Research Article

Differential Expression of TXNIP Isoforms in the Peripheral Leukocytes of Patients with Acute Myocardial Infarction

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Background. Acute myocardial infarction (AMI) is the most serious type of coronary atherosclerotic heart disease (CAD). The pathological changes are characterized by atherosclerosis. Oxidative stress plays an important role in atherosclerosis. Thioredoxin-interacting protein (TXNIP), an endogenous inhibitor and regulator of thioredoxin, could bind thioredoxin to regulate its expression and antioxidant activity negatively. The NCBI data show that there are two isoforms in TXNIP gene, namely, TXNIP1 and TXNIP2. Our previous studies have shown that TXNIP expression levels in patients with unstable angina pectoris (UAP) were increased compared with controls (CTR). However, no upregulation of TXNIP was detected in AMI patients. Methods. The leucocytes were isolated from peripheral venous blood, and total RNA of the leucocytes was extracted. Then, real-time quantitative PCR was performed. Results. mRNA levels of TXNIP2 in AMI were significantly increased compared with CTR (P < 0.05). However, the expression of TXNIP1 was downregulated in AMI, but the difference was not statistically significant (P > 0.05). Logistic regression analysis showed that TXNIP2 mRNA levels were significantly associated with AMI (OR = 2.207, P < 0.05). Conclusions. The expression of TXNIP2, not TXNIP1, is upregulated in leukocytes of AMI patients, indicating that only TXNIP2 in circulating leucocytes may be involved in the pathogenesis of AMI.

1. Introduction

Acute myocardial infarction (AMI) is the most serious type of coronary atherosclerotic heart disease (CAD), which seriously endangers human health [1]. Despite the identification of numerous molecular mechanisms, understanding of the pathophysiology of this clinical syndrome remains incomplete. The main risk factors include cigarette smoking, diabetes mellitus, hypertension, and hyperlipidemia [2, 3]. The pathological changes of AMI are characterized by atherosclerosis. Oxidative stress plays an important role in atherosclerosis. Thioredoxin-interacting protein (TXNIP), an endogenous inhibitor and regulator of thioredoxin, could bind thioredoxin to regulate its expression and antioxidant activity negatively. The NCBI data show that there are two isoforms in TXNIP gene, namely, TXNIP1 and TXNIP2. Our previous studies have shown that TXNIP expression levels in patients with unstable angina pectoris (UAP) were increased compared with controls (CTR). However, no upregulation of TXNIP was detected in AMI patients. Methods. The leucocytes were isolated from peripheral venous blood, and total RNA of the leucocytes was extracted. Then, real-time quantitative PCR was performed. Results. mRNA levels of TXNIP2 in AMI were significantly increased compared with CTR (P < 0.05). However, the expression of TXNIP1 was downregulated in AMI, but the difference was not statistically significant (P > 0.05). Logistic regression analysis showed that TXNIP2 mRNA levels were significantly associated with AMI (OR = 2.207, P < 0.05). Conclusions. The expression of TXNIP2, not TXNIP1, is upregulated in leukocytes of AMI patients, indicating that only TXNIP2 in circulating leucocytes may be involved in the pathogenesis of AMI.

ROS participates in various biochemical reactions and is an essential form of energy. But, excessive ROS can lead to disease in pathological conditions. Thioredoxin (TRX) is a multifunctional protein with redox activity, which can act as a neutralizing agent by combining ROS, protecting cells from oxidative stress.

Thioredoxin-interacting protein (TXNIP), a 46 kDa protein originally found in HL-60 cells, also known as Vitamin D3 upregulated protein 1 (VDUP1), is an endogenous inhibitor and regulator of TRX [4–6]. TXNIP could bind TRX to negatively regulate its expression and antioxidant activity [7–9]. And TXNIP negatively regulates the expression of JNK, P38, and VCAM1, increases vascular inflammation, and accelerates the process of atherosclerosis [10]. The inducible nature of TXNIP under several stress conditions, including UV light, γ rays, heat shock, and high glucose suggested that TXNIP may play a role in the cellular processes of...
sequences of the two regions are completely different. The coding region A contains 250 bases, and the coding region B contains 85 bases. Except for the last base, the nucleotide primers, and the hollow arrows represent the position of the reverse primers. The two forward primers share the same reverse primer. The coding region A contains 250 bases, and the coding region B contains 85 bases. Except for the last base, the nucleotide sequences of the two regions are completely different. That is to say, there are differences in amino acid length and sequence between the N terminal (83 amino acids) of TXNIP1 and the N terminal (28 amino acids) of TXNIP2.

2. Material and Methods

2.1. Patients and Controls. From February 2017 to October 2017, a total of 87 AMI patients were recruited from the Department of Cardiology, Jining No.1 People’s Hospital, Jining, Shandong, China. All patients were diagnosed with angiography. Ninety age- and sex-matched CTR, without cerebrovascular diseases, were recruited from the Health and Physical Examination Center. All the subjects were coded, and the authors could not identify a single subject. This study was approved by the Human Ethics Committee of Jining No.1 People’s Hospital, and informed consents were obtained.

2.2. Sample Collection. Peripheral venous blood (5 mL) of the AMI and CTR was collected into EDTA-K2 anticoagulant tube after an overnight fast. The leucocytes were isolated with human leukocyte isolation system (LTS-1078, Haoyang Biological, Tianjin, China), according to the manufacturer’s protocol. Total cholesterol (TC), high-density lipoprotein cholesterol (HDLC), low-density lipoprotein cholesterol (LDLC), and triglyceride (TG) were determined with an ADVIA® 2400 automated analyzer (Siemens Healthcare Diagnostics, Erlangen, Germany) in the Laboratory of Experimental Medicine.

2.3. Real-Time Quantitative PCR (qPCR). Total RNA of the leucocytes was isolated using TRIZol reagent (15596026, Invitrogen, USA) following the manufacturer’s protocol. cDNA was reverse-transcribed using 1.5 μL total RNA as template with RevertAid H Minus First Strand cDNA Synthesis Kit (K1622, Thermo Scientific, USA). qPCR was performed to detect the mRNA levels using SYBR® Premix Ex Taq™ (RR420A, Takara, China) by Real-Time PCR System (Applied Biosystems 7500, USA). Amplification primers (TXNIP1-F: AGG GGT ATT GAC ATC CAC CA, β-actin-F: GGA CTT CGA GCG TAC AG) were included in the PCR reaction system. The qPCR reaction was set up in a reaction volume of 20 μL containing 10 μL SYBR Premix, 0.5 μL forward primer (10 μM), 0.5 μL reverse primer (10 μM), 2 μL cDNA, and 7 μL nuclease-free water. All samples were performed in triplicate. The PCR operation program was set according to the manufacturer’s protocol. The relative expression was calculated with the following equation: relative expression = 2-ΔΔCt [20]. β-Actin is the reference gene for normalization.

2.4. Statistical Analysis. Data were expressed as means ± standard deviation (X ± SD). Comparisons of means between groups were made by Student’s t-test and comparisons of prevalence by chi-square test. All statistical analysis were
performed with SPSS v22.0. *P < 0.05 was considered statistically significant.

### 3. Results

#### 3.1. Clinical Characteristics

The clinical characteristics are summarized in Table 1. Patients with AMI have a significantly higher prevalence of smoking and a lower HDL-C level compared with CTR. The calculation method of expression level is as follows: first, we calculate the $\Delta Ct (\Delta Ct = Ct_{TXNIP1/2} - Ct_{\beta-actin})$ of all samples and the mean value of $\Delta Ct$ of the CTR group ($\Delta Ct_{CTR}$). Then, the $\Delta Ct$ value is calculated as follows: $\Delta \Delta Ct = \Delta Ct - \Delta Ct_{CTR}$. Finally, the expression of the sample was calculated according to relative expression $= 2^{-\Delta \Delta Ct}$. qPCR results showed that the mRNA expression levels of TXNIP1 and TXNIP2 were inconsistent between CTR and AMI. Compared with CTR, the expression of TXNIP1 was downregulated in AMI, but the difference was not statistically significant. The expression of TXNIP2 was increased, and the difference was statistically significant. *$P < 0.05$.

#### 3.2. TXNIP Isoforms’ mRNA Expression Levels

Our previous studies have shown that TXNIP gene expression levels in patients with UAP were significantly increased compared with CTR but AMI not. So, we detected the mRNA levels of the two variants of TXNIP gene, TXNIP1 and TXNIP2, by qPCR. As shown in Figure 2, mRNA levels of TXNIP2 in AMI patients were significantly increased compared with CTR ($P < 0.05$). However, the mRNA levels of TXNIP1 were completely different from that of TXNIP2. The expression of TXNIP1 was downregulated in AMI, but the difference was not statistically significant ($P > 0.05$).

Further multivariate logistic regression analysis between AMI and TXNIP1 mRNA levels, TXNIP2 mRNA levels, smoking, hypertension, and diabetes was carried out. And the results showed that only TXNIP2 mRNA levels were positively associated with AMI (OR = 2.207, *P < 0.05*). It is important to point out that the mRNA expression levels were redefined in the multivariate logistic regression analysis [21]. The mRNA relative expression “<1.00” was defined as the low expression, and “>1.00” was defined as high expression.

### 4. Discussion

Multiple risk factors lead to AMI, such as smoking, diabetes mellitus, hypertension, hyperlipidemia, inflammatory response, oxidative stress, apoptosis, vascular remodeling, plaque stress, and blood flow shear stress [2, 3, 22]. Our previous studies have shown that the TXNIP gene is upregulated in leukocytes in patients with UAP, but this phenomenon does not exist in AMI patients [19]. The symptoms of UAP are diverse. When the disease just occurs, the symptoms are mild. Most of our subjects had a history of primary hospital visits and began to use certain drugs. Studies have shown that these drugs have an impact on the expression of TRX. So, our previous studies still need to be further validated. Considering the diagnostic criteria and other factors, we only carried out the expression of TXNIP isoforms in AMI.

TXNIP gene is localized on chromosome 1 (1q21.1), with a total of 8 exons (Figure 1). There are two isoforms, TXNIP1 (391 amino acids and 2983 bp mRNA, NCBI data) and TXNIP2 (336 amino acids and 2614 bp mRNA,
only TXNIP2 was detected. Their expression levels showed that the sequence region of the two isoforms (Figure 1), and then detect whether there are different isoforms in the current reagent companies, and we cannot point out that there are no differences in subsequent studies. Moreover, it is necessary to different isoforms, and we will continue to explore their functional differences.

In this study, we designed primers for different isoforms, and we tested these two isoforms, but we had detected only one isoform. In addition, it is necessary to verify whether the two isoforms are completely the same protein after posttranslational modification.

Additionally, there are some defects or limitations in the present study. Firstly, the sample size of this study was relatively small, which could lead to some statistical errors. Secondly, on the choice of the control group, the control population was not validated by angiography, which resulted in asymptomatic coronary heart disease likely to be included in CTR. Thirdly, this study lacks protein level detection, which reduces reliability to some extent. We will complement this study in subsequent studies. Finally, compared to AMI patients, all CTR are negative for risk factors, including smoke, diabetes, and hypertension. Perhaps, the TXNIP2 differences between AMI and CTR are due to risk factors and/or AMI. In our previous study [19], these factors were basically the same, and the difference in the expression level of the TXNIP gene was very close to the results of the TXNIP1 in

Figure 3: Amino acid sequence similarity of the two isoforms among species. The similarity of amino acid sequence of isoform 1 is 94.46% and that of isoform 2 is 74.75%. Black shades represent the same amino acids, while other shades are different amino acids. Isoform 1 refers to TXNIP1 or Txnip1, and isoform 2 refers to TXNIP2 or Txnip2. All capital letters represent genes of Homo sapiens, and the capitalization of the initials is the genes of Mus musculus.

NCBI data), which are evolutionally conserved (Figure 3). The difference between the two proteins is mainly at the N-terminus, and the differential expression of TXNIP2 was in AMI and CTR, indicating that TXNIP1 and TXNIP2 may play different roles in the development of AMI. It is also possible that TXNIP1 and TXNIP2 participate in completely different physiological processes.

At present, there are no reports about these two isoforms, and we will continue to explore their functional differences in subsequent studies. Moreover, it is necessary to point out that there are no specific antibodies to these two isoforms in the current reagent companies, and we cannot detect whether there are differences in the protein levels. In theory, the antibody used in our previous studies could detect these two isoforms, but we had detected only one band. In combination with the trend of the expression level of the two variants in this study, we speculate that the expression of the two isoforms may vary greatly; the expression of TXNIP2 is very low and has not been detected in previous studies. In addition, it is necessary to verify whether the two isoforms are completely the same protein after posttranslational modification.
this study. Therefore, we think that the effects of these risk factors can be ignored, but it still needs further verification.

In conclusion, we found that the mRNA level of TXNIP2, but not TXNIP1, was significantly upregulated in AMI. Further analysis will be continued to verify its role in AMI. These data would shed light on our understandings of the AMI pathogenesis and development of better diagnostic markers.

5. Conclusions

The expression of TXNIP2, not TXNIP1, increased in peripheral blood leukocytes of AMI patients, indicating that only TXNIP2 in circulating leukocytes may be involved in the pathogenesis of AMI.

Data Availability

All data arising from this study are contained within the article.

Conflicts of Interest

All the authors declare no conflict of interest.

Acknowledgments

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References

[1] Centers for Disease Control and Prevention, "Prevalence of coronary heart disease—United States, 2006–2010," MMWR. Morbidity and Mortality Weekly Report, vol. 60, no. 40, pp. 1377–1381, 2011.
[2] A. Cubukcu, I. Murray, and S. Anderson, "What's the risk? Assessment of patients with stable chest pain," Echo Research and Practice, vol. 2, no. 2, pp. 41–48, 2015.
[3] Z. Li, J. Cheng, L. Wang, P. Yan, X. Liu, and D. Zhao, "Analysis of high risk factors and characteristics of coronary artery in premenopausal women with coronary artery disease," International Journal of Clinical and Experimental Medicine, vol. 8, no. 9, pp. 16488–16495, 2015.
[4] I. N. Mohamed, S. S. Hafez, A. Fairaqa, A. Ergul, J. D. Imig, and A. B. el-Remessy, "Thioredoxin-interacting protein is required for endothelial NLRP3 inflammasome activation and cell death in a rat model of high-fat diet," Diabetologia, vol. 57, no. 2, pp. 413–423, 2014.
[5] T. S. Devi, K.-I. Hosoya, T. Terasaki, and L. P. Singh, "Critical role of TXNIP in oxidative stress, DNA damage and retinal pericyte apoptosis under high glucose: implications for diabetic retinopathy," Experimental Cell Research, vol. 319, no. 7, pp. 1001–1012, 2013.
[6] K. S. Chen and H. F. DeLuca, "Isolation and characterization of a novel cDNA from HL-60 cells treated with 1,25-dihydroxyvitamin D-3," Biochimica et Biophysica Acta, vol. 1219, no. 1, pp. 26–32, 1994.
[7] B. F. Wang and J. Yoshioka, "The emerging role of thioredoxin-interacting protein in myocardial ischemia/reperfusion injury," Journal of Cardiovascular Pharmacology and Therapeutics, vol. 22, no. 3, pp. 219–229, 2017.
[8] O. N. Spindel, R. M. Burke, C. Yan, and B. C. Berk, "Thioredoxin-interacting protein is a biomechanical regulator of Src activity: key role in endothelial cell stress fiber formation," Circulation Research, vol. 114, no. 7, pp. 1125–1132, 2014.
[9] L. P. Singh and L. Perrone, "Thioredoxin interacting protein (TXNIP) and pathogenesis of diabetic retinopathy," Journal of Clinical & Experimental Ophthalmology, vol. 4, no. 4, p. 287, 2013.
[10] J. Yoshioka and R. T. Lee, "Thioredoxin-interacting protein and myocardial mitochondrial function in ischemia-reperfusion injury," Trends in Cardiovascular Medicine, vol. 24, no. 2, pp. 75–80, 2014.
[11] N. M. Alhawiti, S. Al Mahri, M. A. Aziz, S. S. Malik, and S. Mohammad, "TXNIP in metabolic regulation: physiological role and therapeutic outlook," Current Drug Targets, vol. 18, no. 9, pp. 1095–1103, 2017.
[12] Y. Yu, K. Xing, R. Badamas, C. A. Kusznyski, H. Wu, and M. F. Lou, "Overexpression of thioredoxin-binding protein 2 increases oxidation sensitivity and apoptosis in human lens epithelial cells," Free Radical Biology and Medicine, vol. 57, pp. 92–104, 2013.
[13] E. Junn, S. H. Han, J. Y. Im et al., "Vitamin D3 up-regulated protein 1 mediates oxidative stress via suppressing the thioredoxin function," Journal of Immunology, vol. 164, no. 12, pp. 6287–6295, 2000.
[14] H. Mogami, S. Yura, E. Kondoh, H. Masutani, J. Yodoi, and I. Konishi, "Differential expression of thioredoxin binding protein-2/Txnip in human placenta: possible involvement of hypoxia in its suppression during early pregnancy," The Journal of Obstetrics and Gynaecology Research, vol. 43, no. 1, pp. 50–56, 2017.
[15] K. P. Xia, H. M. Ca, and C. Z. Shao, "Protective effect of notoginsenoside R1 in a rat model of myocardial ischemia/reperfusion injury by regulation of vitamin D3 upregulated protein 1/NF-xB pathway," Die Pharmazie - An International Journal of Pharmaceutical Sciences, vol. 70, no. 11, pp. 740–744, 2015.
[16] H. J. Zhu, D. G. Wang, J. Yan, and J. Xu, "Up-regulation of microRNA-135a protects against myocardial ischemia/reperfusion injury by decreasing TXNIP expression in diabetic mice," American Journal of Translational Research, vol. 7, no. 12, pp. 2661–2671, 2015.
[17] V. Frodermann and M. Nahrindentorf, "Neutrophil-macrophage cross-talk in acute myocardial infarction," European Heart Journal, vol. 38, no. 3, pp. 198–200, 2017.
[18] Y. Ma, A. Yabluchanskiy, R. P. Iyer et al., "Temporal neutrophil polarization following myocardial infarction," Cardiovascular Research, vol. 110, no. 1, pp. 51–61, 2016.
[19] Y. Zhang, J. Huang, X. Yang et al., "Altered expression of TXNIP in the peripheral leukocytes of patients with coronary atherosclerotic heart disease," Medicine, vol. 96, no. 49, p. e9108, 2017.
[20] X. Liao, Y. Chen, D. Liu, F. Li, X. Li, and W. Jia, "High expression of LAMP3 is a novel biomarker of poor prognosis in patients with esophageal squamous cell carcinoma," International Journal of Molecular Sciences, vol. 16, no. 8, pp. 17655–17667, 2015.
[21] L. Li, W. Wang, R. Zhang et al., “High expression of LAMP2 predicts poor prognosis in patients with esophageal squamous cell carcinoma,” Cancer Biomarkers, vol. 19, no. 3, pp. 305–311, 2017.

[22] Task Force Members, G. Montalescot, U. Sechtem et al., “2013 ESC guidelines on the management of stable coronary artery disease: the task force on the management of stable coronary artery disease of the European Society of Cardiology,” European Heart Journal, vol. 34, no. 38, pp. 2949–3003, 2013.