In Vitro and In Vivo Imaging of Prostate Cancer Angiogenesis Using Anti-Vascular Endothelial Growth Factor Receptor 2 Antibody-Conjugated Quantum Dot

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Objective: Authors aimed to determine the targeting ability of vascular endothelial growth factor receptor 2 (VEGFR2)-conjugated quantum dots (QDs) in vitro, and apply it for a xenograft prostate cancer mouse model.

Materials and Methods: Conjugation reaction of QDs was performed by using the N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide (EDC) and sulfo-(N-hydroxysulfosuccinimide) (Sulfo-NHS). The human umbilical vein cord endothelial cells (HUVECs) were incubated with QDs, conjugated with anti VEGFR2, to see a specific binding in vitro. Fluorescent cell images were taken by a confocal microscope. The human prostate cancer cells (PC3) were injected to five nude mice on hind limbs to make the xenograft tumor model. QD-antiVEGFR2 antibody complex was injected into the tumor model and fluorescence measurements were performed at 1, 4, 9, 12, 15, and 24 hours after the injection.

Results: The specific interaction between HUVECs and QD-antiVEGFR2 antibody was clearly shown in vitro. The in vivo fluorescence image disclosed that there was an increased signal of tumor, 12 hours after the injection of QDs.

Conclusion: By showing endothelial cells binding with QDs-antiVEGFR2 antibody and an experimental application of the antibody for VEGFR2 imaging in the prostate cancer xenograft mouse model, we suggest that the antibody-conjugated QDs can be a potential imaging tool for angiogenesis of the cancer.

Index terms: Quantum dot; VEGFR2; Angiogenesis; Prostate cancer; Near infrared

INTRODUCTION

Angiogenesis is the formation of new capillaries from existing blood vessels, and plays an important role for tumor growth and metastasis by providing oxygen and nutrients to the proliferating tumor cells (1-3). Moreover, the angiogenic activity is known to be related to tumor malignancy (4, 5). Therefore, angiogenic activity is highly relevant to the tumor state, and several probes were suggested to make a possible noninvasive detection. This strategy for tumor diagnosis may be expanded to monitoring of the tumor progression by drug treatment. Recently, the molecular imaging techniques have helped to develop a direct visualization and characterization of the cellular or molecular activation of angiogenesis-related phenomena (6). The role of the molecular imaging are becoming increasingly
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important for studying this process, in both clinical and research settings, ranging from identifying the sites of angiogenesis to confirming the process of biochemistry that lies at the basis of the angiogenic process (7).

Quantum dots (QDs) are semiconductor nanocrystals with a quantum confinement property, which enables them to emit fluorescence from visible to infrared wavelengths on excitation (8). Over the last few years, due to their bright fluorescence, excellent photo-stability, and their narrow and tunable emission spectrum, QDs have gained much interest for biological imaging purposes (7).

Several specific endothelial molecular markers of angiogenesis are overexpressed on tumor vascular endothelial cells. Vascular endothelial growth factor receptor 2 (VEGFR2) is one of the major regulators of angiogenesis. Overexpression of VEGFR2 has been associated with tumor progression and poor prognosis in several tumors, including gastrointestinal tract, pancreas, breast, prostate, and lung cancers (9). Therefore, the imaging modalities, which can directly visualize specific molecular markers of angiogenesis, such as VEGFR2, would be very helpful for monitoring the angiogenic treatments and detection of the tumors.

The purpose of this study is to prepare anti-VEGFR2 antibody, conjugated QDs, in order to confirm the targeting ability in vitro, and to evaluate the feasibility of the use of QDs in xenograft prostate cancer mouse models.

MATERIALS AND METHODS

Conjugation of AntiVEGFR2 Antibody to QDs

One nmol of antiVEGFR2 antibody (Santa Cruz Biotechnology Inc., Santa Cruz, CA, USA) was mixed with 10000 equivalent of N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide (EDC, Sigma-Aldrich, St. Louis, MO, USA) and 12000 equivalent of sulfo-NHS (Molecular Bioscience, Boulder, CO, USA) in 0.1 M 2-(N-morpholino)ethanesulfonic acid buffer (pH 4.7, Pierce, Rockford, IL, USA). After incubation for 30 minutes at 25°C, the excess of reagents were removed by 50 k microcon centrifugal filter (Millipore, Billerica, MA, USA). The activated antiVEGFR2 antibody was then reacted with functionalized QD with amino PEG by the previously reported method (10) in 1 x phosphate buffered saline (PBS, Invitrogen, Eugene, OR, USA) for 3 hours at 25°C. Finally, both QDs530-antiVEGFR2 and QDs800-antiVEGFR2 conjugated were purified, using the Superdex® 200 size exclusion column (GE Healthcare, Pittsburgh, PA, USA), and have been characterized by an agarose gel electrophoresis and dynamic light scattering (DLS). For the in vitro study, QDs530-antiVEGFR2 conjugates were used, and for the animal study, QDs800-antiVEGFR2 conjugates were used. The gel electrophoresis was run in a 0.5 x TBE buffer (Tris/borate/ethylenediaminetetraacetic acid, Sigma-Aldrich, St. Louis, MO, USA), using a 1% agarose gel at 50 V for 1 hour. To identify the QD-antibody conjugate, the gel was post-stained with colloidal blue solution. The number of VEGFR2 per QDs was determined by a bicinchoninic acid (BCA) protein assay (Thermo Scientific, Rockford, IL, USA).

HUVEC Cell Culture

Human umbilical vein cord endothelial cells (HUVECs) were purchased from company (Innopharma Screen, Asan, Korea). The cells were grown in M199 media, supplemented with penicillin-streptomycin 10 mL/L, 25 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES), 10 unit/ml Heparin, 2.2 g/L sodium bicarbonate, and 20% fetal bovine serum (FBS)/b-fibroblast growth factor 20 ng/mL.

In Vitro Cell Imaging Using QDs-AntiVEGFR2 Antibody

The interaction of QDs-antiVEGFR2 antibody with the cells was investigated, using a confocal fluorescence microscope (LSM 5 Live, Carl Zeiss, Oberkochen, Germany), as reported previously (11). HUVEC (10000 cells/well) were seeded in 8-well chambered cover glass (Lab-Tek, Thermo Scientific, Rochester, NY, USA) with 400 µL of cell media. After 24 hour incubation at 37°C, the cells were fixed by 4% paraformaldehyde (Wako, Osaka, Japan) for 20 minutes, and were washed by PBS, several times. A 150 nM of QDs530 and QDs530-antiVEGFR2 antibody conjugates were added to each well. They were incubated for 3 hours. To remove the non-specific binding of QDs, the cells were washed by PBS and 0.01% tween20. Nucleus was stained by 4', 6-diamino-2-phenylindole (DAPI). For excitation of QDs, 488 Ar-ion laser was used with emission band pass filter of 495-555 nm, while the DAPI was excited with 405 nm laser and observed with emission band pass filter of 420-480 nm. A 40 x water objective was used for obtaining all of the fluorescent images and differential interference contrast (DIC) images.

PC3 Prostate Tumor Model

Five male Athymic nude mice of 6-7 weeks old were obtained from an animal facility (Orient, Seoul, Korea) and were housed under a specific pathogen-free environment.
The mice were maintained under controlled conditions (12-hour dark-light cycles; temperature, 20-24.8°C) with free access to sterilized mouse chow.

The PC-3 was purchased from the cell line bank (Korean Cell Line Bank, Seoul, Korea). The cells were grown in RPMI 1640 medium, supplemented with 10% FBS and penicillin-streptomycin. The cells were cultured at 37°C in a humidified 5% CO₂/95% air atmosphere.

To generate the tumor cells, prostate cancer cells (PC3) (1.5 x 10⁶ cells in PBS 0.2 mL) was injected, subcutaneously, into the back or the right flank of the mice. After 4 to 6 weeks, tumors were allowed to grow to the mean maximum diameter of 15 mm (range, 10-15 mm). All animal protocols were approved by the Institutional Animal Care and Use Committee.

**In Vivo Imaging of QDs and QDs-VEGFR2**

Fluorescent images were obtained, using a Maestro in vivo Imaging System (CRi Inc., Woburn, MA, USA) for data acquisition and analysis. Before imaging, the mice were anesthetized by intraperitoneal injection of a solution containing 8 mg/mL ketamine (Ketalar®, Panpharma, Fougères, France) and 0.8 mg/mL xylazine (Rompun®, Bayer Pharma, Puteaux, France) at 0.01 mL/g of body weight.

QDs800-antiVEGFR2 antibody complex (140 pmole/mice) was injected via retro orbital route into three mice. Fluorescence measurements were performed at 1, 4, 9, 12, 15 and 24 hours after the injections. As a control group, control QDs800 without VEGFR antibody complex (140 pmole/mice) was injected into two mice.

In all cases, optical image sets were composed of a Cermax®-type 300 Watt Xenon lamp with a green filter set (a band-pass filter from 503 to 548 nm and 560 nm longpass filter, which were used for excitation and emission, respectively) to acquire one complete image cube. The tunable filter was automatically increased in 10-nm increments from 560 to 750 nm. A camera was used to capture the images at each wavelength, using a constant exposure. The system control was done with LabVIEW (National Instruments cooperation, Austin, TX, USA). The images were transferred to Image J software (http://rsbweb.nih.gov/ij/) as a JPEG format. The regions of interest (ROIs) were drawn on the tumor site, and the hind limb as a signal of background. This method was adapted from the other published study by Schellenberger et al. (12). We measured the ROI values three times from different locations, and the mean values were used for representative values.

After imaging, the mouse was sacrificed and the tumor tissue was prepared for VEGFR2 immunohistochemical staining. Immunostaining of VEGFR2 was performed using polymeric methods (DAKO real envision detection Systems, Glostrup, Denmark). The consecutive sections obtained from formalin-fixed specimen were made into paraffin-embedded blocks, and were deparaffinized. The rehydrated sections were boiled in a 0.01% sodium citrate buffer (pH 6.0), and then incubated with 3% hydrogen peroxide to block the endogenous peroxidase activity. The sections were then probed with monoclonal anti-VEGFR2 antibody (Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA). After incubation with horseradish peroxidase-labelled polymer, they were exposed to chromogen plus 3, 3′-diaminobenzidine substrate, and then counterstained with hematoxylin.

**RESULTS**

Figure 1 illustrates QDs-antiVEGFR2 antibody was synthesized using EDC chemistry. A gel electrophoresis analysis and DLS data of QDs-antiVEGFR2 antibody conjugates are shown in Figures 2, 3. The gel electrophoresis data showed that QDs were first modified with polyethylene glycol (PEG)-NH₂, and anti VEGFR2 antibody was successfully conjugated with QDs-PEG-NH₂ as confirmed with colloidal blue staining. The hydrodynamic diameter of the resulting QDs-antiVEGFR2 antibody conjugate was much bigger than amine-functionalized QDs, which confirmed again that the antiVEGFR2 antibody was attached on QDs. From BCA protein assay (data is not shown here), the number of antiVEGFR2 antibody per QD was about 1.

Human umbilical vein cord endothelial cells were incubated with QDs530-antiVEGFR2 antibody and QDs530 to see a specific binding of QDs-antiVEGFR2 antibody conjugates in the endothelial cells, which express exclusively VEGFR2 on their membrane. Representative
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DIC and fluorescent images were shown in Figure 4. Green fluorescent QDs-antiVEGFR2 antibody was bound to HUVECs, whereas, QDs without the antibody were not observed in the cells. This result showed that there was a specific binding between HUVECs and QDs-antiVEGFR2 antibody.

Figure 5 shows serial fluorescent images after an injection of the antiVEGFR2 antibody conjugated QDs. The images showed that there is an increased signal of tumors after 12 hours of QDs injections. Figure 6 shows a signal to the background ratio, at each time point. As with the findings of optical imaging, the ratio is also increasing after 12 hours of QDs injections.

Figure 7 shows the immunohistochemical staining results of the tumor. There was positive staining of VEGFR2, which confirms that the tumor expressed the VEGFR2.

**DISCUSSION**

Angiogenesis, which is the formation of new functional blood vessels, plays an important role in many disease processes, including ischemia or cancer. Angiogenesis is a critical process in the tumor growth and invasion (13, 14). Many researches, investigating molecular pathways of tumor angiogenesis, have led to the identification of a number of key molecules that are involved in the stimulation of new vessel growth from existing host vasculature. Therefore, it is not surprising that many studies have focused on the angiogenic markers, such as VEGFR or $\alpha_{\nu}\beta_{3}$ receptor (9) as a target for anti-angiogenic drugs.

Quantum dots are dot shape semiconductor particles with diameters on the order of 2-8 nm in visible fluorescent range, which roughly contains 200-100000 atoms. Due to their novel optical and electronic properties, QDs have been intensively studied as a new class of nano probe for molecular, cellular, and *in vivo* imaging (15, 16). Because the emission wavelength of QDs is size-dependent, fluorescence color may be fine-tuned throughout the visible and near infrared (NIR) spectrum, during a chemical
synthesis, by controlling the QD size and composition. The semiconducting nature and the size-dependent fluorescence of QDs have made them very attractive for use in optoelectronic devices and biological detection. In addition, QDs have a continuous absorption band and a narrow Gaussian emission profile. These combined properties enable excitation of multiple-colored QDs with a common excitation source and the detection of multiple colors within a limited spectral window without cross-talk.

Quantum dots conjugated with an antibody have been extensively studied because of their possible application in the bioimaging. QDs conjugated with anti-VEGFR2 antibody have also been studied by Chen et al. for a dual-function positron emission tomography/near-infrared fluorescence probe. Although this research showed QD conjugated with anti-VEGFR2 antibody and its application in tumor imaging, more detailed characterization and specific binding studies will be needed to improve the potential of the QD-antibody conjugates in the area of tumor diagnostics.

The importance of the diagnosis and therapy of prostate cancer is increasing since prostate cancer has become a leading cause of morbidity and mortality in men. The evaluation of angiogenesis is also important in the development and spread of prostate cancer. Several studies have shown the importance of evaluation of tumor neovascularization, because it was definitely correlated with an increased risk of distant metastasis and prostate cancer recurrence after surgery, as well as with poorer overall survival.

The targeted delivery of biocompatible QD conjugates in vivo has been limited to date to detect specific vascular biomarkers in the tissue. Gao et al. used QDs conjugated to monoclonal antibodies directed against prostate-specific membrane antigen to target and image human prostate cancer cells, which are growing in the liver of mice. More recently, arginine-glycine-aspartic acid
peptide-conjugated QDs have been used to specifically target integrin αvβ3 in a murine xenograft model because the integrin αvβ3 is significantly upregulated in the tumor, but not in the normal tissue (17, 24, 25).

In our study, we used NIR QDs for in vivo imaging studies. One of the greatest advantages of QDs for imaging in living tissue is that their emission wavelengths can be
tuned throughout the NIR spectrum, by adjusting their composition and size, resulting in photostable fluorophores that are stable in biological buffers (16). The superiority of NIR QDs has been demonstrated in sentinel lymph node mapping, a common procedure in breast cancer surgery, whereby, the lymph node closest to the organ affected is monitored for the presence of locally disseminated cancer cells (17). However, despite those advantages of NIR emission, locating the fluorescent objects in vivo can still be challenging because tissue absorption and scattering, in addition to limiting the light coming out of the subject, also limit the amount of incident excitation light that reaches the fluorophores. To the best of our knowledge, there is no report regarding imaging of the prostate tumor angiogenesis, using NIR QDs.

With the exciting potential of QDs for clinical application, the potential toxicity of QDs has become a serious concern. Theoretically, the toxicity of QDs can be derived from the leakage of toxic core semiconductor materials (such as CdSe/ZnS), by UV irradiation, and nonspecific uptake through endocytosis, inducing cell apoptosis and the transfer of absorbed optical energy to nearby oxygen molecules, generating reactive oxygen species (ROS) (26-28). It may be of greater concern that QDs, and many other types of nanoparticles, have been found to aggregate, bind nonspecifically to cellular membranes and intracellular proteins, and induce the formation of ROS (15).

In our study, the images of HUVECs showed the targeting ability of QD-antibody conjugates with a specific binding between QDs-antiVEGFR2 and the cells. As shown on Figure 4, confocal microscope images of HUVEC revealed that HUVEC cells with QDs-VEGFR2 showed strong interactions, whereas, the HUVEC with QDs only showed no signals. Our results suggest that we can selectively visualize the cells with high activity of VEGFR2, which can suggest the potential for the diagnosis or therapy onto the specific cells. However, the binding to VEGFR2 cannot be tumor-specific binding, as it can be shown in any tumors that have increased level of VEGFR2 or other conditions with increased VEGFR2.

There are several reports for imaging the angiogenesis in animal and clinical studies (29-31). Jun et al. (29) tried to visualize the tumor angiogenesis using a VEGF receptor 2 antibody conjugated magnetic resonance contrast agent, and Ko et al. (30) evaluated angiogenesis using an ultrasound contrast agent, using animal models. In a clinical study, Kim et al. (31) compared the perfusion parameters in CT with tumor grade and microvessel density.

The in vivo images revealed that there were increased signals from tumors comparing with the surrounding tissues, after 12 hours after injection (Fig. 5). Our findings from in vitro and in vivo images suggest the possibilities that these targeting QDs could be used in living objects, even though many tremendous studies, regarding the optimal dose, toxicity, and biodistribution would be needed.

The main limitation of our study is that the number of in vivo cases is too small to analyze statistically. Out of the five animals used in our study, only one case survived and showed QD signals from tumors, despite its possibilities of NIR QDs in vivo. The remaining four animals were dead during imaging after intraperitoneal anesthesia. We think one of the main reasons is due to the temperature, because there was no heater or warm bed in the chamber of fluorescent imaging system we used. The nude mouse are very vulnerable to the temperature changes or other stress, so caring them required more experience and knowledge concerning caring nude mouse models. The other possibilities include the toxicity of QD. For clinical application of NIR QD, the safety issue needs to be evaluated, thoroughly. Even though the in vivo results are unsatisfactory, we could show the possibilities for imaging tumor in living object with NIR QD from our results from an in vivo study. Further studies, using animal models, would be needed for detailed evaluation in the living object.

In spite of these limitations, we tried to get in vitro and in vivo images regarding VEGFR2 using QDs, which is one of the most potential nanomaterials for molecular imaging.

In conclusion, we showed that the endothelial cells were labeled by QDs-antiVEGFR2 antibody with the specific binding, and in vivo image of the prostate cancer xenograft mouse model indicated a potential application for VEGFR2 imaging in vivo. This approach suggests that antibody conjugated QDs can be a promising detector of cancer angiogenesis in the field of tumor diagnosis.

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