Identification of a Susceptibility Locus for Severe Adolescent Idiopathic Scoliosis on Chromosome 17q24.3

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Abstract

Adolescent idiopathic scoliosis (AIS) is the most common spinal deformity, affecting around 2% of adolescents worldwide. Genetic factors play an important role in its etiology. Using a genome-wide association study (GWAS), we recently identified novel AIS susceptibility loci on chromosomes 10q24.31 and 6q24.1. To identify more AIS susceptibility loci relating to its severity and progression, we performed GWAS by limiting the case subjects to those with severe AIS. Through a two-stage association study using a total of ~12,000 Japanese subjects, we identified a common variant, rs12946942 that showed a significant association with severe AIS in the recessive model (P = 4.00 × 10^{-8}, odds ratio [OR] = 2.05). Its association was replicated in a Chinese population (combined P = 6.43 × 10^{-12}, OR = 2.21). rs12946942 is on chromosome 17q24.3 near the genes SOX9 and KCNJ2, which when mutated cause scoliosis phenotypes. Our findings will offer new insight into the etiology and progression of AIS.

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Introduction

Adolescent idiopathic scoliosis (AIS) is the most common structural deformity of the spine, occurring in 2–3% of healthy children from the age of 10 to skeletal maturity worldwide [1]. A Japanese study showed that the incidence of scoliosis of more than 15 degrees increases linearly with age from 10 (0.07% in boys, 0.44% in girls) to 14 (0.25% in boys, 1.77% in girls), and that most of these cases are AIS [2].

AIS is a multi-factorial disorder, with genetic factors playing an important role in its etiology [3]. Population studies have shown that its familial incidence is higher than that in general populations [4], while twin studies have consistently shown higher concordance in monozygotic compared with dizygotic twins. For example, a meta-analysis of several twin studies revealed 73% monozygotic and 36% dizygotic twin concordance [5]. Using the Danish Twin Registry, Andersen et al. observed 25% proband-wise concordance in monozygotic twins (six of 44 concordant) compared with 0% concordance in dizygotic twins (0 of 91), with an overall prevalence of approximately 1% [6].

Several genetic studies regarding AIS susceptibility have previously been reported. Although genome-wide linkage analyses have revealed some AIS susceptibility loci [7–16], only CHD7 has been identified as a susceptibility gene [13]. Genetic association studies of AIS, however, have identified several predisposition genes. Single nucleotide polymorphisms (SNPs) in ESR1, ESR2, MATX1, MTNR1B, and TPH1 genes are reported to be associated with AIS susceptibility [17–21]. Recently, we used genome-wide association study (GWAS) to identify novel AIS susceptibility locus on chromosomes 10q24.31 near the ESR1 gene [22] and 6q24.1 in the GPR126 gene [23].

We used a common-control design [24,25] in our previous GWAS. However, because undiagnosed general populations or patients with unrelated diseases are used as controls in this design,
there is a potential loss of power associated with the inability to exclude latent diagnoses of the disease. One way of overcoming this is to adopt a more stringent case definition; for example, one based on early age of onset or the identification of a more severe disease phenotype [26]. Severe cases are presumed to have a high influence AIS progression.

### Materials and Methods

We defined Cobb's angle for severe AIS as above 40°. Cobb’s angles were obtained at the time the patient was recruited in this study. A written informed consent was obtained from all subjects participating in the study. The study was approved by the institutional review boards of RIKEN and participating institutions. The subjects for the GWAS were all Japanese females; 554 with severe AIS (aged 10–39) and 1,474 control subjects (aged 7–96) were recruited as previously described [22]. For the Japanese replication study, we recruited an independent set of a case-control study consisting of 268 severe AIS (aged 10–39) and 9,823 controls (aged 20–96) in the same way. For examining the relation between the genotype of the SNPs identified by the case-control association study and the AIS severity (Cobb angle) in Japanese, we collected the data of 1,767 AIS subjects used for the previous GWAS and replication studies who had AIS with a Cobb angle of severe AIS was above 40°. Cobb’s angles were obtained at the time the patient was recruited in this study. For examining the relation between the genotype of the SNPs identified by the case-control association study and the AIS severity (Cobb angle) in Japanese, we collected the data of 1,767 AIS subjects used for the previous GWAS and replication studies who had AIS with a Cobb angle of severe AIS was above 40°. Cobb’s angles were obtained at the time the patient was recruited in this study. For examining the relation between the genotype of the SNPs identified by the case-control association study and the AIS severity (Cobb angle) in Japanese, we collected the data of 1,767 AIS subjects used for the previous GWAS and replication studies who had AIS with a Cobb angle of severe AIS was above 40°. Cobb’s angles were obtained at the time the patient was recruited in this study.
Table 2. Association of rs12946942 with severe AIS in Japanese and Chinese populations.

| Population | Study     | RAF     | P value | Odds ratio       | PBD   |
|------------|-----------|---------|---------|------------------|-------|
|            |           | case    | control |                   |       |
|            |           | 0.274   | 0.203   | 1.95 × 10^−1     | 2.24  |
| Japanese   | GWAS      | 0.274   | 0.203   | 1.95 × 10^−1     | 2.24  |
|            | Replication | 0.224   | 0.213   | 6.09 × 10^−2     | 1.59  |
|            | Total     | 0.258   | 0.211   | 4.00 × 10^−4     | 2.05  |
| Chinese    | Replication | 0.392   | 0.288   | 3.27 × 10^−3     | 2.59  |
| Combined   | Meta-analysis | 0.643   | 0.186   | 6.43 × 10^−12    | 2.21  |

RAF: risk allele (T allele) frequency. CI: confidence interval.

* By x2 test.
† by the Mantel-Haenszel method.

In Chinese, we recruited 571 females with severe AIS (aged 10–19) and 326 female controls (aged 25–83) living in and around Nanjing city, China. All were self-reported Han Chinese.

Genotyping of SNPs and Quality Control

Genomic DNA was extracted from the peripheral blood leukocytes of severe AIS and control subjects using standard protocols. In the GWAS, we genotyped case subjects using the Illumina Human610 Genotyping BeadChip and control subjects with the Illumina HumanHap550v3 Genotyping BeadChip. SNPs common to both platforms were then combined and analyzed as previously described [30]. Inadequate SNPs and subjects were checked and excluded as previously described [22]. In the Japanese replication study, the case subjects were genotyped using the PCR-based Invader assay [31] and controls were genotyped using Illumina HumanHap550v3 Genotyping BeadChip. In the Chinese replication study, all subjects were genotyped using the PCR-based Invader assay as described above.

Statistical Analysis

The association between the SNPs was examined by x2 test for three models (allele model, recessive model and dominant model) and minimum P values in the three models were evaluated. In the same way, incidence of severe AIS, and the Hardy–Weinberg equilibrium (HWE) of the genotypes were examined by x2 test. Data of GWAS and Japanese replication study were combined by addition, and data of total Japanese studies and Chinese study were combined using the Mantel-Haenszel method. The Breslow-Day statistic was used to test homogeneity of the common odds ratio. The associations between the SNP genotypes and the Cobb angle of AIS subjects were evaluated using the Kruskal-Wallis and Mann-Whitney U tests. Imputation was performed using MACH version 1.0.16.c and Minimac software with reference haplotypes from the 1000 Genomes Project June 2011 EAS population as described elsewhere [23]. In the analysis, pairwise r2 values were calculated using the R Bioconductor package snpMatrix (version 1.16.2), and the LD map was created using in-house programs. We performed association analysis of imputed data using the single.snp.tests function in the R package snpStats version 1.3.4 after converting Minimac output to the uncertain genotype data format for snpStats. Regional association plots were generated using R statistical environment version 2.13.0.

Results

After stringent quality control of the subjects and SNPs, we examined the association of 455,121 SNPs with severe AIS using the x2 test for three models (allele model, recessive model and dominant model). No SNP reached the GWAS significance threshold (P < 5 × 10^−8) at this stage (Figure S1). Then, we selected 27 SNPs (Table S1) according to the following criteria: 1) a minimum P value in the three models < 1 × 10−3; 2) a minor allele frequency ≥ 0.1. SNPs in strong linkage disequilibrium (LD) with a correlation coefficient (r2) of 0.8 with other SNPs were excluded from analysis. We checked their association using an independent set of Japanese female case-control subjects and combined all Japanese data.

Six SNPs showed association of genome-wide significance level (P < 5 × 10^−8) (Table 1). Five of them were in the known loci of AIS susceptibility that we previously reported; three SNPs (rs11190870, rs625039 and rs11958564) were close to LBX1 on chromosome 10q24.31 [22] and two SNPs (rs570507 and rs9496346) were on chromosome 6q24.1 in the GPR126 gene [23]. In addition, rs12946942 on chromosome 17q24.3 showed significant association in the recessive model (P = 4.00 × 10^−3, odds ratio [OR] = 2.05). We further examined the relation between the rs12946942 genotypes and the AIS severity (Cobb’s angle) using a total of 1,767 AIS cases. rs12946942 showed significant association (P = 3.02 × 10^−3; by the Kruskal-Wallis test).

We performed a replication study for rs12946942 in a Chinese case-control population. The association of rs12946942 was significant in the Chinese population for all three models. Combined P values from the Mantel-Haenszel method for the Japanese and Chinese studies in the recessive model showed genome-wide significance (P = 6.43 × 10^−12) (Table 2). rs12946942 defined a 130-kb LD block within an approximately 2-Mb region on chromosome 17 (Figure 1). No RefSeq genes have been mapped in this LD block. Twenty common SNPs in the LD block were genotyped in the GWAS, the most significant of which was rs12946942 (Figure 1).

To further characterize the chromosome 17q24.3 locus, we imputed genotypes of additional SNPs in the locus using 1000 Genomes Project’s East Asian population samples’ (EAS) reference haplotypes and tested their association with AIS. SNPs rs12946942 and rs12941471 yielded the strongest evidence for association (Figure S2A), which were in complete LD (r2 = 1) with each other. After conditioning on the top SNP (rs12946942), there was no secondary association signal for AIS within the region (Figure S2B).

Discussion

The region defined by rs12946942 was a gene desert. The closest genes include SOX9 and KCNJ2 [32]. SOX9 (MIM 608160) is a promising candidate gene for AIS as it encodes a transcription factor involved in chondrogenesis [33]. SOX9 mutations cause campomelic dysplasia (MIM 114290), a skeletal dysplasia characterized by bowed, long bones, small scapula, tracheobronchial narrowing, sex reversal and kyphoscoliosis [34]. Very long-range cis-regulatory elements controlling tissue-specific SOX9 expression have been previously reported [35,36]. The LD block containing rs12946942 has recently been defined as a susceptibility locus of prostate cancer in European Caucasians [37]. The block contains...
six enhancer elements, of which the E1 enhancer forms a long-range chromatin loop to SOX9 in a prostate cancer cell line. Two SNPs within the E1 enhancer were shown by in vitro reporter assays to direct allele-specific gene expression. We hypothesize that variants in this region may likewise participate in scoliosis pathogenesis by controlling scoliosis-related tissue-specific expression of SOX9 or other genes.

KCNJ2 (MIM 600681) is another promising candidate gene for AIS. It encodes the inward-rectifying potassium channel Kir2.1, which is a component of the inward rectifier current IK1. IK1 provides a repolarizing current during the most terminal phase of repolarization and is the primary conductance that controls the diastolic membrane potential [38]. KCNJ2 mutations lead to a cardiomyopathic type of periodic paralysis known as Andersen-Tawil syndrome (ATS; MIM 170390) [39], which is characterized by ventricular arrhythmias, periodic paralysis, facial and skeletal dysmorphism including hypertelorism, small mandible, cleft palate, syndactyly, clinodactyly, and scoliosis [38,39]. Furthermore, the 17q24.2-q24.3 micro-deletion syndrome whose deletion area includes KCNJ2 and rs12946942 showed skeletal malformations similar to the ATS phenotype including progressive scoliosis [40]. Interestingly, a similar micro-deletion syndrome including KCNJ2, but not rs12946942, was not associated with a scoliosis phenotype [41].

Thus, through a Japanese GWAS followed by replication studies in Japanese and Chinese populations, we identified a susceptibility locus for severe AIS on chromosome 17q24.3 that showed genome-wide significance. This region contains a few promising candidate genes that may be associated with the disease. Further studies are now necessary to identify the causal gene and its variant in the locus.

Supporting Information

Figure S1 Manhattan plot showing the P values from genome-wide association study (minimum P value in allele, recessive and dominant models). The horizontal line represents the genome-wide significance threshold (P = 5 × 10^{-8}). (TIF)

Figure S2 Regional association plots and recombination rates of AIS susceptibility locus on chromosome 17q24.3. The chromosome position (NCBI Build 37) of SNPs...
against $-\log_{10} [P\text{ value}]$ from a logistic regression analysis is shown. (A) Unconditioned analysis. The SNP with highest association signal (rs12946942) is represented as a purple diamond. Imputed (circles) and genotyped SNPs (squares) are colored according the LD (r²) with rs12946942. (B) Conditioned analysis. Red circles are unconditioned and gray circles are conditioned for rs12946942 (gray triangle).

### Table S1 Association of the 27 SNPs selected from the GWAS

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### Author Contributions

Conceived and designed the experiments: AM MM SL. Performed the experiments: AM Y. Takahashi JD MK. Analyzed the data: AM IK TAJ AT T. Tsunoda. Contributed reagents/materials/analysis tools: YO XQ HJ H. Yan KK NK MU SM H. Yanagida HT NH T. Tsuji TS HS TK IY KW KC Y. Toyama YQ. Wrote the paper: AM SI.