Wnt Protein-mediated Satellite Cell Conversion in Adult and Aged Mice Following Voluntary Wheel Running

Background: Satellite cells are activated in response to muscle injury or mechanical stimuli. Results: Running-induced up-regulation of Wnt/β-catenin signaling activates Myf5 and MyoD transcription. Conclusion: Voluntary wheel running induces satellite cell activation by direct regulation of Wnt/β-catenin signaling. Significance: Focusing on Wnt signaling may help to understand the function and intrinsic ability of satellite cells in adult myogenesis.

Muscle represents an abundant, accessible, and replenishable source of adult stem cells. Skeletal muscle-derived stem cells, called satellite cells, play essential roles in regeneration after muscle injury in adult skeletal muscle. Although the molecular mechanism of muscle regeneration process after an injury has been extensively investigated, the regulation of satellite cells under steady state during the adult stage, including the reaction to exercise stimuli, is relatively unknown. Here, we show that voluntary wheel running exercise, which is a low stress exercise, converts satellite cells to the activated state due to accelerated Wnt signaling. Our analysis showed that up-regulated canonical Wnt/β-catenin signaling directly modulated chromatin structures of both MyoD and Myf5 genes, resulting in increases in the mRNA expression of Myf5 and MyoD and the number of proliferative Pax7+ Myf5+ and Pax7+ MyoD+ cells in skeletal muscle. The effect of Wnt signaling on the activation of satellite cells, rather than Wnt-mediated fibrosis, was observed in both adult and aged mice. The association of β-catenin, T-cell factor, and lymphoid enhancer transcription factors of multiple T-cell factor/lymphoid enhancer factor regulatory elements, conserved in mouse, rat, and human species, with the promoters of both the Myf5 and MyoD genes drives the de novo myogenesis in satellite cells even in aged muscle. These results indicate that exercise-stimulated extracellular Wnts play a critical role in the regulation of satellite cells in adult and aged skeletal muscle.

Satellite cells are muscle-specific stem cells located between the basal lamina and plasma membrane of muscle fibers (1). In adult skeletal muscle, satellite cells represent the source of 2.5–6% of all the nuclei of a muscle fiber and remain in a quiescent state under normal conditions. Satellite cells play an essential role in muscle regeneration and recovery because they have the intrinsic ability to differentiate to generate a large number of new myofibers in few days (2). Exercise alters myofiber composition to increase muscle performance and metabolism and also has a positive effect on the regulation of satellite cells. Several studies have reported that the number of satellite cells increases after exercise in humans and animals (3, 4). In contrast, the number of satellite cells decreases during aging, and this is associated with a decrease in muscle quality and functional properties (5). Loss of skeletal muscle mass with concomitant fibrosis during aging eventually leads to sarcopenia. Sarcopenia is now recognized as a serious health issue affecting millions of aging adults, and exercise is one of the primary treatments used to prevent muscle shrinking and to reduce the risks of sarcopenia because it can increase muscle strength and endurance, even in old age.

Although physical exercise plays an important role in the intervention for sarcopenia, its effectiveness on satellite cells in the skeletal muscle during aging has not been well evaluated. Satellite cells exhibit a quiescent state and an activated proliferative state during skeletal muscle turnover, and each state can be identified with specific stage markers. Both quiescent and activated satellite cells express a stem cell-specific transcriptional factor, Pax7. Quiescent satellite cells are Pax7-positive but are negative for myogenic Myf5 and MyoD. In contrast, activated satellite cells co-express Myf5 and MyoD together with Pax7. Myf5 and MyoD are members of the basic helix-loop-helix transcriptional factors that play essential roles in controlling satellite cell differentiation and skeletal muscle development (6). Although changes in satellite cell numbers upon exercise stimulation have been extensively investigated, not many studies have been performed on the characteristic changes in satellite cells themselves, and the molecular mechanisms of exercise-stimulating extracellular factors controlling the expression profiles of satellite cells remain elusive.

Exercise alters various extracellular signals in skeletal muscles, affecting the fate of satellite cells. Wnt signaling during embryonic myogenesis and postnatal development plays important roles in lineage control (7). Wnt signaling...
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induces satellite cell proliferation during skeletal muscle regeneration (8). The transition from Notch signaling to Wnt signaling controls the transition from proliferation to differentiation in myogenic progenitors during muscle regeneration (9). Wnt signaling has positive effects on skeletal muscle regeneration but negative effects related to the induction of muscle fibrosis (10). Although many studies have indicated that Wnt has multiple functions in skeletal muscle (10–13), the actions of Wnts in response to exercise stimuli have not been well characterized, especially their roles in the regulatory cascade with respect to controlling satellite cells.

Wheel running is an effective exercise model for long-distance running in rodents and is less stressful than other models because it is not a compulsory exercise (14, 15). Wheel running exerts various physiological effects, including decreased body mass, increased maximum oxygen uptake, and metabolic improvement of skeletal muscles (16–20). Remarkably, the number of satellite cells in skeletal muscle increases after wheel running (21). Although the molecular mechanism underlying this effect remains unclear, some studies have reported that Wnt signaling is activated by acute treadmill running and functional overload (22, 23). However, the contribution of Wnt/β-catenin signaling to the regulation of satellite cells in response to exercise is unknown, and parallel examinations of muscle fibrosis following exercise have not been performed.

In this study, we investigated the effects of Wnt/β-catenin signaling on the regulation of satellite cells after voluntary wheel running in both adult and aged mice. Interestingly, exercise significantly up-regulated the expression of Wnt3, Wnt5a, and Wnt5b, and the up-regulation of Wnt signaling altered the quiescent state of satellite cells into the activated state, even in aged skeletal muscle without fibrosis. Canonical Wnt signaling directly modulated the chromatin structures of the Myf5 and MyoD promoters and up-regulated their expression in concert with the activation of satellite cells. To our knowledge, this report is the first to show the Wnt-mediated positive myogenic effects of exercise on satellite cells in adult and aged skeletal muscle and to describe the regulatory mechanism underlying chromatin remodeling of the Myf5 and MyoD genes.

EXPERIMENTAL PROCEDURES

Animal Care—Animal experiments were carried out in a humane manner after receiving approval from the Institutional Animal Experiment Committee of the University of Tsukuba and the Institutional Animal Care and Use Committee (IACUC) of the National Institute of Advanced Industrial Science and Technology. Animals were housed in animal facilities with sufficiently controlled temperature and humidity under a 12/12-h light/dark cycle and had access to chow and water ad libitum. Male C57BL/6J mice (CLEA Japan, Japan) ages 8–12 weeks (adult) and 24 months (aged) were used in this study. To generate positive control samples of muscle injury, 100 μl of cardiotoxin (CTX,4 0.05 mg/ml; Sigma) was injected into the gastrocnemius in 12-week-old mice, and the gastrocnemius was excised 3 days after the injection.

Voluntary Wheel Running and Tissue Collection—Adult and aged mice were divided into a control group and runner group. Runner group mice were housed individually in cages (32 × 35 × 23 cm) equipped with a running wheel (16 cm in diameter) and performed voluntary wheel running for 4 weeks beginning at 8 weeks or 23 months of age. Wheel running activity was monitored continuously with an original program, which was used in our previous studies (24, 25). Daily and total running distances were recorded for each mouse throughout the duration of exercise. Each day at noon, running data were collected and recorded on the computer for each mouse, and the daily distance counter was reset. Control group mice were housed in cages without a running wheel. After 4 weeks of exercise, mice were sacrificed by cervical dislocation. For RNA isolation or protein extraction, the gastrocnemius muscles were dissected out quickly from each mouse. Each sample was then frozen in liquid nitrogen after measurement of the wet weight and kept in a freezer at −80 °C until homogenization. For immunohistochemistry, the chests of sacrificed mice were opened, and their tissues were fixed by circulation of PBS and then 4% paraformaldehyde throughout the body. The gastrocnemius muscles were then dissected from each mouse, and the samples were placed in 4% paraformaldehyde until analysis.

Immunohistochemical Analysis—Fixed samples were placed in a 30% sucrose solution and stored in the refrigerator overnight. Cross-sections were cut to a thickness of 200 μm using a microtome (ROM-380, Yamato Kohki) and stored in tissue collection medium (25% glycerin, 30% ethylene glycol, 0.05 M phosphate) at −20 °C until analysis. We verified that there were no differences between the sections cut after fixation and the frozen sections (data not shown). For pretreatment for the staining of 5-bromo-2′-deoxyuridine (BrdU), sections were incubated in 50% formaldehyde for 90 min at 65 °C. After washing with Tris-buffered saline containing 0.25% Triton X-100 (TBS-T), sections were placed in 1 N HCl for 15 min at 37 °C. For secondary detection, Cy3-conjugated donkey anti-rabbit, -rat, or -goat IgG (1:500, Jackson Immunoresearch), Cy5-conjugated donkey anti-rat IgG (1:500, Jackson Immunoresearch), and AlexaFluor 488-conjugated donkey anti-mouse or -rabbit IgG (1:500, Invitrogen) were used with DAPI (Wako, Osaka, Japan), and sections were incubated overnight at 4 °C. After several careful washes, sections were placed on glass slides and enclosed using

4 The abbreviations used are: CTX, cardiotoxin; GSK-3β, glycogen synthase kinase β; HDAC1, histone deacetylase 1; Lef, lymphoid enhancer factor; p70S6K, p70 S6 kinase; TCF, T-cell factor; qRT, quantitative RT; IHC, immunohistochemistry.
was determined by the $\Delta \Delta Ct$ method and normalized to the expression of GAPDH. We confirmed that there were no differences between the data determined by the $\Delta \Delta Ct$ method and the standard curve method (data not shown).

### Protein Extraction and Western Blot Analysis

—Each tissue was homogenized in lysis buffer (50 mM Heps, pH 7.4, 150 mM NaCl, 10 mM EDTA, 10 mM NaF, 10 mM Na3P2O7, 2 mM Na2VO4, 1% sodium deoxycholate, 1% Nonidet P-40, 0.2% SDS) with protease inhibitor mix (4-(2-aminoethyl)benzenesulfonyl fluoride, aprotinin, E-64, leupeptin hemisulfate monohydrate, bestatin, pepstatin A; Nacalai Tesque, Kyoto, Japan) on ice. Homogenates were then centrifuged at 1,770 × g for 10 min at 4 °C, and the supernatants were collected. After measurement of the protein concentrations using a BCA protein assay kit (Thermo Fisher Scientific, Yokohama, Japan), samples were normalized to 2 µg/µl with SDS-PAGE loading buffer (62.5 mM Tris-HCl, pH 6.8, 2% w/v SDS, 10% glycerol, 50 mM DTT, 0.01% w/v bromphenol blue). Protein extractions were run on SDS-polyacrylamide gels (SuperSep Ace, Wako) and transferred to polyvinylidene difluoride membranes. Membranes were blocked in Blocking One (Nacalai Tesque) for 1 h at room temperature. The primary antibodies used were obtained from Cell Signaling Technology (Danvers, MA) and included the following: rabbit anti-Akt (1:1,000); rabbit anti-phospho-Akt (Ser-473; 1:1,000); rabbit anti-p70S6K (Thr-389; 1:1,000); rabbit anti-phospho-p70S6K (Thr-389; 1:1,000); rabbit anti-Akt (1:1,000); rabbit anti-Myf5 (1:200); rabbit anti-MyoD (1:200), and rabbit anti-GAPDH (1:2,000). Membranes were incubated overnight at 4 °C. Anti-rabbit IgG, HRP-linked whole anti-donkey (1:20,000, GE Healthcare), was used as a secondary antibody, and membranes were incubated for 1 h at room temperature. After careful washes in TBS containing 0.05% Tween 20, membranes were soaked in Pierce Western Thermo blotting substrate (Thermo Fisher Scientific), and the bands were visualized using a chemiluminescence system and an LAS-3000 Mini luminescent image analyzer (FUJIFILM, Tokyo, Japan). Images for each membrane were analyzed using ImageJ software (National Institutes of Health; rsbweb.nih.gov) as described previously (26). Average and standard deviations were calculated based on the signal intensities from each experiment. Myf5, MyoD, and β-catenin immunoreactivity was normalized to that of GAPDH.

### Chromatin Immunoprecipitation (ChiP) Assay

—Approximately 20–50 µg of DNA from each muscle sample was used to perform chromatin immunoprecipitation (ChiP). The following primary antibodies were used for the ChiP assay: rabbit antibody to β-catenin (Sigma), rabbit antibody to histone deacetylase 1 (HDAC1, Millipore, Billerica, MA), rabbit antibody to LEF1 (Millipore, rabbit antibody to TCF), and mouse antibody to heterochromatin protein 1 (HP1) (Millipore). Normal rabbit IgG was used as a negative control in the analysis. Membranes were incubated overnight at 4 °C. Anti-rabbit IgG, HRP-linked whole anti-donkey (1:20,000, GE Healthcare), was used as a secondary antibody, and membranes were incubated for 1 h at room temperature. After careful washes in TBS containing 0.05% Tween 20, membranes were soaked in Pierce Western Thermo blotting substrate (Thermo Fisher Scientific), and the bands were visualized using a chemiluminescence system and an LAS-3000 Mini luminescent image analyzer (FUJIFILM, Tokyo, Japan). Images for each membrane were analyzed using ImageJ software (National Institutes of Health; rsbweb.nih.gov) as described previously (26). Average and standard deviations were calculated based on the signal intensities from each experiment. Myf5, MyoD, and β-catenin immunoreactivity was normalized to that of GAPDH.

### Table 1

| Target gene | Sequence (5’–3’) | Product length |
|-------------|-----------------|----------------|
| GAPDH       | Forward          | 157 bp         |
|             | Reverse          |                |
| IL-6        | Forward          | 92 bp          |
|             | Reverse          |                |
| TNF-α       | Forward          | 105 bp         |
|             | Reverse          |                |
| Pax7        | Forward          | 196 bp         |
|             | Reverse          |                |
| Myf5        | Forward          | 141 bp         |
|             | Reverse          |                |
| MyoD        | Forward          | 140 bp         |
|             | Reverse          |                |
| Myog        | Forward          | 110 bp         |
|             | Reverse          |                |
| Heyl        | Forward          | 165 bp         |
|             | Reverse          |                |
| DLL4        | Forward          | 179 bp         |
|             | Reverse          |                |
| Wnt3        | Forward          | 151 bp         |
|             | Reverse          |                |
| Wnt5a       | Forward          | 164 bp         |
|             | Reverse          |                |
| Wnt5b       | Forward          | 185 bp         |
|             | Reverse          |                |
| TGF-β       | Forward          | 138 bp         |
|             | Reverse          |                |
| Col1a2      | Forward          | 199 bp         |
|             | Reverse          |                |
| Col1a3      | Forward          | 149 bp         |
|             | Reverse          |                |
| Fln1        | Forward          | 88 bp          |
|             | Reverse          |                |
| Postn       | Forward          | 147 bp         |
|             | Reverse          |                |
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ChIP was performed using a commercial kit (Millipore) according to the manufacturer's recommendations. Briefly, chromatin was pre-cleared with a protein A-agarose slurry with salmon sperm DNA by incubation for 1 h at 4 °C on a rotating platform in a volume of ~1 ml with ChIP dilution buffer (0.01% SDS, 1%, Triton X-100, 1.2 mm EDTA, 16.7 mm Tris-HCl, pH 8.1, and 167 mm NaCl). After agarose removal by centrifugation, 1% of the pre-cleared chromatin was saved and used as input DNA. Antibodies were added to the chromatin and incubated at 4 °C with rotation for 1 h. Immunoprecipitation was performed with 1.0 μg of antibodies. Protein A-agarose was added and incubated at 4 °C with rotation for 2 h. Three separate washes with low salt immune complex wash buffer (0.1% SDS, 1% Triton X-100, 2 mm EDTA, 20 mm Tris-HCl, pH 8.1, and 150 mm NaCl) were then performed, followed by separate washes for 15 min with rotation with the following buffers: high salt immune complex wash buffer (0.1% SDS, 1% Triton X-100, 2 mm EDTA, 20 mm Tris-HCl, pH 8.1, and 500 mm NaCl); LiCl salt immune complex buffer (0.25 m LiCl, 1% Igepal-CA630, 1% deoxycholic acid (sodium salt), 1 mm EDTA, and 10 mm Tris, pH 8.0); and two separate washes with TE buffer (10 mm Tris-HCl, 1 mm EDTA, pH 8.0). Chromatin-DNA complexes were eluted (0.1 mm NaHCO3, 1% SDS) from agarose beads, and cross-links were reversed by incubation at 65 °C for 4 h. Samples were RNase-treated (RNase A); protein was digested (proteinase K), and DNA was purified using a spin column (Wizard DNA clean-up system, Promega). Immunoprecipitated DNA for specific genes was analyzed and quantified by PCR. PCR primers used with ChIP samples were as follows: Myf5 promoter 2.4 kb upstream (forward, 5’-TGTGTGGACAAGGTGTTAATTA-3’; reverse, 5’-CACCACCTTCCAGAGTAA-3’); MyoD promoter 2.7 kb upstream (forward, 5’-CTTTCACACTTCCCCACCTGAATG-3’; reverse, 5’-AGGGACATTAAGATCGCTTT-3’). The number of PCR cycles and amount of ChIP DNA were adjusted so that the accumulated product was in the linear range of the exponential curve of the PCR amplifications. PCR products were separated by electrophoresis on agarose gels and stained with Gel Green (Biotium Inc.). The ultraviolet light-induced fluorescence of stained DNA was captured by a Dolphin-View2 imaging system (Kurabo, Osaka, Japan), and intensities of the signals were quantified by ImageJ software, as described previously (26). For each threshold, multiple independent ChIP experiments were performed. Average and standard deviations were calculated based on the signal intensities from each experiment.

**Statistical Analysis**—Experiments were analyzed for statistical significance using a Student's t test, with all error bars expressed as means ± S.D. Differences with p values of less than 0.05 were considered significant.

**RESULTS**

**Voluntary Wheel Running Provided Mild Stimulation for the Gastrocnemius Muscle**—Adult and aged mice performed voluntary wheel running for 4 weeks (Fig. 1A). The running distances were ~13 km/day for adult mice and ~4 km/day for aged mice. There were no changes in body mass or wet weight of gastrocnemius muscles after 4 weeks of running in both adult and aged mice (Fig. 1B). In previous studies of high intensity exercise, such as functional overload leading to muscle hypertrophy, Akt/mammalian target of rapamycin/p70S6K signaling is activated in skeletal muscle to increase the levels of cellular protein synthesis (27, 28). In contrast, in voluntary wheel running, a low intensity exercise without excessive stress, the activation levels of Akt/mammalian target of rapamycin/p70S6K signaling components were relatively low (Fig. 1C). Akt and p70S6K phosphorylation did not change after running, indicating that voluntary wheel running was only a mild physiological stimulus in this study.

Several studies have demonstrated that high intensity exercise also induces muscle inflammation involving acute muscle regeneration, as shown by increases in tumor necrosis factor-α (TNF-α) and interleukin-6 (IL-6) mRNA expression in skeletal muscle (29–32). However, in this study, activation of these typical inflammatory signals and cytokines was not detected, and TNF-α and IL-6 mRNA expression levels were normal. Although TNF-α and IL-6 mRNA expression levels were slightly up-regulated after exercise in both adult and aged runner mice, the fold change was significantly lower than that of the positive control, in which up-regulation of inflammatory cytokines was induced via muscle injury (i.e. injection of CTX; Fig. 1D). These results indicated that voluntary wheel running induced neither dynamic changes in the activities of protein synthesis signaling nor acute exercise-associated inflammation in both adult and aged skeletal muscle after 4 weeks of running.

**Voluntary Wheel Running Activated satellite cells and proliferation in both Adult and Aged Skeletal Muscle**—Satellite cells express Pax7 as a master transcription factor in the skeletal muscle lineage. Unlike the quiescent population, activated satellite cells co-express Myf5 and MyoD together with Pax7. Although it is well known that the number of satellite cells increases following wheel running in adult skeletal muscle, detailed characterization of the stages of these induced satellite cells and parallel examinations in both adult and aged animals have not been investigated. Our study indicated that the number of Pax7+ cells increased following wheel running in both adult and aged mice (data not shown), being consistent with a previous report (21). The number of Pax7+ cells increased more than 3-fold in adult mice and 2-fold in aged mice following 4 weeks of voluntary wheel running. Notably, in the case of aged mice (2 years old), exercise rescued the decrease in the number of Pax7+ cells during aging, and the number of increased Pax7+ cells was almost the same as that in adult mice at 12 weeks of age. Because Pax7+ cells include both quiescent and activated satellite cell populations, immunohistochemical analysis using muscle cross-sections was performed to determine how the satellite cell population changes after running. In adult mice, although Pax7+ MyoD+ -activated satellite cells were rarely observed in the control group, Pax7+MyoD+ -activated satellite cells clearly appeared in the runner group (Fig. 2A). Similarly, in aged mice, activated satellite cells significantly increased in the runner group (Fig. 2B). The number of Pax7+MyoD+ cells increased following running by nearly 6-fold, even in aged mice. Our immunohistochemistry (IHC) analysis suggested that mild running exercise effectively increased the number of satellite cells and that the increased population was predominantly characterized by activated satellite cells rather than quiescent satellite cells.
We next examined the effects of exercise on BrdU incorporation to satellite cells in skeletal muscle using Pax7+/Myf5+/BrdU staining. The mitotic marker BrdU was administered daily during the last week of the exercise period. The number of BrdU+ cells increased more than 2-fold after running in adult and aged mice (data not shown), indicating that exercise stimulated cellular proliferation, consistent with the results of a previous study (33). Importantly, the proportion of Pax7+/Myf5+/BrdU+ and Pax7+/MyoD+/BrdU+ cells to total Pax7+ cells significantly increased after running (Fig. 2, C and D). Because BrdU+ cells

![Image](https://example.com/image.png)
indicate proliferating and newly generated cell populations, Pax7+ Myf5+ BrdU+ cells and Pax7+ MyoD+ BrdU+ cells represented proliferating activated satellite cells.

Because of the finding of de novo proliferative population within activated satellite cells during IHC analysis, we performed qRT-PCR analysis to confirm and compare the expression levels of the stem cell marker genes Pax7, Myf5, and MyoD using skeletal muscle-derived total RNAs in samples from both adult and aged mice. Unexpectedly, the mRNA expression levels of Pax7 were not altered in the adult mouse group (Fig. 2E,
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Voluntary Wheel Running Activated Canonical Wnt/β-Catenin Signaling—To examine the molecular mechanisms underlying satellite cell activation after running, we focused on the Notch and Wnt signaling pathways, which regulate satellite cell self-renewal and myogenesis at the developmental embryonic stage (34). The expression of Delta ligand 4 (DLL4) and HeyL, which are the ligand and target gene, respectively, in Notch signaling, tended to decrease slightly after exercise in both the adult and aged groups (Fig. 3A). Although Notch signaling plays a critical role in controlling the proliferation of satellite cells in their lineage, the decreased expression of the ligand and target gene in our analysis implied that the generation of activated satellite cells via voluntary exercise, and the transcriptional control of the Myf5 and MyoD genes was probably unrelated to the Notch pathway.

Wnts are extracellular ligands that modulate the cell fate choice event of various types of adult stem cells (35–37). As a paracrine factor, the stem cell niches express Wnts by responding to external stimuli (38, 39). In adult skeletal muscles following exercise, Wnt3 ligand was produced from myofibers rather than undifferentiated satellite cells (data not shown); therefore, we analyzed further the expression profiles of Wnt proteins using whole muscle tissue. Interestingly, the expression of Wnt ligands (Wnt3, Wnt5a, and Wnt5b) significantly increased after mild exercise, and their up-regulation was consistently observed in both the adult and aged runner groups (Fig. 3B). The expression of Wnt3 increased more than 3-fold in adult mice and 2-fold in aged mice after 4 weeks of voluntary wheel running, and the promotional effect was consistent with the increase in Pax7+/MyoD−-activated satellite cells in adult and aged runner mice. Wnt signaling plays an important role in further promoting myogenic differentiation in lineage-committed progenitor cells but not in undifferentiated satellite cells in which Notch signaling retains fate control under normal physiological conditions (9). In Wnt signaling, there are canonical and noncanonical signaling routes that depend on the type of Wnt protein, with Wnt3 being canonical signaling members that transduce their signals through intracellular β-catenin. Although Wnt5a and Wnt5b are included in noncanonical signaling members, Wnt5a has been reported to regulate the accumulation of β-catenin in a previous study (8). In fact, the expression of Wnt5a and Wnt5b was increased after running, consistent with Wnt3 expression (Fig. 3B). Thus, we supposed that Wnt5a and Wnt5b have the function as canonical signaling members. The noncanonical Wnt signaling pathway has been found to be associated with myogenic regulation in skeletal muscle (7).

Next, we evaluated intracellular signal transduction because the Wnt member proteins Wnt3, Wnt5a, and Wnt5b were obviously associated with the promotion of activated satellite cells upon exercise. The phosphorylation of glucose synthase kinase 3β (GSK-3β) and β-catenin was investigated by Western blot analysis (Fig. 3C). β-Catenin, which acts in the nucleus with TCF and LEF transcriptional factors, is phosphorylated by GSK-3β, which leads to its ubiquitin-dependent degradation in the absence of Wnt ligands (43). However, when Wnt binds to the frizzled receptor, GSK-3β is phosphorylated (and therefore inactivated), and β-catenin accumulates and can translocate into the nucleus to activate its target genes. The phosphorylation of GSK-3β (Ser-9) tended to increase, and the expression of β-catenin significantly increased after voluntary running in both adult and aged mice, indicating that canonical Wnt/β-catenin signaling was activated upon performing exercise. Based on these results, our data confirmed that voluntary wheel running induced the activation of satellite cells, the activation of Myf5 and MyoD transcription, the expression of canonical Wnt member proteins, and the activation of canonical Wnt/β-catenin signaling in both adult and aged skeletal muscle.

Voluntary Running-dependent Wnt Activation Did Not Induce Skeletal Muscle Fibrosis—Next, we investigated how fibrosis was related to elevated Wnt/β-catenin levels in the skeletal muscle after a voluntary running exercise. IHC analysis using antibodies against ER-TR7, a specific marker of reticular fibroblasts and reticular fibers, and the plasma membrane marker dystrophin showed that there were no significant differences between the control group and runner group in both

**FIGURE 2. Activation of satellite cells following voluntary wheel running.** A and B, immunohistochemistry analysis of satellite cells following exercise. Representative merged images of immunohistochemistry staining for MyoD (red) and Pax7 (green) with DAPI from adult (A) and aged mice (B) are shown. Left panels are from the control group, and right panels are from the runner group. The proportions of Pax7+ MyoD− cells per total Pax7+ cells are shown in the right graph. White arrows indicate Pax7+ MyoD− cells. C, representative merged images of immunohistochemistry staining for Pax7 (red), Myf5 (green), and BrdU (blue) from the adult runner group. Magnification of the area surrounded by the dashed square is shown in the right panels, and white arrows indicate Pax7+ MyoD− BrdU+ cells. D, proliferative activated satellite cells increased following exercise. The proportion of Pax7+ MyoD− BrdU+ cells per total Pax7+ cells in each group was plotted. E and F, expression profiles of stem cell markers of satellite cells. The mRNA expression levels of satellite cells markers (Pax7, left; Myf5, center; MyoD, right) in gastrocnemius were detected by qRT-PCR (E). The protein expression levels of Myf5 (left graph) and MyoD (right graph) in gastrocnemius were detected by Western blot. The right images represent the typical blot patterns of Myf5 (upper), MyoD (middle), and GAPDH (lower) (F). In all graphs, signals are represented by white bars (controls) in cont (adult and aged mice), gray bars (adult runner (Run) mice), and black bars (aged runner mice). The mRNA and protein values were normalized to that of GAPDH and then plotted as the expression ratio relative to the control sample in adult mice. All data are expressed as the mean ± S.D. (n = 4/group). Significant differences are as follows: *, compared with adult control group; **, compared with aged control group (p < 0.05).
adult and aged mice (Fig. 4, A and B). IHC analysis using antibodies against periostin (Postn), a matricellular fibrotic protein, and MHC2a provided similar results, with no fibrotic features in all skeletal muscle tissues examined (Fig. 4 C). Activated satellite cells first differentiate into myoblasts and then into myocytes (myogenin-positive cells), finally giving rise to myofibers (expressing MHC). Wheel running usually produces a shift from type 2b (glycolytic fiber) MHC isoforms to type 2a (oxidative fiber) MHC isoforms in skeletal muscle (19). Clear changes were observed in MHC2a$^{\text{H11001}}$ fibers in the runner group, with an increase observed in the MHC2a$^{\text{H11001}}$ population following wheel running in both adult and aged skeletal muscle, indicating the occurrence of exercise-mediated oxidative metabolism (Fig. 4 C). However, the expression of Postn was unchanged after running, suggesting that fibrosis was not promoted by exercise.

Further IHC analysis using antibodies against collagen 1 showed no differences between the control and runner groups in adult and aged mice (data not shown). Additionally, qRT-PCR analysis of the genes associated with muscle fibrosis (i.e. TGF-β, collagen 1α2 (Col1a2), collagen 5α3 (Col5a3), fibronectin (Fn1), and Postn) confirmed that elevated Wnt expression in the skeletal muscle due to exercise was not correlated with fibrosis because almost no changes were observed in the expression levels of fibrosis-associated genes (Fig. 4D). Taken together, these data demonstrated that activation of Wnt/β-catenin signaling after voluntary running did not cause fibrosis, even in aged skeletal muscle.

**Up-regulated Wnt/β-Catenin Signaling Regulated Satellite Cell Activation**—Our analysis indicated that 4 weeks of voluntary wheel running efficiently promoted the expression of Wnt3, Wnt5a, and Wnt5b, even in aged skeletal muscle (Fig. 3B and C). This was supported by the increased expression of Delta ligand 4 (DLL4) and HeyL mRNAs (Fig. 3A). Western blot analysis revealed an increase in the phosphorylation of GSK-3β (Ser-9) and the accumulation of β-catenin in muscle tissues from exercised mice compared to controls (Fig. 3C). These findings suggests that Wnt/β-catenin signaling is upregulated following voluntary wheel running, which may promote satellite cell activation and muscle repair.
Upon phosphorylation of GSK-3β, high levels of accumulated β-catenin were detected in the skeletal muscles of mice that underwent exercise (Fig. 3B). Although recent studies have implicated Wnt signaling in abnormal repair and fibrogenesis (10), almost no stimulation of fibrosis was observed in runner group samples at both the mRNA and protein levels (Fig. 4).

To better understand the regulatory mechanisms linking the observed activation of the satellite cell population and the elevated Wnt signaling in runner group mice, we next tried to determine the direct target genes of the Wnt signaling pathway following voluntary wheel running. Because cooperative increases in the expression levels of Wnts, Myf5, and MyoD and the appearance of activated Pax7+Myf5+ and Pax7+MyoD+ satellite cells were observed, we simply examined whether the activator complex associated with canonical Wnt signaling bound the promoters of MyoD and Myf5 genes. First, we investigated 5 kb upstream of the Myf5 and MyoD genes of mice, rats, and humans. As shown in Fig. 5A, many consensus

FIGURE 4. Voluntary wheel running did not induce muscle fibrosis. A and B, representative images of immunohistochemistry analysis of ER-TR7 (red) and dystrophin (green) in adult (A) and aged mice (B). Left panels are from the control group, and right panels are from the runner group. C, representative images of immunohistochemistry analysis of periostin (red) and MHC2a (green). The upper panels represent adult mice, and the lower panels represent aged mice (upper panels represent the control group, and lower panels represent the runner group, respectively). D, qRT-PCR analysis of Wnt3, myogenin (Myog), and fibrotic mRNAs (TGF-β, collagen 1α2 (Col1α2), collagen 5α3 (Col5α3), fibronectin (Fn1), and periostin (Postn)). The mRNA expression was normalized to that of GAPDH and then plotted as the expression ratio of each gene relative to the control sample in the adult group. White bars, controls in adult and aged mice; gray bars, adult runner mice; black bars, aged runner mice. All data are expressed as the mean ± S.D. (n = 4/group). Significant differences are as follows: *, compared with adult control group; **, compared with aged control group (p < 0.05). Scale bars, 20 μm in all panels.
sequences of the TCF/LEF regulatory element were present in the mouse, rat, and human Myf5 promoter regions. Interestingly, a sequence (AACAACAACA) containing multiple TCF/LEF regulatory elements and highly conserved in mice, rats, and humans was found 2.5 kb upstream in the Myf5 promoter region (Fig. 5A and supplemental Fig. S1). Furthermore, a similar conserved sequence (ATTCCAAATTTCG) was also found in the mouse, rat, and human MyoD promoters (supplementary Fig. S2). The conserved sequence 2.7 kb upstream of the MyoD gene involved the TCF/LEF regulatory element (Fig. 5B). These findings implied that Wnt signaling retained an indispensable role in the transcriptional regulation of the Myf5 and MyoD genes.

ChIP-PCR analysis was performed to examine the effect of exercise-dependent increases in Wnt expression on the chromatin-mediated regulation of the Myf5 and MyoD genes (Fig. 5, C and D). In the dissected skeletal muscle of adult runner mice, the association of HDAC1 with the TCF/LEF regulatory elements of the Myf5 promoter decreased (>80%). In aged runner mice, the association of HDAC1 further decreased (>90%). A similar dissociation of HP1, a reader of repressive methylation at the level of histone H3 lysine 9, was detected on the Myf5 promoter in adult and aged skeletal muscle. Histone methylation is known to be associated with both transcriptionally active and repressive chromatin states. The enrichment of dimethylated H3 lysine 9 (K9me2) on the Myf5 promoter disappeared after exercise in both adult and aged mice, suggesting that a repressive chromatin state controlled the aged stage. In the dissected skeletal muscle of runner mice, dimethylated histone H3 K4 (K4me2) and acetylated histone H3 (Ac-H3), modified histones associated with gene activation, were apparently enriched on the Myf5 promoter compared with those in control mice. The associations of β-catenin, LEF1, and TCF with the conserved regulatory sequences on the Myf5 promoter were enriched upon voluntary exercise, indicating that the β-catenin-TCF/LEF-activated complex associated in response to Wnt signaling.

Similar results were observed with MyoD. MyoD was expressed in activated satellite cells, as was Myf5, and both basic helix-loop-helix transcription factors retained similar epigenetic profiles following exercise, implying that running stimulation altered the chromatin structure of MyoD and Myf5 genes, even in aged skeletal muscle. The enhancement of β-catenin-TCF/LEF binding to Wnt signaling regulatory sites on the Myf5 promoter was higher than that of the MyoD promoter, suggesting that the binding efficiency reflected the structure of the six TCF/LEF regulatory sites on the mouse Myf5 promoter and of the two TCF/LEF regulatory sites within the detection site used in our ChIP analysis. These results suggested that voluntary wheel running increased the de novo expression of Wnt3 in adult and aged mice and thereby rescued impaired myogenesis in aged mice via the canonical Wnt signaling-mediated transcriptional activation of the Myf5 and MyoD genes, important targets that drive the state of activated satellite cells. Associations of β-catenin, TCF, LEF1, K4me2, and Ac-H3 increased upon exercise, whereas K9me2, HDAC1, and HP1 were dissociated from the Wnt signaling response elements on the Myf5 and MyoD promoters after exercise, suggesting that an active chromatin state controlled myogenesis in adult and aged skeletal muscle upon exercise. Taken together, these findings supported the conclusion that exercise stimulus-associated Wnt expression contributed to genetic chromatin reassembly in adult myogenesis rather than to fibrosis and promoted myogenic proliferation and exit from the quiescent states of satellite cells, which was critical for exercise-dependent myogenesis in adult and aged skeletal muscle.

DISCUSSION

In this study, we investigated the characteristic changes in satellite cells and the molecular mechanisms regulating their activation in mouse skeletal muscle following voluntary wheel running. Our analysis showed that exercise induced the up-regulation of several Wnts in myofibers (Fig. 3) and that activation of the canonical Wnt/β-catenin signaling pathway increased the expression of Myf5 and MyoD in satellite cells (Fig. 2, A–D) through association of the β-catenin-TCF/LEF activator complex with the Myf5 and MyoD promoters (Fig. 5). Myf5 is expressed in only Pax7+ cells, satellite cells but not in myofibers, and MyoD is also expressed in satellite cells or differentiating myoblasts (Fig. 2) (44). Therefore, we focused to assess the regulatory mechanism of Myf5 and MyoD genes in skeletal muscle. In the case of adult neurogenesis, astrocyte cells residing near adult neural stem cells produce Wnt3 ligands (35) and the secreted extracellular Wnts trigger the neurogenesis by promoting the expression of NeuroD basic helix-loop-helix transcription factor (38, 39). Because Wnt3 expression in adult brain flexibly changes by responding to external stimuli such as running exercise (45), stem cell niches act as a sensor to modulate the ability of adult neural stem cells. In the case of adult skeletal muscle, a similar relationship between satellite cells (trigger myogenesis via the basic helix-loop-helix myogenic genes) and residing myofibers (Wnt production) were observed in the current study. Wnt-mediated expression of myogenic genes in satellite cells caused their conversion from the quiescent state into the activated state in both adult and aged mice. Wnt signaling plays an important role in embryonic muscle development (7, 34). The neural tube expresses Wnt1, which activates the Myf5 gene, whereas the dorsal ectoderm expresses Wnt7a, which promotes MyoD expression (46). Although Wnt retains its necessary role in muscle development during the embryonic stage (47, 48), it also contributes to postnatal/adult myogenesis in the muscle regeneration process after injury. In adult skeletal muscle, CTX-induced injury increases the expression of Wnt5a, Wnt5b, and Wnt7a mRNA, and the induction is accompanied by the generation of Myf5+ cells (49). In a study using in vitro myoblast cultures, the up-regulation of myogenin and MHC, mature stage markers, was seen on treatment with lithium chloride (LiCl), a GSK-3β inhibitor that activates Wnt signaling (50). Treatment of myoblasts with Wnt3a also promoted the progression of differentiation accompanied by accelerated expression of desmin, a cytoskeletal intermediate filament protein (9). Our study demonstrated that the exercise-mediated activation of Wnt signaling affected satellite cells in the first step of adult myogenesis rather than lineage progression into the mature stage. The expression levels of Wnt3, Wnt5a, and Wnt5b increased upon voluntary wheel running. These findings indicated the critical role of Wnt proteins in...
Exercise-stimulated Wnt signaling directly triggered the expression of Myf5 and MyoD transcripts in adult/aged skeletal muscle. 

A, schematic representation of Wnt/β-catenin regulatory sites on the mouse, rat, and human Myf5 promoters. The region about 5 kb upstream of the Myf5 gene is shown with the TCF/LEF consensus sequence. Each yellow square box represents a TCF/LEF regulatory element. A highly conserved sequence within species (* indicates the location), AACAAACAAA, was located 2.5 kb upstream of the Myf5 gene, and the conserved sequence contained multiple TCF/LEF regulatory elements.

B, schematic representation of Wnt/β-catenin regulatory sites on the mouse, rat, and human MyoD promoters. The region about 5 kb upstream of the MyoD gene is shown with the TCF/LEF consensus sequence. A highly conserved sequence within species (* indicates the location), ATTTCAAATTTTGC, was located 2.7 kb upstream of the MyoD gene, and the conserved sequence involved the TCF/LEF regulatory element.

C, ChIP analysis of the mouse Myf5 promoter in adult/aged skeletal muscle after voluntary wheel running. PCR primers were designed to surround the TCF/LEF regulatory elements that are conserved within species at 2.5 kb upstream of the Myf5 gene. ChIP results obtained by independent replicate experiments are represented as percentages of the input signal (signal relative to input (%)). Signals are represented by white bars (controls in adult and aged mice), gray bars (adult runner mice), and black bars (aged runner mice).

D, ChIP analysis of the mouse MyoD promoter in adult/aged skeletal muscle after voluntary exercise. PCR primers were designed to surround the TCF/LEF regulatory element contained within the conserved sequence among mouse, rat, and human MyoD promoters at the 2.7-kb upstream locus.
adult myogenesis. Among the various Wnt proteins, specific isoforms are activated in response to different treatments (e.g., drugs, reagents, and stimuli) and stimuli (e.g., exercise, injury, and surgery altering the serum environment); the signaling pathways also differ between canonical and noncanonical signaling.

The positive role of Wnt signaling in adult skeletal muscle differentiation under various types of exercise has been described in several recent studies. Acute treadmill running and functional overload decreases GSK-3β activity and activates canonical Wnt signaling in skeletal muscle (22). Following functional overload, β-catenin, frizzled receptor, and LEF1 mRNA expression increased in the plantaris muscle (23). Moreover, the number of β-catenin-expressing M-cadherin-positive cells increases upon functional overload (51). These exercise types mimic the high intensity resistance exercise required for strengthening muscles. Acute high intensity exercises, such as jogging and running, cannot be performed easily, especially in aged individuals, where the risk of sarcopenia increases. In the case of voluntary wheel running, the exercise intensity is low and less stressful than that of other activities. Similar to a study in which high intensity exercise was enforced (22, 23), the voluntary wheel running used in this study as mild exercise up-regulated the expression of Wnt isoforms effectively in both adult and aged skeletal muscle.

However, in aged mice, the exercise-mediated up-regulation of Wnts and the activation of the canonical Wnt/β-catenin signaling pathway cannot be simply accepted as clinically beneficial because elevated Wnt signaling also stimulates fibrosis (10, 52). A previous parabiosis mouse study reported a decrease in GSK-3β and an increase in β-catenin and axin 2 (downstream targets of Wnt signaling), indicating that the up-regulation of Wnt signaling in aged mice induces fibrosis rather than triggering myogenic differentiation (10). Moreover, Brack et al. (10) reported that most of the satellite cells isolated from aged mice differentiate into fibroblasts, whereas those from young mice do not. Scimè et al. (53) reported that the expression of Wnt inhibitors, such as secreted frizzle-related protein 2 (sFRP2) and Wnt inhibitor factor 1 (Wif1), decreases with age, suggesting a decrease in Wnt antagonism during aging. These data suggest that increased Wnt signaling during aging potentially induces the transdifferentiation of satellite cells into a fibroblastic lineage. However, in this study, both IHC and qRT-PCR analyses showed that there were no differences in the levels of ER-TR7, periostin, or collagen 1 expression between adult and aged skeletal muscle. ER-TR7 is a marker of reticular fibroblasts and reticular fibers and is generally used to detect fibrosis (10, 40). Periostin is a fibrotic protein retaining the role of critical regulator of fibrosis and is located in the extracellular matrix of skeletal muscle and fibroblasts (41). Knock-out of periostin suppresses muscular dystrophy-induced fibrosis (42). Importantly, although voluntary exercise induced significant up-regulation of Wnts and activation of the canonical Wnt/β-catenin signaling pathway in our study, in turn increasing the expression of the Myf5 and MyoD myogenic genes, exercise-induced up-regulation of Wnt signaling was not involved in muscle fibrosis. Therefore, we can speculate the following: 1) exercise-responding Wnts inhibit fibrosis; 2) artificial surgery exposing the aged environment to young skeletal muscle may cause unexpected regulatory responses that do not reflect the normal action of Wnt and related molecules; and 3) other factor(s) co-exist to cause distinct phenotypes under canonical Wnt signaling depending on the physiological conditions (exercise versus inflammation, healthy subjects versus muscle dystrophy/disease, and low stress condition versus stressful condition).

In conclusion, this study revealed the regulatory mechanisms underlying Wnt-mediated myogenesis in satellite cells after exercise. The voluntary wheel running used in this study as mild exercise up-regulated canonical Wnt/β-catenin signaling in skeletal muscle. The expression levels of Wnt3, Wnt5a, and Wnt5b significantly increased after 4 weeks of voluntary running in both adult and aged mice, and the elevated Wnts and downstream signaling cascade did not induce fibrosis, even in aged skeletal muscle. The canonical Wnt/β-catenin signaling pathway directly modulated the chromatin structures of both the MyoD and Myf5 genes and activated their expression at the transcriptional level. Exercise-stimulated chromatin remodeling on the MyoD and Myf5 promoters altered the state of satellite cells from quiescent to activated, resulting in increased numbers of proliferative Pax7+ MyoD+ cells. The associations of β-catenin, TCF, and LEF transcription factors on multiple TCF/LEF regulatory elements, conserved in mouse, rat, and human species, on the promoters of both the Myf5 and MyoD genes drove myogenesis in satellite cells, even in aged muscle, suggesting that exercise-stimulated extracellular Wnts played a critical role in the regulation of satellite cells in adult and aged skeletal muscle. Our study suggested that mild running was effective in preventing the progression of sarcopenia because exercise-induced Wnts did not cause fibrosis in aged skeletal muscle and rescued the activity of satellite cells. Further studies aiming to discriminate between the effects of Wnts in the exercise state and fibrotic state and for determining the modulator(s) controlling signaling are crucial for understanding the functions and intrinsic ability of satellite cells in adult myogenesis.

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