INTRODUCTION

Adolescent idiopathic scoliosis (AIS) is characterized as a puberty-onset spinal deformity that affects millions of children worldwide. Despite great progress made in the study of AIS over the last 20 years, its pathogenesis is far from fully understood due to its genetic complexity. Spinal fusion is recommended for AIS patients with Cobb angles exceeding 45°, but the risks and health economics related to surgical treatments are thorny issues. Clarifying the pathogenesis of AIS is of great significance for its early diagnosis, prevention, and treatment.

Tent5a modulates muscle fiber formation in adolescent idiopathic scoliosis via maintenance of myogenin expression

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Abstract
Objective: Paravertebral muscle asymmetry may be involved in the pathogenesis of adolescent idiopathic scoliosis (AIS), and the Tent5a protein was recently identified as a novel active noncanonical poly(A) polymerase. We, therefore, explored the function of the AIS susceptibility gene Tent5a in myoblasts.

Materials and methods: RNA-seq of AIS paravertebral muscle was performed, and the molecular differences in paravertebral muscle were investigated. Twenty-four AIS susceptibility genes were screened, and differential expression of Tent5a in paravertebral muscles was confirmed with qPCR and Western blot. After the knockdown of Tent5a, the functional effects of Tent5a on C2C12 cell proliferation, migration, and apoptosis were detected by Cell Counting Kit-8 assay, wound-healing assay, and TUNEL assay, respectively. Myogenic differentiation markers were tested with immunofluorescence and qPCR in vitro, and muscle fiber formation was compared in vivo.

Results: The AIS susceptibility gene Tent5a was differentially expressed in AIS paravertebral muscles. Tent5a knockdown inhibited the proliferation and migration of C2C12 cells and inhibited the maturation of type I muscle fibers in vitro and in vivo. Mechanistically, the expression of myogenin was decreased along with the suppression of Tent5a.

Conclusions: Tent5a plays an important role in the proliferation and migration of myoblasts, and it regulates muscle fiber maturation by maintaining the stability of myogenin. Tent5a may be involved in the pathogenesis of AIS by regulating the formation of muscle fiber type I.
The etiology of AIS is multifactorial, including abnormal bone metabolism, endocrine hormone dyscrasia, or neuromuscular system imbalance, and the genetic basis cannot be ignored in the pathogenesis of AIS. A functional variant of AIS, Lbx-As1, was identified in a genome-wide association study based on 4,317 Chinese AIS patients. Many susceptibility genes related to skeletal muscle development, such as Lbx1, Pax3, and Tbx1, have been reported in AIS. Recently, a large number of studies have reported paravertebral muscle asymmetry based on imaging changes, histological characteristics, and electrophysiological differences, which indicates that paravertebral muscle asymmetry may be involved in the pathogenesis of AIS.

Recently, 20 loci significantly associated with AIS have been reported from 79,211 Japanese individuals, and Tent5a was one of the susceptibility genes for AIS. Tent5a was recently identified as a novel active noncanonical poly(A) polymerase. Poly(A) length is highly regulated in the nucleus and cytoplasm, and poly(A) tails help regulate gene expression and probably play a critical role in cell differentiation. Tent5a has previously been found to be associated with a variety of human diseases, including retinitis pigmentosa, skeletal disorders, Alzheimer's disease, and obesity. However, the function of the Tent5a gene in muscle development is still unclear.

Based on the asymmetry of paravertebral muscle in AIS, we explored the function of the AIS susceptibility gene Tent5a in myoblasts, as well as the formation of muscle fibers. We revealed that Tent5a plays an important role in the proliferation and migration of myoblasts and regulates muscle fiber maturation by maintaining the stability of myogenin. These findings shed new light on the etiology of AIS and provide new therapeutic opportunities for AIS patients to correct muscle asymmetry with nonoperative management.

2 | MATERIALS AND METHODS

2.1 | Multifidus specimens of AIS patients

This study was approved by the institutional research ethics board (HX-2019-607). Twenty female AIS patients with a right thoracic curve who underwent spinal corrective surgery between the ages of 12 and 18 were included in this study (Figure 1A). In addition, we collected muscle samples from five teenagers who underwent spinal surgery due to spinal cord trauma as the normal control. Detailed information for curve type, age, gender, and Cobb angle of the AIS patients is presented in Table S1. Muscle tissues (~1.0 x 1.0 x 0.5 cm) were obtained from the multifidus at the apical region of concavity and convexity before tissue cautery. Specimens were divided into two parts. One part was frozen in precooled isopentane for immunofluorescence staining, and the other part was flash-frozen in liquid nitrogen for RNA and protein testing.

2.2 | RNA-seq and differential expression analysis

After total RNA was extracted and quantified, library preparation of each sample was constructed according to the manufacturer’s protocol with 1 μg total RNA. After isolation of the poly(A) mRNA, mRNA fragmentation and priming were performed using NEBNext First Strand Synthesis Reaction Buffer and NEBNext Random Primers. Then, libraries with different indices were multiplexed and loaded on an Illumina HiSeq instrument according to the manufacturer’s instructions (Illumina).

Quality control, mapping, and expression analysis were performed, and the Padj of genes was set to <0.05 to detect differentially expressed genes. GOSeq (v1.34.1) was used to identify Gene Ontology terms that annotate a list of enriched genes with a significant Padj less than 0.05. In addition, topGO was used to plot DAG. We used scripts in-house to enrich significantly differentially expressed genes in KEGG pathways.

2.3 | Expression analysis of AIS susceptibility genes

In this study, 24 susceptibility genes reported in East Asian AIS patients according to two large sample size genome-wide association studies were selected. Zhu et al. reported five susceptibility genes (Lbx1-As1, Pax3, Eph4, Bcl2, and Ajap1) with a sample of 4,317 Chinese AIS patients and 6,016 controls. The other 19 susceptibility genes (Mtmr11, Arf1, Tbx1, Csmd1, Klf24, Bckdhb, Tent5a, Creb5, Nt5dc1, Uncx, Pixna2, Agmo, Meox2, Fto, Lbx1, Bnc2, Abo, Pax1, and Cdh13) were identified from a meta-analysis of three genome-wide association studies consisting of 79,211 Japanese individuals. Detailed information on the 24 susceptibility genes is presented in Table S2.

2.4 | C2C12 cell culture and lentiviral transfection

C2C12 cells were acquired from the Chinese National Collection of Authenticated Cell Cultures (No. SCSP-505). C2C12 cells were cultured in growth medium (DMEM with 10% fetal bovine serum, Gibco), and myogenic differentiation was induced with differentiation medium (DMEM with 2% horse serum, HyClone) when C2C12 cells grew to 90% confluence.

The knockdown and control lentiviruses were purchased from Wuhan BrainVTA Co., Ltd. The density of C2C12 cells was approximately 80% after inoculation for 24 h, the growth medium was replaced, and concentrated lentivirus was added (MOI = 100). Auxiliary infection reagents were also added, and the growth medium was replaced the next day. Puromycin was used to select Tent5a-inhibited C2C12 cells. EGFP fluorescence was detected with fluorescence microscopy 48 h after transfection, and qPCR and Western blotting were used to verify the knockdown effectiveness.
2.5 | Effects of Tent5a knockdown on C2C12 cells

C2C12 cells at 70% confluence were selected and prepared into a single-cell suspension. To detect the effect of Tent5a knockdown on myoblast viability, C2C12 cells were inoculated in 96-well plates, and 2000 cells in 100 µl of growth medium were seeded in each well. After incubation for 24 h, the Cell Counting Kit-8 assay (YZ-CK04, Solarbio) was performed according to the manufacturer’s protocol. The wound-healing assay was used to quantify cell migration. Cells were inoculated with a 6-well plate and scratched when the cell density increased close to 100%. The observation time points of the wound-healing assay included 0, 24, and 48 h. To detect the effect of Tent5a knockdown on myoblast apoptosis, C2C12 cell apoptosis was induced by H2O2, and a TUNEL assay (A113, Vazyme) was conducted according to the manufacturer’s protocol. After culturing in differentiation medium for 5 days, C2C12 cells were fused and formed mature myotubes. Immunofluorescence staining and qPCR were used to detect the effect of Tent5a knockdown on myoblast differentiation.

2.6 | Primary cultures and cell transplantation

Satellite cells were isolated by a method described by Watanabe et al. Briefly, the quadriceps femoris muscles of 4-week-old C57BL/6 mice (Dossy Experimental Animals Co., Ltd.) were cut off, and nonmuscular tissues, including fat, fascia, nerves, blood vessels, and tendons, were carefully removed using an asana microscope. The muscle was repeatedly pruned to approximately 1 mm3 in size with ophthalmic scissors and digested with 0.1% type I collagenase (C8140, Solarbio) for 1 h. After enzymatic digestion, filtration, and resuspension, satellite cells were incubated in high serum growth medium. Satellite cells were further purified by the differential adhesion method and verified by Pax7 staining (Pax7, DSHB). P3-5 myoblasts were selected for lentivirus transfection, and the knockdown efficiency of Tent5a expression was confirmed with qPCR and Western blot.

One day before cell transplantation, 4-week-old C57BL/6 mice were selected to create a muscle injury model. Briefly, 50 µl of cardiotoxin (10 µM) was multipoint injected into the tibialis anterior...
muscle using a 31G insulin syringe. Then, myoblasts transfected with knockdown or control lentivirus were prepared, and 50 µl of single-cell suspension within 5 x 10^6 transfected myoblasts was multipoint injected into the tibial anterior muscle.22 Four weeks after myoblast transplantation, the tibialis anterior muscles were collected for subsequent experiments.

### 2.7 Immunofluorescence staining

The thickness of the frozen section for AIS samples was 8 µm. After fixation with 4% paraformaldehyde and then blocking with 5% BSA (Biofroxx) in 0.025% PBS-Tween 100 (Biofroxx), we incubated the slices with primary antibodies against MHC I (1:20, BA-D5, DSHB) and MHC IIA (1:20, SC-71, DSHB) overnight at 4°C. The secondary antibodies (1:200, Yeasen) were incubated at room temperature and protected from light for 1 h. Muscle samples of mouse tibialis anterior were processed in the same manner as described above.

For satellite cells and C2C12 cells, 4% paraformaldehyde and 0.5% Triton®X-100 (Biofroxx) were used to incubate cells successively. After blocking, they were stained with the following primary antibodies: Pax7 (1:5, Pax7, DSHB), MHC (1:20, MF 20, DSHB), Myh7 (1:100, sc-53090, Santa Cruz), myogenin (1:100, sc-12732, Santa Cruz), and MyoD (1:100, sc-377460, Santa Cruz). The nuclei were stained with DAPI for 5 min. We collected the images using a fluorescence microscope. ImageJ software (Version 1.52 V) was used to analyze the percentage of myosin heavy chain-, Myh7-, myogenin-, and MyoD-positive cells.

### 2.8 qPCR and Western blot

After the muscle samples were pulverized using a mortar and pestle in liquid nitrogen or the cells were washed with PBS, RNAiso (108-95-2, Takara) was added, and total RNA was isolated with Qiagen RNAeasy Mini Kits (1062832, Qiagen) per the manufacturer’s guidelines. The PrimeScript™ RT reagent kit with a gDNA eraser was used to produce cDNA. PowerUp™ SYBR™ green master mix (A25742, Thermo Fisher Scientific) was added to detect the expression of related genes, and the gene-specific primers are listed in Table S3 and Table S4. Quantitative real-time PCRs were performed with an ABI QuantStudio 3 machine (Thermo Fisher Scientific).

RIPA lysis solution with 1% PMSF was used to extract total protein. Protein quantification was performed using the BCA method. Gel electrophoresis and membrane transfer were performed according to standard processes. After nonspecific antigen blocking with 5% BSA, primary antibodies including Tent5a (1:200, A12765, ABclonal), MHC (1:50, MF20, DSHB), MHC I (1:50, BA-D5, DSHB), myogenin (1:500, sc-12732, Santa Cruz), and Gapdh (1:5000) were used to incubate the polyvinylidene fluoride transfer membrane overnight. Images were captured with the ChemiDoc™ Imaging System. The relative expression of Tent5a, MHC, and MHC I was calculated according to the gray values using ImageJ software (Version 1.52 V).

### 2.9 Statistical analysis

Data were analyzed and plotted using GraphPad Prism 8.4.3 (GraphPad Software Inc.), and all quantitative data are presented as the mean ± SD. Statistical analyses were carried out using t-tests or rank-sum tests when comparing continuous variables and the chi-square test or Fisher’s exact test for dichotomous variables. Statistical tests were chosen based on sample size and normality of distribution. A p value of < 0.05 was considered statistically significant.

### 3 RESULTS

#### 3.1 RNA sequencing of AIS paravertebral muscles

To investigate the molecular differences of the paravertebral muscle in AIS, RNA sequencing of convex and concave paravertebral muscles was performed (Figure 1B). The results showed that the expression of 10 genes was upregulated on the concavity side, and 48 genes were downregulated (Figure 1C). KEGG analysis showed that the differentially expressed genes were enriched mainly in the insulin signaling pathway, PPAR signaling pathway, and cAMP signaling pathway (Figure 1D). GO analysis showed that the differentially expressed genes were enriched mainly in extracellular space, glucose metabolic process, and glucose homeostasis (Figure 1E).

#### 3.2 Differential expression of Tent5a in AIS paravertebral muscles

We screened for the expression of previously reported AIS susceptibility genes from RNA sequencing data, including Mtmr11, Arf1, Tbx1, Csmd1, Kif24, Bckdhb, Tent5a, Creb5, Nt5Dc1, Uncx, Plxn2, Agmo, Meox2, Fto, Lbx1, Bnc2, Abo, Pax1, Cdhs13, Lbx1-As1, Pax3, Epha4, Ajap1, and Bcl12.5,13 Only the Tent5a gene showed differential expression in the two-sided paravertebral muscles of AIS patients (Figure 2). The differential reads for Tent5a were also confirmed with the use of the IGV tool (Figure S1). We selected another 10 pairs of paravertebral muscles from AIS patients and verified the differential expression of Tent5a with qPCR (Figure 3A) and Western blot (Figure 3B,C).

#### 3.3 Asymmetry of fiber type in AIS

Compared with the convex side, a significantly lower proportion of type I muscle fibers was found on the concave side (Figure 3F,G), but the proportion of type IIA muscle fibers was higher (Figure 3H,I). In
addition, the expression of MHC and MHC I proteins in the paravertebral muscles of AIS patients was detected by Western blot. There was no significant difference in the relative expression of MHC between the concavity and convex sides (Figure 3B,D), and the MHC I protein on the concave side of the paravertebral muscle was significantly lower than the MHC I protein on the convex side (Figure 3B,E). These results support the asymmetry of fiber type in AIS. In addition, the relative mRNA expressions of myofiber-related genes were compared among the convex side, concave side, and control muscles. The mRNA expressions in the concave side of AIS, but not the convex side, were closer to the normal control, including Myh1, Myh7, Tnn1, Tnn2, Tnnt3, and Tent5a (Figure 3J).

3.4 | Tent5a knockdown inhibited the proliferation and migration of C2C12 cells

To confirm the effect of Tent5a on myoblasts, we produced Tent5a knockdown C2C12 cells by transfection with lentivirus. Cell counting showed that the cell number of the Tent5a knockdown group was significantly lower than the cell number of the Tent5a knockdown group of the negative control group on Day 3 (Figure 4C). In addition, the Cell Counting Kit-8 assay showed lower cell viability in the Tent5a knockdown group (Figure 4D).

The effect of Tent5a knockdown on the migration ability of C2C12 cells was detected with a wound-healing assay. After 48 h of culture, the migration area in the Tent5a knockdown group was significantly lower than the migration area in the Tent5a knockdown group in the control group (Figure 4E,F). A TUNEL assay was used to detect the effect of Tent5a knockdown on the apoptosis of C2C12 cells. The percentage of apoptosis was comparable between the two
groups. (Figure 4G,H). These results suggest that Tent5a knockdown inhibited the proliferation and migration of C2C12 cells.

### 3.5 Tent5a knockdown inhibited myogenic differentiation of C2C12 cells

After 5 days of myogenic differentiation, the expression of the myogenic protein was detected with immunofluorescence staining. Compared with the negative control group, the intensity and density of immunofluorescence for MHC were significantly lower in the Tent5a knockdown group (Figure 5A). The percentage of MHC (+) nuclei was also decreased in the Tent5a knockdown group (Figure 5B). Myogenin, a myogenic transcription factor essential for myogenic differentiation, was decreased in the Tent5a knockdown group (Figure 5D–F). These results suggested that Tent5a knockdown inhibited the myogenic differentiation of C2C12 cells.

Intriguingly, we found that the deregulation of Tent5a led to a significant decrease in the mRNA expression of Myh7, Tnni1, and Tnnc1 (Figure 5G). Furthermore, the percentage of Myh7 (+) nuclei was also decreased in the Tent5a knockdown group (Figure 5C). These results suggested that Tent5a knockdown inhibited mainly myogenic differentiation and especially impaired the maturation of genes related to type I muscle fibers.

### 3.6 Tent5a knockdown inhibited the maturation of type I muscle fibers in vivo

To investigate the effect of Tent5a on myogenic differentiation in vivo, myoblasts were labeled with EGFP using lentiviral transfection and transplanted into the tibial anterior muscle of mice (Figure 6A). EGFP-positive muscle fibers represented the myogenic differentiation of transplanted cells. Four weeks later, we found that the percentage of MHC I/EGFP muscle fibers in the Tent5a knockdown group was significantly lower than the percentage of MHC I/EGFP muscle fibers in the control group (Figure 6B,C). In the Tent5a knockdown group, the relative mRNA expression of Myh7 and Tnnc1 was also significantly decreased, while the relative expression of the Tnnc2 gene was increased (Figure 6D). These results confirmed that
Tent5a knockdown mainly inhibited the maturation of type I muscle fibers in vivo.

### 3.7 Regulatory effects of Tent5a on myogenic regulatory factors

We detected the mRNA expression of Tent5a, myogenic regulatory factors (myogenin, MyoD, Myf5, and Myf6), and myogenic-related genes (Myh4 and Myh7). The relative expression of Tent5a was upregulated from Day 0 to Day 3 and downregulated gradually from Day 4 to Day 5 (Figure 7A). The relative expression of myogenin was significantly upregulated on the first day and maintained at a high level on Day 5 (Figure 7A). The Western blot results showed that the relative expression of Tent5a was upregulated from Day 0 to Day 2 and then maintained at high expression levels during myoblast differentiation (Figure 7C,D), and the relative expression of myogenin was increased progressively from Day 0 to Day 5 (Figure 7C,E).

To investigate the regulatory effects of Tent5a on myogenic regulatory factors, we conducted Tent5a knockdown experiments, and the mRNA expression levels of myogenin, MyoD, Myf5, and Myf6 were detected on Day 3 of myogenic differentiation. The relative expression of myogenin was significantly downregulated in the Tent5a knockdown group (Figure 7B). The previous results also showed decreased myogenin (+) nuclei in the Tent5a knockdown group (Figure 5D–F). These results suggested that Tent5a knockdown mainly restrained the expression of myogenin.
DISCUSSION

Because of the genetic complexity of AIS, its pathogenesis has not been clarified. In contrast to congenital scoliosis, no significant vertebral dysplasia is observed in the early stages of AIS. Recently, an increasing number of studies have speculated that paravertebral muscle asymmetry may be involved in the pathogenesis of AIS. Here, we investigated the role of the susceptibility gene Tent5a in the pathogenesis of AIS, and we found that Tent5a knockdown inhibited the proliferation and migration of C2C12 cells and inhibited the maturation of type I muscle fibers in vitro and in vivo. In addition, the Tent5a gene might maintain myogenic differentiation by promoting the expression of myogenin. Together, these results indicate that the AIS susceptibility gene Tent5a may be involved in the pathogenesis of AIS by modulating muscle fiber formation via maintenance of myogenin expression (Figure 8).

Although paravertebral muscle asymmetry has been reported in AIS, the molecular differences in the paravertebral muscle of AIS are unclear. We conducted RNA-seq to compare the convex and concave paravertebral muscles. Fifty-eight genes were differentially expressed in AIS paraspinal muscle, and these genes were mainly enriched in the insulin signaling pathway, PPAR signaling pathway, and cAMP signaling pathway. The insulin signaling pathway is crucial for the regulation of glucose levels and energy metabolism, and its downstream molecules play important roles in glycogen synthesis, protein synthesis, glucose transport, and cell apoptosis. Glucose metabolism and insulin sensitivity are essential for muscle contraction, and previous evidence has shown that muscle disorders could cause abnormal lipid deposition. Recently, FR030200-Tent5a signaling has been reported to participate in the insulin resistance of the vascular injury. In addition, Tent5a has been validated as a trans-regulator for leptin, and Tent5a knockdown increased insulin-stimulated leptin secretion from the adipocytes. PPAR belongs to
the ligand-activated receptor family of nuclear hormone receptors. One of its subtypes, PPARα, is involved in lipid metabolism and regulates the expression of many myogenic genes.\textsuperscript{31–33} Similarly, scholars recently revealed the differential expression of H19 and ADIPOQ in the paravertebral muscles of AIS patients, and they speculated that transcriptome differences in AIS patients might be the reason for the structural and functional imbalance of paravertebral muscles.\textsuperscript{34}

Investigating the function of AIS susceptibility genes could help us to further understand the pathogenesis of AIS. Recently, several AIS susceptibility genes, such as Lbx1 and Pax3, have been identified to be associated with skeletal muscle function.\textsuperscript{5,8,35,36} We screened 24 AIS susceptibility genes and verified the differential expression of Tent5a in the paravertebral muscle. Furthermore, we knocked down the expression of Tent5a and inhibited the proliferation and migration function of myoblasts.

The myoblasts could fuse and transform into fast-twitch (type I) and slow-twitch (type IIa, type IIb, and type IIx) skeletal muscle fibers in vivo. The function of skeletal muscle depends largely on different proportions of muscle fibers, and its delicate

\textbf{FIGURE 7} Regulatory effects of Tent5a on myogenic regulatory factors. (A) The dynamic expression of Tent5a and myogenic regulatory factors (myogenin, MyoD, Myf5, and Myf6) during myogenic differentiation. (B) qPCR showing decreased mRNA expression of myogenin in the Tent5a knockdown group. Each bar represents mean ± SD. Unpaired t-test. \(*p < 0.05, \**p < 0.01. (C–E) Representative Western blots showing dynamic expression of Tent5a and myogenin during myogenic differentiation. Each bar represents mean ± SD. Unpaired t-test. \(*p < 0.05, \**p < 0.01, \***p < 0.001

\textbf{FIGURE 8} Schematic drawing of Tent5a modulates muscle fiber formation in adolescent idiopathic scoliosis. The poly(A) polymerase Tent5a promotes myogenic differentiation by maintaining the mRNA stability of myogenin, and it is involved in the pathogenesis of AIS by modulating muscle fiber formation.
combination is essential for complex body movement and postural behavior. In AIS patients, we found different proportions of type I and type IIA muscle fibers between the concave and convex sides of the paravertebral muscles, which supported the asymmetry of paravertebral muscle in AIS. In addition, we found that Tent5a knockdown inhibited myogenic differentiation and especially impaired the maturation of genes related to type I muscle fibers. The composition of muscle fibers is regulated by many factors. Exercise could promote an increased proportion of type I muscle fibers and enable the transition from fast to slow muscles. Tbx1 was reported to regulate the muscle fiber type and oxidative metabolism in the myotube and stimulate the differentiation of myoblasts. Taken together, these results indicate that Tent5a may be involved in the pathogenesis of AIS by disturbing muscle fiber formation.

Tent5a has recently been identified as an active atypical poly(A) polymerase. Poly(A) polymerase plays a highly modulatory role in the nucleus and cytoplasm, and different lengths of poly(A) tails could influence the stability, transport, and translation of mRNAs. Recently, Tent5a was identified as a trans-regulator of leptin through proteomic analysis. Patients identified with mutation of Tent5a were associated with severe osteogenesis imperfecta, and skeletal dysplasia was found in the mouse with Tent5a mutation. In addition, Tent5a was expressed in hematopoietic system cells and played an important role in heme-induced hemoglobinization. Deletion of the Tent5a gene led to abnormal formation of eye and body color in Xenopus. In this study, we first reported the function of Tent5a in myoblasts and myogenic differentiation in vitro and in vivo.

As the core member of myogenic regulator factors, myogenin regulates the fate of myoblasts and culminates in the formation of mature muscle fibers. In this study, we found that Tent5a knockdown mainly restrains the expression of myogenin. Previous studies have shown that myogenin processed myoblast fusion and determined the number and size of muscle fibers. MyoD and myogenin are an overlapping group of myogenic regulatory factors that maintain the expression of each other in a feed-forward cycle. Rather than MyoD, late maturation depends mainly on myogenin, and MyoD cannot compensate for the loss of myogenin in vivo. In addition to controlling the maturation of skeletal muscle, myogenin also affected the location and distribution of resident satellite cells, which demonstrated a multilevel contribution of myogenin to muscle homeostasis throughout life. Therefore, the poly(A) polymerase Tent5a might play an important role in maintaining the mRNA stability of myogenin. The risks and health economics related to spinal correction in AIS are thorny issues, and Tent5a might be a new regulatory target for muscle development. Our findings provide new therapeutic opportunities to correct muscle asymmetry for AIS with nonoperative management.

5 | CONCLUSIONS

The poly(A) polymerase Tent5a plays an important role in the proliferation and migration of myoblasts, and it might regulate muscle fiber maturation by maintaining the stability of myogenin. As the susceptibility gene of AIS, Tent5a may be involved in the pathogenesis of AIS by regulating the formation of muscle fiber type I. These findings provide new therapeutic opportunities to correct muscle asymmetry for AIS with nonoperative management.

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CONFLICT OF INTEREST

The authors have declared that no competing interest exists.

AUTHOR CONTRIBUTIONS

YMS and SSH designed the project. ML, HLY, DWW, and XHY collected and analyzed the data. ML and HLY wrote the manuscript. All authors reviewed the manuscript.

DATA AVAILABILITY STATEMENT

I confirm that my article contains a Data Availability Statement even if no data is available (list of sample statements) unless my article type does not require one (e.g., Editorials, Corrections, Book Reviews, etc.).

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