

# Effect of siRNA-mediated downregulation of VEGF in Tca8113 cells on the activity of monocyte-derived dendritic cells

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Received November 1, 2011; Accepted December 28, 2011

DOI: 10.3892/ol.2012.568

**Abstract.** Vascular endothelial growth factor (VEGF) is a tumor angiogenesis factor that is important in immune regulation. In our previous study, we found that VEGF expression in the peripheral blood and neoplasm nest from patients with oral squamous cell carcinoma (OSCC) was positively correlated with the course of disease, while an inverse correlation between VEGF expression and dendritic cells (DCs) was identified in the peripheral blood. Therefore, in the present study, we investigated whether inhibition of human VEGF in the human tongue carcinoma cell line Tca8113 had effects on the activity of monocyte-derived DCs. We knocked down the expression of human VEGF in Tca8113 cells using the small interfering RNA (siRNA) technique. Tca8113 cells pre-transfected with siRNA targeting VEGF were co-cultured with monocyte-derived immature and mature DCs. Cell proliferation was evaluated by a WST-8 assay. Cell apoptosis, cell cycle and cell phenotypes were determined by flow cytometry. The data revealed that downregulation of the human VEGF significantly inhibited the proliferation of Tca8113 cells and increased apoptosis. Inhibition of human VEGF arrested the cell cycle of Tca8113 cells at the G0/G1 phase. Our results showed that the co-culture

of DCs with Tca8113 cells markedly inhibited the expression of the mature markers of DCs including HLA-DR, CD80, CD86, CD40 and CD1a, as well as the immature marker CD83, while inhibition of human VEGF in Tca8113 cells significantly reversed these effects. Therefore, human VEGF in Tca8113 cells may not only regulate the cell proliferation and apoptosis of oral squamous cell carcinoma cells, but may also inhibit DC maturation.

## Introduction

Head and neck squamous cell carcinomas (HNSCCs) are common malignancies. Combinations of chemotherapy, surgery and radiotherapy have shown beneficial effects on the local and regional control of these diseases. However, these strategies have been unsuccessful in the treatment of local recurrence and metastatic diseases (1,2). Therefore, a number of investigators are examining new methods to overcome the shortcomings of these treatments.

Various types of cancer express vascular endothelial growth factor (VEGF), which induces neoangiogenesis and blockade of angiogenesis, a critical step in the development and metastasis of solid tumors, and has become a major useful approach in cancer therapy (3). Our previous studies revealed that VEGF expression in the peripheral blood and neoplasm nest of patients with oral squamous cell carcinoma (OSCC) was positively correlated with the course of this disease and was inversely correlated with dendritic cells (DCs) in the peripheral blood (4). DCs are specialized antigen presenting cells, which acquire, process and present antigens, such as tumor-associated antigens, to T cells for the induction of antigen-specific immune responses. Immature DCs are known to migrate into tumor beds and capture tumor-derived antigens for presentation to specific T cells. However, adaptive immune responses are often hampered in cancer patients (5). Panayiotis *et al* reported that VEGF was associated with worse overall survival in patients with HNSCC (6). We found that a low density of mature DC infiltrated into tumor tissue, which may be caused by the immunosuppressive microenvironment of OSCC (4).

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**Abbreviations:** APC, antigen presenting cell; DC, dendritic cell; GM-CSF, granulocyte macrophage colony-stimulating factor; IL-4, interleukin-4; OSCC, oral squamous cell carcinoma; PBMC, peripheral blood mononuclear cell; PE, phycoerythrin; TNF- $\alpha$ , tumor necrosis factor  $\alpha$ ; VEGF, vascular endothelial growth factor

**Key words:** oral squamous cell carcinoma, vascular endothelial growth factor, RNA interference, dendritic cells

Considering that the blockade of VEGF in a mouse model leads to increased antigen uptake and migration of tumor-associated DCs (7), we speculated that inhibition of human VEGF increases the differentiation and maturation of DCs in OSCC, resulting in an increased inhibition of tumorigenesis. In the present study, we investigated whether inhibition of human VEGF in the human tongue carcinoma cell line Tca8113 had an impact on the activity of monocyte-derived DCs. We downregulated the expression of human VEGF in Tca8113 cells using the small interfering RNA (siRNA) technique. We analyzed the expression of mature markers on DCs following the co-culture of DCs with VEGF-downregulated Tca8113 cells.

## Materials and methods

**Cell line.** Tca8113 cells were kindly provided by the Ninth People's Hospital of Shanghai, China. The cells were cultured in RPMI-1640 medium (Gibco, Grand Island, NY, USA) supplemented with 10% heat-inactivated fetal bovine serum (FBS, HyClone Corp, USA). The cultures were incubated at 37°C in an atmosphere of 5% humidified CO<sub>2</sub>.

**RNA extraction and reverse transcription polymerase chain reaction (RT-PCR).** Total RNA was isolated using the TRIzol<sup>®</sup> reagent (Invitrogen, USA) according to the manufacturer's instructions. For RT-PCR, the following DNA primers were synthesized by Invitrogen: Human VEGF, sense primer 5'-GAGGGCAGAATCATCACGAAGT-3' and anti-sense primer 5'-GGTGAGGTTTGATCCGCATAA-3'; human GAPDH, sense primer 5'-TGGTCTCCTCTGACTTCAAC-3' and anti-sense primer 5'-GTGAGGGTCTCTCTCTTCT-3'. Human GAPDH was used as an internal control. The PCR program included pre-incubation for 3 min at 94°C, followed by 28 cycles of 30 sec at 94°C, 30 sec at 55°C, 30 sec at 72°C, and final extension for 10 min at 72°C. Amplification was performed using the Applied Biosystems Veriti (ABI Corporation, USA). The PCR products were analyzed on an agarose gel and photographed on a UV-transilluminator.

**Protein expression analysis.** Total protein expression was analyzed with western blotting as previously described (8). Briefly, the cells were solubilized in lysis buffer [10 mM Tris-HCl (pH 7.5), 150 mM NaCl, 1% NP-40, 1 mM PMSF, and 10 ng/ml soybean trypsin inhibitor]. The proteins were transferred onto a polyvinylidene difluoride (PVDF) filter membrane (Amersham Biosciences). Following blocking with 5% non-fat dry milk in PBS containing 0.01% Tween-20 (T-PBS), the membrane was blotted with the anti-VEGF (1:200; eBioscience Inc.) or GAPDH (1:1000; Abcam Inc.) polyclonal antibodies. The blots were washed in T-PBS for 15 min each and then developed using peroxidase-labeled rabbit anti-mouse IgG (1:2000) for 60 min at room temperature. The protein was visualized by enhanced chemiluminescence.

**Construction of siRNA expression vectors.** The mammalian cell expression vector pRNATU6.1/Neo, carrying a green fluorescent protein (GFP) reporter gene, was purchased from GenScript Corp. (USA) and was used to generate biologically active siRNAs under the U6 promoter. Four

21-nucleotide sequences targeting VEGF were designed and cloned into pRNATU6.1/Neo generating VEGFsiRNA-V1, VEGFsiRNA-V2, VEGFsiRNA-V3 and VEGFsiRNA-V4. A siRNA sequence that is not homologous to any known human DNA sequence was used as a negative control. The constructs were verified by DNA sequencing (Bioasia, China). The effects of RNA interference on the constructs were investigated by RT-PCR.

**Cell transfection.** A total of 1.0x10<sup>4</sup> Tca8113 cells were plated in each well of a 12-well plate (Costar, Corning, USA). Upon attaining 90-95% confluence, the cells were transfected with 2 µg DNA and 4 µl Lipofectamine 2000 (Invitrogen, USA) per well according to the manufacturer's instructions. G418 (400 ng/ml) (Life Technologies) was added to the medium after 48 h of culture, and the cells were cultured for two weeks to permit selection.

**Cell viability analysis.** Cell viability was assessed with a cell counting kit according to the manufacturer's instructions (Dojin Laboratories, Kumamoto, Japan). Briefly, cells at a density of 3x10<sup>3</sup> cells/well were plated in a 96-well plate. Following 5 days of culture, 10 µl 4-[3-(2-methoxy-4-nitrophenyl)T1]-2-(4-nitrophenyl)-2H-5-tetrazolol-1, 3-benzene disulfonate sodium salt (WST-8) was added to each well. Following incubation for 1, 2, 3 and 4 h, the absorbance was measured at 450 nm using a multi-detection microplate reader (Hynergy<sup>™</sup> HT, Bio-Tek).

**Cell cycle and apoptosis analysis.** The cell cycle was analyzed with propidium iodide (PI) as previously described (9). Briefly, the cells were fixed with 70% ethanol and were subsequently treated with 0.25 mg/ml RNase. Finally, cells were stained with 20 µg/ml propidium iodide (PI). Annexin V/PI was used for the apoptosis analysis. Briefly, cells were stained with Annexin V-APC (Bender MedSystems Inc., USA) for 15 min and then incubated with PI for an additional 10 min. Viable cells were PI<sup>-</sup>Annexin V-APC<sup>-</sup> and non-viable cells, which underwent necrosis, were PI<sup>+</sup>Annexin V-APC<sup>+</sup>. Early apoptotic cells were PI<sup>-</sup>Annexin V-APC<sup>+</sup>, whereas late apoptotic dead cells exhibited PI<sup>+</sup>Annexin V-APC<sup>+</sup>.

**Generation of monocyte-derived DCs.** Peripheral blood mononuclear cells (PBMCs) were isolated from human blood using Ficoll/Hypaque (density 1.077 g/ml, Shanghai Shisheng Cytobiotechnological Corp., China) according to the manufacturer's instructions. Following the last wash, PBMCs were resuspended in RPMI-1640 with 10% FBS and were then plated in a 24-well culture plate. Following 3 h of incubation, monocytes were purified by removing the non-adherent cells in the supernatant. To obtain immature DCs, monocytes were cultured with 100 ng/ml rhGM-CSF and 40 ng/ml rhIL-4 (Peprotech UK) for five days. Mature DCs were obtained following an additional two days of culture in medium supplemented with 100 ng/ml rhGM-CSF, 40 ng/ml rhIL-4 and 20 ng/ml rhTNF-α (Peprotech, UK).

**Cell co-culture.** Cell-to-cell contacting co-culture experiments were performed as previously described (ref.). In one set of experiments, immature DCs were co-cultured with Tca8113 or VEGFsiRNA-treated Tca8113 cells. In another set

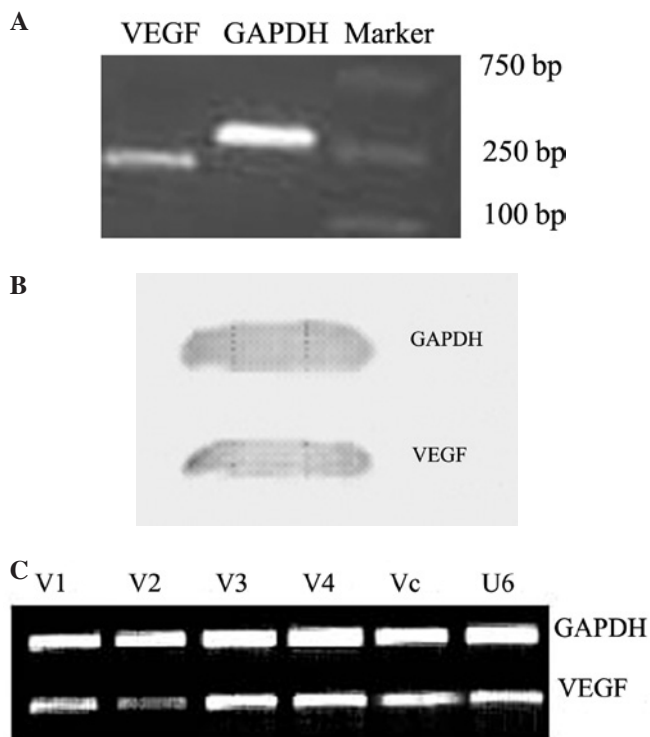


Figure 1. The expression of VEGF in the Tca8113 cell line and selection of the effective VEGF siRNA sequence in Tca8113 cells. The expression of VEGF in Tca8113 was detected by (A) RT-PCR and (B) by western blotting. (C) The efficiency of transfection reached approximately 90% as observed by fluorescence microscopy. (D) VEGF siRNA-V2 effectively inhibited VEGF expression in Tca8113 cells after 48 h by RT-PCR. V1: VEGFsiRNA-V1. V2: VEGFsiRNA-V2. V3: VEGFsiRNA-V3. V4: VEGFsiRNA-V4. Vc: siRNA-Vc. U6: empty vector. VEGF, vascular endothelial growth factor; siRNA, small interfering RNA; RT-PCR, reverse transcription polymerase chain reaction.

of experiments, mature DCs were co-cultured with Tca8113 or VEGFsiRNA-treated Tca8113 cells. The co-culturing conditions were the same for both sets of experiments where RPMI-1640 was supplemented with 10% FBS, 100 ng/ml, rhGM-CSF, 40 ng/ml rhIL-4 and 20 ng/ml rhTNF- $\alpha$ . The ratio of DC to tumor cells was 1:1 at  $1 \times 10^5$  cells/ml.

**Analysis of the DC phenotype with flow cytometry.** The DC phenotype was analyzed with flow cytometry. Mouse anti-HLA-DR-PerCP-cy5.5, CD80 (PE), CD83 (PE), CD1a (PE) and CD14 (APC), as well as corresponding isotypes were used, and murine isotype APC-conjugated, PE-conjugated and PerCP-cy5.5 IgG antibodies were purchased from eBioscience (USA). The antibodies for CD86 (APC) and CD40 (APC) were from Caltag (USA), while the murine isotype served as a negative control.

**Statistical analysis.** The data were analyzed using the SPSS version 13.0 software. Differences between paired groups were analyzed using Student's paired t-test.  $P < 0.05$  was considered to indicate a statistically significant difference.

## Results

**Downregulation of VEGF in Tca8113 cells.** The PCR and western blotting results revealed that Tca8113 cells highly

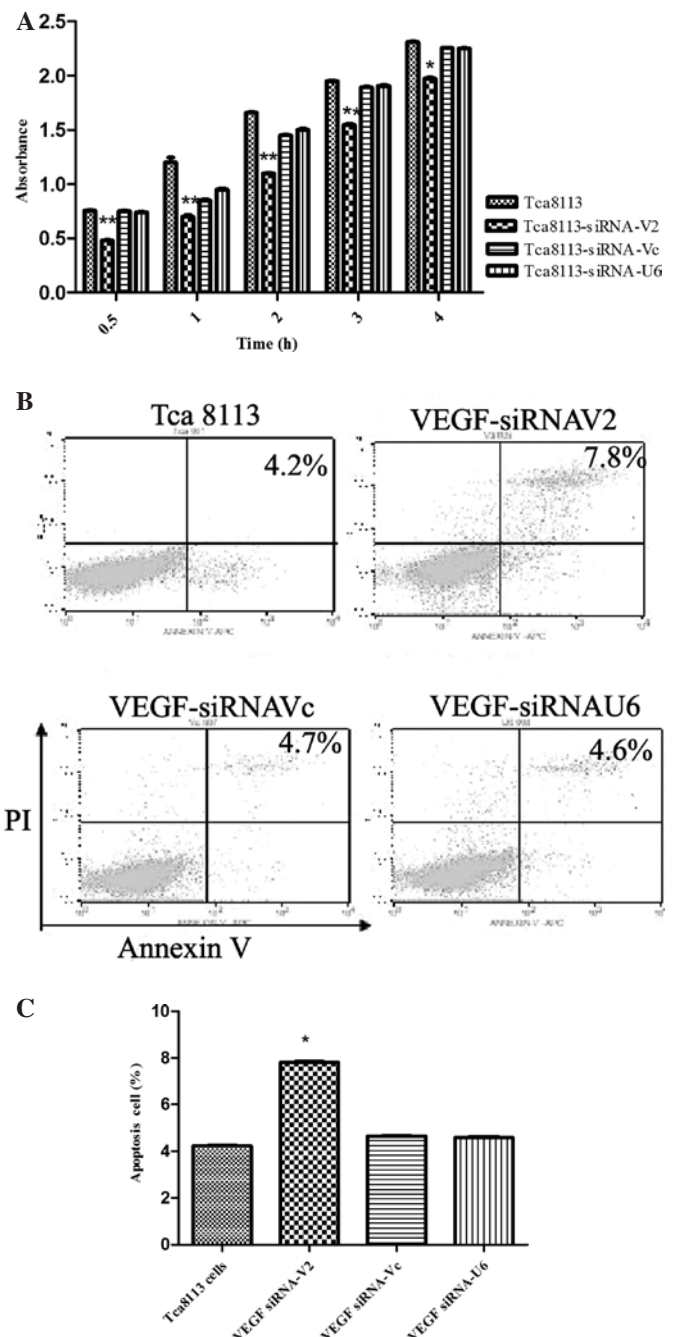
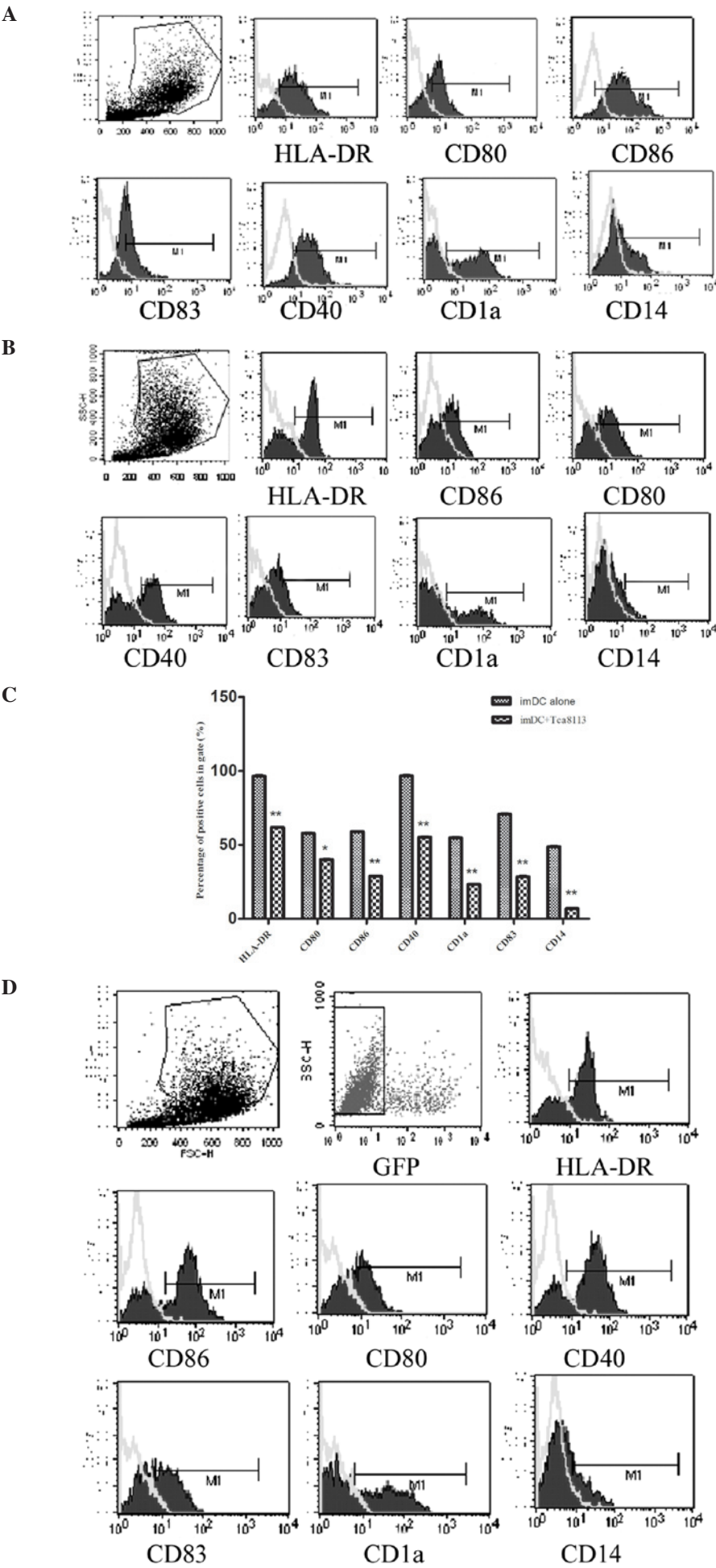


Figure 2. Effects of VEGF siRNA-V2 on the viability and apoptotic state of Tca8113 cells. (A) VEGF siRNA-V2 inhibited Tca8113 cell viability as assayed with a cell counting kit using WST-8 dye. Results are shown as the mean absorbance  $\pm$  SEM of three independent experiments. (B) VEGF siRNA-V2 promoted Tca8113 cell apoptosis. The cells were detected by Annexin V/PI double staining, and representative dot plots of Annexin-V/PI staining are shown: lower left quadrant, viable (double negative) population; lower right quadrant, apoptotic (AnnexinV<sup>+</sup>/PI) population; upper left quadrant, necrosis (Annexin V/PI<sup>+</sup>) population; upper right quadrant, late apoptotic dead cells (Annexin V<sup>+</sup>/PI<sup>+</sup>). (C) The percentages of Annexin V positive cells are shown. Data are shown as the means  $\pm$  SEM of three independent experiments. \* $P < 0.05$  versus control. VEGF, vascular endothelial growth factor; siRNA, small interfering RNA; PI, propidium iodide.

expressed VEGF at the mRNA and protein levels (Fig. 1A and B). We blocked the expression of VEGF in Tca8113 cells by transfection with VEGFsiRNA-V1, VEGFsiRNA-V2, VEGFsiRNA-V3 and VEGFsiRNA-V4, respectively. The transfection efficiency reached approximately 90%. Following





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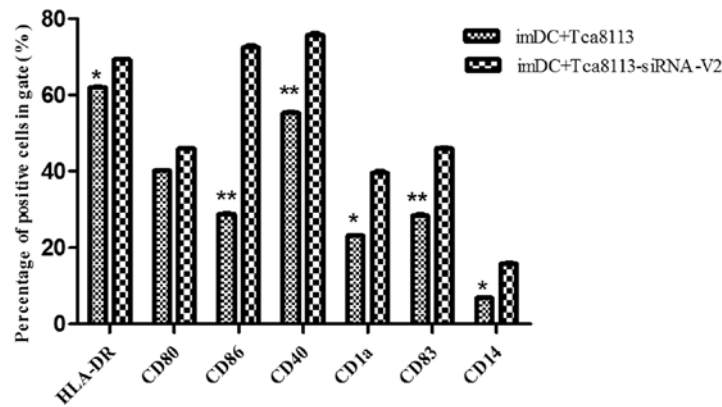


Figure 3. VEGF suppressed the differentiation of immature DCs into mature DCs. Immature DCs were cultured alone or with Tca8113 cells in medium containing GM-CSF, IL-4 and TNF- $\alpha$  for 48 h. (A) The expression of HLA-DR, CD40, CD80, CD83 and CD1a were analyzed by flow cytometry. (B) The expression of HLA-DR, CD40, CD80, CD83 and CD1a decreased when immature DCs were co-cultured with Tca8113 cells for 48 h. (C) The expression of HLA-DR, CD40, CD80, CD83 and CD1a was suppressed in DCs when co-cultured with Tca8113 cells. (D) The expression of HLA-DR, CD40, CD80, CD83 and CD1a increased when immature DCs were co-cultured with VEGFsiRNA-V2-treated Tca8113 cells for 48 h. (E) The expression of HLA-DR, CD40, CD80, CD83 and CD1a was higher on immature DCs when co-cultured with VEGF siRNA-V2-treated Tca8113 cells than with wild-type Tca8113 cells. Data are the mean  $\pm$  SD of triplicate samples. \* $P$ <0.05 versus control. DCs, dendritic cells; VEGF, vascular endothelial growth factor; siRNA, small interfering RNA; GM-CSF, granulocyte macrophage colony-stimulating factor; IL-4, interleukin-4; TNF- $\alpha$ , tumor necrosis factor  $\alpha$ .

Table I. Cell cycle states of Tca8113 cells with or without transfection with siRNA plasmid (%).

Groups	G0-G1	G2-M	S
Tca8113	57.00 $\pm$ 0.98	8.10 $\pm$ 1.22	34.90 $\pm$ 0.85
Tca8113-shRNA-V2	63.06 $\pm$ 0.76	4.95 $\pm$ 1.30	31.99 $\pm$ 0.34
Tca8113-shRNA-VC	58.44 $\pm$ 1.38	7.58 $\pm$ 0.56	33.98 $\pm$ 0.67
Tca8113-shRNA-U6	58.76 $\pm$ 0.51	6.79 $\pm$ 1.36	34.45 $\pm$ 0.67

G418 selection for two weeks, RT-PCR analysis revealed that VEGF was significantly downregulated in Tca8113 cells transfected with VEGFsiRNA-V2, as compared with the controls (Fig. 1C). Therefore, we selected VEGFsiRNA-V2 as the most effective plasmid for use in subsequent experiments.

**Downregulation of VEGF in Tca8113 cells inhibited cell proliferation and cell cycle and induced apoptosis.** To determine whether interference of VEGF regulated cell growth, cell viability was assessed with a cell counting kit as described in Materials and methods. Downregulation of VEGF in Tca8113 cells by transfection with VEGF siRNA V2 significantly inhibited cell proliferation as compared to the negative controls (Fig. 2A). In contrast to the controls, the cell cycle of Tca8113 cells transfected with VEGFsiRNA-V2 was significantly arrested at the G0/G1 phase (Table I).

To examine whether suppression of VEGF expression induced apoptosis, the cells were stained with Annexin V and PI. The data revealed that apoptotic cells increased significantly in Tca8113 cells following downregulation of VEGF (Fig. 2B and C). These results indicated that VEGF inhibited cell growth and induced apoptosis of Tca8113 cells.

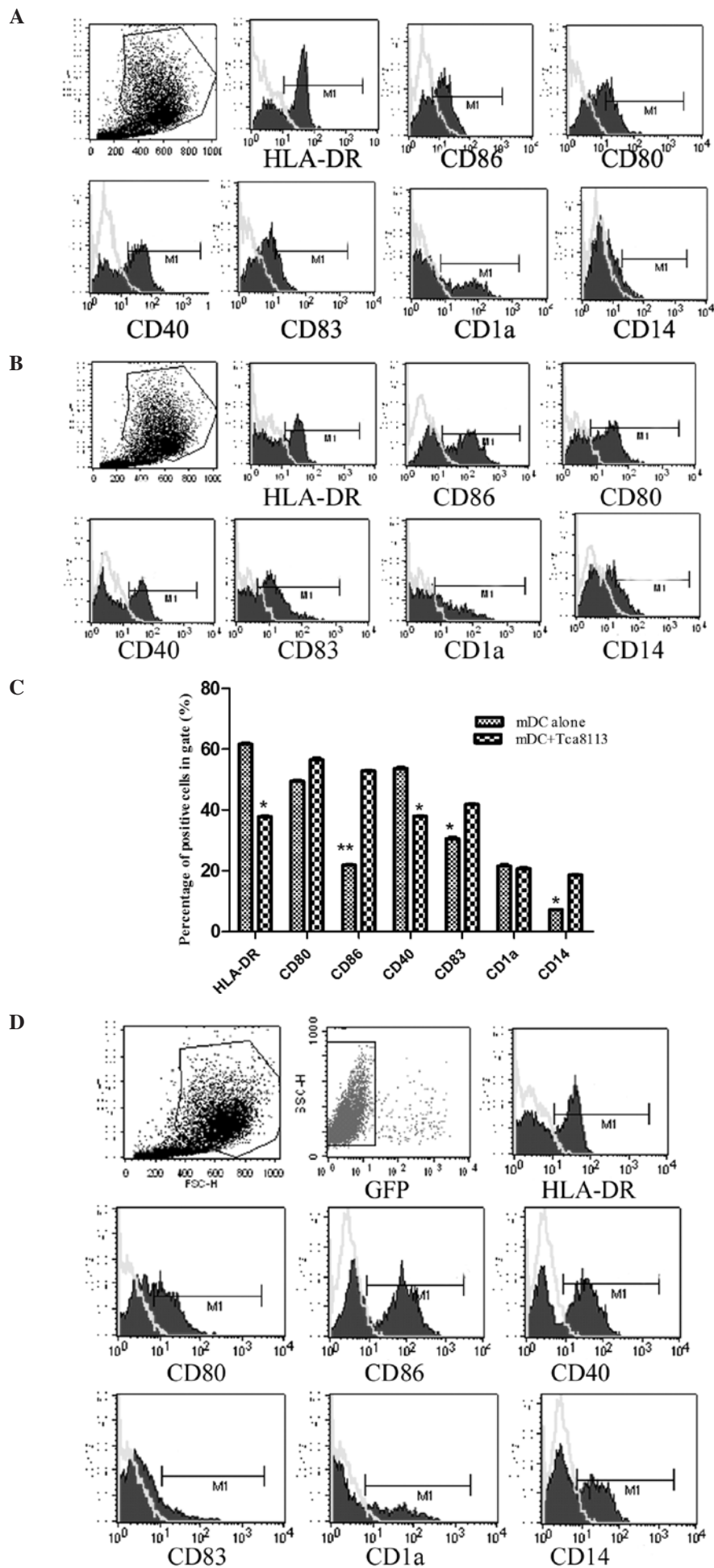
**VEGF inhibited the differentiation of immature DCs into mature DCs.** To analyze whether downregulation of VEGF in

Tca8113 cells had an impact on the differentiation of immature DCs into mature DCs, we co-cultured the immature DCs with Tca8113 cells or VEGF-downregulated Tca8113 cells in the presence of GM-CSF, IL-4 and TNF- $\alpha$ . The mature markers of DCs including HLA-DR, CD80, CD86, CD40 and CD1a, as well as the immature marker CD83, were analyzed by flow cytometry (Fig. 3A). The expression of HLA-DR, CD80, CD86, CD40, CD83, CD1a and CD14 on these DCs was markedly decreased by Tca8113 cells (Fig. 3B and 3C). However, the downregulation of VEGF in Tca8113 cells significantly reversed these effects (Fig. 3D and E). This suggested that VEGF released by Tca8113 had an inhibitory role on the differentiation of immature DCs into mature DCs.

**VEGF-downregulated Tca8113 cells increased the proportion of mature DCs.** To determine whether the downregulation of VEGF in Tca8113 cells regulated the activity of mature DCs, we co-cultured mature DCs with Tca8113 cells or VEGFsiRNA-treated Tca8113 cells. The mature markers of DCs including HLA-DR, CD80, CD86, CD40 and CD1a, as well as the immature marker CD83, were assessed by flow cytometry (Fig. 4A). In contrast to the controls, the expression of HLA-DR, CD80, CD86, CD40 and CD14 on mature DCs following co-culture with Tca8113 cells was markedly decreased, while the immature marker CD83 was increased (Fig. 4B and C). Downregulation of VEGF in Tca8113 cells significantly reversed these effects, whereas the expression of HLA-DR, CD86, CD80, CD40 and CD14 on mature DCs increased (Fig. 4D and 4E). This difference indicated that Tca8113 cells released VEGF, and may regulate the activity of mature DCs.

## Discussion

It is well known that human VEGF is highly expressed in numerous types of cancer. VEGF is particularly upregulated



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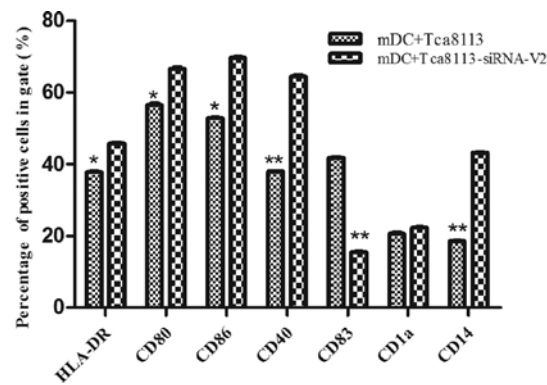


Figure 4. VEGF also suppressed the surface molecules of mature DCs. The mature DCs were cultured alone or co-cultured with Tca8113 cells in the RPMI-1640 medium containing 10% FBS for 48 h. We analyzed the DC surface molecules excluding the Tca8113 cells by setting up two gates. (A) The expression of HLA-DR, CD40, CD80, CD83 and CD1a increased when mature DCs were cultured alone for 48 h. (B) The expression of HLA-DR, CD40, CD80, CD83 and CD1a was decreased when mature DCs were co-cultured with Tca8113 cells for 48 h. (C) The expression of HLA-DR, CD40, CD80, CD83 and CD1a was suppressed in mature DC when co-cultured with Tca8113 cells. (D) The expression of HLA-DR, CD40, CD80, CD83 and CD1a was increased when mature DCs were co-cultured with VEGF siRNA-V2-treated Tca8113 cells for 48 h. (E) The expression of HLA-DR, CD40, CD80, CD83 and CD1a was higher on mature DCs when co-cultured with VEGF siRNA-V2-treated Tca8113 cells than with wild-type Tca8113 cells. Data represent the mean  $\pm$  SD of triplicate samples. \* $P < 0.05$  versus control. DCs, dendritic cells; FBS, fetal bovine serum; VEGF, vascular endothelial growth factor; siRNA, small interfering RNA.

in response to hypoxia, activated oncogenes or inactivated tumor suppression genes (10). Therefore, suppression of VEGF expression at the RNA or protein levels may inhibit tumor growth. It is reported that VEGF influenced the growth of HNSCC (11) and tongue squamous cell carcinoma, one of the most common malignancies in the oral cavity (11,12). However, VEGF expression in Tca8113 cells was previously unknown. In this study, we found that the human tongue carcinoma cell line Tca8113 highly expressed VEGF at the mRNA and protein levels. Therefore, we established the biological basis for inhibiting or blocking the growth and development of tongue squamous cell carcinoma by interfering with VEGF expression.

When VEGF was downregulated in Tca8113 cells by siRNA produced from transfected plasmids, the proliferative ability was decreased and the proportion of cells in the G0 phase and apoptotic state were increased. These findings indicated that VEGF modulated the growth and apoptosis of human tongue squamous cell carcinoma and further validated VEGF as a rational therapeutic target.

In our study, the recombinant vector with green fluorescence protein (GFP) targeting VEGF (VEGFsiRNA) was transfected into the Tca8113 cells. Compared with chemically synthesized siRNA, the vector-produced siRNA persisted for a longer period of time in transfected cells (13). With the GFP reporter gene, we were able to detect the efficiency of transfection by fluorescence microscopy and revealed that VEGF gene expression in Tca8113 cells was significantly silenced. It has been shown that when the expression of VEGF is suppressed, the binding of VEGF with its receptor can be blocked, and the proliferation of tumor cells and endothelial cells decreased (14). Therefore, our results suggest that the decreased proliferation of Tca8113 cells may be attributed to the siRNA-mediated suppression of VEGF production.

Some of the most important strategies in the treatment of malignancies include the induction of cell apoptosis and the regulation of the cell cycle, as suggested by Sasano and Suzuki (15). In our study, after silencing the expression of

VEGF, the proportion of cells in the G0-G1 phase and the apoptotic cells increased, which is in agreement with the findings of Sasano and Suzuki (15). Therefore, VEGF is important in the proliferation of tumors, and inhibition of VEGF expression with siRNA is potentially useful in the treatment of tongue squamous cell carcinoma.

Tumor cells often evoke specific immune responses; however, they fail to eliminate all tumor cells. Our previous studies revealed that in OSCC, the majority of the DCs were immature DCs, while the total amount of DCs decreased in the neoplasm nest and peripheral blood of patients with OSCC (16). To study the effect of VEGF on the differentiation of immature DCs to mature DCs, we co-cultured immature DCs with Tca8113 cells or VEGF-downregulated Tca8113 cells in the normal medium containing GM-CSF, IL-4 and TNF- $\alpha$ , which induced the differentiation of immature DCs into mature DCs. Our results revealed that the expression of HLA-DR, CD86, CD40, CD83, CD1a and CD14 on mature DCs following co-culture with Tca8113 cells was significantly decreased, while the downregulation of VEGF in Tca8113 cells reversed this effect. Therefore, we inferred that the high VEGF expressing Tca8113 cells inhibit the immature DCs differentiating to mature DCs. This result is in agreement with the study by Hasebe *et al*, in which monocyte-derived DCs from cancer patients cultured with GM-CSF plus IL-4 exhibited lower levels of CD11c, CD40, CD86 and HLA-DR expression as compared with those derived from healthy volunteers (10).

In our study, we found that the expression of HLA-DR, CD86, CD40, CD83, CD1a and CD14 increased significantly in the immature DCs co-cultured with VEGF-downregulated Tca8113 cells. This result revealed that DC phenotypes were restored to a certain extent following the inhibition of VEGF. The functional activity of DCs depends, in part, on the expression of HLA-DR and co-stimulatory molecules (17). Reduced expression of these cell surface receptors, as observed in the presence of Tca8113 cells, may thus cause altered functionality of DCs. Alfaro *et al* found that DCs still matured under the

effect of VEGF but they expressed less HLA-DR and CD86, and this effect was suspended by the VEGF inhibitor (18).

VEGF also suppressed the surface molecules of mature DCs. We found that the expression of HLA-DR, CD86, CD80, CD40 and CD14 on mature DCs decreased in the presence of Tca8113 cells. However, when mature DCs were co-cultured with VEGF-downregulated Tca8113 cells, the expression of HLA-DR, CD86, CD80, CD40 and CD14 on the DCs was restored. This observation indicated that Tca8113 cells inhibited mature DCs from maintaining their mature status. Moreover, when we co-cultured DCs with VEGF-downregulated Tca8113 cells, the proportion of mature DCs increased to a certain extent. Therefore, siRNA targeting of the VEGF gene was capable of relieving the inhibition of VEGF on DC maturation and improving the function of DCs. Our results further support other previous findings indicating that an increased VEGF is correlated with the reduced number of DCs in tumor tissue and in the peripheral blood of patients with various types of cancer (10,19). VEGF is known to promote tumor growth and inhibit the activation of nuclear factor  $\kappa$ B (NF- $\kappa$ B) in endothelial progenitor cells, thereby inhibiting endothelial progenitor cells from differentiating into mature DCs (20). We speculated that VEGF released by Tca8113 cells induce monocytes differentiating into endothelial cells but not mature DCs. The VEGF-induced endothelial cells may also be involved in angiogenesis in the cancer tissue.

In conclusion, siRNA targeting the VEGF gene is capable of inhibiting Tca8113 cell growth, inducing apoptosis and relieving the inhibition of VEGF on DC maturation. VEGF siRNA may be a novel and promising therapeutic strategy for the treatment of OSCC.

## Acknowledgements

This study was supported by a grant from the National Natural Science Foundation of China (No. 81072213), Nanjing Medical Science and Research Project (No. YYK11039), the Jiangsu Health Science and Research Project (No. H200944), the Jiangsu Provincial Natural Science Foundation (No. BK2009043) and the Nanjing Science and Technology Development Plan (No. 201001084). The authors thank Shanghai Ninth Hospital for kindly providing the Tca8113 cell line.

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