Research Article

Treatment with Herbal Formula Extract in the hSOD1\textsuperscript{G93A} Mouse Model Attenuates Muscle and Spinal Cord Dysfunction via Anti-Inflammation

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Amyotrophic lateral sclerosis (ALS), a multicomplex neurodegenerative disease, has multiple underlying pathological factors and can induce other neuromuscular diseases, leading to muscle atrophy and respiratory failure. Currently, there is no effective drug for treating patients with ALS. Herbal medicine, used to treat various diseases, has multitarget effects and does not usually induce side effects. Each bioactive component in such herbal combinations can exert a mechanism of action to increase therapeutic efficacy. Herein, we investigated the efficacy of an herbal formula, comprising Achyranthes bidentata Blume, Eucommia ulmoides Oliver, and Paeonia lactiflora Pall., in suppressing the pathological mechanism of ALS in male hSOD1\textsuperscript{G93A} mice. Herbal formula extract (HFE) (1 mg/g) were orally administered once daily for six weeks, starting at eight weeks of age, in hSOD1\textsuperscript{G93A} transgenic mice. To evaluate the effects of HFE, we performed footprint behavioral tests, western blotting, and immunohistochemistry to detect protein expression and quantitative PCR to detect mRNA levels in the muscles and spinal cord of hSOD1\textsuperscript{G93A} transgenic mice. To evaluate the effects of HFE, we performed footprint behavioral tests, western blotting, and immunohistochemistry to detect protein expression and quantitative PCR to detect mRNA levels in the muscles and spinal cord of hSOD1\textsuperscript{G93A} transgenic mice. To evaluate the effects of HFE, we performed footprint behavioral tests, western blotting, and immunohistochemistry to detect protein expression and quantitative PCR to detect mRNA levels in the muscles and spinal cord of hSOD1\textsuperscript{G93A} transgenic mice. To evaluate the effects of HFE, we performed footprint behavioral tests, western blotting, and immunohistochemistry to detect protein expression and quantitative PCR to detect mRNA levels in the muscles and spinal cord of hSOD1\textsuperscript{G93A} transgenic mice. To evaluate the effects of HFE, we performed footprint behavioral tests, western blotting, and immunohistochemistry to detect protein expression and quantitative PCR to detect mRNA levels in the muscles and spinal cord of hSOD1\textsuperscript{G93A} transgenic mice.

1. Introduction

Amyotrophic lateral sclerosis (ALS) is a neurodegenerative disease with various underlying pathological factors, including genetic and environmental factors that induce other neuromuscular diseases, leading to muscle atrophy and respiratory failure. The known pathological mechanisms of ALS are mitochondrial dysfunction, oxidative stress, and neuroinflammation in the glial cells [1]. Activated astrocytes and microglia release proinflammatory cytokines and toxic factors that contribute to neurotoxicity [2], and energy metabolism plays a key role in driving the onset and progression of ALS [3]. In preclinical studies, drugs have targeted these mechanisms for treating ALS; however, most drugs targeting a single process have been unsuccessful in clinical trials [4]. Therefore, there is an urgent need for an effective cure for ALS. Patients with ALS and their families have a poor quality of life. Moreover, only a few drugs, such as riluzole and edaravone, can delay death by 3–4 months. ALS results from several pathological dysfunctions, including metabolic dysfunction in the spinal cord and muscle, and therefore, new therapies and drug discoveries should be aimed at multiple targets in the muscles and spinal cord [5, 6].
Herbal medicines have been used to treat various diseases because they rarely induce side effects and have multitarget effects [7]. In ALS, conventional drugs, such as riluzole and edaravone, which are mostly ineffective and cause side effects, do not greatly prolong the survival of patients; therefore, herbal medicines have been evaluated for use in patients with ALS [8]. Each component of this herbal combination may use a different mechanism of action, thus increasing therapeutic efficacy. According to our previous study, treatment with Bojungikgi-tang showed neuroprotective effects and delayed disease progression in an ALS animal model [9]. *Achyranthes bidentata* Blume strengthens bones and muscles and protects against N-methyl-D-aspartate-induced excitotoxicity in hippocampal neurons [10] and nerve crush injury in mice [11]. *Eucommia ulmoides* Oliver improves metabolic functions by decreasing ATP levels and increasing the use of ketone bodies and glucose in the skeletal muscle [12]. *Paecenia lactiflora* Pallas exerts protective, analgesic, anti-inflammatory, and immunomodulatory effects in vitro and in vivo [13]. Since ALS is a complex disease caused by muscle dysfunction and loss of motor neurons, we examined the effects of an herbal medicine formula containing *A. bidentata* Blume, *E. ulmoides* Oliver, and *P. lactiflora* Pallas to harness the multitarget advantage of herbal medicine on tibialis anterior (TA), gastrocnemius (GC), and spinal cord (SP) of hSOD1<sup>G93A</sup> mice. In this study, we demonstrated that HFE treatment exhibited anti-inflammatory and antioxidative effects and enhanced autophagy dysfunction in the TA, GC, and SP of hSOD1<sup>G93A</sup> mice, suggesting that multitargeted treatment with HFE regulates the ALS-inducing pathological mechanism.

2. Materials and Methods

2.1. Animals. Male B6SJL-hSOD1<sup>G93A</sup> and female B6SJL mice were purchased from Jackson Laboratory (Bar Harbor, ME, USA) and mated to obtain hemizygous male hSOD1<sup>G93A</sup> mice. Hemizygous hSOD1<sup>G93A</sup> mice were selected as described previously [9]. Two to three mice were placed in each cage, habituated to the specific pathogen-free cages, and maintained in the specific pathogen-free animal facility at a temperature of 20 ± 2°C and humidity of 50 ± 10%, with a 12-h:12-h light:dark cycle. Food and tap water were provided *ad libitum*. Behavioral tests were performed by experimenters who were blinded to the experimental groups. All mouse experiments were approved by the Institutional Animal Care and Use Committee of the Korea Institute of Oriental Medicine (KIOM protocol # 17-061). Male hSOD1<sup>G93A</sup> mice were randomly divided into the following groups: nontransgenic mice (*nTg*), *n* = 8; hSOD1<sup>G93A</sup> transgenic mice (*Tg*), *n* = 8; and herbal formula extract (HFE)-treated hSOD1<sup>G93A</sup> transgenic mice (*Tg + HFE*), *n* = 8. HFE (1 mg/g) was orally administered once daily for six weeks, starting at eight weeks of age in hSOD1<sup>G93A</sup> transgenic mice. The *nTg* and hSOD1<sup>G93A</sup> transgenic mice were used as controls and were administered distilled water (Figure 1).

2.2. Treatment with Herbal Extract. The herbal formula composed of *A. bidentata* Blume, *E. ulmoides* Oliver, and *P. lactiflora* Pallas (1:1:1) was purchased from Kwang-myungdang Medicinal Herbs Co. (Ulsan, Republic of Korea). The HFE was prepared as previously described [14].

2.3. Footprint Test. A footprint test was conducted on the day before the mice were sacrificed to measure motor activity. The footprint test was performed as previously described [14].

2.4. Tissue Preparation and Western Blotting. After anesthe-tizing the mice with avertin (250 mg/kg), the tibialis anterior (TA), gastrocnemius (GC), and spinal cord (SP) were collected and homogenized in RIPA buffer (50 mM Tris-Cl [pH 7.4], 1% NP-40, 0.1% sodium dodecyl sulfate [SDS], and 150 mM NaCl) containing a protease and phosphatase inhibitor cocktail (Thermo Fisher Scientific, Waltham, MA, USA). Homogenized tissues were centrifuged at 13,000 rpm for 15 min at 4°C. The total protein was quantified using a bicinchoninic acid assay kit (Pierce, Rockford, IL, USA).

Protein samples for western blotting were heated with SDS sample buffer. Equal amounts of total protein were separated on a 4–12% SDS-PAGE precast gel (Thermo Fisher Scientific) and transferred onto polyvinylidene fluoride membranes. The membranes were incubated with the specific primary antibodies for 12 h at 4°C. The primary antibodies used were anti-β catenin (1:1,000; Cell Signaling Technology, Danvers, MA, USA), anti-P62 (1:1,000; Cell Signaling Technology), anti-LC3b (1:1,000; Cell Signaling Technology), anti-GFAP (1:5,000; Agilent Technologies, Santa Clara, CA, USA), anti-CD11b (1:1,000; Abcam, Cambridge, MA, USA), anti-BAX (1:1,000; Santa Cruz Biotechnology, Dallas, TX, USA), anti-HO1 (1:1,000; Abcam), anti-feritin (1:1,000; Abcam), anti-tubulin (1:1,000; Abcam), anti-ferritin (1:1,000; Abcam), anti-Iba1 (1:2,000; Fuji film Wako Chemical, Richmond, VA, USA), and anti-GFAP (1:5,000; Agilent Technologies). The membranes were then incubated with horseradish peroxidase-conjugated secondary antibodies (anti-rabbit or anti-mouse; Santa Cruz Biotechnology) and washed with TBST. The membranes were probed with enhanced chemiluminescence reagents (Thermo Fisher Scientific) and visualized using an imaging system (Bio-Rad, Hercules, CA, USA) for chemiluminescence detection. All immunoblots were quantified using the ImageJ software (National Institutes of Health, Bethesda, MD, USA).

2.5. Immunohistochemistry. The spinal cord was fixed in 4% paraformaldehyde and rinsed in 0.01 M PBS. Sectioned paraffin tissues were deparaffinized, and immunohistochemistry was performed as described previously [9]. The primary antibodies used were anti-CHAT (1:500; Thermo Scientific, Wilmington, DE, USA), anti-Iba1 (1:2,000; Fujifilm Wako Chemical, Richmond, VA, USA), and anti-GFAP (1:5,000; Agilent Technologies), and the secondary antibody was an anti-horseradish peroxidase-conjugated mouse or rabbit IgG (GenDEPOT, Katy, TX, USA). The spinal cord slides
were incubated with anti-mouse or rabbit IgG biotinylated secondary antibodies (Vector Laboratories, Burlingame, CA, USA) using diaminobenzidine (Vector Laboratories). Diaminobenzidine-stained slides were dehydrated with serial ethanol and xylene solutions and mounted with a coverslip and mounting solution. Images were captured using an Olympus BX51 microscope, and the intensity of the primary antibodies was quantified using ImageJ software.

2.6. RNA Extraction and Real-Time Reverse Transcription PCR. Total RNA was extracted from the muscle tissues using the RNA extraction kit (Intron Biotechnology, Seongnam-Si, Korea). RNA extraction and the RT-PCR were performed as previously described [15]. The primer sequences used in this study are shown in Table 1. The relative mRNA levels of the target genes were presented as fold values.

2.7. Statistical Analysis. All data are reported as the mean ± standard error (SEM). Statistical analyses were performed using GraphPad Prism 9 software (GraphPad, Inc., La Jolla, CA, USA). The results were analyzed with a one-way analysis of variance followed by Tukey’s test for multiple comparisons. Statistical significance was set at \( p < 0.05 \).

3. Results

3.1. HFE Treatment Improved Motor Activity and Reduced Inflammation-Related Protein Levels in the TA and GC of hSOD1\textsuperscript{G93A} Mice. To investigate the effect of HFE on motor activity, we measured the stride length of the mice in a footprint test. As shown in Figure 2, HFE improved motor function by 1.3-fold compared with the Tg group (\( p = 0.0049 \)).

Next, to examine the biological mechanism involved in improving motor function following HFE treatment, we evaluated the effect of HFE on inflammation and oxidative stress in the TA and GC of hSOD1\textsuperscript{G93A} mice. Levels of inflammation-related proteins such as β-catenin, CD11b, and GFAP were increased by 3.4-, 6.6-, and 2.0-fold, respectively, compared with those in the nTg group (\( p = 0.0044, p = 0.0037, p = 0.0173 \)). However, HFE treatment significantly reduced these levels by 1.9-, 2.4-, and 1.6-fold, respectively, compared with those in the Tg mice (\( p = 0.0194, p = 0.0118, p = 0.0378 \), Figures 3(a) and 3(c)).

In addition, HFE treatment reduced the mRNA level of IL18 by 1.5-fold compared with that in the TA of nTg mice (\( p = 0.0052 \), Figure 3(b)). Oxidative stress is involved in inflammation [16]. Therefore, we also determined the effect of HFE on the expression levels of oxidative stress-related proteins in the TA of the nTg, Tg, and Tg-HFE groups. The levels of oxidative stress-related proteins, including HO1, ferritin, transferrin, and Bax, were increased by 4.1-, 6-, 1.5-, and 5-fold, respectively, in the TA of the Tg groups compared with those in the nTg group (\( p = 0.0004, p = 0.0074, p = 0.0468, p = 0.0004 \), Figure 3(d)). Furthermore, HFE treatment dramatically reduced HO1, ferritin, transferrin, and Bax levels in the TA of the Tg group by 3.7-, 2.6-, 1.5-, and 1.5-fold, respectively (\( p = 0.0006, p = 0.0438, p = 0.0223, p = 0.0275 \), Figure 3(d)). Furthermore, the mRNA level of COX IV1a was significantly reduced by 1.2-fold in the TA of HFE-treated mice compared with that in Tg mice (\( p = 0.0416 \), Figure 3(e)). These findings suggest that HFE treatment improved motor activity by regulating inflammatory reactions in the skeletal muscle of hSOD1\textsuperscript{G93A} mice.

3.2. HFE Treatment Regulates Autophagy Function in TA and GC of hSOD1\textsuperscript{G93A} Mice. Autophagy dysfunction is a pathological feature of the muscles and spinal cord in ALS. In addition, because muscle atrophy is caused by autophagy dysfunction, we investigated the expression of autophagy-related proteins, such as p62 and LC3b, and atrophy-related proteins, including TGFβ and SMAD2, in the TA and GC of hSOD1\textsuperscript{G93A} mice. As shown in Figure 4, the levels of p62 and LC3b proteins were reduced by 2.0- and 1.5-fold (\( p = 0.0016, p = 0.0441 \)) in the HFE-treated TA and 1.5- and 2.0-fold in the HFE-treated GC, respectively, compared with those in Tg mice (\( p = 0.0056, p = 0.0003 \), Figures 4(a) and 4(c)).

In addition, the expression of TGFβ and SMAD2 was decreased by 2.3- and 1.7-fold (\( p = 0.0024, p = 0.0083 \)) in the HFE-treated TA and 1.8- and 1.5-fold in the HFE-treated GC (\( p = 0.0413, p = 0.0048 \), respectively, compared with those in Tg mice (Figures 4(b) and 4(e))). Furthermore, HFE treatment significantly attenuated the mRNA levels of muscle denervation-related genes, myogenin (Myog), and cholinergic receptor nicotinic gamma subunit (Chrnγ), by 1.5- and 1.3-fold (\( p = 0.0478, p = 0.025 \)), respectively, in the TA compared with those in the Tg group (Figure 4(d)).

3.3. HFE Treatment Attenuates Motor Neuronal Cell Death in the Spinal Cord of hSOD1\textsuperscript{G93A} Mice. To examine the effect of HFE on motor neuron death, we investigated the immunoreactivity of CHAT, Iba1, and GFAP in the spinal cord of hSOD1\textsuperscript{G93A} mice. The number of motor neuron cells positive for CHAT increased by 6.3-fold in the ventral horn of the spinal cord in HFE-treated mice than in hSOD1\textsuperscript{G93A} mice (\( p = 0.0189 \), Figure 5(a)). In addition, the expression
respectively, in the spinal cord of hSOD1 G93A mice compared with those in Tg mice (Figure 5(b)).

**4. Discussion**

Thus far, an effective drug that can improve the quantity of life of patients with ALS is yet to be developed. Therefore, we examined the effects of herbal medicines against the multiple pathological mechanisms of ALS, including metabolic and muscle dysfunction, inflammation, autophagy dysfunction, and oxidative stress [17–19] in the muscle and spinal cord of an ALS animal model. We found that the HFE (A. bidentata Blume, E. ulmoides Oliver, and P. lactiflora Dallas) improved muscle function by reducing inflammation and autophagy dysfunction in the muscle and spinal cord of hSOD1 G93A mice.

Neuroinflammation or inflammation is critical in neurodegenerative diseases, including Alzheimer’s disease, Parkinson’s disease, and ALS. In ALS, microglia activation and activated astrocytes are increased in the spinal cord, inducing motor neuron death [20]. In addition, neuroinflammation in hSOD1 G93A mice is involved in the upregulation of NLRP3-inflammasome-related proteins [21]. However, based on these findings, most drugs target a single mechanism, such as those with antiglutamatergic (ceftriaxone, memantine, and talampanel), anti-inflammation (celecoxib, erythropoietin, and NP001), antioxidative stress, and neurotrophic effects, have failed in clinical trials [4]. TAR DNA-binding protein 43 (TARDBP, TDP-43) and ORF 72 on chromosome 9 (C9orf72) mutation were detected in sporadic and familiar ALS cases and involved in neuroinflammation [22, 23].

ALS is associated with muscular disease, including muscle weakness, inflammation, and denervated atrophy [24]. Specifically, inflammation and oxidative stress affect muscle homeostasis and myogenesis following FOXO activation, leading to muscle atrophy. Lawler et al. showed that muscle atrophy is induced by Foxo3 activation under oxidative stress and inflammation conditions. Huang et al. also demonstrated that oxidative stress and inflammation suppression attenuated denervation-induced muscle atrophy [25].

Autophagy is important for maintaining muscle functions; however, dysfunction in autophagy induces muscle degeneration, inflammation, oxidative stress, and mitochondrial dysfunction in ALS [26–29]. In ALS, inflammatory events in the skeletal muscle induce motor weakness, neuro-muscular junction impairment, and motor neuron death. Autophagy dysfunction induces oxidative stress [30] and antioxidants through redox signaling, and the Nrf2-Keap1 pathway regulates autophagy function [31]. However, it is unclear whether autophagy activation or inactivation causes ALS. Zhang et al. reported that autophagy activation decreased the accumulation of misfolded proteins in the motor neurons of hSOD1 G93A mice [32]. However, Bhattacharya et al. showed that autophagy activation augmented motor neuron degeneration and did not extend the life span of hSOD1 G93A mice [33]. These results suggest that other mechanisms such as oxidative stress and inflammation should be considered therapeutic targets, as ALS is a complex disease. In this study, we demonstrated that HFE

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<th>Abbreviation</th>
<th>Primer</th>
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<td>Chrng</td>
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Figure 3: Herbal formula extract (HFE) reduced the expression of inflammatory proteins and oxidative stress-related proteins in the tibialis anterior (TA) and gastrocnemius (GC) of hSOD1<sup>G93A</sup> mice. (a) Representative immunoblots of inflammatory proteins and their quantification in the TA of nontransgenic (nTg), transgenic (Tg), and HFE-Tg mice. n = 3–4 per group. (b) mRNA expression level of IL18 in the TA of each group. n = 4–5 mice per group. (c) Representative blot and quantification of the level of GFAP protein in the GC of nTg, Tg, and HFE + Tg mice. n = 3–4 per group. (d) Levels of oxidative stress-related proteins (HO1, ferritin, transferrin, and BAX) were reduced by HFE treatment in the TA of hSOD1<sup>G93A</sup> mice. n = 3–4 per group. (e) mRNA expression level of COX IV 1a in the TA of each group. n = 4–5 mice per group. *p < 0.05, **p < 0.01, ***p < 0.001. nTg: nontransgenic mice, Tg: transgenic mice, Tg + HFE: HFE-treated transgenic mice.
**Figure 4:** Herbal formula extract (HFE) regulated expression of autophagy- and muscle atrophy-related proteins in hSOD1<sup>G93A</sup> mice. (a) Representative immunoblots and quantification of autophagy-related proteins (p62 and LC3b) and muscle atrophy-related proteins (TGF-β and SMAD2) in the tibialis anterior (TA) (b) and gastrocnemius (GC) (c, e) of each group. *n* = 3–4 mice per group. (d) mRNA level of muscle denervation-related genes (*Myo* and *Churg*) in each group. *n* = 3–5 per group. Data are presented as the mean ± SEM. *p* < 0.05, **p** < 0.01, ***p** < 0.001. nTg: nontransgenic mice, Tg: transgenic mice, Tg + HFE: HFE-treated transgenic mice.
Figure 5: Herbal formula extract (HFE) reduced neuroinflammation in the spinal cord of hSOD1<sup>G93A</sup> mice. (a) Representative images of ventral horn of spinal cord showing diaminobenzidine staining using anti-CHAT, anti-Iba1, and anti-GFAP. HFE administration increased CHAT-positive cells and reduced Iba1- and GFAP-positive cells in the ventral horn of the spinal cord in the Tg group. Quantification of positive intensity of primary antibody. <i>n</i> = 3–4 mice per group. Scale bars: 2 mm. (b) Representative image of SMAD2, p62, and ferritin expression detected using western blotting analysis in the nTg, Tg, and Tg-HFE groups. Tubulin was used as the loading control. Quantification of immunoblots of GFAP, SMAD2, P62, and ferritin. Data are presented as the mean ± SEM. <i>n</i> = 3–4 mice per group. *<i>p</i> < 0.05, **<i>p</i> < 0.01, ***<i>p</i> < 0.001. nTg: nontransgenic mice, Tg: transgenic mice, Tg + HFE: HFE-treated transgenic mice.
treatment exhibited anti-inflammatory and antioxidative effects and attenuated autophagy dysfunction in the TA, GC, and SP of hSOD1<sup>G93A</sup> mice, suggesting that multitargeted treatment with HFE regulates the ALS-inducing pathological mechanism.

In the skeletal muscle, denervation-induced muscle atrophy and neuromuscular junction impairments occur with motor neuron loss, and mitochondria are involved in inducing apoptosis for denervation [34]. TGF-β, which causes muscle fibrosis, Smad, and histone deacetylase 4 (HDAC4), is related to ALS progression and is upregulated in ALS [35, 36]. Furthermore, HDAC4 activates the synaptic acetylcholine receptors MuSK to promote muscle reinnervation, and HDAC4 deletion contributes to neurogenic atrophy [37]. Consistently, we found that the expression of TGF-β and Smad proteins was increased in the muscle of hSOD1<sup>G93A</sup> mice; however, treatment with HFE significantly reduced the levels of these proteins. Furthermore, denervation-related genes (Myog and Chrng) were reduced by HFE in the TA of hSOD1<sup>G93A</sup> mice. Thus, treatment with HFE may prevent muscle atrophy and delay motor neuron loss in hSOD1<sup>G93A</sup> mice.

Muscle metabolism is involved in body energy homeostasis, and denervation of muscle and atrophy is related to muscle metabolism. Therefore, metabolic dysfunction is a key factor in ALS. Dobrowony et al. reported a relationship between metabolic changes and disease progression in a hSOD1<sup>G93A</sup> mouse model [38] and suggested that the muscle is a critical therapeutic target in ALS. Muscle metabolism is related to mitochondrial function. Mitochondrial abnormalities with dysregulation of respiratory complexes, such as complexes I and IV, were observed in the muscles of patients with ALS and an hSOD1<sup>G93A</sup> mouse model [39, 40]. We previously observed that herbal medicine improved motor activity and inhibited mitochondria cristae disruption in hSOD1<sup>G93A</sup> mice [9]. Therefore, herbal medicines may improve motor function by regulating muscle metabolism via inhibition of mitochondrial dysfunction in hSOD1<sup>G93A</sup> mice.

5. Conclusions

The HFE treatment improved motor function through anti-inflammation, antioxidative, and autophagy regulation effects in the TA, GC, and SP of hSOD1<sup>G93A</sup> mice. These findings suggest that HFE are therapeutic candidates for treating patients with ALS or inhibiting disease progression. However, this study has some limitations. The effects of HFE treatment on neuromuscular junction impairment in skeletal muscles and motor neurons should be further investigated in an ALS animal model. In addition, muscle metabolism and mitochondria dysfunction in hSOD1<sup>G93A</sup> mice should be examined, as HFEs improve motor function. Chang et al. had reported that TDP43-induced aggregation was reduced by berberine, herbal medicine, by the regulation of mTOR-autophagy signaling pathway [41]. Therefore, studies are also needed to investigate the efficacy of HFE in various ALS models, such as mice modeling TDP43 and C9orf72 ALS, as this experiment was performed only in hSOD1<sup>G93A</sup> mice before it can be established that HFE are therapeutic candidates for patients with ALS.

Data Availability

All the data are available within the article.

Ethical Approval

The study was conducted according to the guidelines of the Korea Institute of Oriental Medicine, and the study was approved by the Institutional Animal Care Committee of the Korea Institute of Oriental Medicine (protocol number: 17-061).

Conflicts of Interest

The authors declare no conflict of interest.

Authors’ Contributions

EJY contributed to the study conception and design. Material preparation, data collection, and analysis were performed by EJY, SHL, and MC. The first draft of the manuscript was written by EJY and all authors commented on previous versions of the manuscript. All authors read and approved the final manuscript. Eun Jin Yang and Sun Hwa Lee contributed equally to this work.

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