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Sirtuin 1 Prevents Arterial Endothelial Cell Injury Triggered by Oxidized Low-density Lipoprotein via Modulating the TXNIP/ NLRP3 Signaling Pathway

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ABSTRACT The researchers aimed to explore the protective mechanism of sirtuin 1 (Sirt1) against arterial endothelial cell injury triggered by oxidized low-density lipoprotein (Ox-LDL) via modulating the TXNIP/NLRP3 signaling pathway. Lentiviral infection or siRNA transfection was employed to establish human umbilical vein endothelial cells (HUVECs) presenting stably overexpressed Sirt1 or with interfered Sirt1 expression. Compared with control cells, Ox-LDL-treated HUVECs with overexpressed Sirt1 had significantly enhanced proliferative activity and decreased apoptosis rate, while HUVECs with control cells, HUVECs with overexpressed Sirt1 had significantly decreased TXNIP and NLRP3 protein expressions (P<0.05), whereas HUVECs with interfered Sirt1 expression had significantly decreased TXNIP and NLRP3 protein expressions (P<0.05). Adding SR1-37330 significantly elevated the expression level of Sirt1 in cells. In the case of Ox-LDL-induced damage to HUVECs, Sirt1 is capable of safeguarding vascular endothelial cell functions by suppressing the activity of the TXNIP/NLRP3 signaling pathway.