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Potential involvement of dietary advanced glycation end products in impairment of skeletal muscle growth and muscle contractile function in mice

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Abstract

Diets enriched with advanced glycation end products (AGE) have recently been related to muscle dysfunction processes. However, it remains unclear whether long-term exposure to an AGE-enriched diet impacts physiological characteristics of skeletal muscles. Therefore, we explored the differences in skeletal muscle mass, contractile function and molecular responses between mice receiving a diet high in AGE (H-AGE) and low in AGE (L-AGE) for 16 weeks. There were no significant differences between L-AGE and H-AGE mice with regard to body weight, food intake or epididymal fat pad weight. However, extensor digitorum longus (EDL) and plantaris (PLA) muscle weights in H-AGE mice were lower compared with L-AGE mice. Higher levels of N\textsuperscript{ε}(carboxymethyl)-L-lysine, a marker for AGE, in EDL muscles of H-AGE mice were observed compared with L-AGE mice. H-AGE mice showed lower muscle strength and endurance in vivo and lower muscle force production of PLA muscle in vitro. mRNA expression levels of myogenic factors including myogenic factor 5 and myogenic differentiation in EDL muscle were lower in H-AGE mice compared with L-AGE mice. The phosphorylation status of 70-kDa ribosomal protein S6 kinase Thr\textsuperscript{389}, an indicator of protein synthesis signalling, was lower in EDL muscle of H-AGE mice than that of L-AGE mice. These findings suggest that long-term exposure to an AGE-enriched diet impairs skeletal muscle growth and muscle contractile function, and that these muscle dysfunctions may be attributed to the inhibition of myogenic potential and protein synthesis.

Key words: Muscle strength: Muscle fatigue resistance: Muscle force production: Myogenesis: Protein synthesis

Skeletal muscles, which account for approximately 40\% of total body mass, are the largest tissue in the body and play a crucial role in locomotion and metabolism. Both skeletal muscle mass and function are regulated in response to changes in nutrition, physical activity, environment and pathological conditions. Maintenance of muscle mass and function is an important public health issue, as low muscle quality is associated with clinical problems including poor glycaemic control, limited mobility and even mortality\textsuperscript{1,2}.

Advanced glycation end products (AGE), which are produced by a non-enzymatic reaction between glucose and proteins, are closely involved in the pathogenesis of diabetes\textsuperscript{3,4}, Alzheimer disease\textsuperscript{5}, cancer\textsuperscript{6} and hypertension\textsuperscript{7}. Moreover, elevated AGE in blood or skin are related to low muscle mass, grip strength and walking speed in elderly and diabetic patients\textsuperscript{8-11}. Thus, a negative relationship between AGE and skeletal muscle function has been suggested.

AGE are endogenously formed but are also absorbed from exogenous sources such as foods, especially when prepared at elevated temperatures\textsuperscript{12}. As dietary AGE are poorly absorbed (approximately 10\%)\textsuperscript{13}, their potential action in human health has gone largely unnoticed. However, recent systematic reviews of randomised-controlled trials have shown that exposure to AGE-enriched diets may be responsible for the development of chronic diseases such as diabetes, CVD and chronic kidney disease through increased inflammation and oxidative stress\textsuperscript{14}, and that dietary restriction of AGE alleviates insulin resistance, oxidative stress and endothelial dysfunction\textsuperscript{15}. In addition, an AGE-enriched diet induced insulin resistance in mouse skeletal muscle\textsuperscript{16}. Thus, we hypothesised that dietary AGE might impair skeletal muscle function. However, whether dietary AGE affect physiological characteristics of skeletal muscle is not yet fully understood. Therefore, our aim was to evaluate the effect of long-term exposure to AGE-enriched diets on skeletal muscle function.

Abbreviations: AGE, advanced glycation end products; CML, N\textsuperscript{ε}(carboxymethyl)-L-lysine; EDL, extensor digitorum longus; H-AGE, high in AGE; L-AGE, low in AGE; LC3, microtubule-associated protein light-chain 3; p70S6K, 70-kDa ribosomal protein S6 kinase; Pax7, paired box 7; PLA, plantaris; MHC, myosin heavy chain; mTOR, mammalian target of rapamycin; MyoD, myogenic factor 5; Myf5, myogenic differentiation; SOL, soleus.

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exposure to an AGE-enriched diet on skeletal muscle mass and contractile functions.

Myogenesis plays an important role in postnatal growth and skeletal muscle development. Sequential activation and expression of paired box 7 (Pax7) and myogenic regulatory factors including myogenic factor 5 (Myf5), myogenic differentiation (MyoD) and myogenin are essential for the initiation and progression of myogenesis.[21]. An in vitro study revealed that AGE treatment for 48h suppressed myogenin expression in C2C12 myotubes.[18]. Further, an in vitro study showed that long-term intake of a high-fructose diet decreased MyoD expression in mouse skeletal muscle, which was attenuated by supplementation with pyridoxamine, an AGE formation inhibitor.[20]. The receptor for AGE also regulates muscle differentiation through Pax7 and myogenin.[20]. These studies suggest that AGE impact the expressions of myogenic factors and inhibit myogenesis.

Skeletal muscle growth occurs when the rate of protein synthesis exceeds that of degradation. A central signalling pathway regulating protein synthesis is the Akt/mammalian target of rapamycin (mTOR)/70-kDa ribosomal protein S6 pathway regulating protein synthesis.[21] In addition, a recent study demonstrated that AGE treatment was reported to induce autophagy by suppressing the Akt/mTOR pathway in neonatal cardiomyocytes.[24]. In vitro study revealed that AGE inhibition decreased the phosphorylation status of Akt at Ser473, which is a key enzyme in the ubiquitin–proteasome system, in C2C12 myotubes.[18]. These findings suggest that AGE inhibit protein synthesis as well as promote protein degradation in skeletal muscle. Therefore, our secondary aim was to evaluate the influence of dietary AGE on the regulatory systems of myogenesis and protein turnover in skeletal muscle.

We examined the differences in muscle mass, contractile properties and molecular responses between mice that received a diet high in AGE (H-AGE) or one low in AGE (L-AGE) for 16 weeks. We found that N-(carboxymethyl)-L-lysine (CML), a well-characterised compound that serves as a marker for AGE,[23] accumulated in fast-type extensor digitorum longus (EDL) muscle, and EDL muscle weight was lower in H-AGE mice compared with L-AGE mice. The contractile properties were also poorer in H-AGE mice. Furthermore, mRNA expressions of myogenic factors and phosphorylation status of mTOR/p70S6K signalling pathway were lower in EDL muscles of H-AGE mice compared with L-AGE mice.

Methods

Animals and treatment

Male ICR mice (5 weeks old) were purchased from Shimizu Breeding Laboratories (Kyoto, Japan). Mice were placed in a room maintained at 22–24°C with a 12 h light–12 h dark cycle and fed a standard diet (AIN-93G; Oriental Yeast) and water ad libitum. After 1 week of adjustment, mice were randomised to receive a diet either low in AGE (AIN-93G unbaked, n 10) or high in AGE (AIN-93G baked for 1 h at 160°C, n 10) consistent with previous studies.[26–28]. The H-AGE diet contained five times as much AGE compared with the L-AGE diet without changes in nutrient composition.[28]. Both groups were followed-up for 16 weeks. Body weight and food intake were measured every week and every 2 weeks, respectively. At the end of the study period, animals were fasted for 4 h and anaesthetised intraperitoneally with mixtures of medetomidine hydrochloride (0.3 mg/kg), midazolam (4.0 mg/kg) and butorphanol (5.0 mg/kg). Under anaesthesia, soleus (SOL), EDL and plantaris (PLA) muscles, tibial, epididymal fat pads and blood samples were collected. All animal protocols were carried out in accordance with the Guide for the Care and Use of Laboratory Animals by the National Institutes of Health (Bethesda, MD, USA) and were approved by the Kyoto University Graduate School of Human and Environmental Studies.

Grip strength test

The grip strength test was performed 5d before the end of the 16-week study. Grip strength was measured using a force transducer (DS2-50N; IMADA) according to the published protocol (DMD_M.2.2.001). In brief, mice were allowed to rest on a horizontal mesh with their forelimbs and hindlimbs (four paws) and were then gently pulled back until their grip was broken. Three consecutive measurements were obtained within 1 min, and were averaged to determine mean grip strength. Data were normalised to body weight and expressed as N/g.

Wire-hanging test

To evaluate muscle fatigue resistance, the wire-hanging test was carried out 4d before the end of the 16-week study consistent with the published protocol.[29]. In brief, mice were suspended by their forelimbs from a metallic wire (2-mm thick, 50-cm long) suspended 40 cm above soft ground. As soon as the mice were suspended, the timer was started. The timer was stopped when the animal fell and re-started after it was placed again on the wire. The number of falls from the wire was recorded, and the test was stopped after ten falls or 180 s of suspension. Data were expressed as the average fall score: each animal was given a score of 10 at the beginning of the test, which was reduced by 1 after each fall. The average fall score was calculated at a given time during the test as \( \frac{10n - x}{n} \), where \( n \) is the number of animals and \( x \) the cumulative number of falls. The longest holding time was also recorded, and the holding impulse was calculated as \( s \times g \), where \( s \) is the longest holding time and \( g \) the body weight.

In vitro muscle contraction

In vitro muscle contraction was analysed as described previously.[30,31]. Isolated PLA muscles were mounted on an incubation apparatus with tension set to 0·5 g and rested in Krebs–Ringer bicarbonate buffer (117-mM NaCl, 4·7-mM KCl, 2·5-mM CaCl₂, 1·2-mM KH₂PO₄, 1·2-mM MgSO₄, 24·6-mM NaHCO₃ containing 2-mM pyruvate for 30 min. All buffers were continuously bubbled with 95% O₂–5% CO₂ and maintained at 37°C. For measuring force production, muscles were stimulated using
an electric stimulator (SEN-3401; Nihon Kohden) at frequencies of 5, 10, 25, 50, 75 and 100 Hz (train duration: 1 s; pulse duration: 0.1 ms; voltage: 10 V) with a 2-min rest period between contractions(32). Subsequently, for measuring muscle fatigability, muscles were stimulated at 100 Hz, 1 s on 1 s off, for 30 s. Force was monitored using a force transducer (TRN001; Kent Scientific) and recorded (U-228-2P-500; Pantos). Force production data were normalised to muscle weight and expressed as N/g.

Real-time RT-PCR analyses

Real-time RT-PCR analyses were performed as previously described(33). In brief, total muscle RNA was extracted using a miRNeasy Mini kit (Qiagen) according to the manufacturer's protocol. RNA was reverse-transcribed to complementary DNA (cDNA) using PrimeScript RT Master Mix (Takara Bio); synthesised cDNA was then subjected to real-time RT-PCR (StepOne, Applied Biosystems) using Takara SYBR Premix Ex Taq II (Takara Bio). The relative fold change of expression was calculated by the comparative CT method using StepOne software version 2.3. To normalise the amount of total RNA present in each reaction, glyceraldehyde 3-phosphate dehydrogenase was used as an internal standard. The primer sequences are shown in Table 1.

Measurement of myosin heavy chain isoform composition

Analysis of myosin heavy chain (MHC) isoform composition (I, IIa, IIx and IIb) was carried out using a modification of previously described methods(34). In brief, muscles were homogenised in ice-cold lysis buffer (1:40, v/v) containing 20-mM Tris-HCl (pH 7.4), 1% Triton X, 50-mM NaCl, 250-mM sucrose, 50-mM NaF, 5-mM sodium pyrophosphate, 2-mM dithiothreitol, 4-mg/l leupeptin, 16000 µg/l trypsin inhibitor, 0.1% Triton X, 50-mM NaCl, 250-mM sucrose, 50-mM NaF, 5-mM sodium pyrophosphate, 2-mM dithiothreitol, 4-mg/l leupeptin, 50-mg/l trypsin inhibitor, 0.1-mM benzamidine and 0.5-mM phenylmethylsulphonyl fluoride (buffer A) and centrifuged at a constant speed of 16000 x g for 40 min at 4°C. Extracted proteins were solubilised in Laemmli sample buffer and boiled for 5 min. The sample proteins were separated by SDS-PAGE using 10% polyacrylamide gel at a constant current of 40 mA/gel for 90 min. After SDS-PAGE, the proteins were transferred to polyvinylidene fluoride membranes (Hybond-P, GE Healthcare) by using trans-blot cell (Bio-Rad) at a constant voltage of 100 V for 1 h at 4°C. After the transfer, the membranes were blocked for 1 h using fish gelatin (Takara Bio). Next, the membranes were incubated overnight at 4°C with primary antibodies: Akt (9272; Cell Signaling Technology), Akt Ser473 (9271; Cell Signaling Technology), p70S6K Thr389 (9206; Cell Signaling Technology), p70S6K Thr389 (9206; Cell Signaling Technology) and microtubule-associated protein light-chain 3 (LC3) (2775; Cell Signaling Technology). The membranes were washed with Tris-buffered saline with 0.1% Tween 20 (TBS-T, pH 7.5) and then allowed to react with anti-rabbit IgG (Cell Signaling Technology) for 1 h at room temperature. After the final wash with TBS-T, protein bands were visualised using chemiluminescence (Wako). Signal density was measured using Light-Capture (AE-6971; ATTO Corporation).

Western blot analyses

Western blot analysis was performed as described previously(35,36). Extracted samples in buffer A were solubilised in Laemmli’s sample buffer and boiled for 5 min. The samples (10-µg of protein) were separated by SDS-PAGE using 10% polyacrylamide gel at a constant current of 40 mA/gel for 90 min. After SDS-PAGE, the samples were transferred to polyvinylidene fluoride membranes and blocked for 1 h using fish gelatin (Takara Bio). Next, the membranes were incubated overnight at 4°C with primary antibodies: Akt (9272; Cell Signaling Technology), Akt Ser473 (9271; Cell Signaling Technology), p70S6K Thr389 (9206; Cell Signaling Technology), p70S6K Thr389 (9206; Cell Signaling Technology) and microtubule-associated protein light-chain 3 (LC3) (2775; Cell Signaling Technology). The membranes were washed with Tris-buffered saline with 0.1% Tween 20 (TBS-T, pH 7.5) and then allowed to react with anti-rabbit IgG (Cell Signaling Technology) for 1 h at room temperature. After the final wash with TBS-T, protein bands were visualised using chemiluminescence (Wako). Signal density was measured using Light-Capture (AE-6971; ATTO Corporation).

Measurement of Nε-(carboxymethyl)-L-lysine content

The muscle sample was homogenised in buffer A, and the supernatant was collected after centrifugation at 16000 g for 40 min at 4°C. Blood samples were collected from the carotid artery into tubes containing heparin as anticoagulant, and plasma was separated by centrifugation at 8000 g for 15 min. The content of CML in muscles and plasma was measured using a CircuLex CML/Nε-(carboxymethyl) lysine ELISA kit (Medical & Biological Laboratories) according to the manufacturer's protocol. Protein concentration of the supernatant of muscles and plasma was determined using the Bradford technique, and the CML content in samples was expressed as µg/mg protein.

Statistical analyses

Values are expressed as means with their standard errors. Statistical significance was assessed using parametric Student’s t test or non-parametric Mann–Whitney U test (Tables 2–4 and Fig. 1) and two-way ANOVA with diet and frequencies (Fig. 2) or diet and muscles (Fig. 3 and Fig. 4) as main factors. In the event of significant main effects and/or interactions, post hoc Tukey–Kramer tests were performed. Differences between groups were considered statistically significant at P < 0.05. The sample size was calculated by power analysis at 0.8 and a level of significance of 0.05 with a hypothesised effect size of 1.2 and a level of significance of 0.05 with a hypothesised effect size of 1.2

Table 1. List of primer sequences (forward, reverse) used for real-time RT-PCR analyses

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<tr>
<th>Protein (gene)</th>
<th>Sequences (5' → 3')</th>
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<tr>
<td>Pax7 (Pax7)</td>
<td>ATTACGGCGAGTGTCAAGAATCAA</td>
</tr>
<tr>
<td>Myf5 (Myf5)</td>
<td>GAGGTCCGCGCTTGTCAATCCA</td>
</tr>
<tr>
<td>MyoD (Myod1)</td>
<td>CTTATCAAGCCTGCGCGCGAGA</td>
</tr>
<tr>
<td>Myogenin (Myogenin)</td>
<td>GACAAGCAGCTCAAGCTGGAACA</td>
</tr>
<tr>
<td>MuRF1 (Trim63)</td>
<td>GGACCCAGAGAATGGTGGATAG</td>
</tr>
<tr>
<td>Atrogen-1 (Fbxo32)</td>
<td>ATCATGAAGTGCCTGCTGTCACCA</td>
</tr>
<tr>
<td>GAPDH (Gapdh)</td>
<td>TGTGCGGATGATGCGGTTCA</td>
</tr>
</tbody>
</table>

Pax7, paired box 7; Myf5, myogenic factor 5; MyoD, myogenic differentiation; MuRF1, muscle RING finger 1; GAPDH, glyceraldehyde 3-phosphate dehydrogenase.

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previous study\(^{57}\), which estimated the sample size to be 10. All statistical analyses were performed using Ekusen-Toukei 2012 software (Social Survey Research Information).

**Results**

**Dietary advanced glycation end products suppressed skeletal muscle growth**

The characteristics of mice exposed to L-AGE and H-AGE are shown in Table 2. There were no significant differences between L-AGE and H-AGE mice in body weight, tibial length, food intake and epididymal fat pad weight. EDL muscle weight was lower in H-AGE mice compared with L-AGE mice when normalised to body weight and tibial length. PLA muscle weight was lower in H-AGE mice when normalised to tibial length and tended to be lower when normalised to body weight \((P=0.096)\). There were no significant differences in SOL muscle weight between L-AGE and H-AGE mice.

**Dietary advanced glycation end products increased \(N\)-carboxymethyl-L-lysine accumulation in skeletal muscle**

To examine the effect of the H-AGE diet on muscle performance, we assessed muscle strength and fatigue resistance in vivo. The grip strength test was used to measure muscle strength. Grip strength in all four limbs in H-AGE mice was weaker than in L-AGE mice (Fig. 1(a)). To measure fatigue resistance, we carried out the wire-hanging test. The average fall score during the wire-hanging test was 3-7 at the end of the experiment in L-AGE mice, whereas the score fell faster in H-AGE mice and reached 1-4 (Fig. 1(b)). The longest holding time during the wire-hanging test was shorter by 30 % in H-AGE mice compared with L-AGE mice; holding impulse was significantly lower in H-AGE mice than in L-AGE mice (Fig. 1(c)).

**Dietary advanced glycation end products deteriorated muscle force production in vitro**

We next measured force production in skeletal muscle \textit{in vitro} to determine the effect of H-AGE on muscle contractile property. PLA muscle of H-AGE mice was found to produce significantly lower force at 75 and 100 Hz compared with L-AGE mice (Fig. 2(a)). There were no significant differences in the patterns of force decline between L-AGE and H-AGE mice (Fig. 2(b)).

**Dietary advanced glycation end products did not affect myosin heavy chain isoform composition in skeletal muscle**

As muscle fibre type composition is a determinant of muscle strength and fatigue resistance, we evaluated the effect of H-AGE on MHC isoform composition in skeletal muscles. No significant differences were observed in MHC isoform composition in either SOL or EDL muscles between L-AGE and H-AGE mice (Table 4).

**Dietary advanced glycation end products decreased mRNA expressions of myogenic factors in skeletal muscle**

To determine the effect of H-AGE on regulatory factors related to skeletal muscle growth, we evaluated the expression levels of mRNA involved in myogenesis. Myf5 mRNA expression level was significantly decreased in EDL muscle of H-AGE mice (Fig. 3). In addition, MyoD mRNA expression levels were lower in H-AGE mice compared with L-AGE mice. Pax7 and myogenin mRNA expression levels did not differ significantly based on an H-AGE diet.

**Dietary advanced glycation end products suppressed protein synthesis signalling in skeletal muscle**

To examine the effect of H-AGE on protein synthesis, we evaluated the phosphorylation status of Akt/mTOR/p70S6K.

**Table 2. Characteristics of low in advanced glycation end products (L-AGE)- and high in AGE (H-AGE)-fed mice**

(Mean values with their standard errors; \(n=10\) per group)

<table>
<thead>
<tr>
<th></th>
<th>L-AGE Mean</th>
<th>L-AGE SE</th>
<th>H-AGE Mean</th>
<th>H-AGE SE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial body weight (g)</td>
<td>31.5 0.8</td>
<td>31.5 0.8</td>
<td>31.5 0.8</td>
<td>31.5 0.8</td>
</tr>
<tr>
<td>Final body weight (g)</td>
<td>53.6 2.5</td>
<td>53.6 2.5</td>
<td>59.7 4.0</td>
<td>59.7 4.0</td>
</tr>
<tr>
<td>Tibial length (cm)</td>
<td>2.01 0.02</td>
<td>2.06 0.02</td>
<td>2.01 0.02</td>
<td>2.06 0.02</td>
</tr>
<tr>
<td>Food intake (kJ/d)</td>
<td>79.1 1.7</td>
<td>76.2 1.7</td>
<td>78.2 1.7</td>
<td>78.2 1.7</td>
</tr>
<tr>
<td>Food intake (kcal/d)</td>
<td>18.9 0.4</td>
<td>18.7 0.4</td>
<td>18.7 0.4</td>
<td>18.7 0.4</td>
</tr>
<tr>
<td>Tissue weight normalised to tibial length (mg/cm)</td>
<td>5.8 0.2</td>
<td>5.5 0.2</td>
<td>5.5 0.2</td>
<td>5.5 0.2</td>
</tr>
<tr>
<td>SOL</td>
<td>7.8 0.3</td>
<td>7.0 0.2</td>
<td>7.0 0.2</td>
<td>7.0 0.2</td>
</tr>
<tr>
<td>EDL</td>
<td>12.3 0.3</td>
<td>11.4* 0.4</td>
<td>11.4* 0.4</td>
<td>11.4* 0.4</td>
</tr>
<tr>
<td>PLA</td>
<td>107.7 137-4</td>
<td>1215.3 136-4</td>
<td>1215.3 136-4</td>
<td>1215.3 136-4</td>
</tr>
<tr>
<td>Epididymal fat pad weight</td>
<td>0.22 0.01</td>
<td>0.20 0.01</td>
<td>0.20 0.01</td>
<td>0.20 0.01</td>
</tr>
<tr>
<td>EDL</td>
<td>0.30 0.02</td>
<td>0.25* 0.02</td>
<td>0.25* 0.02</td>
<td>0.25* 0.02</td>
</tr>
<tr>
<td>PLA</td>
<td>0.46 0.01</td>
<td>0.41 0.03</td>
<td>0.41 0.03</td>
<td>0.41 0.03</td>
</tr>
<tr>
<td>Epididymal fat pad weight</td>
<td>39.7 4.4</td>
<td>41.6 3.84</td>
<td>41.6 3.84</td>
<td>41.6 3.84</td>
</tr>
</tbody>
</table>

\(\text{SOL, soleus; EDL, extensor digitorum longus; PLA, plantaris.}\)

\(^{*}P<0.05\) v. L-AGE.
signalling pathway. The phosphorylation status of Akt at Ser473 was not affected by the H-AGE diet, but the expression of phosphorylated p70S6K Thr 389, an indicator of mTOR activity, was lower in H-AGE mice compared with L-AGE mice (Fig. 4).

Dietary advanced glycation end products had no effect on protein degradation pathways in skeletal muscle

To analyse the effect of H-AGE on protein degradation, we evaluated the expression levels of molecules involved in autophagy and ubiquitin–proteasome systems. The relative expression levels of LC3II/LC3I, an indicator of autophagy activity, were not affected by the H-AGE diet (Fig. 5). There were also no differences between L-AGE and H-AGE mice with regard to mRNA expressions of atrogin-1 and muscle RING finger 1 (MuRF1), another key enzyme of the ubiquitin–proteasome system (Fig. 5).

Table 4. Myosin heavy chain (MHC) isoform composition (Mean values with their standard errors; n 10 per group)

<table>
<thead>
<tr>
<th></th>
<th>SOL</th>
<th></th>
<th>EDL</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>MHC I (%)</td>
<td>62.0</td>
<td>2.0</td>
<td>60.3</td>
<td>1.5</td>
</tr>
<tr>
<td>MHC IIAx (%)</td>
<td>38.0</td>
<td>2.0</td>
<td>39.7</td>
<td>1.5</td>
</tr>
<tr>
<td>MHC IIB (%)</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>MHC I (%)</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>MHC IIAx (%)</td>
<td>32.7</td>
<td>0.9</td>
<td>33.6</td>
<td>1.4</td>
</tr>
<tr>
<td>MHC IIB (%)</td>
<td>67.3</td>
<td>0.9</td>
<td>66.4</td>
<td>1.4</td>
</tr>
</tbody>
</table>

L-AGE, low in advanced glycation end products; H-AGE, high in advanced glycation end products; SOL, soleus; EDL, extensor digitorum longus.
production, without changes in MHC isom form composition. Third, mRNA expressions of myogenic factors, Myf5 and MyoD, in skeletal muscle were lower in H-AGE mice compared with L-AGE mice. Fourth, phosphorylation status of p70S6K Thr^{389} was lower in H-AGE mice than in L-AGE mice.

Recently, an epidemiological study has suggested a relationship between AGE accumulation and skeletal muscle mass in humans\cite{11}. In addition, an experimental study showed that AGE treatment in cultured muscle cells suppressed MyoD and promoted atrophy\cite{18}. That study also revealed that skeletal muscle atrophy in diabetic mice was attenuated by treatment with alagebrum chloride, a breaker of AGE-based cross-links\cite{18}. These observations indicate that AGE are potent factors that influence muscle mass. In the present study, H-AGE mice showed smaller muscle masses of EDL and PLA compared with L-AGE mice (Table 2), which was accompanied by elevated CML content (Table 3). This is the first report of long-term exposure to an AGE-enriched diet promoting AGE accumulation in skeletal muscles and impaired skeletal muscle growth.

AGE are also related to muscle dysfunction. It has been reported that elevated blood CML is associated with poor grip strength\cite{39} and slow walking speed\cite{39}. Furthermore, men with higher skin AGE levels had lower muscle strength and power\cite{10}. In the present study, we found that grip strength and wire-holding performance were poorer in H-AGE mice compared with L-AGE mice (Fig. 1). In addition, we showed that in vitro muscle force production normalised to muscle weight was lower in H-AGE mice than in L-AGE mice (Fig. 2(a)). These findings indicate that dietary AGE affect muscle contractile properties in addition to muscle mass.

In this regard, some reports have shown that myofibrillar proteins are subject to glycation modification. Glycation of actomyosin increased with ageing in mice and rats\cite{42}. Glycation of myosin altered filament formation\cite{40}, structure and ATPase activity\cite{41,42}. AGE-modified actin is also accumulated with ageing\cite{42}. Therefore, the modification of muscle

**Discussion**

The results of the present study revealed several novel findings about the effects of dietary AGE on skeletal muscle functions. First, mice fed an H-AGE diet for 16 weeks exhibited smaller muscle mass than L-AGE mice, which was accompanied by CML accumulation in skeletal muscle. Second, exposure to H-AGE induced skeletal muscle dysfunction, including muscle strength, fatigue resistance and in vitro muscle force production, without changes in MHC isom form composition. Third, mRNA expressions of myogenic factors, Myf5 and MyoD, in skeletal muscle were lower in H-AGE mice compared with L-AGE mice. Fourth, phosphorylation status of p70S6K Thr^{389} was lower in H-AGE mice than in L-AGE mice.

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contractile proteins such as myosin and actin is a possible mechanism inducing the loss of muscle contractile properties.

Muscle strength is also affected by muscle fibre type composition\(^{44}\). The fibre types based on MHC isoforms essentially reflect muscle contractile properties\(^{45}\). In the present study, MHC isoform composition did not differ between L-AGE and H-AGE mice (Table 4), indicating that dietary AGE-induced muscle dysfunction is not attributed to the alteration of muscle fibre type composition.

In the present study, dietary AGE affected muscle mass more in fast-type muscles, EDL and PLA, than in slow-type muscles, SOL (Table 2). This indicates that the susceptibility to AGE might be dependent on fibre type. Indeed, CML accumulated in EDL muscle but not in SOL muscle during the 16-week treatment period (Table 3). In addition, previous reports have shown that AGE accumulation in diabetic rat skeletal muscle was greater in fast-type fibres\(^{46}\). These findings suggest that dietary AGE promote AGE accumulation preferentially in fast-type fibres of whole-body muscle, and thereby induce decreased muscle performance. However, some reports have shown that the presence of AGE did not differ between fibre types in diabetic\(^{47}\) or ageing rats\(^{48}\). Thus, additional studies are required to elucidate any dependency on fibre type for AGE influence.

Postnatal skeletal muscle growth is regulated by the sequential expressions of myogenic factors such as Pax7, Myf5, MyoD and myogenin. Pax7 is the specific marker for quiescent and activated muscle satellite cells, which are the resident skeletal muscle stem cells\(^{49,50}\). Myf5 is expressed during embryonic development and integrates various developmental signalling pathways to initiate myogenesis\(^{51}\). MyoD is considered a master control gene for MyoD\(^{52}\); myogenin participates in later stages of myogenesis and is directly associated with the differentiation process that leads to myotube formation and maturation\(^{53,54}\). In the present study, Myf5 mRNA expression level in EDL muscle was lower in H-AGE mice than in L-AGE mice (Fig. 3). Furthermore, MyoD mRNA expression level was suppressed in H-AGE mice compared with L-AGE mice. These findings suggest that dietary AGE inhibit muscle formation during a relatively early stage of myogenesis, resulting in muscle growth deterioration.

MyoD is also a crucial modulator of muscle fibre type. MyoD binds to an enhancer box (E-box) element in the MHC promoter and induces its transcription\(^{55}\). In particular, MyoD expression is high in fast-type muscle and is involved in the induction of fast-type fibre phenotypes\(^{56}\). Over-expression of MyoD has been shown to up-regulate the expression of fast-type MHC\(^{57}\). Conversely, MyoD-knockout mice showed a shift towards slower fibre phenotype in fast-type muscles\(^{56,58}\). These observations raise the possibility that the down-regulation of MyoD mRNA expression in H-AGE mice induced fibre-type transition towards a slower muscle phenotype. However, dietary AGE did not affect MHC isoform composition in the present study (Table 4). In this respect, some studies have reported inconsistent shifts in MHC with MyoD expressions\(^{59,60}\). Thus, the alteration of MyoD expression may not always be linked to the shift in MHC isoform distributions.

In addition to myogenesis, protein synthesis and subsequent accumulation of protein are necessary for skeletal muscle growth. A previous study indicated that AGE induced myotube atrophy via inhibition of the Akt pathway\(^{18}\). In the present study, the phosphorylation status of p70S6K Thr\(^{389}\) was lower in EDL muscle of H-AGE mice compared with L-AGE mice, although no change was observed in Akt Ser\(^{473}\) phosphorylation (Fig. 4). These results suggest that dietary AGE affect the mTOR/p70S6K signalling pathway independently of Akt in skeletal muscle. On the other hand, AGE may possibly modulate protein degradation as noted by previous studies, which have shown that AGE treatment in cultured skeletal and cardiac muscles activated autophagy and ubiquitin–proteasome pathways\(^{18,24}\). However, no influences on the ratio of LC3II:Lc3I and mRNA expressions of atrogin-1 and MuRF1 were observed (Fig. 5), which suggests that dietary AGE do not affect protein degradation. Therefore, it is suggested that dietary AGE deteriorate skeletal muscle growth by inhibiting protein synthesis.

AGE are generally found in several foods prepared at high temperatures\(^{12}\). Therefore, animal models exposed to heat-treated foods are often used to evaluate the influence of AGE in vitro\(^ {16,26–28,61–65}\). However, a limitation in using these models is that the heat applied may produce unexpected products in diets. From this perspective, our study cannot exclude the possibility that products other than AGE may affect muscle dysfunction. Therefore, further investigations using inhibitors of AGE accumulation should be conducted to understand the direct links between AGE and muscle functions.

In conclusion, in the present study, we showed that exposure to an AGE-enriched diet for 16 weeks promoted CML accumulation in skeletal muscle. Furthermore, we observed impairment in skeletal muscle functions including skeletal muscle mass, muscle strength and fatigue resistance in H-AGE mice compared with L-AGE mice. AGE-enriched diet-associated skeletal muscle dysfunction might be due to a failure of postnatal myogenesis and protein synthesis. The present study shows a possibility that long-term exposure to an AGE-enriched diet impairs postnatal growth and skeletal muscle development.

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