and molecular levels through the variation in its composition.[18, 19] Molecular components found in other body fluids like serum and urine are also present to a certain extent in the saliva; thus, saliva has been used as a diagnostic tool for various diseases.[20] Saliva is composed of 98% water, while the remaining 2% consists of electrolytes, glycoproteins, and antibacterial substances like immunoglobulins and lysosomal enzymes.[19, 21] It contains a broad spectrum of biomarkers to detect various diseases. Saliva collection is a non-invasive procedure; its easily stored and transported, making it economical and efficient.[22] Recent studies and technologies have shown that saliva can be used as a tool for the diagnosis of cancer, immunodeficiency, hormone imbalances, and liver function.[5] Salivary electrolyte composition varies in individuals due to age, oral health status, and adverse habits like smoking and consumption of alcohol. In OSCC, the oral cavity environment is compromised, impacting the salivary composition and alteration of certain micro-nutrients. Assessing the altered levels of such micro-nutrients in saliva can diagnose various oral cancers.[21]

Saliva lies nearby of oral tissues, and any modification in oral tissues certainly reflects in the salivary composition, and estimating salivary Mg in patients with OSCC can be a sensitive aid in diagnosis.[16] Thus, the present study aimed to evaluate salivary magnesium levels in patients with OSCC and determine whether it can serve as a biomarker in the diagnosis and prognosis of OSCC.

Materials and Methods

This case-control study was performed on patients diagnosed with OSCC. Patients visiting the oral pathology and microbiology department for biopsy procedures were taken as subjects after microscopic evaluation for the presence of OSCC. The Institutional Ethics Committee at Dr. D. Y. Patil Dental College and Hospital, Pune, India, approved the experimental protocol (DYPDCH/IEC/123/129/19) for undertaking this study.

The sample consisted of 36 individuals, calculated assuming a confidence interval level (almost 95%). Out of the 36 participants, 18 patients were those diagnosed with OSCC histopathologically, and the remaining 18 were otherwise healthy patients visiting the hospital for routine dental procedures. The study participants were subjected to detailed clinical examination to identify other potential lesions. A detailed case history was obtained from each individual before sample collection. The entire procedure of the study was explained to every participating subject, and informed consent was taken from everyone accordingly. The study was conducted between June 2020 and June 2021.

Sample collection was performed in the morning from 10:00 AM to 12:00 PM to avoid diurnal variation. Participants were exempted from consuming any food or drinks for 2 hours before the collection period.[9] Whole unstimulated saliva around 3 ml was collected from every individual where participants were asked to pool saliva on the floor of the mouth for a minute, and by the drooling method, the accumulated saliva was collected in a sterile glass tube. The collected samples were subjected to centrifugation at 2500 rpm for 15 mins at 4 degrees Celsius. The supernatants were stored at -20 degrees for further analysis of Mg. Salivary Mg was estimated using the xylidyl blue method and the values were obtained in mg/dl as described previously.[23] The obtained mean values were compared among the cases and controls.

Statistical analysis to determine the difference in salivary Mg was performed using Unpaired t-Test and One-way ANOVA in IBM SPSS software version 20. Data were expressed as Mean±SD, and significance levels were determined; the p-value of <0.05 was considered statistically significant.

Results and Discussion

A total of 36 individuals were enrolled in the present study. Out of which, 18 individuals were histopathologically proven cases of OSCC, and 18 were healthy individuals taken as the control group. After performing a detailed case history, it was observed that all OSCC patients reported an adverse habit history of consumption of tobacco in smoking or non-smoking form. The summary of habit history is mentioned in (Table 1).

**Table 1. Types of adverse habits among OSCC patients.**

<table>
<thead>
<tr>
<th>Type of habit</th>
<th>Number of patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tobacco chewing only</td>
<td>6</td>
</tr>
<tr>
<td>Tobacco chewing, areca nut consumption, Mishri, Pan consumption</td>
<td>1</td>
</tr>
<tr>
<td>Tobacco chewing and smoking</td>
<td>3</td>
</tr>
<tr>
<td>Tobacco chewing and alcohol</td>
<td>4</td>
</tr>
<tr>
<td>Mishri only</td>
<td>1</td>
</tr>
<tr>
<td>Areca nut only</td>
<td>2</td>
</tr>
<tr>
<td>Tobacco chewing and Mishri</td>
<td>1</td>
</tr>
</tbody>
</table>

Mean salivary Mg levels were more significant in OSCC patients than in the control group. However, a statistically significant difference was not evident according to the unpaired t-test. The salivary Mg levels in OSCC and control groups are presented in (Tables 2-4).

**Table 2. Salivary Mg levels in OSCC and control groups estimated from the present study.**

<table>
<thead>
<tr>
<th>Groups</th>
<th>Mean</th>
<th>Standard Deviation</th>
<th>Median</th>
<th>95% Confidence Interval [Lower bound]</th>
<th>95% Confidence Interval [Upper bound]</th>
</tr>
</thead>
<tbody>
<tr>
<td>OSCC</td>
<td>0.5778</td>
<td>0.36551</td>
<td>0.5</td>
<td>0.3960</td>
<td>0.7595</td>
</tr>
<tr>
<td>Control</td>
<td>0.4333</td>
<td>0.28901</td>
<td>0.45</td>
<td>0.2896</td>
<td>0.5771</td>
</tr>
</tbody>
</table>
The spread of oral cancer is turning into an alarming epidemic. OSCC is the leading cause of mortality in terms of oral cancer. The etiological association of OSCC is usually with carcinogens in tobacco and its related products consumed by people through the oral cavity. To name a few are nitrosamines and polynuclear aromatic hydrocarbons. Recent studies point towards the association of various parallel factors playing an important role in carcinogenesis despite all these facts. These parallel factors are nothing but trace elements mentioned above that undergo alterations during the progression of OSCC. Assessing the alterations in the body fluids is a critical step toward early diagnosis. A higher incidence of OSCC is witnessed in industrialized countries where tobacco and alcohol consumption is more considerable. Consumption of tobacco in smoking or non-smoking form and areca nut has shown alterations in body fluids of microelements such as Mg, Zn, Cu, and Ca.

Saliva is an essential body fluid. Its close approximation with oral cavity saliva exhibits micro-molecular changes associated with oral tissues. The non-invasive collection of saliva as a sample for investigation has further enhanced the potential use of saliva as a diagnostic tool in any diseased condition. Of all the trace elements found in the body, Mg counts as the most abundant mineral essential during the cell cycle. Its regulatory role during protein synthesis and DNA replication has been investigated. In the present study, we evaluated the salivary magnesium levels of patients with OSCC and compared them with those obtained from an equal number of healthy controls to study its association with the disease. The mineral composition found in individuals consuming tobacco-based products and those without adverse habits is variable. Dziewulska et al. studied salivary mineral composition in patients with oral cancer and found no significant difference in the salivary magnesium levels between oral cancer patients and healthy control following the results observed in the present study. Kolte et al. emphasized that smoking influences the salivary mineral composition, increasing the saliva's magnesium levels. These findings are applicable in the present study, where patients with OSCC have been noted to have tobacco-related adverse habits. The increased magnesium levels in our study group can be associated with tobacco consumption in smoking or non-smoking form. Similar findings were noted by Kode et al. while evaluating the levels of trace elements in the saliva of patients with oral submucous fibrosis (OSMF). Shpitzer et al. found higher magnesium concentrations in the saliva of 25 patients with OSCC associated with the lateral aspect of the tongue, which correlated with the present study. However, contrasting results were obtained by Al-Rawi and his associates, where statistically significant lower salivary magnesium levels were estimated in patients with oral cancer. Estimation of salivary magnesium levels in patients with premalignant lesions has concluded with a significantly lower level of mean salivary magnesium in the diseased state than in the healthy state. In a quantitative serum analysis for magnesium in patients with OSMF and OSCC, Hosthor et al. found lower serum magnesium levels in their diseased subjects. The conflicting results can be explained based on adverse habits (consumption of tobacco) of patients with OSCC and fewer subjects involved.

Although Mg is considered an epiphenome concerning the cell cycle and proliferation, its potential role in the progression of OSCC remains unfolded. The variation in results between studies can be attributed to inconsistent experimental observations and epidemiologic data regarding different cancers. In the present study, the lower levels of Mg in the control group can be attributed to lower dietary intake. Furthermore, aside from dietary factors, an individual's lifestyle and socioeconomic status impact Mg levels. The conflicting results among various authors can be assigned to the variation in the number of involved subjects, the stage of

<table>
<thead>
<tr>
<th>Group</th>
<th>No. of subjects</th>
<th>Mean</th>
<th>Std. Deviation</th>
<th>Std. Error</th>
<th>95% Confidence Interval for Mean</th>
<th>Lower Bound</th>
<th>Upper Bound</th>
<th>Minimum</th>
<th>Maximum</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>18</td>
<td>0.5778</td>
<td>0.3655</td>
<td>0.08615</td>
<td>0.0548</td>
<td>-0.2890</td>
<td>0.5771</td>
<td>0.2896</td>
<td>0.7595</td>
</tr>
<tr>
<td>OSCC</td>
<td>18</td>
<td>0.3680</td>
<td>0.3655</td>
<td>0.08615</td>
<td>0.0548</td>
<td>-0.2890</td>
<td>0.5771</td>
<td>0.2896</td>
<td>0.7595</td>
</tr>
<tr>
<td>Total</td>
<td>36</td>
<td>0.5056</td>
<td>0.3655</td>
<td>0.08615</td>
<td>0.0548</td>
<td>-0.2890</td>
<td>0.5771</td>
<td>0.2896</td>
<td>0.7595</td>
</tr>
</tbody>
</table>

Table 3. Statistical data obtained using unpaired t-test.

Table 4. Statistical data obtained using One-way ANOVA.
the OSCC of the patient, the method of collection of saliva, and its analysis for Mg levels. In the present study, the sample size was 18 per group, comparatively less than other studies. Thus, the author recommends using a greater sample size and standardizing saliva collection and analysis for salivary Mg.

Conclusion

The present study did not find any significant difference in the mean salivary Mg levels between the OSCC and healthy group in the present study. In conclusion, the author would like to emphasize that the comparison of salivary Mg in patients with OSCC and healthy individuals did not provide sufficient evidence of Mg to serve as an alleged diagnostic tool or a potential biomarker for OSCC. Differences in the cancer cases studied so far, and the techniques used to evaluate the Mg levels can be accounted for in this conflict. Moreover, an individual’s nutritional status also held a potential role in the varying levels of Mg and was left unattended by a majority of the studies. Thus, the author suggests future studies focus on the existing lacunae and evaluate the nutritional socioeconomic status of the individual while estimating both serum and salivary Mg concentrations in an optimal sample size.

Acknowledgments

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Conflict of interest

None.

Financial support

None.

Ethics statement

The Institutional Ethics Committee at Dr. D. Y. Patil Dental College and Hospital, Pune, India, approved the experimental protocol (DYPDCH/IEC/123/129/19) for undertaking this study.

Informed consent was obtained from all individual participants included in the study.

References


