

Loss of epidermal growth factor receptor expression in oral squamous cell carcinoma is associated with invasiveness and epithelial-mesenchymal transition

著者	Kimura Iyo, Kitahara Hiroko, Ooi Kazuhiro, Kato Koroku, Noguchi Natuyo, Yoshizawa Kunio, Nakamura Hiroyuki, Kawashiri Shuichi
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# Loss of epidermal growth factor receptor expression in oral squamous cell carcinoma is associated with invasiveness and epithelial-mesenchymal transition

IYO KIMURA, HIROKO KITAHARA, KAZUHIRO OOI, KOROKU KATO, NATUYO NOGUCHI, KUNIO YOSHIZAWA, HIROYUKI NAKAMURA and SHUICHI KAWASHIRI

Department of Oral and Maxillofacial Surgery, Division of Cancer Medicine, Kanazawa University Graduate School of Medical Science, Kanazawa, Ishikawa 920-8640, Japan

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**Abstract.** Inhibition of epidermal growth factor receptor (EGFR) signaling has emerged as a novel therapeutic strategy for the treatment of oral squamous cell carcinoma (OSCC). The EGFR-directed inhibitor cetuximab is currently the only approved targeted therapy for the treatment of OSCC. EGFR status may affect the patient response to cetuximab treatment. In the present study, via analysis of the immunomarker for EGFR, it was revealed that 58.3% of the total cases investigated stained positively for EGFR expression, and furthermore, that invasiveness was inversely correlated with EGFR expression. Expression levels of EGFR were quantified, and the correlation between EGFR expression and cetuximab sensitivity was investigated using three varying grades of invasive human OSCC line. EGFR expression in high-grade invasive cells was significantly downregulated compared with that of low-grade invasive cells. There was no significant antiproliferative effect in the high-grade invasive cells treated with various concentrations of cetuximab. The EMT-associated genes, N-cadherin, vimentin and Snail, were upregulated in the high-grade invasive cells. The low-grade invasive cells exhibited characteristics of typical epithelial cells, including the expression of E-cadherin and absence of the expression of N-cadherin, vimentin and Snail. Transforming growth factor- $\beta$  induced low-grade invasive cells to undergo an epithelial-mesenchymal transition (EMT)-associated gene switch, which resulted in low levels of EGFR expression. The results of the present study suggested that loss of EGFR expression in OSCC was associated with EMT, and may have functional implications with regard to tumor invasiveness and the resistance to cetuximab treatment.

## Introduction

Epidermal growth factor receptor (EGFR/ErbB1/HER1) overexpression is found in the majority of oral squamous cell carcinoma (OSCC) tumors, and associations have been made between increased expression levels and an aggressive phenotype, poor prognosis and resistance to anticancer therapy (1). A number of anti-EGFR antibodies (including cetuximab and panitumumab) and small molecule tyrosine kinase inhibitors (including gefitinib and erlotinib), which target EGFR, have been developed (2). However, only the monoclonal antibody cetuximab (Erbix<sup>®</sup>) is currently approved for the treatment of advanced OSCC (3). Cetuximab therapy has been observed to be effective in the treatment of recurrent and metastatic OSCC, as a first-line treatment in combination with platinum-based chemotherapy (4-7), and as a second-line therapy in patients exhibiting platinum-resistant disease (8-10). In clinical practice, EGFR expression may be evaluated by utilizing a standardized immunohistochemistry (IHC) assay, which is designed to assess cell membrane staining (11). IHC studies have demonstrated EGFR overexpression in >75% of the total analyzed OSCC cases, demonstrating values of 73.42% (12), 87.5% (13) and 72% (14). However, the cetuximab response rate is typically not greater than 20% (15), therefore the role of EGFR expression as a predictor of patient response to EGFR-targeted therapeutic agents remains to be fully elucidated.

The sensitivity of certain tumors to cetuximab may be explained by the presence of mutations in the EGFR tyrosine kinase domain (16,17). However, such mutations are rare in OSCC (18). Thus, there is a requirement to elucidate the mechanisms underlying the differential drug response of cancer cells with wild-type EGFR. This may assist with the identification of patients who may respond and clinically benefit from cetuximab treatment, and furthermore may aid in the development of novel therapeutic strategies, in order to circumvent the *de novo* or acquired resistance of tumors to cetuximab. Tumor resistance mechanisms to cetuximab may also be targetable, and may thus present a potential opportunity to increase the efficacy of cetuximab therapy. In general, tumors that are frequently therapy-resistant may originate from tumor cells exhibiting an epithelial-mesenchymal

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*Correspondence to:* Dr Hiroyuki Nakamura, Department of Oral and Maxillofacial Surgery, Division of Cancer Medicine, Kanazawa University Graduate School of Medical Science, 13-1 Takara-machi, Kanazawa, Ishikawa 920-8640, Japan  
E-mail: hnak@me.com

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transition (EMT) phenotype (19). During EMT, epithelial cells lose their characteristic polarity and cell-cell contact, and acquire a more migratory mesenchymal phenotype (20). EMT has additionally been demonstrated to promote stem cell properties. Furthermore, the *in vitro* sensitivity of head and neck cancer cell lines to anticancer agents has been revealed to be affected by EMT-associated gene expression (21,22).

In the present study, the expression of EGFR in human OSCC tissues and cell lines was evaluated and the association between EGFR expression and clinicopathological features in these tissues and cells was investigated. Furthermore, the OSCC cells and tissues were analyzed with regard to their EMT characteristics, and the OSCC cell lines were also examined for their intrinsic sensitivity to cetuximab treatments.

## Materials and methods

**Tissue samples.** The present study was approved by the Ethics Committee of the Kanazawa University Graduate School of Medical Science (Kanazawa, Japan) and written informed consent was obtained from each patient. The subjects consisted of 24 patients exhibiting primary OSCC who underwent surgical resection of their tumors at the Department of Oral and Maxillofacial Surgery at Kanazawa University Hospital (Kanazawa, Japan), between 1998 and 2008. The patients ranged from 43-89 years old (mean  $\pm$  standard deviation, 64.4 $\pm$ 11.2 years). The tumor-node-metastasis categories were assigned according to the Union for International Cancer Control system (23). The tumor differentiation grade was determined according to the classification criteria proposed by the World Health Organization (24). The mode of tumor invasion was assessed according to the criteria proposed by Yamamoto *et al* (25).

**IHC.** Each specimen was fixed in 10% buffered formalin (Sigma-Aldrich Japan K.K., Tokyo, Japan) and subsequently embedded in paraffin for the preparation of serial sections. Immunohistochemical staining was performed using an EGFR pharmDx™ kit (K1492; Dako Japan K.K., Tokyo, Japan). The specificity of the staining was confirmed using Universal Negative Control for IS-Series Rabbit Primary Antibodies (cat no. IS600; Dako Japan K.K.) rather than primary antibody as a negative control. The images of 3 separate stained regions were captured under a light microscope at x100 magnification and evaluated based on the number of stained tumor cells, according to the method described by Bernardes *et al* (26). EGFR expression was evaluated based on the extent of immunolabeling in the tumor cell membranes, and was classified on a 4-point scale as follows: 0, no labeling or labeling in <10% of tumor cells; 1, weak labeling, homogeneous or patchy, in >10% of tumor cells; 2, moderate labeling, homogeneous or patchy, in >10% of tumor cells; and 3, intense labeling, homogeneous or patchy, in >10% of tumor cells. The initial scores were subsequently grouped into two final groups: Low expression (0 or 1) and high expression (2 or 3).

**Cell lines.** A total of 3 human OSCC cell lines, established from tumor biopsies with varying grades of invasive ability, were used: OSC-20 and OSC-19, low-grade invasive cells,

and HOC313, high-grade invasive cells. OSC-20 is a cell line derived from a 58-year-old female exhibiting tongue cancer that metastasized to the cervical lymph nodes (27). OSC-19 was derived from a 61-year-old male exhibiting tongue cancer that metastasized to the cervical lymph nodes (28). HOC313 was derived from a 51-year-old female exhibiting squamous cell carcinoma (involving the mandibular gingiva and oral floor) that metastasized to the cervical lymph nodes (29). Additionally, normal human dermal fibroblasts (NHDFs; cat. no. PCS-201-010), obtained from the American Type Culture Collection (Manassas, VA, USA), served as a control.

**Proliferation assay and cetuximab sensitivity.** A 3-(4, 5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay was used in order to investigate the ability of cetuximab to inhibit the proliferation of the 3 OSCC lines (OSC-19, OSC-20 and HOC313) *in vitro*. Initially, 2,500 cells/well were cultured in Dulbecco's modified Eagle's medium (DMEM; D7440; Sigma-Aldrich Japan K.K.) supplemented with Hyclone™ 10% fetal bovine serum (FBS; GE Healthcare, Logan, UT, USA), in 96-well tissue culture plates. Following 24 h of culture, the cells were treated with varying concentrations of cetuximab (Erbix™; Merck Serono Co., Ltd., Tokyo, Japan; 0, 0.005 and 0.01  $\mu$ g/ml) diluted in DMEM supplemented with 2% FBS. In order to measure the number of metabolically active cells following 24, 48 and 72 h of incubation, an MTT assay was performed and the results were quantified according to the manufacturer's protocols. The optical density at 450 nm was measured using a microplate reader (iMark Microplate Absorbance Reader; Bio-Rad Laboratories, Inc., Hercules, CA, USA).

**RNA extraction, complementary DNA (cDNA) synthesis and quantitative polymerase chain reaction (qPCR).** The messenger RNA (mRNA) levels of EGFR, E-cadherin, vimentin, Twist and Snail were analyzed using a Rotor-Gene Q 2plex HRM System (9001560; Qiagen GmbH, Hilden, Germany) with FAM/ZEN/IBFQ probes (DNA sequence unopened; Integrated DNA Technologies, Inc., Coralville, IA, USA). Total RNA was extracted using the RNeasy Protect Mini kit (74124; Qiagen GmbH), and cDNA was obtained by utilizing the PrimeScript 1st Strand cDNA Synthesis kit (6110A; Takara Bio, Inc., Otsu, Japan). All reactions and conditions were performed according to the manufacturer's protocols. Ribosomal RNA (18S) was amplified as an internal standard using HEX/ZEN/IBFQ probes (DNA sequence unopened; Integrated DNA Technologies, Inc.). Data were analyzed using the comparative Cq method, which presented the data as a fold-change in expression level relative to a calibrator sample (in the present case, the mean expression of the 3 replicates of the NHDF cell line).

**Enzyme-linked immunosorbent assay (ELISA).** For enzyme-linked immunosorbent assay analysis of EGFR and phosphorylated EGFR, 3 human oral cancer cell lines (OSC-20, OSC-19 and HOC313) were cultured. Following incubation for 24 h, the cells were seeded into a 96-well tissue culture plate (BD Biosciences, Tokyo, Japan) and fixed for 5 min by 4% formaldehyde solution (Sigma-Aldrich Japan K.K.) in phosphate-buffered saline (Sigma-Aldrich Japan K.K.). The expression levels of EGFR and phosphorylated EGFR were quantified using the human EGFR In-Cell ELISA kit (ab126419;

Table I. Correlation between EGFR expression and clinicopathological features of oral squamous cell carcinoma patients.

Clinicopathological feature	EGFR expression		P-value
	High (n=14)	Low (n=10)	
Tumor category			0.991
T1	3	2	
T2	8	6	
T3	1	0	
T4	2	2	
Node category			0.456
Negative	13	7	
Positive	1	3	
Local recurrence			0.071
Negative	10	4	
Positive	4	6	
Differentiation			0.137
Well	10	7	
Moderate	3	0	
Poor	1	3	
Mode of invasion, grade			0.039 <sup>a</sup>
3	7	1	
4C	5	3	
4D	2	6	

<sup>a</sup>P<0.05. EGFR, epidermal growth factor receptor.

Abcam, Cambridge, MA, USA) according to the manufacturer's protocols. The data were presented as a fold-change in expression level relative to the mean expression of the 3 replicates of the NHDF cell line.

**EMT induction.** In order to induce EMT, OSC-20 and OSC-19 cells were seeded at 70% confluence into a 10-cm plastic dish (BD Biosciences) and cultured for 48 and 72 h in DMEM to induce EMT (Sigma-Aldrich Japan K.K.) with 0.5% FBS. Recombinant human transforming growth factor- $\beta$ 1 (240-B; R&D Systems, Inc., Minneapolis, MN, USA) was added at a final concentration of 5 ng/ml.

**Statistical analysis.** Statistical analysis was performed using JMP<sup>®</sup> version 9.0 (SAS Institute, Inc., Cary, NC, USA). The association between EGFR protein expression and clinicopathological features was examined using Pearson's  $\chi^2$  test. Data from the ELISA and MTT assays are presented as the mean  $\pm$  standard error. Differences between groups were tested for statistical significance using the two-tailed Mann-Whitney U test. P<0.05 was considered to indicate a statistically significant difference.

## Results

**EGFR expression is associated with certain clinicopathological features.** EGFR expression was detected in the cell membrane and cytoplasm of the tumor cells (Fig. 1). According to the criteria for EGFR immunohistochemical evaluation (26), 58.3%

(14/24) of the total cases investigated were positive for EGFR expression. Table I illustrates the correlations between EGFR expression and certain clinicopathological features. EGFR expression in high-grade invasive OSCC (mode of invasion, grade 4D) was significantly downregulated compared with that of low-grade invasive OSCC (mode of invasion, grades 3 and 4C; P=0.039). By contrast, EGFR expression did not correlate with tumor size (P=0.991), local recurrence (P=0.071) histological differentiation (P=0.137) or lymph node metastasis (P=0.456).

**EGFR is differentially expressed in the 3 different grades of invasive OSCC cell lines.** As demonstrated by immunohistochemical analysis, only invasiveness (mode of invasion) was inversely correlated with the expression of EGFR (Table I). Therefore, EGFR expression in the 3 different grades of invasive human OSCC lines (OSC-19, OSC-20 and HOC313) was analyzed at the mRNA and protein level using qPCR and ELISA, respectively. Data are presented as a fold-change relative to the mean of 3 replicates of the NHDF cell line (Fig. 2). Among the three OSCC cell lines (OSC-19, OSC-20 and HOC313), the EGFR mRNA expression level ranged from 0.7- to 3.7-fold higher compared with that of the NHDFs (Fig. 2A). The expression of EGFR mRNA in the 2 low-grade invasive cell lines (OSC-20 and OSC-19) was 3.7- and 2.3-fold higher compared with the NHDFs, respectively. The EGFR mRNA expression in the high-grade invasive cells (HOC313) was markedly reduced and similar to that of the NHDFs. By contrast, the 2 low-grade invasive cell lines (OSC-20 and OSC-19) possessed EGFR



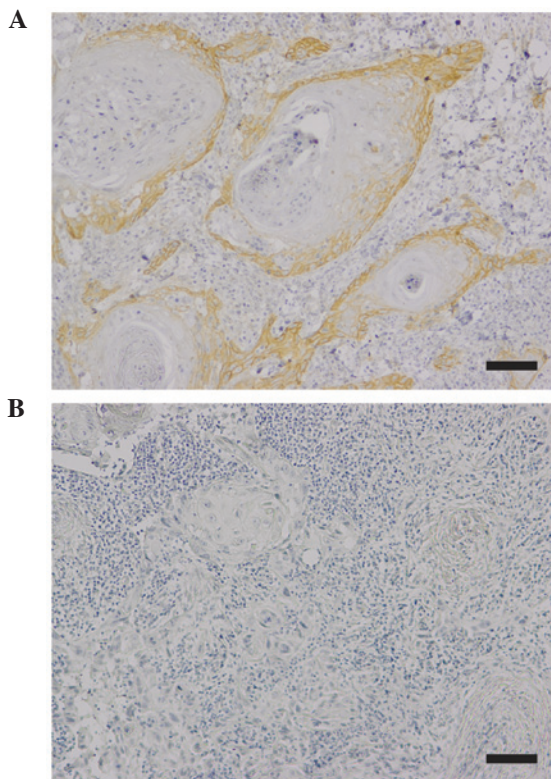


Figure 1. Immunohistochemical staining of EGFR in human OSCC tissues. (A) Positive immunoperoxidase staining for EGFR in OSCC. (B) Specificity of the staining was confirmed by using non-immune antibody instead of primary antibody as a negative control. Scale bar, 100  $\mu$ m. EGFR, epidermal growth factor receptor; OSCC, oral squamous cell carcinoma.

protein expression levels 3.5- and 2.1-fold higher compared with those of the NHDFs, respectively (Fig. 2B). The high-grade invasive cells (HOC313) exhibited the lowest levels of EGFR protein expression. The quantities of phosphorylated EGFR were assessed in order to determine the activation status of EGFR. Expression levels of phosphorylated EGFR ranged from 2.2- to 0.87-fold higher compared with those of the NHDFs (Fig. 2C). Expression patterns of phosphorylated EGFR were similar to the expression patterns of EGFR mRNA and protein.

#### *Cetuximab sensitivity is dependent on OSCC invasiveness.*

In order to evaluate the significance of EGFR status in relation to cetuximab sensitivity, the cetuximab sensitivity of 3 human OSCC lines exhibiting varying grades of invasive ability (OSC-19, OSC-20 and HOC313) was evaluated. Cells were seeded at clonal density, and the antiproliferative effects of cetuximab were determined using an MTT assay (Fig. 3). Antiproliferative effects of cetuximab were observed in the OSC-20 and OSC-19 cell lines. In the OSC-20 and OSC-19 cells (low-grade invasive OSCC), following incubation with cetuximab (0.005  $\mu$ g/ml and 0.01  $\mu$ g/ml) for 48 and 72 h, proliferation was significantly inhibited compared with non-treated cells ( $P < 0.05$ ). However, the varying concentrations of cetuximab exerted no significant antiproliferative effects on the HOC313 cell line ( $P > 0.05$ ).

*An EMT phenotype is detectable in the high-grade invasive OSCC cell line.* In order to evaluate whether EGFR expression was associated with the features of EMT, levels

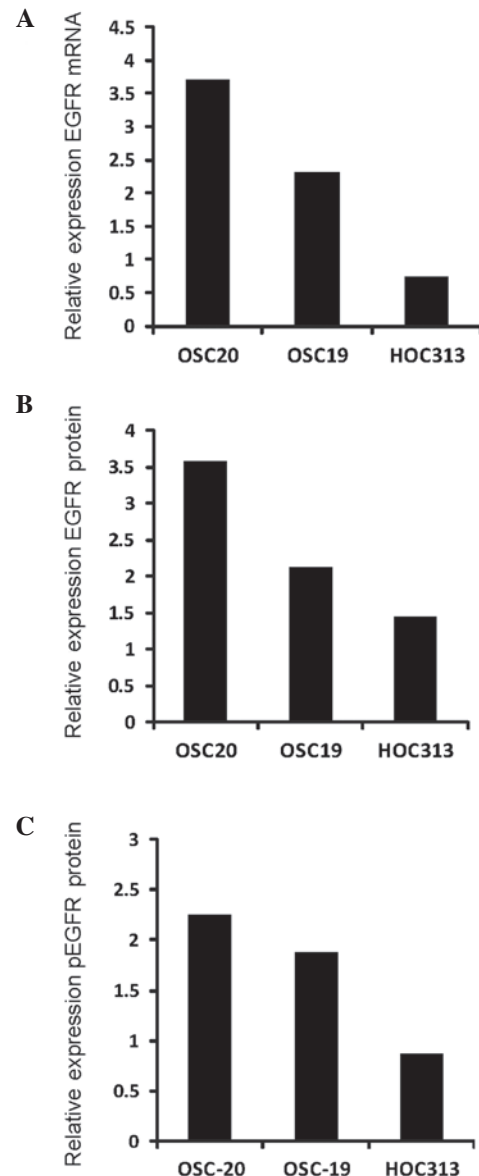


Figure 2. EGFR expression in 3 varying grades of invasive human oral squamous cell carcinoma line. The OSC-19, OSC-20 and HOC313 cell lines were evaluated for variations in (A) EGFR mRNA, (B) EGFR protein and (C) pEGFR expression levels. Data are presented as a fold difference in expression level relative to the mean expression of the normal human dermal fibroblast cells. EGFR, epidermal growth factor receptor; p, phosphorylated; mRNA, messenger RNA.

of N-cadherin, E-cadherin, vimentin and Snail were investigated in 3 different invasive human OSCC cell lines (OSC-19, OSC-20 and HOC313; Fig. 4A). The expression levels of EMT-associated genes in the OSC-19 and HOC313 cells are presented as fold-changes relative to the OSC-20 cells. The HOC313 cells (high-grade invasive cells with loss of EGFR expression) exhibited a mesenchymal phenotype manifested through loss of E-cadherin and acquisition of N-cadherin, vimentin and Snail expression compared with the OSC-20 and OSC-19 cell lines (Fig. 4A). In order to examine whether TGF- $\beta$  altered the expression of the EMT-associated genes and EGFR, the 2 low-grade invasive cell lines (OSC-20 and OSC-19) were exposed to TGF- $\beta$  (Fig. 4B). Expression levels are presented as fold differences relative to the control vehicle at 48 and 72 h, respectively. Serial examination of EMT

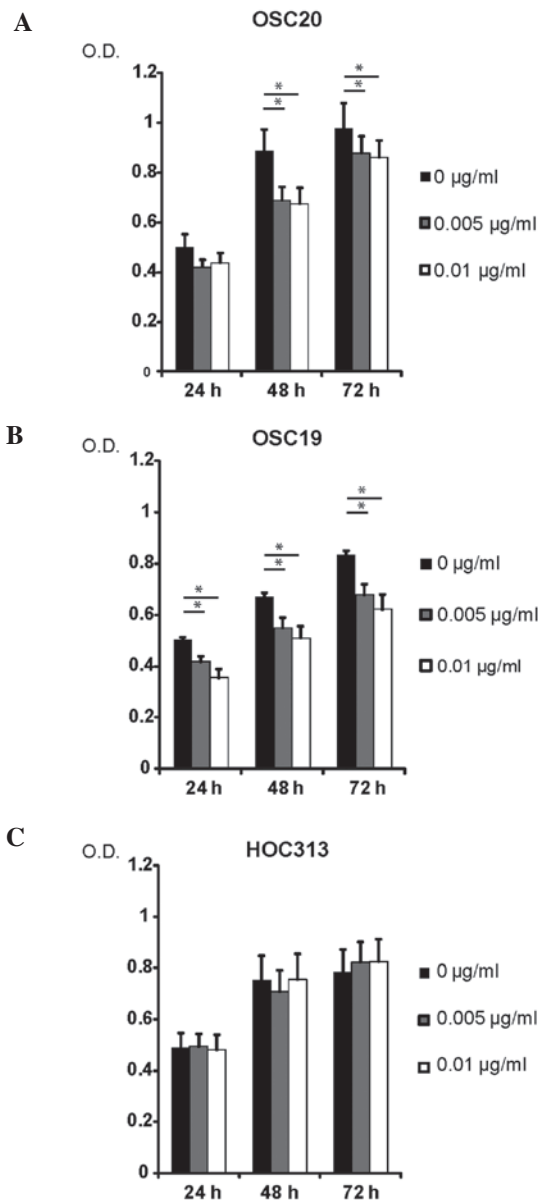


Figure 3. Cetuximab sensitivity of 3 varying grades of invasive human oral squamous cell carcinoma line. (A) OSC-20, (B) OSC-19 and (C) HOC313 cells were treated with various concentrations of cetuximab as indicated, and the antiproliferative effects of cetuximab treatment were determined by 3-(4, 5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide assay. Each data point represents the mean of three independent experiments. Data are presented as standard deviations. Error bars correspond to the standard error. \*P<0.05. O.D., optical density.

markers (loss of E-cadherin and upregulation of N-cadherin, vimentin and Snail) over a specific time course (48 and 72 h) revealed that TGF-β treatment induced EMT in the OSC-20 cells (Fig. 4B). By contrast, TGF-β was not effective in the induction of EMT in the OSC-19 cells. Notably, total EGFR mRNA in the OSC-20 and OSC-19 cells was reduced following EMT induction by TGF-β treatment.

**Discussion**

EGFR (a tyrosine kinase receptor from the ErbB family) is an oncogene that has been identified in a number of malignancies, including cancer of the breast, prostate, pulmonary system,

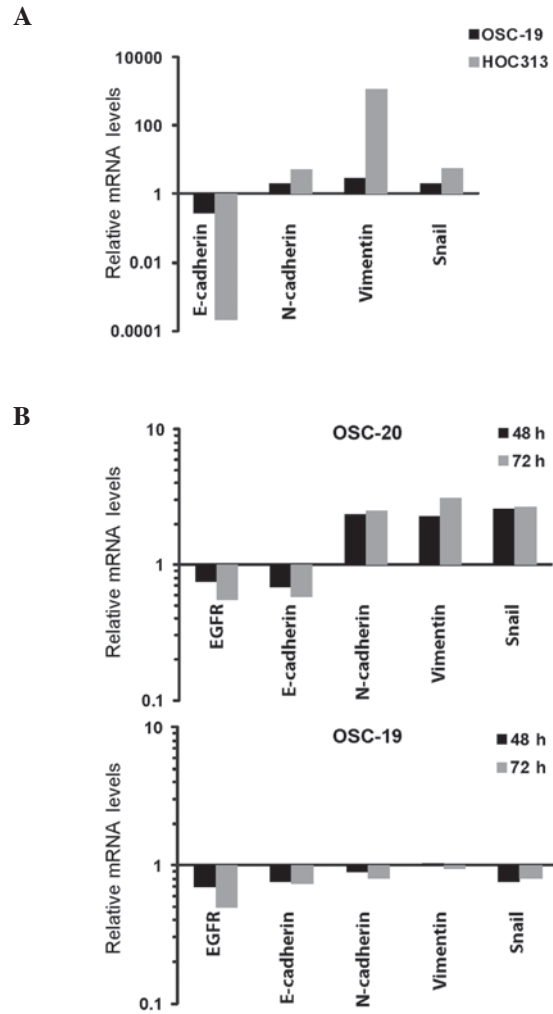


Figure 4. EGFR and EMT-associated gene expression in 3 varying grades of invasive human OSCC line. (A) Relative mRNA expression levels of EMT-associated genes, E-cadherin, N-cadherin, vimentin and Snail, in OSC-20, OSC-19 and HOC313 cell lines. The high-grade invasive OSCC cell line (HOC313) demonstrated upregulation of EMT-associated genes. Expression levels in OSC-19 and HOC313 are displayed as fold-changes relative to the expression levels in OSC-20. (B) Relative mRNA expression levels of EGFR and EMT-associated genes, E-cadherin, N-cadherin, vimentin and Snail, in OSC-20 and OSC-19 cell lines. TGF-β induced EMT-associated gene expression in low-grade invasive OSCC cell lines (OSC-20 and OSC-19). The cells were treated with human TGF-β or a control vehicle for 48 and 72 h. Expression levels are displayed as fold-changes relative to the control vehicle at 48 and 72 h, respectively. EGFR, epidermal growth factor receptor; EMT, epithelial-mesenchymal transition; OSCC, oral squamous cell carcinoma; mRNA, messenger RNA; TGF-β, transforming growth factor β.

bladder, and head and neck (30). EGFR is overexpressed and appears to be the dominant controlling factor underlying the malignant phenotype in head and neck squamous cell carcinomas, via adjustment of the molecules involved in invasive angiogenic and lymphangiogenic processes (12). By analyzing the immunomarker for EGFR in the present study, positive staining for EGFR expression was identified in 58.3% of the total cases analyzed. The results of the present study were consistent with previously reported data (12-14). Furthermore, to the best of our knowledge, the present study demonstrated for the first time that invasiveness (mode of invasion) was inversely correlated with EGFR expression in OSCC patients. The present study also revealed that there

is a loss of EGFR expression in the high-grade invasive OSCC line (HOC313 cells). The results of the present study suggested that loss of EGFR expression was associated with acquirement of an invasive phenotype in OSCC. This potential association will require further investigation in future studies.

The present study analyzed the correlation between EGFR expression and the sensitivity to cetuximab using 3 human OSCC cell lines (OSC-19, OSC-20 and HOC313), possessing varying grades of invasive abilities. Variability in the responses of certain cell lines to cetuximab treatment was identified. Cetuximab was observed to suppress cell growth by ~23% in the most treatment-sensitive cell line (OSC-20), however, it exerted no effect on the HOC313 cell line. The OSC-20 cell line demonstrated a marked increase in EGFR mRNA and protein levels compared with the HOC313 cell line. However, the HOC313 cell line, which was unaffected by cetuximab treatment, demonstrated the lowest levels of EGFR expression. When evaluating the affect of EGFR expression on the cetuximab treatment responses, a significant difference in EGFR expression levels was observed between cetuximab-sensitive (OSC-20) and -resistant cells (HOC313). Cetuximab-resistant cell lines exhibited reduced levels of EGFR expression, indicating that the cetuximab treatment response may be dependent on EGFR expression status. In addition, increased levels of phosphorylated EGFR correlated with cetuximab sensitivity, revealing a potential role of activated EGFR in patient responses to cetuximab. A previous study has suggested that cells may be sensitive to the inhibition of EGFR only if they are dependent on EGFR activation for cell survival and growth (31). In agreement with these previous results, the levels of phosphorylated EGFR have been utilized to predict gefitinib (an EGFR tyrosine kinase inhibitor) sensitivity in OSCC patients (32).

The response to EGFR-targeted agents is inversely correlated with EMT in multiple tumor types without known EGFR mutations, including non-small cell lung cancer, and carcinoma of the bladder, colorectal region, pancreas and breast (33-37). In the present study, the EMT-associated genes, N-cadherin, vimentin and Snail, were upregulated in cetuximab-resistant OSCC tumor cells (HOC313). The present study additionally demonstrated that TGF- $\beta$  was able to induce a low-grade invasive OSCC cell line (OSC-20) to undergo an EMT-associated gene switch, which resulted in a loss of EGFR expression. The results of the present study indicated that the net effect of TGF- $\beta$  signaling was the loss of EGFR activity, accompanied by a concomitant EMT-associated gene switch of tumor cells, as well as acquirement of a migratory mesenchymal phenotype. Further studies are required in order to elucidate the precise molecular mechanisms underlying the association between EMT and the regulation of EGFR expression.

In the present study, it was demonstrated that EGFR status affected the response to cetuximab treatments, therefore, the level of invasiveness or presence of an EMT phenotype may potentially assist in the prediction of EGFR status. EMT-associated alterations in gene expression, including the upregulation of N-cadherin, vimentin and Snail, have additionally been proposed as potential biomarkers for the indication of EGFR status and the prediction of patient

response to cetuximab treatment. However, further *in vivo* studies using clinical samples are required in order to confirm these results.

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