RNF213 gene silencing upregulates transforming growth factor β1 expression in bone marrow-derived mesenchymal stem cells and is involved in the onset of Moyamoya disease

CHANGSHUI WANG1, CUILIAN SUN1, YUESHU ZHAO1, HUIMIN SONG1, ZHENGYOU LI2, FENG JIN1 and CHANGMENG CUI1

1Department of Neurosurgery, Affiliated Hospital of Jining Medical University, Jining, Shandong 272000; 2Department of Neurosurgery, Shandong Province Western Hospital, Shandong Province ENT Hospital, Jinan, Shandong 250022, P.R. China

Received August 27, 2020; Accepted December 12, 2020

DOI: 10.3892/etm.2021.10456

Abstract. Moyamoya disease (MMD) is a chronic and progressive cerebrovascular occlusion disease, the precise etiology of which is poorly understood. Ring finger protein 213 (RNF213) has been previously identified as a susceptibility gene that serves an important role in angiogenesis, where it has been shown to be closely associated with the onset of MMD. Patients with MMD exhibit increased expression levels of various pro-inflammatory molecules and angiogenic factors. Under certain conditions, bone marrow mesenchymal stem cells (BMSCs) have the ability to differentiate to form neuron-like and microglia-like cells. In the present study, a total of 40 MMD patients and 40 healthy individuals were enrolled. ELISA assays revealed that the expression of serum vascular endothelial growth factor (VEGF) and transforming growth factor β1 (TGF-β1) were higher than that in healthy controls. Furthermore, rat BMSCs (rBMSCs) were isolated and cultured using the whole bone marrow adherence method, which were then phenotyped using flow cytometry. Osteogenic and adipogenic differentiation were determined by using Alizarin red and oil red O staining, respectively. RNF213 was knocked-down using a lentivirus-mediated short hairpin RNA system in passage three rBMSCs, and successful transfection of the RNF213 was confirmed by RT-qPCR and fluorescence imaging. The expression levels of VEGF and TGF-β1 in these rBMSCs were measured on days 7 and 14, respectively. The results demonstrated that RNF213 knockdown upregulated TGF-β1 at both protein and mRNA levels, but did not exert any effect on VEGF gene expression. In conclusion, these findings suggested that that RNF213 knockdown may contribute to aberrant TGF-β1 expression via a pathway that remains to be unidentified, indicating that quantitative changes in RNF213 gene expression may serve an important role in the pathogenesis of MMD.

Introduction

Moyamoya disease (MMD) was first defined by Suzuki and Takaku in 1999 (1) and presents as the presence of an abnormal vascular network at the base of the brain. MMD is characterized by progressive stenosis and even occlusion of the terminal portion of the bilateral internal carotid and intracerebral arteries (2). The most common symptoms of MMD include headache, ischemia and hemorrhage (3). The incidence of MMD in Japan has increased from 0.35 per 100,000 individuals in 1995 to 0.94 per 100,000 individuals in 2006 (4). MMD has mostly been reported to exhibit a high incidence in various East Asian populations, including Chinese, Japanese and Korean (5). Although basic studies, including those using genomic and proteomic approaches, have been performed, the precise etiology of MMD remains unclear.

An increasing number of studies have demonstrated that, although the occurrence and development of MMD are multifactorial, genetic factors are closely associated with the pathophysiology of MMD (6,7). More recently, a variety of loci were found to be involved in MMD, including 17q25, 8q23, 6q25 and 3p24-p26 (8-11). In particular, genome-wide and locus-specific association studies have identified ring finger protein 213 (RNF213) in the 17q25-ter region as a novel susceptibility gene for MMD (12). RNF213 has been shown to serve an important role in angiogenesis and is closely associated with the onset of MMD (13-15). A previous study has revealed that the large trunk arteries have irregular diameters and abnormal sprouting occurs in zebrafish following RNF213 knockdown (16). In addition, MMD is also frequently accompanied with hypertension (17). A possible reason for this is that changes in RNF213 expression can result in increased blood pressure, which aggravates...
intracranial hemodynamic disorders further and induce the formation of 'smoky' blood vessels (18,19). A homozygous variant of RNF213 is considered to be the most pathogenic and was found to be significantly associated with severe manifestations of MMD (20). The mutation rate of the rs112735431 locus of RNF213 was high among Japanese patients with familial and sporadic Moyamoya (21), with similar findings observed in the Han Chinese population (22,23). RNF213 has also been recognized as an effective biomarker for predicting MMD prognosis (24). Alternatively, a variety of biomarkers and angiogenic factors, including vascular endothelial growth factors (VEGF), cytokines, such as matrix metallopeptidase 9, have been previously reportedly implicated in MMD (25).

Based on this evidence, it was hypothesized that the RNF213 gene silencing may promote the abnormal expression of MMD-associated factors through a specific gene pathway, leading to the development of MMD.

Mesenchymal stem cells (MSCs) possess high self-renewal abilities and multidirectional differentiation potentials (26). In certain conditions, they can differentiate to form neuron-like and microglia-like cells (27,28) MSCs has been reported not only to increase endothelial cell growth, but also to promote skin wound healing through vascular endothelial growth factor-C-mediated angiogenesis (29,30). In addition, MSCs have the advantage of being easily obtained and reportedly exert neuroprotective effects against ischemic brain damage (31), making their use increasingly popular for basic research and clinical studies. Cerebral hemorrhage and ischemia are common pathophysiological states in neurosurgical diseases, including MMD and subarachnoid hemorrhage (32). After cerebrovascular events, neurocognitive function is impaired (33). MSCs has been shown to confer beneficial effects against cerebrovascular diseases (34). It has also been reported that there were no significant complications within 5 years of MSC treatment for ischemic stroke (35). However, to the best of our knowledge, no studies have focused on the effects of the susceptible gene RNF213 on MMD by using MSCs. The rat bone marrow mesenchymal stem cells (rBMSCs) were therefore selected for the present in vitro study, where the aim was to determine the effects of RNF213 silencing on the expression of VEGF and transforming growth factor β1 (TGF-β1) in rBMSCs and to investigate the association between the RNF213 gene expression and MMD occurrence.

Materials and methods

Subjects. A total of 40 patients with MMD were enrolled into the Departments of Neurosurgery and Neurology of the Affiliated Hospital of Jining Medical University (Jining, China) between September 2014 and April 2015. The selection criteria were as follows: i) Digital subtraction angiography (DSA) confirming MMD and patients meeting the Guidelines for Diagnosis and Treatment of Moyamoya Disease (Spontaneous Occlusion of the Circle of Willis) (36); ii) CT or MRI examination confirming cerebral infarction and cerebral hemorrhage or ischemia; iii) patients with complete clinico-pathological data. The exclusion criteria were as follows: i) Patients with a history of autoimmune disease or neuropsychiatric diseases; ii) patients with intracranial tumors, severe brain injury and cardiovascular disease; iii) use of any medications that could affect cognitive function. An additional 40 healthy individuals were recruited as the control group from the physical examination center of the Affiliated Hospital of Jining Medical University during the same period as aforementioned, and the individuals were confirmed to exhibit no abnormality by laboratory and imaging examinations. The study protocol and sample collection procedures were approved by the Ethics Committee of the Affiliated Hospital of Jining Medical University and written informed consent was obtained from each participant.

Determination of serum VEGF, TGF-β1 and RNF213 levels. Morning fasting blood samples (5 ml each) were collected from each subject by elbow venipuncture. Following collection, the samples were left to stand at room temperature for 1 h and then centrifuged at 3,000 x g at 4°C for 20 min. The supernatant was removed and centrifuged for a second time at 3,000 x g at 4°C for 10 min to obtain serum, which was stored at -80°C until analysis. The concentrations of VEGF (cat. no. DVE00; R&D Systems, Inc.), TGF-β1 (cat. no. DB100B; R&D Systems, Inc.) and RNF213 (cat. no. JL19472; Shanghai Jianglai Biological Technology Co., Ltd.; http://www.jonln.com) were determined by ELISA, according to the manufacturer's protocols.

Animals and rBMSC culture. A total of 25 male Sprague-Dawley rats (weight, 90-100 g; age, 4 weeks) were purchased from Jinan Pengyue Experimental Animal Breeding, Co., Ltd. The rats were housed under standard conditions of temperature (22±2°C), light (12:12 h light/dark cycle) and humidity (50±5%) with free access to food and water. The rats were acclimatized for 7 days before the experiment. All animal procedures were conducted in accordance with Guide for the Care and Use of Laboratory Animals with the approval of Affiliated Hospital of Jining Medical University. rBMSCs were isolated according to the following method. Briefly, the rats were anesthetized by 80% CO2 (45 sec), sacrificed by cervical dislocation and then immersed in 75% alcohol for 10 min for disinfection. Following removal of the double lower limb skin, the femur and tibia were aseptically isolated and then soaked in PBS with 1% penicillin-streptomycin for 1 min. The samples were transferred to 15-ml centrifuge tubes. The ends of the bones were removed and the bone marrow cavities exposed. The marrow cavity was washed 5-10 times with complete medium, namely 89% Dulbecco's modified Eagle's medium/Nutrient Mixture F-12 medium (Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% fetal bovine serum and 1% penicillin-streptomycin (Gibco; Thermo Fisher Scientific, Inc.) until the medium was clear. All the flushing fluid was then inoculated into a culture flask and incubated at 37°C and 5% CO2. The medium was replaced after 48 h. Based on assessment of proliferation, the medium was changed every 2 days. At ~80% confluence, the cells were harvested with 0.25% trypsin-EDTA and sub-cultured at a ratio of 1:2.

Alizarin red and oil red O staining. Cells were seeded into 6-well plates at a density of 2x10⁴ cells/well and incubated overnight at 37°C with 5% CO2 in StemXVivo™
Osteogenic/Adipogenic Base Media (cat. no. CCM007; R&D Systems, Inc.). After reaching 100% confluence, the medium was replaced with StemXVivo® Adipogenic Differentiation Media (cat. no. CCM011; R&D Systems, Inc.) to induce adipogenesis. After reaching 70% confluence, the medium was replaced with StemXVivo® Osteogenic Differentiation Media (cat. no. CCM009; R&D Systems, Inc.) to induce osteogenesis. The differentiation medium was replaced every 3 days. After 14 days of adipogenic induction or 21 days of osteogenic induction, MSCs were harvested and fixed with 4% paraformaldehyde for 30 min at 4˚C. After washing twice in PBS, cells were stained with Alizarin red for 25–30 min with antibodies against the following cell surface antigens: CD29 (dilution, 1:50; cat. no. AF2405; R&D Systems), CD106 (dilution, 1:200; cat. no. ab134047; Abcam), and CD34 (dilution, 1:200; cat. no. ab81289; Abcam) antibodies. Cells were washed three times and incubated with FITC-conjugated anti-goat (dilution, 1:200; cat. no. ab6881; Abcam) or FITC-conjugated anti-rabbit (dilution, 1:200; cat. no. ab6717; Abcam) secondary antibodies. Flow cytometry was performed using a CytoFLEX flow cytometer (Beckman Coulter, Inc.), and the data were analyzed with FlowJo software (version 10.4.2; FlowJo LLC).

Immunofluorescence. Cells were fixed with 4% paraformaldehyde for 15 min at room temperature and permeabilized with 0.2% Triton X-100 for 20 min at room temperature. After rinsing with PBS, cells were then incubated with CD106 (dilution, 1:200; cat. no. ab134047; Abcam) and CD34 (dilution, 1:200; cat. no. ab81289; Abcam) antibodies for 2 h at room temperature. After three washes with PBS, cells were incubated with FITC-conjugated anti-rabbit (dilution, 1:200; cat. no. ab6717; Abcam) secondary antibody. DAPI was used for nuclear staining for 3 min at room temperature. The fluorescence staining was captured with an inverted fluorescence microscope (Olympus IX73; Olympus Corporation).

Lentiviral transfection. pSIH1-H1-copGFP was used to construct the lentiviral vectors. The lentiviral vectors encoding the RNF213 short hairpin RNA (RNF213-shRNA) were designed by Hanbio Biotechnology Co., Ltd. A scramble sequence, named negative-shRNA, was used as the negative control. The rBMSCs were divided into the following three groups: i) RNF213-shRNA; ii) negative-shRNA; and iii) control. The control group was not transfected with plasmids. For lentiviral infection, rBMSCs in the logarithmic growth phase were seeded into 12-well plates at a density of 5x10⁴ cells/well. After 24 h, lentivirus was added at a multiplicity of infection of 100. The cells were maintained at 37˚C (5% CO₂) for an additional 24 h, after which the medium was replaced with 1 ml fresh complete medium (containing 10% fetal bovine serum and 1% penicillin-streptomycin). The cells were incubated for another 24 h and then used for further analysis. After 3 days, green fluorescent protein (GFP) expression was assessed using an inverted fluorescence microscope (magnification, x100).

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total RNA was extracted from rBMSCs using an EZ-10 spin column total RNA isolation kit (cat. no. B610583; Sangon Biotech Co., Ltd.), according to the manufacturer's protocol. cDNA was generated using the Quant one step qRT-PCR Kit (cat. no. KR118; Tiangen Biotech Co., Ltd.). The reverse transcription reaction was carried out at 42˚C for 15 min, followed by 95˚C for 3 min. qPCR was subsequently performed in a Bio-Rad CFX96 instrument (Bio-Rad Laboratories, Inc.) using a SYBR SuperReal PreMix Plus (cat. no. PP205; Tiangen Biotech Co., Ltd.) and gene-specific primers (Table I). The thermocycling conditions were as follows: Initial denaturation at 95˚C for 2 min, followed by 40 cycles of amplification at 95˚C for 10 sec and 60˚C for 32 sec. β-actin was used as the internal standard and each cDNA was tested in triplicate. The 2^ΔΔCq method was used for quantitative analysis (37).

Western blotting (WB). For WB, total protein was isolated from the rBMSCs cells with the use of RIPA and PMSF buffer (Beijing Solarbio Science & Technology Co., Ltd.) according to the manufacturer's instructions. Protein concentration was determined using the bicinchoninic acid protein assay kit (Beijing Solarbio Science & Technology Co., Ltd.). The samples were separated by 12% SDS-PAGE with ~50 µg protein in each lane and transferred onto PVDF membranes.

### Table I. Primer sequences used for reverse transcription-quantitative PCR analysis.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer sequence (5'-3')</th>
<th>Reverse primer sequence (5'-3')</th>
</tr>
</thead>
<tbody>
<tr>
<td>RNF213</td>
<td>CAGCGTGTAGGCGACATCAA</td>
<td>TTGTACTGGGCTGTAGTAGC</td>
</tr>
<tr>
<td>VEGF</td>
<td>CTGCTGTGGAGCTTGAGTGG</td>
<td>CAAACAGACTCCGGCTCCTC</td>
</tr>
<tr>
<td>TGF-β</td>
<td>CAATTCCTGGGGCTACCTTG</td>
<td>AGCCCTGTATCCGTCGCTC</td>
</tr>
<tr>
<td>β-actin</td>
<td>TCAGGTCATCATACTCCGAAT</td>
<td>AAAGAAAGGGTGTAACCGA</td>
</tr>
</tbody>
</table>

RNF213, ring finger protein 213; VEGF, vascular endothelial growth factor; TGF-β1, transforming growth factor β1.
Non-specific binding was blocked by 5% skimmed milk in TBS-0.1% Tween-20 (TBST) buffer and incubated overnight at 4°C with the following primary antibodies: TGF-β1 (dilution, 1:1,000; cat. no. ab215715; Abcam), VEGF (dilution, 1:1,000; cat. no. ab231260; Abcam) and β-actin (dilution, 1:1,000; cat. no. ab8227; Abcam). Following washing with 0.1% TBST, the membranes were subsequently probed with horseradish peroxidase-conjugated secondary antibody for 1 h at room temperature (dilution, 1:2,000; cat. no. ab205718; Abcam). Western blot data were detected using enhanced chemiluminescence solution (Beijing Solarbio Science & Technology Co., Ltd.) and quantified using ImageJ software (version 1.8.0; National Institutes of Health). β-actin was used as the loading control.

Statistical analysis. The results are presented as the mean ± SD and were analyzed using the SPSS statistics software (version 23.0; IBM Corp.). Sex distribution was compared using the χ² test. Other groups were compared using an independent t-test for normally distributed variables. Differences between groups were assessed by one-way analysis of variance followed by Duncan's test. Pearson correlation analysis was used to analyze the correlation between various factors. P<0.05 was considered to indicate a statistically significant difference.

Results

Demographic and clinical characteristics of MMD. The typical morphology of MMD, as determined by DSA, is shown in Fig. 1. The demographic and clinical patient characteristics are presented in Table II. The age of MMD onset was 15-61 years, where the hemorrhage types involved included intraventricular, cerebral lobe and subarachnoid; and the ischemic types involved included cerebral infarction and insufficiency of cerebral blood supply. No significant associations in age, sex and body mass index were observed between the healthy control (aged 20-64 years) and MMD groups.

Serum VEGF and TGF-β1 levels. For patients with MMD, the VEGF level was 450.06±299.78 pg/ml, whilst that in the control group was 271.43±124.42 pg/ml. Similarly, increased levels of TGF-β1 were observed in the MMD group, compared with those in the healthy controls (Fig. 2A). The TGF-β1 level was 62.93±12.95 pg/ml in patients with MMD and 40.08±8.94 pg/ml in the control group (Fig. 2B). Accordingly, the levels of VEGF and TGF-β1 were significantly higher in the patient group, as compared with those in the control group (P<0.01; Fig. 2). The expression levels of RNF213 in the MMD and healthy control groups were also measured. However, there was no difference between the two groups (Fig. S1A). In addition, correlation between RNF213 and TGF-β1 in serum was analyzed, where there was no significant correlation (Fig. S1B).

rBMSC morphological alterations and identification. After 12 h of culture, the rBMSC began to adhere to the culture flask, displayed acceptable proliferation and exhibited a round or fusiform appearance with a clear outline (Fig. S2A). Following culture for 48 h, the rBMSCs exhibited a polyhedral shape with evidence of cell clustering, forming colonies after 3 days (Fig. 3A and B). After 3 days, the cell fusion rate had reached 80-90%. Passage three rBMSCs underwent...
adipogenic induction for 14 days, which appeared as red lipid droplets with oil red O staining (Fig. 3C). Osteogenesis was also induced in third-passage rBMSCs for 21 days, presenting as red, clear, densely-calcified nodules following Alizarin red staining (Fig. 3D).

Subsequently, flow cytometry confirmed that CD29 (Fig. 4A) and CD106 (Fig. 4B) were positively expressed, with no expression of CD34 (Fig. 4C) and CD45 (Fig. 4D). The immunofluorescence analysis indicated that CD106 were highly expressed, but not CD34 (Fig. 4E). All the results indicated that rBMSCs were successfully isolated in high purity.

RNF213 silencing in rBMSCs. At 2 days post-transfection, the rBMSCs exhibited normal morphology under an optical microscope (Fig. 5A), where the expression of GFP was detected by fluorescence microscopy (Fig. 5B). RNF213 mRNA expression analysis was subsequently conducted. Compared with control group, rBMSCs transfected with RNF213-shRNA displayed a significantly lower RNF213 expression on days 7 and 14, and there was no significant difference between the negative-shRNA and control groups (Fig. 5C). These data suggested that the successful silencing of the RNF213 gene was achieved in rBMSCs.
Figure 4. Expression of rat bone marrow mesenchymal stem cell markers, as detected by flow cytometry and immunofluorescence. CD29 and CD106 were positively expressed, whilst no CD34 or CD45 expression was observed. (A) CD29, (B) CD106, (C) CD34 and (D) CD45. (E) Immunofluorescence analysis of CD106 and CD34 expression in rBMSCs (scale bar, 200 µm).

Figure 5. Successful transfection and silencing of RNF213 expression in rBMSCs. (A) Visualization of rBMSCs under normal culture conditions 2 days after the transfection, as determined by optical microscopy (magnification, x100). (B) Expression of green fluorescent protein in rBMSCs 2 days after the transfection under an inverted fluorescence microscope (magnification, x100). (C) Relative RNF213 mRNA levels were determined by reverse transcription quantitative polymerase chain reaction, where β-actin was used as the internal control. Data are presented as the mean ± SD and were analyzed by one-way ANOVA followed by Duncan's test. **P<0.001 vs. Control. RNF213, ring finger protein 213; rBMSCs, rat bone marrow mesenchymal stem cells; shRNA, short harpin RNA.
Figure 6. RNF213 silencing increases TGF-β1 but not VEGF expression. (A and B) mRNA expression as measured using reverse transcription-quantitative PCR. β-actin was used as the internal control. (A) TGF-β1 mRNA expression in rBMSCs was significantly increased on days 7 and 14 post-RNF213 silencing. (B) VEGF mRNA expression in rBMSCs had not changed on days 7 and 14 post RNF213-silencing. *P<0.001 vs. Control. RNF213, ring finger protein 213; TGF-β1, transforming growth factor β1; VEGF, vascular endothelial growth factor; rBMSCs, rat bone marrow mesenchymal stem cells; shRNA, short hairpin RNA.

**Effects of RNF213 silencing on TGF-β1 and VEGF mRNA expression.** As shown in Fig. 6, the TGF-β1 expression levels in the RNF213-shRNA group were significantly higher compared with those in the control groups, and there was no significant difference between the negative-shRNA and control groups. The expression level of TGF-β1 was > three times greater compared with that in the control group on day 7, and > two times greater compared with that in the control group on day 14 (Fig. 6A). However, no significant differences in the VEGF expression were observed between the RNF213-shRNA and the control groups on both days (Fig. 6B).

**Effects of RNF213 knockdown on TGF-β1 and VEGF protein expression.** RNF213 silencing increased TGF-β1 expression, as compared with that in the control group on days 7 and 14 after transfection, respectively, and there was no significant difference between the negative-shRNA and control groups (Fig. 7A, B and D). The VEGF protein expression level in the negative-shRNA group and RNF213-shRNA group did not significantly differ from the control group. (Fig. 7C, E and F).

**Discussion**

MMD is a chronic and progressive cerebrovascular occlusion disorder, where the etiology and pathogenic mechanism of which remain poorly understood. The basic pathological feature of MMD is an abnormal vascular network formed at the base of the skull (38). The age of onset distribution for MMD has been suggested to have two peaks, one at 5 and another at ~40 years of age (39). Regarding gender, two previous studies mentioned the female-to-male ratio. One survey conducted in hospitals throughout Japan reported that the ratio was 1.8 (40), whilst another survey conducted in Hokkaido showed that the ratio was 2.18 (41). Epidemiological analyses have reported distinctive features of familial Moyamoya disease, where the male to female ratio at 5.0 in familial cases (42). Surgical treatment constitutes the main available treatment for MMD with no pharmacological options currently available (43). In a previous study that included 10 years of follow-ups, anti-platelet medications were not found to affect the incidence of cerebral infarction in patients with Moyamoya disease (44). Surgical revascularization is the most successful therapy used to improve cerebral hemodynamics and reduce the risk of subsequent strokes (45). Typical revascularization surgeries for MMD include direct, indirect and combined procedures (46). However, due to the heterogeneity of MMD, the optimal surgical procedures for both ischemic and hemorrhagic MMD remain unclear.

Previous studies have successively determined that VEGF and TGF-β1 are abnormally expressed in the plasma of patients with MMD (47-49). To the best of our knowledge, VEGF is the most prominent promoting factor of angiogenesis, with the ability to induce endothelial cell proliferation and increase vascular permeability (50). It has also been reported to stimulate the proliferation of neuronal precursors both in vitro and in vivo (51). Vascular endothelial growth factor is a key positive regulator of both physiological and pathological angiogenesis (52,53). Therefore, VEGF serves an important role in promoting the formation of new ‘smoke-like’ blood vessels and the establishment of collateral circulation in the brain. Perl et al (54) found that the expression of VEGF and its receptor were significantly increased in the brains of patients with ischemic cerebrovascular disease following long-term ischemia and hypoxia stimulation. In the present study, VEGF was found to be more highly expressed in the serum of patients with MMD compared with that in the control group. This suggests that serum VEGF elevation may be involved in the development of abnormal vascular networks at the base of the brain.

TGF-β1 serves an important role in promoting angiogenesis and neuroprotection, in addition to promoting anti-inflammatory and chemotactic processes (55,56). A previous study reported that the delivery of a TGF-β1 expression plasmid into the arteries resulted in intimal and medial hyperplasia (57). This phenomenon is similar to the pathological changes observed on the arteries of patients with MMD. The expression of TGF-β1 was found to be significantly higher in the superficial temporal artery smooth muscle cells of patients with MMD compared with those in patients with cerebral arteriosclerosis and healthy controls, but, there was no difference in TGF-β1 expression in the superficial temporal artery smooth muscle cells between patients with...
cerebral arteriosclerosis and healthy individuals (58). These aforementioned studies suggested that TGF-β1 promotes angiogenesis and intimal hyperplasia in MMD. The results of the present study revealed that TGF-β1 expression was significantly higher in patients with MMD compared with that in healthy individuals, suggesting an influence on the formation of smoke-like blood vessels. The RNF213 gene is a susceptibility gene for MMD, with rs112735431 particularly relevant to the occurrence and development of MMD (59). In addition to rs112735431, numerous variant sites have been identified, including rs148731719, rs371441113 and rs138130613 (60,61). One possible explanation for the lack of a significant correlation between RNF213 and TGF-β1 is that RNF213 polymorphism in patients with MMD is a gain-of-function mutation (62).

In scientific research, the use of stably transfected cell lines remains to be the most reliable routine method for delivering genes of interest. Although several types of stem cells, such as embryonic stem cells have also been used, their use comes with several limitations, including the length of time required to obtain a sufficient number of stem cells, as well as legal and ethical restrictions surrounding transplantation and gene therapy (63). Therefore, stem cells do not fully meet current experimental requirements. In this regard, BMSCs were a suitable alternative for the present study, with advantages including a lower rejection rate, convenience and no moral or ethical restrictions (64). It has also been reported that BMSCs can promote central nervous system repair (65). However, there is no standard protocol for the isolation, culture and identification of BMSCs. Currently, four primary methods are used to isolate and purify BMSCs, namely the whole bone marrow adherent method, density gradient centrifugation, flow cytometric separation and immunomagnetic beads (66).
to the indeterminate phenotype and susceptibility of BMSCs to damage, the latter of the two methods are rarely used for practical application (67). Although higher-purity BMSCs can be obtained by density gradient centrifugation, this method is complex in practice and delivers the lowest yield of all four methods (68). Whole bone marrow adhesion is easy to conduct and most accurately simulates the natural BMSC environment (69). Therefore, in the present study, the whole bone marrow adherent method was used to isolate rBMSCs. Currently, there are three methods used to identify MSC (70). In the present study, changes in rBMSC morphology were observed under a microscope, whilst surface antigen expression (CD29, CD106, CD34 and CD45) was assessed by flow cytometry. Furthermore, adipogenic and osteogenic differentiation of rBMSCs was assessed by oil red O and Alizarin red staining, respectively. The results from the present study indicated that the whole bone marrow adherent method was successfully used to isolate and culture BMSCs in rats.

In the present study, the RNF213 gene was silenced in rBMSCs using lentiviral vectors, which was confirmed by RT-qPCR. The RNF213 gene encodes a 5,256-amino acid protein containing a zinc finger and an ATPase domain, which exhibit E3 ubiquitin ligase and energy-dependent chaperone activities, respectively (71). Previous studies have shown that RNF213 is associated with the onset and severity of MMD and may therefore be a potential marker for the evaluation of MMD prognosis (72-74). Animal experiments have highlighted that the intima-media is more fragile following the ligation of the common carotid arteries in RNF213-knockdown mice compared with that in normal mice, with altered hemodynamics (75). The pathological changes to the lesion vessels in MMD primarily exhibit an irregular smooth muscle cell shape, vacuoles, in addition to fragile vessels with a reduced wall thickness (3). Smooth muscle cell proliferation is the major cause of lesion vessel thickening and medial thinning in patients with MMD (76). Recent data also suggested that various proinflammatory molecules (such as C-reactive protein, interleukin-6) and angiogenic factors are involved in the pathogenesis of MMD (77,78). Combined with suggestions from the present study on MMD etiology, it was hypothesized that the aberrant expression of VEGF and TGF-β1 following genetic and/or environmental alterations promotes abnormal smooth muscle cell proliferation, resulting in arterial stenosis and occlusion. This ultimately leads to the clinical symptoms of hemorrhage or ischemia. To verify this speculation, the mRNA and protein expression levels of VEGF and TGF-β1 were measured in rBMSCs (79). The authors declared that they have no competing interests.

**References**


64. Park MG, Shin JH, Lee SW, Park HR and Park KP: RNF213 as the major susceptibility factor by 4-hexylresorcinol is mediated by transforming growth factor β1 gene into arteries stimulating angiogenesis. Theranostics 8: 1607-1623, 2018.


