Original Article
OKN-007 decreases VEGFR-2 levels in a preclinical GL261 mouse glioma model

Patricia Coutinho de Souza1,2, Nataliya Smith1, Richard Pody1, Ting He1, Charity Njoku1, Robert Silasi-Mansat3, Florea Lupu3, Bill Meek4, Hong Chen3, Yunzhou Dong3, Debra Saunders1, Albert Orock1, Erik Hodges1, Sarah Colijn1, Nadezda Mamedova1, Rheal A Towner1,2

1Advanced Magnetic Resonance Center, Oklahoma Medical Research Foundation, Oklahoma City, OK; 2Department of Veterinary Pathobiology, College of Veterinary Medicine, Oklahoma State University, Stillwater, OK; 3Cardiovascular Biology Research Program, Oklahoma Medical Research Foundation, Oklahoma City, OK, 4Center for Health Sciences, Oklahoma State University, Tulsa, OK

Received February 20, 2015; Accepted April 20, 2015; Epub June 15, 2015; Published July 1, 2015

Abstract: Angiogenesis is essential to tumor progression, and the precise imaging of the angiogenic marker vascular endothelial growth factor receptor 2 (VEGFR-2) may provide an accurate evaluation for angiogenesis during a therapeutic response. With the use of molecular magnetic resonance imaging (mMRI), an in vitro cell assay indicated significantly decreased T1 relaxation values when tumor endothelial cells (TEC), which positively expressed VEGFR-2 (Western blot), were in the presence of the VEGFR-2 probe compared to TEC alone (P < 0.001). For in vivo mMRI evaluations, we assessed VEGFR-2 levels in untreated and OKN-007-treated GL261 mouse gliomas. Regarding treatment response, OKN-007 was also able to significantly decrease tumor volumes (P < 0.01) and increase survival (P < 0.001) in treated animals. Regarding in vivo detection of VEGFR-2, OKN-007 was found to significantly decrease the amount of VEGFR-2 probe (P < 0.05) compared to an untreated control group. Fluorescence imaging for the VEGFR-2 probe indicated that there was colocalization with the endothelial marker CD31 in an untreated tumor bearing mouse and decreased levels for an OKN-007-treated animal. Immuno-fluorescence imaging for VEGFR-2 indicated that OKN-007 treatment significantly decreased VEGFR-2 levels (P < 0.0001) when compared to untreated tumors. Immuno-electron microscopy was used with gold-labeled anti-biotin to detect the anti-VEGFR-2 probe within the plasma membrane of GL261 tumor endothelial cells. This is the first attempt at detecting in vivo levels of VEGFR-2 in a mouse GL261 glioma model and assessing the anti-angiogenic capability of an anticancer nitrone. The results indicate that OKN-007 treatment substantially decreased VEGFR-2 levels in a GL261 glioma model, and can be considered as an anti-angiogenic therapy in human gliomas.

Keywords: Molecular magnetic resonance imaging, glioma, VEGFR-2, angiogenesis, in vivo, GL261

Introduction

Angiogenesis, the formation of new blood vessels from preexisting vasculature, is a hallmark of cancer which plays pivotal roles in tumor development and metastasis [1-3]. Physiological angiogenesis is tightly regulated by pro-endothelial and anti-endothelial growth factors (i.e. endogenous inhibitors) and occurs by a series of complex and interrelated steps [2, 4, 5]. One of the most potent proangiogenic agents is the well-characterized vascular endothelial growth factor (VEGF) [5]. VEGF acts on vascular endothelial cells through compatible receptor tyrosine kinases VEGF receptor (VEGFR)-1, -2, and -3 [5]. VEGFR-1 is required for the recruitment of haematopoietic stem cells and the migration of monocytes and macrophages, VEGFR-2 regulates vascular endothelial function, and VEGFR-3 regulates lymphatic endothelial cell function [5]. In addition, VEGFR1 and VEGFR2 are highly expressed in different types of cancers such as gliomas [6]. Non-invasive and quantitative imaging of tumor angiogenesis is critical for lesion detection, patient stratification, and monitoring the therapeutic response of cancer patients, including brain tumors [7, 8].

Of the many different types of gliomas, glioblastoma (GBM) is the most common, accounting for about 40% of all primary brain tumors [9].
GBM is also one of the most vascular [10] and deadly cancers, with a very low 5-year survival rate of 6% [11]. A number of studies have shown that overexpression of VEGF and/or VEGFRs correlated with poor prognosis in multiple cancer types, including gliomas [12]. Furthermore, it has been also reported that intra-tumoral levels of VEGF and VEGFRs correlate with the histological grade of gliomas [13]. Knowing the dynamic distribution of VEGFRs, especially VEGFR-2, will lead to better understanding of the mechanisms underlying tumor angiogenesis, as well as provide a better tool for cancer diagnosis and treatment [14-15]. Currently, anti-angiogenic agents are routinely used for the treatment of patients with gliomas [16]. However, despite advances in pharmacological and surgical therapy, gliomas remain an incurable disease [16], highlighting the need for new treatment approaches.

OKN-007 (2, 4-disulfophenyl-PBN; or disodium 4-[(tert-butyl-imino) methyl] benzene-1,3-disulfonate N-oxide or disufenton; also previously known as NXY-059) is a very effective compound against in vivo adult glioma models [17-19], and it is currently undergoing clinical trial assessment as a new investigational drug for recurrent adult GBMs. OKN-007 is a small molecule that can traverse the blood-brain barrier and has also anti-inflammatory, antioxidant, pro-apoptotic [20, 21], and anti-angiogenic properties [18, 19].

Recently, our group used molecular magnetic resonance imaging (mMRI) to evaluate the heterogeneous expression of VEGFR-2 in C6 and RG2 rat glioma models, using a specific VEGFR-2 contrast agent [22]. The study indicated that the C6 gliomas have more VEGFR-2 occurring in the relatively large vessels within the tumor periphery, whereas the RG2 glioma may have increased VEGFR-2 in the microvessels within the tumor interior [22].

In this study, we wanted to assess the anti-angiogenic potential of the anti-cancer agent, OKN-007, using VEGFR-2 targeted mMRI in a mouse GL261 glioma model. Fluorescence imaging to confirm that the VEGFR-2 probe is associated with endothelial cells and that OKN-007 directly affects VEGFR-2 levels, as well as in vitro binding affinity assessment of the VEGFR-2 probe for tumor endothelial cells (TEC), was also conducted.

### Materials and methods

#### Synthesis of anti-VEGFR-2 MRI contrast agent

The contrast agent, biotin-BSA (bovine serum albumin)-Gd-DTPA, was prepared as previously described by our group [23], based on the modification of the method developed by Dafni et al. [24]. The estimated molecular weight for the biotin-albumin-Gd-DTPA moiety is ~80 kDa. It is estimated that there are 1.3 biotin and 23 Gd-DTPA groups bound to each BSA molecule. Briefly, anti-VEGFR-2 mAb (Santa Cruz Biotech, Inc., CA, USA) was conjugated to the albumin moiety through a sulfo-NHS (N-succinimidyl-S-acetylthioacetate)-EDC (N-succinimidyl 3-(2-pyridyldithio)-propionate) link according to the protocol of Hermanson [25]. Each animal was injected with 200 μl anti-VEGFR-2-BSA-Gd-DTPA-biotin (VEGFR-2 probe) [22, 26, 27] intravenously via the tail vein with an amount estimated to be 200 μg anti-VEGFR-2 and 100 mg biotin-BSA-Gd-DTPA per injection. The estimated molecular weight of the VEGFR-2 probe is 232 kDa (see Figure 1A). To verify the molecular weight of the VEGFR-2 probe, electrophoresis was done. Briefly, the probe was mixed 1:1 with sample buffer, and boiled for 2 min. The sample was then pipetted into a gel (TGX gradient 4-15% gel from Bio-Rad), and the gel was run at 100 V constant for 90 min. The protein bands were visualized using Coomassie Brilliant Blue. For the immuno-electron microscopy study, an IgG contrast agent was used for a non-specific isotype control [22, 27].

#### In vitro assessment of the anti-VEGFR-2 probe

Tumor endothelial cells (TEC) were isolated according to a protocol described by Miltenyi Biotech (Miltenyi Biotech Inc., San Diego, CA). Briefly, Lewis lung Carcinoma (LLC) tumors (~1 cm diameter) were isolated from s.c graft tumors from C57BL/6 mice. Isolated tumor tissue was enzymatically digested (mixture of collagenases, DNAse and Dispases) for 1 hour, followed by depletion of CD45+ cells using CD45 microbeads (Miltenyi Biotech Inc.). In order to separate tumor endothelial cells, the cells were further passed through anti-CD31 microbeads (Miltenyi Biotech Inc.), and CD31 positive cells were eluted and cultured in endothelial medium (Lonza Group Ltd., Allendale, NJ). Cultured TEC cells were confirmed using an endothelial specific marker, vWF (Abcam,
OKN-007 decreases VEGFR-2 in GL261 gliomas


Cambridge, MA, and VEGFR2 (Cell Signaling Technology, Danvers, MA) expression was confirmed with fluorescence staining and Western blot assessments. Mouse GL261 cells were grown in flasks in complete growth medium (DMEM Media with 10% fetal bovine serum (FBS), Invitrogen, Grand Island, NY, USA) to confluence. Two to three hours before treatment, the growth medium was replaced with serum-free medium.

For analysis of VEGFR2 expression, tumor endothelial cells (TEC) or bovine aortic EC were lysed with a lysis buffer (50 mM Tris-HCl, pH 6.8, 5% β-mercaptoethanol, 2% SDS, and protease inhibitors). After addition of 4× SDS sample buffer and incubation at 95°C for 5 min, samples were subjected to SDS-polyacrylamide gel electrophoresis and transferred to Amersham Protran 0.45 μm NC membrane (GE Healthcare Life Sciences). The membranes were then probed with anti-VEGFR2 antibody (Cell Signaling Technology) visualized by SuperSignal West Pico Chemiluminescent Substrate (Thermo Scientific) as described previously [28].

For in vitro MRI assessment of the anti-VEGFR2 probe, vials were prepared containing TEC (10^6) alone, TEC with the anti-VEGFR2 probe, or water (no cells). In the samples that contained the probe, 2 μg of the anti-VEGFR2 probe (based on antibody calculation), and cells were incubated for 45 min. Following incubation, cells were collected, washed with PBS, centrifuged (500 rpm), and the pellet was resuspended in PBS for MR imaging. Each measurement was repeated 6 times per treatment group. MRI signal intensities were obtained using RARE sequence (Rapid Acquisition with Relaxation Enhancement) [repetition time (TR) 1300 ms, echo time (TE) 9.0 ms, refocusing flip angle 180°, 256 × 256 matrix, 4 signal averages per acquisition, 3.50 × 3.50 cm² field of view (FOV), 1 mm slice thickness], T₁ relaxation times were obtained using a RARE-variable TR (TR 200, 400, 800, 1200, 1600 and 2000 ms; TE 15 ms; 256 × 256 matrix; 2 signal averages per acquisition; 3.50 × 3.50 cm² FOV; slice thickness 1.0 mm). T₁ relaxation times were calculated using Paravision (5.0) software.

**Intracerebral glioma cell implantation**

All animal studies were conducted with the approval of the Oklahoma Medical Research Foundation Institutional Animal Care and Use Committee. As a model for orthotopic intracranial brain tumors, a GL261 mouse glioma model was used (n = 17). GL261 mouse glioma cells were implanted intracerebrally in C57BL6/J mice. The heads of anesthetized...
mice were immobilized (stereotaxic unit; Kopf Instruments, Tujunga, CA), and with aseptic techniques, a 1 mm burr hole was drilled in the skull 1 mm anterior and 2 mm lateral to the bregma on the left side. A 20 μL gas-tight Hamilton syringe was used to inject $2 \times 10^4$ GL261 cells (in 10 μL of PBS) into the left frontal lobe at a depth of 1.5 mm relative to the dural surface in a stereotaxic unit. The cell lines were maintained and expanded immediately prior to inoculation. Following injection, the skin was closed with surgical sutures.

**OKN-007 treatment**

OKN-007 (2, 4-disulfophenyl-N-tert-butyl nitro- ne, Ryss Laboratories, Union City, CA) was administered to the mice in their drinking water at a concentration of 150 mg/kg/day (0.20% w/v for a 20 g mouse). The treatment started at 13-15 days after glioma cell implantation, when the tumor volumes were between 10 and 15 mm$^3$. The treatment was administered continuously until the end of the study. Mice receiving normal drinking water were used as UT controls. The amount of OKN-007 consumed by each mouse, which were housed in separate cages, was determined by weighing water bottles each day. No significant deviation was observed in the volume of liquid uptake of compound in these mice. The average intake of OKN-007 was approximately 140-150 mg/kg/day. Mortality was recorded daily along the study period to calculate the percentage survival of all animals.

**In vivo magnetic resonance techniques**

**Morphological imaging:** Nude mice were anesthetized and positioned in a stereotaxic cradle. A 30-cm horizontal bore Bruker Biospin magnet operating at 7 Tesla (T; Bruker BioSpin GmbH, Karlsruhe, Germany), was used with a S116 gradient set to perform all MRI experiments. A mouse head coil was used for signal detection and a 72 mm quadrature volume coil for transmission. Multiple-slice, multiple echo (MSME) imaging (FOV = 2.50 × 2.50 cm$^2$, TR = 2000 ms, TE = 17.5 or 58.2 ms, matrix = 192, averages = 2, slices = 16, slice thickness = 1 mm) was used to calculate tumor volumes and to inspect tumor morphology. Starting ten days after the GL261 tumor cells inoculation, each mouse brain was imaged in vivo every 2-3 days until the end of the study. Tumor volumes were measured independently by at least three individuals for each dataset.

**Molecular MRI:** Molecular MRI was performed when the tumor volumes were close to their maximum tumor volumes (120-180 mm$^3$). A variable-TR RARE sequence (rapid acquisition with refocused echoes, with multiple TRs of 200, 400, 800, 1200 and 1600 ms, TE of 15 ms, FOV of 3.5 × 3.5 cm$^2$, matrix size of 256 × 256 and a spatial resolution of 0.137 mm) was used to obtain $T_1$-weighted images before and after administration of probe or control contrast agents.

**Calculation of relative probe concentration:** Relative probe (contrast agent) concentrations were calculated to assess the levels of VEGFR-2 in each animal. A contrast difference image was created from the pre- and (120 minutes) postcontrast datasets for the slice of interest, by computing the difference in signal intensity between the post-contrast and the pre-contrast image on a pixel basis. From a difference image, ten regions of interest (ROI) of equal size (0.05 cm$^2$) were drawn within areas with the highest $T_1$ relaxation in the tumor parenchyma and contralateral side of the brain for each animal after the VEGFR-2 probe injection at the TR 800 ms. The values obtained from the ROIs in the tumor regions were normalized to the corresponding contralateral sides. The $T_1$ relaxation value of the specified ROIs was computed from all the pixels in the ROI by the following equation [29] (processed by ParaVision 4.0, Bruker): $S \ (TR) = S_0 \ (1 - e^{-TR/T1})$, where TR is the repetition time, $S_0$ is the signal intensity (integer machine units) at $TR \gg T_1$ and $TE = 0$, and $T_1$ is the constant of the longitudinal relaxation time. An overlay of the contrast difference image and $T_1$-weighted image was generated using the 3D Analysis Software for Life Sciences Amira® (Fei, Hillsboro, Oregon).

**Fluorescence staining**

The whole brain of each mouse from each treatment group was removed and fixed in 4% paraformaldehyde solution in phosphate buffered saline, then embedded in Optimal Cutting Temperature (OCT) compound and frozen in liquid nitrogen. Frozen tissue blocks were sectioned at 10 μm, mounted on positive-charged slides, and air-dried for 30 minutes. The tissue samples were permeabilized with 0.01% sapo-
OKN-007 decreases VEGFR-2 in GL261 gliomas

nin in PBS (PBS/SAP) (10 minutes at 20°C) and incubated in a mixture of 3% bovine serum albumin and 5% normal donkey serum (host animal of the conjugated detection antibody) in PBS/SAP (1 hour at 20°C). Saponin was kept in all incubation buffers throughout the staining procedure, to ensure a proper penetration of the antibodies. Saponin exclusively removes cholesterol molecules from lipid-containing structures, leaving stable holes large enough for penetration of antibodies.

Next, the tissue sections were incubated with anti-CD31 antibody (BD Pharmingen Purified Rat anti-mouse CD31, catalog number: 550274) overnight at 4°C. The samples were washed 3 × 10 minutes in PBS/SAP, and incubated for 1 hour at 20°C with combinations of appropriate detection antibodies conjugated with fluorescein isothiocyanate and Cy3 diluted 1:100 in 1% bovine serum albumin in PBS/SAP. After washing as above, the sections were mounted between glass slides and coverslips using Vectashield hardset mounting medium (Vector Laboratories, Burlingame, CA) containing TO-PRO-3 iodine (Molecular Probes, Eugene, OR) as nuclear counterstain.

As negative controls for polyclonal antibody staining, the primary antibodies were replaced with equivalent amounts of mouse nonimmune serum. Specimens were examined by epifluorescence confocal imaging using a Nikon C1 confocal laser-scanning unit equipped with a three-laser launcher (488, 543, and 633 nm emission lines) installed on an Eclipse TE200-U inverted microscope (Nikon, Melville, NY). Images were taken with a ×20 plan achromat objective (NA 0.46). Image collection parameters (neutral density filters, pinhole, and detector gains) were kept constant during image acquisition, to make reliable semiquantitative comparisons between the linear and branched regions of the arteries.

The measurement of fluorescence intensity was done as previously described [30]. In brief, 10 images (12-bit, 4095 gray levels/pixel) were collected for each experimental condition, and the mean fluorescence intensity (MFI) of the whole image or 15 to 20 regions of interest (ROIs) per image was integrated using the EZ-C1 software (Nikon). Image collection parameters (neutral density filters, pinhole, and detector gains) were kept constant during image acquisition to make reliable comparisons between specimens. Unpaired t-test statistical analysis of the data was done using Prism 6.0b (GraphPad Software).

Gold labeling immunoelectron microscopy

Brain tissue was obtained from GL261 glioma bearing mice that were administered either VEGFR-2 probe or a non-specific IgG contrast agent with gadolinium (Gd)-DTPA (diethylene triamine penta acetic acid)-albumin construct. Tissues were removed and cut to size of about 2 cm and 3-5 mm thick then fixed in 4% paraformaldehyde + 0.1% glutaraldehyde in 0.1 M phosphate buffer (PB) for 1 hour at room temperature or at 4°C, washed thoroughly with thoroughly with PB (pH 7.3). Tissue was trimmed further to 1-2 mm pieces. Osmication was omitted for labelling procedure. Dehydration was in an ethanol series (50%, 70%, 80%, at 15 min each). Tissue was placed in 2:1 LR White resin (Electron Microscopy Sciences, PA) to 70% ethanol for 1 hour to avoid tissue shrinkage, and then further infiltrated in 100% LR White for 30 minutes. Tissue was added to gelatin capsules and LR White polymerized at 50°C for 24 hours. Gold-silver sections were placed on nickel grids coated with formvar or colloidal. Sections on grids were protein blocked in Aurion goat gold conjugate (Electron Microscopy Sciences, PA) for 30 minutes, washed in PBS-0.2% bovine serum albumin 3x for 5 minutes at pH 7.4 (Electron Microscopy Sciences, PA). Aurion ultra small gold conjugate-goat anti-Biotin (Electron Microscopy Sciences, PA) in PBS-0.2% BSA-c was applied at 1:100 for overnight at 4°C or for 4 hours at room temperature. Grids were washed in PBS-0.2% BSA-c (pH 7.4), 6x for 5 minutes; in PBS, 3x for 5 minutes; in distilled water 5x for 2 minutes. Silver enhancement was done by placing grids on a droplet of Aurion R-Gent SE-EM enhancement mixture (Electron Microscopy Sciences, PA) for 90 minutes, and then washed in distilled water. Controls involved sections with no secondary antibody (anti-Biotin) and no silver enhancement. Sections were stained with uranyl acetate for 15 minutes followed by lead citrate for 3 minutes. Sections were viewed with a Zeiss T109 electron microscope operated at 80 kV and equipped with a Gatan digital camera and micrograph acquisition software.
OKN-007 decreases VEGFR-2 in GL261 gliomas

San Diego, CA, USA. Tumor volumes and the VEGFR-2 levels were reported as the mean ± standard deviation. Student t-tests (independent-samples, two-tailed t-test) were used to assess the differences between means of untreated and OKN-007 treated GL261 glioma mice. For statistical analysis, Kaplan-Meier survival curves and Log-rank (mantel-cox) test were used to compare the survival times among the untreated and OKN-007 treated groups. All p values < 0.05 were considered statistically significant.

Results

An in vitro assay was conducted to assess if TEC, which express VEGFR-2, can specifically bind the VEGFR-2 probe. First of all it was determined that TEC do express VEGFR-2, as determined by a Western blot (Figure 1B). Next, in vitro MRI assessment of the anti-VEGFR-2 probe in TEC indicated that the VEGFR-2 probe significantly decreased T1 relaxation values in cells with the probe (P < 0.001) compared to cells alone (Figure 1C, 1D).

Once it was established that the VEGFR-2 probe could be used to detect VEGFR-2, in vivo mMRI was used with the VEGFR-2 targeting probe to assess the anti-angiogenic effect of OKN-007 in a mouse GL261 glioma model. Initially we established if there was a therapeutic effect of OKN-007 in GL261 glioma-bearing mice. Starting ten days after GL261 tumor cell

Statistical analysis

Statistical analyses were performed using the GraphPad Prism 6 (GraphPad Prism 6 Software, San Diego, CA, USA). Tumor volumes and the VEGFR-2 levels were reported as the mean ± standard deviation. Student t-tests (independent-samples, two-tailed t-test) were used to assess the differences between means of untreated and OKN-007 treated GL261 glioma mice. For statistical analysis, Kaplan-Meier survival curves and Log-rank (mantel-cox) test were used to compare the survival times among the untreated and OKN-007 treated groups. All p values < 0.05 were considered statistically significant.

Results

An in vitro assay was conducted to assess if TEC, which express VEGFR-2, can specifically bind the VEGFR-2 probe. First of all it was determined that TEC do express VEGFR-2, as determined by a Western blot (Figure 1B). Next, in vitro MRI assessment of the anti-VEGFR-2 probe in TEC indicated that the VEGFR-2 probe significantly decreased T1 relaxation values in cells with the probe (P < 0.001) compared to cells alone (Figure 1C, 1D).

Once it was established that the VEGFR-2 probe could be used to detect VEGFR-2, in vivo mMRI was used with the VEGFR-2 targeting probe to assess the anti-angiogenic effect of OKN-007 in a mouse GL261 glioma model. Initially we established if there was a therapeutic effect of OKN-007 in GL261 glioma-bearing mice. Starting ten days after GL261 tumor cell

Statistical analysis

Statistical analyses were performed using the GraphPad Prism 6 (GraphPad Prism 6 Software, San Diego, CA, USA). Tumor volumes and the VEGFR-2 levels were reported as the mean ± standard deviation. Student t-tests (independent-samples, two-tailed t-test) were used to assess the differences between means of untreated and OKN-007 treated GL261 glioma mice. For statistical analysis, Kaplan-Meier survival curves and Log-rank (mantel-cox) test were used to compare the survival times among the untreated and OKN-007 treated groups. All p values < 0.05 were considered statistically significant.

Results

An in vitro assay was conducted to assess if TEC, which express VEGFR-2, can specifically bind the VEGFR-2 probe. First of all it was determined that TEC do express VEGFR-2, as determined by a Western blot (Figure 1B). Next, in vitro MRI assessment of the anti-VEGFR-2 probe in TEC indicated that the VEGFR-2 probe significantly decreased T1 relaxation values in cells with the probe (P < 0.001) compared to cells alone (Figure 1C, 1D).

Once it was established that the VEGFR-2 probe could be used to detect VEGFR-2, in vivo mMRI was used with the VEGFR-2 targeting probe to assess the anti-angiogenic effect of OKN-007 in a mouse GL261 glioma model. Initially we established if there was a therapeutic effect of OKN-007 in GL261 glioma-bearing mice. Starting ten days after GL261 tumor cell

Statistical analysis

Statistical analyses were performed using the GraphPad Prism 6 (GraphPad Prism 6 Software, San Diego, CA, USA). Tumor volumes and the VEGFR-2 levels were reported as the mean ± standard deviation. Student t-tests (independent-samples, two-tailed t-test) were used to assess the differences between means of untreated and OKN-007 treated GL261 glioma mice. For statistical analysis, Kaplan-Meier survival curves and Log-rank (mantel-cox) test were used to compare the survival times among the untreated and OKN-007 treated groups. All p values < 0.05 were considered statistically significant.

Results

An in vitro assay was conducted to assess if TEC, which express VEGFR-2, can specifically bind the VEGFR-2 probe. First of all it was determined that TEC do express VEGFR-2, as determined by a Western blot (Figure 1B). Next, in vitro MRI assessment of the anti-VEGFR-2 probe in TEC indicated that the VEGFR-2 probe significantly decreased T1 relaxation values in cells with the probe (P < 0.001) compared to cells alone (Figure 1C, 1D).

Once it was established that the VEGFR-2 probe could be used to detect VEGFR-2, in vivo mMRI was used with the VEGFR-2 targeting probe to assess the anti-angiogenic effect of OKN-007 in a mouse GL261 glioma model. Initially we established if there was a therapeutic effect of OKN-007 in GL261 glioma-bearing mice. Starting ten days after GL261 tumor cell

Statistical analysis

Statistical analyses were performed using the GraphPad Prism 6 (GraphPad Prism 6 Software, San Diego, CA, USA). Tumor volumes and the VEGFR-2 levels were reported as the mean ± standard deviation. Student t-tests (independent-samples, two-tailed t-test) were used to assess the differences between means of untreated and OKN-007 treated GL261 glioma mice. For statistical analysis, Kaplan-Meier survival curves and Log-rank (mantel-cox) test were used to compare the survival times among the untreated and OKN-007 treated groups. All p values < 0.05 were considered statistically significant.

Results

An in vitro assay was conducted to assess if TEC, which express VEGFR-2, can specifically bind the VEGFR-2 probe. First of all it was determined that TEC do express VEGFR-2, as determined by a Western blot (Figure 1B). Next, in vitro MRI assessment of the anti-VEGFR-2 probe in TEC indicated that the VEGFR-2 probe significantly decreased T1 relaxation values in cells with the probe (P < 0.001) compared to cells alone (Figure 1C, 1D).

Once it was established that the VEGFR-2 probe could be used to detect VEGFR-2, in vivo mMRI was used with the VEGFR-2 targeting probe to assess the anti-angiogenic effect of OKN-007 in a mouse GL261 glioma model. Initially we established if there was a therapeutic effect of OKN-007 in GL261 glioma-bearing mice. Starting ten days after GL261 tumor cell

Statistical analysis

Statistical analyses were performed using the GraphPad Prism 6 (GraphPad Prism 6 Software, San Diego, CA, USA). Tumor volumes and the VEGFR-2 levels were reported as the mean ± standard deviation. Student t-tests (independent-samples, two-tailed t-test) were used to assess the differences between means of untreated and OKN-007 treated GL261 glioma mice. For statistical analysis, Kaplan-Meier survival curves and Log-rank (mantel-cox) test were used to compare the survival times among the untreated and OKN-007 treated groups. All p values < 0.05 were considered statistically significant.

Results

An in vitro assay was conducted to assess if TEC, which express VEGFR-2, can specifically bind the VEGFR-2 probe. First of all it was determined that TEC do express VEGFR-2, as determined by a Western blot (Figure 1B). Next, in vitro MRI assessment of the anti-VEGFR-2 probe in TEC indicated that the VEGFR-2 probe significantly decreased T1 relaxation values in cells with the probe (P < 0.001) compared to cells alone (Figure 1C, 1D).

Once it was established that the VEGFR-2 probe could be used to detect VEGFR-2, in vivo mMRI was used with the VEGFR-2 targeting probe to assess the anti-angiogenic effect of OKN-007 in a mouse GL261 glioma model. Initially we established if there was a therapeutic effect of OKN-007 in GL261 glioma-bearing mice. Starting ten days after GL261 tumor cell
OKN-007 decreases VEGFR-2 in GL261 gliomas

implantation, all animals from each group (untreated vs. OKN-007 treated) were imaged every 2-3 days until the end of the study. The tumor volumes of the OKN-007-treated animals were significantly smaller ($p = 0.0054$; $32.95 \pm 7.130$, $n = 4$) than the untreated group ($103.0 \pm 16.96$, $n = 7$) at 21 days following cell injection (Figure 2A-C). Additionally, OKN-007-treated mice were found to survive significantly longer ($P = 0.0005$; median survival = 30 days; $n = 7$), compared with untreated animals (median survival = 22 days; $n = 10$) (Figure 2D). These results indicate that OKN-007 had an anti-cancer effect in GL261 tumors.

In vivo VEGFR-2 targeted mMR images were then obtained to assess the effect of OKN-007 on VEGFR-2 levels. Representative examples of images obtained in GL261 glioma-bearing mice are shown in Figure 3. $T_2$-weighted and $T_1$-weighted morphological images obtained before injection of the VEGFR-2 probe are shown in Figure 3A and 3B respectively. A difference image, which is the subtraction between the $T_1$ images of pre- and 2 hrs after injection of the contrast agent, shows increased anti-VEGFR-2 probe accumulation in Figure 3C, compared to the untreated animals ($10.12 \pm 2.25$) (Figure 4C). These results showed that OKN-007 was able to decrease the detection of the VEGFR-2 probe in vivo in GL261 tumors.

It was then important to verify in ex vivo GL261 tumor samples that the VEGFR-2 probe was present using fluorescence imaging. The fluorescence agent Cy3 (red) attached to streptavidin was used to target the biotin moiety of the anti-VEGFR-2 probe in ex vivo excised tumor tissue from a GL261 glioma-bearing mice administered the anti-VEGFR-2 probe in vivo. It was also necessary to demonstrate that the VEGFR-2 probe is associated with endothelial cells. The fluorescence images detecting the VEGFR-2 probe were superimposed with immunofluorescence images for CD-31 (endothelial marker). Co-localization (represented as yellow-orange regions) of the anti-VEGFR-2 probe (red) with CD31 (green) in some areas of glioma tissue from GL261 glioma-bearing mice administered the anti-VEGFR-2 probe was observed (Figure 5), and found to be less for OKN-007-treated samples (Figure 5B) compared to untreated tumors (Figure 5A). Ex vivo VEGFR-2 levels in untreated and OKN-007-treated GL261 tumor samples were also assessed to verify that OKN-007 actually did affect VEGFR-2 directly. Immunofluorescence imaging for VEGFR-2 indicated that the untreated GL261 gliomas had high levels of VEGFR-2, which were significantly higher than GL261 gliomas treated with OKN-007 ($P < 0.0001$) (Figure 5C-E). These results verified that: (1) the VEGFR-2
OKN-007 decreases VEGFR-2 in GL261 gliomas

Electron microscopy was used to detect the possible subcellular location of the VEGFR-2 probe in GL261 glioma cells. Ex vivo immunoelectron microscopy of GL261 mouse glioma tissues after post-embedding incubation with gold-labeled anti-biotin antibody was performed to determine the sub-cellular localization of the anti-VEGFR-2 probe (Figure 6). The gold-labeled anti-biotin was found to be distributed within the endothelial cell and glioma cell plasma membrane/cytoplasm, as well as the nuclei of the GL261 tumor cells in mice administered the anti-VEGFR-2 in vivo (Figure 6A). A control was found to have post-embedding immunogold labeling only in the endothelial cell nucleus (Figure 6B). These results indicate that the VEGFR-2 probe is internalized into the cell membrane/cytoplasm.

Discussion

Angiogenesis, the formation of new blood vessels, is a critical step during tumorigenesis and represents a pathological hallmark of cancer [1], and plays a crucial role in glioma development and growth [31]. Briefly, the start of the angiogenesis process requires some angiogenic exogenous stimulus, such as hypoxia, current metabolic requirements, and tumour growth. More than 25 different growth factors and cytokines related to the induction of angiogenesis have been reported [32]. The production of these proangiogenic factors is a result of genetic alterations or is induced by hypoxia. Intratumour hypoxia occurs at the time when there is an probe was higher in untreated GL261 gliomas compared to those treated with OKN-007, (2) the VEGFR-2 probe colocalized with endothelial cells, and (3) VEGFR-2 levels in OKN-007-treated tumors were lower than those detected in untreated tumors.

Figure 4. mMRI detection of VEGFR-2 in a GL261 mouse glioma model. Representative MR images from: (A) Untreated and (B) OKN-007 treated GL261 mouse gliomas. (Aii, Bii) T1-weighted images overlayed with a difference T1-weighted image (red), which was the subtraction between the 2 hr post-contrast and the pre-contrast images after injection of either the anti-VEGFR-2 probe contrast agent. Overlays of the contrast difference images and the T1-weighted images were generated using the 3D Analysis Software for Life Sciences Amira®. (C) OKN-007 was found to significantly decrease (P = 0.4137) (2.54 ± 2.02) the levels of VEGFR-2 in the treated group compared to the untreated animals (10.12 ± 2.25). Values are represented as means ± SD. Asterisks indicate statistically significant differences (*P < 0.05). UT: untreated animals.
OKN-007 decreases VEGFR-2 in GL261 gliomas

Figure 5. Ex vivo detection of the anti-VEGFR-2 probe, colocalization with endothelial cell marker CD31, and detection of VEGFR-2 in untreated and OKN-007-treated GL261 mouse gliomas. Fluorescence images of the anti-VEGFR-2 probe (red) and the endothelial marker CD31 (green) in untreated (Ai-Aiii: 3 separate samples in the mid-tumor regions) and OKN-007-treated (Bi-Biii: 3 separate samples in the mid-tumor regions) GL261 tumor-bearing animals. Note colocalization (yellow-orange) of the anti-VEGFR-2 probe with some endothelial cells in the untreated group. Representative VEGFR-2 immunofluorescence images in untreated (C) and OKN-007-treated (D) GL261 tumor-bear-
OKN-007 decreases VEGFR-2 in GL261 gliomas

Imbalance between oxygen supply and demand, due to irregular and chaotic blood flow [33]. Furthermore, when a solid tumor, such as a brain tumor, grows larger than a critical size (1-2 mm in diameter), it must recruit new blood vessels to supply the required oxygen and nutrition levels necessary for its survival and proliferation [16, 34]. This process comprises the formation of new blood vessels from preexisting ones, and is a crucial step in the progression of cancer from a small and localized neoplasm to a highly aggressive tumor [16]. Additional details about the angiogenesis process in gliomas can be found in a number of outstanding review articles [16, 31, 35-37]. Gliomas are highly vascularized tumors [37]. Furthermore, the formation of abnormal tumor vasculature and glioma cell invasion along white matter tracts are proposed to be the major causes of the therapeutic resistance of these tumors [16]. Thus, inhibition of angiogenesis has emerged as a promising strategy for the treatment of brain cancers.

In this study, we demonstrated through conventional and molecular MRI that OKN-007 inhibited the tumor levels of VEGFR-2, tumor growth, and increased survival time in treated GL261 mouse gliomas when compared to the untreated group. Our group has already established that OKN-007 is an effective anti-cancer agent in rodent pre-clinical models for adult GBMs, including F98 [18], U87 [18], and C6 [17, 19] glioma models. Furthermore, in an orthotopic rat F98 glioma model and a human U87 xenograft model in athymic rats, OKN-007 significantly increased the survival of treated versus untreated rats [19]. Similarly to our previous studies, OKN-007 was also able to decrease the tumor volumes in the GL261 mouse glioma model. In the present study, the tumor volumes of OKN-007-treated GL261 gliomas were significantly smaller ($P = 0.0054$) when compared to the untreated group (Figure 2C). MRI is an ideal method for accurately measuring tumor volumes, as it was previously demonstrated that there is a very high correlation when measuring pre-clinical intracranial tumor volumes with MRI compared to histology [38].

The survival time of GL261-tumor bearing mice was also significantly increased after OKN-007
OKN-007 decreases VEGFR-2 in GL261 gliomas

therapy, as we also have previously reported in other adult glioma models [18, 19]. The OKN-007-treated group (n = 7) had a median survival of 30 days, which was significantly longer (P = 0.0005) than the median survival of 22 days for the untreated group (n = 10) (Figure 2D). OKN-007 was able to increase the survival time by 8 days in the treated group compared to the untreated animals. Considering that one mouse month is equivalent to ~2.6 human years, based on a 4-year maximum life span [39], OKN-007 could potentially increase survival time in adult GBM patients by ~7-8 months.

There are a number of excellent recent reviews on the use of molecular MRI applications to assess angiogenesis [40-42]. Recently, our group reported the in vivo characterization and distribution of VEGFR-2 in C6 [22, 27] and RG2 [22] glioma models using molecular magnetic resonance imaging (mMRI) and an anti-VEGFR-2 gadolinium-based contrast agent. MRI signal enhancement of the VEGFR-2 probe in the RG2 tumor interior was more than twice that for the C6 tumor interior, suggesting more VEGFR-2 from microvessels inside the RG2 tumor [22]. However, compared to C6 gliomas, RG2 gliomas have a relatively more homogeneous pattern for VEGFR-2 levels in the tumor overall [22]. In the C6 glioma, distinctively higher levels of VEGFR-2 were found in peri-tumor and perinecrotic regions compared to tumor interior regions [27]. Other groups that have used mMRI probes for VEGFR-2 have used aptamer-modified magnetic nanoprobes [43], and a low molecular weight peptoid-(Gd)8-dendron [44]. There are several other MRI methods that can be used to assess vascular alterations, such as perfusion imaging or microvascular MRI [40, 45], and hyperpolarized 13C spectroscopy [46], however these methods do not directly target VEGFR-2.

It was also previously demonstrated that OKN-007 is an effective anti-angiogenic compound in vivo, by directly decreasing microvessel density (CD-31) and HIF-1α levels in both F98 and U87 glioma models [18], and ex vivo by directly decreasing the levels of VEGFR-2 in C6 rat gliomas, as measured by Western blots [19]. In the present study we confirmed using the mMRI technique, that OKN-007 was also able to decrease the levels of VEGFR-2 in a preclinical GL261 mouse glioma model (Figure 4).

Specifically, we combined mMRI with a Gd-DTPA-albumin-based contrast agent for signal detection with the specificity of an antibody for VEGFR-2 (anti-VEGFR2 probe), to detect in vivo VEGFR-2 levels in untreated and OKN-007 treated adult GL261 mouse gliomas. To calculate the Ti relaxation ratio, we normalized the Ti relaxation values from the tumor area to the contralateral side of the brain for each of the animal datasets. We used the contralateral side of the brain to normalize the Ti relaxation values of the tumor region as we detected no significant difference (P = 0.1824) in the Ti relaxation values in the two treatment groups within the contralateral brain tissue, 120 minutes after the administration of the anti-VEGFR-2 probe.

We have already demonstrated with immunofluorescence that the anti-VEGFR-2 probe binds to the VEGFR-2 present on both endothelial and tumor cells, via the co-localization of the anti-VEGFR-2 probe with VEGFR-2 expression in a rat C6 glioma model [27]. Co-localization of the anti-VEGFR-2 probe with the endothelial marker, laminin, was also very high in C6 gliomas [27]. In the present study for the mouse GL261 glioma model, fluorescence staining for the biotin moiety of the VEGFR-2 probe at 2 hours post administration of the probe confirmed the distribution of the VEGFR-2 probe similarly to our previous study [27]. Furthermore, the probe also showed co-localization with the endothelial marker CD31 (Figure 5A) indicating that it does target endothelial cells. In addition, in vitro assessment of the binding specificity of the VEGFR-2 probe was shown in TEC (Figure 1C, 1D), with a significant increase in binding to TEC with the probe (P < 0.0001) compared to cells alone. Immunofluorescence imaging for VEGFR-2 levels indicated that OKN-007 was able to specifically decrease the levels of VEGFR-2 (Figure 5C, 5E).

Here we also used immunoelectron microscopy to demonstrate the presence of the anti-VEGFR-2 probe in the cell membrane/cytoplasm and cell nuclei of the endothelial cells and GL261 glioma tumor cells (Figure 6A). During this technique, anti-biotin-gold was used to bind specifically to the biotin group of the anti-VEGFR-2 probe. The IgG contrast agent, which also has a biotin moiety, was not detected at very high levels with the anti-biotin-gold in the...
OKN-007 decreases VEGFR-2 in GL261 gliomas

Figure 7. Schematic representation of possible anti-angiogenic mechanisms of action for the anti-cancer agent OKN-007. (1) VEGF is ubiquitously expressed in almost all tumors. Glioma cells have been demonstrated to secrete VEGF, which leads to increased angiogenesis [52]. (2) VEGF exerts its action through binding to VEGFR-2, leading to autophosphorylation of tyrosine residues in the cytoplasmic domain of VEGFR-2 and driving downstream pathway such as PI3K/AKT and MAPK to promote endothelial cells proliferation and migration (3). VEGF also stimulates reactive oxygen and nitrogen species (RONS) production via Rac1-mediated NADPH oxidase activation (4) [53] and also increases mitochondria-derived hydrogen peroxide (H₂O₂) (5) [54]. (6) RONS, in turn, potentiate VEGFR phosphorylation [52]. RONS can also upregulate VEGF secretion and VEGFR expression through induction of transcription factors HIF-1α (hypoxia-inducible factor-1α) (7) [54-56]. (8) Under hypoxic conditions, the active HIF-1α recognizes and binds to cis elements in the promoter of genes that mediate angiogenesis, such as the gene encoding VEGF [57]. (9) NF-κB (nuclear factor of kappa light polypeptide gene enhancer in B-cell) is normally under strict regulation...
OKN-007 decreases VEGFR-2 in GL261 gliomas

by its sequestration in the cytoplasm in association with a heterotrimeric complex of NF-κB with the inhibitor of κB (IkB) [57]. Generation of reactive oxygen and nitrogen species (RONS) and activation of PI3 kinase-AKT pathway can converge and activate NF-κB. NF-κB also activates several genes that mediate angiogenesis, such ELR+ CXC chemokines [57]. OKN-007 decreases levels of VEGFR-2 [19] and HIF-1α [18] in rat gliomas, and nitrones such as PBN (α-phenyl-tert-butyl nitrone) decrease NF-κB [13] in a mouse endotoxemia model [50]. There is also evidence that OKN-007 effectively traps free radicals [14] in vitro [44]. Consequently, based on the possible mechanisms-of-action and our data reported in this study, OKN-007 is an effective agent that can inhibit tumor angiogenesis [15]. The schematic is a modification of combined schemes obtained from Oxidative Medicine and Cellular Longevity, Article ID 374963, Zhou Y et al. (2013) [Copyright (2013) with permission from Hindawi Publishing Corporation] [58], and from Nature Medicine, 11 (9): 925-7, Strieter RM. (2005) [Copyright (2005) with permission from Nature Publishing Group] [57].

Conclusion

This is the first attempt at detecting in vivo levels of VEGFR-2 in a mouse GL261 glioma model and assessing the anti-angiogenic capability of an anticancer nitrone. Our study confirmed that in vivo VEGFR-2 levels can be monitored and can be used to assess the efficacy of an anti-angiogenic therapy.

This method can potentially be applied towards other types of cancers for the in vivo assessment of VEGFR-2 levels, and assess various anti-angiogenic agents. The results indicate that OKN-007 treatment substantially decreased VEGFR-2 levels in a GL261 glioma model, and can be considered as an anti-angiogenic therapy in human gliomas.

Acknowledgements

We gratefully acknowledge the Peggy and Charles Stephenson Cancer Center at the University of Oklahoma, Oklahoma City, OK, for use of their Histology and Immunohistochemistry Core for cryosectioning, and funding by an Institutional Development Award (IDeA) from the National Institute of General Medical Sciences of the National Institutes of Health under grant number P20 GM103639 (R.A.T.). This work was also supported by the Oklahoma Medical Research Foundation (R.A.T.).

Disclosure of conflict of interest

None.

Abbreviations

BSA, bovine serum albumin; DTPA, diethylene triamine penta acetic acid; FOV, field of view; GBM, glioblastoma; Gd, gadolinium; HIF-1α, hypoxia-inducible factor-1α; IkB, inhibitor of κB; mMRI, molecular magnetic resonance imaging; MSME, Multiple-slice multiple echo imaging; NF-κB, nuclear factor of kappa light polypeptide...
OKN-007 decreases VEGFR-2 in GL261 gliomas

gene enhancer in B-cells; PBN, α-phenyl-tert-butylnitron; OCT, optimal cutting temperature; OKN-007, (2, 4-disulfophenyl-PBN; or disodium 4-[(tert-butylimino)methyl]benzene-1,3-disulfonate N-oxide or disufenton; also previously known as NXY-059); RARE, rapid acquisition with refocused echoes; ROI, region of interest; RONS, reactive oxygen and nitrogen species; Sap, saponin; TE, echo time; TEC, tumor endothelial cells; TR, relaxation time; UT, untreated; VEGF, vascular endothelial growth factor; VEGFR, vascular endothelial growth factor receptor.

Address correspondence to: Rheal A Towner, Advanced Magnetic Resonance Center, Oklahoma Medical Research Foundation, 825 NE 13th St, Oklahoma, OK 73104 USA. Tel: +1-405-271-7383; E-mail: Rheal-Towner@omrf.org

References

OKN-007 decreases VEGFR-2 in GL261 gliomas


OKN-007 decreases VEGFR-2 in GL261 gliomas


