E-Cadherin gene expression in oral cancer: Clinical and prospective data

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Abstract

Background: Low protein expression of E-cadherin in oral squamous cell carcinoma (OSCC) has been associated with clinical and histopathological traits such as metastases, recurrence, low survival and poor tumor differentiation, and it is considered a high-risk marker of malignancy. However, it is still unknown whether low expression of E-cadherin is also present at the mRNA level in OSCC cases. Objective: The aim of this study was to compare E-cadherin mRNA expression in OSCC patients and controls and to correlate the expression with clinical and prospective characteristics.

Material and Methods: Forty patients and 40 controls were enrolled. E-cadherin mRNA expression was evaluated by quantitative real-time polymerase chain reaction using TaqMan probes.

Results: E-cadherin mRNA expression was significantly decreased in OSCC patients compared to that of controls (p<0.001). Whereas no significant association between clinical parameters and E-cadherin expression levels was observed, we noted lower E-cadherin expression levels in patients with positive lymph node metastasis.

Conclusions: E-cadherin mRNA expression was markedly diminished in OSCC, in agreement with previous results that examined E-cadherin expression at the protein level. E-cadherin is downregulated in the early clinical stages of OSCC, and its mRNA levels do not change significantly in the advanced stages, suggesting that there is limited usefulness of this parameter for predicting disease progression.

Key words: Oral cancer, E-Cadherin, gene expression.
Introduction
Global demographic transitions suggest that there will be an increase in cancer burden over the next several decades, particularly in middle- and low-income countries (1). In 2012, GLOBOCAN reported that oral cancer made up approximately 2.1% of newly registered cancer cases. According to this study, approximately 145,000 deaths occurred because of oral cancer 77% of which occurred in less developed regions (2).

Oral squamous cell carcinoma (OSCC) is the most common histological cancer type of the oral cavity (3), and it is considered to have a multifactorial etiology due to the synergy of multiple risk factors (4,5).

In the advanced stages, neoplastic cells undergo a dedifferentiation process, which is followed by a loss of intercellular adherence. Intercellular adhesion is primarily maintained by E-cadherin adhesion molecules, and specifically by E-cadherin in the epithelial tissue (6). E-cadherin is a 120 kDa transmembrane glycoprotein encoded by the CDH1/E-cadherin gene located on chromosome 16q22.1, and it is considered a tumor suppressor gene due to its ability to negatively regulate cell proliferation (7).

E-cadherin downregulation normally occurs during embryogenesis in the context of an epithelial-mesenchymal transition (EMT) (8-10), which allows cells to migrate to other embryonic layers. Similarly, neoplastic cells migrate and reach distant organs in malignant tumors of epithelial origin after undergoing an EMT that involves decreased E-cadherin expression (8).

Accordingly, low expression of E-cadherin in OSCC has been associated with clinical and histopathological features of malignancy, such as metastasis, recurrence, low survival and poor tumor differentiation (11-13), and it has been referred to by some authors as a “high-risk marker of malignancy” (14,15). However, in most studies, E-cadherin expression has been assessed at the protein level using immunohistochemistry (12,15-23).

Cancer is a genomic disease (24), and gene expression at the mRNA level of markers involved in carcinogenesis, such as E-cadherin, needs to be evaluated in OSCC at the mRNA level of markers involved in carcinogenesis, such as E-cadherin, needs to be evaluated in OSCC.

The aim of this study was to quantify the gene expression levels of E-cadherin in oral cancer patients and controls and to evaluate their association with tumor, node, metastasis (TNM) stage, persistence, recurrence, death, and response to therapy.

Material and Methods
A total of 40 cases of primary OSCC were diagnosed over a period of 3 years (2012-2015) (24 males and 16 females, aged 60.8 ± 15.7 years) at the Instituto Jaliscience de Cancerología (Jaliscience Cancer Institute); no anticancer therapy had been initiated in any of these cases. The 40 control samples were obtained from patients (22 male and 18 female subjects aged 58.7 ± 17.5 years) who visited the maxillofacial surgery service at the Hospital Civil Nuevo (New Civil Hospital) for tooth extraction with prosthetic and orthodontic indications.

For each patient, data regarding the following clinico-pathological factors were collected: gender, age, smoker status, alcohol consumption, TNM stage, recurrence, persistence and date of death. The exclusion criteria for both groups were having or having had any type of cancer other than OSCC, being diagnosed with an autoimmune disease that affected the oral cavity, such as pemphigus vulgaris or lichen planus, or not signing the informed consent. Protocol was reviewed and accepted by Universidad de Guadalajara biosecurity and etic committee number CI-02112.

- Clinical and prospective data
  The stage of the disease was evaluated at the Surgical Head and Neck Oncology Department of the Jaliscience Cancer Institute, and each tumor was analyzed clinically and by computerized tomography. The disease stages were then determined according to the TNM system (AJCC) (25).

  Due to the finding that the diameter of the tumor and invasion into lymphatic nodes are regarded as important clinical markers involved in prognosis, the lymphatic nodes were considered positive if patients presented with metastasis in one or more lymphatic nodes. T stage and total TNM scores were assigned in accordance with the National Cancer Institute guidelines, and each case was determined to be early- or advanced-stage disease (NCI, 2016) (26).

  According to this system, advanced T stage refers to tumors that are larger than 4 cm, whereas advanced TNM stage also incorporates tumors of a smaller size that are accompanied by positive lymphatic nodes with or without distant metastasis.

  Prospective data were taken and updated each year after diagnosis and at the beginning of anticancer treatment. The average follow-up was 43.02 ± 13.48 months, with a minimum of 21 months and a maximum of 65 months.

  Persistence was diagnosed in patients who showed tumor activity based on imaging studies after a 9 to 12 month period after the conclusion of therapy, according to the criteria previously described by DeSouza et al. (27).

  Additionally, recurrence of the disease was defined as, despite there being a period of time with eradication of clinical and imaging manifestations of disease, also known as disease-free period or disease-free interval, the reappearance of cancer, whether in the location of the original tumor (local recurrence), in regional lymph nodes or in adjacent tissues (regional recurrence), or if it had disseminated to any other body part (distant recurrence) (28).

  A patient was considered to be a non-responder to anticancer therapy if they experienced recurrence and/or persistence.

- Sample retrieval and storage
  A fresh tissue sample that was at least 5 mm in diameter was obtained at the time of the surgical procedure,
during biopsy in the OSCC group and tooth extraction in the control group. Samples were stored in a -80°C freezer.

-RNA isolation and cDNA synthesis
RNA extraction from tissue samples was performed using TRIzol (Invitrogen®) reagent coupled with Polytron® (Kinematica Inc.) homogenization at room temperature (RT). Afterwards, 200 µl of chloroform was added for every 750 µl of TRIZOL. Samples were shaken vigorously using a vortex for 15 seconds, incubated at room temperature for 5 minutes, and centrifuged at 11000 rpm for 10 minutes at 4°C. The upper aqueous phase was transferred carefully to precipitate the RNA by mixing with 1 volume of isopropanol. Samples were incubated at RT for 15 minutes and centrifuged at 14000 rpm for 10 minutes at 4°C. Next, samples were washed with 80% ethanol in diethyl pyrocarbonate water, and the RNA pellet was air-dried for 5-10 minutes. RNA concentration and quality were measured using a NanoDrop Spectrophotometer (NanoDrop Technologies, Wilmington, DE). cDNA was synthesized from 100 ng of total RNA using a High Capacity cDNA Reverse Transcription Kit (Applied Biosystems) according to the manufacturer’s instructions. cDNA samples were stored at −20°C.

-Relative quantification by Real-Time Polymerase Chain Reaction (qRT-PCR)
Before quantification of the gene of interest, selection of an endogenous gene for normalization was performed by construction of standard curves. In this stage, a total of 100 ng cDNA was taken from 6 OSCC cases and 6 controls to verify that the endogenous gene had no variation between study groups. Five serial dilutions of each cDNA were made to generate a concentration range from 25 ng to 200 ng. The qRT-PCR reactions to evaluate the 18S rRNA, B2M, GAPDH and HPRT1 endogenous genes were performed in a final volume of 10 µL, 1X TaqMan Universal PCR Master mix, 1X TaqMan Gene Expression Assay primer and probe mix (Applied Biosystems). The qPCR reactions were performed on a StepOne Plus Real-Time PCR System (Applied Biosystems, Foster City, CA) according to the manufacturer’s instructions. Standard curves were compared using Student’s t-test. The gene 18S rRNA was selected because it had the lowest significance.

The relative mRNA expression levels of E-cadherin were determined using a TaqMan® Gene Expression Assay (Hs001023894_m1; Applied Biosystems). The TaqMan probe corresponded to the cdh1/E-cadherin gene joining at exons 2 and 3 from the 284 bp position and amplified a fragment of 61 bp. During the quantification of E-cadherin gene expression using the qRT-PCR assay, 20 ng of cDNA from each patient sample was used in duplicate for each PCR (according to the mixing conditions previously described). The Cq values were normalized to those of the 18S-rRNA endogenous gene, and gene expression was calculated as the fold-change relative to the average expression of the controls using the 2^ΔΔCt method.

-Immunohistochemistry (IHC)
To confirm the protein presence in the tumoral tissue, immunohistochemical technique was performed. Tissue samples were processed and evaluated. Three-µm-thick sections were cut, placed on poly-L-lysine-coated slides, deparaffinized in a 60°C oven for 30 minutes and incubated in xylol for 5 minutes. The sections were then rehydrated in decreasing alcohol concentrations (absolute, 90%, 70%, and 50%) and washed in distilled water. To retrieve the epitopes, the tissue sections were incubated in 10 mM sodium citrate solution, pH 9, in a microwave oven at a maximum power of 750 W for two cycles of 5 minutes each. The sections were then cooled to room temperature and washed with distilled water. The endogenous peroxidases were blocked with 0.9% hydrogen peroxide, and the samples were washed again with distilled water and phosphate-buffered saline solution (PBS, pH 7.4). The sections were incubated for 30 minutes with primary antibodies against E-cadherin (Clone NCH-38, 1:50, Dako Corp., Carpinteria, CA, USA). Next, the sections were incubated with a biotinylated anti-mouse/anti-rabbit secondary antibody and a streptavidin/peroxidase complex (LSAB+-labeled streptavidin-biotin, Dako Corp.) for 30 minutes each, and the reaction products were detected with 3,3′-diaminobenzidine-H2O2 (Dako Corp.).

For microarray analysis, negative and positive controls were obtained from paraffin blocks containing samples of healthy mucosa and fibroma using a 5-µm punch and were placed on slides along with the samples. The immunopositivity and immunolocalization were analyzed by an oral pathologist. The E-cadherin membranal protein (mP) was expressed on the cell membrane in all tumor tissues, regardless of the high or low expression status.

-Statistical Analysis
Comparisons were made between OSCC cases and controls by Student’s t-test. To perform Chi-square or Fisher’s exact test, the conversion of categorical variables was necessary; zero was considered to be the cutoff point for under- or overexpression. Comparisons between mRNA E-cadherin expression levels, mP E-cadherin and disease characteristics, such as TNM and TNM clinical stage, and prospective data were calculated by the Mann-Whitney U-test and Chi square. Statistical analysis was performed using SPSS statistics v23 software, and ≤ 0.05 was considered to be statistically significant.

Results
Sociodemographic and risk factor profiles from the
OSCC case and control groups were obtained. Both groups were similar in regard to gender and age distribution as well as smoking status, whereas alcohol consumption was higher in the control group. With the purpose of assessing the influence of risk factors on E-cadherin gene expression, a comparative analysis between E-cadherin mRNA levels and alcohol or tobacco consumption was made in both case and control groups, without showing significant differences (t-student, p-values are shown in Table 1).

The description of the clinical characteristics and prospective data of patients with oral cancer showed that a majority of the patients presented tumors larger than 4 cm, with regional lymph node metastasis and advanced TNM clinical stage (Table 2). However, none of the patients showed dissemination to distant organs. From the prospective data, it should be highlighted that most of the patients responded to the therapy and were still alive at the time that this paper was completed.

The difference in gene expression of E-cadherin between OSCC cases (-4.3 ± 5.6) and controls (-0.3 ± 4.7) was significant (p < 0.001) (Fig. 1A), showing that E-cadherin was downregulated at the mRNA level in OSCC. Neither tumor size, lymph node status nor TNM stage showed a significant correlation with the E-cadherin mRNA level (Fig. 1B,C,D). However, we noted that patients with negative lymph node metastasis had a higher level of E-cadherin mRNA than that of patients with positive lymph node metastasis (-1.8 ± 5.6 vs -5.6 ± 5.3, respectively). Spearman’s rho was calculated with the aim of correlating the levels of mRNA E-cadherin with clinical parameters. Although the correlation was significant for all parameters evaluated, the rho value indicated a positive weak correlation for tumor size (p = .003 rho = .332) and invasion of lymph nodes (p = .019 rho = .268) but, strikingly, the correlation becomes weakly negative for the global clinical stage (p = .001 rho = -.389), where the trend line shows that as the clinical stage increases, the mRNA levels decrease.

Prospective data (recurrence, persistence, response to therapy and death) were also not significantly related to E-cadherin mRNA levels (Fig. 2A-D), although there was a trend for lower values of E-cadherin expression in deceased (-6.8 ± 7.1) vs. living patients (-3.79 ± 5.36). The quantitative expression of E-cadherin was dichotomized in high and low expression groups using a value of 0 as the cutoff and was used to generate survival analysis curves. The average survival in the high expression group was 48.7 ± 3.3 months, and survival was reduced to 44.8 ± 3.3 months in the low expression group, although this difference was not significant. However, it can be seen in the survival curves that deaths occurred earlier in patients with low expression of E-cadherin (Fig. 2E).

E-cadherin at the cell membrane (mP E-cadherin) was marked by immunohistochemistry (Fig. 3A,B). Using Mann Whitney U test, a statistically significant relation between mP E-cadherin and E-cadherin mRNA levels in OSCC patients was observed (p=.03), as tissues with low mP E-cadherin and E-cadherin mRNA levels in OSCC patients was observed (p=.03), as tissues with low mP E-cadherin expression mostly had low levels of E-cadherin mRNA, and conversely tissues with high mP E-cadherin expressed high levels of E-cadherin mRNA (Fig. 3C). As expected, therefore, similarly to E-cadherin mRNA levels, using the Chi-square test we found that mP E-cadherin was not significantly associated with the clinical and prospective parameters T (p=

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**Table 1:** Sociodemographic, risk factor descriptive data and expression mRNA E-cadherin difference from OSCC patients and the control group.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Case</th>
<th>Control</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Age</td>
<td>60.8</td>
<td>15.7</td>
</tr>
<tr>
<td>No.</td>
<td></td>
<td>%</td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>17</td>
<td>42.5</td>
</tr>
<tr>
<td>Male</td>
<td>23</td>
<td>57.5</td>
</tr>
<tr>
<td>Smoker</td>
<td></td>
<td>p value</td>
</tr>
<tr>
<td>No</td>
<td>19</td>
<td>47.5</td>
</tr>
<tr>
<td>Yes</td>
<td>21</td>
<td>52.5</td>
</tr>
<tr>
<td>Alcohol</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>19</td>
<td>47.5</td>
</tr>
<tr>
<td>Yes</td>
<td>21</td>
<td>52.5</td>
</tr>
</tbody>
</table>

Abbreviation: OSCC, oral squamous cell carcinoma SD: Standard deviation. *T-student l.C. 95%.
Table 2: Clinical and prospective data from 40 OSCC patients.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Clinical</th>
<th>Prospective</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No.</td>
<td>%</td>
</tr>
<tr>
<td>T stage</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Advanced</td>
<td>25</td>
<td>62.5</td>
</tr>
<tr>
<td>Not advanced</td>
<td>15</td>
<td>37.5</td>
</tr>
<tr>
<td>Lymph node metastasis</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive</td>
<td>27</td>
<td>67.5</td>
</tr>
<tr>
<td>Negative</td>
<td>13</td>
<td>32.5</td>
</tr>
<tr>
<td>TNM stage</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Advanced TNM stage</td>
<td>26</td>
<td>77.5</td>
</tr>
<tr>
<td>Early TNM stage</td>
<td>14</td>
<td>22.5</td>
</tr>
</tbody>
</table>

Abbreviations: OSCC, oral squamous cell carcinoma. Tumor advanced, Lymph node metastasis and clinical TNM stages were assigned in accordance with the guidelines of the NCI.

Discussion

E-cadherin downregulation at the mRNA level occurs due to several mechanisms and plays a key role in the cell motility that is required for the invasion and metastasis cancer hallmark to occur (29). Despite the crucial role of E-cadherin downregulation in cancer, only two previous studies, to our knowledge, have analyzed the mRNA expression of E-cadherin in OSCC tumors. Wang et al. found that the expression of E-cadherin mRNA in the invasive tumor front was lower than in the central region (18). Their study did not report a comparison of the mRNA expression of E-cadherin with the clinical and prospective data of oral cancer patients. Yao et al. (23) analyzed the mRNA levels of E-cadherin by qRT-PCR using the 2- ΔΔCt method in 20 samples of oral cancer, but they used normal tissue adjacent to the tumor as the control group. In their study, they reported significant differences between tumor tissue adjacent to the tumor as the control group. In their study, they reported significant differences between tumor tissue and adjacent normal tissue, but they did not report associations between E-cadherin mRNA expres-
Fig. 2: Comparison of the E-cadherin gene expression values and prospective data. A) Recurrence, no vs yes: mean -1.81 ± 5.32 vs -4.05 ± 6.38, minimum -13.00 vs -12.00, maximum 6.7 vs 5.4, respectively. B) Persistence, no vs yes: mean -4.19 ±5.30 vs -4.16 ± 7.16, minimum -13.95 vs -12.00, maximum 6.38 vs 5.40. C) Therapy response, no vs yes: mean -4.49 ± 6.53 vs -3.60 ± 5.15, minimum -12.00 vs -13.95, maximum 5.40 vs 6.38, respectively. D) Death, no vs yes: mean -3.79 ± 5.36 vs -6.80 ± 7.18, minimum -15.46 vs -13.95, maximum 6.38 vs 3.83, respectively. E) Kaplan–Meier survival curve for overall survival according to high/low E-cadherin mRNA expression and survival time. Survival with low E-cadherin mRNA expression (orange line) decreases earlier than with high E-cadherin mRNA expression (blue line).

Fig. 3: All tumors showed membranal protein (mP) E-cadherin expression at different intensities. A) High mP expression of E-cadherin by immunohistochemistry in well-differentiated OSCC. B) Low mP expression of E-cadherin by IHC in OSCC (magnification 40x). Mann-Whitney tests. C) Tissue with low mP E-cadherin: mean -5.48 ± 4.8, minimum -12.00, maximum .92. Tissue with high mP E-cadherin: mean .39 ± 5.47, minimum -10.19, maximum 6.38.

SIP1 (31), the microRNA family that includes miR200, include the SNAIL superfamily (Snail 1 and Snail 2) (30), and in vitro and metastatic behavior of squamous cell carcinomas (22,37). These E-cadherin regulatory mechanisms in the neighborhood cancer cells, may carry some of the alterations in genes and signaling networks operating in the neighboring cancer cells, making such tissues an unreliable control.

In this study, E-cadherin mRNA expression was markedly diminished in OSCC tumors, in agreement with previous data on E-cadherin protein levels (11-13). However, low E-cadherin expression at the mRNA level occurred in the early clinical stages and did not continue to decrease significantly in the advanced stages, so its potential usefulness for OSCC staging seems limited. Nevertheless, we observed a trend towards further decreases in E-cadherin mRNA levels with positive lymph node metastasis and a weak negative correlation with TNM clinical stage.

One limitation of this type of studies is, on the one hand that E-cadherin gene expression, although constitutive, can be regulated by causes other than the carcinogenesis process, such as wound repair, inflammatory processes and immune response. On the other hand, there might be variations in the proportion of epithelial or mesenchymal tissue present in the tissue extracted from each sample, as well as variations in the precise tumor area from which the sample is obtained (front of tumoral advance or central tumor area), which constitutes another possible limitation of these studies.

Future studies with larger sample sizes will be needed to ascertain whether the E-cadherin mRNA level could eventually constitute a predictor of lymph node metastasis in OSCC. Such studies would also allow us to confirm our results regarding the non-significant correlations of E-cadherin mRNA levels with prospective variables such as persistence, recurrence and therapy response.

E-cadherin is regulated by several transcriptional factors that have emerged as important transcriptional repressors of E-cadherin specifically in oral squamous cell carcinomas in vitro (33-35), as well as the coexpression of ErbB-2 (36). Likewise, gene expression is also regulated epigenetically, regulating gene expression through changes in the structure of chromatin that results from the transfer of methyl groups.

Several authors have chosen to study concomitantly the expression of E-cadherin with some of the regulatory proteins in situ (22,37); thus, it would be interesting for future studies to include an evaluation of various transcriptional or regulatory factors and the epigenetics involved in the expression of E-cadherin specifically in oral squamous cell carcinomas.

In conclusion, our study confirms the importance of gene expression E-cadherin levels in OSCC and shows that its downregulation at the mRNA level is likely to be an early and constant event in the evolution of this disease. Further studies will be necessary to establish whether E-cadherin levels continue to decrease significantly as OSCC progresses.

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Conflict of interest

The authors have declared that no conflict of interest exist.