Side population cells have the characteristics of cancer stem-like cells/cancer-initiating cells in bone sarcomas

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**BACKGROUND:** Several human cancers have been found to contain cancer stem-like cells (CSCs) having cancer-initiating ability. However, only a few reports have shown the existence of CSCs in bone and soft tissue sarcomas. In this study, we identified and characterised side population (SP) cells that showed drug-resistant features in human bone sarcoma cell lines.

**METHODS:** In seven osteosarcoma cell lines (OS2000, KIKU, NY, Huo9, HOS, U2OS and Saos2) and in one bone malignant fibrous histiocytoma (MFH) cell line (MFH2003), the frequency of SP cells was analysed. Tumourigenicity of SP cells was assessed in vivo and in vitro. Gene profiles of SP cells and other populations (main population; MP) of cells were characterised using cDNA microarrays.

**RESULTS:** SP cells were found in NY (0.31%) and MFH2003 (5.28%). SP cells of MFH2003 formed spherical colonies and re-populated into SP and MP cells. In an NOD/SCID mice xenograft model, 1 × 10^3 sorted SP cell-induced tumourigenesis. cDNA microarray analysis showed that 23 genes were upregulated in SP cells.

**CONCLUSIONS:** We showed that SP cells existed in bone sarcoma cell lines. SP cells of MFH2003 had cancer-initiating ability in vitro and in vivo. The gene profiles of SP cells could serve as candidate markers for CSCs in bone sarcomas.

**Keywords:** cancer stem-like cell; cancer-initiating cell; osteosarcoma; bone malignant fibrous histiocytoma; side population

Over the past three decades, there have been remarkable advances in the treatment of bone and soft tissue sarcomas. These include the introduction of adjuvant chemotherapy, establishment of guidelines for adequate surgical margins and the development of post-excision reconstruction (Fletcher et al., 2002; Lewis, 2007). However, the prognosis of non-responders to chemotherapy is still poor and the mechanisms of tumourigenesis of bone and soft tissue sarcomas remain to be demonstrated.

Generally, cancer masses are considered to be a complex of heterogeneous but equally malignant cell populations. However, recent stem cell research on the development of normal organs has drawn attention to the existence of a ‘cancer stem-like cell (CSC)’ counterpart, which is characterised by its self-renewal capacity, differentiation potential, and cancer-initiating ability (Visvader and Lindeman, 2008). On the basis of these characteristics, CSCs have been postulated to be responsible for driving the growth of tumours and for the recurrence of neoplasms after current therapeutic modalities are used.

Initial attempts to characterise CSCs were accomplished using cell surface molecules in acute myeloid leukaemia. Several groups that found CSCs capable of initiating leukaemia were found in the CD34^+ CD38^- fraction (Warner et al., 2004). Recently, CSCs have been isolated from several human solid tumours that have markers for putative normal stem cells, including breast cancer (CD44^+ CD24 ESA^-) (Al-Hajj et al., 2003), pancreatic cancer (CD44^+ CD24 ESA^-), CD133^+ CXCR4^- (Hermann et al., 2007; Li et al., 2007), brain cancer (CD133^-) (Singh et al., 2004), prostate cancer (CD44^-/CD133^-) (Collins et al., 2005), hepatocellular carcinoma (CD133^-) (Yin et al., 2007) and colon cancer (CD133^-) (Ricci-Vitiani et al., 2007).

On the other hand, in the analysis of haematopoietic stem cells, a sub-population that effluxes the DNA-binding dye Hoechst 33342 out of the cell membrane through an ABC-binding cassette (ABC) transporter was recognised as a stem cell population (Goodell et al., 1996; Zhou et al., 2002; Robinson et al., 2005). This cell population expressing the ABC transporter was defined as side population (SP) cells, which were distinguished from cells of the other population (main population; MP). Recent studies demonstrated that SP cells could be characterised as CSCs in primary tissues of gastrointestinal cancers (Haraguchi et al., 2006) and ovarian cancer (Szotek et al., 2006). SP cells were also shown in established tumour cell lines with different origins, such as glioma (Kondo et al., 2004), breast (Krugel et al., 2006) and thyroid cancer monoclonal cell lines (Mitsutake et al., 2007).
To date, however, distinct molecular markers on CSCs are still lacking in many cancers. Moreover, only a few reports have shown the existence of CSCs in bone and soft tissue sarcomas (Gibbs et al., 2005; Wu et al., 2007; Tirino et al., 2008). In this study, with the goal of determining specific markers of CSCs, we identified and characterised SP cells having cancer-initiating ability in osteosarcoma and bone malignant fibrous histiocytoma cell lines.

MATERIALS AND METHODS

Cell lines and culture

Seven human osteosarcoma (OS) cell lines (OS2000, KIKU, NY, Huo9, HOS, U-2OS and Saos2) and one bone human malignant fibrous histiocytoma (MFH) cell line (MFH2003) were used. OS2000, KIKU and MFH2003 were established in our laboratory (Wada et al., 1988; Nabeta et al., 2003; Tsukahara et al., 2006). The other cell lines were kindly donated by or purchased from the Japanese Collection of Research Bioresources Cell Bank (Tokyo, Japan) and from the American Type Culture Collection (Manassas, VA, USA). MFH2003 and OS2000 were cultured with Iscove’s modified Dulbecco’s Eagle’s medium (IMDM; GIBCO BRL, Grand Island, NY, USA) containing 10% FBS and the others were maintained in Dulbecco’s modified Eagle’s medium (DMEM; Sigma-Aldrich, St Louis, MO, USA) containing 10% FBS in a 5% CO₂ incubator at 37°C.

Identification of side population

Cell suspensions were labelled with dye Hoechst 33342 (Cambrex Bio Science Walkersville Inc., MD, USA) using the methods described by Goodell et al (1996) with some modifications. Briefly, cells were trypsinised and re-suspended in pre-warmed DMEM supplemented with 5% FBS at a concentration of 1 × 10⁶/ml. Hoechst33342 dye was added at a final concentration of 2.5 or 5.0 μg/ml in the presence or absence of verapamil (50 or 75 μM; Sigma-Aldrich) as an inhibitor of the ABC transporter. The cells were incubated at 37°C for 90 min with continuous shaking. At the end of the incubation, the cells were washed with ice-cold PBS with 5% FBS, centrifuged at 4°C and resuspended in ice-cold PBS containing 5% FBS. Propidium iodide (at a final concentration of 5 μg/ml; Molecular Probes–Invitrogen, Eugene, OR, USA) was used to gate viable cells. Flow cytometry and cell sorting were performed using FACS Vantage SE (BD Biosciences, Bedford, MA, USA), EPICS ALTRA (Beckman-Coulter, Fullerton, CA, USA) and FACS Aria II (BD Biosciences). The Hoechst 33342 dye was excited at 357 nm and its fluorescence was analysed using dual wave-lengths (blue, 402–446 nm; red, 650–670 nm).

RNA preparation

Total RNAs were extracted from SP cells and MP cells using the RNeasy Mini kit (Qiagen, Hilden, Germany) according to the manufacturer’s protocol. Using an Amino Allyl MessageAmp aRNA Kit Ver. 2 (Sigma-Aldrich Japan, Ishikari, Japan), amino allyl-modified aRNAs were prepared from total RNAs from SP and MP cells as previously described (Takeuchi et al., 2008).

Real-time PCR analysis

Real-time PCR was reverse transcribed using the SuperScript III reverse transcriptase enzyme (Invitrogen) according to the manufacturer’s instructions. Real-time PCR was performed with SYBR Green Real-time Core Reagent (Applied Biosystems, Foster City, CA, USA) according to the manufacturer’s instructions on an ABI Prism 7900 Sequence Detection System (Applied Biosystems). Primers were designed to generate a PCR product of ∼200 bp. The thermal cycling conditions were 94°C for 2 min, followed by 35 cycles of 15 s at 94°C, 30 s at 60°C and 30 s at 72°C. Levels of expression were normalised to the glyceraldehyde-3-phosphate dehydrogenase (G3PDH) housekeeping gene.

Spherical colony formation assay

Spherical colony formation assay was performed as described by Gibbs et al (2005) with some modifications. Briefly, cells were plated at 2000 cells per well in six-well ultra-low attachment plates (Corning Inc., Corning, NY, USA). Mesenchymal Stem Cell Basal Medium (MSCBM) and MSCBM SingleQuots (Takara Bio Inc., Ohtsu, Japan) were used for cell culture. Fresh aliquots of epidermal growth factor and basic fibroblast growth factor were added every other day. On day 14, the numbers of colonies were counted.

Gene expression profiling using cDNA microarrays

The aRNAs from SP cells were labelled with Cy5 dye and those from MP cells were labelled with Cy3 dye. The dye-labelled aRNA samples were hybridised to a 29 138-spot Human Panorama Micro Array (Sigma-Aldrich) for 16 h at 45°C. The intensities of Cy5 and Cy3 fluorescence for every gene spot on the hybridised array were measured with a GenePix 4000B scanner (Axon Instrument, Austin, TX, USA), and were analysed with GenePix Pro 5.0 software (Axon Instrument). Global normalisation of the resultant data was carried out using Excel 2004 (Microsoft, Redmond, WA, USA). As a result, 24 730 genes were available for further analysis. The average expression ratio of Cy5 to Cy3 was obtained for each gene. A dye-swap experiment (labelling SP and MP cells with Cy3 and Cy5, respectively) was also performed. An average ratio of more than 2.0, reproducible in two experiments, was determined to indicate differential upregulation in SP cells.

RESULTS

Identification of SP cells in human bone sarcoma cell lines

To identify the CSCs of bone sarcomas, we tried to detect side population (SP) cells in bone and soft tissue sarcoma cell lines. As depicted in Figure 1A, the NY and MFH2003 cell lines included 0.38 and 5.28% SP cells, respectively. In each case, the percentage of SP cells was markedly diminished by treatment with verapamil, which is an inhibitor of the pumps responsible for the exclusion of Hoechst dye, indicating that this population truly represented SP cells (Figure 1B). Therefore, MFH2003, containing the highest proportion of SP cells, was selected and further analysed.

Isolation of SP cells and their repopulation as both SP and MP cells

After excluding dead cells and cellular debris on the basis of scatter signals and propidium iodide fluorescence, MFH2003 cells were sorted into SP and MP cells. As shown in Figure 2A, the G1 gate showed the SP cells that had negative/low patterns of staining with Hoechst dye, indicating that this population truly represented SP cells. The G2 gate showed the main population cells that were positively stained with Hoechst 33342. To ascertain the
purity of sorted cells, the obtained SP (G1) cells and MP (G2) cells were re-analysed. The purity levels were 92.86 and 90.78% in the SP population and MP population, respectively. These results supported the specificity for further analysis using the resultant SP and MP cells.

To examine whether SP cells could generate both SP and MP cells, sorted SP and MP cells were further cultured in vitro. On day 14, the cells were re-stained with Hoechst 33342 and analysed by flow cytometry. We observed that SP cells re-populated both SP and MP cells. The ratio of SP cells to MP cells was still much higher than that before sorting. In contrast, SP cells were not detected in sorted MP cells.

Expression of ABCG2 mRNA, which is a marker of SP cells, was increased in SP cells (Figure 2B). These results also supported the specificity for further characterisation of SP cells, especially with regard to their cancer-initiating ability.

Spherical colony formation

We next evaluated the ability of SP and MP cells to generate spherical colonies. A total of 2000 SP and MP cells were sorted and cultured immediately under conditions of serum starvation, providing an anchorage-independent environment. On day 14, SP cells showed spherical colony formation (Figure 3A).
other hand, as shown in Figure 3B, most MP cells died and the others formed a few small colonies. We removed spherical colonies from the suspension culture and attempted again to determine whether the cells could attach to a substratum. As shown in Figure 3C, cells were seen expanding from the sphere. In Figure 3D, the number of colonies is shown, indicating clearly that, among MFH2003 cells, SP, but not MP, cells had the potential for spherical colony formation.

Cancer-initiating ability of SP and MP cells in vivo

To address the issue of whether tumourigenic activity differed between SP and MP cells, $1 \times 10^2 - 1 \times 10^5$ SP and MP cells sorted from MFH2003 were injected into NOD/SCID mice (Figure 4A). To rule out the effects of the toxicity of Hoechst, we routinely performed (i) depletion of dead cells by PI staining and (ii) a viability check using trypan-blue staining after cell sorting. Almost all MP cells were viable as SP cells. Subcutaneous tumour formation was induced by the injection of $1 \times 10^3$ SP cells (Table 1). We also observed that $1 \times 10^4$ SP cells formed a larger tumour mass than did $1 \times 10^3$ SP cells (data not shown). In contrast, at least $1 \times 10^5$ MP cells were required to give rise to a tumour. Macroscopic and microscopic findings of a tumour derived from SP cells are shown in Figure 4A and B. These results supported the hypothesis that SP cells have a high cancer-initiating ability, similar to CSCs. At 8 weeks after xenotransplantation, the frequencies of SP and MP cells in a formed tumour derived from $1 \times 10^4$ SP cells were analysed ex vivo. SP cells were hardly detected in the tumour. Most SP cells re-populated into MP cells in vivo in 8 weeks (data not shown).

Identification of upregulated genes in SP cells

Finally, we performed gene expression profiling of SP cells using cDNA microarrays. As shown in Table 2, 23 genes were found to be upregulated in SP cells, compared with MP cells. Although the functions of upregulated genes varied, these results suggested that the factors connected to DNA transcription, transport of substrates, cell proliferation and apoptosis might have a role in the cancer-initiating ability of SP cells. In addition, the increased expression of ABCG2 in SP cells confirmed the accuracy of the gene expression profiling analysis.

DISCUSSION

In this study, we showed that (i) small SP populations existed in one osteosarcoma cell line and one bone MFH cell line; (ii) SP cells derived from MFH2003 could re-populate both SP and MP cells in vitro; (iii) SP cells could form spherical colonies and re-populate into SP and MP cells; (iv) SP cells had tumourigenesis in an in vivo xenograft model; and (v) factors regarding transcription, cell proliferation and apoptosis were upregulated in SP cells.
We observed proportions of SP cells of 0.31 and 5.28% in NY and MFH2003, respectively. The proportions of SP cells we observed were similar to those in most previous reports, with 2% noted in human breast cancer cell line MCF7, 0.4% in rat G6 glioma, 1.2% in human HeLa carcinoma (Kondo et al, 2004) and 4–37% in neuroblastoma cell lines (Hirschmann-Jax et al, 2004).

The SP cells were defined by the efflux of Hoechst 33342, a cell-permeable DNA-specific bisbenzimidazole dye, through an ABC transporter. Therefore, SP cells are considered to be resistant to multi-chemotherapeutic drugs and to confer malignant phenotypes to tumours (Dean et al, 2005). Hence, the characterisation of SP cells might be a useful tool for analysis of CSCs, especially when specific CSC surface markers are unknown.

We found that SP cells could re-populate both SP cells and MP cells in vitro. These results suggested that SP cells were capable of self-renewal and also generated MP cells by asymmetric division. This indicated that a tumour hierarchy might exist in bone MFH.

Table 1  Tumorigenesis of SP and MP cells in NOD/SCID mice

| Cell number for injection | SP cells | MP cells |
|--------------------------|----------|----------|
| $1 \times 10^2$          | 0/5      | 0/5      |
| $1 \times 10^3$          | 1/5      | 0/5      |
| $1 \times 10^4$          | 2/5      | 0/5      |
| $1 \times 10^5$          | 1/2      | 1/2      |

SP and MP cells were isolated separately and injected into the backs of the subcutaneous space of NOD/SCID mice. Tumour formation was observed for 12 weeks after injection.
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Sarcoma (Gibbs candidates for CSC-specific markers in bone and soft tissue tumour-infiltrating lymphocytes. Kimura effective isolation of CSCs from bone and soft tissue sarcomas profile of SP cells might help to expand the possibility of an IRX3 Iroquois-related homeobox 3 BC023667 Regulation of transcription 3.3 2.9 NR4A2 Nuclear receptor subfamily 4, group A, member 2 NM_006186 Regulation of transcription 2.5 4.1 MCL1 Myeloid cell leukaemia sequence 1 NM_021960 Apoptosis 2.1 3.9 APOE Apolipoprotein E NM_000041 Lipid transport 2.1 3.8 NRA42 Nuclear receptor subfamily 4, group A, member 2 NM_006186 Regulation of transcription 2.5 4.1 IRX3 Iroquois-related homeobox 3 BC023667 Regulation of transcription 3.3 2.9 GPER G-protein coupled receptor protein signaling 2.1 5.0 NLRP12 NLR family, pyrin domain containing 12 NM_144687 Apoptosis 2.1 2.3 PTPN PTPN11 tyrosine phosphatase 1 NM_002902 Neurogenesis 2.0 2.6 ABCG2 ATP-binding cassette, sub-family G, member 2 NM_004827 Transport 2.2 2.9 APOL1 Apolipoprotein L NM_145343 Lipid transport 2.2 2.6 MDFI MyoD family inhibitor NM_005586 Unknown 2.8 3.1 PRSS15 PrSS15 protease, serine, 15 NM_004793 ATP-dependent proteolysis 2.1 2.5 MSX1 Mouse homeobox homolog 1 NM_002348 Regulation of transcription 2.1 2.8 LDLR Low density lipoprotein receptor NM_000527 Cholesterol metabolism 2.1 3.1 LMNA Lamin A/C NM_00170707 Cellular morphogenesis 2.1 2.9 MVK Mevalonate kinase BC016140 Cholesterol biosynthesis 2.2 2.1

Table 2 List of genes upregulated in SP cells of MFH2003

| Gene symbol | Gene name | Accession no. | Gene ontology | Cy5/Cy3 | Cy3/Cy5 |
|-------------|-----------|---------------|---------------|--------|--------|
| ANKRD11 | Ankyrin repeat domain 11 | NM_013275 | Electron transport | 2.1 | 2.7 |
| SLC2A4 | Solute carrier family 2, member 4 | NM_001042 | Carbohydrate transport | 2.2 | 2.5 |
| KIAA1440 | KIAA1440 | AB037861 | Unknown | 2.2 | 3.3 |
| SURF6 | Surfeit 6 | NM_006753 | Ribosomogenesis | 2.3 | 2.8 |
| VPF | Vascular permeability factor | M27281 | Cell proliferation | 3.0 | 3.5 |
| C2orf14 | Chromosome 2 open reading frame 14 | NM_012469 | RNA processing | 2.1 | 3.6 |
| PHLDA3 | Pleckstrin homology-like domain, family A, member 3 | NM_012396 | Physiological processes | 2.1 | 2.4 |
| ZNF19 | Zinc finger protein 19 | NM_152907 | Regulation of transcription | 2.8 | 2.4 |
| MCL1 | Myeloid cell leukaemia sequence 1 | NM_021960 | Apoptosis | 2.9 | 3.9 |
| APOE | Apolipoprotein E | NM_000041 | Lipid transport | 2.1 | 3.8 |
| NRA42 | Nuclear receptor subfamily 4, group A, member 2 | NM_006186 | Regulation of transcription | 2.5 | 4.1 |
| IRX3 | Iroquois-related homeobox 3 | BC023667 | Regulation of transcription | 3.3 | 2.9 |
| MSX1 | Mouse homeobox homolog 1 | NM_002348 | Cholesterol metabolism | 2.1 | 3.1 |
| PTPN | PTPN11 tyrosine phosphatase 1 | NM_002902 | Apoptosis | 2.1 | 2.3 |
| ABCG2 | ATP-binding cassette, sub-family G, member 2 | NM_004827 | Cholesterol biosynthesis | 2.0 | 2.6 |
| APOL1 | Apolipoprotein L | NM_145343 | Transport | 2.2 | 2.9 |
| MDFI | MyoD family inhibitor | NM_005586 | Unknown | 2.8 | 3.1 |
| PSS15 | PSS15 protease, serine, 15 | NM_004793 | ATP-dependent proteolysis | 2.1 | 2.5 |
| MSX1 | Mouse homeobox homolog 1 | NM_002348 | Regulation of transcription | 2.1 | 2.8 |
| LDLR | Low density lipoprotein receptor | NM_000527 | Cholesterol metabolism | 2.1 | 3.1 |
| LMNA | Lamin A/C | NM_00170707 | Cellular morphogenesis | 2.1 | 2.9 |
| MVK | Mevalonate kinase | BC016140 | Cholesterol biosynthesis | 2.2 | 2.1 |

Genes showing the ratio more than 2.0, which were reproducible in two experiments, were listed.

Normal stem cells (Kondo et al., 2004; Singh et al., 2004). On the other hand, we also observed that most SP cells xenotransplanted in NOD/SCID mice re-populated into MP cells in vivo. SP cells might hardly be maintained in vivo for long time, that is, more than 8 weeks. In other words, the niche of the mouse model might not be adequate for the maintenance of SP cells derived from bone sarcoma cell lines, such as MFH2003.

The ability of SP cells to generate spherical colonies was higher than that of MP cells. This is consistent with previous studies (Patrawala et al., 2005; Mitsutake et al., 2007). We recognised that the difference was not a consequence of longer retention of Hoechst dye in MP cells, because MP cells were viable after staining with the dye, followed by sorting and maintenance in a culture medium with FBS. However, we cannot completely rule out the possibility that the difference was due to some effect of the culture medium in FBS. We could detect a higher tumourigenic potential of SP cells than of MP cells in vivo using a NOD/SCID xenograft model. In the field of bone and soft tissue sarcoma, normal stem cells, only Wu et al. (2007) succeeded in showing the in vivo cancer-initiating ability of CSCs derived from bone and soft tissue sarcomas, using SP cells isolated from fresh primary tumour tissues. The ability to consistently isolate MFH2003-derived SP cells allowed us to conduct SP cell-specific gene profiling. Moreover, it might become possible to identify CEL-defined CSC-specific tumour antigens for immunotherapy targeting CSCs. Thus far, we have been trying to identify CEL-defined tumour antigens by forward and reverse immunological approaches and have carried out antigen peptide vaccination trials in bone and soft tissue sarcomas (Sato et al., 2002; Ida et al., 2004; Tsukahara et al., 2004, 2008a, b, 2009; Kagawuchi et al., 2005; Kimura et al., 2008). Currently, we are trying to establish autologous CTL clones recognising SP cells of MFH2003 from tumour-infiltrating lymphocytes.

Thus far, only Oct3/4, Nanog and CD133 were reported to be candidates for CSC-specific markers in bone and soft tissue sarcoma (Gibbs et al., 2005; Tirino et al., 2008). Therefore, the gene profile of SP cells might help to expand the possibility of an effective isolation of CSCs from bone and soft tissue sarcomas using these specific markers. In the current gene expression profiling, 23 genes with various functions were upregulated in SP cells. Among them, eight genes (VPF, c20orf14, MCL1, NRA42, IRX3, NLRP12, PTN and LMNA) might be considered to be potential tumourigenic factors in malignancies. VPF, generally known as vascular endothelial growth factor, regulates vascular permeability, angiogenesis, cell migration and apoptosis in tumours (Nagy et al., 2008). C20orf14 is upregulated in lymphoma (Su et al., 2008) and HPV16/18-positive cervical cancer (Vazquez-Ortiz et al., 2007). MCL1 is a member of the B-cell lymphoma (BCL) family. MCL-1 negatively regulates pro-apoptotic factors (Bak and Bax) (Chen et al., 2007) and accelerates leukaemogenesis (Beverly and Varmus, 2009). NAA42 belongs to the steroid nuclear hormone receptor superfamily and has a role in cell transformation in cervical cancer (Ke et al., 2004). IRX3 is epigenetically inactivated by methylation in CpG islands in brain tumours (Ordway et al., 2006) and prostate cancer (Morey et al., 2006). NLRP12, also known as RNO2/mornarch-1, is reported to activate inflammation in humans (Ye et al., 2008). PTN is an angiogenic factor that stimulates tumour-associated vascular formation in many malignancies. (Perez-Pinera et al., 2008). LMNA is reported to encode lamin A, which is a putative colonic epithelial stem cell marker and is also a prognostic factor in colorectal cancer (Willis et al., 2008). On the other hand, four genes (ANKRD11, PHLDA3, APOL1 and MSX1) are known as tumour-suppressor factors. ANKRD11 is a p53-interacting protein and activates the transcription of p53 in breast cancer. PHLDA3 is a positive regulator of Fast-dependent death signalling, related to cisplatin-mediated apoptosis (Kerley-Hamilton et al., 2005). APOL1 is classically thought to be involved in lipid transport and metabolism and has rarely been characterised with regard to cell survival. Although the structure of APOL1 is similar to that of the anti-apoptotic proteins of the Bcl-2 family (Vanhollebeke and Pays, 2006), it can induce autophagic cell death (Wan et al., 2008). MSX1, a homeobox gene important for embryonic neural crest development, can induce the inhibition of tumour-initiating ability in soft agar in vitro. Taken together, the gene expression profiling in SP cells derived from MFH2003 containing various tumour-proliferative and tumour-suppressive factors might indicate the complexity of maintaining the...
characteristics of SP cells as tumour initiators. However, considering the characteristics of SP cells for cancer-initiating ability, the apoptosis-related molecules among these genes (MCL-1, ANKRDL1, PHLD3A and APOL1) might have roles in the proliferation of SP cells. Moreover, these molecules could be candidates for specific markers and, in addition, molecular therapeutic targets.

In conclusion, we identified SP cells in established human bone sarcoma cell lines. Moreover, we demonstrated that bone MHF-derived SP cells can re-populate both SP and MP cells and have cancer-initiating ability in vitro and in vivo. These findings supported the idea that bone sarcomas might contain a certain population of CSCs. Gene profiling of SP cells could serve to elucidate candidate markers for specific bone and soft tissue sarcomas.

Thus, further studies for the characterisation of CSCs in human bone and soft tissue sarcomas might contribute to the elucidation of the mechanisms of tumourigenesis and to the establishment of novel therapeutic strategies.

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