PROX1 Gene is Differentially Expressed in Oral Cancer and Reduces Cellular Proliferation

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Abstract: Homeobox genes are a family of transcription factors that play a pivotal role in embryogenesis. Prospero homeobox 1 (PROX1) has been shown to function as a tumor suppressor gene or oncogene in various types of cancer, including oral squamous cell carcinoma (OSCC). We have previously identified PROX1 as a downregulated gene in OSCC. The aim of this study is to clarify the underlying mechanism by which PROX1 regulates tumorigenicity of OSCC cells. PROX1 mRNA and protein expression levels were first investigated in 40 samples of OSCC and in nontumor margins. Methylation and amplification analysis was also performed to assess the epigenetic and genetic mechanisms involved in controlling PROX1 expression. OSCC cell line SCC9 was also transfected to stably express the PROX1 gene. Next, SCC9-PROX1-overexpressing cells and controls were subjected to proliferation, differentiation, apoptosis, migration, and invasion assays in vitro. OSCC samples showed reduced PROX1 expression levels compared with nontumor margins. PROX1 amplification was associated with better overall survival. PROX1 overexpression reduces cell proliferation and downregulates cyclin D1. PROX1-overexpressing cells also exhibited reduced CK18 and CK19 expression and downregulation of cyclin D1. Our results suggest that PROX1 functions as a tumor suppressor gene in oral carcinogenesis.

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INTRODUCTION

Oral squamous cell carcinoma (OSCC) is the most common malignancy of the oral cavity and a major cause of cancer morbidity and mortality worldwide. Oral carcinogenesis is a multifactorial process associated with cumulative genetic mutations that alter proto-oncogenes and tumor suppressor gene function, resulting in disturbed cellular proliferation and cell differentiation.

Homeobox genes encode transcriptional factors that control cellular proliferation and differentiation during embryonic development. These genes have been aberrantly expressed in solid tumors, including OSCC. The prospero homeobox 1 (PROX1) gene encodes a nuclear transcription factor (prospero homeobox protein 1/PROX1) that plays a major role during embryonic lymphangiogenesis, differentiation of the central nervous system, lens fiber elongation, and hepatocyte proliferation. PROX1 gene inactivation results in abnormal cell cycle proliferation, probably because of downregulation of cell cycle inhibitors.

In human cancers, PROX1 gene acts in a tissue-dependent manner, as a transcriptional activator or repressor, leading to variable effects on cellular proliferation and differentiation. PROX1 overexpression promotes aggressive behavior of many endothelial tumors, colon cancer, and gliomas. However, in hepatocellular carcinoma, high PROX1 expression inhibits transforming activity and cellular proliferation and is associated with well-differentiated tumors and better prognosis. Hagiwara et al also found PROX1 overexpression to suppress cell growth and tumor formation in HeLa cells, partially mediated by protein kinase C β. Additionally, it was also demonstrated that PROX1 strongly inhibits the proliferation of neuroblastoma cell lines as well as cyclin D1, cyclin-A, and cyclin B1, consistent with a role in cell cycle arrest.

In contrast, loss of PROX1 function has been detected in hematologic malignancies, sporadic breast cancer, and carcinomas of the biliary system. Mutations and DNA methylation appear to be the major causes behind loss of PROX1 function in some tumors. Recently, an antimitastatic role of PROX1 was observed in PROX1-silenced hepatocarcinoma cell lines via TWIST1 gene inhibition. In OSCC, Sasahira et al demonstrated that PROX1 and FOXC2 act as oncogenes by inducing lymphangiogenesis and angiogenesis. Additionally, PROX1 was associated with tumor progression (pT and
clinical stage), nodal metastasis, and lymphovessel density. These studies suggest that PROX1 may function as an oncogene or a tumor suppressor gene in a cancer type-specific manner. Interestingly, a previous microarray study done by our group revealed that PROX1 transcripts were downregulated in OSCC when compared with tumor-free margins. However, the underlying mechanism by which PROX1 acts in oral cancer is still unclear. In this study, we analyzed the expression levels of PROX1 transcripts and proteins as well as PROX1 amplification and methylation status in OSCC tissues and tumor-free surgical margins. We also investigated how PROX1 affects cell proliferation, differentiation, survival, migration, and invasion in a squamous cell carcinoma cell line.

**METHODS**

**Tumor Samples**

Specimens were obtained by surgical resection from OSCC patients (men, ≥40 years old) admitted for diagnosis and treatment at the Arnaldo Vieira de Carvalho Cancer Institute, Heliópolis Hospital, and Hospital das Clínicas (School of Medicine, University of São Paulo, Brazil). Histopathological diagnosis was performed according to World Health Organization classification for tumors. Clinopathological staging was determined by the TNM classification of the International Union Against Cancer. All patients have provided written informed consent to participate in this study that was approved by the Brazilian National Ethics Committee (Process #16491) and meets the Declaration of Helsinki.

Forty fresh surgical samples of primary OSCC and their corresponding nonneoplastic margin tissues were immediately snap-frozen in liquid nitrogen. After histological confirmation, all tissue samples were checked prior to RNA extraction so each OSCC sample contained at least 70% tumor cells. The corresponding surgical margins were reported as ‘tumor-free.’ The GENCAPO (Head and Neck Genome Project) consortium was responsible for collecting samples and initial processing, collecting clinical data, performing histopathological analysis, and obtaining informed consent from each patient.

**Cell Culture**

The human OSCC cell lines SCC4, SCC9, and SCC25 were obtained from the American Type Culture Collection (ATCC, Manassas, VA) and cultured (DMEM) and Han F12 medium (DMEM/F12; Invitrogen, Carlsbad, CA) supplemented with 10% fetal bovine serum (FBS), 400 ng/mL hydrocortisone (Sigma-Aldrich, St. Louis, MO), 100 μg/mL penicillin, and 100 μg/mL streptomycin. Cell lines were authenticated using the AmpFLSTR Identifiler PCR Amplification kit (Life Technologies) at the Special Techniques Laboratory of the Israelita Albert Einstein Hospital (LATE-HIAE), São Paulo, Brazil. Cell line matched (100% identity) the STR profile described by the ATCC database.

**Plasmid DNA Transfection for PROX1 Expression**

SCC9 cells were transfected with pCMV6 empty vector (mock) or pCMV6-PROX1 expression vector (OriGene, Rockville, MD), containing full-length human PROX1 cDNA, using Lipofectamine 2000 (Invitrogen), following the manufacturer’s instructions. Transfected cells were selected in medium with 300 μg/mL G418 (Invitrogen).

**Real-Time RT-PCR**

Total RNA was extracted from cells using TRizol (Invitrogen, Carlsbad, CA) according to the manufacturer’s instructions. First-strand cDNAs were synthesized with a High Capacity cDNA Archive kit (Applied Biosystems, Grand Island, NY, USA). Quantitative RT-PCR (RT-qPCR) reactions were performed using an Applied Biosystems 7500 Real-Time PCR System with SYBR Green I Dye (Applied Biosystems) according to the manufacturer’s instructions. The following primers were used to detect PROX1 and endogenous “housekeeping” gene HPRT: PROX1 (forward 5'-CTCCGTGGAACTCAGGCCCC-3' and reverse 5'-GCCGGCTTAAAGGCGGTGCTG-3') and HPRT (forward 5'-CCACCACCTGTGGCTGTA-3' and reverse 5'-TCCCCTGTGTA CTGGTCA-3').

The following primers were used to investigate genes identified as deregulated in oral cancer (data not shown): GATA3 (forward 5'-GTCCCTGGCGAAGCTCA-3' and reverse 5'-GCTCCCATGCTTCGCTC-3'), NOTCH1 (forward 5'-GGTTAGACTGTCGCTCTAGATC-3' and reverse 5'-GGATGTCAGTCGCTCAAGCGTGA-3'), E2F1 (forward 5'-CATCCAGGTGATCTTCCTC-3' and reverse 5'-GACACA GCGGTTCCTTGCTC-3'), and WISP3 (forward 5'-ACTGTAGGCTGGAACCATTACT-3' and reverse 5'-TGTCACCTGTGTT AGATATCCC-3').

All RT-qPCR reactions were performed in total volume of 25 μL, containing 1 μL of cDNA sample, 400 nM of each primer, 12.5 μL of SYBR Green Master Mix (Applied Biosystems). The PCR amplifications were performed under the following conditions: initial denaturation for 10 minutes at 95°C, followed by 40 cycles of 10 seconds at 95°C and 1 minute at 60°C. Melting curve analysis was performed at the end of each cycle to confirm amplification specificity and lack of primer dimer. Quantitative analysis was performed according to the Pfaffl mathematical model.

The differential expression of PROX1 in OSCC samples was divided into high and low according to the value obtained from RT-qPCR normalized with the respective tumor-free margins. The cut-off value was set up at the median expression level of OSCC samples.

**Gene Amplification Assay**

Gene amplification analysis of the OSCC tissue and tumor-free margins was performed by qPCR using an Applied Biosystems 7500 Real-Time PCR System with SYBR Green I Dye. The following primers were used to detect PROX1 (F: 5'-AGCCTCCTGGGAACCTAGCCAG-3' and R: 5'-CCACCACGAGGAAAGAAAAA-3') and ZNF80 (F: 5'-CTGAGGACCGTGCAGGCTCATCCTCCT-3' and R: 5'-TAAATGCTCTGACAGGTGA CTGAGGTCG-3'). ZNF80 was used as a reference gene to normalize the qPCR data. Next, both primer sets generated were validated in silico. Quantitative analysis was performed according to the Pfaffl mathematical model. Briefly, we calculated the ΔCq value for the difference between the unknown and calibrator samples. The relative quantity was calculated as RQ = 2-ΔCq and the normalized relative quantity (NRQ) as NRQ = RQPROX1/RQZNF80. Further, we calculated relative copy number (RCN) as RCN = NRQ/RF, where RF is rescaling factor, which is the geometric mean of the NRQ values of a set of 17 normal oral mucosa bearing 2 PROX1 copies. RCN values close to 1 indicate the presence of 2 PROX1 copies and RCN values > 1 indicate PROX1 positive amplification.
**Methylation Assay**

Methylation status of PROX1 in OSCC tissue and tumor-free margins was evaluated by the EpiTect Methyl Custom qPCR Array technology according to the manufacturer’s instructions (Qiagen, Valencia, CA). Thirty fresh OSCC samples and 17 corresponding nonneoplastic margins were treated with a simple DNA methylation-sensitive and methylation-dependent restriction enzyme digestion without bisulfite conversion. After digestion, the remaining DNA was quantified using DNA methylation real-time PCR arrays in an ABI 7500 Real-time system. Methylation status was expressed as percentage of inputted DNA that was methylated, as determined using EpiTect Methyl DNA methylation PCR data analysis (Qiagen, http://www.sabiosciences.com/dna_methylation_data_analysis.php), following the manufacturer’s instructions. The methylated DNA status in the samples was divided into 2 groups based on the following cut-off values: >50% (hypermethylated) or <50% (hypomethylated).

**Immunohistochemistry**

Immunohistochemical analysis of PROX1 protein expression was performed using the polymer-linked method in 30 OSCC samples and 8 nonneoplastic tissues. Four-μm-thick tissue sections were cut, deparaffinized, and subjected to antigen recovery treatment with 1 mM EDTA buffer target retrieval solution, pH 8.0 at 95°C, in a water bath for 20 minutes. Endogenous peroxidase activity was blocked by incubating with phosphate-buffered saline (PBS) and 3% hydrogen peroxidase for 30 minutes. After washing with PBS, the sections were treated with protein block (Dako, X0909, Carpenteria, CA) and then incubated with monoclonal rabbit anti-PROX1 (Abcam, clone 5G10, Cambridge, UK) diluted 1:50, followed by the Envision Dual Link System horseradish peroxidase (HRP) method (Dako, K4061). Reactions were initiated by incubating the sections with 3,3′-diaminobenzidine tetrahydrochloride (Dako, K3468). The negative controls were obtained by substituting the primary antibody with nonimmune serum. To quantify PROX1 protein expression in OSCC samples and nonneoplastic margins, immunohistochemistry (IHC) scores were used as described by Yama-toji et al. with minor modifications. Next, the percentage of PROX1 positive epithelial cells was evaluated at x100 magnification throughout each entire section and scored as follows: 1, 0% to 25%; 2, 25% to 50%; 3, 25% to 75%; and 4, 75% to 100%. The percentage of PROX1 positive tumor cells and staining intensity were then multiplied to obtain each PROX1 IHC score. We considered these samples positive when PROX1 IHC score was >7.37 (average score for nonneoplastic margins).

**Western Blotting**

Cells were lysed in RIPA buffer (50 mM Tris–HCl pH 7.4, 150 mM NaCl, 1 mM EDTA, 1% NP-40, 1% deoxycholic acid, 0.5% sodium dodecyl sulfate, 1 mM phenylmethylsulfonyl fluoride, 1 mM N-ethylmaleimide, 1 mM dithiothreitol, 10 μg/mL soybean trypsin inhibitor, 1 μg/mL leupeptin, and 1 μg/mL aprotinin). We revealed 30 μg of total protein per sample using 8% sodium dodecyl sulfate (SDS) polyacrylamide gel electrophoresis. The primary antibody to detect PROX1-1 (Abcam, clone 5G10), p21 (Abcam, ab7960), cyclin D1 (Santa Cruz, sc-20044, Dallas, Texas, USA) and actin (Sigma, clone CA-74) were incubated overnight at 4°C at the dilution 1:300. The nitrocellulose membranes were developed using a chemilumi-nescent Western blot system (Enhanced Chemiluminescent Western blot kit; GE Healthcare, Vienna, Austria).

**Bromodeoxyuridine Labeling Index**

Cells were plated in chamber slides (30,000 cells/well) and serum starved for 48 hours. Next, cell culture medium supplemented with 10% FBS was added for 24 hours followed by bromodeoxyuridine reagent incubation for 2 hours at 37°C. Bromodeoxyuridine (BrdU) incorporation in proliferating cells was estimated using an immunohistochemical analysis kit (Invitrogen, BrdU Staining Kit). The BrdU-labeling index, expressed as the percentage of cells labeled with BrdU, was determined by counting 1000 cells in 3 independent reactions using the Kontron 400 image analysis system (Zeiss Axiom Imager A1, Dublin, California, USA).

**Ki-67 Index**

Cells were cultured as described above and fixed after incubation with culture medium supplemented with 10% FBS for 24 hours. Cells were incubated with monoclonal antibodies against Ki-67 (Clone MIB-5, Dako), followed by the Envision Dual Link System HRP (Dako). The Ki-67 index was calculated using an image analysis system by counting labeled nuclei of 1000 cells in 3 independent reactions, expressed as percentage of Ki-67-positive cells.

**Proliferation Curves**

Cells were seeded in 24-well plates in DMEM/F12 containing 10% FBS and incubated in serum-free medium for 48 hours. After this period, DMEM/F12 10% FBS was added back and cells from triplicate wells were trypsinized after 24, 48, 72, 96, 120, and 144 hours and counted in the Neubauer chamber.

**Flow Cytometry**

For cell cycle analysis, SCC9 cells were seeded in 100-mm dishes and serum starved for 48 hours. Next, the cell culture medium supplemented with 10% FBS was added and the cells were collected after 24 hours. Cells were fixed in cold 70% ethanol, stored at −20°C, washed in cold PBS, and treated with 10 mg/mL of RNase for 1 hour at 37°C. After staining with 50 μg/mL of propidium iodide for 2 hours at 4°C, cell distribution in the cell cycle was analyzed by flow cytometry.

Cells were analyzed on a FACSCalibur flow cytometer equipped with an argon laser (Becton-Dickinson, San Jose, CA). At least 10,000 events were analyzed in each sample. Quantitative flow cytometric analysis was performed using CellQuest software (Becton-Dickinson).

**Immunocytochemistry**

Immunostaining was performed on cytokeratins (CK) 1, 10, 13, 14, 16, 18, and 19 using the Envision Dual Link System HRP (Dako, K4061). Cells were incubated with mouse anti-CK10 (1:100, DEK-10, BioGenex, Fremont, California, USA), anti-CK16 (1:300, ab8741, Abcam), anti-CK18 (1:1500, DC10, Dako), anti-CK19 (1:2000, RCK108, Dako), rabbit anti-CK1 (1:200, ab24643, Abcam), anti-CK13 (1:100, ab92551, Abcam), and anti-CK14 (1:2000, ab7800, Abcam) followed by the polymer-link method. The slides were scanned into high-resolution images using an Aperio ScanScope XT (Aperio Technology, Inc, Vista, CA). The images were then viewed in an Image Scope (Aperio Technology, Inc) and immunostaining was quantified using the Positive Pixel Count algorithm, v9 (Aperio).
Apoptosis Analysis

Apoptosis index was determined by annexin V-FITC labeling. The cells were collected, washed with PBS, and resuspended in the binding buffer (10 mM HEPES pH 7.4, 150 mM NaCl, 5 mM KCl, 1 mM MgCl₂, and 1.8 mM CaCl₂) containing annexin V-FITC at 1:500. After 20 minutes incubation in the dark at room temperature, the cells were also stained with propidium iodide (PI, Sigma-Aldrich). Apoptosis was analyzed on a FACSCalibur flow cytometer equipped with an argon laser (Becton Dickinson) and quantified as the number of annexin V-FITC positive and PI negative cells divided by the total number of cells. At least 10,000 events were analyzed in each sample.

Migration and Invasion Assays

The in vitro cell migration and invasion assays were performed in 24-well plates (Corning, Inc, New York, NY) using modified Boyden chamber inserts with a polycarbonate filter membrane containing 8-μm pores. For the invasion assay, the BD BioCoat Matrigel Invasion Chamber was used in 24-well plates (BD Biosciences). The cells (1 × 10⁵) were suspended in 200 μL of serum-free DMEM/F12 and seeded onto the upper compartment of the transwell chamber and DMEM/F12 containing 10% FBS was used in the lower chamber for stimulation. After 24 hours of incubation for the migration and invasion analysis, the medium in the upper chamber was removed, and the filters were fixed in 4% paraformaldehyde following 20 minutes in methanol 100%. The cells on the lower surface were stained with toluidine blue solution in 1% borax for 15 minutes. A cotton swab mechanically removed cells that did not migrate through the pores. The die was eluted using 1% SDS and the absorbance was measured at 560 nm. Two independent experiments were performed with triplicates.

Statistical Analysis

To analyze PROX1 mRNA expression levels in tissue samples and OSCC cell lines, we used Wilcoxon nonparametric test and one-way analysis of variance (ANOVA) with Tukey post-test, respectively. The different PROX1 expression levels in tissue samples were divided into 2 groups (high vs low) and the median expression level was used as the cut-off value. We used one-way ANOVA with Tukey post-test to analyze proliferation of PROX1-overexpressing SCC9 and control cells, CK immunoeexpression, and apoptosis index. We used Fisher exact test to estimate the statistical difference between PROX1 gene expression levels and clinicopathological parameters such as mean age, tumor location, tumor size-pT, nodal metastasis-pN, pathological grade, lymphatic and/or perineural invasion, and recurrence. When clinical information was not described in the files, it was considered as missing data. Kaplan–Meier product-limit estimation with log-rank (P < 0.05) was used for survival analysis from lifetime data according to gene expression levels. The GraphPad Prism 5 statistical package (GraphPad Software, Inc, La Jolla, CA, USA) was used for statistical analysis.

RESULTS

PROX1 Gene and Protein Expression is Downregulated in OSCC Compared With Normal Oral Mucosa

We performed RT-qPCR amplifications on 40 fresh surgical samples of primary OSCC and their corresponding nonneoplastic margin tissues and OSCC cell lines to assess the PROX1 mRNA expression level. Significantly higher PROX1 expression levels were found in nonneoplastic margins (Min–Max, 10.59–1010) compared with OSCC samples (Min–Max, 1.19–22; 5.17-fold, P < .001; Wilcoxon test) (Figure 1A). OSCC cell line SCC-9 exhibited low PROX1 mRNA expression levels when compared with SCC-4 and SCC-25 cell lines (Man–Whitney, P < .001) (Figure 1B). When PROX1 expression levels (higher vs lower) were correlated with clinicopathological features and disease outcome (Table 1), there was no significant association with age group, tumor location, pTNM classification, pathological grade, lymphatic and/or perineural invasion, and local recurrence.

IHC was also performed to confirm lower PROX1 expression in OSCC samples compared with nonneoplastic margins. Immunoreactivity for PROX1 was observed as a nuclear and cytoplasmatic expression, predominantly in suprabasal layers of nonneoplastic margins (Figure 1C). Only 3 OSCC samples were considered positive for PROX1 expression after IHC score analysis. Similar to mRNA expression, OSCC's showed significantly lower PROX1 immunoeexpression than nonneoplastic margins (P < .001; Man–Whitney test) (Figure 1D and E). Although no statistical difference was found, well-differentiated OSCC showed high PROX1 expression in well-differentiated areas mostly represented by keratin pearls.

PROX1 Amplification, but not Methylation Status, Is Associated With Better Survival Rate

To investigate a possible mechanism for PROX1 down-regulation, methylation analysis was performed on 30 fresh OSCC samples. No difference between OSCC samples and normal oral tissues (data not shown) was found because only 4 tumors and 2 margins presented 50% mean frequency of promoter methylation. PROX1 amplification was seen in 16/23 OSCC samples. There was a significant difference in PROX1 amplification between OSCC and tumor-free margin samples (P < .001, Figure 1F).

PROX1 amplification and methylation showed no significant association with clinicopathological features such as tumor location, pTNM classification, pathological grade, and lymphatic and/or perineural invasion (Table 1). However, the P value for the survival curve, determined by the log-rank test, was significantly different in the survival rates between positive and negative PROX1 amplification groups (P = 0.08, log-rank). Patients with positive PROX1 amplification had much more favorable prognosis than those with negative amplification (Figure 1G). PROX1 methylation status was unrelated to overall survival (P = 0.34, log-rank). We also performed a correlation analysis between PROX1 expression and amplification and found no significant difference. However, there was a positive correlation (r = 0.02, P = 0.90; Spearman correlation), suggesting that PROX1 expression tends to be higher in tumors with PROX1 amplification (data not shown).

PROX1 Overexpression Promotes an Antiproliferative Phenotype in OSCC Cells

To determine whether PROX1 overexpression contributes to proliferation, differentiation, apoptosis, migration, and invasion of OSCC cells, PROX1 was stably expressed in SCC9 cell line because it showed the lowest PROX1 mRNA endogenous levels. PROX1 overexpression was confirmed by increased expression of both PROX1 mRNA and protein in SCC9-PROX1 clones but not in the control (SCC9-control, Figure 2A and B).

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Three stable PROX1-overexpressing clones and 1 control clone were chosen for further analysis.

SCC9-PROX1 cells proliferation significantly decreased when compared with SCC9 and SCC9-control, as assessed by proliferation curves ($P = .04$), BrdU incorporation ($P < .001$), and Ki-67 expression ($P < .001$) (Figure 2C, E, and F). SCC9-PROX1 cell clones also showed increased G0-G1 population in cell cycle analysis (Figure 2D). Also, a decrease in cyclin D1 protein expression was observed when compared with SCC9-control cells (Figure 2G). There was no significant difference in p21 expression in SCC9-PROX1 and control cells (data not shown).

FIGURE 1. PROX1 expression is downregulated in OSCC samples. Normalized expression of PROX1 transcripts in nontumor margins and OSCC samples (A) and in OSCC cell lines (B). The full line corresponds to the median value for each group. Asterisk indicates statistically significant difference between OSCC and nontumor samples ($P < 0.001$, Wilcoxon) (A). SCC9 showed reduced PROX1 mRNA expression levels (B). Immunohistochemical detection of PROX-1 in nontumoral margins (C) and OSCC (D). PROX-1 expression was found in the nucleus and cytoplasm of the epithelial cells located in suprabasal layers for the nontumor margin samples. The PROX-1 protein expression was significantly lower in OSCC samples than nontumoral margins ($P < .001$, Mann–Whitney $U$ test). The IHC scores for nontumoral margins and OSCC range from 1.00 to 12.00 (median, 6.0) and from 1.00 to 9.00 (median, 1.50), respectively (E). Kaplan–Maier estimation of overall survival of OSCC patients according to PROX1 amplification ($P$ value according to log-rank test) (F). IHC = immunohistochemistry, OSCC = oral squamous cell carcinoma, PROX1 = prospero homeobox 1.
Genes related to cell proliferation were selected arbitrarily from a previous microarray analysis (unpublished data). qRT-PCR was used to investigate selected genes related to cell proliferation: GATA3 and WISP3 were upregulated (2.79 and 3.77 fold-change mean, respectively), whereas E2F1 and NOTCH1 were downregulated (-11.90 and -1.82 fold-change mean, respectively) in SCC9-PROX1 cell clones compared with SCC9-control (Figure 3).

To further characterize the effects of PROX1 overexpression, in vitro assays to measure apoptosis, migration, and invasion were carried out. PROX1 overexpression did not significantly modulate those events when compared with control cell (Supplemental file 1, 2, and 3, http://links.lww.com/MD/A82).

**PROX1 Overexpression Reduces CK Expression Associated With Poorly Differentiated OSCC**

To assess the effects of PROX1 overexpression on OSCC differentiation, immunocytochemical analysis on CKs 1, 10, 13, 14, 16, and 19 were performed. Immunocytochemical evaluation showed that all CK reactivity was homogeneously restricted to the cytoplasm of positive cells. There was discrete reduced expression for CK1 and 13 in SCC9-PROX1 cells compared with the control cells (data not shown). CK18 and 19 expression were significantly reduced in SCC9-PROX1 cell clones when compared with SCC9 and SCC9 control cells (P < .001, one-way ANOVA, Figure 4). Expression of CK10, 14, and 16 were expressed in almost all cells with no observed difference between SCC9-PROX1 cell clones and control cells (data not shown).

**DISCUSSION**

PROX1 was previously reported by us as downregulated in OSCC after microarray analysis. This prompted us to further investigate PROX1 role in this neoplasm. mRNA expression levels was found lower in OSCC samples when compared with their nontumoral margins. Its protein expression levels were also decreased in OSCC samples when compared with nontumoral margins, suggesting that loss of PROX1 expression in OSCC is associated with a malignant phenotype. Additionally, only well-differentiated tumors showed on IHC a weak positivity for PROX-1 protein. In agreement with these findings, in vitro overexpression of PROX1 was associated with reduced cell proliferation of OSCC cells. Also, a more favorable prognosis was seen in patients with positive PROX1 amplification. All these findings favor the hypotheses that PROX1 may act as a tumor suppressor gene in OSCC.

**TABLE 1.** PROX1 gene expression levels and amplification associated with OSCC clinicopathological features (P value Fisher exact test)

| Clinicopathological Features | PROX1 Expression | PROX1 Amplification |
|-----------------------------|------------------|---------------------|
| Age *                      |                  |                     |
| 40–60 y old                | High 11, Low 14  | P Value 0.29        |
| >60 y old                  | High 6, Low 3    | P Value 1.00        |
| Tumor location *          |                  |                     |
| Tongue                     | High 7, Low 6    | P Value 1.00        |
| Floor of the mouth        | High 10, Low 11  | P Value 1.00        |
| pT classification *       |                  |                     |
| T1/T2                      | High 7, Low 7    | P Value 1.00        |
| T3/T4                      | High 10, Low 10  | P Value 1.00        |
| pN classification *       |                  |                     |
| N+                         | High 7, Low 7    | P Value 0.74        |
| N0                         | High 9, Low 11   | P Value 0.74        |
| Pathological grade *      |                  |                     |
| Well-differentiated        | High 9, Low 6    | P Value 0.74        |
| Moderately differentiated  | High 8, Low 11   | P Value 0.74        |
| Lymphatic invasion *      |                  |                     |
| LI+                        | High 5, Low 7    | P Value 0.72        |
| LI-                        | High 11, Low 11  | P Value 1.00        |
| Perineural invasion *     |                  |                     |
| PI+                        | High 10, Low 9   | P Value 0.73        |
| PI-                        | High 7, Low 8    | P Value 0.73        |
| Local recurrence *        |                  |                     |
| Present                    | High 2, Low 4    | P Value 0.68        |
| Absent                     | High 14, Low 14  | P Value 0.68        |

L1 = lymphatic invasion, OSCC = oral squamous cell carcinoma, PNI = perineural invasion.

* Missing data of 6 patients for PROX1 expression and 5 patients for PROX1 amplification.
the expression of genes that promote or inhibit proliferation and cell cycle progression. In some cancer types as breast and hepatocarcinoma, PROX1 is inactivated because of neoplastic transformation and progression of cancer cells, indicating a tumor suppressor function. Moreover, PROX1 plays a tumor-promoting role in colon cancer, malignant astrocytoma, and kaposiform hemangioendothelioma. As mentioned above, our findings suggest that decreased or loss of PROX1 expression is evident after epithelial malignant transformation. Consistent with this finding, PROX1...
amplification is seen in patients with a better prognosis, although PROX1 expression was not related with clinical features presented by our patients, probably because of the low number of cases. Interestingly, Sasahira et al.26 found PROX1 overexpression in OSCC samples, which was associated with local progression, clinical stage lymphovessel density, nodal metastasis, and worse prognosis. These opposite findings in the same histologic tumor may be because of different patient profiles, and also regional differences between populations. It is well known that OSSC tumors may have different behavior profiles, and also regional differences between populations. It is well known that OSSC tumors may have different behavior when considering patient gender, age, localization, and etiological factors.27 Therefore, future studies confirming the clinical and prognostic value of PROX1 in OSSC should consider evaluating a large cohort of patients with the same clinical profile.

Decreased expression of PROX1 was also found in other types of tumors as lymphomas, sporadic breast cancer, hematological malignancies, and carcinomas of the liver, biliary duct, and pancreatic biliary system.22–24,38,39 In hepatocellular carcinomas, reduced PROX1 expression is associated with poorly differentiated tumors and worst prognosis.18 This study did not find association of PROX1 expression levels with prognosis and survival in OSCC.

Epigenetic silencing is one of the mechanisms responsible for PROX1 inactivation in tumors.22,23 Hypermethylation of CpG islands was identified as a mechanism for PROX1 inactivation in breast carcinomas and carcinomas of the biliary system.22,23 This encouraged us to investigate if PROX1-decreased expression in OSCC was related to promoter DNA methylation. We performed methylation assays in 30 OSCC and 17 nontumoral margins, and there were no significant differences in promoter methylation between the samples. However, it is a common knowledge that gene expression is a complex process with multiple levels of control, and besides methylation, homeobox genes are also under microRNA and translational regulation control; for example, genetic alterations could also be an alternative mechanism leading to PROX1 downregulation in OSCC. These genetic alterations have been previously found in carcinomas of the biliary system, cell lines derived from hematological malignancies, and hepatocellular carcinomas.18,21,22

We investigated PROX1 amplification in OSCC and nontumoral margins because gene amplification can lead to increased gene expression levels. It was observed that OSCC samples showed higher gene copy number when compared with nontumoral margins. To investigate if tumors with higher PROX1 expression levels also exhibited PROX1 amplification, we performed correlation analysis between amplification and expression in the same tumor samples. Although we found no significant differences, the positive correlation suggests that PROX1 expression tends to be higher in tumors with PROX1 amplification. Neither methylation nor amplification analysis showed any significant correlation.

FIGURE 3. PROX1 transcriptionally regulates the expression of NOTCH1, E2F1, GATA3, and WISP3. NOTCH1 (A) and E2F1 (B) were downregulated in SCC9-PROX1 cell clones compared with control cells as well as GATA3 (C) and WISP3 (D) that were upregulated. PROX1 = prospero homeobox 1.
inhibits in vitro cell proliferation of OSCC, which corroborates with previous results in other tumor cells from esophageal squamous cell carcinoma and hepatocellular carcinoma. Under normal development, this gene has been associated with the regulation of cell proliferation by downregulating cell cycle inhibitors CDKN1B (p27) and CDKN1A (p21). In this study, PROX1 overexpression reduced cyclin D1 expression, without interfering with p53 and p21 expression. Overexpression of cyclin D1 produces shorter G1 phase and less dependency on growth factors, resulting in abnormal proliferation. Foskolou et al recently demonstrated that PROX1 was sufficient to decrease Cdc25A and induce p27-Kip1 but not p21-Cip1 or p53, negatively regulating neuroblastoma carcinogenesis. Additionally, PROX1 also suppresses proliferation of hepatocarcinoma cells via inhibiting Twist to trigger p53-dependent senescence. Possibly PROX1 regulates different cell cycle proteins, which could account for the context dependent function of this gene in cancer pathogenesis. We believed that PROX1 could contribute to reduce the proliferation of OSCC-altering cyclin D1 expression.

We also found that PROX1 transcriptionally regulates the expression of genes related to cell proliferation, as demonstrated by upregulated genes WISP3 and GATA3 and downregulated genes NOTCH1 and E2F1 in SCC9-PROX1 cells. WISP3 is downregulated in aggressive cancers and lost in most invasive carcinomas, including breast carcinoma, where it is tumor-inhibitory and seems to suppress breast tumor growth and invasiveness of tumor cells. Loss of GATA3 expression was found to be a reliable indicator of poor prognosis in breast cancer, and lack of GATA3 function is associated with lower survival, more malignant histological attributes, metastasis, and increased tumor mass. E2F1 plays a pivotal role in regulating the expression of genes involved in G1-S transition and DNA synthesis, and it is the most well-known transcription factor regulated by the cyclin/Cdk/Rb pathway. E2F1 regulatory binding sites are also present in the PROX1 gene promoter. Additionally, E2F1 is overexpressed in OSCC and silencing of E2F-1 inhibits proliferation and induces apoptosis. Thus, PROX1 may act in altering E2F1 expression and downregulating cyclin D1, resulting in reduced cell proliferation in PROX1 overexpressing cells.
CKs are structural components of the epithelial cytoskeleton and constitute major proteins for cellular differentiation. Alterations in the expression pattern of some specific CKs have been reported to contribute to progression of OSCC.\(^5\)\(^0\)\(^5\)\(^1\)\(^2\) PROX1 overexpressing cells showed significantly reduced expression of CK18 and CK19. These CKs were previously associated with worse prognosis, poorer pathologic differentiation grade, tumor recurrence, and metastasis to lymph nodes in OSCC.\(^5\)\(^0\)\(^5\)\(^2\) Thus, the PROX1 gene may contribute in the differentiation of OSCC by downregulating these CKs because differentiated cells did not express CK18 and CK19. Interestingly, PROX1 also contributes to tumor differentiation in hepatocellular carcinomas and pancreatic carcinoma.\(^3\)\(^8\)

In summary, our results provide evidence that PROX1 is differentially expressed between nontumoral margins and OSCC, its overexpression reduces proliferative activity and contributes to the differentiation of OSCC cells, and its amplification is related to better prognosis of OSCC samples, suggesting that the downregulation of this gene in oral cancer contributes to the malignant phenotype.

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