### Abstract
The purpose of this study was to examine the effects of irradiation by 125I seeds in human lung cancer xenograft model and to determine the underlying mechanisms involved, with a focus on angiogenesis. A group of 40 mice bearing A549 lung adenocarcinoma xenografts was randomly separated into 4 groups: control group (n=10), sham seed (0 mCi) implant group (n=10), 125I seed (0.6 mCi) implant group (n=10) and 125I seed (0.8 mCi) implant group (n=10), respectively. The body weight and tumor volume were recorded every four days until the end of the study. At 30 days after irradiation, the microvessel density, proliferative index and apoptotic index were evaluated by quantitative morphometric analysis of the expression of CD34, proliferating cell nuclear antigen (Ki-67) and *in situ* terminal transferase-mediated fluorescein deoxy-UTP nick-end labeling (TUNEL), respectively. The changes in the expression of hypoxia inducible factor-1α (HIF-1α) and vascular endothelial growth factor (VEGF) were detected by real-time PCR and western blot analysis. Consequently, 125I seed irradiation suppressed the growth of lung cancer xenografts in nude mice, while inhibiting cell proliferation and angiogenesis and inducing apoptosis as demonstrated by Ki67, CD34 and TUNEL staining. HIF-1α and VEGF mRNA and protein expression levels were substantially downregulated following 125I seed irradiation. Collectively, our data suggest that irradiation by 125I seeds is a promising new option for lung cancer treatment.

### Introduction
Lung cancer is the most common cancer worldwide, and more than 1.8 million new cases and almost 1.6 million deaths were estimated in 2012, with most of these cases (80-85%) categorized as non-small cell lung cancer (NSCLC) (1). In China, lung cancer is the most frequently diagnosed cancer and the leading cause of cancer-related death (2). NSCLC treatments include surgery, chemotherapy, and radiation therapy, which are determined by the type and stage of cancer. Surgery is the only curative treatment of NSCLC; however, less than 20% of NSCLC patients who were not diagnosed at advanced stages of the disease are potentially curable with surgical resection (3). Numerous clinical studies have confirmed the effectiveness of chemotherapy combined with external beam irradiation (EBRT) in the treatment of patients with NSCLC presenting with locally advanced malignancy (4-7). To our disappointment, they have not greatly affected outcomes, and the gain often comes with substantial severe toxicity (myelosupression, nausea, vomiting and radiation pneumonitis), especially affecting important organs and tissues (heart, esophagus, and large blood vessels) (8,9). A great number of patients with advanced NSCLC also cannot tolerate the currently available treatment modalities mainly owing to their poor general condition, especially with other complication. Therefore, novel therapeutic approaches are urgently needed to effectively prolong survival time and obviously improve the quality of life in advanced patients.

In recent years, 125I implantation has been widely used to treat advanced and inoperable prostate cancer, lung cancer, pancreatic cancer, colorectal cancer and head and neck cancer because of accurate positioning, little trauma, high doses in the target volume, few normal tissues exposed and minimal complications (10-14). The advantages of 125I have led to its widespread application in China (11,15).

Although many clinical trials have reported that 125I seed irradiation is a feasible adjuvant treatment to control local...
symptoms and prolong survival for advanced or unresectable NSCLC (16,17), its biological effects and underlying molecular mechanisms are far from fully understood. 

Qu and colleagues (18) demonstrated that a continuous low-dose rate of irradiation (CLDR) induced stronger growth inhibition in A549 cells than single EBRT due to the aggravation of G2/M arrest, and increased apoptosis. Takabayashi and colleagues (19) demonstrated that 125I seed irradiation induced apoptosis and suppressing proliferation in histologically varied gastric cancers to exert antitumor effects. The tumor microenvironment is often hypoxic, and tumor hypoxia is associated with formation of new capillaries and resistance to radiation therapy. HIF (e.g. HIF-1α, HIF-2α, HIF-3α) are activated in response to hypoxia, which upregulate numerous genes affecting angiogenesis. VEGF as the best known and the most efficient angiogenic factor is constitutively activated. It is thus inferred that the HIF upregulation of VEGF is most likely responsible for angiogenesis in tumors (20,21). It is well known that angiogenesis is absolutely required for tumor growth. Therefore, inhibition of angiogenesis can prevent tumor growth (22). This study was designed to investigate 125I-induced cell apoptosis, suppressing proliferation and angiogenesis and the possible role of HIF-1α and VEGF in the process of 125I brachytherapy in lung carcinoma. Herein, we examined its biological effects on adenocarcinoma cells, the most common pathological type of NSCLC, using xenograft models.

Materials and methods

Cell line and cell culture. The human lung adenocarcinoma cell line A549 was purchased from the American Type Culture Collection (ATCC, Manassas, VA, USA). The cells were cultured in RPMI-1640 (Gibco, Carlshad, CA, USA) medium supplemented with 10% heat-inactivated fetal bovine serum and 1% pen-strep (100 U/ml penicillin and 100 mg/ml streptomycin) (Gibco) in a 37°C humidified incubator containing 5% CO2. All experiments were performed using cells grown to 60-80% confluence. For in vivo injections, suspensions with >95% viability were used, as determined by trypan blue exclusion.

Animal model. Female BALB/c nude mice, 4-6 weeks old and weighing 17-20 g, were purchased from Institute of Chinese Academy of Medical Sciences and allowed to acclimatize for 1 week under specific pathogen-free conditions (23±2˚C and 55±5% humidity) in the Animal Care Facility before any intervention was initiated. The mice were housed and maintained under specific pathogen-free conditions in facilities approved by the Animal Care and Use Committee of Qingdao University School of Medicine. A549 cells (5x10⁶) in 0.2 ml PBS were injected subcutaneously into the right dorsal flank of each mouse. Mice were monitored daily for tumor development and total body weight, tumor incidence and mass, were recorded every four days till the end of study. The tumor volume (V) was calculated by the following formula: V (mm³)=LxW²x1/2 (L, length of tumor; W, width of tumor).

125I seeds. The 125I seeds were provided by Invasive Technology Department of The Affiliated Hospital of Qingdao University, with a diameter of 0.8 mm, and a length of 4.5 mm. The average energy was 27.4 to 35.5 keV, with a half-life of 59.6 days. After decaying into the organs, the 125I seeds released continuous low-dose γ-ray and soft X-ray (5% of 35 keV and 95% of 28 keV, respectively). The internal irradiation was relatively long-acting, 93-97% of the brachytherapy dose was delivered within 8-10 months.

125I brachytherapy seed implant. As tumors reached a size of approximately 300 mm³ in about 24 days, mice were randomly divided into 4 groups (n=10/group). Four groups were grouped as follows: non-implanted control group; sham seed implant group; 125I seed (0.6 mCi) implant group; and 125I seed (0.8 mCi) implant group. Before in vivo implantation, each mouse was anesthetized via inhalation of diethyl ether (provided by Pharmacy Department of the Affiliated Hospital of Qingdao University). The visible mass in mice was punctured by the 18-gauge needles of the Mick-applicator through which seeds were implanted. After 30 days of treatment, all mice were humanely sacrificed, and tumors were harvested and weighed, then fixed in 4% paraformaldehyde (PFA) or flash frozen in liquid nitrogen.

Immunohistochemistry for Ki67 and CD34. Immunohistochemical procedures of Ki67 and CD34 were done as described previously by Lee and colleagues (23). In brief, samples of the tumors were fixed in 4% PFA for 24 h, embedded with paraffin, then sectioned (4 µm) longitudinally and stained with hematoxylin and eosin (H&E). Following staining cell proliferation and microvessel density was assessed by quantitative morphometric analysis of Ki67 and CD34 expression, respectively. For detecting CD34 and Ki67 immunoreactivity, tissue sections were deparaffinized with sequential washing of xylene, rehydrated with descending grades of ethanol, followed by antigen retrieval and endogenous peroxidase treatment. After blocking endogenous peroxides and nonspecific proteins, sections were incubated at 4˚C overnight with rabbit anti-mouse monoclonal anti-Ki67 antibody (Abcam, Cambridge, MA, USA) at a 1:100 dilution and rabbit anti-mouse monoclonal anti-CD34 antibody (Cell Signaling Technology, Boston, MA, USA) at a 1:85 dilution, respectively. Subsequently, slides were washed three times in Tris-buffered saline (TBS), and incubated for 20 min at room temperature with peroxidase-conjugated goat anti-rabbit IgG secondary antibodies (Abcam). Negative controls were incubated with PBS instead of the primary antibody. Visualization was achieved with a dianinobenzidine (DAB) substrate system in which nuclei with DNA fragmentation or endothelial cells were stained brown.

TUNEL staining. For the detection of apoptosis, tumor specimens were subjected to a TUNEL assay using the In Situ Cell Death Detection kit (Roche, Basel, Switzerland), in accordance with the manufacturer’s instructions. Briefly, 4% PFA fixed tissues were permeabilized with Proteinase K working solution (provided in the kit) at 37˚C for 30 min. The slides were washed twice in PBS and incubated with 50 µl of TUNEL reaction mixture (provided in the kit) in a dark and humid atmosphere for 60 min at 37˚C, followed by incubation with Converter-POD solution (provided in the kit) for 30 min at 37˚C. Slides were developed using DAB, mounted under glass coverslip, and then analyzed under a light microscope.
Microscopic quantitative analyses of Ki67, CD34, and TUNEL. To quantify MVD, 5 random fields at x400 magnification per slide were examined for each tumor (one slide per mouse, 5 slides of each treatment group) and microvessel (MV) counts per field were done by two investigators in a blinded fashion. A single MV was defined as a cluster clearly separate from adjacent MVs or single cell stained positive for CD34 with the presence of a lumen. To quantify Ki67 expression, the number of Ki67-positive cells and the total number of tumor cells were counted in 5 random fields at x400 magnification. Proliferative index was calculated by the percentage of positive cells to total tumor cells. For quantification of TUNEL positive cells, the number of TUNEL-positive and total tumor cells was counted in 5 random fields at x400 magnification. The apoptosis index was calculated as the ratio of TUNEL-positive to total tumor cells. DAB-stained sections were examined via a BX51T-PHD-J11 microscope (Olympus Co., Tokyo, Japan). Images were captured and analyzed using Image-Pro Plus software (Media Cybernetics, Rockville, MD, USA).

Total RNA extraction and real-time PCR. Total RNA was extracted from flash-frozen tumor tissue using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's instructions. Then 5 µg of total RNA was reverse-transcribed in a 50 µl reaction by TIANscript RT kit (Tiangen Biotech, Beijing, China). Real-time PCR was performed with Super Real PreMix Plus (Tiangen Biotech) in the ABI 7500 Real-time PCR System (Applied Biosystems, Foster City, CA, USA). The ther-mocycling conditions were as follows: pre-denaturation at 95°C for 30 sec, followed by 40 cycles at 95°C for 10 sec, 58°C for 30 sec, and 70°C for 30 sec. We determined the expression of VEGF and HIF-1α mRNAs, using the following PCR primers: for VEGF, forward 5'-GGAGCGTTCACTGTGAGC-3', reverse 5'-GCCGAGTCTGTGTTTTTGC-3', and amplified fragment length of 96 bp; for HIF-1α, forward 5'-CTGGAAACGAGTGAAAGG-3', reverse 5'-ATGCTAAATCGGAGGGTA-3', and amplified fragment length of 86 bp; for β-actin, forward 5'-GGCACACCACCTTCTAC-3', reverse 5'-CTGGGTCATCTTTCAC-3', and amplified fragment length of 107 bp. The housekeeping gene β-actin served as an internal control. Relative mRNA expression levels were calculated by the 2-∆∆Ct method in the ABI 7500 Sequence Detection system (Applied Biosystems), according to the manufacturer's protocol.

Protein extraction and western blotting assay. Total protein was extracted from tumor tissues with lysis buffer (Beyotime
Institute of Biotechnology, Shanghai, China). The lysates were ultra-sonicated and centrifuged at 12,000 x g at 4°C for 10 min. Subsequently, the supernatant was transferred into new tubes. Protein concentrations were determined by BCA assay (Beyotime Institute of Biotechnology). Protein lysates (20 µg) were then separated by 10% SDS-PAGE, and transferred to PVDF membranes. After blocking with TBS/5% skim milk, the membranes were incubated with rabbit anti-mouse monoclonal anti-VEGF antibody (1:1000; Abcam) and rabbit anti-mouse monoclonal anti-HIF-1α antibody (1:1000; Abcam) for 2 h at room temperature, respectively. They were then washed with TBST, and incubated with HRP-conjugated goat anti-rabbit IgG secondary antibodies (1:10,000; CoWin Biotech, Beijing, China) for 1.5 h at room temperature. As a loading control, the blots were also probed with rabbit anti-mouse monoclonal anti-β-actin antibody (1:1,000; Abcam). Proteins were then visualized using ECL Plus Detection Reagents (Beyotime Institute of Biotechnology) with UVP GDS-8000 System (Thermo Scientific Inc., Waltham, MA, USA). Bands were quantified using ImageJ software (National Institutes of Health, Bethesda, MD, USA).

Statistical analysis. The data were obtained from at least three independent experiments and each experiment used three parallel samples. All results are presented as mean ± standard error. Differences between means in more than two groups were analyzed by ANOVA followed by Student-Newman-Keuls test. Differences between means in two groups were compared using the unpaired-sample t-test. All statistical analyses were done using SPSS software, version 22.0 for Windows (SAS Institute, Cary, NC, USA). Differences were considered significant at P-values <0.05.

Results
Effect of 125I seed irradiation on tumor growth of lung cancer. Tumor xenografts consisting of transplanted human lung adenocarcinoma A549 cells were used to evaluate the antitumor effects of the 125I seeds. When tumors reached approximately 300 mm³ at day 24, twenty seeds with the radio dosage of 0.6 or 0.8 mCi were implanted into mice of respective groups. In addition, to account for possibility of local effects, we also implanted sham seed with the radio dosage of 0 mCi on a separate group of animals. All seeds were well located without loss and displacement and removed. To confirm its enhanced tumor suppression, we tested tumor growth. As shown in Fig. 1A, there were no significant changes in tumor volume during the first 3 weeks after seed implantation, but after that, tumors of 0.6 or 0.8 mCi group were much smaller than 0 mCi or the
control group and significant differences in tumor volumes were observed between the 0.6 or 0.8 mCi group and the control group (all $P<0.05$). Tumor volumes on day 44 (day 20 after treatment) in 0.6 mCi (627.517±116.251) or 0.8 mCi (525.763±81.141) groups were significantly suppressed when compared with control group (1021.040±183.301; $P=0.012$, $P=0.006$), with an average decrease in tumor size of about 38% or 49% (Fig. 1B). Tumor volumes in 0 mCi or control groups increased exponentially with no regression, and no significant differences in tumor volumes were observed on day 54 between 0 mCi and control groups ($P=0.124$). 0.6 or 0.8 mCi group also decreased tumor weight by 49 and 62% compared with control group (all $P<0.05$; Fig. 1C). This indicated that 125I seed irradiation significantly inhibited growth of the tumors during the 4- to 5-week treatment. Besides, none of the mice died during the treatment, and no significant complications associated with seed implantation were observed in vital organs. These results underscore the safety of 125I seed treatment.

Effect of 125I seed irradiation on morphology of lung cancer.

To investigate the effect of the 125I seed irradiation on the histology of the A549 xenografts, tumor sections obtained from mice in four groups were stained using H&E. As shown in Fig. 2, the histological appearance of the tumors in 0 mCi and control groups was quite different from that in 0.6 and 0.8 mCi groups. In 0 mCi and control groups, the cancer cells were densely arranged with large darkly stained nuclei and obvious karyokinesis (Fig. 2A and B). Whereas, the cancer cells around the 125I seed in 0.6 and 0.8 mCi groups revealed broad necrosis after treatment (Fig. 2C and D). The cancer cells adjacent to the necrotic region were loosely arranged with condensed nuclei and reduced eosinophilic cytoplasm. These results indicated that 0.6 or 0.8 mCi seed implantation caused growth inhibition of cancer cells in the A549 xenografts.

**Figure 3.** Proliferation in tumor tissues is decreased in lung cancer xenograft irradiated with 125I seeds. (A) Immunohistochemical images of proliferation marker Ki-67 in tumors harvested from animals. Representative sections are depicted (x400). (B) Quantification of Ki-67-positive cells as described in Materials and methods. Error bars represent SEM. *$P<0.05$, **$P<0.01$, ***$P<0.001$ compared with control group.

Effect of 125I seed irradiation on proliferation, MVD, and apoptosis.

To determine possible mechanisms underlying 125I seed-mediated suppression of tumor growth, we hypothesized that 125I seed mediates its antitumor effects via inhibition of proliferation. Therefore, we first examined its anti-proliferative effects by performing Ki67 immunohistochemistry on tumors obtained at necropsy from the four groups. The results in Fig. 3A and B showed that the proliferation indices were significantly decreased in 0.6 mCi (46.200±8.349) and 0.8 mCi (38.600±6.025) group compared with control group (71.000±10.000; $P<0.001$, both). However, there was no significant difference in the proliferation indices between 0 mCi group (63.200±6.221) and control group ($P=0.059$). The tumor tissues were stained with CD34,
an endothelial marker, to measure MVD and thus ascertain whether tumor angiogenesis might be affected following 125I seed therapy. MVD, calculated as a measure of angiogenesis, was significantly lower in 0.6 mCi (7.400±1.140) and 0.8 mCi (5.800±1.924) groups compared with control group (12.400±1.517; P<0.001, both; Fig. 4A and B). Treatment with 0.6 or 0.8 mCi seed implantation resulted in a 40% or 53% decrease in MVD compared with controls, respectively. Treatment with 0 mCi (11.600±1.140) seed implantation did not result in a significant decrease in MVD compared with controls (P=0.366). To further elucidate the mechanism by which 125I seed irradiation elicits its antitumor effects, we examined tumor cell apoptosis using the TUNEL assay. As shown in Fig. 5, the apoptotic indices in control and 0 mCi groups were 27.00±4.69% and 35.50±3.42%, respectively. Further, the apoptotic indices in 0.6 and 0.8 mCi groups were 50.00±2.58% and 62.33±4.51%. There was statistically significant difference in apoptotic indices between 0.6 and 0.8 mCi groups and control group (P<0.001, both). These findings suggest that treatment with 125I seed irradiation causes apoptosis in tumor cells.

125I seed irradiation downregulates the expression of VEGF and HIF-1α in tumors. It has been demonstrated that a high level of HIF-1α in the tumor microenvironment leads to enhanced angiogenesis and proliferation. Given that its gene target VEGF is also a critical regulator for neovascularization (24,25), we questioned whether 125I seeds mediate its effects through modulation of HIF-1α and VEGF. Therefore, we determined the HIF-1α and VEGF mRNA expression by RT-PCR in A549 xenograft after treatment in all four cohorts of animals. Expression of HIF-1α and VEGF mRNA in the 0.6 mCi (0.370±0.086, 0.279±0.066) and 0.8 mCi (0.278±0.065, 0.215±0.062) groups were significantly lower than in the control group (1.055±0.078, 1.073±0.103, all P<0.001; Fig. 6A). Moreover, HIF-1α and VEGF mRNA expression did not differ between 0 mCi group (0.810±0.006, 0.923±0.089) and control group (P=0.150, 0.236). These data suggest that 125I seed irradiation significantly affects the expression of HIF-1α and VEGF mRNA. Expression of HIF-1α and VEGF were also examined by western blotting (Fig. 6B). As shown in Fig. 6C, HIF-1α and VEGF protein expression decreased in the 0.6 mCi (0.554±0.0585, 0.396±0.010) and 0.8 mCi (0.502±0.0582, 0.385±0.0842) groups compared with the control group (0.852±0.0395, 0.810±0.0593; all P<0.001). However, there were no significantly statistical differences in HIF-1α and VEGF protein expression between 0 mCi group (0.810±0.006, 0.923±0.089) and control group (P=0.157, P=0.191). These data suggest that 125I irradiation significantly affects HIF-1α and VEGF protein expression.

Figure 4. Angiogenesis in tumor tissues is decreased in lung cancer xenografts irradiated with 125I seeds. (A) Sections of tumors developing in nude mice were stained with anti-CD34 monoclonal antibody (x400). (B) Quantification of MVD in tumors as described in Materials and methods. Error bars represent SEM. *P<0.05, **P<0.01, ***P<0.001 compared with control group.
Discussion

NSCLC is the most prevalent type of lung cancer. Despite therapeutics advances in surgery, chemotherapy, and radiotherapy, the five-year overall survival rate is poor. Radiotherapeutics approach has recently started to play an important role in the treatment of advanced lung cancer. However, the adverse effects of conventional EBRT on surrounding organs pose a major problem. The advantages of 125I seed radiation are the low dose rates and conformal irradiation which increases the dose applied within the target area, thus decreasing the incidental radiation injury to normal tissues and the attendant complications (26,27). Several recent studies suggest that apoptosis and proliferative inhibition may have important roles in the therapeutic effects of 125I seeds (18,28), but the mechanism in the treatment of lung cancer, especially in animal models, is not known completely. In addition, the HIF-1α and VEGF may be involved in the lung cancer tumorigenesis (29). Irradiation therapy may downregulate the expression of HIF-1α and VEGF in tumors, thereby affecting angiogenesis and then affecting tumor growth (30,31). However, comprehensive knowledge on this topic, particularly at the molecular level, is still lacking. In this study, we assessed the radiobiological effects of 125I seeds on human lung adenocarcinoma cells in vivo.

A549 lung cancer cells were cultured ex vivo and implanted subcutaneously into the nude mice to create the animal model. We examined the effects of irradiation on tumor growth and found that radioactive 125I seed significantly inhibited tumor growth in nude mice. By observing H&E-stained slides, large necrotic regions were observed around the 125I seed in tumors. Immunohistochemistry staining indicated that the expression of Ki67 and microvessel density (CD34) was inhibited by 125I seed irradiation in tumor tissues. The Ki67, which is a proliferative cell marker expressed in all phases of the cell cycle except the G0 stage, is considered to be a reliable index of the proliferation rate (32). Angiogenesis which is required for the delivery of oxygen and nutrients in the tumor tissues, possibly resulting in the apoptosis in vivo, was inhibited by 125I seeds. To test the effect of 125I seed radiation on apoptosis, we assayed apoptosis using TUNEL staining. Our data showed that 125I seed significantly induced apoptosis in tumor tissues, indicating that 125I seed may indeed inhibit tumor growth through the inhibition of angiogenesis. These data suggest that 125I seed radiation is an effective radiotherapeutics approach for lung cancer through inhibiting cell proliferation, angiogenesis and inducing apoptosis in vivo.

Hypoxia is known to be a hallmark of solid tumors and increased angiogenesis (33); under hypoxia condition, HIF-1

Figure 5. Apoptosis in tumor tissues is increased in lung cancer xenografts irradiated with 125I seeds. (A) Representative TUNEL staining images of tumor cells are depicted. (B) The average ratio of TUNEL-positive cells in the 0.6 and 0.8 mCi implanted groups were significantly increased over those in the control group and 0 mCi seed implanted groups (all P<0.001). All images were taken at original magnification x400. Error bars represent SEM. """"P<0.001 compared with control group.
regulates expression of numerous angiogenic genes and stimulate angiogenesis as a long-term solution to the hypoxic conditions (34,35). HIF-1 is a heterodimeric transcription factor consisted of HIF-1α and HIF-1β subunits (36). HIF-1α is an important mediator induced by hypoxia, growth factors, and oncogenes (37). As a transcription factor involved in the process of gene related hypoxic adaptation of neoplasm, HIF-1α is often upregulated in human cancers. HIF-1α activates the transcription of many genes such as VEGF, endothelin-1, and inducible nitric oxide synthase, which are involved in vasodilatation, neovascularization, and tumor metastasis (38). Many investigations have revealed that the relationships between HIF-1α, VEGF and angiogenesis resulted from hypoxic condition (39,40). To test whether 125I seed radiation affects HIF-1α levels, we analyzed HIF-1α expression by real-time PCR and western blotting and showed that 125I seed specifically inhibited HIF-1α expression in cancer cells.

VEGF is known to be an important angiogenic factor for endothelial cells. Many studies recently showed that VEGF levels correspond with advanced lung cancer. The increased VEGF levels were found to be associated with poor prognosis in patients with NSCLC (41,42). In addition, the activation of VEGF expression can be induced by HIF-1α, as indicated above. Our data showed that 125I seed radiation inhibited VEGF expression at the transcriptional level and protein level. HIF-1 expression is known to play an important role in VEGF transcriptional activation in response to hypoxia (34). Thus, 125I seed radiation may inhibit VEGF transcriptional level through the decrease of HIF-1 expression in cancer cells. This result is consistent with the data showing the inhibition of angiogenesis by 125I seed because HIF-1α and VEGF are important for tumor angiogenesis. Our study demonstrated that HIF-1α and VEGF play an important role in the therapeutic effects of continuous low-energy 125I irradiation and are involved in the mechanism of the 125I seed implantation therapy process.

Collectively, it was shown that 125I seed radiation can elicit significant anticancer effects by inhibiting proliferation and inducing apoptosis in vivo. In the tumor microenvironment, reducing the levels of HIF-1α and VEGF induced by 125I seed might inhibit angiogenesis. These findings suggest that the 125I seed radiation may be a promising therapeutics approach against lung cancer.

References