Original Article

Association study between growth differentiation factor 5 polymorphism and non-contact anterior cruciate ligament rupture in Chinese Han population

Biao Chen, Bin Li, Yong-Jian Qi, Kai Tie, Liao-Bin Chen

Department of Orthopaedic Surgery, Zhongnan Hospital, Wuhan University, Wuhan, China

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Abstract: Background: Anterior cruciate ligament (ACL) rupture is the most common ligamentous injury for active adolescents and young adults each year. However, the precise etiologies of ACL injury are not fully understood. The present study was to investigate +104T/C polymorphism of growth differentiation factor 5 (GDF5) gene in patients with ACL rupture, and evaluate the effects of polymorphism on GDF5 mRNA levels in ligament of patients with ACL rupture in central China. Methods: A total of 286 Chinese patients with ACL rupture and 500 healthy controls were enrolled in this study. The +104T/C polymorphism in GDF5 gene were genotyped by DNA sequencing. GDF5 mRNA expressions levels in ligament were determined by quantitative PCR. Results: The frequency of the TT genotype tended to be higher in ACL rupture group than in control group (62.6% vs. 48.0%, P<0.001, OR = 1.81, 95% CI: 1.35-2.44). T allele of the GDF5 +104T/C polymorphism was more common in ACL rupture group than in control group (P<0.001). Patients carrying TT genotype expressed lower levels of GDF5 mRNA than C carriers (P = 0.005) among ACL rupture. Conclusion: Our study indicated that GDF5 +104T/C polymorphism was associated with ACL rupture patients in central China. This is likely from decreased expressions of GDF5 mRNA. Further studies are necessary to explore the functional implication of the GDF5 +104T/C polymorphism in Chinese ACL rupture patients.

Keywords: Anterior cruciate ligament, rupture, growth differentiation factor 5, polymorphism, expression

Introduction

The anterior cruciate ligament (ACL) rupture is one of the most common knee injuries in sports and often results in joint effusion, muscle weakness, altered movement, and reduced functional performance [1, 2]. Individuals who participate in these high risk sports, such as football, basketball, and skiing, are more likely to suffer from ACL rupture [3]. ACL rupture occurs mainly during non-contact events, and the long-term consequence of ACL injury may lead to the development of knee osteoarthritis [4]. Due to the outcome and high incidence of ACL rupture, it has always been an intense area of focus in the field of sports medicine [5, 6].

Although intrinsic and extrinsic factors for ACL rupture have been identified, the exact etiology of this injury is not yet fully understood [7]. Many researchers have suggested that genetic elements should be considered as an intrinsic risk factor for ACL rupture [8]. Consistent with these reports, more and more studies have shown that single nucleotide polymorphisms (SNP) within the COL1A1, COL5A1, COL12A1 and MMP3 genes are associated with an increased risk of ACL rupture [8-12].

Growth differentiation factor 5 (GDF5), also known as cartilage-derived morphogenetic protein 1 (CDMP1) or bone morphogenetic protein 14 (BMP14), belongs to the transforming growth factor -beta (TGF-β) superfamily [13]. It has been extensively studied in literatures, and found to be involved in musculoskeletal processes, affecting endochondral ossification, synovial joint formation, tendon and ligament repair, and bone formation [13-17]. Especially, GDF5 is essential for the regulation of ligament homeostasis. In rabbits ACL fibroblasts, a higher level of GDF5 enhanced the expression of type I collagen (the predominant protein in ligaments), changed the stress fiber formation and cellular adhesion by modulating the distribution of integrin alpha2 [18]. Brachypodism mice exhibited developmental failure of the condyles
GDF5 and anterior cruciate ligament rupture

Several SNPs in GDF5 gene have been identified to be associated with sports injury related diseases such as osteoarthritis (OA), anterior cruciate ligament (ACL) and tendon rupture, and meniscus injury. However, some of these associations have not been confirmed, and there are even some contradictory data in different populations [19-25]. Interesting, GDF5 +104T/C polymorphism identified to be associated with OA in a meta-analysis of European and Asian cohorts was associated with ACL rupture and was a functional polymorphism [26]. The +104T/C polymorphism could alter the expression levels of GDF5 within a wide range of connective tissues [21, 27, 28]. Andrew et al. found that GDF5 +104T/C polymorphism was predicted to affect transcription factor binding and it might therefore highlight a regulatory site that could be exploited to manipulate GDF5 expression and alleviate the detrimental effect mediated by the T-allele of rs143383 by using reporter constructs and electrophoretic mobility shift assays [29]. Although the association studies mentioned above suggested a role for the rs143383 variant and the predisposition to various phenotypes, the specific biological role of GDF5 in ligament is still not fully understood [25].

To our knowledge, there have been few studies investigating the relationship between the functional SNP in GDF5 and ACL rupture. The purpose of this study is to investigate the possible association role of SNP rs143383 in GDF5 gene and ACL rupture in Chinese Han population.

**Methods**

**Subjects**

A total of 286 Chinese patients of Han ethnicity, surgically diagnosed as primary noncontact and intra-articular ligament of the knee joints [14].

Table 1. Baseline characteristics of the patients with ACL rupture and healthy controls

<table>
<thead>
<tr>
<th>Variable</th>
<th>ACL rupture</th>
<th>Healthy Controls</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n = 286 (%)</td>
<td>n = 500 (%)</td>
</tr>
<tr>
<td>Female</td>
<td>103 (36%)</td>
<td>170 (34%)</td>
</tr>
<tr>
<td>Age (yrs) [Mean ± SD]</td>
<td>28.7 ± 6.6</td>
<td>26.2 ± 8.1</td>
</tr>
<tr>
<td>BMI (kg/m²) [Mean ± SD]</td>
<td>23.6 ± 5.4</td>
<td>22.9 ± 4.8</td>
</tr>
</tbody>
</table>

Note: BMI, body mass index.

A total of 500 healthy controls were selected from the medical staff of Zhongnan Hospital, Wuhan University, as well as from healthy volunteers in the same geographic area of Wuhan city. The controls were ethnically matched and had no history of ligament or tendon injury. Informed consent was taken from each study subject. The study protocol was approved by the ethics committees of Zhongnan Hospital of Wuhan University.

**DNA extraction and genotyping**

A blood sample was obtained from each case and control for DNA extraction. Genomic DNA was isolated from 5 ml EDTA anticoagulated venous blood according to the standard protocol of the QiAamp DNA Blood Midi kit (Qiagen, Hilden, Germany). The quality of DNA samples was checked by measuring the A260/280 ratio using a fluorescence spectrophotometer F-4500 (Hitachi, Tokyo, Japan). The ratio fell between 1.8 and 2.0 for all samples.

The +104T/C polymorphism (rs143383) in the GDF5 gene was genotyped and determined by DNA sequencing analysis. The region containing +104T/C polymorphism site was amplified using a single primer set: forward 5'-CAG CAT TAC GCC ATT CTT CC-3' and reverse 5'-CGC TGA ATG ACA CCA AAG AGA-3' [30]. The PCR was performed as follows: initial denaturing at 95°C for 5 min, followed by 35 cycles of 95°C for 30 s, 57°C for 30 s, and 72°C for 45 s, and final extension at 72°C for 5 min. The PCR products were purified using QiAquick PCR Purification kit (Qiagen, Hilden, Germany) according to the manufacturer’s instructions, and the purified PCR products were sequenced using an ABI BigDye Terminators v3.1 Cycle Sequencing Kit (Applied Biosystems, Foster City, CA, USA) and run on an ABI 3100 automatic sequencer (Applied Biosystems, Foster City, CA, USA).

For quality control purposes, negative and positive controls were processed with each batch of samples and all experiments were repeated twice to ensure consistency.
Quantification of GDF5 mRNA

Ligament tissue was taken from 62 patients with ACL rupture undergoing arthroscopic-assisted ACL reconstruction after written informed consent. Total RNA was extracted from fresh ligament tissue of ACL rupture patients, with Trizol reagent according to the manufacturer’s instruction (Invitrogen, Carlsbad, CA, USA). A total of 1 µg of RNA was reverse-transcribed into cDNA using a First Strand cDNA Synthesis Kit (Fermentas, Burlington, ON, Canada).

Expression levels of GDF5 mRNA in ligament tissue were detected by quantitative PCR analytsor with a Rotor-Gene 3000 system (Corbett research, Concorde, NSW, Australia) using the Quantitect SYBR Green PCR Kit (Takara, Shiga, Japan) following the manufacturer’s guidelines, employing the β-actin gene as a reference gene. The following primer pairs were used: GDF5 forward 5’-CTG TGA TTC CAG GAG TGC AG-3’, and reverse 5’-ATC CTC TTC ATT GAC TCT GCC-3’ [31], β-actin forward 5’-CGA GAT CGT GCG GGA CAT-3’ and reverse 5’-CAG GAA GGA GGG CTG GAA C-3’ [32]. The quantitative PCR reaction was carried out in a 25 µl volume containing 2 × SYBR® Premix Ex Taq™ (Takara, Shiga, Japan) 12.5 µl, 0.2 µM of each primer, and 1 µl cDNA. PCR conditions were as follows: an initial denaturation at 95°C for 10 min, followed by 45 cycles of 95°C for 15 s, 56°C for 15 s, and 72°C for 15 s, and final extension at 72°C for 10 min. Gene expressions of GDF5 in all samples were analyzed by applying the 2^ΔΔCt relative quantification method [33].

Statistical analysis

The statistical analysis was done by SPSS 18.0 software (SPSS Inc., Chicago, USA). The comparisons of the genotype and allelic frequencies were performed using the chi-square (χ^2) test with Yates’ correction or Fisher’s exact test. Odds ratios (OR) and 95% confidence intervals (CI) were calculated for the disease in carriers of the specific alleles. Bonferroni multiple corrections were used for the corrected P (Pc) by multiplying the P value by the number of the statistical tests in order to avoid the α1 error. Continuous variables were reported as means ± standard deviation, and statistical comparisons were performed with the two-tailed, Student’s t-test and one-way ANOVA analysis. A P value < 0.05 was considered to be significant.

The statistical power to detect genetic association study was calculated using the QUANTO software (http://hydra.usc.edu/gxe). At the 0.05 level of significance with the two-sided test for GDF5 +104T/C polymorphism, our study had 87% power to detect an effect with a relative risk of 2.0 in the group of ACL rupture patients and healthy controls.

Results

Distribution of GDF5 allele and genotype frequencies in patients with ACL rupture and in healthy controls

In all participants, genotype distributions were in Hardy-Weinberg equilibrium for the +104T/C (rs143383) polymorphisms (P = 0.190 in ACL rupture and 0.067 in controls). The sex ratio...
and age distribution did not differ between ACL rupture and controls (P = 0.568; P = 0.849, respectively).

The distribution of allele and genotype frequencies of +104T/C (rs143383) in GDF5 gene in patients with ACL rupture and in controls was shown in Table 2. The prevalence of the TT genotype tended to be higher in ACL rupture than in controls, reaching a significant difference (62.6% vs. 48.0%, P< 0.001, OR = 1.81, 95% CI: 1.35-2.44). Similarly, the frequency of the T allele was increased in ACL rupture cases compared to the controls (79.9% vs. 70.5%, P< 0.001, OR = 1.67, 95% CI: 1.30-2.13).

A significant association was found when the patients were divided into subgroups according to gender (Table 2). In female participants, the frequencies of T allele and TT genotype of +104T/C were significantly higher in ACL rupture than in controls (78.6% vs. 69.4%, P = 0.019, OR = 1.81, 95% CI: 1.08-2.43; 60.2% vs. 46.5%, P = 0.028, OR = 1.74, 95% CI: 1.06-2.86, respectively). In male participants, the distribution of +104T/C allele and genotype frequencies did not differ between ACL rupture and the healthy controls.

**Association of +104T/C polymorphism with GDF5 mRNA expression levels in patients with ACL rupture**

As illustrated in Figure 1, the +104T/C polymorphism seemed to affect GDF5 mRNA expression levels in patients with ACL rupture. The GDF5 mRNA expression levels were lower in TT genotype of +104T/C (n = 34) than those in subgroups of C carrier (n = 28, P = 0.005).

**Discussion**

Although there has been an emerging interest in the role of GDF5 as the new target in the regulation of progress in sports injury related diseases, relatively limited information exists to date on the relationship between ACL rupture and GDF5 genetic polymorphisms. This study provided the evidence for the association between the +104T/C polymorphism in GDF5 gene and ACL rupture in the Han Chinese population in Central China. We found that the frequencies of T allele and TT genotype of +104T/C were significantly higher in ACL rupture group than in control group. Moreover, a significant association was found when the patients were further divided into subgroups according to gender. Our results suggested that in female participants the frequency of TT genotype of +104T/C were significantly higher in ACL rupture group than in control group, and GDF5 +104T/C polymorphism was likely to have some effect on disease prevalence in female subjects.

No other studies in the Chinese population regarding the +104T/C polymorphism in relation to ACL rupture have been published so far to compare our data. However, there were several studies showed similar results in connective tissue pathologies: it was worth mentioning that the GDF5 rs143383 TT genotype was previously associated with a 2-fold increase in risk of Achilles tendon ruptures and chronic tendinopathy [34]. The study in Asian populations and Japanese population indicated that the +104T/C polymorphism in GDF5 gene showed significant association with osteoarthritis [28]. Williams et al. found that the frequency of the TT genotype of +104T/C was significantly increased in patients with lumbar disc degeneration than in healthy controls [35]. Raleigh et al. showed that GDF5 rs143383 variant had no association with ACL rupture in Caucasians [25]. This difference might be explained by the limited power, as well as by ethnicity or gene - environment interactions present in Central China but not in Caucasians.

The current study found that individuals with ACL rupture who had TT genotype expressed...
lower levels of GDF5 mRNA compared with subjects who had C carriers (TC+CC). Our findings corroborated those of Egli et al. and Dodd et al. [13, 29]. GDF5 rs143383 has recently been reported to be associated with OA susceptibility, with lower expression of the risk-associated T allele observed in vitro and in vivo. The in vivo studies were performed on cartilage tissue from OA patients. A luciferase reporter assay in the Asian study demonstrated a direct functional effect of the rs143383 on the expression of a reporter construct in a chondrogenic cell line, with the OA associated T allele showing reduced expression relative to the C allele [28]. Our findings suggested that a slight reduction in the expression of GDF5 in ACL tissue increased the individual’s risk of developing ACL rupture.

Some studies have shown that ACL rupture may be linked to the transmission of heritable factors [36, 37]. Recently, the gene encoding GDF5 was shown to associate with Achilles tendon pathology (both ruptures and chronic tendinopathy) [34]. GDF-5 expression is an important regulator of ligament homeostasis, higher levels of GDF-5 stimulate the production of type I collagen (the predominant protein in ligaments) subunit genes in rabbit ACL by modulating integrin α2 activity [18]. So, the GDF5 gene has been proposed to affect ligament biology and as it associates with other musculoskeletal soft tissue injuries.

In conclusion, our study suggests that the T allele of +104T/C polymorphism in GDF5 gene were associated with patients with ACL rupture in the Han Chinese population in central China. Patients carrying TT genotype expressed lower levels of GDF5 mRNA than C carriers among ACL rupture patients. Our results suggested that the T allele of +104T/C may be a risk factor for ACL rupture patients. This is likely from decreased expressions of GDF5 mRNA. We recognized that our current study had limitations of small sample size and lack of functional characterization of the T mutation at this position in the promoter activity and T cell activation. Therefore, further studies with a larger sample size in independent cohorts from the same ethnic group are necessary to determine the relation of the GDF5 +104T/C polymorphism with ACL patients, and to investigate the function of the GDF5 gene polymorphism.

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Disclosure of conflict of interest

None.

Address correspondence to: Dr. Liao-Bin Chen, Department of Orthopaedic Surgery, Zhongnan Hospital, Wuhan University, Donghu Road 169, Wuhan 430071, Hubei Province, PR China. Tel: +86-027-67813585; Fax: +86-027-67812892; E-mail: lbchen@whu.edu.cn

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