Hypoxic response elements and Tet-On advanced double-controlled systems regulate \( h\text{VEGF}_{165} \) and \( \text{angiopoietin-1} \) gene expression in vitro

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Abstract

Angiogenesis in ischemic tissue is a complex and multi-gene event. In the study, we constructed hypoxic response elements (HRE) and the Tet-On advanced double-controlled systems and investigated their effects on the expression of \( h\text{VEGF}_{165} \) and \( \text{angiopoietin-1} \) (Ang-1) genes in rat cardiomyocytes exposed to hypoxia and pharmacologic induction. We infected neonatal rat cardiomyocytes with recombinant rAAV-rtTA-Rs-M2/rAAV-TRE-Tight-Ang-1 and rAAV-9HRE-\( h\text{VEGF}_{165} \). Our results indicated that the viral titer was \( 1 \times 10^{12} \) vg/mL and the viral purity exceeded 98%. \( h\text{VEGF}_{165} \) expression was induced by hypoxia, but not by normoxia (\( P < 0.001 \)). Ang-1 expression was evident under doxycycline induction, but undetectable without doxycycline induction (\( P < 0.001 \)). Immunofluorescence staining showed that positively stained \( h\text{VEGF}_{165} \) and Ang-1 protein appeared only under both hypoxia and doxycycline induction. We demonstrate here that HRE and the recombinant Tet-On advanced double gene-controlled systems sensitively regulate the expression of \( h\text{VEGF}_{165} \) and Ang-1 genes in an altered oxygen environment and under pharmacological induction in vitro.

Keywords: gene control, vascular endothelial growth factor, angiopoietin-1, hypoxia, doxycycline, cardiomyocyte

INTRODUCTION

Therapeutic angiogenesis is a therapeutic intervention that introduces exogenous angiogenic growth factors to enhance collateral blood flow in the infarcted myocardium and improve global heart performance\(^{[1,2]} \). Although increasing interest in therapeutic angiogenesis has focused on vascular endothelial growth factor (VEGF), one of the most potent angiogenic factors, some studies indicate that VEGF, as an early-phase angiogenic factor, often produces immature vessels that are unable to create functional collateral development\(^{[3-5]} \). Moreover, uncontrolled long-term expression of VEGF delivered by recombinant adenovirus (rAAV) vector in vivo may result in side effects such as hemangioma formation, retinopathy or arthritis\(^{[6]} \).

It is well documented that angiogenesis is a complex event in which multiple angiogenic factors may exert their actions at different phases of angiogenesis\(^{[7]} \). Therefore, for therapeutic angiogenesis, in addition to early angiogenic factors, introduction of late phase angiogenic factors to modify immature vascu-
lature and enhance neovessel function has attracted considerable interest. Data indicate that angiopoietin-1 (Ang-1) plays a critical role in promoting maturation of VEGF-induced vessels in the late phase of angiogenesis\(^\text{[8]}\). However, uncontrolled early Ang-1 expression in ischemic tissues is likely inadequate to remodel VEGF-induced capillaries and does not benefit functional angiogenesis\(^\text{[9,10]}\). Clinically, due to limitations in gene delivery technology, multiple genes are usually transferred simultaneously. However, expression of multiple angiogenic genes at the same time is not consistent with the physiology of angiogenesis. Therefore, timely, controlled expression of these genes in vivo is required. Thus far, the ideal multi-gene expression control system in ischemic heart is still under development.

Our preliminary experiment confirmed that the hypoxic response element (HRE), as an effective gene switch, reliably induced the expression of the hVEGF\(_{165}\) gene expression under hypoxia and re-oxygenation both in vivo and in vitro\(^\text{[11-13]}\). In the present study at the cellular level, we employed recombinant Tet (tetracycline)-On together with the HRE system and investigated the feasibility of regulated expression of hVEGF\(_{165}\) and Ang-1 genes in cardiomyocytes in an altered oxygen environment and under pharmacological induction. We aimed to explore a new approach of therapeutic gene control for further selected multiple gene therapy in ischemic heart disease in vivo.

**MATERIALS AND METHODS**

**Tissues and animals**

A healthy adult lung was obtained from the National Human Genome Center in Shanghai, China. Total human RNA was extracted using TRIzol Reagent (Invitrogen, USA).

Neonatal Sprague-Dawley rats (1-3 d old, weighing 5-7 g, mean 6.3±0.6 g) were obtained from The Experimental Animal Centre of Xuzhou Medical College. The study protocol was approved by the local Institutional Review Board at the authors affiliated institution. Acquisition of human tissue specimens was carried out in accordance with the institution guide line. All animals received humane care in compliance with the Guideline for Care and Use of Laboratory Animals published by Jiangsu Province, China.

**Expression vectors and plasmid construction**

The rAAV-9HRE-hVEGF\(_{165}\) plasmid was a gift from Dr. Hua Su (the Cardiovascular Research Institute, University of California, USA). Recombinant AAV vectors were prepared with a three-plasmid cotransfection system as described previously\(^\text{[13,14]}\). The titer of rAAV-9HRE-hVEGF\(_{165}\) was \(2 \times 10^{12}\) vector genomes (vg/mL).

For the Tet-On advanced system, a reverse Tet Transactivator (rtTA)-Rs-M2, a p-tetracycline response element (pTRE)-Tight, a pTRE-Tight-luc and AAV expression vectors (pSNAV) were purchased from Clonetech, Inc. (USA) and Vector Gene Technology Ltd (Beijing, China). All vector constructs were confirmed by DNA sequencing.

Briefly, total RNA from human lung tissue was isolated using TRlZol Reagent (Invitrogen, USA); 2 \(\mu\)g of total RNA was reverse-transcribed with SuperScript™ III Reverse Transcriptase (Invitrogen, USA) according to the manufacturer’s instructions. PCR amplification was performed using the cDNA isolated from total cellular RNA with the PrimeSTAR™ HS DNA polymerase kit (TaKaRa, Japan). The primers for first-round PCR were F0: 5’-ACGTCACAGAGAGAAACATC-3’ and R0: 5’-CAGCTCTCCTGACATTCTTTGT-3’. The primers for second-round PCR were Ang-1-F: 5’-GGCGTATCCATGAGTTTCCTCTTCC-3’ and Ang-1-R: 5’-GCAAGCTTTCGTCG-3’, which were designed based on the sequence of the Ang-1 gene (GenBank, No. NM 001146) and were synthesized by Invitrogen, USA.

To facilitate cloning, F and R primers contained a BamHI site at the 5’ end of the coding sequence of F and a HindIII site at the 5’ end of R (underlined above). After being recovered from the gel using an agarose gel DNA purification kit (TaKaRa, Japan), the PCR-amplified DNA was added with a poly "A" tail with the DNA A-tailing kit (TaKaRa, Japan) and ligated into the pMD18-T simple vector (TaKaRa, Japan). The primary structure of the insert was confirmed by direct sequencing. The fragment of coding Ang-1 was released from the pMD18-T-Ang-1 by digestion with BamHI and HindIII and subcloned into the expression vector pTRE-Tight as described previously\(^\text{[15]}\). The obtained recombinant eukaryotic expression vector pTRE-Tight-Ang-1 was identified by PCR and then analyzed by the restriction enzymes BamHI/HindIII.

The fragments encoding rtTA-Rs-M2 (765bp), TRE-Tight-Ang-1 (1.8kb) and TRE-Tight-Luc (2kb) were released, respectively, from pTet-On advanced, pTRE-Tight-Ang-1 and pTRE-Tight-Luc by digestion with EcoRI/Bgl II, XhoI/SalI or XhoI/NheI. Afterwards, they were ligated to EcoRI/Bgl II, XhoI/SalI or XhoI/NheI sites of the empty vectors (pSNAV), respectively. The recombinant eukaryotic expression vectors pSNAV-rtTA-Rs-M2, pSNAV- Tight-Ang-1 and pSNAV-TRE-Tight-Luc were identified by PCR...
and digested with EcoRI/BgI II, XhoI/SalI or XhoI/NheI, respectively.

**Viral production determination**

Three plasmid constructs (rtTA-Rs-M2, pTRE-Tight, pTRE-Tight-Luc) were transfected into BHK-21 cells in six-well plates with Lipofectamine 2000 (Invitrogen, USA). The cells were maintained with G418 (800 μg/mL) for 24 h. Three cell lines were designated as BHK/Tet-On, BHK/TRE-Tight-Ang-1 or BHK/TRE-Tight-Luc and were infected with HSV1/rc/ΔUL2 at a multiple of infection (MOI) of 0.1 for 48 h until all the cells exhibited cytopathic effect. The PEG/NaCl precipitated chloroform extraction technique was used to purify the vectors of rAAV-rtTA-Rs-M2, rAAV-TRE-Tight-Ang-1 and rAAV-TRE-Tight-Luc.

The purified plasmids were used as standards, and the probes were labeled by PCR. The serially diluted virus stock suspension and standards preparations were determined by the dot-blot method using digoxigenin-labeled gene as probe. A viral titer was generated from the detected hybridization signals. The protein traps were obtained from 10% SDS-PAGE electrophoresis and the viral purity was also determined.

**Cardiomyocyte culture and identification**

Following ether inhalation anesthesia, thoracotomy was performed. The animal heart was rapidly collected, cut into small pieces and digested with trypsin (1.25 mmol/L) for 10 min. The separated cells were centrifuged at 200 g for 7 min and then suspended in DMEM for 2 h (1% CO₂ and 99% air). To inhibit fibroblast growth, unattached cells were treated with 5-bromodeoxyuridine (BrdU, 25 mmol/L) for 24 h before infection and then maintained in DMEM supplemented with 10% FBS.

For determination of cellular purity, cardiac troponin-I (cTnl) staining was completed with a rabbit polyclonal antibody against cTnl (Chemicon, USA) and secondary goat anti-rabbit IgG antibody (Sigma, USA). After hematoxylin and eosin (HE) staining, cardiomyocytes chosen randomly in six visual fields were visualized under a light microscope. The number of cTnl positively stained cells (N2) and nuclei with HE positive staining (N1) were counted. Cardiomyocyte purity was determined based on the following formula: cellular purity = N2/N1 × 100%.

**Viral infection and treatments**

To assay the optimal proportion of the virus for Ang-1 transfection, rAAV-rtTA-Rs-M2 was mixed with a fixed amount of rAAV-TRE-Tight-Ang-1 (10⁸ vg) in different ratios (1:1, 1:2, 1:3, 1:4, 1:5 and 1:6). After 12 h infection, the cells were treated with 1 μg/mL doxycycline hydrochloride (Dox, Sigma, USA) for 12 h.

An optimal Dox concentration for Ang-1 expression induction was determined. rAAV-rtTA-Rs-M2 and rAAV-TRE-Tight-Ang-1 at a ratio of 1:4 were infected into the cultured cells (10⁶ ug). Ang-1 expression was induced by addition of Dox at different concentrations (0.01-10.0 μg/mL). The cells were harvested after 12 h incubation.

For the determination of an optimal concentration of rAAV-TRE-Tight-Ang-1 for infection, the cells were infected with different concentrations of rAAV-TRE-Tight-Ang-1 (10¹⁰, 5 × 10⁹, 10⁸, 5 × 10⁶ and 10⁵) for 12 h. Then, the cells were treated with 1 μg/mL Dox for 12 h, 10⁹ vg was also used as a control without Dox induction. Similarly, to identify the optimal concentration of rAAV-9HRE-hVEGF165, the cells were infected with different concentrations of rAAV-9HRE-hVEGF165 (10¹¹, 5 × 10¹⁰, 10⁹, 5 × 10⁸, and 10⁶ vg and PBS) for 8 h and then cultured in a hypoxic incubator (1% O₂) for an additional 12 h[12].

The cardiomyocytes were divided into eight groups. Group A: The cells were infected by rAAV-9HRE-hVEGF165 and cultured under hypoxia (1% O₂). Group B: The cells were co-infected with rAAV-9HRE-hVEGF165 and rAAV-rtTA-Rs-M2 / rAAV-TRE-Tight-Ang-1 (1:4) under hypoxia without Dox induction. Group C: The cells were co-infected with rAAV-9HRE-hVEGF165 and rAAV-rtTA-Rs-M2 / rAAV-TRE-Tight-Ang-1 under hypoxia with Dox induction (1 μg/mL). Group D: The cells were co-infected with rAAV-9HRE-hVEGF165 and rAAV-rtTA-Rs-M2 / rAAV-TRE-Tight-Ang-1 under normoxia with Dox induction. Group E: The cells were infected with rAAV-rtTA-Rs-M2 / rAAV-TRE-Tight-Ang-1 under normoxia with Dox induction. Group F: The cells were infected with rAAV-9HRE-Laz under hypoxia, as vector control. Group G: The cells were infected with rAAV-rtTA-Rs-M2 and rAAV-TRE-Tight-Luc under normoxia with Dox induction, as vector control. Group H: An equal volume of PBS was added to the cells under hypoxia with Dox induction, as blank control.

**Western blotting studies**

The cells were solubilized in lysis buffer (100 mmol/L Tris-HCl, 4% SDS, 20% glycerine, 200 mmol/L DTT and protease inhibitors, pH 6.8). Total cellular protein was denatured by boiling for 10 min with an equal volume of 2 × Tris-glycine SDS buffer. Protein was separated by 10% SDS-PAGE and transferred to nitrocellulose membrane (Millipore, USA). After blocking with 5% non-fat milk/PBS-T for 3 h
at room temperature, the membranes were incubated with a goat anti-Ang-1 antibody (Santa Cruz, USA) and a mouse anti-hVEGF<sub>165</sub> antibody (Sigma, USA), respectively. Then, fluorescently labeled secondary antibody (Rockland, USA) was added for 1 h and subsequently scanned by the Odyssey Infrared Imaging System (Li-Cor Biosciences, USA).

**RT-PCR**

Total RNA from cultured cells was extracted using RNAiso Reagent. RT-PCR was performed by using 1 μg of total RNA as described by the M-MLV (RNase H) procedure. The primer sequences of hVEGF<sub>165</sub> were 5′-CTTGCCCTTGCTGCTTACCT-3′ for the forward primer and 5′-CCTGCAACGGGAGTCTGT-3′ for the reverse primer. The primer sequences of Ang-1 were 5′-GCGGATCCATGAGTTCTTTCT-3′ for the forward primer and 5′-GCAAGCTTTCAAAAATCTAAAGGTCG-3′ for the reverse primer.

**Immunofluorescence analysis**

Total mixed virus (rAAV-9HRE-hVEGF<sub>165</sub> and rAAV-rtTA-Rs-M2 / rAAV-TRE-Tight-Ang-1, at 1∶1 ratio, 4×10<sup>10</sup> vg) was added to FBS-free culture medium for 6 h. Then, the cells were cultured under hypoxic conditions (1% O<sub>2</sub>) for 12 h. The cells were then treated with 1 μg/mL Dox under re-oxygenation (95% air / 5% CO<sub>2</sub>) up to 12 h. The harvested cells were stained with a mouse anti-hVEGF<sub>165</sub> antibody (Sigma, USA) and a goat anti-Ang-1 antibody (Santa Cruz, USA) at 4°C for 48 h. The secondary antibodies, conjugated with fluorescein isothiocyanate for VEGF (a goat anti-mouse FITC fluorescent antibody 1∶150, Sigma, USA) or rhodamine for Ang-1 (a rabbit anti-goat FITC fluorescent antibody, Boster LTD, China), were added. The cells were observed under a fluorescence microscope (Olympus, Japan).

**Statistical analysis**

All data are expressed as mean±SD and analyzed by one-way ANOVA and Student-Newman-Keuls test. P values less than 0.05 were considered statistically significant.

**RESULTS**

**Recombinant virus identification and cardiomyocyte purity determination**

A structure of the Tet-On advanced system for Ang-1 gene expression control is shown in **Fig. 1A**. PCR products of pSNAV-TRE-Tight-Ang-1 (1.8 kb) and pSNAV-rtTA-Rs-M2 (765 bp) are shown in **Fig. 1B**. pSNAV-rtTA-Rs-M2 was digested with EcoRI/BglII and two bands (7.1 kb/765 bp) were obtained. Moreover, pSNAV-Tight-Ang-1 was digested with XhoI/SalI and two bands (6.3kb/1.8kb) were available (**Fig. 1B**). The results of the PCR and restriction enzyme analyses indicated that pSNAV-rtTA-Rs-M2, pSNAV-TRE- Tight-Ang-1 and pSNAV-TRE-Tight-Luc were successfully constructed (**Fig. 1B and 1C**). The titer of recombinant virus was 1×10<sup>12</sup> vg/mL and the viral purity of rAAV-rtTA-Rs-M2, rAAV-TRE-Tight-Ang-1 and rAAV-TRE-Tight-Luc was greater than 98%.

The representative photo for the purity of myocardial cells was provided to indicate the identification of myocardial cells (**Fig. 2**), and the purity of the cardiomyocytes was (90±3)%.

**Optimal Dox concentration for induction of Ang-1 expression**

Western blotting studies showed that a low basal level of Ang-1 protein expression was detected in the absence of Dox. A higher level of Ang-1 protein expression was achieved with 1 μg/mL Dox induction. Ang-1 expression was found in a dose-dependent manner with Dox concentration (**Fig. 3A**).

**Optimal proportion of rAAV-rtTA-Rs-M2 and rAAV-TRE-Tight-Ang-1**

We found that when the ratio of rAAV-rtTA-Rs-M2 and rAAV-TRE-Tight-Ang-1 was set to 1∶4, a high level of Ang-1 protein expression was achieved. Even at the optimal ratio (1∶4) of rAAV-rtTA-Rs-M2 and rAAV-TRE-Tight-Ang-1, there was no Ang-1 protein expression without Dox induction (**Fig. 3B**).

**Optimal concentrations of rAAV-TRE-Tight-Ang-1 and rAAV-9HRE-hVEGF<sub>165</sub>**

Under Dox induction, a high level of Ang-1 protein expression was evident when the total amount of rAAV-TRE-Tight-Ang-1 reached 10<sup>10</sup> vg. Ang-1 protein expression was decreased following reduction of rAAV-TRE-Tight-Ang-1. There was no Ang-1 protein expression in the absence of Dox regardless of the amount of rAAV-TRE-Tight-Ang-1 (**Fig. 3C**). A higher level of hVEGF<sub>165</sub> protein expression was obtained with 10<sup>10</sup> vg virus (rAAV-9HRE-hVEGF<sub>165</sub>). Following decrease of rAAV-9HRE-hVEGF<sub>165</sub>, hVEGF<sub>165</sub> protein expression was reduced simultaneously (**Fig. 3D**).

**hVEGF<sub>165</sub> and Ang-1 expression**

RT-PCR determination indicated that hVEGF<sub>165</sub> mRNA bands of about 484 bp were found in group A, B and C, but not in the other groups (P < 0.001). Ang-1 mRNA bands about 1497 bp were found in group C,
D and E where Dox was administered, but not identified in group F, G and H \((P < 0.001)\). Both \(h\text{VEGF}_{165}\) and \(\text{Ang-1}\) mRNA expression was found in group C only (Fig. 4A).

Western blotting determination indicated that \(h\text{VEGF}_{165}\) protein expression was detected in group A, B and C, but not in the other groups \((P < 0.001)\), while \(\text{Ang-1}\) protein expression was found in group C, D and E, but not identified in the other groups \((P < 0.001)\). Both \(h\text{VEGF}_{165}\) and \(\text{Ang-1}\) protein expression appeared in group C only (Fig. 4B).

Immunofluorescence analyses also showed that both \(h\text{VEGF}_{165}\) and \(\text{Ang-1}\) protein immunofluorescence could be observed in group C. The expressed sites of \(h\text{VEGF}_{165}\) and \(\text{Ang-1}\) protein were located in the cytoplasm rather than in the nucleus (Fig. 5).

**DISCUSSION**

It is generally recognized that angiogenesis is a complex and multi-gene event. Making therapeutic angiogenesis more closely resemble physiological processes in tissue is extremely important. Thus, multi-angiogenic gene control is technically required to achieve this goal. Physiologically, angiogenesis in tissue relies on two sequential phases including an early phase that strictly depends on the presence of an early angiogenic factor (i.e. VEGF). During the VEGF-dependent period, endothelial cells, mainly originating from the local pre-existing vasculature, become activated to form a set of immature and irregularly shaped vessels surrounded by a thin endothelial layer. This is
followed by a late phase where vessel maturation induced by a late angiogenic factor is required to ensure the proper acquisition of functional competence of the newly formed vasculature. During this period, VEGF is not only obsolete but is even detrimental for vessel functionality and pericyte regeneration\cite{16,17}. Therefore, the approach developed here is an appealing alternative to the simultaneous delivery of VEGF together with other angiogenic factors. Ang-1, one of the late angiogenic factors, has been proven to be capable of promoting angiogenesis and, in synergy with VEGF, maintaining stability and integrity of the mature vasculature by mediating interaction between endothelial cells and their underlying support cells\cite{10,21}. Furthermore, Ang-1 is the first angiogenic factor identified to exert a protective effect on the vascular endothelial barrier by blocking the action of permeability-increasing mediators, such as VEGF\cite{8}.

Lee et al.\cite{22} reported that VEGF and Ang-1 expression was upregulated in human ischemic myocardium. VEGF expression progressively increased over time after the onset of acute ischemia, peaked at 6 weeks and was followed by Ang-1 expression. Therefore, during therapeutic angiogenesis with VEGF and Ang-1 genes, the accurate and timely control of Ang-1 gene expression behind early hypoxia-mediated VEGF expression may be a key point for multiple gene-induced angiogenesis in an ischemic heart\cite{23,24}. So far, an ideal in-

**Fig. 3 Determination of inducible gene expression of Ang-1.** A: High levels of Ang-1 expression were found with 1 μg/mL Dox induction. Compared with groups 0, 0.01, 0.1, 5.0 and 10.0, \(P < 0.05\). B: There was no significant difference among ratios of 1:1 to 1:4 (rAAV-rTA-Rs-M2 / rAAV-TRE-Tight-Ang-1). Compared with groups 1:1, 1:2, 1:3 and 1:4, \(P < 0.05\). C: There was no significant difference among concentrations of 10\(^{11}\), 5×10\(^{10}\) and 10\(^9\) (rAAV-TRE-Tight-Ang-1). Compared with groups 10\(^{11}\), 5×10\(^{10}\) and 10\(^9\), \(P < 0.05\). D: There was no significant difference among concentrations of 10\(^{11}\), 5×10\(^{10}\) and 10\(^9\) (rAAV-9HRE-hVEGF\(_{hs}\)). Compared with groups 10\(^{11}\), 5×10\(^{10}\) and 10\(^9\), \(P < 0.05\).
ducible expression control system for \textit{Ang-1} in vivo is the so-called TRE-regulated gene system. The Tet-On advanced system has several advantages over other regulated mammalian gene expression systems: low basal expression, high inducibility, no pleiotropic effects and high absolute expression levels. In our study, we successfully constructed a new Tet-On advanced system for \textit{Ang-1} gene inducible expression. We found that, at the cellular level, the Tet-On advanced system effectively activated \textit{Ang-1} gene expression with the addition of Dox, a tetracycline (Tc) derivative, to the culture medium and was rapidly turned off upon withdrawal of the antibiotic. Furthermore, \textit{Ang-1} expression was tightly regulated dose-dependently in response to varying concentrations of Dox and was independent of the absolute amount of rAAV-TRE-Tight-Ang-1. Dox is a commonly used broad-spectrum antibiotic. In biology, Dox is usually employed in the inducible expression of Tet-on / Tet-Off system or as an inhibitor for suppression of target gene expression\cite{25,26}. Our result suggests that Dox induction was fully necessary for enabling gene control over \textit{Ang-1} expression. We noted that the background gene expression of \textit{Ang-1} was extremely

\textit{Fig. 4} Gene expression of hVEGF\textsubscript{165} and \textit{Ang-1} in cardiomyocytes. A: hVEGF\textsubscript{165} mRNA (484bp) was evident in group A, B and C under hypoxia. \textit{Ang-1} mRNA (1,497bp) was found in group C, D and E with Dox induction, but was undetected in the other groups. Compared with group D, E, F, G and H, $^*$\textit{P} < 0.001. Compared with group A, B, F, G and H, $^\#$\textit{P} < 0.001. B: Expression of hVEGF\textsubscript{165} protein was detected in group A, B and C under hypoxia. \textit{Ang-1} protein expression appeared in group C, D and E with Dox induction, but was undetected in the other groups. Compared with group D, E, F, G and H, $^*$\textit{P} < 0.001. Compared with groups A, B, F, G and H, $^\#$\textit{P} < 0.001.

\textit{Fig. 5} Representative figures of immunofluorescence staining. rAAV-9HRE-hVEGF\textsubscript{165} and rAAV-(rtTA-Rs-M2/TRE-Tight-Ang-1) were co-infected into cardiomyocytes in group C under hypoxia and Dox induction. A: Red fluorescence (\textit{Ang-1}). B: Green fluorescence (hVEGF\textsubscript{165}). C: Merged figure of A and B illustrating overexpression of both hVEGF\textsubscript{165} and \textit{Ang-1} proteins (Scale bar = 50 \textmu m).
low in the absence of Dox due to the combined effect of the modified rtTA and TRE-Tight, while the maximal level of Ang-1 gene expression was comparable with that obtained from strong, constitutive mammalian promoters such as cytomegalovirus (CMV) [27,28]. This Tet-On advanced system gave us ready access to tightly regulated, high level Ang-1 gene expression.

As for rAAV-rtTA-Rs-M2 and rAAV-TRE-Tight-Ang-1, an optimal proportion should meet the need of obtaining the highest level of Ang-1 expression with the lowest dose of Dox. We found no significant difference of Ang-1 expression in the range of 1:1 to 1:4 (rAAV-rtTA-Rs-M2/rAAV-TRE-Tight-Ang-1). In our recombinant Tet-On advanced system, rtTA-Rs-M2 acted as a control element for Dox. Therefore, under the premise of keeping the highest level of Ang-1 expression, our selected ratio of 1:4 guaranteed that the exact amount of rtTA-Rs-M2 was minimized. From the clinical point of view, a low dose of Dox for target gene induction would further decrease possible side effects and improve safety of antibiotic induction.

Concerning the combined efficiency of targeted gene control, hVEGF<sub>165</sub> and Ang-1 expression could be induced by altered oxygen and Dox administration, respectively. We noted that one gene control system had no significant interference on the inducible expression of the other target gene, although expression of hVEGF<sub>165</sub> or Ang-1 in the co-infected group was lower than that in the single gene infected group.

Based on our results, following early hypoxia-mediated hVEGF<sub>165</sub> expression, it is feasible to realize timely control over Ang-1 gene expression with a Tet-On advanced system. Our findings present a strategy of enhanced therapeutic angiogenesis with the sequentially controlled expression of multiple angiogenic genes transferred simultaneously in vitro.

Nevertheless, although HRE and the Tet-On advanced systems acted as effective double gene switches in vitro, our new system needs to be validated in animal models in which actual ischemic myocardial pathologies can be reproduced. Additionally, the combined efficiency of VEGF and Ang-1-driven therapeutic angiogenesis should be further evaluated.

In conclusion, HRE and the Tet-On advanced systems have been verified to be new promising gene switches for VEGF and Ang-1-inducible expression control in vitro. Thus, this innovative double gene control system leads to a broader prospect for effective and safe angiogenic therapy in ischemic heart disease in vivo.

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