Protective effects of Brazilian propolis supplementation on capillary regression in the soleus muscle of hindlimb-unloaded rats

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Abstract
The protective effects of Brazilian propolis on capillary regression induced by chronically neuromuscular inactivity were investigated in rat soleus muscle. Four groups of male Wistar rat were used in this study; control (CON), control plus Brazilian propolis supplementation (CON + PP), 2-week hindlimb unloading (HU), and 2-week hindlimb unloading plus Brazilian propolis supplementation (HU + PP). The rats in the CON + PP and HU + PP groups received two oral doses of 500 mg/kg Brazilian propolis daily (total daily dose 1000 mg/kg) for 2 weeks. Unloading resulted in a decrease in capillary number, luminal diameter, and capillary volume, and an increase in the expression of anti-angiogenic factors, such as p53 and TSP-1, within the soleus muscle. Brazilian propolis supplementation, however, prevented these changes in capillary structure due to unloading through the stimulation of pro-angiogenic factors and suppression of anti-angiogenic factors. These results suggest that Brazilian propolis is a potential non-drug therapeutic agent against capillary regression induced by chronic unloading.

Keywords Muscle atrophy · Capillary regression · Oxidative stress · Brazilian propolis · Hindlimb unloading · Anti-angiogenic factors

Introduction
The capillary network in skeletal muscle changes in response to physiological or pathological conditions. An increase in muscle activity, such as exercise [1, 2], and electrical stimulation [3] contribute to angiogenesis, while a decrease in neuromuscular activity, such as muscle unloading [4–7] and cast immobilization [8], and some diseases, such as cancer cachexia [9] and diabetes [10–12], leads to capillary regression.

Hindlimb unloading (HU) is a well-established rodent model of hypokinesia and hypodynamia [13] and results in capillary regression, especially within a muscle having a high percentage of slow oxidative fibers, such as the rat soleus muscle [4–7]. Various authors have reported that HU induces a decrease in the size and number of capillaries within the unloaded soleus muscle [14, 15]. These alterations have been attributed to a loss of endothelial cells due to apoptosis [7].

A chronic decrease in neuromuscular activity induces oxidative stress within the unloaded skeletal muscle. Oxidative stress has been reported to play a role in the upregulation of negative angiogenic regulators, such as thrombospondin-1 (TSP-1). Anti-angiogenic TSP-1 acts as angiogenic inhibitor through its anti-proliferative and pro-apoptotic effects [16–19]. The essential role of TSP-1 in capillary regression has been reviewed by Olfert et al. [20]. TSP-1 is also a key participant in capillary regression within the rat soleus muscle induced by HU [4]. Muscle capillarity is tuned by pro-angiogenic signaling, such as vascular endothelial...
growth factor (VEGF). Pro-angiogenic VEGF plays a role in angiogenesis, stimulating the formation of new vascular networks by recruiting and promoting the differentiation of endothelial cells [21]. Additionally, under the condition of muscle unloading, increased VEGF has a positive effect on angio-protection within the skeletal muscle [5, 15]. Taken together, these data suggest that pro- and anti-angiogenic factors are associated with the protection or progression of capillary regression observed in unloaded skeletal muscle.

In earlier studies, we found that supplementation with an antioxidant was an effective strategy to prevent capillary regression within unloaded muscle, with the effect achieved through modulation of the imbalance between pro- and anti-angiogenic factors [5, 6, 22]. Propolis, a substance produced by bees from the resin collected from trees and shrubs, has long been used as a non-drug therapeutic agent. It has various physiological and pathological properties, such as antiviral, antibacterial, antimutagenesis, and antioxidant effects [23]. Brazilian propolis contains flavonoids and cinnamic acid derivatives, both of which have been revealed to possess powerful antioxidant effects [24–26]. Therefore, we hypothesized that the administration of Brazilian propolis to hindlimb unloaded rats would attenuate oxidative stress within the unloaded muscle, resulting in protection of the associated capillary regression associated with muscle atrophy. The aim of this study was to investigate the effects of propolis supplementation on capillary regression within the soleus muscle of the hindlimb-unloaded rat.

Materials and methods

Experimental design

Twenty-eight adult male Wistar rats (age 13 weeks; Japan SLC, Hamamatsu, Japan) weighing between 250 and 300 g were randomly divided into four groups: (1) control (CON), (2) control plus propolis supplementation (CON + PP), (3) hindlimb unloading (HU), and (4) hindlimb unloading plus propolis supplementation (HU + PP). HU in the rats in the HU and HU + PP groups was obtained by tail suspension for 2 weeks, as previously described [13]. Briefly, after a string was fixed to the tail with tape, a swivel hook was placed through the string just distal to the tip of the tail. The hindquarters were suspended by the tail just high enough to prevent any weight bearing of the hindlimbs on the floor of the cage. The forelimbs were allowed to maintain contact with the floor of the cage, and the suspended rats had access to water and food ad libitum. All rats were housed individually in an isolated chamber maintained at room temperature (22 ± 2 °C) under a 12:12-h light–dark cycle. This study was approved by the Institution Animal Care and Use Committee and adhered to the Kobe University Animal Experimentation Regulations (Kobe, Japan). All experimental and animal care procedures were conducted in accordance with the National Institutes of Health Guide for the Care and Use of Laboratory Animals (National Research Council, Washington DC; 1996).

Propolis supplementation

Brazilian propolis (Yamada Bee Farm, Okayama, Japan) diluted by 5% arabic gum-containing solution was administered orally by a feeding needle twice a day (500 mg/kg, with a 6-h interval between the two supplements; total daily dose 1000 mg/kg) for 2 weeks to the rats in the CON + PP and HU + PP groups. The rats in the CON and HU groups were treated similarly with a 5% arabic gum-containing solution without propolis. The non-toxicity of diet supplementation with 1000 mg/kg/day of Brazilian propolis had been confirmed in a preliminary 28-day toxicity test performed by the supplier (Yamada Bee Farm), thereby supporting our use of this dose to study disuse-induced capillary regression.

Muscle preparation

At the end of the experimental period, the rats were anaesthetized deeply by inhalation of 2% isoflurane gas via an anesthetic mask. For the histological and biological measurements, the left soleus muscle was removed and excess fat and connective tissue cleaned off, following which it was weighed, mounted on cork, and rapidly frozen in dry ice-cooled acetone. To visualize the three-dimensional capillary architecture of the soleus muscle, we examined the right soleus muscle using a confocal laser scanning method as described previously [7, 27, 28]. Briefly, a catheter was inserted into the right iliac artery, and 0.9% physiological saline containing heparin (1000 IU/l) at 37 °C was injected to wash out the intravascular blood and induce maximal vasodilation. A contrast medium (8% gelatin solution containing 1% fluorescent material) was then injected. After reflux with 50 ml of a contrast medium, the right ankle joint was held in a maximum plantar-flexion position and the entire right hindlimb was quickly immersed into ice-cooled saline. After 15 minutes, the right soleus muscle was removed and excess fat and connective tissue cleaned off; the right soleus muscle was then mounted on cork, and rapidly frozen in dry ice-cooled acetone, as described for the left soleus muscle. All frozen samples were stored at −80 °C until subsequent analyses.

Histological analyses

The mid-belly of the left soleus muscle was mounted onto a specimen chuck using Tissue-Tek OCT compound (Sakura Finetechnical, Tokyo, Japan), and transverse
sections (thickness 12 µm) were cut on a cryostat microtome (CM3050S; Leica Microsystems, Heidelberg, Germany). The transverse sections were stained for alkaline phosphatase (AP) to visualize capillaries; the sections were incubated in 5-bromo-4-chloro-3-indolyl phosphate/nitro blue tetrazolium for 45 min at 37 °C and fixed with 4% paraformaldehyde. The capillary-to-fiber (C/F) ratio was determined by counting capillaries and myofibers on each cryosection using microscopic images of AP staining.

In situ detection of reactive oxygen species (ROS) generation was evaluated with the fluorescent probe dihydroethidium (DHE) which detects oxidative stress by emitting light following its interaction with O2⁻ to form oxyethidium. DHE is cell permeable and interacts with nucleic acids to emit a light-red color that is detectable qualitatively by fluorescence microscopy (model BX51; Olympus, Tokyo, Japan). This staining method in skeletal muscle tissues has been applied in previous studies [5, 22]. Briefly, the transverse sections (thickness 10 µm) were incubated with 5 × 10⁻⁶ mol/l DHE (Wako Pure Chemicals, Osaka, Japan) for 30 min at 37 °C in the dark box, following which the sections were rinsed with phosphate-buffered saline (PBS; 37 °C) and observed under a fluorescent microscope equipped with a filter (excitation at 545 nm). The fluorescence intensity was determined by the fluorescence area fraction using gray scale analysis in five images per muscle. All measurements were performed using Image J software program (National Institutes of Health [NIH], Bethesda, MD, USA).

Three-dimensional architecture of the capillary network

The capillary volume and luminal diameter within the right soleus muscle were determined by confocal laser scanning microscopy (model TCS-SP; Leica Microsystems) with an argon laser (488 nm). This three-dimensional (3D) imaging was performed as previously described [7, 12, 15, 27]. Briefly, the longitudinal 100-µm-thick sections were scanned to a depth of 50 µm at a slice thickness of 1 µm. Stacking in 50-slice file was converted into digital images in order to visualize the 3D architecture of the capillary network. Finally, the capillary volume (× 10⁻² mm³/mm³) was measured in a 3D image (250 × 250 × 50 µm, length × width × depth), and the capillary luminal diameter (µm) was measured using the Image J software program (NIH, Bethesda, MD, USA).

Western blot analysis

A portion (approx. 20 mg) of each left soleus muscle was homogenized in RIPA lysis buffer containing 1 mM Na₃VO₄, 1 mM NaF, and protease inhibitor cocktail (1:100; P8340; Sigma-Aldrich, St. Louis, MO, USA). The total protein concentration of the supernatants was determined according to Bradford [29] using the protein assay kit of Bio-Rad Laboratories (Hercules, CA, USA). The proteins were loaded (20 µg/lane) and separated on a 12.5% sodium dodecyl sulfate–polyacrylamide gel, then blotted onto polyvinylidene difluoride membranes, and blocked for 1 h with 5% skimmed milk in PBS with Tween 20 (PBST). The membranes were incubated using antibodies against superoxide dismutase 1 (SOD-1; 1:200 in PBST; catalog no. sc-8637, Santa Cruz Biotechnology), or p53 (1:200 in PBST; catalog no. sc-98; Santa Cruz Biotechnology), or VEGF (1:200 in PBST; catalog no. sc-2670, Santa Cruz Biotechnology), or p53 (1:200 in PBST; catalog no. sc-98; Santa Cruz Biotechnology) overnight at 4 °C, then incubated in a solution with horseradish peroxidase (HRP)-conjugated anti-goat (1:10000 in PBST) or HRP-conjugated anti-mouse (1:10000 in PBST) for 1 h. The proteins were detected using the EZWestLumi one enhanced chemiluminescence solution (ATTO, Tokyo, Japan). Finally, images were analyzed with the LAS-1000 (Fujifilm, Tokyo, Japan) chemiluminescent image analyzer and quantified using the Multi-Gauge Image Analysis Software program (Fujifilm) against the relative concentration of β-actin (1:1000 in PBST; catalog no. sc-47778; Santa Cruz Biotechnology) as an internal control.
Quantitative PCR analysis

Total RNA was extracted from approximately 10 mg of each left soleus muscle using an extraction kit (QuickGene RNA tissue kit SII; Fujifilm). Reverse transcription was carried out using the high-capacity cDNA archive kit (Applied Biosystems, Foster City, CA, USA), and cDNA samples were stored at −20 °C until use. Expression levels of TSP-1 (Rn01513693_m1) were quantified by TaqMan Gene Expression Assays (Applied Biosystems). Each TaqMan probe and primer set was validated by performing a quantitative real-time PCR with a series of cDNA template dilutions to obtain standard curves of threshold cycle time against relative concentration using the housekeeping gene 18S as an internal standard. All samples and non-template control reactions were performed in a 7500 Fast Sequence Detection System (Applied Biosystems).

Statistical analysis

All values are expressed as mean ± standard error of the mean. Overall significant differences between the four experimental groups were determined using one-way analysis of variance followed by Tukey post hoc tests to determine specific group differences. Statistical significance was set at \( P < 0.05 \).

Results

Body and soleus muscle weight and muscle fiber cross-sectional areas

Absolute body weight was significantly lower in the HU and HU + PP groups than in the CON and CON + PP groups; similarly, both absolute and relative soleus muscle weight were significantly lower in the HU and HU + PP groups than in the CON and CON + PP groups. In addition, the relative muscle-to-body weight in the HU + PP group was significantly higher than that in the HU group (Table 1). The fiber cross-sectional area (FCSA) of soleus muscle was significantly decreased in the HU and HU + PP groups compared to the CON and CON + PP groups. There was no significant difference in FCSA between the HU and HU + PP groups (Table 1).

The C/F ratio

The representative images of soleus muscle stained for AP are shown for each group in Fig. 1a–d. The C/F ratios in the HU and HU + PP groups were significantly lower than those in the CON and CON + PP groups (Fig. 1e). However, the C/F ratio in the HU + PP group was significantly higher than that in the HU group (Fig. 1e).

Capillary volume and luminal diameter

The representative confocal laser scanning microscopic images of 3D capillary architecture in each group are shown in Fig. 2a–d. The capillary network in the longitudinal section of the soleus muscle appeared to be less complex in the HU group (Fig. 2c) than in the CON (Fig. 2a), CON + PP (Fig. 2b), and HU + PP (Fig. 2d) groups. The mean capillary diameter and volume in the HU group were significantly lower than those in the CON, CON + PP, and HU + PP groups (Fig. 2e, f).

The frequency distribution of capillary luminal diameter in the HU (Fig. 3c) group was shifted towards capillaries of smaller diameter compared with the CON (Fig. 3a), CON + PP (Fig. 3b), and HU + PP (Fig. 3d) groups. The percentage of capillaries with a diameter of < 2.5 μm, through which red blood cells can pass [30], increased to a high value in the HU group (17.5%) compared with the other groups (CON 4.3%, CON + PP 3.9%, HU + PP 4.9%; Fig. 3a–d).

Table 1  Body weight, soleus muscle weight, and fiber cross-sectional area

| Study groups \( ^a \) | Body weight (g) | Soleus muscle absolute weight (mg) | Soleus muscle weight/body weight (mg/100 g) | Fiber cross-sectional area (μm\(^2\)) |
|-----------------------|-----------------|-----------------------------------|---------------------------------------------|-------------------------------------|
| CON                   | 295 ± 3         | 116 ± 2                           | 39.4 ± 0.9                                  | 3.029 ± 159                        |
| CON + PP              | 288 ± 1         | 115 ± 2                           | 39.7 ± 0.9                                  | 3.242 ± 54                         |
| HU                    | 256 ± 5\( ^* \) | 56 ± 2\( ^* \)                   | 21.9 ± 0.4\( ^* \)                         | 1.226 ± 44\( ^* \)               |
| HU + PP               | 253 ± 4\( ^* \) | 62 ± 1\( ^* \)                   | 24.5 ± 0.5\( ^* \)                         | 1.427 ± 53\( ^* \)               |

Values in table shown as the mean ± standard error of the mean (SEM)

\( ^* , ^{\dagger} , \) and \( ^{\ddagger} \) indicate a significant difference from the CON, CON + PP, and HU groups, respectively, at \( P < 0.05 \)

\( ^a \) CON Control group, CON + PP control plus propolis supplementation group, HU hindlimb unloading group, HU + PP hindlimb unloading plus propolis supplementation group
ROS production and SOD-1 protein levels

Intracellular ROS production was measured by DHE fluorescence in the nucleus. The representative images of the soleus muscle stained with DHE in each group are shown in Fig. 4a–d. DHE fluorescence in the HU group and capillary volume (f) in the soleus muscle of each group. Values are shown as the mean (bars) ± standard deviation of the mean (SEM; error bars). The asterisk (*), dagger (†), and double dagger (‡) indicate a significant difference from the CON, CON + PP, and HU groups, respectively, at $P < 0.05$
was significantly higher than that in the CON, CON + PP, and HU + PP groups (Fig. 4e). The value in the HU + PP group was consistently similar to that in the CON and CON + PP group.

The protein level of the primary cellular enzymatic antioxidant SOD-1 was determined by western blotting. The expression level of SOD-1 protein was significantly higher in the HU group than in the CON, CON + PP, and HU + PP groups, whereas the level in the HU + PP group did not differ from that in the CON and CON + PP groups (Fig. 4f).

**Protein expression level of p53 and mRNA expression level of TSP-1**

The protein expression level of p53, one of anti-angiogenic factors, was significantly increased in the HU group compared with the CON group, while the value in the HU + PP group was maintained at the CON level (Fig. 5a). Similarly, the mRNA expression level of TSP-1, one of the anti-angiogenic factors tested, was significantly increased in the HU group compared with the other groups, while the expression level in the HU + PP group was maintained at the CON level (Fig. 5b).

**Protein expression level of VEGF**

The expression level of VEGF protein, one of the pro-angiogenic factors tested, in the HU group did not differ from that in the CON group. However, the level in the HU + PP group was significantly higher than that in the HU group (Fig. 6).

**Apoptosis of endothelial cells**

Apoptosis of endothelial cells was identified by staining with TUNEL and CD-31 on transverse section of soleus muscle. The representative images of TUNEL-positive endothelial cells were shown in the Fig. 7a–c. The frequency of TUNEL-positive endothelial cells in the HU group was significantly higher than that in the CON, CON + PP, and HU + PP groups, and the value in the HU + PP was maintained at the CON level (Fig. 7d).

**Discussion**

The novel findings of this study are that the administration of Brazilian propolis prevented both overexpression of ROS due to a chronic decrease in neuromuscular activity and capillary regression by suppressing anti-angiogenic signaling and stimulating pro-angiogenic signaling within the atrophied soleus muscle. Based on these results, we suggest that Brazilian propolis treatment is an effective therapy for capillary regression in the chronically unloaded skeletal muscle.

The capillary network in skeletal muscle is susceptible to structural changes in response to various conditions. Previous studies have reported that neuromuscular inactivity results in not only a reduction of capillary number [4, 14], but also a decrease in capillary luminal diameter and volume within the unloaded soleus muscle [5, 7, 15]. Additionally, capillaries with a diameter < 2.5 μm, through which erythrocytes can pass [30], were observed more frequently...
in unloaded skeletal muscle than in the skeletal muscle of grounded rats. This result indicates that unloading increases dysfunctional capillaries within the skeletal muscle. Taken together, these results are consistent with those observed in the present study.

Oxidative stress is increased within an unloaded muscle due to an overproduction of ROS [31–33]. Additionally, oxidative stress associated with unloading has been shown to participate in the pathophysiology of capillary regression within the skeletal muscle, as shown in studies using antioxidant treatment [5, 22]. Therefore, we also investigated oxidative stress in the present study. As a marker for

![Fig. 4](image)

**Fig. 4** Transverse sections from each group (a CON, b CON + PP, c HU, d HU + PP) stained with the fluorescent probe dihydroethidium (DHE) that detects oxidative stress. e, f Mean levels of DHE fluorescence (e) and superoxide dismutase 1 (SOD-1) protein expression (f) in the soleus muscle of each group. Scale bar: 50 μm. Values are shown as the mean (bars) ± SEM (error bars). The asterisk (*), dagger (†), and double dagger (‡) indicate a significant difference from the CON, CON + PP, and HU groups, respectively, at P < 0.05. AU Arbitrary units

![Fig. 5](image)

**Fig. 5** Mean levels of p53 protein expression (a) and mean levels of thrombospondin-1 (TSP-1) mRNA (b) in the soleus muscles of each group. Values are shown as the mean (bars) ± SEM (error bars). The asterisk (*), dagger (†), and double dagger (‡) indicate a significant difference from the CON, CON + PP, and HU groups, respectively, at P < 0.05

![Fig. 6](image)

**Fig. 6** Mean levels of vascular endothelial growth factor (VEGF) protein expression in the soleus muscles of each group. Values are shown as the mean (bars) ± SEM (error bars). The asterisk (*), dagger (†), and double dagger (‡) indicate a significant difference from the CON, CON + PP, and HU groups, respectively, at P < 0.05
oxidative stress, we examined DHE fluorescence intensity and SOD-1 protein expression within the unloaded soleus muscle [5, 22, 34]. Our results show that HU enhanced both DHE fluorescence intensity and SOD-1 protein expression, thereby reflecting increased oxidative stress within the unloaded soleus muscle, which is consistent with the findings of earlier studies [5, 22, 34]. Oxidative stress increases protein expression of transcriptional factor p53 [35], known as a tumor suppressor protein [36]. p53 plays a key role in regulating various signaling pathways to maintain skeletal muscle health [37]. It has been reported that the increase in p53 protein expression induces endothelial cell apoptosis and then suppresses angiogenesis [38, 39]. Indeed, p53 protein expression has been found to increase within the unloaded soleus muscle [40]. Roudier et al. demonstrated that HU-related p53 protein expression participates in capillary regression by upregulating TSP-1 protein expression [4], leading to endothelial cell apoptosis. However, Roudier et al. also reported that p53 does not play a role in the decreased VEGF protein expression in response to unloading [4]. The results of our present study on the expression of these proteins are consistent with those reported by Roudier et al. [4]. In addition, in a review of the literature, Olfert et al. found that capillary regression depends on increasing TSP-1 expression but not on decreasing VEGF expression [20]. Our present results clearly suggest that HU-induced oxidative stress leads to capillary regression via the upregulation of p53 and its downstream TSP-1 expression, with support provided by our observation of increased endothelial cell apoptosis, as expressed in TUNEL-positive endothelial cells within the unloaded soleus muscle.

Antioxidants can be regarded as countermeasures for disuse-induced microangiopathy. Results from earlier studies suggest that supplementation with anti-oxidative food components could attenuate disuse-induced capillary regression [5, 22]. Brazilian propolis contains a variety of bioactive components and possesses many biological and pharmacological properties [23], and it has been used since ancient times as a non-drug therapeutic agent. Brazilian propolis also has antioxidant activity due to its content of flavonoids [24] and cinnamic acid derivatives [25, 26]. Our study of the effects of Brazilian propolis on unloading-induced oxidative stress revealed that Brazilian propolis suppressed the ROS production observed in DHE fluorescence intensity and SOD-1 protein expression studies within the unloaded soleus muscle to near control levels. These results suggest that Brazilian propolis bestows protective effects on unloading-induced oxidative stress within the muscle. We also found that Brazilian propolis attenuated the unloading-induced changes in anti-angiogenic factors and capillary architecture within the skeletal muscle. These results are in agreement with those reported in our previous studies [5, 22] in which we used the powerful antioxidant astaxanthin, leading use to presume that Brazilian propolis administered daily into
disused muscle exhibits antioxidant and resultant vascular protective effects equivalent to those with daily antioxidant astaxanthin supplementation. Antioxidant treatment has been found to attenuate the p53-related anti-angiogenic pathway through scavenging ROS [41, 42]. Some of the components of Brazilian propolis have been reported to be associated with pro-angiogenic effects via modulation of oxidative stress [43, 44]. Therefore, the prevention of capillary regression by Brazilian propolis observed in the study can likely be explained by the regulation of oxidative stress and related negative angiogenic regulators, such as p53 and TSP-1. This notion is supported by the observation of the suppression of endothelial cell apoptosis within the unloaded muscle treated with Brazilian propolis.

We observed an increase in VEGF protein expression within the unloaded muscle treated with Brazilian propolis. Decreased VEGF expression may have little involvement in capillary regression [20], but increased VEGF expression prevents unloading-induced capillary regression [15, 22]. Thus, Brazilian propolis may prevent capillary regression by increasing pro-angiogenic VEGF protein expression, which is a novel effect of Brazilian propolis. However, further investigations are needed to clarify the underlying mechanism in Brazilian propolis-related VEGF protein expression observed in the present study.

Anti-tumorigenic effects of Brazilian propolis have been demonstrated in various cancers in both animal and in vitro models [45–47]. The anti-tumorigenic effect can be explained mainly by the pro-apoptotic [48] and anti-angiogenic activities [49] of the active components in Brazilian propolis. In the present study, however, we found that Brazilian propolis did not induce endothelial cell apoptosis or capillary regression within the soleus muscle of rats treated with or without HU, which seems to suggest a role for endothelial cytoprotection in skeletal muscle. These results may indicate that Brazilian propolis exerts both pro- and anti-apoptotic effects depending on cellular status. Additionally, an in vitro toxicological study of Brazilian propolis by Xuan et al. showed that a high concentration of Brazilian propolis increases ROS production and p53-dependent apoptosis in human umbilical vein endothelial cells but that a low concentration decreases both processes [50]. This result means that Brazilian propolis has both pro-cytotoxic and anti-cytotoxic activity depending on its concentration. In our study, we administered a daily oral dose of Brazilian propolis of 1000 mg/kg to rats for 2 weeks; the safety of this dose in terms of toxicity and frequency of administration had been confirmed in preliminary in vivo experiments by the manufacturer. Therefore, our results support the possibility that Brazilian propolis acts as an anti-cytotoxic agent within the soleus muscle. However, we did not investigate the dose-dependent effects of Brazilian propolis on capillary regression in disused muscle. Further studies are needed to clarify this issue.

In conclusion, the results of our study indicate for the first time that the administration of Brazilian propolis has protective effects on the capillary regression that accompanies chronic muscle unloading. The pro-angiogenic effects of Brazilian propolis may be attributed to its bioactive components which lead to a suppression of the anti-angiogenic pathway and stimulation of the pro-angiogenic pathway. Therefore, Brazilian propolis seems to be a non-drug therapeutic agent that is applicable to the treatment of skeletal muscle angiopathy that occurs in various situations, such as bed rest, cast immobilization, diabetes, and cancer cachexia.

Conclusion

In conclusion, we found that Brazilian propolis supplementation was an effective treatment to attenuate capillary regression in soleus muscle associated with prolonged mechanical unloading. Our results suggest that Brazilian propolis can be administered as a therapeutic intervention for the treatment of skeletal muscle angiopathy due to disuse-induced muscle atrophy.

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Author contributions MT, MK, and HF conceived and designed the experiments. MT and MK performed the experiments. MT, MK, and HF analyzed the data. MK, NM, HK, AI, and HF contributed by providing regents, materials and analysis tools. MT, HK, AI, and HF interpreted the data and wrote the paper. All authors approved the final version of the manuscript.

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Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

Ethical approval All applicable international, national, and/or institutional guidelines for the care and use of animals were followed. All procedures performed in studies involving animals were in accordance with the ethical standards of the institution or practice at which the studies were conducted. This article does not contain any studies with human participants performed by any of the authors.
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