Correlation analysis of ADAMTS-4, VCAM-1, and TAK1 expression in cartilage tissue from osteoarthritis patients.

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Abstract

Objective: Osteoarthritis (OA) is a chronic progressive lesions caused by multiple factors. Previous finding showed that the expressions of ADAMTS-4, VCAM-1, and TAK1 were increased in the cartilage tissue from OA compared with normal tissue. ADAMTS-4 was positively correlated with TAK1 level. VCAM-1 expression was related to severity of OA. It was found that ADAMTS-4 and VCAM-1 can influence the dynamic balance of ECM, while TAK1 may participate in OA through MAPKs and NF-κB signaling pathways. However, the correlation of their expressions in cartilage tissue from patients with OA is still unclear. This study intends to analyse their correlation in OA cartilage through the detection of expressions in OA and normal cartilage.

Patients and methods: A total of 72 OA patients were enrolled and divided into two groups: severe OA and mild-to-moderate OA groups. Another 22 healthy individuals were selected as normal control. Western blot was applied to detect ADAMTS-4, VCAM-1, and TAK1 expression in cartilage tissue. Their correlations were then analysed.

Results: There were 35 patients in severe group with mean age at 58.64 ± 4.32 y old, while 37 subjects in mild-to-moderate group with average age at 57.32 ± 5.21 y old, the ages of which were no significant different. The levels of ADAMTS-4, VCAM-1, and TAK1 in severe group were obviously higher than that in mild-to-moderate group and normal control. The levels of these three genes presented positive correlation with each other.

Conclusions: The expressions of ADAMTS-4, VCAM-1, and TAK1 were elevated and exhibited positive correlation in cartilage tissues of patients with OA, which provides academic basis for the future biomarkers in the therapy of OA.

Keywords: Osteoarthritis, ADAMTS-4, TAK1, Cartilage tissue, VCAM-1.

Introduction

With the growth of aging population in China, the incidence of Osteoarthritis (OA) keeps on increasing, which seriously influence the patient's psychological health, physical health, and normal life [1]. As a type of chronic progressive joint cartilage degenerative lesions, OA occurrence is accompanied by a series of physiological and pathological processes including inflammation and immune reaction. It is mainly featured as cartilage Extracellular Matrix (ECM) degradation and articular cartilage loss in clinic, of which the dynamic imbalance of bone ECM metabolism is the major cause [2].

It was showed that a disintegrin and metalloproteinase with a thrombospondin (ADAMTS) can promote the degradation of ECM and participate in the maintenance of ECM dynamic balance. However, the aberrant expression of ADAMTS may directly induce imbalance of ECM. Moreover, it was found that ADAMTS-4 was overexpressed in OA compared with normal people [3-6].

Vascular Cell Adhesion Molecule 1 (VCAM-1) is a kind of important cell surface glycoprotein and is characterized as regulating cell adhesion between cells. It was revealed that VCAM-1 level was significantly higher in serum and cartilage tissue of patients with OA than that in normal ones. Meanwhile, VCAM-1 expression showed certain correlation with the severity of OA and could be treated as one of the predictors of the onset risk of OA [7-9].

Accumulative studies demonstrated that as one of the member of Mitogen-Activated Protein Kinase (MAPKs) family, transforming growth factor β activated kinase 1 (TAK1) was not only related to the immune and inflammation, but also involved in regulation of cell proliferation, differentiation, and apoptosis. Furthermore, it participates in the regulation of a
variety of signaling pathways, such as MAPKs and NF-κB. Experimental data demonstrated that TAK1 was highly expressed in the cartilage tissue of OA patients, thus related to the course of OA [10-13].

As the expressions of ADAMTS-4, VCAM-1 and TAK1 showed certain association with OA occurrence and development, this study aimed to clarify the interaction and correlation among the three proteins by detecting their expressions in the cartilage tissue from patients with OA, thus to provide effective biomarkers for the early diagnosis of OA.

Materials and Methods

Materials

Goat anti-human monoclonal antibodies for TAK1, ADAMTS-4 and VCAM-1 were bought from Xinlebio. Mouse anti-goat secondary antibody was purchased from Santa Cruz. Goat anti-human monoclonal antibodies for TAK1, ADAMTS-4 and VCAM-1 were bought from Xinlebio. Mouse anti-goat secondary antibody was purchased from Santa Cruz. Anti-goat secondary antibody was purchased from Santa Cruz. The automatic exposure meter was from ImageQuant LAS 500 integration imaging.

Clinical information

A total of 72 OA patients in Linyi People's Hospital from October 2015 to October 2016 were enrolled. Exclusion criteria: patients with tumor, acute or chronic arthritis, tuberculosis, diabetes, autoimmune diseases such as rheumatoid arthritis and ankylosing spondylitis, obvious narrowed knee joint space, significant joint changes such as advanced joint osseous stiffness and intra-articular fracture, in pregnancy and lactation, allergic constitution, or other reactive arthritis, etc. Another 22 patients with normal cartilage tissue were also enrolled. The cartilage tissue in normal group presents moderate chondrocytes number with normal cell structure under the microscope. The cartilage tissue was isolated and kept on ice. The tissue with size of around 1 mm³ was moved to 1.5 ml EP tube every 0.1 g tissue and stored at -80°C. The stored sample was rewarmed on ice for 10 min and then treated by the mixture of 490 μl lysis and 10 μl protease inhibitor for 30 min. Next, the tissue was grinded and centrifuged at 10 min at low temperature. The supernatant was moved to a new EP tube and boiled for 15 min together with 50 μl 10X loading buffer.

Cartilage tissue isolation and extraction

The cartilage tissue was isolated and kept on ice. The tissue with size of around 1 mm³ was moved to 1.5 ml EP tube every 0.1 g tissue and stored at -80°C. The stored sample was rewarmed on ice for 10 min and then treated by the mixture of 490 μl lysis and 10 μl protease inhibitor for 30 min. Next, the tissue was grinded and centrifuged at 10 min at low temperature. The supernatant was moved to a new EP tube and boiled for 15 min together with 50 μl 10X loading buffer.

HE staining

The slice was dewaxed by xylene for 20 min and then gradient dehydrated by ethanol (100%, 95%, 85% and 70%). After being washed by ddH₂O, the slice was stained by hematoxylin for 5 min. Next, the slice was treated by 1% hydrochloric-alcohol solution for 15 s. After stained by eosin for 5 min, the slice was observed under the microscope to analyse the cartilage tissue morphology.

Western blot

A total of 50 μg protein was separated by 8% SDS-PAGE at 60 V for 45 min followed by 125 V for 1.5-2 h. Then the protein was transferred to membrane at 35 V for 40 min using the semidry method. Next, the membrane was incubated in primary antibody at 1:100 at 4°C overnight and secondary antibody at 1:1000 at room temperature. The membrane was treated by ECL and detected by Bio-Rad automatic detector. Protein expression was normalized to β-actin and the expressional folds were determined.

Statistical analysis

All statistical analysis was performed using SPSS 13.0 software. The data was depicted as mean ± standard deviation and analysed by Spearman rank correlation analysis. A statistical difference was presented as P<0.05.

Results

Morphologic observation and scoring of OA cartilage tissue

The cartilage tissue was stained by HE for microscopic observation according to Mankin criteria. As shown in Figure 1, normal articular cartilage exhibited smooth surface, more cell number in alignment, uniform color, clear and visible cartilage cell nucleus, and pink cartilage matrix. Comparatively, the cartilage in OA patients presented coarse surface, collagen fibrosis and crack formation in shallow cartilage matrix layer, degenerate and hypertrophic chondrocytes in fascicular alignment, and irregular cartilage matrix staining. Additionally, uneven surface, decreasing cell number, irregular shape, interstitial fibrosis, and crack phenomenon were observed in mild-to-moderate group. The disappearance of cartilage surface and extreme reduction of cell number were found in severe group.
Protein expression in cartilage tissue

Western blot demonstrated that the expressions of ADAMTS-4, VCAM-1 and TAK1 protein in cartilage tissue from OA patients were statistically higher than that in normal control (P<0.05, Figure 2). Moreover, their levels in severe group were significantly higher than that in mild-to-moderate group (P<0.05). It suggested that TAK1, ADAMTS-4, and VCAM-1 changes in cartilage tissue of OA were positively related to OA process.

Correlation analysis of ADAMTS-4, VCAM-1 and TAK1 expression

Pairwise comparison was performed to analyse the correlation of ADAMTS-4, VCAM-1, and TAK1 expressions in cartilage tissue. Our data showed positive correlation among the levels of these three genes in OA patients. The levels of ADAMTS-4, VCAM-1, and TAK1 exhibited significant difference between mild-to-moderate group and normal control (P=0.032, P=0.041, P=0.026, respectively), severe group and normal control (P=0.014, P=0.0092, P=0.0054, respectively), and severe group and mild-to-moderate group (P=0.043, P=0.045, P=0.036, respectively).

Discussion

OA is a kind of progressive disease, the causes of which are complicated. It was reported that the environmental factors and aberrant expressions of genes were both directly involved in the occurrence and development of OA. Hysteretic treatment due to insensitivity of early diagnosis leads to the degeneration of articular cartilage and even cartilage loss, which needs artificial joint replacement and causes serious harm to the patients. At present, there is still lack of effective prediction, diagnosis for the timely treatment of OA. Thus, it is of great significance to determine novel biomarkers for the diagnosis, prevention and control of OA [1-5].

In clinic, it was confirmed that cartilage degeneration was the main pathological features of OA, while ECM homeostatic imbalance was an important reason. ADAMTS-4 shows potent effect to degrade aggrecan and its elevation can induce ECM imbalance, resulting in cartilage degeneration and OA. Therefore the upregulation of ADAMTS-4 was proposed to be treated as a predictor for OA [14,15]. Of note, as an important cell adhesion factor, VCAM-1 is involved in regulating cell-cell interaction and cell surface glycoprotein. The elevation of VCAM-1 may inhibit local tissue degradation [16,17]. TAK1 plays a critical role in MAPKs and NF-κB signaling pathways. It was revealed to participate in the pathological process of OA. Similarly, our clinical data demonstrated that the expressions of ADAMTS-4, VCAM-1, TAK1 in cartilage tissue from OA patients were altered, which were consistent with the previous findings [18-20].

Intriguingly, the levels of ADAMTS-4, VCAM-1 and TAK1 from patients were gradually enhanced as the OA aggravated. Correlation analysis demonstrated that ADAMTS-4, VCAM-1, and TAK1 expressions had significant correlation in both severe group and mild-to-moderate groups, indicating that the expressions of ADAMTS-4, VCAM-1 and TAK1 could be used as predictor for the OA diagnosis and evaluation of OA severity.

Conclusion

The expressions of ADAMTS-4, VCAM-1 and TAK1 were increased in cartilage tissue of patients with OA and exhibited positive correlation, which could be further used as indicators in the prediction of occurrence and development of OA.

Disclosure of Conflict of Interest

None

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References


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