Preclinical trial of the multi-targeted lenvatinib in combination with cellular immunotherapy for treatment of renal cell carcinoma

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Abstract. Lenvatinib is an oral, multi-targeted tyrosine kinase inhibitor of vascular endothelial growth factor receptors 1-3, fibroblast growth factor receptors 1-4, platelet-derived growth factor receptor β, RET and KIT. Cellular immunotherapy has the potential to be a highly targeted treatment, with low toxicity to normal tissues and a high capacity to eradicate tumor tissue. The present study assessed the safety, maximum tolerated dose (MTD) and preliminary antitumor activity of lenvatinib and cellular immunotherapy in a murine model of renal cell carcinoma (RCC). The present study used a therapeutic dose of 0.12 mg lenvatinib and/or 10⁴ rat uterine cancer adenocarcinoma (RuCa)-sensitized lymphocytes administrated once daily continuously in 7-day cycles. Tumor regression was observed in mice with RCC following treatment with lenvatinib and 10⁴ RuCa-sensitized lymphocytes. MTD was established as once daily administration of 0.18 mg lenvatinib and 10⁵ RuCa-sensitized lymphocytes. The most common treatment-related adverse effects observed were fatigue (40%), mucosal inflammation (30%), proteinuria, diarrhea, vomiting, hypertension and nausea (all 40%). Combination therapy using lenvatinib and cellular immunotherapy enhanced the antitumor effect induced by single treatments and prolonged the survival of mice with RCC compared with either of the single treatments. Treatment with lenvatinib (0.12 mg) combined with 10⁴ RuCa-sensitized lymphocytes was associated with manageable toxicity consistent with individual agents. Further evaluation of this combination therapy in mice with advanced RCC is required. In conclusion, cellular immunotherapy and oncolytic therapy for cancer may be improved by the synergistic effects of lenvatinib and sensitized lymphocytes. In the present study, the inherent antineoplastic and immune stimulatory properties of the two agents were enhanced when used in combination, which may provide a basis for clinical treatment of patients with RCC.

Introduction

Standard therapies including chemotherapy and radiotherapy for various types of human cancer are toxic and often ineffective (1). Cellular immunotherapy has the potential to be a highly targeted alternative to these therapies, with a high capacity to eradicate tumors and low toxicity to normal tissues (2-4). Cellular immunotherapy often employs active immunization with cells, peptides, proteins or nucleic acids, as well as utilizing adoptive transfer of effector cells that directly target antigens on malignant cells (5). A variety of these approaches have been demonstrated to be successful in treating ovarian and lung cancer (6,7). In the present study, approaches that generate cellular therapies by combining active immunization with cells that use adoptive transfer of effector cells, which directly target RCC cells in xenograft mice, were investigated.

Lenvatinib is an oral, multi-targeted tyrosine kinase inhibitor of vascular endothelial growth factor receptors 1-3 (VEGFR1-3), fibroblast growth factor receptors 1-4 (FGFR1-4), platelet-derived growth factor receptor β (PDGFR-β), RET and KIT. Previous studies have identified VEGF, FGFR1-4, PDGFR-β, RET and KIT-mediated angiogenesis to be key factors in the development of renal cell carcinoma (RCC) (8,9). Lenvatinib has potent anti-tumor activity against various human tumors in murine xenograft models, mediated by the inhibition of angiogenesis (10). Various agents that inhibit the VEGF pathway have demonstrated clinical benefit in metastatic RCC (11). Lenvatinib, which targets the VEGFR1-3, FGFR1-4, PDGFR-β, RET and KIT pathways, has also demonstrated clinical benefit in RCC (11). A previous study indicated that lenvatinib combined with everolimus significantly extended overall survival compared with everolimus alone in patients with metastatic RCC (12).

Research suggests that tumors become resistant to therapy via feedback mechanisms that compensate for targeted inhibition (13). The upregulation of hypoxia-inducible factor 1 target genes, such as VEGF, has been implicated in RCC (12).
Additionally, genetic alterations resulting in the constitutive activation of cellular immunotherapy signaling pathways have been reported to be associated with RCC (14). Therefore, a combination of agents that target VEGF and cellular immunotherapy-mediated pathways may simultaneously block two critical signaling pathways that are activated in RCC, and may overcome an aspect of resistance to single-agent therapy.

RCC, caused by mutations in renal tubular cells, is one of the most common urologic neoplasms (15). Increasing research has focused on the development of comprehensive treatments for RCC (16,17). The present study demonstrated the synergistic effects between cellular immunotherapy and lenvatinib in a murine model of RCC and the landscape of current targeted and cellular immunotherapy for RCC was examined. Our results indicated that cellular immunotherapy combined with targeted agents, such as lenvatinib, is undergoing preclinical trials in RCC. The results of the present study demonstrated that cellular immunotherapy combined with lenvatinib inhibited tumor growth in xenograft mice. Furthermore, it was demonstrated that cellular immunotherapy combined with lenvatinib significantly prolonged the survival of tumor-bearing mice (P<0.05). These results indicated that cellular immunotherapy combined with lenvatinib treatment may be applied in clinical practice for treatment of patients with RCC.

Materials and methods

Ethics statement. The present study was conducted in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health (13). Experimental protocols were approved by the Chinese Association for Laboratory Animal Sciences (Guangzhou, China) and Animal Health Products and the Committee on the Ethics of Animal Experiments Defense Research (Guangzhou, China). All surgery and euthanasia were performed under sodium pentobarbital anesthesia (30 mg/kg, Sunbio Inc., Shanghai, China), and all efforts were made to minimize suffering.

Cell culture and reagents. Cells from a human kidney adenocarcinoma cell line (ACHN), renal carcinoma cell line (786-0) and rat uterine cancer adenocarcinoma cell line (RuCa) cell line were obtained from the National Cancer Institute (NCI) Frederick Cancer Research Facility, the NCI Division of Cancer Treatment Tumor Repository (both Frederick, MD, USA) and the American Type Culture Collection (Manassas, VA, USA), respectively. ACHN, 786-0 and RuCa cells were cultured in Dulbecco's modified Eagle's medium supplemented with 10% heat-inactivated fetal bovine serum (BioWhittaker; Lonza Walkersville, Inc., Walkersville, MD, USA), 3 mM L-glutamine (Meilon Biotechnology Technology Co., Ltd., Shanghai, China), 50 µg/ml gentamicin (BioWhittaker; Lonza Walkersville, Inc.) and 1% penicillin/streptomycin. Normal human astrocyte cells (catalogue no. CC-2565; Clonetics™; Lonza) were purchased from Lonza Walkersville, Inc., and maintained in an AGM Astrocyte Growth Bullet kit (Lonza Walkersville, Inc.), according to the manufacturer's instructions.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total cellular RNA was extracted from cells using an RNeasy mini kit (Qiagen, Inc., Valencia, CA, USA), according to the manufacturer's instructions. cDNA (10 ng) was used for qPCR with the SYBR-Green Master Mix system (Bio-Rad Laboratories, Inc., Hercules, CA, USA). RNA samples (1 µl) were reverse transcribed using the iScript cDNA synthesis kit (catalogue no. 1708890; Bio-Rad Laboratories, Inc., Hercules, CA, USA), according to the manufacturer's instructions. RNA enzyme-free ultrapure water (dilution, 1:10) and the absorbency at wavelengths of 260 and 280 nm were recorded using an ultraviolet spectrophotometer (UV-1100; Shanghai Mapada Instruments Co., Ltd., Shanghai, China) to determine purity and concentration. The cDNA sample (1 µl) was subjected to a PCR reaction, performed in an iCycler thermal cycler using IQ SYBR Green Supermix (both Bio-Rad Laboratories, Inc.). The Fast Real-Time PCR system was used with the following protocol: 25°C for 5 min, 42°C for 1 h and 70°C for 5 min for a total of 35 cycles. GAPDH levels were measured as an internal control. The forward and reverse primers are as follows: VEGFR, forward 5'-GCCCCGCCT TATGATTCCTGC-3' and reverse 5'-CTCGCGCCGTTC GTATAGTC-3'; PDGFR, forward 5'-CAGCAGTATGAA AGCGTGG-3' and reverse 5'-GGAAGAAAAGGTTGGCAC AG-3'; RET, forward 5'-GCAGGACCGGTCAATGAT-3' and reverse 5'-GACTGACCCCGTCGAGC-3' and KIT, forward 5'-TGCAAGACAACTGTAAGA-3' and reverse 5'-AAC TGGGAAAATGACATCTGG-3'. Primers were synthesized by Invitrogen (Thermo Fisher Scientific, Inc., Waltham, MA, USA). The expression level of mRNA was denoted as mRNA/GAPDH, assessed using the 2^ΔΔCq protocol, following detection of expression levels in all three cell types (18). PCR was repeated three times in triplicate.

MTT cytotoxicity assays. ACHN, 786-0 and RuCa cells (5x10^3/ml) were cultured in 96-well plates for 24 h at 5% CO₂ and 37°C and incubated with lenvatinib (0.1 mg/ml; Sunbio Inc.) and/or sensitized lymphocytes (1x10^5/ml; Sunbio Inc.) for 48, 72 and 96 h at 37°C in triplicate for each condition. Phosphate buffered saline (PBS) was added instead of the lenvatinib or sensitized lymphocytes to act as a control. At each time point, 20 µl MTT reagent (5 mg/ml) in PBS solution was added to each well and the plates were incubated at 37°C for 4 h. The majority of the medium was removed and 100 µl dimethyl sulfoxide was added into the wells to solubilize the formazan crystals. The optical density was measured using a microplate reader (Bio-Rad Laboratories, Inc.) at a wavelength of 450 nm.

Induction of apoptosis in RCC cells. ACHN, 786-0 and RuCa cells were grown at 37°C in 5% CO₂ until 80% confluence was reached. Apoptosis was assessed by incubation of these cells with lenvatinib (0.1 mg/ml) and/or sensitized lymphocytes (1x10^5/ml) for 72 h at 5% CO₂ and 37°C. Following incubation with the lenvatinib and/or sensitized lymphocytes, the cells were trypsinized and collected. Subsequently, the cells were washed twice for 30 min in cold PBS, adjusted to 1x10^6 cells/ml with PBS and labeled with annexin-V-fluorescein isothiocyanate (FITC) and propidium iodide (PI) using an annexin-V-FITC kit (BD Biosciences, San Jose, CA, USA). Apoptotic cells were quantified using a FACSScan flow cytometer (BD Accuri™ C6 Plus; BD Biosciences, San Jose, CA, USA).
Purification of blood derived cells and cytotoxicity assay. Murine peripheral blood mononuclear cells (PBMCs) were isolated from theuffy coat of 40 healthy male mice purchased from the Animal Center of Nanjing Medical University (Nanjing, China) between 8-9 weeks of age with a body weight of 20-25 g by Ficoll-Paque (GE Healthcare Life Sciences, Chalfont, UK) density gradient centrifugation. Mice were maintained in a room with constant temperature (22±1°C) and a 12-h light/dark cycle and housed in cages (<5 mice per cage) with ad libitum access to food and water. T cells were purified from PBMCs via magnetic bead separation using antibody-coated Dynabeads (Invitrogen; Thermo Fisher Scientific, Inc.) containing monoclonal antibodies against cluster of differentiation (CD) 14, CD16, CD56, human leukocyte antigen class II antigen D related (HLA-DR)/DP and beads coated with an Fe-specific human immunoglobulin (Ig) G4 against mouse IgG, or using anti-CD3 MACS microbeads (Miltenyi Biotec GmbH, Bergisch Gladbach, Germany), according to the manufacturer’s guidelines. The purified T cells were stimulated at 37°C for 72 h with ACHN, 786-0 or RuCa cells. All cells were recovered from the culture, counted, and used as effector cells.

Tumor cells (2x10^6) were labeled with 200 μCi chromium 51 (Zen-bio Inc., Shanghai, China; ^51Cr) for 1 h at 37°C, washed four times with PBS and resuspended at 2x10^5 cells/ml in PBS. To quantify T-cell induced cell death, tumor cells (10^4 cells/well) were incubated at 37°C with varying numbers of T cells (10^4, 10^5, 10^6 and 10^7) for 48 h. The assay was performed in 96-well round-bottom plates in triplicate, and the percentage of specific lysis was calculated after analysis of cytotoxicity in the supernatants of the wells using the following formula: 100 x [experimental counts per minute (cpm)-spontaneous cpm]/(total cpm-spontaneous cpm)]. Spontaneous and total ^51Cr release values were determined in the presence of either PBS alone or 10% Triton X-100. RuCa-sensitized lymphocytes were used in animal experiments.

Enzyme-linked immunosorbent assay (ELISA). The production of interferon-γ (IFN-γ) from the cells was analyzed using an ELISA assay. ELISA was performed using an sandwich ELISA kit (catalogue no. ab193969; Abcam, Cambridge, UK) to determine the IFN-γ protein expression level in the mice spleen, obtained during surgery following sacrifice. Briefly, the spleen was homogenized in 1 ml of PBS containing 1% Triton X-100 using the plunger of a 10-ml syringe, and centrifuged at 7155 x g for 5 min at 4°C, and the total protein concentration of the supernatant was determined using a biocinchoninic acid assay as previously described (19). IFN-γ antibody (catalogue no. ab175878; Abcam, Cambridge, UK), horseradish peroxidase-streptavidin and TMB One-Step Substrate (Abcam) reagent were incubated in turn and in 96-well plates at room temperature according to the manufacturer's protocol. Stop solution (Abcam) was then added and the optical density (OD) was read on a microplate reader at 450 nm (Thermo Fisher Scientific, Inc.).

Animal experiments. A total of 40 male BALC/c mice (mean age of 24 months and a body weight of 20-25 g) with RCC were purchased from the West China Experimental Animal Center of Sichuan University (Sichuan, China). Mice were maintained in a room with constant temperature (22±1°C) and a 12-h light/dark cycle, and cages in groups <5 per cage with ad libitum access to food and water. The mice were split into 4 groups, containing 10 mice. Each mouse in the treatment group received 0.12 mg lenvatinib or/and 10^6 RuCa-sensitized lymphocytes administered once daily continuously in 7-day cycles. The mice in the control group received 2 ml PBS. Tumor dimensions were measured 10 times, every 2 days. Tumor volumes were calculated according to the following formula: Length x width^2 x 0.52. On day 30 after the final inoculation, tumor tissues were extracted and analyzed using RT-qPCR assays (as described). Mice were kept alive for 180 days to assess the effects of the treatments on survival.

Efficacy safety assessment. Efficacy assessment included the maximum toxicity dose in mice models and the dose-limiting toxicity in the presence and absence of cellular immunotherapy and lenvatinib. Safety assessments included the incidence rates (>10%) of the most frequent treatment-emergent adverse events in the 30-day treatment period.

Statistical analysis. All data are presented as the mean ± standard deviation. Analysis was performed using SPSS software version 20.0 (IBM Corp, Armonk, NY, USA). Statistical significance of differences between mean values was assessed using Student's t-test for unpaired data. Comparisons of data between multiple groups were performed with one-way analysis of variance, followed by Newman Keuls post hoc test. P<0.05 was considered to indicate a statistically significant difference.

Results

Antineoplastic evaluation of lenvatinib and sensitized lymphocytes in vitro. In order to investigate the effects of cellular immunotherapy and lenvatinib on renal cancer cells, RuCa, ACHN and 786-0 cells were treated with 0.12 mg lenvatinib or/and 10^4 sensitized lymphocytes. The results in Fig. 1 demonstrated that the cytotoxicity of RuCa, ACHN and 786-0 cells when treated with lenvatinib or/and sensitized lymphocyte increased in a time dependent manner after 48, 72 and 96 h of treatment. The results of the MTT assay indicated that the cytotoxicity of RuCa, ACHN and 786-0 cells were inhibited after treatment with sensitized lymphocytes and/or lenvatinib (Fig. 1).

Duration of treatment, dose-limiting toxicities and maximum tolerated dose (MTD). Median overall duration of treatment was 7 days across all dosing cohorts. The dosing cohorts included 0.08, 0.12, 0.16, 0.18, 0.20 and 0.24 mg of lenvatinib and 10^3, 10^4, 10^5, 10^6 and 10^7 of sensitized lymphocytes. In our results, 0.18 mg lenvatinib once daily and 10^6 sensitized lymphocytes once daily was identified as the MTD.

Mice received at least one dose of 0.18 mg lenvatinib and 10^6 with post baseline safety evaluation. Following the last lenvatinib and sensitized lymphocyte dose, the most common treatment-related adverse events (AEs) were hypertension, diarrhea, constipation, weight loss, loss of appetite, proteinuria and rash (Table I). The most common treatment-related AEs with a toxicity criteria grade ≥3 were hypertriglyceridemia, proteinuria, diarrhea and fatigue (10% each; Table II).
Effect of cellular immunotherapy and lenvatinib treatment on apoptosis of RCC cells. The morphological changes induced by cellular immunotherapy and lenvatinib treatment in RuCa, ACHN and 786-0 cells indicated that treatment with sensitized lymphocytes and lenvatinib inhibited the growth of renal tumors. Therefore, the therapeutic effects of sensitized lymphocytes and lenvatinib in vitro on human and mouse RCC cells were investigated. Results demonstrated that apoptosis of ACHN, 786-0 and RuCa cells significantly increased after treatment with lenvatinib compared with the control group, and the apoptosis of ACHN, 786-0 and RuCa cells significantly increased after treatment with combined lenvatinib and sensitized lymphocyte treatment group compared with the lenvatinib and sensitized lymphocyte groups (P<0.01; Fig. 2). The mRNA expression levels of several genes associated with lenvatinib-mediated apoptosis, including VEGFR,
PDGFR, RET and KIT, were analyzed using RT-qPCR. As demonstrated in Fig. 3, mRNA expression levels of VEGFR, PDGFR, RET and KIT were significantly decreased in RCC cells following treatment with lenvatinib compared with the control and they also were significantly decreased in RCC cells following treatment with combined lenvatinib and sensitized lymphocyte treatment group compared with lenvatinib group (P<0.01). These results suggest that lenvatinib inhibited the expression of angiogenesis-related genes.

**Effect of cellular immunotherapy and lenvatinib treatment of RCC on survival in vivo.** To explore whether cellular
immunotherapy and lenvatinib are effective anticancer agents in vivo, the anti-tumor activity of the sensitized lymphocytes and lenvatinib in a murine model of RCC was investigated. Results demonstrated that the tumor volume in mice treated with sensitized lymphocytes plus lenvatinib was significantly lower than that of the mice treated with PBS or a single agent (P<0.01; Fig. 4A). Furthermore, long-term survival was assessed for 180 days after treatment with cellular immunotherapy and lenvatinib treatment. As demonstrated in Fig. 4B, cellular immunotherapy and lenvatinib (n=10 in each group) significantly prolonged the survival of mice compared with the control group (P<0.01). Furthermore, the combined treatment of cellular immunotherapy and lenvatinib was significantly more effective at prolonging survival of the mice than either of the treatments alone (P<0.01; Fig. 4B). These results indicate that combined treatment of sensitized lymphocytes plus lenvatinib is strong enough to partially protect the animals and eliminate the tumor cells, which translated into long-term survival. The maximum percent tumor change from baseline to post baseline nadir is demonstrated by the ELISA results presented in Fig. 5 and illustrates the immune responses of the mice to combined therapy.

**Discussion**

Therapeutic targeting of VEGF-mediated pathways has increased the number of available treatment options for human cancer (10,20,21). A number of challenges remain to allow the treatment to overcome the resistance to single-agent therapies (22-24). Combinations of anticancer agents have previously been demonstrated to have strong therapeutic effects (25-27). Optimal treatment schemes using these agents have been
defined and clinical development of combinations of anticancer agents have demonstrated low toxicity (28). The most evident and efficient effects are induced by cellular immunotherapy and targeted therapy (28). However, there is a continued requirement for more effective combinations of targeted agents and cellular immunotherapies that demonstrate improved efficacy, are manageable and have low toxicity (29-31).

A previous study demonstrated that the MTD and recommended phase II dose in this phase I experiment was confirmed to be 18 mg lenvatinib in combination with 5 mg everolimus once daily for treatment of RCC (12). In the present study, cellular immunotherapy and targeted therapy was introduced to treat RCC in a xenograft murine model. At the MTD and lower-dose cohort, the combination of lenvatinib and cellular immunotherapy was associated with manageable toxicity. Treatment-related AEs were consistent with class effects typical of VEGFR-targeted therapy and immunotherapy (32). A previous study indicates that fatigue, mucosal inflammation, proteinuria, hypertension, as well as gastrointestinal toxicity, occurred at a low frequency (32). Another previous study indicated that proteinuria, hypertriglyceridemia, diarrhea, and fatigue were the most common treatment-related TEAEs (33).

A phase I study of lenvatinib, which is a multi-targeted tyrosine kinase inhibitor, in patients with advanced solid tumors demonstrated that lenvatinib is an efficient anticancer agent (34); however, only 9% of patients had a partial response to treatment and so the results of the study suggested that lenvatinib as a monotherapy had limited efficacy for cancer patients. A previous study by Yamada et al (35) reported that lenvatinib has manageable toxicity when administered in a 2-week-on/1-week-off cycle and demonstrated preliminary activity for durable disease control. However, result analysis suggested that antiangiogenic activity correlated with anti-tumor activity in patients with partial therapeutic effects (35). In a phase II trial combining the treatment doses of bevacizumab and everolimus, median progression-free survival in previously treated patients was 7.1 months (23%) (36). Therefore, the combination regimen was associated with a high cure rate compared with single agent treatment (37,38). The responses observed with these combination therapies confer an advantage over single-agent therapy; however, further study on the use of these combination therapies is required (39).

In the present exploratory analysis of tumor therapy, treatment with lenvatinib (0.12 mg) and 10^4 RuCa-sensitized lymphocytes once daily resulted in a mean survival rate of 80%, and median progression-free survival of ~180 days in mice with RCC. In conclusion, the present study used two anticancer agents, lenvatinib (0.12 mg) combined with 10^4 RuCa-sensitized lymphocytes, once daily to treat RCC in mice. The majority of treatment-related AEs were consistent with positive effects of immunotherapy and multi-targeted tyrosine kinase inhibitors, and were managed effectively by administration of lenvatinib and sensitized lymphocytes continuously in 7-day cycles. Notably, beneficial effects of cellular immunotherapy and targeted therapy may further elucidate the clinical value of combination therapy for cancer treatment.

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References


Figure 5. Treatment with lenvatinib plus SLYS enhances immune responses against tumor cells in vivo. (A) CTL response against renal cell carcinoma. (B) Tumor-specific IFN-γ released from stimulated splenocytes in a murine model of renal cell carcinoma. Data are presented as the mean ± standard deviation of triplicate samples. One-way analysis of variance revealed a significant effect. *P<0.01 vs. the PBS control. PBS, phosphate buffered saline; SLYS, sensitized lymphocytes; CTL, cytotoxic T lymphocyte; IFN, interferon.
The role of somatic genetic events and the challenges for targeted therapies.


