The proliferation of malignant melanoma cells could be inhibited by ranibizumab via antagonizing VEGF through VEGFR1

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Purpose: Angiogenesis is an important mediator in tumor progression. Vascular endothelial growth factor (VEGF) is one of the major cytokines that can influence angiogenesis. However, the potential mechanism of tumor growth inhibition through anti-VEGF agents is still unclear. This study was performed to examine whether ranibizumab could inhibit malignant melanoma growth in vitro and to determine the safety of ranibizumab on human adult retinal pigment epithelium cell line (ARPE-19 cells).

Methods: Malignant melanoma cells obtained from a clinic were cultured in vitro. VEGF concentrations secreted by malignant melanoma cells and the ARPE-19 cells were examined by enzyme-linked immunosorbent assay (ELISA). The two kinds of cells were both treated with VEGF and its antagonist, ranibizumab. The dynamic changes of the two types of cells were monitored by real-time cell electronic sensing (RT-CES) assay. The effect of ranibizumab on both types of cells was verified by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl (MTT) assay. The expression of VEGF receptor 1 (VEGFR1) RNA in uveal melanoma was further investigated through the PCR technique.

Results: The levels of VEGF secreted by malignant melanoma cells were much higher than those of ARPE-19 cells, and were markedly decreased in the action of 0.1 mg/ml ranibizumab. However, there was no obvious reduction of VEGF in the presence of ranibizumab for ARPE-19 (p>0.05). Meanwhile, RT-CES showed that the viability of malignant melanoma cells increased greatly in the presence of VEGF. When VEGF was 20 ng/ml, viability of the malignant melanoma cells increased by 40% compared with the negative control. There was no evident effect on proliferation of ARPE-19 (p>0.05). Furthermore, the growth of malignant melanoma cells was obviously inhibited after ranibizumab intervention. When ranibizumab was administered at 0.25 mg/ml, the survival rate of the malignant melanoma cells decreased to 57.5%. Nevertheless, low-dose exposure to ranibizumab had only a slight effect on the growth of ARPE-19, and PCR result demonstrated that VEGFR1 plays a role in this tumor tissue rather than VEGFR2.

Conclusions: Ranibizumab can selectively inhibit malignant melanoma cell proliferation by decreasing the expression of VEGF; the possible mechanism of the inhibitory effect may involve VEGFR1 antagonism.

Vascular endothelial growth factor (VEGF) was first described as a molecule that could increase the permeability of blood vessels. Additionally, VEGF promotes the proliferation of new blood vessels, and is essential for normal embryonic development and wound healing. There is an obvious correlation between intensity of VEGF and tumor prognosis [1]. VEGF encompasses a family of proteins that include placenta growth factor (PIGF), VEGF-A, VEGF-B, VEGF-C, VEGF-D, and VEGF-E. The VEGF receptor (VEGFR) family in mammals contains three members, namely VEGFR1, VEGFR2, and VEGFR3. These factors directly participate in the genesis of blood capillaries and lymphatic vessels [2-7].

Three anti-VEGF agents—pegaptanib, bevacizumab, and ranibizumab [8]—have been used for the treatment of patients with neovascularization pathology. Ranibizumab (Lucentis®, Genentech, Inc., South San Francisco, CA) is a recombinant humanized immunoglobulin designed for intraocular use which can bind to and inhibits the biological activity of human VEGF-A [9]. It has been shown to be safe and effective when given intravitreally to patients with neovascular wet age-related macular degeneration (AMD). In addition, ranibizumab has recently been approved for diabetic macular edema (DME) therapy [10]. Bevacizumab (Avastin®, Genentech, Inc.) a full-length, humanized, monoclonal antibody against all types of VEGF, is the most commonly used drug in the United States for the treatment of neovascular
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Solution. Paraffin sections from the tissue were examined. A part of the tumor tissue was fixed by 10% formaldehyde for 15 months previously; it was confirmed that intravitreal bevacizumab could ameliorate the decline in visual acuity caused by radiation maculopathy [13].

Tumor growth is angiogenesis dependent, and therapy targeting tumor vasculature is an attractive alternative or adjunct to conventional therapy. VEGF is important in several malignant and nonmalignant pathologies. Previously, it was shown that selective inhibition of VEGF binding to VEGFR2 with a fully human monoclonal antibody (r84) is sufficient for effective control of tumor growth in a preclinical model of breast cancer [14]. One report demonstrated the effectiveness of anti-VEGF therapy as a modulator of immune cell infiltration, as well as intratumoral and serum cytokine levels, in multiple preclinical models of breast cancer [15]. In metastatic colorectal cancer, an objective response rate of 3.3% was observed among chemotherapy-pretreated patients receiving monotherapy with bevacizumab [16]. Trials of bevacizumab with chemotherapy as the first-line treatment for metastatic non-small-cell lung cancer have yielded the results of improving patient outcomes [17,18].

We were interested in exploring whether ranibizumab, the anti-VEGF agent, would result in novel efficacy against ocular tumor. In addition, we intended to evaluate the safety of ranibizumab. In the present study, the effects of VEGF on malignant melanoma cells of the ciliary body and on the human adult RPE cell line (ARPE-19) were investigated to determine whether the growth of two kinds of cells is VEGF dependent, and the inhibitory mechanism of ranibizumab on the growth of tumor cells and ocular safety was also explored.

METHODS

The study was performed in the Affiliated Eye Hospital of Shandong University of Traditional Chinese Medicine. All measurements adhered to the tenets of the Helsinki agreement.

Primary culture of ocular malignant melanoma and culture of the ARPE-19 cell line: Malignant melanoma of the ciliary body was derived from a female inpatient in our hospital. A part of the tumor tissue was fixed by 10% formaldehyde solution. Paraffin sections from the tissue were examined with hematoxylin and eosin (H&E) staining. Immunohistochemical staining of HMB-45, S-100 protein, and Melan-A (Golden Bridge Biotechnology Company Ltd., Peking, China) were performed for identification of melanoma. The other part of the tumor tissue was cultured in RPMI 1640 medium (HyClone, Tianjin, China) with 10% fetal bovine serum (FBS; HyClone) after digestion with 0.05% trypsin. The following experiments were carried out with three to six cell passages.

The human RPE cell line (ARPE-19) was obtained from the American Type Cell Culture (ATCC, Manassas, VA); it matched for the ATCC human cell line CRL-2302. The ARPE-19 cell was verified by the ATCC Cell Line Authentication Service (Promega, Madison, WI) using short tandem repeat analysis and an amelogenin gender-determining locus, as shown in Table 1. The cells were used between passages 3 and 6. Cells were maintained in the same medium as melanoma, that is, RPMI 1640 with 10% FBS. All cells were cultured at 37 °C in a 5% CO₂ incubator with a humidified environment.

Malignant melanoma cells of the ciliary body were cultured primarily in the laboratory and cells between the third and the sixth generation were used in the experiments. When they had reached 80–90% confluence, cells were digested with 0.05% trypsin. After washing twice with PBS, the cell pellet was smeared onto three pieces of slides for immunocytochemistry staining with HMB-45, S-100 protein, and Melan-A, respectively.

VEGF secretion detected by ELISA: To determine the VEGF level secreted by malignant melanoma cells and ARPE-19, 200 µl supernatant was collected from the media after 24 h culture in the absence and presence of 0.1, 0.25, 0.5, and 1 mg/ml of ranibizumab. The concentration was measured using a VEGF ELISA kit (R&D, Minneapolis, MN) following the suggested protocol. The optical densities were determined within 30 min and recorded with a microplate reader (BioTek, EXL800, Winooski, VT) at 450 nm.

Real-time cell electronic sensing assay: First, $1 \times 10^{4}$ malignant melanoma cells or ARPE-19 were seeded into each well of a 16-well plate containing 100 µl of culture medium and incubated for 20 h in a 5% CO₂ incubator. After removing the medium, the VEGF solution (Pepro Tech Inc., Suzhou, China) was diluted with fresh RPMI 1640 medium and then added into each well with final concentrations of VEGF 0, 1, 5, 10, and 20 ng/ml. Each concentration was set in triplicate.

Second, the medium was removed after 20 h of incubation for the two kinds of cells. The final concentrations of ranibizumab in malignant melanoma cells and ARPE-19 were
0, 0.1, 0.25, 0.5, and 1 mg/ml, respectively. Each concentration was set in triplicate.

Analyses for both experiments were conducted using an RT-CES analyzer (ACEA Biosciences Inc., Hangzhou, China) for at least 80 h and the cell culture medium volume used for detection was 200 µl. The electrical impedance, displayed as a normalized cell index (NCI), was monitored and the dynamic changes induced by the interaction between cells and VEGF or ranibizumab was recorded.

RT-CES system monitors cellular events in real time by measuring the electrical impedance between microelectrodes integrated into the bottom of custom-made tissue culture plates (E-Plates). Since cells have a very high electrical resistance, the more cells are attached to the bottom of the E-Plates, the higher the electrical impedance will be. Thus, the electrical impedance, displayed as NCI, can be used to monitor cell viability, number, and adhesion in a large number of cell culture wells simultaneously, at any given frequency, and over any desired period of time without taking the plates out of the incubator. The RT-CES assay has been proven to be a valuable and reliable approach to real-time monitoring of dynamic changes induced by cell-chemical interaction [19-22].

**MTT assay:** Based on the observations above, the potential effect of ranibizumab on melanoma cells and ARPE-19 cells was further investigated via MTT assay. First, 1×10⁴ cells were seeded into each well containing 200 µl cell culture medium in 96-well plates and incubated for 24 h in a 95% air and 5% CO₂ incubator. The ranibizumab solution was diluted with fresh RPMI 1640 medium and then added to the cultivated wells, where the final concentrations of ranibizumab in both cells’ wells (in triplicate) were 0, 0.1, 0.25, 0.5, and 1 mg/ml, respectively. Wells without ranibizumab were used as the control experiments. After incubation for 24 h, 20 µl of 5 mg/ml of MTT was added to each well, followed by further incubation for 4 h. The survival rate (%) was expressed as follows: survival rate (%) = ([A] ranibizumab/[A]control) × 100, where A is the absorbance value of the relative sample at 490 nm. Every experiment was repeated at least three times.

### Table 1. STR genotype and amelogenin gender-determining locus of ARPE-19 cell used.

| Loci          | Query profile: ARPE-19(UCLA-19) | ATCC reference database profile |
|---------------|---------------------------------|----------------------------------|
| D3S1358       | 14                              | 15                               |
| THO1          | 6                               | 9.3                              |
| D21S11        | 28                              | 29                               |
| D18S51        | 12                              | 16                               |
| Penta_E       | 7                               | 11                               |
| D5S818        | 13                              | 13                               |
| D13S317       | 11                              | 12                               |
| D7S820        | 9                               | 11                               |
| D16S539       | 9                               | 11                               |
| CSF1PO        | 11                              | 11                               |
| Penta_D       | 11                              | 13                               |
| Amelogenin    | X                               | Y                                |
| vWA           | 16                              | 19                               |
| D8S1179       | 13                              | 19                               |
| TPOX          | 9                               | 11                               |
| FGA           | 23                              | 11                               |
| D19S433       | 12                              | 13                               |
| D2S1338       | 19                              |                                  |

Number of shared alleles between query sample and database profile: 16
Total number of alleles in the database profile: 16
Percent match between the submitted sample and the database profile: 100
Extraction of total RNA from formalin-fixed, paraffin-embedded tissue: Total RNA from the tumor was extracted from formalin-fixed, paraffin-embedded tissue using a modification of the method described by Rupp and Locker [23]. Briefly, RNA was extracted from ten 8 μm sections, and further paraffin was removed by extracting twice with 1.5 ml of xylene for 10 min followed by rehydration through subsequent washes with 100, 90, and 70% ethanol diluted in RNase-free water; then the tissue was collected by centrifugation at 16,000 × g for 5 min. After the final step with 70% ethanol rinse, the pellet was dried, resuspended in 200 ml of RNA lysis buffer containing 10 mmol/l Tris (pH 8.0), 0.1 mmol/l EDTA (pH 8.0), 2% sodium dodecyl sulfate (pH 7.3), and 500 mg/ml proteinase K (Sigma, Deisenhafen, Germany), and incubated at 60 °C for 16 h until the tissue was completely solubilized. The precipitate was removed and resuspended in 1 ml Trizol reagent (Aidlab Biotechnologies Co., Ltd., Peking, China) and incubated at 4 °C for 30 min. RNA was purified by phenol and chloroform extractions followed by precipitation with an equal volume of isopropanol for 30 min. The RNA pellet was washed once in 75% ethanol, dried, and resuspended in 20 μl of RNase-free water; it was then quantified by a micro-spectrophotometer (K5600, Beijing Kaiao Technology Development Co., Ltd., Peking, China).

PCR determination: The total RNA was reverse transcribed using Moloney murine leukemia virus (M-MLV) reverse transcriptase (Aidlab Biotechnologies Co., Ltd.). PCR was performed on a TP600 reverse-transcription PCR (RT–PCR) system (Takara, Japan), according to the manufacturer’s instructions. The primer sequences for human VEGF receptor 1 (VEGFR1) were 5′-TTT AAA AGG CAC CCA GCA CAT-3′ (forward) and 5′-CTT ACC ATT TCA GGC AAA GAC-3′ (reverse); primer sequences for human VEGFR2 were 5′-GGC CCA ATA ATC AGA GTG GCA-3′ (forward) and 5′-TGT CAT TTC CGA TCA CTT TTG GA-3′ (reverse). The amplification conditions were as follows: 95 °C for 30 s and 40 cycles of 95 °C for 5 s and 57 °C for 34 s. PCR products were analyzed using agarose gel electrophoresis.

Statistical analysis: Data were expressed as the mean ± SD (standard deviation) from at least three independent experiments. One-way analysis of variance (ANOVA) was used for significance testing, and p<0.05 was considered statistically significant.

RESULTS

Histopathological observation of malignant melanoma of the ciliary body: The tumor was composed of fusiform and epithelioid malignant melanoma cells. The cytoplasm was abundant with melanin granules. The nuclei were big with prominent nucleoli (Figure 1A). Additionally, paraffin sections were depigmented by KMnO4 (potassium permanganate) before H&E staining (Figure 1B), and the tumor showed histopathological features of infiltrative growth. Immunohistochemical staining indicated that the tumor tissue expressed HMB-45 and S-100 positively (Figure 1C,E). Melan-A was weakly positive (Figure 1D). The positive expression was displayed as brown yellow in cytoplasm. Melanin granules appeared dark green after cytoplasm was counterstained with Giemsa [24].

Malignant melanoma cell culture and immunocytochemical staining: The malignant melanoma cells were cultured at 37 °C in a 5% CO2 humidified incubator. The cells appeared to have attached characteristics with a spindle shape, and were rich in melanin granules. During the course of culturing, the cells depigmented gradually (Figure 2A).Passaging to the fifth generation, the melanin granules disappeared completely (Figure 2B). Immunocytochemical staining indicated positive expression of S-100 (Figure 2E) and weak positive expression of HMB-45 and Melan-A (Figure 2C, D). The positive expression is located in the cytoplasm with a brown-yellow color. PBS solution was used as the primary antibody in the negative control for immunohistochemical staining and immunocytochemical staining.

Malignant melanoma cells secrete more VEGF than ARPE-19 cells: We determined the levels of VEGF of the supernatant from the two kinds of cells with or without ranibizumab by ELISA. Untreated malignant melanoma cells expressed 1533.4±7 pg/ml VEGF (Figure 3); the mean levels of VEGF of malignant melanoma cells after 24 h exposure to 0.1, 0.25, 0.50, and 1 mg/ml of ranibizumab were 822.55±9.3 pg/ml, 875.02±7.8 pg/ml, 836.13±6.3 pg/ml, and 844.8±6 pg/ml, respectively. ARPE-19 without ranibizumab expressed 908.75±9.1 pg/ml VEGF, and the mean levels of VEGF of ARPE-19 cells after 24 h exposure to 0.1, 0.25, 0.50, and 1 mg/ml ranibizumab were 891.1±7.3 pg/ml, 828.5±5 pg/ml, 767±4.5 pg/ml, and 870±6.2 pg/ml, respectively.

RT-CES dynamic study for the effect of VEGF and ranibizumab on malignant melanoma cells and ARPE-19 showed that tumor cells are more inclined to be inhibited by ranibizumab: Malignant melanoma and ARPE-19 cells were treated with 1, 5, 10, and 20 ng/ml of VEGF. After adding VEGF to the malignant melanoma cells (Figure 4A), NCI became higher kinetically following the culture time compared with the cells without VEGF. What is more, NCI became higher kinetically following the culture time (p<0.05). VEGF had a significant influence on the viability of malignant melanoma cells. When VEGF was 5, 10, and 20 ng/ml, the malignant melanoma cells’ viability increased by 18%, 20%, and 40%, respectively, compared with the
untreated cells (Figure 4B). This increase effected by VEGF was dose dependent.

There was no evident effect of VEGF on cell proliferation of ARPE-19 cells. As shown in Figure 4C,D, we observed that NCI decreased slightly compared with the untreated cells when VEGF was 1, 5, 10, or 20 ng/ml.

The observations demonstrated that when ranibizumab was added to the malignant melanoma cell system, the NCI was lower compared to that of cells without ranibizumab treatment (Figure 5A). Moreover, the NCI decreased following the culture time (p<0.05). The results indicate that ranibizumab has an inhibitory effect on the growth of malignant melanoma cells. Furthermore, this inhibitory effect induced by ranibizumab was also dose dependent. When ranibizumab...
was 0.1 mg/ml, 0.25 mg/ml, and 0.5 mg/ml, the survival rates of malignant melanoma cells decreased to 12%, 57.5%, and 72% compared with the untreated cells (Figure 5B).

When ranibizumab was only 0.1 mg/ml, the electrode impedance decreased slightly compared with ARPE-19 without ranibizumab (p>0.05). There was a decrease when ranibizumab increased to 0.25 mg/ml, and no evident difference in ARPE-19 proliferation treated with 0.5 mg/ml and 1 mg/ml ranibizumab; the survival rates of ARPE-19 were 55.6% and 42.8%, respectively (Figure 5C,D). The results indicate that a high dose of ranibizumab has an inhibitory effects on the growth of ARPE-19.

**MTT assay for effects of ranibizumab on uveal melanoma and ARPE-19 cells:** As shown in Figure 6, it was observed that the effect of ranibizumab on the two kinds of cells occurred in a dose-dependent manner, which was similar with the results for RT-CES. When ranibizumab was 0.1 mg/ml, 0.25 mg/ml, 0.5 mg/ml, and 1 mg/ml, the survival rates of malignant melanoma cells were 98.1%, 91.05%, 88.55%, and 79.2%, respectively, and the survival rates of ARPE-19 were 98.96%, 96.1%, 89.6%, and 88.03%, respectively (Figure 5C,D). The results indicate that a high dose of ranibizumab has an inhibitory effects on the growth of ARPE-19.

**VEGF** was originally identified as an endothelial cell–specific growth factor that can stimulate angiogenesis and enhance vascular permeability. VEGF-A is the most well-studied member of the VEGF family and is a key target for antiangiogenic therapy [25]. VEGFR1 regulates endothelial cell function indirectly through macrophage recruitment [26], followed by deposition of angiogenic growth factors by these cells. In monocytes, VEGFR1-specific ligands VEGF-B and PIGF both induce signaling pathways known to operate downstream of most tyrosine kinase receptors, such as extracellular signal-regulated kinase (ERK)/mitogen-activated protein kinase (MAPK), phosphatidylinositol-3-kinase.

**DISCUSSION**

VEGF was originally identified as an endothelial cell–specific growth factor that can stimulate angiogenesis and enhance vascular permeability. VEGF-A is the most well-studied member of the VEGF family and is a key target for antiangiogenic therapy [25]. VEGFR1 regulates endothelial cell function indirectly through macrophage recruitment [26], followed by deposition of angiogenic growth factors by these cells. In monocytes, VEGFR1-specific ligands VEGF-B and PIGF both induce signaling pathways known to operate downstream of most tyrosine kinase receptors, such as extracellular signal-regulated kinase (ERK)/mitogen-activated protein kinase (MAPK), phosphatidylinositol-3-kinase.
(PI3K)/protein kinase B (PKB/AKT), and the stress kinase p38MAPK [27], in which VEGFR1-mediated signaling pathways are essential for VEGFR1 biology in vivo. PI GF is also dispensable for embryonic and adult physiological angiogenesis [28], but promotes pathological angiogenesis in several diseases. Moreover, PI GF binds to neuropilin-1 (NRP-1). VEGFR1 is expressed by a broad range of cell types, including human tumor cells [29].

It now appears that VEGF also has autocrine functions acting as a survival factor for tumor cells, protecting them from stresses such as hypoxia, chemotherapy, and radiotherapy [30]. In cancer, tumor angiogenesis contributes to tumor growth and metastasis. An important step in antiangiogenic cancer therapy was taken when the anti-VEGF blocking antibody bevacizumab showed remarkable results in the treatment of metastatic colorectal cancer, and it has been approved by the Food and Drug Administration (FDA) for treatment of metastatic colorectal cancer [31, 32]. Ranibizumab is an affinity-matured antigen-binding fragment (Fab) derived from bevacizumab, and thus has a higher affinity for VEGF-A [33]. Ranibizumab was developed specifically for intravitreal administration to treat vascular eye diseases. Intravitreal injection of 0.5 mg of ranibizumab is recommended for patients with subfoveal choroidal neovascularization (CNV). However, the effect of ranibizumab on tumors has not been clearly identified.

Figure 4. Dynamic response of malignant melanoma and ARPE-19 cell exposure to vascular endothelial growth factor (VEGF). A and B represent the dynamic response of malignant melanoma cells with different concentrations of VEGF, as follows: untreated malignant melanoma cells (control, 0 ng/ml); 1 ng/ml VEGF; 5 ng/ml VEGF, 10 ng/ml VEGF, and 20 ng/ml VEGF. B: Comparison of increases in cell activity in the presence of VEGF after malignant melanoma cells were incubated in the cell system for about 72 h. *p<0.05 compared to untreated cells. C and D show the dynamic response of ARPE-19 with different concentrations of VEGF, as follows: untreated cells (control); 1 ng/ml VEGF; 5 ng/ml VEGF, 10 ng/ml VEGF, and 20 ng/ml VEGF. D: Comparison of increases in cell activity in the presence of VEGF after ARPE-19 were incubated in the cell system for about 72 h. *p<0.05 compared to untreated cells. Data represents three independent experiment and all data points plotted as mean values±SD (*p<0.05).
Uveal melanoma (UM) is an intraocular malignant tumor occurring mainly in Caucasian adults that usually originates from melanocytes of the choroid, iris, and ciliary body [34,35]. UM has an incidence of about 2–8 per million per year in Caucasians [36]. Immunohistochemistry has been the primary tool to distinguish melanomas from other tumors; it is also an adjunct tool to distinguish benign and malignant melanocytic tumors. Useful markers for melanoma include S-100 protein, which is highly sensitive, as well as HMB-45, Melan-A, tyrosinase, and microphthalmia-associated transcription factor (MITF), which are generally more specific [37]. In our study, the results of immunocytochemical staining of malignant melanoma cells and the primary tissue were basically similar; they both showed diffuse staining of S-100 protein and HMB-45, while Melan-A was weakly positive.

The UM cells we cultured were abundant in pigmentation, but lacked pigment after several passages. It has been reported that the production of melanin pigment of human melanoma cells is dependent on tyrosine levels in medium [38]. The results from a nonspecific melanoma experimental model (B16 melanoma) suggested that primary tumors contain cells with variable growth characteristics and metastatic potential [39]. One study [40] focused on the isolation of seven human melanoma cell lines. In the cases derived from pigmented biopsy samples, cellular pigmentation increased as the cells became confluent; this observation was most striking in the multilayered clumps of cells that formed. This
increase in melanin synthesis with increased cell density was accompanied by a parallel increase in tyrosinase synthesis. Cells grown in RPMI 1640 showed less pigmentation than those cultured in DME. As previously reported [41], the density-dependent increase in the rate of tyrosinase synthesis by UCT-Mel 2 could be inhibited by $10^{-6}$ M retinoic acid. Permanent cell lines (UCT-Mel 1 through 7) were established from biopsies of metastatic tissue taken from seven patients with malignant melanoma. University of Cape Town melanoma cell line 2 is the full name of UCT-Mel 2. Although derived from tissue that was pigmented in vivo, UCT-Mel 4 cells could lose the pigment in vitro.

It has long been accepted that angiogenesis plays an important role in tumor growth, invasion, and eventually metastasis. VEGF was shown to be secreted by tumors and to stimulate angiogenesis [42]. In this study, we cultured the tumor specimen obtained from our clinic in vitro, and investigated whether ranibizumab played an inhibitory role in the growth of malignant melanoma. Using ELISA, we examined the VEGF expression levels in malignant melanoma cells and ARPE-19 affected by ranibizumab. The results showed that

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**Figure 6.** MTT assay for effects of ranibizumab on uveal melanoma cells and ARPE-19 cells. A: The dynamic response of malignant melanoma cells with different concentrations of ranibizumab: untreated cells (control); 0.1 mg/ml; 0.25 mg/ml; 0.5 mg/ml and 1 mg/ml. B: The dynamic response of ARPE-19 with different concentrations of ranibizumab: untreated cells (control); 0.1 mg/ml; 0.25 mg/ml; 0.5 mg/ml and 1 mg/ml. Date represents four independent experiment and all data points plotted as mean values±SD (*p<0.05).

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**Figure 7.** VEGF Receptor expression in human malignant melanoma of ciliary body. Total RNA from malignant melanoma of ciliary body was extracted from formalin-fixed, paraffin-embedded tissue and the expression of VEGFR1 and VEGFR2 was examined by RT-PCR.
the levels of VEGF secreted by malignant melanoma cells were much greater than those of ARPE-19, and the levels of VEGF of malignant melanoma cells declined markedly via the action of ranibizumab. On the other hand, there was no evident decrease of VEGF in the presence of ranibizumab for ARPE-19. When VEGF was added to the malignant melanoma cells, the viability of these cells increased greatly, and this increased kinetically following the culture time. However, there was no evident effect of VEGF on the cell proliferation of ARPE-19. Further research showed that the growth of malignant melanoma cells was obviously inhibited when ranibizumab intervened.

The signaling responses at the molecular level in UM need further investigation. It has been reported that the function of VEGFR family proteins is largely restricted to angiogenesis and regulation of vasculogenesis [43]. In this study, the expressions of VEGFR1 and VEGFR2 were examined by PCR.

Our findings indicate that malignant melanoma cells can express more VEGF, and the growth of tumor cells is VEGF dependent. In cell culture, the proliferation of malignant melanoma of the ciliary body can be restrained effectively by anti-VEGF agents. The possible mechanism of the inhibitory effect may be antagonizing VEGF through VEGFR1. In contrast, low-dose exposure of ranibizumab had a slight effect on the growth of ARPE-19. Only high-dose ranibizumab has an inhibitory effect on ARPE-19 cells.

In summary, our study suggests that ranibizumab can selectively inhibit malignant melanoma cell proliferation by decreasing the levels of VEGF. The possible mechanism of the inhibitory effect may be antagonizing VEGFR1. Moreover, our observations demonstrate that although ranibizumab can greatly inhibit ARPE-19 proliferation in vitro at higher concentrations, it has little effect on ARPE-19 at lower concentrations. As this is an in vitro study, which may not simulate the complicated human internal environment perfectly, further in vivo investigations are necessary for exploration of this mechanism.

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