Abstract. Mitotically quiescent cancer stem cells (CSC) are hypothesized to exhibit a more aggressive phenotype involving greater therapeutic resistance and metastasis. The aim of our study was to develop a method for identifying quiescent CSC in esophageal squamous cell carcinoma (ESCC) based on their expression of the p75 neurotrophin receptor (p75NTR) and other proposed CSC markers, such as CD44 and CD90. Double immunostaining of surgical ESCC specimens revealed that the mean Ki-67-labeling index of the p75NTR-positive cells was significantly lower than that of the p75NTR-negative cells. Real-time PCR analysis of sorted fractions of ESCC cell lines (KYSE cells) revealed that stem cell-related genes (Nanog, p63 and Bmi-1) and epithelial-mesenchymal transition (EMT)-related genes (N-cadherin and fibronectin) were expressed at significantly higher levels in the p75NTR-positive fractions than in the CD44-positive or CD90-positive fractions. In addition, the p75NTR-positive fractions exhibited significantly higher colony formation in vitro, significantly enhanced tumor formation in mice, and significantly greater chemoresistance against cisplatin (CDDP) than the CD44-positive or CD90-positive fractions. Furthermore, in both the cultured cells and those from the mouse xenograft tumors, the p75NTR-positive/CD44-negative and p75NTR-positive/CD90-negative KYSE cell fractions contained significantly higher proportions of mitotically quiescent cells. These results suggest that the mitotically quiescent CSC population in ESCC can be identified and isolated based on their p75NTR expression, providing researchers with a novel diagnostic and therapeutic target.

Introduction

Recent studies have demonstrated that some cancer stem cells (CSC) remain in a mitotically quiescent state, which enhances their chemoresistance and results in a more invasive and aggressive phenotype (1,2). The p75 neurotrophin receptor (p75NTR) is expressed in the quiescent basal layer of the esophageal epithelium (3,4), and in esophageal squamous cell carcinoma (ESCC). It is also expressed in populations of cells that exhibit enhanced colony-forming abilities and xenograft tumorigenicity (5,6). CD44 (7) and CD90 (8) have also been reported to be markers of populations of esophageal CSC that display enhanced xenograft tumorigenicity and/or metastatic potential. However, none of these studies investigated the mitotic status of the examined CSC populations. The aim of this study was to develop a method for identifying mitotically quiescent CSC in ESCC based on their p75NTR, CD44 and CD90 expression patterns.

Materials and methods

Tissue microarray. A total of 56 tumor specimens from ESCC patients who had undergone surgery at our hospital from 1990 to 2008 were analyzed using a tissue microarray, as described previously (9). All of the patients underwent R0 resections, and none of them died in hospital. The median follow-up time was 29 months. The subjects included 50 male and 6 female patients, and their mean age was 62.8 years. The patients' TNM stages (ver. 6) were as follows: stage I, 6 patients; stage IIA, 15 patients; stage IIB, 5 patients; stage III, 24 patients; and stage IV, 6 patients. All of the M1 cases involved only distant lymph node metastases, which were surgically removed. Thirty-six patients underwent postoperative cisplatin-based chemotherapy. The institutional review board at the University of Toyama approved this study (#20-57).
**Immunohistochemistry.** The antibodies used for the immunohistochemical staining were as follows: anti-Ki-67 antibody (dilution 1:100), anti-cytokeratin Oscar in vitro diagnostic antibody (dilution 1:200; Abcam Ltd., Cambridge, UK), anti-human CD44 monoclonal antibody (156-3Cl1; dilution 1:400; Cell Signaling Technology, Beverly, MA, USA), anti-human CD90 monoclonal antibody (EPR3132; dilution 1:100), and human p75NTR monoclonal antibody against p75NGER (NGER5; dilution 1:100) (both from Abcam). The immunostaining was performed using Envision Plus kits, horseradish peroxidase, or 3,3′-diaminobenzidine (DAB; Dako Cytomation, Kyoto, Japan) as recommended by the supplier. Counterstaining was performed with Mayer’s hematoxylin.

The numbers of p75NTR-, CD44- or CD90-positive cells and all tumor cells were counted in three random fields of each section. Then, the proportions of each cell type in each tumor were calculated. We classified tumors as positive when >5% of the tumor cells were stained.

Double staining of Ki-67 (brown)/p75NTR (red) or Ki-67 (brown)/CD44 (red) was performed using Bond III automated immunostainers (Leica Biosystems). The Ki-67 labeling index was defined as the number of tumor cells that exhibited positive nuclear immunostaining divided by the total number of tumor cells.

**Human esophageal cancer cell lines and culture conditions.** Human ESCC cell lines (KYSE-30, KYSE-140, KYSE-150, KYSE-220, KYSE-510, KYSE-520 and KYSE-790) were established by Shimada et al. and cultured in Ham’s F12/Roswell Park Memorial Institute (RPMI)-1640 medium (Wako, Osaka, Japan) supplemented with 2% fetal calf serum (FCS) (Gibco, Grad Island, NY, USA), according to a previously reported method (10).

**Analysis of cancer stem cell surface antigens and cell sorting.** The surface antigen markers of the samples were analyzed using a FACSCant II flow cytometer and were sorted using a FACSaria II cell sorter and BD FACSDiva software (both from BD Biosciences, San Jose, CA, USA). Fluorescein isothiocyanate (FITC) or allophycocyanin (APC)-conjugated monoclonal mouse anti-human CD44 (clone BDH105 and p75NTR (clone ME20.4-1.H4) antibodies were purchased from Miltenyi Biotec (Bergisch Gladbach, Germany). FITC-conjugated monoclonal mouse anti-human CD90 (clone 5E-10) antibodies were obtained from BD Biosciences. FITC-conjugated antibodies were used to detect CSC surface markers. Isotype-matched APC- or FITC-conjugated antibodies (Miltenyi Biotec) were used as controls. Cultured cells were washed once with phosphate-buffered saline (PBS) (-), before being dissociated from the culture plates using 0.25% trypsin-ethylenediaminetetraacetic acid (Invitrogen, Carlsbad, CA, USA) and centrifuged. Single cells were resuspended in PBS (-) containing 2% FBS, and then the FITC-conjugated antibody or the isotype control antibody was added, before the cells were incubated at 4°C for 30 min. After being washed twice with PBS (-) containing 2% FBS, the cells were resuspended in PBS (-) containing 2% FBS. The cells were also stained with 7-aminoactinomycin D (7-AAD; Bio-Rad Laboratories, Richmond, CA, USA) to exclude dead cells. The samples were analyzed and sorted using a flow cytometer. We classified cells as positive when >1% of their cells were stained.

The KYSE-30 cells were sorted into p75NTR-positive/CD44-negative, p75NTR-negative/CD44-positive, and p75NTR-negative/CD44-negative fractions. The KYSE-140 cells were sorted into p75NTR-positive/CD90-positive, p75NTR-negative/CD90-negative, p75NTR-negative/CD90-positive and p75NTR-negative/CD90-negative fractions.

**RNA extraction, cDNA synthesis and real-time PCR.** Total RNA was extracted using the TRizol reagent (Invitrogen, Carlsbad, CA, USA). cDNA was synthesized using the PrimeScript First Strand cDNA synthesis kit (Takara Inc., Kyoto, Japan) and used for the quantitative polymerase chain reaction (qPCR) analysis. The cDNA samples were amplified in an Mx3000P real-time qPCR system (Agilent Technologies, Santa Clara, CA, USA) using SYBR Premix Ex Taq II (Takara Inc.), as described previously (11). Each mRNA expression level was normalized to that of the reference gene GAPDH. The primer sequences are shown in Table I. The expression level of each mRNA molecule was evaluated using the ΔΔCt method.

**Cell cycle analysis.** The cell cycle analysis was performed using the CycleTest Plus DNA reagent kit (Becton-Dickinson Inc., San Jose, CA, USA) as recommended by the supplier and a FACSCant II, and the resultant data were analyzed with the software FCS4 Express Cytometry (Becton-Dickinson Inc.).

<table>
<thead>
<tr>
<th>Gene product</th>
<th>Forward and reverse primers (5′-3′)</th>
<th>Size (bp)</th>
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| Nanog | F: ATGCCCTCACACGGAGACTGT  
R: AAGTGAGTTGTTGGCCTTTTG | 83 |
| p63 | F: CAGACCTTGGCAATCATCC  
R: CAGCATGTTGCTTCTTTCAGC | 220 |
| Bmi-1 | F: CCACCTGATGTGTTGTGCTTTTG  
R: TTCATGATGCTCGTCTTGT | 162 |
| Involucrin | F: TGTTCCTCCTCCAGTGCAATCCC  
R: ATTCTCTCAGTGTCCGCCAGTG | 227 |
| E-cadherin | F: GTCTGTCTAGGAGGTGTCT  
R: TACGAGTCTGCTCTTC | 370 |
| N-cadherin | F: AGCACAACCTTAACAGGGAGGT  
R: GGCAAGTGATGGAGGAGGATC | 136 |
| Fibronectin | F: AGGAAGCAGGTTTTAATGT  
R: AGAGCCCTCAATGATGGTACC | 106 |
| DPD | F: TCAAGCAGACGACTTGTGCTGT  
R: CATACATTCACAAAGTCAGACC | 205 |
| ERCC-1 | F: GCTCCCGTACCACACACCT  
R: TCTCTCTGTGTAGCAGC | 313 |
| GAPDH | F: ACCAAGTCCATGCGACCACAC  
R: TCCACCACCTGTGTGCTTGTA | 452 |

F, forward; R, reverse.
Anticancer drug resistance assay. Each cell population was cultured in Dulbecco's modified Eagle's medium/Ham's F-12 containing 5% FCS at a density of 3,200 cells/well in a 96-well plate (Thermo Scientific, Yokohama, Japan) under a humidified atmosphere of 5% CO₂ at 37°C. After being allowed to adhere overnight, the cells were treated with various concentrations of cisplatin (Wako, Osaka, Japan) or were left untreated (control). The medium in each well was changed at 2 days after the initial treatment. Cell viability was determined at 3 days after the initial treatment using the standard 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay (Trevigen, Gaithersburg, MD, USA) in accordance with the manufacturer's instructions, and the optical absorbance of the supernatant in each well at a wavelength of 595 nm was measured using a FilterMax F5 plate reader (Molecular Devices, Tokyo, Japan). Cell viability was defined as the relative absorbance of the control cells versus that of the treated cells. All experiments were performed in triplicate.

Colony formation assay. After being sorted with a FACSAria II, 1,000 cells from each KYSE-30 or KYSE-140 subset were plated in 60-mm tissue culture dishes (Thermo Scientific). After the cells had been cultured for 14 days, any colonies were stained with crystal violet (0.5% crystal violet dissolved in 20% methanol). The numbers of colonies of >3 mm in diameter were counted.

Tumorigenicity assay in nude mice or NOD/SCID mice. This part of the study protocol was approved by and conducted in accordance with the Committee of the Use of Live Animals in Teaching and Research at the University of Toyama. Five- to 6-week-old mice were used for the tumorigenicity assay; athymic nude mice (BALB/CAN.Cg-Foxn1<−/−>CrCrlj) were used for the experiments involving KYSE-30 cells and NOD/SCID mice (NOD.CB17-Prdk<−/−>/Crlj) were used for those involving KYSE-140 cells. All mice were purchased from Charles River Laboratories (Yokohama, Japan). After being sorted, 1,000 to 30,000 KYSE cells were subcutaneously injected into the bilateral lumbar regions of the mice. After 4 weeks, the mice were sacrificed, and their subcutaneous tumors were fixed with 10% buffered formalin and embedded in paraffin, before being subjected to immunohistochemical staining.

In animals that were injected with cancer cells, but did not exhibit any signs of a tumor burden, the injection sites were opened up to confirm that no tumor had developed.

Statistical analysis. Statistical analyses were performed using JMP v.11 (SAS Institute Inc., Cary, NC, USA). The Chi-square test and Fisher's exact test were used for the statistical analyses, and p-values of <0.05 were considered to be statistically significant.

Results

Expression of p75NTR, CD44 and CD90 in the ESCC specimens. We first assessed the expression of p75NTR, CD44 and CD90 in 56 primary ESCC tumors by immunohistologically staining the tumors using a tissue microarray (Fig. 1A). p75NTR was expressed in 19 of the 56 (33.9%) tumors, in which the first few layers nearest to the tumor's infiltrative margin exhibited positive staining. CD44 was expressed in 31 of the 56 (55.4%) tumors and demonstrated a diffuse distribution. On the other hand, CD90 was not expressed in any of the 56 ESCC tumors despite the fact that it was detected in hepatocellular carcinoma tissue using the same procedure (data not shown). p75NTR expression was not correlated with any of the examined clinicopathological factors, such as gender,
age, tumor site, or pTNM stage (Table II), while CD44 expression was correlated with the depth of invasion and lymph node metastasis (Table II). Both p75NTR and CD44 were expressed in 12 of 56 (21.4%) cases, although only some of the diffusely distributed CD44-positive cells were p75NTR-positive. The expression of p75NTR or CD44 alone was observed in 7 of the 56 (12.5%) and 19 of the 56 (33.9%) cases, respectively. Neither p75NTR nor CD44 was detected in 13 of the 56 (23.2%) cases.

An analysis of the Ki-67 labeling index based on double immunostaining revealed that the majority of p75NTR-positive cells were in the resting phase of the cell cycle, while most of the CD44-positive cells were actively proliferating (Fig. 1B). The whole cells, the p75NTR-positive cells, and the CD44-positive cells had mean Ki-67 labeling indices of 0.407, 0.155 and 0.446, respectively (Fig. 1C).

Expression of p75NTR, CD44 and CD90 in ESCC cell lines. To identify cell subsets based on the combined expression of CSC markers in ESCC, we examined the expression of p75NTR, CD44 and CD90 in 10 ESCC cell lines (KYSE-30, KYSE-70, KYSE-140, KYSE-150, KYSE-180, KYSE-220, KYSE-450, KYSE-510, KYSE-520 and KYSE-790) using flow cytometry (Fig. 2A and Table III). p75NTR-positive cells were detected in 6 of the 10 (60.0%) cell lines, in which the proportion of positive cells ranged from 6.4 to 50.4%. CD44-positive cells were detected in 9 of the 10 (90.0%) cell lines, in which the proportion of positive cells ranged from 2.4 to 97.8%. CD90-positive cells were detected in only 1 of the 10 (10.0%) cell lines (positive proportion, 50.9%). p75NTR and CD44, or p75NTR and CD90, were co-expressed in 5 and 1 of the 10 cell lines, respectively (Table III). Epithelial cell adhesion molecule was expressed in all of the examined ESCC cell lines, in which the proportion of positive cells ranged from 97.5 to 99.9% (data not shown). Based on the expression patterns of p75NTR, CD44 and CD90 displayed by the 10 cell lines (Table III), we selected KYSE-30 and KYSE-140 cells for the dual-color flow cytometric analysis. Among the KYSE-30 cells, which expressed p75NTR and/or CD44, p75NTR-positive/CD44-positive cells, p75NTR-positive/CD44-negative cells, p75NTR-negative/CD44-positive cells and p75NTR-negative/CD44-negative cells accounted for 1.1, 14.9, 23.2 and 60.8% of the cells, respectively (Fig. 2B).
Among the KYSE-140 cells, which expressed p75NTR and/or CD90, p75NTR-positive/CD90-positive cells, p75NTR-positive/CD90-negative cells, p75NTR-negative/CD90-positive cells and p75NTR-negative/CD90-negative cells accounted for 20.4, 12.6, 32.3 and 35.2% cells, respectively (Fig. 2B).

Expression of CSC-related genes in fractionated cell subsets. We compared the expression patterns of stem cell-, keratinocyte differentiation- and epithelial-mesenchymal transition (EMT)-related genes in each fraction using real-time PCR. Among the KYSE-30 cells, Nanog, p63 and Bmi-1 were expressed at significantly higher levels in the p75NTR-positive/CD44-negative fraction than in the p75NTR-negative/CD44-positive or p75NTR-negative/CD44-negative fraction (Fig. 3A). In addition, the expression of involucrin was significantly lower in the p75NTR-positive/CD44-negative fraction than in the p75NTR-negative/CD44-positive or p75NTR-negative/CD44-negative fraction (Fig. 3A).

The p75NTR-positive/CD44-negative and p75NTR-negative/CD44-positive fractions demonstrated significantly lower E-cadherin expression than the p75NTR-negative/CD44-negative fraction (Fig. 3C). In addition, the p75NTR-positive/CD44-negative fraction displayed significantly higher N-cadherin and fibronectin expression than the p75NTR-negative/CD44-positive and p75NTR-negative/CD44-negative fractions (Fig. 3C).

Among the KYSE-140 cells, Nanog, p63 and Bmi-1 were expressed at significantly higher levels in the p75NTR-positive/CD44-negative fraction than in the p75NTR-negative/CD44-positive or p75NTR-negative/CD44-negative fraction (Fig. 3A). In addition, the expression of involucrin was significantly lower in the p75NTR-positive/CD44-negative fraction than in the p75NTR-negative/CD44-positive or p75NTR-negative/CD44-negative fraction (Fig. 3A).

The p75NTR-positive/CD44-negative and p75NTR-negative/CD44-positive fractions demonstrated significantly lower E-cadherin expression than the p75NTR-negative/CD44-negative fraction (Fig. 3C). In addition, the p75NTR-positive/CD44-negative fraction displayed significantly higher N-cadherin and fibronectin expression than the p75NTR-negative/CD44-positive and p75NTR-negative/CD44-negative fractions (Fig. 3C).

Among the KYSE-140 cells, Nanog, p63 and Bmi-1 were expressed at significantly higher levels in the p75NTR-positive/CD90-positive fraction than in the p75NTR-negative/CD90-positive or p75NTR-negative/CD90-negative fraction (Fig. 3B). In addition, the expression of involucrin was significantly lower in the p75NTR-positive/CD90-negative fraction than in the p75NTR-negative/CD90-positive or p75NTR-negative/CD90-negative fraction (Fig. 3B).

The p75NTR-negative/CD90-negative fraction exhibited significantly higher involucrin expression than the other fractions (Fig. 3B). Significantly greater E-cadherin expression was observed in the p75NTR-negative/CD90-negative fraction than in the p75NTR-positive/CD90-positive, p75NTR-positive/CD90-negative or p75NTR-negative/CD90-positive fraction (Fig. 3D). The p75NTR-positive/CD90-positive fraction exhibited significantly higher N-cadherin and fibronectin expression than the other fractions (Fig. 3D).
Cell cycle analysis. Flow cytometric cell cycle analysis revealed that the p75NTR-negative/CD44-positive KYSE-30 cells were a mitotically active subpopulation and exhibited the lowest proportion of cells in the G0/G1 phase. On the other hand, the p75NTR-positive/CD44-negative and p75NTR-negative/CD44-negative fractions were relatively quiescent cell fractions, in which >67% of the cells were in the G0/G1 phase (Fig. 4A). Among the KYSE-140 cells, the p75NTR-positive/CD90-positive fraction contained the most actively proliferating cells, while the p75NTR-positive/CD90-negative fraction was composed of quiescent cells, 42.6% of which were in the G0/G1 phase (Fig. 4B).

Comparison of drug-resistant ability. We compared the drug resistance of each cell population using the MTT assay. Among the KYSE-30 cells, the p75NTR-negative/CD44-positive cells exhibited significantly lower viability than the p75NTR-positive/CD44-negative cells after being treated with CDDP (Fig. 4C). Among the KYSE-140 cells, the p75NTR-negative/CD90-negative cells displayed significantly lower viability than the p75NTR-positive or CD90-positive cells after being treated with CDDP (Fig. 4C).

In vitro colony formation ability. The ability of the KYSE-30 cells to form colonies exhibited the following
order: p75NTR-positive/CD44-negative > p75NTR-negative/CD44-positive > p75NTR-negative/CD44-negative (Fig. 5A). As for the KYSE-140 cells, their ability to form colonies displayed the following order: p75NTR-positive/CD90-negative > p75NTR-positive/CD90-positive > p75NTR-negative/CD90-positive > p75NTR-negative/CD90-negative (Fig. 5A).

**In vivo tumor formation.** Among the KYSE-30 cells, the subcutaneous injection of as few as 3x10³ p75NTR-positive/CD44-negative cells or p75NTR-negative/CD44-positive cells into nude mice resulted in the development of tumors at 4/4 (100%) and 2/4 (50%) of the injection sites, respectively, at 4 weeks after the injection procedure. In contrast, the injection of 1x10⁴ p75NTR-negative/CD44-negative cells did not result in tumor formation (Table IV).

The tumors that developed after the injection of 1x10⁴ p75NTR-positive/CD44-negative cells weighed significantly more than those derived from the p75NTR-negative/CD44-positive cells (Fig. 5B). Histological examinations
Figure 4. The cell cycle distribution and chemoresistance of KYSE-30 and KYSE-140 cell subsets. (A and B) Approximately 2x10⁴ live cells were analyzed using the CycleTest Plus DNA reagent kit and a FACSCant II. The results were then analyzed using FCS4 express cytometry software. The proportion of cells in each phase of the cell cycle is shown in each panel. The error bars represent the standard error of the mean. (C) Fractionated KYSE-30 and KYSE-140 cells in 96-well plates (3,200 cells/well) were treated with cisplatin (25, 50, 100 or 200 µM) (or left untreated) for 3 days, and then their viability was determined using the MTT assay.
of the xenograft tumors demonstrated that their phenotypes resembled those of primary ESCC tumors (Fig. 5C). p75NTR was expressed in a small number of the cancer cells in the first few layers nearest to the tumor infiltrating margins. CD44 was diffusely expressed throughout the tumors, except in the cells residing at their margins. A detailed examination of 4 µm-thick serial sections detected p75NTR-positive/CD44-negative cells at the tumor infiltrating margins. Analyses of the Ki-67 labeling index based on double immunostaining revealed that 86.9% of the p75NTR-positive cells were in the resting phase of the cell cycle, while 74.8% of the CD44-positive cells were actively proliferating.

Among the KYSE-140 cells, the subcutaneous injection of as few as 3x10³ p75NTR-positive/CD90-positive cells or p75NTR-positive/CD90-negative cells into NOD/SCID mice resulted in tumors forming at 2/4 (50%) and 1/4 (25%) of the injection sites, respectively, at 4 weeks after the injection procedure (Table IV). On the other hand, the injection of 1x10⁴ p75NTR-negative/CD90-positive cells resulted in the establishment of tumors at 2/4 (50%) of the injection sites. In contrast, even the injection of 3x10⁴ p75NTR-negative/CD90-negative cells did not result in tumor formation (Table IV). The tumors that developed after the injection of 1x10⁴ p75NTR-positive/CD90-positive
cells weighed significantly more than those derived from the p75NTR-positive/CD90-negative (p<0.001) or p75NTR-negative/CD90-positive cells (p<0.001) (Fig. 5B). In addition, the tumors derived from the p75NTR-positive/CD90-negative cells were larger than those derived from the p75NTR-negative/CD90-positive cells (p=0.002) (Fig. 5B). A histological examination of the xenograft tumors showed that p75NTR was expressed in the cancer cells at their infiltrating margins. An analysis of the Ki-67 labeling index based on double immunostaining revealed that 59.6% of the p75NTR-positive cells were in the resting phase of the cell cycle, while 80.4% of the tumor cells were actively proliferating. No CD90 expression was detected in any of the cancer cells in the xenograft tumors (p=0.017) (Fig. 5D).

Discussion

Our immunohistochemical investigation of surgical ESCC specimens for previously reported CSC markers, such as p75NTR, CD44 and CD90, revealed that the expression of p75NTR alone, the expression of CD44 alone, and the expression of both p75NTR and CD44 were detected in 21.4, 25.0 and 30.4% of the tumors, respectively. In addition, none of the reported CSC markers were detected in 23.2% of the tumors, demonstrating the heterogeneity of the cell surface marker expression of ESCC tumors, as has been shown for various other types of tumors (12).

p75NTR was expressed at the tumor infiltrating margins, while CD44 was diffusely expressed throughout the tumors, which is compatible with the findings of previous studies (5,7). Our double immunostaining-based analysis of the Ki-67 labeling index revealed that most of the p75NTR-positive cells were in a mitotically quiescent state, while the majority of the CD44-positive cells were actively proliferating. These findings indicated that quiescent CSC can be found within...
ESCC and that it may be possible to identify different CSC populations based on the combination of cell surface markers that they express.

No CD90 expression was detected in any of the tumor cells in the ESCC specimens. However, we observed positive CD90 staining in the stromal cells surrounding the cancer cells in all of the ESCC specimens, which is compatible with the findings of a previous report, in which CD90 expression was detected in the stromal cells of head and neck squamous cell carcinoma (13). We also observed positive CD90 staining in the tumor cells of hepatocellular carcinoma specimens (positive controls) (7). Although a previous study detected CD90 expression in ESCC specimens by qPCR and isolated CD90-positive cells from fresh cancer tissues by flow cytometry (8), no previous studies have detected CD90 expression in tumor cells from ESCC specimens.

p75NTR, CD44 and CD90 expression were detected at various cell frequencies in 6, 9 and 1 of the 10 examined KYSE cell lines, respectively, further indicating the heterogeneity of the cell surface marker expression seen in ESCC.

Contrary to the finding we obtained during the immunohistochemical examination of the ESCC specimens, CD90 was expressed in 50.9% of the KYSE-140 cells. It has been reported that CD90 expression can be induced by various biological agents, such as thymopoietin, prostaglandins, interleukin-1, tumor necrosis factor-α, vascular endothelial growth factor, and nerve growth factor (NGF) (14,15). Combining the above-mentioned finding with the fact that an NGF autocrine loop was detected in KYSE cells (16), it is suggested that CD90 expression can be induced in certain cell culture conditions. Among the KYSE-30 cells, the p75NTR-positive CD44-negative fraction exhibited strong stem cell-related gene expression, weak keratinocyte differentiation marker expression, anticaner drug resistance, in vitro colony formation and in vivo tumorigenicity; i.e., they displayed a CSC phenotype. Furthermore, the majority of the p75NTR-positive CD44-negative KYSE-30 cells were in a mitotically quiescent state, indicating that a quiescent CSC population is maintained even in actively proliferating cells, whereas, the CD44-positive cells were actively proliferating and exhibited a less marked CSC phenotype. Among the KYSE-140 cells, the p75NTR-positive cells displayed a CSC phenotype; i.e., strong stem cell-related gene expression, weak keratinocyte differentiation marker expression, anticaner drug resistance, in vitro colony formation, and in vivo tumorigenicity, regardless of whether they expressed CD90.

Furthermore, an analysis of the KYSE-140 cells based on their p75NTR/CD90 expression patterns identified two distinct subpopulations in terms of their cell cycle status. The p75NTR-positive/CD90-negative subset was mitotically quiescent cells while the p75NTR-positive/CD90-positive subset was actively proliferating cells, further indicating the existence of a quiescent CSC population in ESCC.

Recent studies have demonstrated that dormant CSC are even present in solid tumors, such as melanoma (17) and ovarian (18), breast (2) and pancreatic tumors (19). In addition, it was reported that these cells enhance chemoresistance, invasiveness and metastasis, as well as the risk of a late relapse after curative surgery (1,2,12,20).

Several studies of leukemia have examined the molecular mechanisms that regulate stem cell quiescence. These studies suggested that inducing leukemia cells to enter the cell cycle may enhance their chemosensitivity and that it is important to identify molecules that will enable us to directly target quiescent CSC (21,22).

Recently it was reported that the NGF/proNGF/p75NTR axis plays a critical role in regulating the self-renewal of quiescent CSC in breast cancer (23). Combining our results with the findings of a previous report in which NGF overexpression and its autocrine loop was shown to enhance cell proliferation in KYSE cells (16), it is possible that p75NTR signaling also plays a role in the regulation of quiescent CSC in ESCC.

We used negative immunohistochemical staining of the proliferation marker Ki-67 and flow cytometric measurements of DNA content to detect cells in the G0/G1 phase of the cell cycle, as it was previously reported that hematopoietic stem cells exist in the quiescent state (according to dual staining using the DNA-binding dye Hoechst 33342 and anti-Ki-67 antibody) (24). Moreover, label incorporation-based assays, such as 5-bromo-2-deoxyuridine (BrdU)-based studies, have also been used to isolate putative quiescent CSC based on their dynamic cycling kinetics (25). Further studies of self-renewal, asymmetric cell division and the duration of the G0/G1 phase in the p75NTR-positive cell subset may facilitate the elucidation of the dynamic kinetics of the CSC present in ESCC.

In the present study, we found that EMT markers were strongly expressed in the p75NTR-positive/CD44-negative and p75NTR-positive/CD90-positive fractions of the KYSE-30 and KYSE-140 cells, respectively. Previous studies have demonstrated that EMT processes play a role not only in the acquisition of a migratory mesenchymal phenotype, which enhances invasiveness and metastasis, but also in the acquisition of stem cell and tumorigenic characteristics in CSC (20,26). Combining these results with the findings of a recent report in which it was demonstrated that NGF promoted EMT in the breast cancer stem cell compartment (23), it is possible that p75NTR signaling is involved in the regulation of EMT and stem cell properties. The identification of the molecular mechanisms that regulate cell stem cell properties, cell cycle status and EMT processes in the p75NTR-positive cell subset of KYSE cells may provide a basis for the development of novel treatment strategies for ESCC.

In conclusion, by examining surgical tumor specimens and cultured cell lines we found that ESCC exhibit heterogeneous expression patterns of cell surface markers such as p75NTR, CD44 and CD90. In addition, an analysis of the Ki-67 labeling indices in the tumor specimens revealed that most of the p75NTR-positive cells were in a mitotically quiescent state, while the majority of the p75NTR-negative cells were actively proliferating. Among KYSE cells, p75NTR-positive cells represent a CSC population that exhibits strong stem cell-related gene expression, weak keratinocyte differentiation marker expression, chemoresistance, in vitro colony formation, and in vivo tumorigenicity, regardless of whether they express CD44 or CD90.

Furthermore, p75NTR-positive/CD44-negative and p75NTR-positive/CD90-negative fractions KYSE cells were found to be mitotically quiescent CSC both in vitro and in vivo.
These results demonstrate that it is possible to detect and isolate quiescent CSC from ESCC, providing researchers with a target that will aid the development of novel therapeutic strategies, as well as diagnostic tools for patient selection.

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