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Unilateral Heat Accelerates Bone Elongation and Lengthens Extremities of Growing Mice

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Abstract

Linear growth failure results from a broad spectrum of systemic and local disorders that can generate chronic musculoskeletal disability. Current bone lengthening protocols involve invasive surgeries or drug regimens, which are only partially effective. Exposure to warm ambient temperature during growth increases limb length, suggesting that targeted heat could noninvasively enhance bone elongation. We tested the hypothesis that daily heat exposure on one side of the body unilaterally increases femoral and tibial lengths. Mice ($N = 20$) were treated with 40 °C unilateral heat for 40 min/day for 14 days post-weaning. Non-treated mice ($N = 6$) served as controls. Unilateral increases in ear (8.8%), hindfoot (3.5%), femoral (1.3%), and tibial (1.5%) lengths were obtained. Tibial elongation rate was > 12% greater (15 $\mu\text{m}/\text{day}$) on the heat-treated side. Extremity lengthening correlated with temperature during treatment. Body mass and humeral length were unaffected. To test whether differences persisted in adults, mice were examined 7-weeks post-treatment. Ear area, hindfoot, femoral, and tibial lengths were still significantly increased ~6%, 3.5%, 1%, and 1%, respectively, on the heat-treated side. Left-right differences were absent in non-treated controls, ruling out inherent side asymmetry. This model is important for designing noninvasive heat-based therapies to potentially combat a range of debilitating growth impediments in children.

Keywords

growth plate; endochondral ossification; temperature; phenotypic plasticity; noninvasive therapeutic bone lengthening

Bone elongation disorders have multiple underlying causes, ranging from injury and illness to genetic bone disease. Advancing insight into linear growth regulation at the molecular level¹ has outpaced development of strategies to offset short stature and/or leg length

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discrepancy caused by childhood growth failure. Limb length inequality can lead to disabling health conditions in adulthood, such as scoliosis, chronic back pain, and osteoarthritis.² Alternatives are needed because existing limb-lengthening procedures involve invasive surgery and/or drug regimens, which are only partially effective.³ A major obstacle to successful bone lengthening by noninvasive means is difficulty in targeting therapeutics to cartilaginous growth plates, which do not have a direct blood supply. Experimental drug delivery approaches include surgically implanted catheters and localized injections into specific growth plates.^{4,5}

Data from our lab and others demonstrate that exposure to warm ambient temperature during growth increases bone blood supply and length in young mice.^{6,7} While continuous whole body heating does not effectively translate to the clinic, intermittent-targeted heating could be accomplished with a heating pad or temperature cuff. Localized heat could be an alternative to surgery and a supplement to systemic bone-lengthening drugs to noninvasively achieve limb length equalization.

The objective of this project is to test a unilateral heating model to increase length of specific bones without surgical or drug intervention. We hypothesize that daily heat exposure on one side of the body will unilaterally increase femoral and tibial lengths on the heat-treated side. Our goal is to develop a low-cost, noninvasive method for lengthening bones that can translate into practical therapy to offset linear growth impediments in children.

MATERIALS AND METHODS

Animals and Experimental Design

Procedures were approved by the Institutional Animal Care and Use Committee of Marshall University (Protocol 558). A total of $N = 26$ male and female C57BL/6 mice were obtained from a commercial vendor or on-site breeding colony at 3 weeks weaning age.

Animals were singly caged at 21 °C in order to minimize external temperature fluctuations due to huddling.^{8,9} Group housing can be a major source of variation due to increased cage activity.¹¹ Singly housed mice have reduced variance in bone and body composition,¹⁰ which is important for experimental consistency.

Mice ($N = 20$) were treated once daily to a unilateral heating regimen for 14 days post-weaning. Mice were anesthetized with 1.5% isoflurane and placed in lateral recumbency on a 40 °C heating pad for 40 min each day (Fig. 1). Temperature of the procedure room was 19 °C. Heating was deliberately scheduled at the same time each day at the light cycle start when growth plate height and growth rate are maximal.^{12,13} The 3–5 week age interval is a time of rapid, temperature-sensitive growth in mice.¹⁴ By comparison, this period could be considered roughly similar to human development between toddler age and entry to middle school.

Left and right sides were heat-treated in separate trials to rule out potential side variation. Limbs on the heat-treated side were wrapped in custom fitting thermal booties to ensure

uniform heat distribution (Fig. 3C inset). Foam separators prevented heat transfer to the non-treated side. Ear and hindfoot temperatures were taken three times during each daily treatment using infrared thermometry.⁷ Core temperature and respiration were recorded at least twice per treatment. Mice were weighed daily. A separate group of singly housed mice ($N=6$) that were not subject to anesthesia or heat served as treatment controls.

Mice were given a single intraperitoneal injection of oxytetracycline (OTC) (7.5 mg/kg, Norbrook 200 mg/ml) at the treatment midpoint to measure tibial elongation rate. OTC is a calcium chelator that becomes permanently incorporated into mineralizing tissue and leaves a band of fluorescence that has long been used as a standard for quantifying bone elongation rate in young growing animals.¹⁵⁻¹⁷

Tissue Collection and Elongation Rate Analyses

Experimental 5-week-old mice ($N=14$ females) were euthanized for tissue harvest 1 day after the last heat-treatment. Control mice ($N=6$ females) were euthanized at the same endpoint. Experimental 12-week-old mice ($N=6$, mixed sex) were euthanized 49 days post-heating to evaluate persistent limb length differences at skeletal maturity (Fig. 1). In addition to limb length, cartilaginous ears were measured to document a treatment effect because ear size increases with ambient temperature.^{7,18}

Tibial elongation rate was measured for all 5-week-old control mice ($N=6$) and a subset of the 5-week-old experimental mice ($N=8$). Tibiae from the remaining 5-week experimental mice ($N=6$) and all 12-week-old mice ($N=6$) were kept intact for lengths. Femora and humeri from all mice ($N=26$ total) were reserved for length.

The proximal tibial growth plate was selected to measure elongation rate because its relatively flat contour yields a uniform growth rate across the epiphysis.¹⁷ The adjacent distal femoral growth plate was not used due to its undulating shape (with varied growth rate) and irregular geometry that changes with age,¹⁹ which introduces sampling error and measurement inconsistency. The proximal femoral and distal tibial growth plates were also not used because they contribute least to total limb lengthening,²⁰ with correspondingly reduced growth activity.²¹

The OTC label was visualized in unfixed slab sections of bisected tibiae. One half of each bone was placed in a specialized holder on a glass slide, and cover-slipped with glycerol in PBS. The other half was reserved for a separate histological study. Fluorescence was visualized using a UV filter on a fluorescence stereomicroscope. Brightfield (to delineate the chondro-osseous junction) and fluorescence images were captured in tandem.

Images were calibrated manually in ImageJ software (version 1.44, National Institutes of Health, USA) from a 2mm stage micrometer, and then two lines were drawn on each image. The first line was drawn across the metaphyseal chondro-osseous junction (COJ), marked by invading vasculature at the lower edge of the growth plate (single arrowheads in Fig. 3B). The second line was drawn across the leading (proximal) edge of the OTC band in metaphyseal bone (double arrowheads in Fig. 3B). The vertical distance between the lines was measured at 5 equidistant points across the growth plate using ImageJ. Measurements

were averaged and divided by the 7-day labeling period to estimate daily elongation rate ($\mu\text{m}/\text{day}$).

Long Bone, Ear, and Hindfoot Measurements

Long bones (femora, tibiae, and humeri) were dissected, cleaned, dried overnight, and scanned on a flatbed scanner. Calibrations were obtained from an included metric ruler. Measurements were acquired from the scanned and calibrated images by drawing a line between proximal and distal landmarks on the articular ends: Femoral length was measured parallel to the shaft from the proximal-most point on the greater trochanter to the distal-most point on the medial condyle; tibial length was measured between the proximal articular surface and the distal-most point on the medial malleolus; and humeral length was the distance between the most proximal and distal articular surfaces. This scan-based method of bone measurement has been previously described as a technique for reliably obtaining limb length data from mice.^{22,23}

Left and right ears were removed by cutting along the concave (inner) base of the pinnae. Ears were placed between glass slides, scanned, and calibrated as above. Areas were acquired by manually tracing the scanned ears in ImageJ.

Hindfoot measurements were collected from digital photographs calibrated with a metric ruler. Limb positions were standardized. To account for the natural bend in the digits, hindfoot length was measured as the sum of two connecting lines drawn between the proximal-most end of the skin overlying the calcaneus (heel) to the point of metatarsophalangeal flexion (line one) and the point of metatarsophalangeal flexion to the tip of the third digit (line two).

Data Processing and Statistical Analysis

Measurements were performed separately by at least two different individuals. To minimize potential bias since observers were not blinded, at least one of the observers was trained in the methodology but was not involved in conducting experiments. No significant differences were found between observers, and so observer means were used.

Statistical analyses were performed using SPSS 21.0 software (IBM Corporation, Armonk, NY) with $\alpha = 0.05$ as accepted significance. Body mass comparisons were made between non-treated control mice and heat-treated experimental mice using two-tailed independent samples *t*-tests. Left-right side comparisons of the heat-treated mice were done using one-tailed paired *t*-tests. One-tailed tests were performed because of the a priori hypothesis that the heat-treated side would be larger. Two-tailed comparisons were done in control mice to assess natural left-right variation since there was no a priori expectation of right-left asymmetry in non-treated animals. Temperature-size relationships were assessed using Pearson's correlation.

Data are reported as mean \pm standard deviation (SD) in tabular format and as mean \pm standard error (SE) in graphical format to facilitate viewing. Sample sizes (minimum of $N=6$ mice per variable, unless otherwise stated) were determined a priori by estimating the effect size and data variability to yield a statistical power of 80% at $\alpha = 0.05$. Missing data

($N = 2$ femora and $N = 1$ tibiae pair) are the result of sample loss due to dissection damage and were excluded from statistical testing on an analysis-by-analysis basis (see Table 1).

RESULTS

Skin temperatures of heat-treated hindfeet and ears averaged 40 °C during treatments. Non-treated side temperatures averaged 30 °C with no major fluctuations (Fig. 2). Core temperature and respiration were 36 °C and 60 breaths/min, respectively, under anesthesia (Fig. 2). When non-treated and heat-treated sides were analyzed in aggregate, there were significant positive correlations between hindlimb temperature and tibial elongation rate (Pearson's $r = 0.60$, $p = 0.007$) (Fig. 2A), as well as between ear temperature and ear area (Pearson's $r = 0.66$, $p < 0.001$) (Fig. 2B). Core temperature, plotted in Figure 2 against left-right side averages for reference purposes only, was not included in the analyses. Correlations were not significant in the hindlimb when each side was examined separately; however, within-side relationships were significant in the ear for both non-treated (Pearson's $r = 0.56$, $p = 0.0019$) and heat-treated sides (Pearson's $r = 0.42$, $p = 0.040$) (Fig. 2B).

Tibial elongation rate was over 12% greater on the heat-treated side (Figs. 2A, 3A-B; Table 1). The average growth acceleration was nearly 15 $\mu\text{m}/\text{day}$ (paired $t = 4.12$, $p = 0.002$). Ear area (paired $t = 7.19$, $p < 0.001$) and hindfoot length (paired $t = 5.49$, $p < 0.001$) increased 8.8% and 3.5%, respectively, compared to the non-treated contralateral side at 5-weeks (Fig. 2B, Table 1). Femoral (paired $t = 6.70$, $p < 0.001$) and tibial (paired $t = 3.44$, $p = 0.013$) lengths were increased 1.3% and 1.5%, respectively (Table 1). Humeral length did not differ (paired $t = 0.35$, $p = 0.365$) (Table 1).

To test whether left-right differences were evident at skeletal maturity, mice were examined 49 days after the last treatment at 12-weeks age. Importantly, ear area (paired $t = 3.98$, $p = 0.006$), hindfoot (paired $t = 5.20$, $p = 0.002$), femoral (paired $t = 2.20$, $p = 0.040$), and tibial (paired $t = 4.02$, $p = 0.005$) lengths were still significantly increased on the heat-treated side of adults (Figs. 3C-D). Heat-treated ears were over 6% larger; hindfeet were 3.5% longer; tibiae were 1% longer; and femora were nearly 1% longer when compared to the non-treated contralateral side. Humeral length did not differ (paired $t = 0.18$, $p = 0.431$).

To assess potential side variation, mice were heat-treated on left and right sides in separate trials. Figure 4 shows paired comparisons of non-treated and heat-treated sides of individual 5-week-old mice that were treated on the left (Fig. 4A) and right (Fig. 4B). The average increases in femoral length on the heat-treated side were 1% and 1.6%, respectively, in the left and right side groups. Although the heat response appears to be slightly less pronounced in the left side cohort (paired $t = 4.56$, $p = 0.003$) relative to the right side group (paired $t = 7.08$, $p < 0.001$), the difference was not significant. An important technical note is that the left cohort was among the first groups examined when methods were not as well established.

When experimental mice were compared with non-treated controls, there were no differences in 5-week ending body mass (independent samples $t = 0.29$, $p = 0.777$) or average gain in mass ($t = 0.17$, $p = 0.870$). Control mice also showed no significant left-right differences in femoral length (Fig. 4C) (paired $t = 0.82$, $p = 0.45$), ear area (paired $t = 0.02$, p

= 0.98), humeral length (paired $t = 0.94$, $p = 0.39$), hindfoot length (paired $t = 0.11$, $p = 0.91$), or tibial elongation rate (paired $t = 0.88$, $p = 0.42$), ruling out inherent side asymmetry (Table 2). Control and experimental mice were harvested in separate trials, so differences in total extremity size are the result of variation between cohorts.

DISCUSSION

The goal of this study was to establish a model system using targeted intermittent heat exposure to permanently increase extremity length in mice. Our data support the hypothesis that daily unilateral limb heating increases femoral and tibial lengths on the heat-treated side. Extremity lengthening was correlated with temperature during treatment, particularly in temperature-sensitive cartilaginous ears (Fig. 2B). The length effect persisted at skeletal maturity after only 14 days of post-weaning treatment. A significant right-left difference was measured without impacting overall body mass, and left-right differences were absent in normal non-treated control mice. Core temperature and respiration were in a physiological range under anesthesia. These results suggest that daily unilateral heating is an effective way to model temperature-enhanced hindlimb elongation in young, rapidly growing mice.

The rationale for the model is to develop methods for increasing bone length with minimally invasive methods that can apply to many different growth-limiting conditions. With these baseline data, our model will allow us to move forward and test mechanisms of heat-enhanced bone lengthening to better tailor future clinical therapies. For example, using in vivo multiphoton imaging, we have shown that short-term (30-min) hindlimb heating increases molecular uptake in mouse tibial growth plates.²⁴ Heat could potentially be applied on a scheduled regimen with systemic bone lengthening drugs to target their delivery to specific skeletal growth plates. Routine heat exposure developed in the model here could thus provide a method for augmenting drug-induced limb elongation.

Interestingly, the humerus did not respond to heat-treatment in any of four independent trials conducted for this study (Table 1). Although it is unclear why humeral length did not differ, one potential explanation is the warmer starting temperature of the forelimb when compared to the hindlimb. The knee joint capsule is normally at least 3–4 °C lower than body core.^{25,26} Our treatments elevated hindlimb temperature by 10 °C on the heat-treated side (Fig. 2). However, skin temperatures in the humeral region more closely resembled body core, consistent with thermal maps for humans showing that 37 °C core temperature extends into the shoulder region, while extremity temperatures progressively decrease in a proximal-distal gradient.²⁷

Our working hypothesis is that heat-treatments do not impact temperature of the humerus due to the proximity of this joint to the body core, and the disproportionately large volume of warm blood delivered to the shoulder region through the large subscapular artery.²⁸ Since most elongation of the humerus occurs at its proximal growth plate (shoulder), versus in the distal (knee) growth plate of the femur,^{20,29-31} it is possible that the left-right symmetry in humeral length reflects its relatively constant temperature. This could be tested by decreasing shoulder temperature with cold, which stunts limb elongation in a dose-dependent manner.³²

Although heat effects on extremity lengthening have been documented for over a century,⁶ temperature is still under-recognized for its ability to modulate bone elongation in growth plates. Brookes and May³³ demonstrated up to a 20% increase in bone growth rate for every 1 °C increase in incubation temperature in growing chicks. Doyle and Smart³⁴ used daily application of short-wave diathermy (heat generating treatment) near the growth plate in rats and showed an increase in femoral and tibial lengths. Granberry and Janes³⁵ were not able to replicate these findings using microwave diathermy in dogs; however, their 100 watt treatment produced bone damage that may have prevented heat-related growth acceleration.

Here we found that unilateral exposure of mild, non-damaging 40 °C heat for 40-min per day for only 14 days permanently increased ear area and hindlimb length on heat-treated sides of young mice. No treatment-related damage was observed morphologically or histologically. These results suggest that heat could be a promising strategy for enhancing elongation potential of specific growth plates without affecting the entire skeleton.

One caveat is that the width of the mouse growth plate is only a fraction of that of a human growth plate. It will be important to replicate these results in a larger animal model to ensure that heat can fully penetrate a larger growth plate, so as to avoid potential angular growth deformities. This should not be problematic, however, since whole body heat-effects on bone length have already been demonstrated in experiments using large animals.^{18,32} Treatments could be optimized for a more robust effect by using a temperature cuff to target the most active growth plates. The distal femoral and proximal tibial growth plates contribute most to lower limb lengthening in humans.³¹ Use of a localized heating device around the knee could be a clinically useful tool to augment existing drug therapies (i.e., growth hormone injections) and to potentially avoid invasive surgical limb length correction. A practical strategy in children could be to wrap a noninvasive heating device around the knee during the sleep period when growth occurs¹³ and growth hormone naturally peaks.³⁶

In conclusion, we believe that such heat-based therapies resulting from the model developed here will have advantages over traditional methods that are potentially painful and invasive. Our next step is to establish the effects in a larger animal model, and test them alongside pharmaceutical interventions. This approach could ultimately lead to development of alternative treatment modalities with better outcomes by reducing costs and side effects of surgery and high-dose systemic pharmaceuticals.

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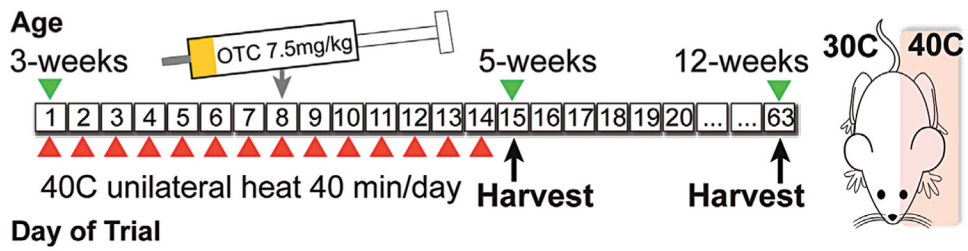


Figure 1. Unilateral heating schematic. Mice were treated with 40 °C unilateral heat for 40 min per day for 14 consecutive days. Oxytetracycline (OTC) was given at the study midpoint to quantify tibial elongation rate.

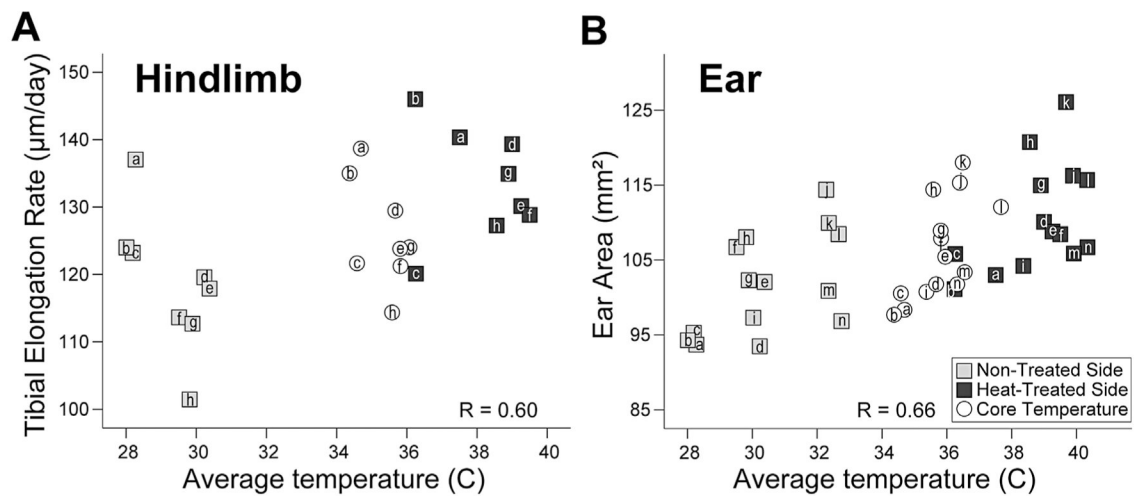


Figure 2.

Extremity temperature correlates with extremity growth. Scatter plots of average temperatures of the hindlimb (A) and ear (B) taken during treatments show a significant positive relationship with growth of non-treated (gray squares) and heat-treated (black squares) sides. Individual 5-week-old mice are denoted by unique letters, which correspond to the same mouse in both graphs. Legend is in (B). For reference, core temperature during treatment (open circles) is plotted against the average value of heat-treated and non-treated sides for each variable. Core was not included in correlation statistics. See text for details.

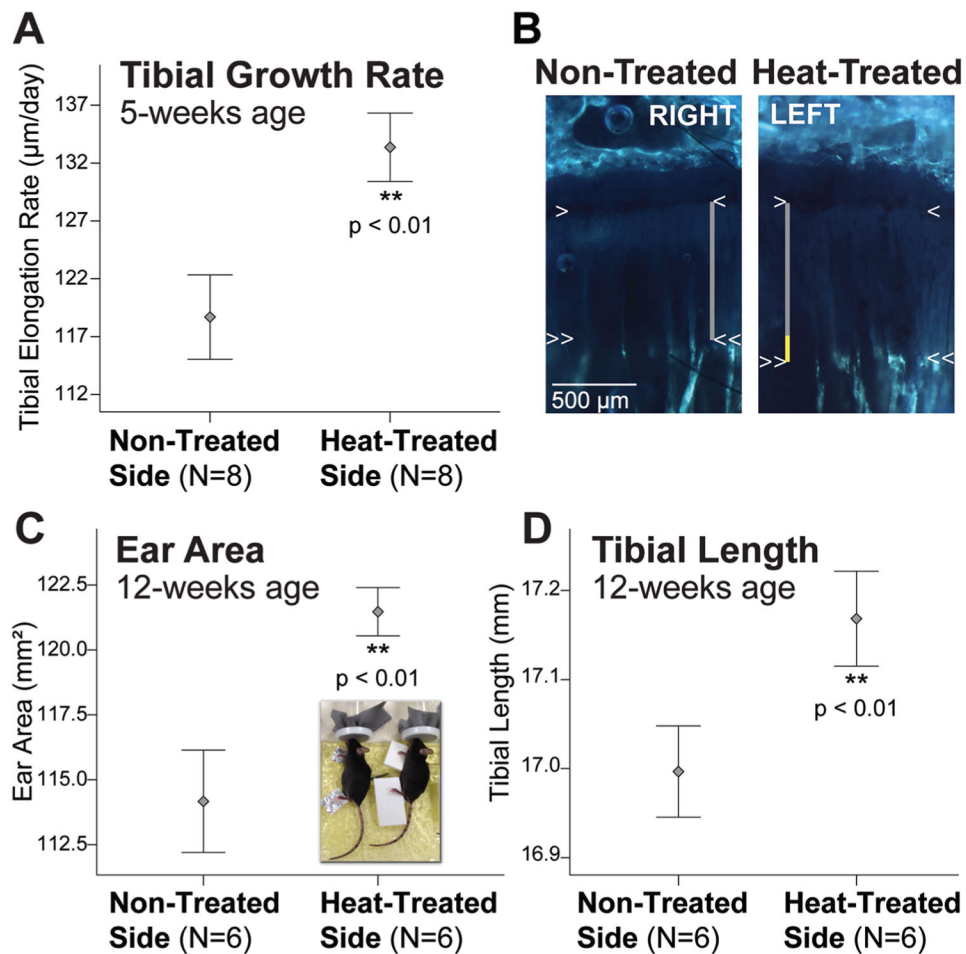


Figure 3. Extremities are lengthened on the heat-treated side. (A) Error bar plots show > 12% increase in tibial elongation rate on the heat-treated side. (B) Right-left tibial slab sections from the same mouse labeled with OTC. The metaphyseal chondro-osseous junction (COJ) is indicated by the single arrowheads. Double arrowheads show OTC band in metaphyseal bone. Growth rate was calculated by measuring the vertical distance between the arrowheads (gray lines). Yellow segment of the vertical line on the heat-treated side shows the total difference in length measured over 7-days. (C) Ear area and (D) tibial length remain significantly increased on the heat-treated side in skeletally mature adults after only 14 days of juvenile heat exposure (see Fig. 1). Mean \pm 1 standard error plotted.

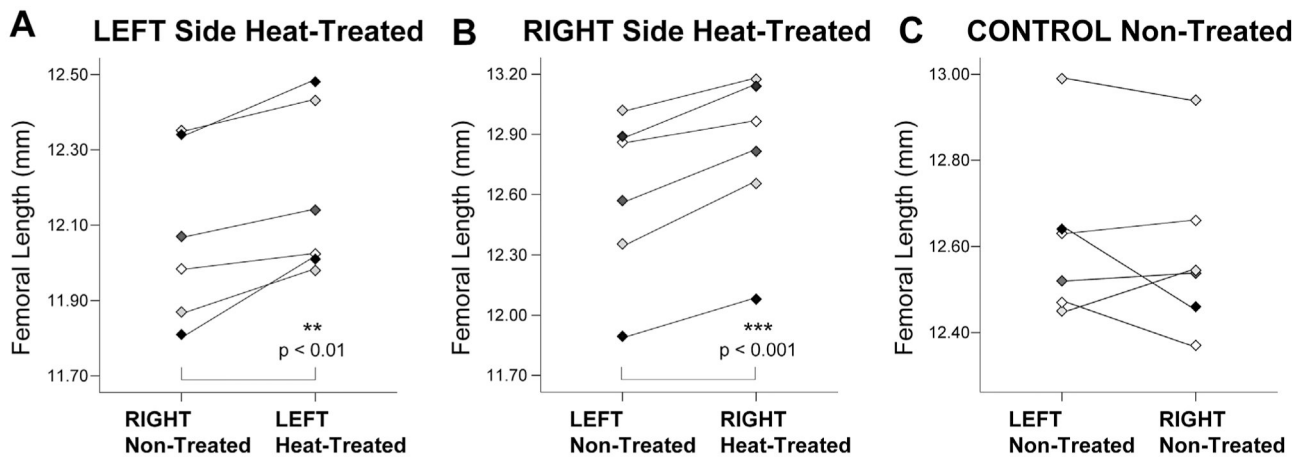


Figure 4.

Comparisons of individual mice treated on left and right sides versus non-treated control mice rule out potential side asymmetry. Paired comparisons of non-treated and heat-treated sides of mice that were treated on left (A) and right (B) show an average increase in femoral length on the heat-treated side of 1% and 1.6%, respectively. In contrast, left-right differences were absent in control mice (C).

Table 1. Comparison of Non-Treated and Heat-Treated Sides of 5-Week-Old Experimental Mice

Parameter	Non-Treated (30 °C)	Heat-Treated (40 °C)	Percent Increase	N
Ear Area (mm ²)	101.7 (6.9)	110.6 (7.2) ^{***}	8.8	14
Humeral Length (mm)	10.06 (0.24)	10.06 (0.25) ^{ns}	0	14
Femoral Length (mm)	12.33 (0.43)	12.49 (0.45) ^{***}	1.3	12
Tibial Length (mm)	14.96 (0.29)	15.18 (0.27) [*]	1.5	5
Hindfoot Length (mm)	17.54 (0.63)	18.15 (0.42) ^{***}	3.5	14
Tibial Elongation Rate (µm/day)	118.7 (10.3)	133.4 (8.4) ^{**}	12.4	8

Values are mean (standard deviation). Sample size (N) is number of left-right pairs. Significantly larger on heat-treated side by one-tailed paired *t*-test: * $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$; ns, non-significant.

Table 2.

Comparison of Left and Right Sides of Non-Treated 5-Week-Old Control Mice

Parameter	Non-Treated Left	Non-Treated Right	Percent Difference	N
Ear Area (mm ²)	98.7 (4.4)	98.7 (4.7)	0	6
Humeral Length (mm)	10.07 (0.20)	10.08 (0.19)	0	6
Femoral Length (mm)	12.62 (0.20)	12.59 (0.20)	0.2	6
Hindfoot Length (mm)	17.90 (0.22)	17.91 (0.25)	0	6
Tibial Elongation Rate ($\mu\text{m}/\text{day}$)	120.8 (6.6)	122.7 (7.6)	1.5	6

Values are mean (standard deviation). Sample size (N) is number of left-right pairs. There were no significant left-right differences by two-tailed paired *t*-tests (all $p > 0.40$).