Up-regulation of stromal cell-derived factor-1α and its receptor CXCR4 expression accompanied with epithelial-mesenchymal transition in human oral squamous cell carcinoma

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Abstract. Stromal cell-derived factor 1α (SDF-1α) and its receptor CXCR4 have been implicated in the tumorigenesis, proliferation, and lymph node metastasis of cancer. Here, we report that highly invasive squamous cell carcinoma (SCC) cells with a spindle cell morphology show a strong expression of both SDF-1α and CXCR4. CXCR4 expression and cell migratory activity were further up-regulated by treatment with SDF-1α or TGF-ß1 in these cells. When epithelial-mesenchymal transition (EMT) was induced by Snail over-expression in SCC cells with an epithelial phenotype, an increased expression of SDF-1α was observed. Furthermore, SDF-1α and TGF-ß1 up-regulated the expression of CXCR4 and cell migratory activity in these cells. These results indicate that SDF-1α and CXCR4 expressions are possible markers of highly-invasive SCC and regulated by EMT.

Introduction

Chemokines organize a superfamily of chemotactic cytokines that possesses a wide range of biological activities, including the regulation of leukocyte trafficking, homing of T- and B-lymphocytes, cytoskeletal rearrangement, and adhesion to extracellular matrix molecules (1-3). They are locally produced in the tissues and act on various cells through selective membrane-bound G-protein-coupled receptors. The CC chemokines, RANTES, and macrophage inflammatory protein-1α (MIP-1α) are ligands for CCR5, and the CXC chemokine, stromal cell-derived factor-1 (SDF-1), is known to be a ligand for CXCR4 and regulate T-cell activation. Expression of SDF-1 is enhanced by stimulation with tumor necrosis factor-α (TNF-α), interferon-γ (IFN-γ), and trans-forming growth factor-ß (TGF-ß) in fibroblasts (4). Furthermore, there is accumulating evidence that the chemokines and their receptors regulate the motility and metastasis of cancer cells. Especially, SDF-1α and CXCR4 have been reported to be important mediators of cancer progression (5-8). CXCR4 inhibition can affect tumor cell dissemination in non-Hodgkin's lymphoma and melanoma (9,10) and cancer cells expressing CXCR4 invade the extracellular matrix and metastasize to organs expressing SDF-1α (11).

On the other hand, the increased motility and invasiveness of cancer cells have been implicated in epithelial-mesenchymal transition (EMT), a phenomenon occurring during embryonic development including gastrulation and neuro-epithelium formation (12-15). During the process of EMT, the loss of E-cadherin expression, an adhesion molecule of epithelial cells, is one of the key steps towards the invasive phenotype (16,17). Several transcription factors including zinc-finger proteins of the Snail/Slug family (18,19), δEF1/ZEB1 (20), SIP-1/ZEB2 (21), and E12/E47 (22), that contain the basic helix-loop-helix motif, have been reported to be involved in this repression through binding to the E-box localized in the promoter of E-cadherin. Furthermore, TGF-ß1 has also been reported to regulate EMT in several cell types (23-25).

In the present study, we analyzed SDF-1α and CXCR4 expressions in SCC cells with both epithelial and mesen-chymal phenotypes and in EMT-induced cells by Snail-over-expression. We found that SDF-1α and CXCR4 expressions in SCC cells were up-regulated in conjunction with increased activities of migration by EMT.

Materials and methods

Cells and cell culture. The human oral SCC cell lines, OM-1, HOC719-PE, HOC719-NE, HOC313, their stable Snail over-

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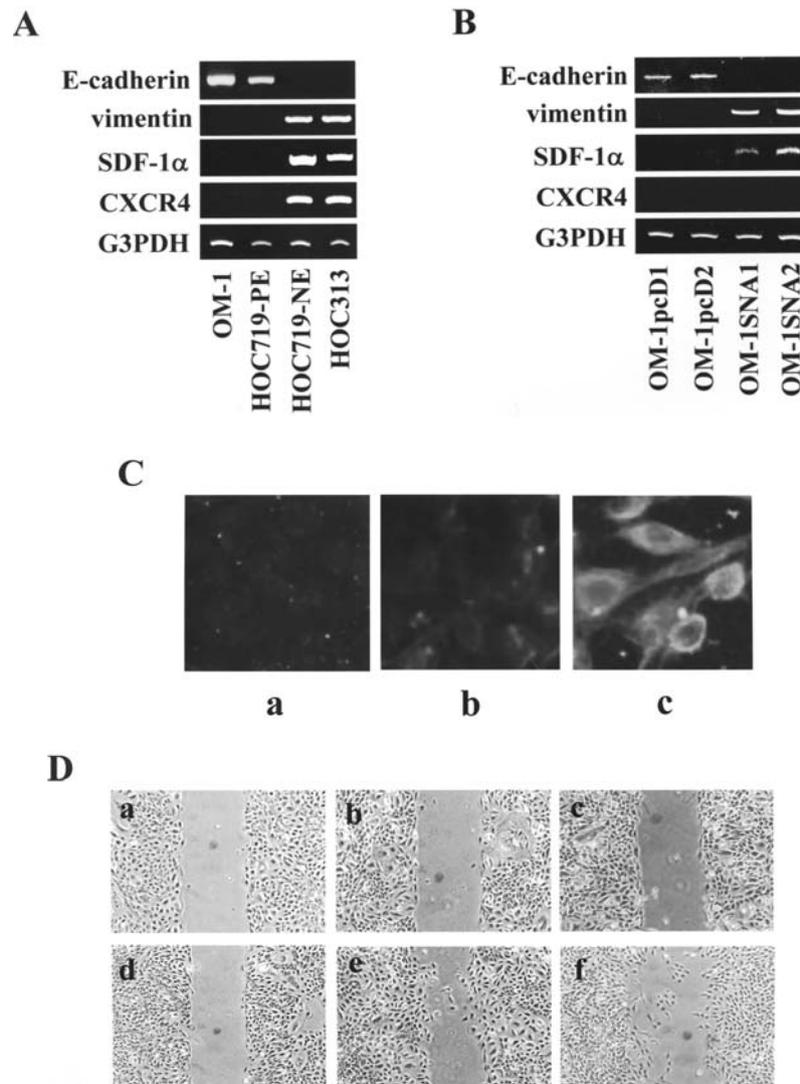Figure 1. Expressions of SDF-1α and CXCR4 in SCC cells. (A) E-cadherin, vimentin, SDF-1α, and CXCR4 expressions in SCC cells with epithelial (OM-1, HOC719-PE) and mesenchymal (HOC719-NE, HOC313) phenotypes. Total-RNAs were isolated from SCC cells and analyzed by RT-PCR. (B) Expression of mRNAs for E-cadherin, vimentin, SDF-1α and CXCR4 in Snail over-expressing (OM-1SNA1, OM-1SNA2) and control (OM-1pcD1, OM-1pcD2) clones. (C) OM-1pcD1 (a), OM-1SNA1 (b), and HOC313 (c) cells were stained with mouse anti-human CXCR4 antibody and visualized by FITC-conjugated anti-mouse IgG secondary antibody. (D) The migratory behavior of OM-1pcD1 (a, d), OM-1SNA1 (b, e), and HOC313 (c, f) cells was analyzed in an *in vitro* wound model. Photographs of culture cells were taken under a phase-contrast microscope in a wound healing assay performed at a magnification of x100. Photographs were taken just after incision (a-c) and 24 h post-incision (d-f).

expressing clones of OM-1SNA1 and OM-1SNA2, and control pcDNA3-transfected clones OM-1pcD1 and OM-1pcD2, have been described previously (19,26,27). HOC719-PE and HOC719-NE cells were isolated from HOC719 cells expressing E-cadherin heterogeneously (19). All cells were cultured at 37˚C in a humidified atmosphere of 5% CO$_2$ in air and maintained with DMEM (Sigma, St. Louis, MO, USA) supplemented with 10% fetal bovine serum (Sigma).

Immunofluorescent staining. Cells cultured on Lab-Tek II Chamber Slides (Nalge Nunc, Tokyo, Japan) were fixed with 4% paraformaldehyde and treated with 0.1% Triton X-100 for 5 min, PBS containing 1% bovine serum albumin (BSA) for 30 min, and a 1:200 dilution of anti-human CXCR4 monoclonal antibody (R&D Systems, Minneapolis, MN, USA) in PBS containing 1% BSA at 4˚C overnight. Detection of CXCR4 protein was performed using 1:1000 dilutions of

FITC-labeled anti-mouse IgG (Dako, Kyoto, Japan). Vectashield (Vector Laboratories, Burlingame, CA, USA) was used as a mounting medium. Images were captured using an Optiphot-2 (Nikon, Tokyo, Japan) configured with fluorescent excitation filters at 800 nm.

Reverse transcriptase-polymerase chain reaction (RT-PCR). Total-RNA was isolated from the cells using TRIzol (Invitrogen, Carlsbad, CA, USA). RT-PCR analysis was performed as described previously (19). The RNA samples were first treated with deoxyribonuclease I (Invitrogen) and converted into cDNA using random hexamer primers and reverse transcriptase (Invitrogen). PCR consisting of 30 cycles of denaturing at 94˚C for 30 sec, annealing for 30 sec, and extension at 72˚C for 1 min was carried out using PCR MASTER (Boehringer Mannheim GmbH, Germany). Amplified products were analyzed on 1.8% agarose gels. The
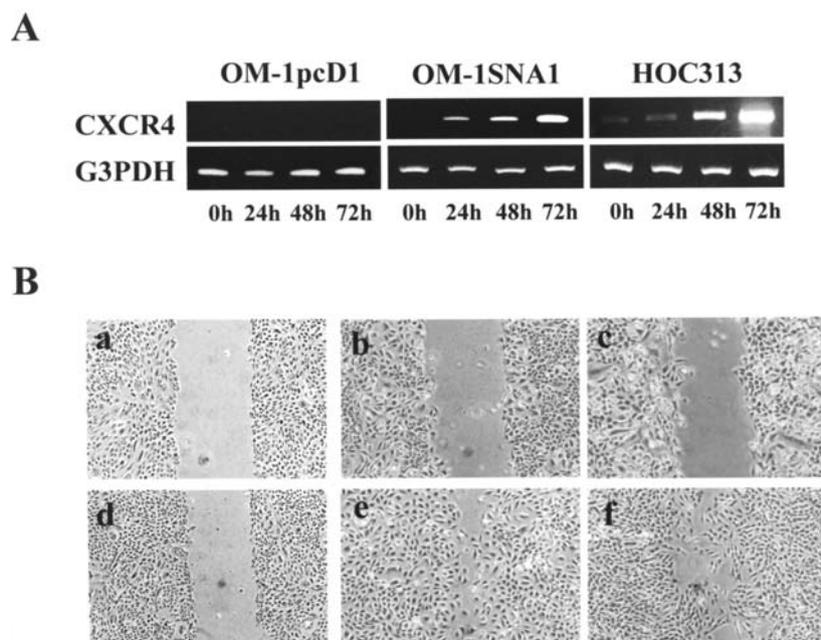
A

B

Figure 2. Effects of SDF-1α on CXCR4 expression in SCC cells. (A) CXCR4 expression in OM-1pcD1, OM-1SNA1 and HOC313 cells treated with 50 ng/ml SDF-1α for the indicated times. Total-RNAs were isolated from SCC cells and analyzed by RT-PCR. (B) The migratory behavior of OM-1pcD1 (a, d), OM-1SNA1 (b, e), and HOC313 (c, f) cells cultured with 50 ng/ml SDF-1α was analyzed in an *in vitro* wound model. Photographs of culture cells were taken under a phase-contrast microscope in a wound healing assay performed at a magnification of x100. Photographs were taken just after incision (a-c) and 24 h post-incision (d-f).

product size, annealing temperature, and primer sequences for SDF-1α and CXCR4 were: SDF-1α, 239 bp, 58˚C, 5'-tgaa cgccaaggtcgtggtcgtg-3' (forward), 5'-ctccaggtactcctgaatccac-3' (reverse); CXCR4, 343 bp, 58˚C, 5'-ctggtattgtcatcctgtcctgc-3' (forward), 5'-tggaccctctgctcacagaggtg-3' (reverse). The primers for E-cadherin, vimentin, and glyceraldehyde-3-phosphate dehydrogenase (G3PDH) were previously described (22,26,27). All primers were obtained from Hokkaido System Science Co., Ltd. (Sapporo, Japan).

Wound healing assay. Cells were seeded in 6-well culture dishes at $3x10^5$ cells per well and incubated until confluent. After confluence, SDF-1α and TGF-ß1 were added to the medium for 48 h. A wound was incised with a pipette tip in the central area of the confluent culture on the dish. Detached cells were removed carefully with PBS and migrated cells in wound areas were observed utilizing a phase-contrast microscope.

Results

Expression of SDF-1α and migratory activity in SCC cells with a mesenchymal phenotype. Expressions of SDF-1α and its receptor, CXCR4, were analyzed in cells with a cuboidal morphology and E-cadherin expression (OM-1 and HOC719-PE) and cells with a spindle morphology, loss of E-cadherin, and strong expression of vimentin and Snail (HOC719-NE, HOC313) (26,27). RT-PCR analysis indicated strong expressions of SDF-1α and CXCR4 in HOC719-NE and HOC313 cells but not in OM-1 and HOC719-PE cells (Fig. 1A). Snail over-expressing OM-1 cells, OM-1SNA1 and OM-1SNA2, showed the down-regulation of E-cadherin and up-regulation

of vimentin expression, as previously described (26). Increased expression of SDF-1α but not CXCR4, was observed in OM-1SNA1 and OM-1SNA2 cells (Fig. 1B). Immunofluorescent staining using a monoclonal antibody showed a strong CXCR4 expression on the cell surface in HOC313 cells (Fig. 1C-c), whereas no significant fluorescence staining was detected in OM-1pcD1 cells (Fig. 1C-a) and OM-1SNA1 cells (Fig. 1C-b). In a wound healing assay, HOC313 (Fig. 1D-f), but not OM-1pcD1 (Fig. 1D-d) cells, were detected in the wound area 24 h post-incision. Increased migratory activity was detected in OM-1SNA1 (Fig. 1D-e) compared to OM-1pcD1 (Fig. 1D-d) cells. OM-1SNA1 and HOC313 cells moved to the wounded area without the formation of cell-to-cell contact.

Increased CXCR4 expression and cell migration by SDF-1α in SCC cells with a mesenchymal phenotype. Cells were cultured with 50 ng/ml of SDF-1α for 72 h and CXCR4 expression was analyzed (Fig. 2A). SDF-1α induced CXCR4 expression in OM-1SNA1 and HOC313 cells but not in OM-1pcD1 cells. Treatment with 50 ng/ml of SDF-1α for 24 h enhanced the migratory properties of OM-1SNA1 (Fig. 2B-e) and HOC313 (Fig. 2B-f) cells but not those of OM-1pcD1 (Fig. 2B-d) cells.

Increased CXCR4 expression and cell migration by TGF-ß1 in SCC cells with a mesenchymal phenotype. Cells were cultured with 2 ng/ml of TGF-ß1 for 72 h and the expressions of SDF-1α and CXCR4 were analyzed (Fig. 3A). TGF-ß1 induced CXCR4 expression in OM-1SNA1 and HOC313 cells, but not in OM-1pcD1 cells. However, SDF-1α expression was induced by TGF-ß1 in all OM1-SNA1, OM-1pcD1, and
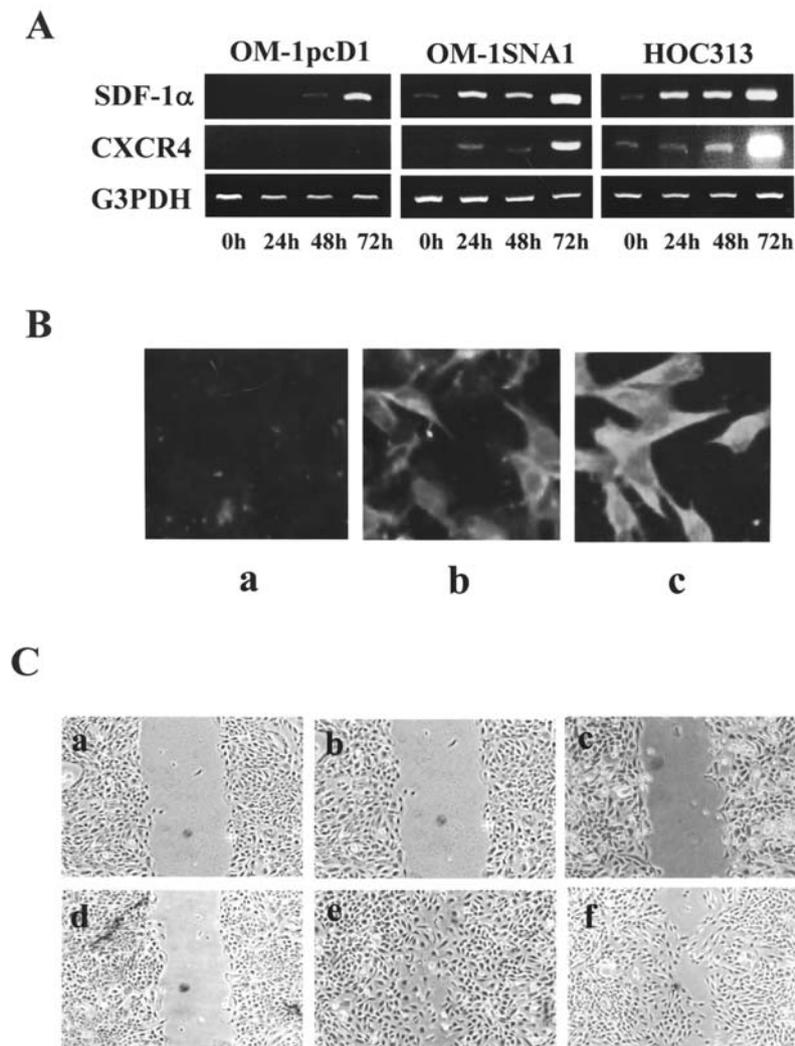
Figure 3. Effects of TGF-ß1 on SDF-1α and CXCR4 expression in SCC cells. (A) Expression of mRNAs for SDF-1α and CXCR4 in OM-1pcD1, OM-1SNA1, and HOC313 cells cultured with 2 ng/ml TGF-ß1 for the indicated times. (B) OM-1pcD1 (a), OM-1SNA1 (b), and HOC313 (c) cells were stained with mouse anti-human CXCR4 antibody and visualized by FITC-conjugated anti-mouse IgG secondary antibody. Before staining, SCC cells were cultured with TGF-ß1 for 72 h. (C) The migratory behavior of OM-1pcD1 (a, d), OM-1SNA1 (b, e), and HOC313 (c, f) cells cultured with 2 ng/ml TGF-ß1 was analyzed in an *in vitro* wound model. Photographs of culture cells were taken under a phase-contrast microscope in a wound healing assay performed at a magnification of x100. Photographs were taken just after incision (a-c) and 24 h post-incision (d-f).

HOC313 cells. In immunofluorescent staining, treatment with TGF-ß1 for 72 h resulted in increased CXCR4 expression in OM-1SNA1 and HOC313 cells, but not OM-1pcD1 cells, similarly to the results in Fig. 3A (Fig. 3B). In the wound healing assay, treatment with TGF-ß1 for 24 h further enhanced the migratory properties of OM-1SNA1 (Fig. 3C-e) and HOC313 (Fig. 3C-f) cells.

Discussion

In previous studies, we showed that OM-1 and HOC719-PE cells had a cuboidal cell morphology with E-cadherin expression, and HOC719-NE and HOC313 cells showed a spindle cell morphology with a loss of E-cadherin and strong expression of vimentin, MMP-2, and Snail (19,26-28). We also reported that Snail over-expression and TGF-ß1 treatment in SCC cells with an epithelial phenotype resulted in a change to the spindle cell morphology and down-regulation

of E-cadherin expression (28). In the present study, we showed strong expressions of SDF-1α and CXCR4 and migratory activity in HOC719-NE and HOC313 cells exhibiting the spindle morphology, but not in OM-1 and HOC719-PE cells with cuboidal shapes. Over-expression of Snail in OM-1 cells resulted in a change to the spindle morphology and the up-regulation of SDF-1α expression accompanied with enhanced migratory activity. Treatment with SDF-1α induced CXCR4 expression in Snail over-expressing and HOC313 cells but not in control cells. TGF-ß1 also induced CXCR4 expression in these cells. Furthermore, SDF-1α expression in all SCC cells was induced by TGF-ß1. Strong migratory activity was detected in HOC313 and OM-1SNA1 cells, and both SDF-1α and TGF-ß1 further promoted the migratory properties of these cells.

SDF-1α has been reported to increase the migratory activity of cells depending on their expression of CXCR4 (29,30). The migration induced by SDF-1α was mediated

Akt activation in epithelioid carcinoma cells (31). Tatsumi *et al* reported that a specific inhibitor of the CXCR4 receptor completely inhibited lung metastasis in a mouse transplantation model and suggested that CXCR4 was involved in the metastatic process of osteosarcoma cells (32). Muller *et al* reported high levels of CXCR4 expression in invasive ductal carcinoma but not normal mammary gland ductal cells using immunostaining (5). They also mentioned a strong CXCR4 expression in tumor cells in lymph node and distant metastases of the lung and liver. Furthermore, Li *et al* reported that expression levels of CXCR4 were correlated with the overall survival rate in breast cancer (33). Chen *et al* observed that TGF-ß1 affected CXCR4 expression through the stimulation of SDF-1α and ERK1/2 phosphory-lation in human monocyte-derived macrophages (34). On the contrary, secreted TGF-ß1 from cholangiocarcinoma cells decreased the expression of SDF-1 in fibroblasts and then inhibited the invasion of cholangiocarcinoma cells (35). We previously reported that Snail expression was strongly increased by TGF-ß1 in highly-invasive SCC cells with the mesenchymal phenotype, but not epithelial phenotype. In the present study, we suggested that the regulation of CXCR4 expression was different with regard to the condition of EMT in SCC cells. Further analysis concerning the regulation of CXCR4 expression accompanying EMT might be useful for the treatment of metastatic lymph nodes of SCC.

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