Interval Training Is Not Superior to Endurance Training With Respect to Bone Accrual of Ovariectomized Mice

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Physical exercise is considered to delay bone loss associated with post-menopausal estrogen deficiency in women. However, the optimal training regimen for maximal bone accrual has not yet been defined. We, therefore, turned to ovariectomized (OVX) C57BL/6 mice and directly compared a low intensity endurance training on the treadmill to medium and high intensity interval trainings tailored to the individual performance limits. Trainings lasted 30 min each and were performed five times/week. After a 5-week training period, mice were sacrificed, and the hind legs were analyzed for assessment of (i) biomechanical stability (three-point bending test), (ii) bone microarchitecture [micro-computed tomography (μCT)], (iii) mineral apposition rate (MAR; histomorphometry), and (iv) muscle volume (MRI). Increased running speeds and quadriceps femoris muscle volumes in trained mice confirmed positive impacts on the cardiopulmonary system and myoinduction; however, none of the treadmill training regimens prevented ovariectomy induced bone loss. Our results provide evidence that treadmill training impacts differentially on the various members of the musculoskeletal unit and call for further experiments investigating frequency and duration of training regimens.

Keywords: osteoporosis, treadmill, bone, micro-computed tomography, mouse, ovariectomy, interval training

INTRODUCTION

Osteoporosis is a major health care problem that is characterized by a loss of bone mass that can lead to pain, immobility, and fragility fractures that, in turn, result in increased mortality (WHO Scientific Group Meeting on Prevention and Management of Osteoporosis, 2004). Osteoporosis is primarily a disease of the elderly; in six European countries (Italy, Spain, Sweden, France, Germany, and UK), a total of 16 million women and 4 million men suffered from osteoporosis in 2015 (International Osteoporosis Foundation, 2019). Approximately 3.2% of women over the age of 50 are affected by fragility fractures every year. The lifetime risk...
Latza et al. Interval Training of Ovariectomized Mice

MATERIALS AND METHODS

Mice and Experimental Design
A total of 40 7–8-week old C57BL/6J female mice were purchased from Charles River (Charles River Laboratories, Research Models and Services, Sulzfeld, Germany). In our facility and throughout the experiments, all animals had free access to pellet food and water. They were housed in cages in small groups up to four individuals under a 12/12 h light/dark cycle. After a week of acclimatization, the mice were randomized into five groups of equal numbers (n = 8). The first group (I) was sham-operated and served as a control while the other four (II–V) were ovariectomized. After 1 week of recovery, mice in the groups III–V started their 2 weeks familiarization to the treadmill. At the end of this period, all mice accustomed to the treadmill (groups III–V) took part in an augmentation run test (ART) that served to determine the individual maximum velocity (V_max). For the next 5 weeks, mice in groups III–V were trained on the treadmill, and the ART was repeated twice (after 2.5 weeks and at the end) in order to monitor training effects as changes in the individual V_max and to adapt the training accordingly as running speed can be used to tailor training intensities (Høydal et al., 2007). Nine and two days before sacrificing the mice, tetracycline and demeclocycline, respectively, were injected intraperitoneally for subsequent assessment of mineral apposition rate (MAR). At the end of the training period, the volumes of the quadriceps femores’ muscles were measured using 7-Tesla magnetic resonance imaging (MRI). Thereafter, all mice were sacrificed, blood was aspirated, uteri were prepared, and femora and tibiae were preserved. In-vivo blood samples could not be taken without weakening the mice and thus influencing the study’s course.

An overview of the study design is shown in Figure 1. The local state’s animal care committee (Landesamt für Landwirtschaft, Lebensmittelsicherheit und Fischerei M-V; www.lallf.de) approved all experiments (7221.3-1-021/17), and all experiments were carried out in accordance with the relevant guidelines and regulations.

Ovariectomy
All animals were anesthetized with ketamine/xylazine (100 mg ketamine and 5 mg xylazine per kg body weight) and kept at 36°C on a warming plate. An ointment was applied to the eyes to prevent dehydration. The lower back was shaved, and a bilateral cut was placed into the skin and peritoneum. The skin of sham-operated mice was re-sutured, while mice of the groups II–V were ovariectomized via open exposure: preparation not only allows for axial loading, but also for differential energy conditions as low and medium intensities guarantee aerobic conditions while high intensity trainings that were tailored to the individual performance limits aim at an anaerobic state (Alizadeh et al., 2015). We here concentrated on C57BL/6 females for their superior endurance capacity and the availability of previous publications (Wallace et al., 2007, 2009; Banu et al., 2008; Hollinski et al., 2008; Kvedaras et al., 2017).

Consensus seems to exist that rodents can model the human skeleton (Frost and Jee, 1992). There is a limitation though because quadrupedal animals – in contrast to humans – experience greater mechanical loading on the appendicular than on the axial bones (Iwamoto et al., 2005). Effects of running exercise on the vertebral bodies of rodents can therefore not be transferred to humans.

Apart from that, the literature on the effect of treadmill training on murine bone shows mixed findings. For example, short term and moderate training periods of 3 weeks led to significantly increased tibial cortical area and periosteal perimeter in male C57BL/6 mice while 8 weeks of a strenuous sprint training on the vertebral bodies of rodents can therefore experience greater mechanical loading on the appendicular skeleton (Alizadeh et al., 2015). We here concentrated on C57BL/6 females for their superior endurance capacity and the availability of previous publications (Wallace et al., 2007, 2009; Banu et al., 2008; Hollinski et al., 2008; Kvedaras et al., 2017).

All of these studies used different mouse strains, different sexes, and different training regimens, suggesting that firm comparisons can only be drawn from identical experimental set-ups.

We here set out to investigate different training intensities and decided on the model of ovariectomized (OVX) females, as a previous study suggested that OVX within a short time frame led to bone loss, which could partially be prevented by a moderate treadmill exercise (Wu et al., 2001). Performing treadmill training...
of the uterus, ovariic tube, and ovary, then cauterization of the ovariic tube, and excision of the ovary. All wounds were sutured. For post-operative pain relief, metamizole was added to the drinking water for 5 days.

**Augmentation Run Test**
The ART served to determine the individual $V_{\text{max}}$ of mice in groups III–V (Hollinski et al., 2008). The treadmill started with a speed of 0.17 m/s for 3 min. It thereafter accelerated over a period of 2 min to 0.2 m/s. This speed was kept for another 3 min before it accelerated again over a period of 2 min to 0.25 m/s. This cycle of constant speed for 3 min and accelerating over a period of 2 min to a speed 0.05 m/s faster than before was maintained until mice no longer kept pace with the treadmill belt and were swept onto the platform behind the running belt. The highest speeds achieved without any external influence represented the individual $V_{\text{max}}$.

**Training Modalities**
Training was performed daily from Monday to Friday, always in the afternoon, and sessions lasted for 30 min. Training for group III consisted of a low intensity endurance run at a constant speed of 0.2 m/s and a 10° incline. Trainings for groups IV and V consisted of medium and high intensity interval trainings without any inclination, respectively. In that regard, mice started at 40% of their $V_{\text{max}}$ for 6 min, followed by four repeats of medium (60% of $V_{\text{max}}$) or high (80% of $V_{\text{max}}$) speed for 3 min interspersed by 1.5 min of recovery at 40% of $V_{\text{max}}$ (Figure 2A).

**Magnetic Resonance Imaging**
Magnetic resonance imaging was used to quantify the volumes of the quadriceps femores muscles. As the muscle volume also depends on length, the ratio of total volume to femur length [calculated as the numbers of micro-computed tomography (μCT) image slices including parts of the femur] was used for statistical analysis. MRI was carried out on a 7-Tesla BioSpec 70/30 (Bruker, Billerica, MA, United States) using a T2 weighted Turbo Rapid Acquisition with Relaxation Enhancement (TurboRARE) sequence with the following settings: TE/TR 25.25/3227 ms, field of view: 28 × 21 mm, matrix size 233 × 175, slice thickness 0.85 mm, resolution 120 × 120 μm, and RARE-factor: 8, capacity spool 72 mm. Mice were anesthetized with isoflurane and were placed on the back with the hind legs fixed.

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**FIGURE 1 |** Study design and timeline (time in days). OVX, ovariectomy; SHAM, sham-operation; Familiarization, 2 weeks of familiarization with the treadmill; ART, augmentation run test; ip, injection of tetracycline and demeclocycline intraperitoneal; pm analyses, start of post mortem analyses.

**FIGURE 2 |** Left panel (A) shows the different training modalities for trained groups III – V. The training phase lasted five weeks with five sessions per week, 30 min each (plotted on the horizontal axis). Group III did low intensity training with constant velocity at a 10° incline. Groups IV and V did an interval training with peak intensities of 60% or 80% of the maximum velocity ($V_{\text{max}}$, plotted on the vertical axis) and active recovery at 40% of the $V_{\text{max}}$. Box plots on the right (B) consist of boxes that represent 25th to 75th percentiles, the medians and bars that represent the 5th and 95th percentiles of the data. Right panel shows the differences in $V_{\text{max}}$ between the ARTs in all the groups (statistics are calculated with Kruskal-Wallis Test with post-test). From first to second ART, all groups increased their $V_{\text{max}}$ significantly (III: $p = 0.008$; IV, V: $p < 0.001$). From second to third ART, none of the groups increased their $V_{\text{max}}$ significantly.
T2 sequence of both hind legs was run. Contours of the quadriceps femores muscles were redrawn manually (Figure 3A), and resulted surfaces were totaled to volumes automatically using ITK-SNAP software (ITK-SNAP 3.4.c 70/30).

**Histomorphometry**

For determination of the MAR, left tibiae were fixed in 4% PFA, transferred into ethanol, and stored in the dark before embedding into methyl methacrylate (MMA). Embedding, preparation and laser microtomy were implemented by LLS ROWIAK LaserLabSolutions GmbH (Hannover). A decalcification was not necessary for their TissueSurgeon laser, so structures like growth lines could be analyzed without overlay effects. Furthermore, the contact free cutting caused fewer artifacts. With a cutting speed of approximately 1 mm²/s and a diameter of the laser pulse of approximately 1–5 μm, a cutting thickness of 10–100 μm could be accomplished. TissueSurgeon worked with a near infrared laser (1,030 nm). We used fluorochromes with shorter wavelengths, so they were not bleached by the laser.

The sections of 10 μm thickness were then analyzed with a fluorescence microscope (Leica CTR 4000) at the Department of Pediatrics (Rostock University Medical Center). Received data were processed with Leica LAS AF Lite software (Leica Microsystems). Therefore, at 50 different locations within the proximal tibiae, the distance between the two labels (tetracycline and domelocycline) was measured, displaying the MAR per week (Fischer et al., 2011; Neuerburg et al., 2019).

**X-Ray Micro-Computed Tomography**

Cortical and trabecular bone parameters were analyzed using a SkyScan 1076 In-Vivo μCT (Bruker, Billerica, MA, United States; Antwerp, Belgium, Software Version 4.2, 0.5 mm Al-filter, integration time of 1.5 s, isotropic voxel size of 9 μm at 49 kV and 200 μA, rotation step of 0.5°, and averaging frame of 3). To that extent, the right femora and tibiae were fixed in 4% PFA, preserved in ethanol. The samples were washed three times in distilled water and equilibrated with the scanning medium, 0.9% saline solution, for at least 12 h for rehydration before measurements. The specimens were placed into the μCT surrounded by a cylindrical polystyrene tube with styrofoam bedding and aligned with the vertical axis of the scanner. At every run, one Eppendorf tube filled with two calcium hydroxide apatite bone mineral density (BMD) calibration rods with densities of 0.25 and 0.75 g/cm³ for calibration was supplemented. For reducing artifacts, it was scanned for a full 360°.

The images were processed with NRecon reconstruction software (Bruker, Billerica, MA, United States), DataViewer (Bruker, Billerica, MA, United States), and CT Analyser (Bruker, Billerica, MA, United States). The μCT images were first processed by NRecon software using a Gaussian filter with a smoothing kernel of 2, a ring artifact reduction of 6, a beam hardening correction of 30%, and a defect pixel masking of less than 20%. Afterward, the bones were vertically orientated and rotated in the same way by using DataViewer software. Finally, image analysis was done by CT Analyser software using a global threshold of 64–255 and manufacturers 2D and 3D algorithms for the analyses of cortical and trabecular bone, respectively. The regions of interest (ROIs) for image analyses were determined between two reference levels: For the femora, a lower reference level was set at the distal metaphyseal growth plate and the upper one at the fusion zone of greater trochanter and the femoral head. In case of the tibia, the reference levels were defined between the tibia-fibular syndesmosis (lower reference level) and the proximal epiphyseal growth plate (upper reference level). For both femora and tibiae, the cortical bone was analyzed by defining 10 intervals (ROIs) of 10% each within the upper and lower reference levels. For tibia, we complemented one ROI of identical size below the lower reference level. The segmentation of trabecular and cortical bone regions was performed using automated algorithms, and all ROI interpolations were checked visually before further analysis.

Every ROI in tibia and femora was assayed for cortical area fraction (Ct.Ar/Tt.Ar) and cortical thickness (Ct.Th). Trabecular bone analyses at the femoral metaphyses contained...
the most distal and proximal 20% within the reference levels. Same rules applied for tibia analyses, just complemented by the lowest ROI up to an interval of 30%. For evaluation of trabecular bone microarchitecture, bone volume fraction (BV/TV), trabecular number (Tb.N), trabecular thickness (Tb.Th), and trabecular bone mineral density (Tb.BMD) were calculated based on 3D calculations (Bouxsein et al., 2010; Behrendt et al., 2016). The calibration for Tb.BMD measurements was performed with the aforementioned phantom rods, which were scanned and reconstructed as described above. The attenuation coefficient was entered into the calibration dialogue window in CT Analyser for determination of Tb.BMD.

Three-Point Bending Test
Left femora were wrapped in gauze moistened with 0.9% sodium chloride solution and were frozen at −20°C. Before the testing, the bones were thawed at room temperature and then positioned with the posterior condyles downward on a three-point bending machine with a 500 N load cell (zwickiLine Z2.5, Zwick GmbH, Ulm, Germany). The tests were executed at the Department of Material Science and Medical Engineering (Rostock University, Germany). The femoral midshaft was placed directly under the testing stamp with a span length of 6 mm. Applying a gradually increasing bending force with 1 mm/min, the femora were loaded to failure (Jepsen et al., 2015). Bending strength (MPa), postyield displacement (mm), maximum load (N), breaking load (N), and Young’s modulus (MPa) were calculated using the recorded load-displacement curves.

Statistics
Statistical analyses were performed using IBM SPSS Statistics 25 (IBM, NY, United States), and graphs and figures were generated using SigmaPlot 13.0 (Systat Software, CA, United States). Due to small sample sizes, we used non-parametric tests only – Mann-Whitney for the (non-paired) comparisons of two groups and Kruskal-Wallis with post-hoc tests for multiple comparisons. Values of $p$ lower than 0.05 were considered significant.

RESULTS

Treadmill Training Led to Increased Running Speeds and Accumulated Muscle Volume
Five weeks of treadmill training at a frequency of five times per week were well-accepted, and all mice were developed physiologically over the experimental period. They all gained weight continuously, and this weight gain was independent of the various training modalities. Importantly, this exercise period showed significant training effects as documented 2 folds:

i. The first 2.5 weeks led to significant increases in the maximum running speeds ($V_{\text{max}}$) as monitored via ARTs. In the first ART, all groups had median $V_{\text{max}}$ of 0.45 m/s, confirming comparable fitness at the onset of the experiments (Figure 2).

Already at the second ART, $V_{\text{max}}$ had increased to medians of 0.53 m/s in the low intensity endurance group (group III) and to 0.55 and 0.6 m/s in the medium and high intensity interval training groups (groups IV and V), respectively. The intensities of the trainings were thus reflected in the increases of running speeds. However, running speeds more or less plateaued after the first 2.5 weeks of training. There was hardly any additional increase between the second and third ARTs for groups III and IV while all but one mouse in the high intensity group V reached a $V_{\text{max}}$ of 0.6 m/s (Figure 2B).

ii. Around 5 weeks of continuous treadmill training also resulted in accumulated muscle volume. We here concentrated on the hind legs and the quadriceps femoris muscle and compared its volume among OVX mice that either did or did not participate in the treadmill training. To that extent, we performed MRI and normalized our results to the length of the respective femora. Figure 3 therefore compares areas between the trained groups (groups III–V) with means of 10.61 mm$^2$ (group III), 10.41 mm$^2$ (group IV), 10.74 mm$^2$ (group V), and the non-trained controls (group II) with means of 8.99 mm$^2$ (group II), respectively. Of note, there were no differences between the various training regimens. In summary, we here showed that 5 weeks of continuous treadmill exercise resulted in significant training effects with respect to muscle volume and running speeds.

Treadmill Training Did Not Prevent OVX Induced Bone Loss
In order to control effective ovariectomy, we analyzed the uteri at the end of the experimental period. Not only were the uteri of OVX mice (groups II–V) less vascularized, they were also significantly smaller as measured by weight (Figure 4). From here on, we concentrated on those mice only that presented a sufficient degeneration of the uteri and therefore excluded nine OVX individuals from further analyses. In detail, three mice were excluded from group II (OVX, no training), three mice from group III (OVX, low intensity training), two from group IV (OVX, medium intensity training), and one from group V (OVX, high intensity training). While the mean uterus weight was 0.37% of the body weight in the sham-operated control mice, OVX resulted in mean relative weights of 0.059%, confirming sustained estrogen-deficiency.

We next assessed the impact of OVX on changes to the cortical and trabecular bones. To that extent, we performed μCT of femora and tibiae. While trabecular bone can only reliably be quantified at the distal and proximal ends of these long bones, this is different for the cortical shell. However, as we were curious whether potential cortical changes would occur evenly distributed over the entire length of the long bones, we divided the femora into 10 and the tibiae into 11 equally sized sections that were analyzed individually. Indeed, we observed the most pronounced changes for the cortical bone in the distal parts of the tibiae and the diaphyseal parts of the femora. For the trabecular bone, the effects were very
prominent for the proximal parts of both femora and tibiae. Example results are presented in Figure 5 for the diaphyseal femoral region (10% of the midshaft) and the most proximal femoral 20% section. As expected, OVX resulted in a significant loss of cortical and trabecular bones as shown by the comparison between groups I and II–V for Ct.Th, Ct.Ar/Tt.Ar, BV/TV, and Tb.N (Figure 5). While the loss of BV/TV could mostly be attributed to a loss of trabecular numbers, a reduction in cortical bone area (Ct.Ar) and total cross-sectional area (Tt.Ar) indicated periosteal bone loss yet did not allow for conclusions at the endosteum. At proximal and distal regions, Tb.Th stayed unaffected by the ovariectomy while Tb.BMD showed a significant decrease in the groups II–V compared to group I in both tibiae and femora. In summary, OVX led to a significant loss of both cortical and trabecular bones, and none of our various training regimens was capable of preventing or reverting this loss.

### Treadmill Training Decreased Bending Strength

To determine the effects of OVX and treadmill training on the bone’s mechanic stability, three point bending tests were carried out. Neither maximum load, breaking load, bending strength, nor post yield displacement were significantly impacted by OVX, as shown in Figure 6. However, treadmill training had a significant negative impact on the Young's modulus (Figure 6A) and on bending strength (Figure 6B). Accordingly, the post yield displacement increased significantly (Figure 6C). Of note, there were no significant differences between the various training regimens.

### Ovariectomy but Not the Treadmill Training Impacted on the Mineral Apposition Rate

To determine the impact of ovariectomy and treadmill training on the MAR, tetracycline and demeclocycine were injected 9 and 2 days before sacrificing the animals. Histomorphometric analyses of tibial laser sections revealed the fluorescent label...
near the endosteum. Importantly, significant increases of MAR were found in all OVX groups (II–V) and resulted in a mean of 13.51 μm/week compared to a mean of 10.61 μm/week in the control mice. However, among the OVX groups, there were no differences between trained and non-trained mice nor among the various training regimens (Figure 7).

**DISCUSSION**

We here investigated treadmill running for its potential to prevent or delay OVX induced bone loss in C57BL/6 mice. In detail, we compared three different training regimens, a moderate endurance as well as medium and high intensity interval trainings, and indeed, the main purpose of this study was to identify the most adequate strain for maximal bone accrual. Our experimental design was based on the two assumptions that (i) increased mechanical loads would lead to periosteal bone growth as well as mechanically more efficient structures and that (ii) higher stresses would result in more bone formation (Robling and Turner, 2009). However, while increased running speeds and accumulated muscle volume of the hind legs confirmed positive impacts on the cardiopulmonary system and myoinduction, none of the treadmill training regimens prevented OVX induced bone loss nor did they improve the biomechanical properties of the bone.

The deterioration of biomechanical properties could be attributed to OVX, and this was in line with previous publications (Lambers et al., 2012; Zhang et al., 2016). Likewise, an enhanced MAR at the endosteum as a consequence of
OVX has been reported before (Wu et al., 2001). However, we could not repeat previous reports on mice and rats alike that demonstrated bone accrual in response to treadmill training (Iwamoto et al., 1998; Wu et al., 2001; Hamrick et al., 2006; Wallace et al., 2009). Individual results are difficult to compare though, as the published studies not only differ with respect to animal species (rats vs. mice) but also to strain, age, sex, nutritional, and hormonal statuses. While our results are disappointing at first sight, there are several implications that are worth pointing out.

i. Our μCT data suggested that even though treadmill running did not prevent OVX induced bone loss, more Ct:Th and Ct.Ar/Tt.Ar were preserved under more strenuous training regimens (Figure 5). These results are supportive of previous publications showing that high and odd impacts lead to the greatest bone accrual as our interval training included acceleration and deceleration as irregular impacts (Tenforde and Fredericson, 2011; Weidauer et al., 2014). It is along these lines that we would like to address the conflict between our results and those by Wu et al. (2001). They established exercise regimens similar to our moderate endurance training yet used a different mouse strain and performed daily exercise. Strain specific genetics as well as frequencies may therefore play a role.

ii. Our μCT data further suggested that even though OVX decreased the trabecular bone mass, treadmill training had no additional impact, neither positive nor negative. These observations are in line with previous publications, showing a lack of exercise-associated changes at the proximal tibia and the vertebral bodies alike (Iwamoto et al., 1999; Bennell et al., 2002; Hollinski et al., 2008). This distinctive feature of the trabecular bone is difficult to explain and has in the past been interpreted as a consequence from females accumulating trabecular bone mass in excess of the optimal bone mass, in order to provide a buffer against the bone loss during lactation (Bennell et al., 2002). It may therefore turn out worthwhile to concentrate on the cortical bone only when optimizing a training regimen (Holzer et al., 2009).

iii. We and others have evidence that longer training periods may be detrimental to the bone (Yeh et al., 1993; Iwamoto et al., 1999; Bennell et al., 2002). The exact kinetics still need to be investigated as is the question of whether or not the physiological bone mass measured in non-OVX controls can be exceeded by treadmill training. However, a clear correlation between strain gradients and levels of bone formation has been reported, and future training regimen aiming at maximal bone formation will require a careful adjustment (Stadelmann et al., 2015).

As it was recently reported that the bones of the hind legs respond better to treadmill training even though the fore legs experience higher external forces, we are confident that we did not miss-out positive responses by concentrating on the hind legs only (Wallace et al., 2015). Nonetheless, there are limitations to our study, and these include the rather small numbers of animals that were calculated based on large effect sizes (Wu et al., 2001). Furthermore, there was no control for total training volume but a control for the amount of time by comparing different training intensities. The total training volume was bound to be different in terms of intensities and distances run. Moreover, taking serum samples throughout the training period would have allowed to monitor systemic parameters, indicating bone formation and resorption. Assuming that bones need to adapt to increased strain and that an initial resorption merges with a subsequent formation phase, serum parameters would allow to detect metabolic changes to the bone before results can be quantified via imaging.

In summary, our results provide evidence that treadmill training impacts differentially on the various members of the musculoskeletal unit and call for follow-up experiments that will identify training regimens as well as factors that will benefit muscles and bones alike.

DATA AVAILABILITY STATEMENT
The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

ETHICS STATEMENT
The animal study was reviewed and approved by Landesamt für Landwirtschaft, Lebensmittelsicherheit und Fischerei M-V; www.lallf.de (7221.3-1-021/17).

AUTHOR CONTRIBUTIONS
TM, BM-H, SB, and D-CF contributed to the conception and design of this study. MW, TL, and RH contributed to the experimental work. MO and JL contributed most to the experimental work. TM is guarantor. BM-H drafted the article. All authors contributed to the analysis of the data. All authors
contributed to the critical revision of the article and approved the final manuscript for publication.

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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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