Intratibial injection of patient-derived tumor cells from giant cell tumor of bone elicits osteolytic reaction in nude mouse

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Abstract. There have been various reports in the literature of an in vivo model for giant cell tumor of bone (GCTB). However, few suitable animal models of GCTB have been established, due to the fact that GCTB contains three histologically different cell types. To the best of our knowledge, injection of patient-derived GCTB cells into bone environment has not been reported until now. In the present study, the biological behavior of GCTB cells in nude mice was investigated through intratibial injection of patient-derived GCTB cells. Patient-derived GCTB cells were obtained from 5 patients who had not undergone chemo- and radiotherapy. Once isolated, the cell suspension was injected into the tibias of nude mice. The growth process was monitored by weekly observation and photographic documentation using X-ray. Four months after injection, nude mice were sacrificed and the injected tibial samples were fixed, and further analyzed using micro-computed tomography (micro-CT), standard histology, tartrate-resistant acid phosphatase (TRAP) staining and mitochondrial immunofluorescence staining. X‑ray, micro‑CT and standard histology revealed osteolytic destruction in the proximal end of the tibia. TRAP staining identified TRAP-positive, osteoclast-like cells distributed in the bone marrow interface of the lesion area. Anti-human mitochondrial immunofluorescence staining confirmed that the surviving cells in the osteolytic destruction were of human GCTB cell origin. These findings indicate that intratibial injection of patient-derived GCTB cells may elicit osteolytic destruction in nude mice. The results of the current study present a novel animal model for GCTB, opening new perspectives to investigate this disease and develop therapeutic agents.

Introduction

Giant cell tumor of bone (GCTB) is an aggressive osteolytic tumor that typically originates in the epimetaphyseal region of a long bone and frequently occurs in the distal end of the femur or the proximal end of the tibia (1,2). GCTB is clinically defined as a benign bone tumor, but is characterized by locally aggressive growth and usually leads to an extensive bone lesion. Radiographically, the lesion of GCTB is purely lytic, exhibiting geographic bone destruction (3). Although it rarely causes mortality, the recurrence rate for GCTB has been reported as 18-50% (4-8).

Histologically, GCTB is composed of three major cell types: Multinucleated, osteoclast-like giant cells; monocytic round-shaped, macrophage-like cells; and spindle-shaped, fibroblast-like stromal cells (9-11). The stromal cells are considered to be the neoplastic component of GCTB as they are the only proliferating cell component in long-term cultures (12,13) and express oncogenes (14-16). The macrophage-like cells, which are recruited by stromal cells, are considered to be osteoclast precursors, and are able to fuse into multinucleated osteoclast-like giant cells, with the latter eventually causing aggressive bone resorption and skeletal destruction (17-19). As stable cell lines of the multinucleated giant cells and monocyte-derived macrophages have not been established, individual GCTB cell components cannot be maintained in culture. As such, it has not been possible to determine the efficacy of antitumor agents on GCTB in vitro (20,21).

Although several in vivo GCTB models are available, studies on this tumor are restricted as the development of the in vivo model is incomplete, associated with a short survival time or does not induce osteolytic lesion (22-24). Stromal cells injected subcutaneously into immunocompromised mice do not produce giant cells (22,24,25). Furthermore, tumor tissues grown on chick chorioallantoic membranes do not appear to recruit chicken monocytes to synthesize new giant cells despite increased vascularization from the membrane, and the survival time is typically 10 days (23). Thus, it was speculated that GCTB monocytes do not originate from the
circulation, but rather arise from the bone marrow. Therefore, it is necessary to directly inject GCTB cells into the bone environment (1).

The intratibial injection method is one of the most widely used murine models to investigate cancer cell growth within the bone environment. It has been used to establish orthotopic models for investigating osteosarcoma biology within the bone environment (26). It has also been used to research bone cancer pain and cancer bone metastasis in prostate (27), and breast cancer (28,29). In addition, the intratibial injection method leads to a reproducible and valuable model for drug testing (27).

Thus far, attempts to grow GCTB in animal models and derive suitable cell lines from primary tumors have failed. This has limited research in the pathobiology of GCTB and the development of specific anti-GCTB agents. In the present report, this problem was addressed by examining whether it was possible to establish an orthotopic model of GCTB in nude mice following intratibial injection of patient-derived tumor cells.

Materials and methods

Ethics statement and patient samples. The use of all patient-derived tumor specimens was approved by the Institutional Review Board and the Research Ethics Committee of Changzheng Hospital (Shanghai, China). Written informed consent was obtained individually from each patient. The pathological diagnosis of GCTB was verified by biopsy prior to surgical excision. Specimens were obtained at the time of surgery from patients undergoing tumor resection, and the diagnosis of GCTB was verified postoperatively by a bone pathologist. Tissue samples from 5 cases (2 male: 3 female) of GCTB were used in the present study, and all experiments were performed with three mice per patient sample. The mean age of the five study patients was 24.4±1.5 years (median age, 25 years; range, 19-28 years). Patients were underwent pathological examination for pathological diagnosis and patients with GCTB were selected for the present study.

Reagents and established cell lines. The human fetal osteoblast hFOB1.19 cell line was obtained from the Institute of Biochemistry and Cell Biology of Shanghai (Shanghai, China). Dulbecco's modified Eagle's medium-F12 (DMEM-F12), DMEM, penicillin/streptomycin and fetal bovine serum (FBS) were purchased from Thermo Fisher Scientific, Inc. (Gibco; Waltham, MA, USA). Collagenase B was obtained from Roche Diagnostics GmbH (Mannheim, Germany). DAPI and the tartrate-resistant acid phosphatase (TRAP) staining kit was purchased from Merck KGaA (Sigma-Aldrich; Darmstadt, Germany). The primary antibody against human Mitochondria (HuMi) was obtained from EMD Millipore (Chemicon; Billerica, MA, USA).

Patient-derived GCTB cells collection and cells culture. Patient-derived GCTB cells were isolated from tumor samples from tumor resections at the Changzheng Hospital as aforementioned. The tissue was mechanically cut into small pieces, and digested with 1.5 mg/ml collagenase B for 3 h at 37°C in DMEM containing 4.5 g/l glucose supplemented with 10% FBS, 100 U/ml penicillin and 100 µg/ml streptomycin. Cells were collected by filtration (100-µm diameter filter), then centrifuged at 200 x g for 5 min at room temperature and washed twice in PBS. The cells were counted using a hemocytometer and resuspended at a density of 5x10⁵ cells/20 µl PBS. For morphological observation and TRAP staining, 1x10⁵ cells were seeded in 6-well plate and cultured in DMEM supplemented with 10% FBS, 100 U/ml penicillin and 100 µg/ml streptomycin in a 37°C humidified incubator with 5% CO₂ (30).

The hFOB1.19 cells were maintained in DMEM-F12 supplemented with 10% FBS, 100 U/ml penicillin and 100 µg/ml streptomycin in a humidified atmosphere with 5% CO₂ at 34°C. The medium was replenished every 2-3 days. After reaching 70-80% confluence, cells were passaged with treatment with 0.25% trypsin. Prior to intratibial injection, the cells were detached from the petri dish with 0.25% trypsin and centrifuged at 200 x g for 5 min at room temperature. Cells were resuspended into a 5x10⁵ cell/20 µl cell suspension in PBS.

Morphological observations. Cells (2x10⁵ cells/well) were incubated for 48 h in 24-well plates. Following incubation, the medium was removed and the cells were washed once with PBS. Morphology was observed using a phase contrast inverted microscope (Leica Microsystems GmbH, Wetzlar, Germany). For detection of osteoclasts, cells (2x10⁵ cells/well) were incubated for 48 h in 24-well plates. Following incubation, the medium was removed and cells were washed twice with PBS. Cultures were fixed for 10 min at room temperature with 4% paraformaldehyde in PBS and then washed four times with PBS. Fixed cells were stained for TRAP using the Acid Phosphatase Leukocyte (TRAP) kit, according to the manufacturer's protocol (cat. no. 387A-1KT; Sigma-Aldrich; Merck KGaA). Staining was observed using a phase contrast inverted microscope at a x400 magnification (Olympus Corporation, Tokyo, Japan).

Experimental animals. A total of fifteen male 4-5-week-old BALB/c nu/nu mice (weighing 18-20 g) were purchased from Shanghai SLAC Laboratory Animal Co., Ltd. (Shanghai, China). The animals were housed in the specific pathogen-free animal facilities with free access to food and water. The temperature was maintained at 20-26°C and a relative humidity of 40-70%. Lighting conditions, 12 h a day fluorescent light and 12 h of darkness.

Intratibial injection of patient-derived GCTB cells. Patient-derived GCTB cells and hFOB1.19 cells were resuspended in PBS at a working concentration of 5x10⁵ cells/20 µl. Patient-derived GCTB cells were prepared from the fresh giant cell tumor of bone specimens without any culture. The intratibial bone injection was performed in 4- to 5-week-old BALB/c nu/nu mice anesthetized with pentobarbital (50 mg/kg) via a percutaneous approach (27). Briefly, the proximal end of the left tibia was exposed surgically, the knee was maintained in a flexed position, and 20 µl of PBS-containing patient-derived GCTB cells were injected into the bone marrow space with a 26-gauge needle. Control mice were injected with hFOB1.19 cells or PBS in the same manner. Four months after injection, nude mice were sacrificed and the injected tibial
samples were fixed, and further analyzed as described below. Procedures involving animals and their care were conducted in conformity with the National Institutes of Health guidelines (31), and were approved by the Animal Care and Use Committee of the Second Military Medical University.

Detection of bone lesions by radiography. Animals were anesthetized and radiographs were obtained using a Kodak DXS 4000 Pro system (Kodak, Rochester, NY, USA) for 1 min at 35 kV on day 0, and weekly following intratibial injection. The radiographs were analyzed for the presence and type of bony lesion (osteolytic, osteoblastic or mixed lytic/blastic).

Three-dimensional, micro-computed tomography (micro-CT). Micro-CT was performed using a SkyScan 1076 Desktop X-ray microtomograph (Bruker microCT, Kontich, Belgium). The excised mouse legs were secured in polystyrene imaging tubes and scanned [x50 magnification; 5.86 µm resolution; 7.5 sec exposure time (40 kV and 258UA); 0.45 rotation step; 180°; and a 1 mm Aluminum filter]. Image reconstruction was performed with volumetric reconstruction software (version 2.4.0.0) and 3D image analysis was performed with CT Analyzer software (version 1.8.1.2) and 3D Visualization software (version 2.0.0.4) (all from Bruker microCT).

The bone volume (trabecular bone volume/tissue volume, BV/TV), and the bone lesion area (bone lesion/normal bone, BL/NB) were determined by three-dimensional structural parameters analysis.

Histological analysis of bone destruction. Following X-ray analysis, the tibial specimens were processed by fixation with 4% paraformaldehyde in PBS for 24 h at 4°C, followed by decalcification in 0.5 M EDTA. Paraffin-embedded sections were cut with a microtome into 4-µm thick sections, then stained with hematoxylin and eosin (H&E) for histological examination of tumor growth, and general form of the tibia. For H&E staining, tissue slices underwent standard rehydration and deparaffinization procedures (32) and were then stained with hematoxylin for 10 min and eosin for 3 min at room temperature. The H&E stained sections were imaged using the Leica DM 4000B microscope at x400 magnification (Leica Microsystems GmbH).

Detection of TRAP activity. Unstained tibia sections were deparaffinized and rehydrated for detection of TRAP activity as an osteoclast marker. Briefly, specimens were equilibrated for 20 min at room temperature in 0.2 M sodium acetate buffer containing 50 mM L-(-)-tartaric acid in deionized water, pH 5.0. The sections were incubated in the TRAP staining reagent for 1 h at 37°C and then at room temperature for 20-30 min until the color reaction was complete. The sections were air-dried and mounted with Eukitt medium. TRAP stained sections were analyzed using the Olympus DP71 phase contrast inverted microscope at x400 magnification (Olympus Corporation).

Cell tracking using anti-HuMi immunofluorescence. Unstained tibia sections were deparaffinized, rehydrated and processed for human specific antigens (mouse anti-human mitochondria; cat. no. MAB 1273; HuMi, 1:50; Chemicon, Temecula, CA, USA) to determine the presence of human cells. Sections were incubated overnight at 4°C with the aforementioned primary antibodies diluted in PBS containing 10% normal goat serum. The following day, sections were treated with the appropriate secondary antibodies (goat anti-mouse IgG; cat no. ab150113; 1:1:000; Abcam, Cambridge, MA, USA) for 1 h at room temperature. Negative controls involved the same procedures with the omission of the primary antibody. DAPI was used for counterstaining nuclei for 5 min at room temperature. The immunostained sections were analyzed using the Olympus BX60 and IX71 epifluorescence microscopes with Olympus DP-70 digital acquisition system at x400 magnification (Olympus Corporation). Images were obtained using 40X objectives and processed with Adobe Photoshop CS3 version 10.0.1 software (Adobe Systems, Inc., San Jose, CA, USA).

Statistical analysis. All of the quantitative experiments were performed more than three times. A Student's t-test was used to determine significant differences between various groups. P<0.05 was considered to indicate a statistically significant difference. All statistical analysis was carried out using SPSS version 18.0 (SPSS, Inc., Chicago, IL, USA).

Results

Clinical characteristics and cellular analysis. The clinical characteristics of the 5 patients included in the present study are presented in Table I. The H&E and TRAP staining of patient tumor tissues are presented in Fig. 1 (Fig. 1A-E and F-J, respectively). Patient-derived GCTB cells were isolated from patient samples using the collagenase B digestion method. They contained the three major cell types, namely mesenchymal stromal cells, round monocytes of macrophage lineage, and multinucleated giant cells (Fig. 1K-O). The multinucleated giant cells were positively stained with TRAP to confirm this characteristic osteoclast feature (Fig. 1P-T). The components, morphology and characteristics of patient-derived GCTB cells were consistent with the clinical pathology reports and H&E staining.

X-ray and micro-CT analysis of bone lesions. The sites of intratibial injections in mice were visible on weekly X-rays. Radiographic data revealed the appearance of osteolytic bone lesions 2 months following intratibial injection of patient-derived GCTB cells (Fig. 2). The intensity of osteolysis gradually increased as time passed (Fig. 2A). Notably, the BL/NB was significantly increased in GCTB mice (Fig. 2C). In control mice that received injections with hFOB1.19 or PBS, there was no evidence of bone damage in the proximal end of the tibia (Fig. 2A and C).

Micro-CT analysis confirmed marked osteolytic destruction in GCTB mice (Fig. 2B). The analysis revealed that GCTB tumors had almost completely destroyed the tibial bone architecture 4 months after the introduction of primary tumor cells, whereas the tibial bone architecture was intact in PBS alone or hFOB1.19 control mice. The ratio of BV/TV was significantly reduced in GCTB mice compared with the control group (Fig. 2D).

Histological analysis and TRAP staining. Paraffin histology revealed a cavity within the tibias of mice injected with primary...
GCTB cells. The visible cavities were surrounded with tumor cells arranged in a barrier around the opening (Fig. 3A). Multinucleated TRAP-positive cells, which are osteoclast-like cells, were clearly visible within the bone matrix (Fig. 3B and C).

**GCTB cells survival at the lesion site.** Four months after transplantation, human GCTB cells were detected with an antibody against HuMi at the bone lesion site, indicating the presence of transplanted cells amongst a background of unstained mouse cells (Fig. 3D-F).

**Discussion**

The recent development of *in vitro* culture of human GCTB cells has contributed to the understanding of the phenotypes and physiology of GCTB. Although the stromal cell subcutaneous injection model and chick chorioallantoic membrane model have been utilized (22,23,25), little is known regarding the orthotopic behavior of human GCTB cells due to the difficulties in establishing a stable cell line that allows for detailed investigation of their biology. In the present study, patient-derived tumor cells isolated from GCTB were injected into the bone marrow of nude mice in order to establish an orthotopic murine model for human GCTB, a method proven to aid engraftment and develop bone lesions in the murine bone marrow microenvironment. Successful engraftment of GCTB-originating cells and subsequent bone lesions were observed in bone environment of mice.

In the current study, histological analysis of GCTB xenograft tissue sections were performed using anti-HuMi, H&E and TRAP staining. Histological analysis of H&E and TRAP staining revealed that the patient-derived GCTB cells survived, and caused an osteolytic reaction in the bone environment. Furthermore, previous studies have reported that

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**Table I. Epidemiological and pathological characteristics of 5 patients with GCTB.**

<table>
<thead>
<tr>
<th>Case</th>
<th>Age (years)</th>
<th>Gender</th>
<th>Site</th>
<th>Grade</th>
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<td>Tibia</td>
<td>I</td>
<td>Primary</td>
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<tr>
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<td>Primary</td>
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<td>Tibia</td>
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<td>Femur</td>
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<td>Primary</td>
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<tr>
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<td>25</td>
<td>Female</td>
<td>Sacrum</td>
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<td>Primary</td>
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GCTB, giant cell tumor of bone.
Figure 2. Digital radiography and micro-CT image of osteolytic destruction in the tibias of nude mice. (A) Mice received intratibial injections of PBS alone or 5x10^5 hFOB1.19 cells or GCTB primary cells. Extensive bone lysis was observed in mice injected with patient-derived GCTB cells (→). (B) Serial 2D cross-sectional and longitudinal micro-CT images of PBS-, hFOB1.19- or GCTB-injected tibiae acquired at 20 µm resolution. Osteolytic destruction (→) was observed in cross-sectional and longitudinal micro-CT sections following primary GCTB cell injection. Representative 3-D reconstruction of micro-CT images of the tibiae from each treatment group, revealing osteolytic lesions in the proximal tibia of mice that received GCTB-injection. (C) BL/NB was significantly increased in GCTB-injection mice. *P<0.05. (D) BV/TV of bone lesions was calculated using a CT analyzer. The percentage of BV/TV was significantly reduced in GCTB-injected mice. *P<0.05; **P<0.01. Micro-CT, micro-computed tomography; GCTB, giant cell tumor of bone; BL/NB, bone lesion/normal bone ratio; BV/TV, bone volume/tissue volume ratio; CS, cross-sectional; L, longitudinal.

Figure 3. Immunofluorescence microphotographs revealing the pattern of human cell markers at the bone lesion site. Tissue sections (4-µm thick) were taken from the tibial bone 120 days after injection with patient-derived GCTB cells. (A) H&E stained demonstrated that animals injected with patient-derived GCTB cells exhibited osteolytic destruction at the proximal tibia. (B) TRAP-positive osteoclasts were observed at the bone lesions in mice injected with patient-derived GCTB cells. (C) Osteoclasts (*) were distributed within the trabecular bone marrow interface and the bone matrix. (D) Anti-HuMi (green) staining revealed the presence of transplanted cells amongst a background of non-staining mouse cells. (E) Cell nuclei were visualized with DAPI staining (blue). The overlay of human mitochondria fluorescence and DAPI staining are presented as merged images (F). Bars represent 50 µm. H&E, hematoxylin and eosin; TRAP, tartrate-resistant acid phosphatase; GCTB, giant cell tumor of bone; HuMi, human mitochondria.
stromal cells can differentiate into mature osteoblasts, adipocytes and chondrocytes (11,12,33-36). Observed differences between the results of the present study and others may indicate that stromal cells undergo multidirectional differentiation, which may be influenced by the tissue microenvironment. As such, different environmental conditions may produce varied differentiation features and functions.

The X-ray and micro-CT features of bone destruction in the present model were similar to the clinical imaging characteristics, revealing pure lytic lesions without any degree of matrix calcification or ossification and surrounding reactive sclerosis (3). However, biopsies of nude mice tibia did not exhibit the same clinicopathological features as the chick chorioallantoic membrane model, which is characterized by the presence of numerous multinucleated giant cells that are uniformly distributed amongst mononuclear spindle-like stromal cells and other monocytes (19). In the present study, a large number of TRAP-positive cells were identified surrounding the surviving stromal cells, and were distributed at the surface of the tibia trabecular and matrix bone. As reported in the literature, stromal cells are capable of secreting a variety of cytokines, including stromal cell-derived factor-1 (37), macrophage colony-stimulating factor (38) and monocyte chemotactic protein-1 (39). The release of these cytokines results in the recruitment of monocytes and further secretion of interleukin (IL)-6 (40), matrix metalloproteinases (41,42), and parathyroid hormone-related protein (43) to promote osteoclast fusion and bone resorption.

A limitation of the current study was the use of patient-derived GCTB cells containing all types of cells from GCTB tissue. Isolation of the different subtypes of GCTB primary cells using flow cytometry to compare the survival and tumorigenicity of different cell populations within the bone environment was not performed. Therefore, additional studies are required to evaluate the survival and tumorigenicity of the various cell types or combinations. Previously, it has been reported that GCTB stromal cells transfected with green fluorescence protein survived 52 weeks in vivo (24). However, isolation of stromal cells from GCTB in vitro results in the rapid loss of expression of receptor activator of nuclear factor-kB ligand (44). This suggests that the pathogenesis of GCTB requires cellular interactions, which provide a suitable microenvironment for tumor growth.

In conclusion, the present study described the establishment of a novel xenograft model of GCTB using patient-derived tumor cells. This murine xenograft GCTB model produced visible osteolytic destruction in the proximal tibias of nude mice. Anti-HuMi immunofluorescence staining confirmed that the surviving cells in the osteolytic destruction were of human GCTB cell origin. These findings may contribute to the understanding of the pathogenesis of GCTB, and thus aid in the improvement of diagnosis, treatment and patient outcome.

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Availability of data and materials

All data generated or analyzed during this study are included in this published article.

Authors' contributions

LX conducted the experiment studies. ZW and ZZ performed the statistical analysis and drafted the manuscript. JX and XY made substantial contributions to conception and design of this work and critically revised the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The use of all patient-derived tumor specimens was approved by the Institutional Review Board and the research ethics committee of Shanghai Changzheng Hospital (approval no. 2011/081), which appeared in the proceedings of the meeting of the Ethics Committee on November 18, 2011.

Patient consent to participate

Written informed consent was obtained individually from each patient.

Competing interests

The authors declare that they have no competing interests.

References


