Silencing protease-activated receptor 2 alleviates ox-LDL-induced lipid accumulation, inflammation, and apoptosis via activation of Wnt/β-catenin signaling

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Abstract. Macrophages conversion to foam cells strongly promoted atherosclerosis progression by plaque formation and plaque rupture. Macrophages swallow oxidized-low density lipoprotein (ox-LDL) to promote foam cell formation. Protease-activated receptor 2 (PAR2) has been reported to take part in atherosclerotic development. However, the effects of PAR2 in macrophages were rarely investigated. In this study, human monocyte, THP-1 was induced to macrophages by using phorbol 12-myristate 13-acetate (PMA). Subsequently, an in vitro model was arranged by using ox-LDL to treat the macrophages. The data showed that inhibition of PAR2 reduced ox-LDL-induced foam cell formation, inflammation, and apoptosis. Additionally, ox-LDL increased PAR2 and inhibited Dickkopf-related protein 1 (DKK1) expression, which is a Wnt signaling inhibitor. PAR2 knocked-down decreased DKK1 and enhanced expression of Wnt3a, β-catenin. Meanwhile, DKK1 overexpression reversed the effects of PAR2 on foam cell formation, inflammation, and apoptosis. In summary, PAR2 is essential for the formation of foam cells, inflammation, and apoptosis in macrophages which plays a critical role during atherosclerosis. PAR2 plays roles in macrophages treated with ox-LDL via DKK1/Wnt/β-catenin signaling.

Key words: Atherosclerosis — Foam cells — Inflammation — Apoptosis — Protease-activated receptor 2

Introduction

Atherosclerosis occurs in arteries with plaque accumulation. It mainly leads to stroke, heart attack or even death. Plaque is formed by fat, cholesterol, calcium and other substances in the blood. The stages of atherosclerosis including intimal thickening, plaque progression and plaque rupture, which is resulted by immune cell infiltration. Furthermore, it is initiated by the formation of foam cells induced by which macrophages engulf low-density lipoprotein cholesterol (LDL) (Tabas et al. 2007). The foam cell’s apoptosis and inflammation accelerate atherosclerosis progression (Wang et al. 2018). In the advancement of atherosclerotic plaques, free cholesterol in foam cells is accumulated to result in endoplasmic reticulum stress, which triggers inflammatory responses and subsequently foam cells apoptosis (Zhou et al. 2005; Cai et al. 2015).

Protease-activated receptors (PAR) mediate cell activation by binding serine proteases which are key activators in the coagulation cascade (Antoniak and Mackman 2014). PAR family (PAR1–4) is expressed omnipresently in vascular cells and is associated with atherosclerosis. PAR1 inhibits cholesterol efflux and promotes atherogenesis (Raghavan et al. 2018). Thrombin-induced platelet activation, which could induce platelet-dependent atherosclerosis, is failing in PAR4 deficient mice (Hamilton et al. 2009). PAR1, PAR3 and PAR4 can be activated by thrombin crucial in hemostasis and thrombosis (Popovic et al. 2010). PAR2 is stimulated by serine proteases including trypsin, trypptase and factor
Xa (FXa) except by thrombin. Our previous investigation showed that PAR2 expression was upregulated in patients with atherosclerosis. PAR2 deficiency in mice attenuated atherosclerosis by reducing macrophage accumulation and collagen deposition (Jones et al. 2018). PAR2 activates M1-like polarization with a significant increase of IL-1β, IL-6, and MCP-1 (Chen et al. 2019). Research demonstrates that FXa activates PAR2 to increase inflammation and lipid uptake (Hara et al. 2018), which is essential for plaque formation and further atherosclerosis.

Oxidized-LDL (ox-LDL) is generated during atherogenesis and is essential for the formation of foam cells (Steinberg 1997). In human nonalcoholic fatty liver disease, it has been found that PAR2 controls cholesterol homeostasis and lipid metabolism (Rana et al. 2019). Therefore, we hypothesized that PAR2 participated in the foam cell formation by regulating the ox-LDL uptake. Furthermore, we evaluated the effects of PAR2 in cell apoptosis and inflammation in macrophages stimulated with ox-LDL.

Materials and Methods

Cell culture

Human THP-1 cells were purchased from the Cell Culture Center of the Chinese Academy of Medical Sciences (Beijing, China). Cells were incubated in Dulbecco’s Modified Eagle Medium (DMEM; Gibco, Thermo Fisher Scientific, USA) with 10% fetal bovine serum (FBS; Gibco, Thermo Fisher Scientific). THP-1 cells were differentiated with 50 nM phorbol 12-myristate 13-acetate (PMA) for 48 h to macrophage. The differentiated cells were used for the following experiments. Cells (75% confluence) were transfected with siRNA-PAR2-1, siRNA-PAR2-2 or control siRNA (siRNA-NC) which were synthesized by Gene Pharma (Shanghai, China) according to the previous investigation (Yeong et al. 2010). The silence sequence is: sense, 5’-CAUCCACAGAAAAGACUGUTT, and anti-sense, 5’-ACAGUCUUUCUGUGGAUGTT. The pcDNA3.0-Flag-DKK1 and empty pcDNA3.0 plasmid (Gene Pharma, Shanghai, China) were co-transfected with siRNA-PAR2 into cells. Macrophages were co-transfected with shRNA-NC, shRNA-PAR2-1 or shRNA-PAR2-2 for 48 h. Further, macrophages were co-transfected with shRNA-NC, shRNA-PAR2-1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h, or macrophages were co-transfected with shRNA-PAR2 and pcDNA3.0-DKK1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h. Other macrophages were treated with ox-LDL (100 μg/ml) or co-transfected with pcDNA3.1-PAR2 or pcDNA3.1 empty plasmid for 48 h. Proteins were obtained from treated THP-1 cells by using RIPA lysis buffer (Thermo Fisher Scientific, USA). Gel electrophoresis was performed to separate proteins. An equal amount of proteins (30 μg) was loaded into the wells of SDS-PAGE (10%) and transferred to PVDF membranes. The membranes were subsequently treated as previously described (Zahid et al. 2019). The primary antibodies were used as follows: PAR2, NLRP3, ASC, caspase-1, Bid, Bcl-2, cleaved-caspase3, caspase-3, cleaved-caspase9, caspase-9, Wnt3a, GSK3β, β-catenin, and DKK1 (Cell Signaling Technology, USA). The secondary HRP-conjugated anti-rabbit antibody (Cell Signaling Technology, USA) was used in this investigation.

RT-qPCR

Macrophages were treated with ox-LDL at a concentration of 25, 50, 100 and 150 μg/ml for 24 h. In next experiments, macrophages were transfected with shRNA-NC, shRNA-PAR2-1 or shRNA-PAR2-2 for 48 h. Further, macrophages were co-transfected with shRNA-NC, shRNA-PAR2-1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h, or macrophages were co-transfected with shRNA-PAR2 and pcDNA3.0-DKK1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h. Other macrophages were treated with ox-LDL (100 μg/ml) or co-transfected with pcDNA3.1-PAR2 or pcDNA3.1 empty plasmid for 48 h. RNA was extracted by TRizol reagent (Thermo Fisher Scientific, USA) in treated cells. The RNA was quantified using NanoDrop ND-2000 (Thermo Fisher Scientific). The relative mRNA expression was determined by real-time PCR using SYBR-Green Real-time PCR Master Mix (Thermo Fisher Scientific). GAPDH was performed as an internal standard. Primers used in this experiment were shown in Table 1. The PCR reaction conditions were as follows: 95°C 10 min for 1 cycle; 95°C 15 s, 60°C 30 s, and 72°C 30 s, for 40 cycles; 95°C 15 s, 60°C 60 s, 95°C 15 s, 60°C 15 s.

Oil Red O staining

Macrophages were transfected with shRNA-NC, shRNA-PAR2-1 for 48 h and treated with ox-LDL (100 μg/ml) for reagent for 3 h at 37°C. The absorbance was detected at a wavelength of 450 nm. Triplicate measurements were repeated.
another 24 h. Further, macrophages co-transfected with shRNA-PAR2 and pcDNA3.0-DKK1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h. The cell culture medium was removed and washed with PBS three times. Cells were fixed with 4% paraformaldehyde for 15 min. Cells were subsequently stained with Oil Red O dissolved in isopropanol and water (3:2) for 20 min. Red-stained lipid droplets in cells were observed by an inverted microscope (Olympus, Japan). Finally, the stained cells were exacted by invert microscopy with ×400 magnification.

**Measurement of total cholesterol and free cholesterol**

Macrophages were transfected with shRNA-NC, shRNA-PAR2-1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h. Further, macrophages co-transfected with shRNA-PAR2 and pcDNA3.0-DKK1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h. The total cholesterol and free cholesterol levels were detected using the commercial kit (E1015, E1016; Applygen Technologies Inc., Beijing, China) following the instructions. Cells were treated with lysis buffer for 10 min and centrifuged at 1500 × g for 10 min. The supernatant was collected and detected at 550 nm.

**Levels of TNF-α, IL-1β, IL-6, and MCP-1 by ELISA kit**

Macrophages were transfected with shRNA-NC, shRNA-PAR2-1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h. Further, macrophages co-transfected with shRNA-PAR2 and pcDNA3.0-DKK1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h. Other macrophages treated with ox-LDL (100 μg/ml) or transfected with pcDNA3.1-PAR2 or pcDNA3.1 empty plasmid for 48 h. After stimulation, the culture medium was collected and centrifuged at 1000 × g for 15 min. The levels of tumor necrosis factor-alpha (TNF-α), interleukin (IL)-1β, IL-6 and monocyte chemoattractant protein-1 (MCP-1) in cell supernatant were assessed by commercial kits (all from Abcam, USA).

| Gene          | Sense                  | Anti-sense             |
|---------------|------------------------|------------------------|
| PAR2 (NM_005242) | GGGTGTGGAATAGACAGGC    | GGGAAAGCATGAACAGAGAG   |
| DKK1 (NM_012242) | AGGTTGTAACCGCATAGGAAAA | TTAGTGCTCTGCAAGTGAGAC  |
| GAPDH (NM_014364) | CAATGACCCCTCCATTGACC   | TTGATTTGGAGGGATCTCG    |

**Table 1. Primers for qRT-PCR of PAR2**

Figure 1. PAR2 expression in ox-LDL-induced THP-1 cells. A. The cell viability in macrophages incubated with different doses of ox-LDL as indicated. The mRNA (B) and protein (C) expression of PAR2 in macrophages treated with different concentrations of ox-LDL.

* p < 0.05, ** p < 0.01, *** p < 0.001 vs. Control group.
Cell apoptosis analysis

Macrophages were transfected with shRNA-NC, shRNA-PAR2-1 for 48 h and treated with ox-LDL (100 μg/ml) for another 24 h. Or macrophages were co-transfected with shRNA-PAR2 and pcDNA3.0-DKK1 for 48 h and treated with ox-LDL for another 24 h. Or macrophages treated with ox-LDL (100 μg/ml) or transfected with pcDNA3.1-PAR2

Figure 2. Effects of PAR2 in ox-LDL-induced macrophages. RT-qPCR (A) and Western blotting (B) were performed to detect transfected efficacy of shRNA-PAR2 in macrophages. * p < 0.05, ** p < 0.01, *** p < 0.001 vs. siRNA-NC group. C. Representative pictures of Oil Red O staining were exhibited in different groups (magnification 400×). D. The levels of total cholesterol and free cholesterol in macrophages treated with ox-LDL were detected by the commercial assay kit. E. The inflammatory mediators, TNF-α, IL-1β, IL-6 and MCP-1, were detected by the ELISA kit. ** p < 0.01, *** p < 0.001 vs. Control group; # p < 0.05, ### p < 0.001 vs. ox-LDL group; ΔΔΔ p < 0.001 vs. ox-LDL+siRNA-NC group.
or pcDNA3.1 empty plasmid for 48 h. Cells were counted with 1×10⁶ cells/ml and resuspended in 1× Binding Buffer. Cell apoptosis was measured by PE Annexin V Apoptosis Detection Kit I (BD Pharmigen) using manual instructions. FlowJo software was performed to analyze apoptotic data.

**Statistical analysis**

Student’s t-test and One-way ANOVA followed with Turkey’s test in GraphPad 6.0 were performed to analyze the statistical differences. A value p < 0.05 was shown as significant difference.

**Results**

**PAR2 knockdown hindered the formation of foam cells, cholesterol accumulation, and inflammation**

According to CCK-8 data, ox-LDL dramatically induced cell death as the concentration increasing (Fig. 1A). The mRNA and protein expression of PAR2 was increased by the treatment of ox-LDL with various doses (Fig. 1B and C). 100 μg/ml ox-LDL was performed to do the following experiment.

To determine the role of PAR2, it is knocked-down in THP-1 cells. Both mRNA and protein levels were downregulated (Fig. 2A and B). siRNA-PAR2-1 was used in the following analysis. Figure 2C showed that ox-LDL treatment dramatically induced lipid accumulation and PAR2 inhibition reversed the effects of ox-LDL. This revealed that PAR2 may participate in the formation of foam cells. Subsequently, total cholesterol and free cholesterol levels were increased significantly in macrophages stimulated with ox-LDL, but PAR2 knocked-down inhibited the expression (Fig. 2D). Besides, ox-LDL dramatically induced the expression of pro-inflammatory cytokines including TNF-α, IL-1β, IL-6, and MCP-1, and inhibition of PAR2 reduced the expression (Fig. 2E). Also, the inflammasome concluding NOD-like receptor 3 (NLRP3), the apoptosis-associated speck-like protein containing CARD (ASC) and caspase-1 was activated by ox-LDL and reversed by PAR2 inhibition (Fig. 3). These data indicate that PAR2 is essential for regulating macrophages conversion to foam cell and inflammation activating.

**Downregulation of PAR2 suppressed ox-LDL-induced apoptosis**

Due to foam cell apoptosis is further induced lipid stack in the blood vessel, we supposed that PAR2 may affect the apoptosis. Data elucidated that ox-LDL dramatically induced apoptosis in macrophages while PAR2 silence reduced apoptosis (Fig. 4A). Later, protein expression data further confirmed the effects of PAR2 that anti-apoptotic Bcl-2 was dramatically increased after inhibition of PAR2 in macrophages stimulated with ox-LDL (Fig. 4B). The pro-apoptotic Bid, cleaved-caspase3, and cleaved-caspase9 were decreased by inhibition of PAR2 in macrophages stimulated with ox-LDL (Fig. 4B).

**Figure 3.** The inflammasome expression in macrophages treated with ox-LDL. Western blotting and protein bands analysis were performed for the determination of the expression of inflammasome complex NLRP3/ASC/caspase-1. ***p < 0.001 vs. Control group; **p < 0.05, ###p < 0.001 vs. ox-LDL group; Δp < 0.05, ΔΔΔp < 0.001 vs. ox-LDL+siRNA-NC group.
Lower levels of PAR2 activated Wnt/β-catenin signaling pathway and inhibited DKK1

As shown in Figure 5, ox-LDL also enhanced the expression of DKK1 in macrophages. DKK1 is an inhibitor for Wnt signaling. Results showed that Wnt3a was prominently annihilated in macrophages treated with ox-LDL. The downstream GSK3β was dramatically enhanced and β-catenin was reduced. However, PAR2 silence inhibited DKK1 expression and activated Wnt/β-catenin signaling.

To investigate whether PAR2 plays roles in macrophages treated with ox-LDL via DKK1/Wnt/GSK3β/β-catenin signaling, we overexpressed DKK1 in PAR2 silenced macrophages using a pcDNA3.0-DKK1 plasmid (Fig. 6A). Results showed that overexpression of DKK1 promoted lipid accumulation and cholesterol levels in macrophages treated with ox-LDL (Fig. 6B and C). Also, overexpression of DKK1 promoted levels of TNF-α, IL-1β, IL-6, and MCP-1 in PAR2 silenced macrophages with ox-LDL treatment (Fig. 6D). The inflammasome key points including NLRP3, ASC, and caspase-1 were also increased by overexpression of DKK1 (Fig. 6E). After then, the enhancement of DKK1 promoted apoptosis (Fig. 7A). Pro-apoptotic protein like Bid, caspase-3 and caspase-9 was activated, yet anti-apoptotic protein Bcl-2 was inhibited (Fig. 7B). These data elucidate that PAR2 participates in ox-LDL induced foam cell formation, inflammation and apoptosis mediated by Wnt/β-catenin signaling pathway.

PAR2 overexpression enhanced the inflammation and apoptosis by regulating DKK1/Wnt/β-catenin signaling

To further confirm that ox-LDL induced apoptosis and inflammation by regulating PAR2 expression, PAR2 was
Roles of PAR2 in ox-LDL-treated macrophages

Roles of PAR2 in ox-LDL-treated macrophages overexpressed in macrophages (Fig. 8A). The overexpression of PAR2 in macrophages promoted the TNF-α, IL-1β, IL-6, and MCP-1 production compared to empty plasmid control (NC), which was similar as the influence of ox-LDL on these pro-inflammatory factors (Fig. 8B). Also, overexpression of PAR2 robustly enhanced the expression of ASC, caspase-1, and NLRP3 (Fig. 8C). The expression levels of pro-apoptotic protein Bid, cleaved-caspase 3 and cleaved-caspase 9 were increased, whereas the anti-apoptotic protein Bcl-2 was decreased both in ox-LDL group and PAR2 overexpressed cells (Fig. 8D). The higher levels of DKK1 both in ox-LDL and PAR2 than in control or NC indicated that ox-LDL enhanced PAR2 positively regulates DKK1 expression (Fig. 8E). Following that, the decrease of Wnt3a and β-catenin, and the increase of GSK3β in ox-LDL group and PAR2 overexpressed group revealed that ox-LDL stimulated PAR2 expression and DKK1/Wnt/β-catenin signaling (Fig. 8E). These data directly indicated that PAR2 regulates DKK1/Wnt/β-catenin signaling to promote inflammation and apoptosis.

Discussion

Atherosclerosis is initiated by the foam cell formation which will lead to inflammation and apoptosis. Atherosclerosis is a lipid-driven inflammatory disease. Under the progression of atherosclerosis, the monocytes adhere to endothelial cells and migrate into the vulnerable area of the artery, and subsequently differentiate into macrophages to promote uptake of ox-LDL. Conversion of macrophages to foam cells is mainly resulted by the increase of the extra cholesterol and decrease of lipid efflux. Researches and our previous investigation showed that PAR2 levels were enhanced in human coronary atherosclerotic lesions (Napoli et al. 2004). Plaque formation involves endothelial dysfunction cells, smooth muscle cells, leukocytes, foam cells, and extracellular matrix. A current study found that PAR2 expression was upregulated in ox-LDL induced in macrophages. The inhibition of PAR2 reduced the conversion of macrophages to foam cells. The dysfunction of cholesterol metabolism is a key point for atherosclerosis progression. Accumulation of cholesterol in macrophages promoted foam cell formation (Fan et al. 2019). The cholesterol efflux is an important part of the regulation of cholesterol homeostasis to avoid cholesterol accumulation and foam cell formation.

Secretion of MCP-1 recruits macrophages and vascular smooth muscle cells migration to intima (Chae et al. 2012). Lipid deposition by macrophages plays a key role in plaque formation during atherosclerotic progression. ox-LDL uptake produces foam cells to release pro-inflammatory cytokines, including IL-1β, TNF-α, and IL-6 (Feng et al. 2019). Our investigation also showed the same results. Moreover,
we found that knock-down of PAR2 reversed the ox-LDL induced inflammation and PAR2 overexpression showed the similar pathological effects as ox-LDL treatment. Caspase-3 was activated in neointimal macrophages after arterial injury which related to apoptosis in lipid-rich plaque (Hutter et al. 2004). In the presence of ox-LDL, macrophages

Figure 6. DKK1 overexpression reversed the effects in PAR2 silenced macrophages with ox-LDL treatment. A. Transfected efficacy of DKK1 in THP-1 cells. B. Oil Red O staining was performed to investigate the effect of DKK1 on lipid accumulation (magnification 400×). C. The levels of total cholesterol and free cholesterol in macrophages were detected by the commercial assay kit. D. ELISA detection of TNF-α, IL-1β, IL-6, and MCP-1 in the supernatant of macrophages. E. The inflammasome expression was detected by Western blotting. * * * p < 0.001 vs. ox-LDL+siRNA-PAR2-DKK1 group.
Roles of PAR2 in ox-LDL-treated macrophages

form to foam cells and further induce macrophage apoptosis which plays role in the formation and accumulation of the lipid core (Hegyi et al. 1996). PAR2 silence dramatically reduced ox-LDL-induced apoptosis and activation of caspase-3 and caspase-9. The Bid is a Bax similar protein, which translocases from the cytosol to mitochondria, leading to apoptosis (Billen et al. 2008), and Bcl-2 inhibits translocation. Inhibition of PAR2 upregulated Bcl-2 expression and downregulated Bid in macrophages treated with ox-LDL. PAR2 overexpression showed the similar biological effects as ox-LDL treatment on apoptosis, suggesting that ox-LDL induced apoptosis by PAR2 upregulation.

DKK1 expression is upregulated in ApoE-/- mice resulting in enlarged and destabilized atherosclerotic lesions (Di et al. 2017). A previous investigation showed that inhibition of DKK1 inhibits the intracellular lipid droplet accumulation compared with control under ox-LDL treatment (Zhang et al. 2015). Also, overexpression of DKK1 in JEG3 cells suppressed the accumulation of lipid and downregulation of peroxisome proliferator-activated receptor δ (Strakovsky and Pan 2012). Our study showed that silenced PAR2 in macrophages treated with ox-LDL suppressed the expression of DKK1. The increasing lipid droplets accumulation and high levels of cholesterol indicated that overexpression of DKK1 promotes the macrophage conversion to foam cells. It has been proved that Wnt pathway activation participates in macrophage motility and lipid internalization (Borreill-Pages et al. 2011). Suppressing Wnt/β-catenin signaling pathway dramatically inhibited the cholesterol efflux in THP-1 macrophage-derived foam cells (Liu et al. 2020). We found that PAR2 overexpression suppressed the expression of DKK1. The DKK1 gene encodes a member of DKK family proteins which bind to the LRP6 co-receptor and inhibits β-catenin-dependent Wnt signaling. Therefore, PAR2 suppressed DKK1 to be able to activate the subsequent signaling Wnt/β-catenin regulating conversion of macrophage to foam cell. Besides, Wnt/β-catenin dramatically reduced activity of GSK-3β that is a regulator of NF-κB regulating inflammation (Schaale et al. 2011). The production of pro-inflammatory factors

Figure 7. DKK1 overexpression increased apoptosis in PAR2 silenced macrophages with ox-LDL treatment. A. Apoptosis was determined by flow cytometry. B. The apoptotic proteins were estimated by Western blotting. ** p < 0.01, *** p < 0.001 vs. ox-LDL+siRNA-PAR2 group.
Figure 8. PAR2 overexpression in macrophages showed the similar biological effects as ox-LDL treatment. A. Transfected efficacy of PAR2 in THP-1 cells. *** p < 0.001 vs. Control group; ### p < 0.001 vs. NC group. B. ELISA detection of TNF-α, IL-1β, IL-6, and MCP-1 in the supernatant of macrophages. C. Western blotting and protein bands analysis were performed for the determination of the expression of inflammasome complex NLRP3/ASC/caspase-1. **D.** The apoptotic proteins were estimated by western blotting. E. Western blotting and protein bands analysis were performed for the determination of the expression of Wnt/β-catenin/GSK3β/DKK1 signaling. *** p < 0.001, ### p < 0.001 vs. Control group; ΔΔΔ p < 0.001 vs. NC group. NC, the macrophages transfected with empty pcDNA3.1 plasmid; PAR2, the macrophages transfected with pcDNA3.1-PAR2 plasmid.
including TNF-α, IL-1β, and IL-6 was also enhanced by overexpression of DKK1. Inhibition of PAR-2 also alleviated the expression of DKK1 and GSK-3β activity; however, DKK1 reversed this effect. Therefore, the effects of PAR-2 in macrophages treated with ox-LDL may mediate by DKK1/Wnt/β-catenin signaling pathway.

In conclusion, our investigation suggests that PAR2 participates in alterations in macrophages stimulated with ox-LDL by suppressing the activity of the Wnt/β-catenin signaling pathway. PAR2 affects the formation of foam cell, inflammation, and apoptosis via DKK1/Wnt/β-catenin pathways. DKK1 enhances lipid accumulation, inflammation and apoptosis in macrophages stimulated with ox-LDL. Previous investigations have found that PAR2 silencing suppresses atherosclerotic progression in mice. Our study provides a more theoretical basis for the future clinical study of PAR2 silencing therapeutic strategy.

Conflict of interests. The authors declare that there are no conflicts of interest.

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