MiR-542-5p regulates the progression of diabetic retinopathy by targeting CARM1

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INTRODUCTION

Diabetic retinopathy (DR), as the most frequent microvascular complication of diabetes mellitus (DM), causes vision loss and blindness in adults worldwide with increasing incidence. MicroRNAs (miRNAs) are involved in the regulation of DR. However, the role of miR-542-5p is still unknown. Here, we demonstrate that miR-542-5p is down-regulated in patients with DR and in high-glucose (HG)-treated retinal pigment epithelial cells. Moreover, miR-542-5p overexpression inhibits apoptosis in retinal pigment epithelial cells exposed to HG. The interaction between miR-542-5p and co-activator-associated arginine methyltransferase 1 (CARM1) is confirmed. MiR-542-5p mimics decrease the CARM1 level and miR-542-5p inhibitor increases the CARM1 level. Additionally, CARM1 overexpression promotes the miR-542-5p-mediated apoptosis in HG-treated retinal pigment epithelial cells. In summary, the data suggest that miR-542-5p may suppress apoptosis in retinal pigment epithelial cells via targeting CARM1, which provides a new therapeutic target for the treatment of patients with DR.

Key words: Diabetic retinopathy, miR-542-5p, CARM1, apoptosis

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Abbreviations: CARM1, co-activator-associated arginine methyltransferase 1; DR, diabetic retinopathy; DM, diabetes mellitus; ECL, enhanced chemiluminescence; HG, high-glucose; miRNAs, microRNAs; NDR, non-diabetic retinopathy; PI, propidium iodide; qRT-PCR, quantitative real-time PCR; UTR, untranslated region

MATERIALS AND METHODS

Clinical samples. A total of 79 patients with DM (34 patients without DR and 45 patients with DR) and 40 healthy volunteers were from Traditional Chinese Medical Hospital of Xinjiang Uygur Autonomous Region. The blood samples from patients were collected and then stored at −80°C until use. In this study, all patients signed informed consent. The experiments conform to the World Medical Association Declaration of Helsinki and were approved by the Ethics Committee of Traditional Chinese Medical Hospital of Xinjiang Uygur Autonomous Region.

Cell culture, treatment and transfection. The human retinal pigment epithelial line ARPE-19 was purchased from ATCC (Manassas, VA, USA), and cultured
in DMEM containing 10% fetal bovine serum at 37°C under 5% CO₂. The cells were treated with 5 mM glucose (Control) or 25 mM glucose (HG) for 24 h. Additionally, the cells were transfected with miR-542-5p mimics or negative control mimics (NC mimics) for 24 h and then exposed to HG for another 24 h.

Quantitative real-time PCR (qRT-PCR). Total RNA was extracted using TRIzol reagent (Takara, Dalian, China). Then, cDNAs were synthesized through Taqman MicroRNA assays (Beyotime, Shanghai, China). The qRT-PCR was performed, and miR-542-5p and CARM1 levels were determined via SYBR Green (Takara, Dalian, China). The relative expression of miR-542-5p and CARM1 was calculated via the 2^−ΔΔCT method.

Western blot analysis. Protein was extracted via RIPA lysis buffer (Beyotime, Shanghai, China). Then, the protein concentration was quantified using BCA Kit (Takara, Dalian, China). Proteins were separated by SDS-PAGE and wet-transferred onto PVDF membrane (Beyotime, Shanghai, China). After blocking in 5% non-fat milk for 1 h, the membrane was incubated with primary antibodies such as CARM1 (1:500), Cleaved caspase 3 (1:500), caspase 3 (1:1000), Bcl-2 (1:1000) and β-actin (1:5000) (all from Abcam, Shanghai, China), at 4°C overnight. β-actin was used as the internal control. Subsequently, the membrane was incubated with the secondary antibody (1:3000, Beyotime, Shanghai, China) for 1 h at 37°C. The bands were visualized via enhanced chemiluminescence (ECL) reaction solution (Takara, Dalian, China).

Cell apoptosis analysis. Cells (3×10^4/ml) were collected and resuspended in 1× Annexin binding buffer (500 μl), and stained with 5 μl Annexin-V-FITC and 5 μl propidium iodide (PI) for 25 min. Then, cell apoptosis was measured via flow cytometry (BD, Japan).

Luciferase reporter assay. TargetScan software (http://www.targetscan.org/vert_72/) predicted that miR-542-5p could target CARM1. The wild-type 3′-UTR sequence of CARM1 containing the miR-542-5p binding site, and the mutant 3′-UTR sequence of CARM1 lacking the miR-542-5p binding site were inserted into the pmirGLO reporter vector (pmirGLO-CARM1-WT and pmirGLO-CARM1-MUT). The ARPE-19 cells were co-transfected with pmirGLO-Report constructs and miR-542-5p mimics or NC mimics using Lipofectamine 2000 (Beyotime, Shanghai, China) for 48 h. The firefly and Renilla luciferase activities were examined through the Dual-Luciferase Reporter Assay System. The Renilla/firefly luciferase ratio was analyzed.

Statistical analysis. Data were presented as mean ± S.D. GraphPad Prism 6.0 was used to carry out statistical analysis. Statistical significance was assessed using Student’s t-test and one-way ANOVA followed by the Bonferroni test. The correlation between the two groups was confirmed through Pearson’s correlation analysis. P<0.05 was considered statistically significant.

RESULTS

The effect of HG treatment on apoptosis of retinal pigment epithelial cells

We first measured the level of miR-542-5p in patients with DR and HG-treated retinal pigment epithelial cells. qRT-PCR showed that the miR-542-5p level is lower in diabetes mellitus (DM)+non-diabetic retinopathy (NDR) group than in the Healthy group, and miR-542-5p is reduced in patients with DM and DR compared with patients with DM. Moreover, the miR-542-5p in DM+NDR group is reduced compared with that in the Healthy group (Fig. 1A). The results from qPCR assay also show that miR-542-5p is reduced in HG-treated ARPE-19 cells compared with control (Fig. 1B).
apoptosis analysis reveals that apoptosis is enhanced in HG-treated ARPE-19 cells (Fig. 1C). Western blot analysis indicates that HG treatment causes the elevation of Cleaved caspase 3 protein level and the reduction of Bcl-2 protein level in ARPE-19 cells (Fig. 1D). These findings imply that miR-542-5p might be related with apoptosis in HG-treated retinal pigment epithelial cells.

**MiR-542-5p inhibits apoptosis in HG-treated retinal pigment epithelial cells**

To explore the effect of miR-542-5p on the apoptosis of HG-treated retinal pigment epithelial cells, we overexpressed miR-542-5p in HG-treated ARPE-19 cells. The qRT-PCR assay demonstrates that HG treatment results in the decrease of miR-542-5p, whereas miR-542-5p mimics significantly elevate miR-542-5p level in HG-treated ARPE-19 cells (Fig. 2A). Annexin-V-FITC/PI staining shows that cell apoptosis is enhanced after HG treatment, whereas miR-542-5p mimics repress cell apoptosis in ARPE-19 cells treated with HG (Fig. 2B). Consistently, western blot analysis reveals that miR-542-5p mimics inhibit the HG-induced increase of Cleaved caspase 3 and HG-induced reduction of Bcl-2 protein level in ARPE-19 cells (Fig. 2C). These data indicate that miR-542-5p suppresses apoptosis of HG-treated retinal pigment epithelial cells.

**CARM1 is verified as a target of miR-542-5p**

We predict that CARM1 is a target of miR-542-5p using Targetscan analysis (Fig. 3A). The interaction be-

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**Figure 2. MiR-542-5p inhibits apoptosis in HG-treated retinal pigment epithelial cells.**

(A) The qRT-PCR was used to measure miR-542-5p mRNA level. (B) Annexin-V-FITC/PI staining was carried out to examine cell apoptosis. (C) Cleaved caspase 3 and Bcl-2 protein levels were measured by western blot analysis. n=3, **P<0.01.

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**Figure 3. CARM1 is verified as a target of miR-542-5p.**

(A) Targetscan analysis predicted that CARM1 was a target of miR-542-5p. (B) The interaction between miR-542-5p and CARM1 was confirmed through luciferase reporter assay. (C) MiR-542-5p mRNA level was measured via qRT-PCR. (D, E) CARM1 mRNA level and protein level were measured using qRT-PCR and western blot analysis, respectively. n=3, **P<0.01.
tween miR-542-5p and CARM1 is identified using the luciferase reporter assay. The results show that the luciferase activity is decreased in ARPE-19 cells co-transfected with miR-542-5p mimics and CARM1 WT 3’-UTR luciferase reporter vector, whereas there is no effect on the luciferase activity of ARPE-19 cells co-transfected with miR-542-5p mimics and CARM1 MUT3’-UTR luciferase reporter vector (Fig. 3B). qRT-PCR results demonstrate that miR-542-5p mimics increase the miR-542-5p level, whereas miR-542-5p inhibitor decreases the miR-542-5p level (Fig. 3C). Western blot analysis and qRT-PCR show that miR-542-5p mimics repress the CARM1 level and miR-542-5p inhibitor increases the CARM1 level (Fig. 3D, E). These results indicate that CARM1 is a target of miR-542-5p and is negatively regulated by miR-542-5p.

CARM1 overexpression attenuates the miR-542-5p-mediated apoptosis in HG-treated retinal pigment epithelial cells

To test whether CARM1 can participate in the regulation of apoptosis induced by miR-542-5p in HG-treated retinal pigment epithelial cells, we performed Annexin-V-FITC/PI staining. The results show that miR-542-5p mimics inhibit apoptosis, and CARM1 overexpression promotes apoptosis in HG treated ARPE-19 cells. Moreover, CARM1 overexpression enhances miR-542-5p-mediated apoptosis in HG-treated ARPE-19 cells (Fig. 4A). Additionally, western blot analysis proves that miR-542-5p mimics lower the CARM1 and Cleaved caspase 3 protein levels, and increase the Bcl-2 protein level, whereas CARM1 overexpression has the opposite effect on these protein levels. Further, CARM1 overexpression rescues these protein levels induced by miR-542-5p mimics in HG-treated ARPE-19 cells (Fig. 4B). The data imply that CARM1 overexpression elevates the miR-542-5p-mediated apoptosis in HG-treated retinal pigment epithelial cells.

DISCUSSION

Here, we demonstrate that miR-542-5p is reduced in patients with DR and in HG-treated retinal pigment epithelial cells. Functional analysis indicated that miR-542-5p represses apoptosis in retinal pigment epithelial cells in the presence of HG. Moreover, the interaction between miR-542-5p and CARM1 was identified. Further, CARM1 overexpression promotes miR-542-5p-mediated apoptosis in retinal pigment epithelial cells exposed to HG. The data imply that miR-542-5p may function in the apoptosis of retinal pigment epithelial cells, which contributes to the treatment of patients with DR.

Increasing evidence has revealed that abnormal expression of miRNAs has a close association with DR. Some miRNAs are upregulated. For example, miR-218 expression is increased, and it represses proliferation as well as stimulates apoptosis in human retinal pigment epithelial cells (Yao et al., 2019). MiR-29b-3p is elevated in patients with DR, and enhances the ratio of Bax/Bcl-2 in human retinal microvascular endothelial cells (Zeng et al., 2019). Conversely, some miRNAs have the opposite role. For instance, miR-7 has been demonstrated to inhibit apop-
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... and alleviates HG-induced damage in retinal pigment epithelial cells (Yang et al., 2019). MiR-142-5p is decreased in retinal tissues of DR rats and HG-treated human retinal endothelial cells, and improves pathological effects in retinal tissues (Liu et al., 2020). Like the later studies, our research verifies that miR-542-5p is suppressed in patients with DR and in HG exposed ARPE-19 cells. Moreover, miR-542-5p represses HG-induced apoptosis in ARPE-19 cells, accompanied by the reduction of the Cleaved caspase 3 protein level and elevation of the Bcl-2 protein level. These findings imply that miR-542-5p may be closely connected with the development of DR.

Previously, miRNAs were found to regulate the development of DR through targeting the 3'-UTR of mRNAs. For example, Fu and Ou (Fu & Ou, 2020) found that miR-152 is reduced and represses HG-induced angiogenesis in human retinal endothelial cells through targeting LIN28B. Chen and others (Chen et al., 2019) discovered that miR-126 is decreased, and stimulates proliferation as well as suppresses apoptosis in HG-induced human retinal endothelial cells via targeting interleukin-17A. Interestingly, our study shows that miR-542-5p could target CARM1. Additionally, miR-542-5p down-regulates the CARM1 expression level. More importantly, accumulating evidence has shown that the function of miR-542-5p may vary in different diseases. For example, miR-542-5p has been found to alleviate fibroblast activation, and suppress proliferation and migration through targeting integrin α6 (Yuan et al., 2018). Conversely, Cheng and others (Cheng et al., 2015) have reported that miR-542-5p stimulates proliferation in osteosarcoma through targeting HUWE1. These findings suggest that miR-542-5p may regulate development of some diseases via targeting the target gene. Further analysis in this study proves that CARM1 overexpression increases the miR-542-5p-mediated apoptosis in retinal pigment epithelial cells treated with HG. Notably, CARM1 was reported to be enhanced in HG-treated retinal pigment epithelial cells, and to promote apoptosis in retinal pigment epithelial cells (Kim et al., 2014). Moreover, Porta and others (Porta et al., 2019) demonstrated that the CARM1 expression level is high in type 2 diabetes. Our study is consistent with previous research. The data suggest that miR-542-5p may take part in the inhibition of apoptosis in retinal pigment epithelial cells through targeting CARM1.

However, other miRNAs may also participate in regulating the progression of DR by targeting CARM1. Moreover, the clinical application of miR-542-5p was not evaluated, and the effect of miR-542-5p on apoptosis in patients with DR by targeting CARM1 was not explored in our study. Therefore, to further investigate the mechanism underlying miR-542-5p in patients with DR, more experiments will be carried out in the future.

In summary, the current study shows that miR-542-5p is down-regulated and CARM1 is up-regulated in HG-treated retinal pigment epithelial cells and in patients with DR. MiR-542-5p participates in the progression of DR via targeting CARM1 (Fig. 5), which could be useful for the treatment of patients with DR.
ments, Qian Bu, Yong Zhao and Lei Yang performed the experiments

REFERENCES


