

1 **Title** – The mechanics and energetics of hopping with elastic ankle exoskeletons

2 **Authors** – Dominic James Farris<sup>1</sup> & Gregory S. Sawicki<sup>1</sup>

3 **Author Contributions** - DJF & GSS both contributed to study conception, study design, data  
4 collection, data analysis and manuscript preparation.

5 **Affiliations** - <sup>1</sup>Joint Department of Biomedical Engineering, University of North Carolina-Chapel Hill &  
6 North Carolina state University, Raleigh, North Carolina, USA.

7 **Running header** - mechanics and energetics of hopping with elastic ankle exoskeletons

8 **Corresponding author** – Dominic Farris, Dept. Biomedical Engineering, NC State University, EB 3, 911  
9 Oval Drive, Raleigh, NC 27695-7115

10 Email: [djfarris@ncsu.edu](mailto:djfarris@ncsu.edu) Phone: (919)-928-2289 Fax: (919)-513-3814

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**13 Abstract**

14 The spring-like mechanics of the human leg during bouncing gaits has inspired the design of passive  
15 assistive devices that use springs to aid locomotion. The purpose of this study was to test if a passive  
16 spring-loaded ankle exoskeleton could reduce the mechanical and energetic demands of bilateral  
17 hopping on the musculoskeletal system. Joint level kinematics and kinetics were collected with  
18 electromyographic and metabolic energy consumption data for 7 participants hopping at 4  
19 frequencies (2.2, 2.5, 2.8 & 3.2 Hz). Hopping was performed without an exoskeleton; with a spring-  
20 less exoskeleton; and with a spring-loaded exoskeleton. Spring-loaded ankle exoskeletons reduced  
21 plantar-flexor muscle activity and the biological contribution to ankle joint moment (15-25%) and  
22 average positive power (20-40%). They also facilitated reductions in metabolic power (15-20%)  
23 across frequencies from 2.2-2.8 Hz compared to hopping with a spring-less exoskeleton. Reductions  
24 in metabolic power compared to hopping with no exoskeleton were restricted to hopping at 2.5 Hz  
25 only (12%). These results highlighted the importance of reducing the rate of muscular force  
26 production and work to achieve metabolic reductions. They also highlighted the importance of  
27 assisting muscles acting at the knee joint. Exoskeleton designs may need to be tuned to optimise  
28 exoskeleton mass, spring stiffness and spring slack length to achieve greater metabolic reductions.

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30 **Keywords** – metabolic power, spring-loaded, locomotion, power, EMG

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### 33 **Introduction**

34 When humans utilise bouncing gaits such as hopping and running, the supporting limb(s) during  
35 ground contact can be considered to act like a compressive spring (3, 13, 29). This has led to such  
36 motions commonly being modelled as a spring-mass system (12, 14, 19-20, 29). The spring-like  
37 behaviour of the lower limb makes it possible for the muscle-tendon units acting around joints to  
38 store and return energy in elastic structures (1, 6, 25, 28). Predominantly, energy is stored and  
39 returned in tendons acting in series with the contractile elements of muscles (16, 31). The result of  
40 this is a reduction in the mechanical work requirements of contractile elements and substantial  
41 metabolic energy savings (2, 7). Some metabolic cost is still incurred owing to the contractile  
42 elements having to produce force to stretch tendons (2, 31).

43 The way in which the muscle-tendon units of the leg utilise elastic energy cycling to efficiently power  
44 bouncing gaits has inspired the design of orthoses and exoskeletons for the leg that seek to passively  
45 store and return energy during movement (8, 18, 21). Such devices might be useful for rehabilitating,  
46 restoring and/or augmenting human locomotor performance by reducing musculo-skeletal loading  
47 and reducing metabolic energy consumption. Loading of the leg is expected to be reduced by  
48 wearing assistive exoskeletons because the device bears some of the load normally taken by the  
49 limb (18, 21). By using this loading to store and return energy in a spring, such devices may reduce  
50 the mechanical demands on biological tissues which may, in turn, reduce metabolic energy  
51 consumption (21).

52 Previous studies of the effects of passive lower-limb exoskeletons on human movement have  
53 typically used hopping motions because of the simple biomechanical goals of the task (8) and its  
54 similar spring-mass mechanics to running but, less complicated joint kinematics (18). Grabowski and  
55 Herr (21) showed that a full leg exoskeleton crossing hip, knee and ankle joints significantly reduced  
56 the metabolic power requirement of bilateral hopping (up to 30%) over a range of frequencies from  
57 2.0-2.6 Hz. Their participants reduced their biological contribution to leg stiffness to keep the

58 combined exoskeleton and biological stiffness similar to leg stiffness when hopping without an  
59 exoskeleton. This helped maintain consistent centre of mass mechanics which appears to be a goal  
60 of movement control for bouncing gaits (19-20).

61 A large proportion ( $\approx 45\%$ ) of the overall positive mechanical power output generated at leg joints  
62 during human running comes from muscle-tendon units acting at the ankle joint (17). Also, leg  
63 stiffness during hopping is predominantly determined by ankle joint stiffness (15). This highlights the  
64 importance of the ankle joint in maintaining the mechanics of bouncing gaits. Therefore, providing  
65 exoskeletal assistance at only the ankle joint could provide a significant proportion of the benefits  
66 achieved with a full leg device, while simplifying its design.

67 Both Ferris et al. (18) and Chang et al. (8) investigated the effects of wearing an ankle-foot orthosis  
68 that was spring-loaded to assist plantar-flexion on joint and limb stiffness during unilateral hopping.  
69 Both studies showed that hoppers were able to reduce biological ankle stiffness to maintain overall  
70 limb stiffness when wearing the devices. Ferris et al. (18) also showed a concurrent reduction in the  
71 amplitude of electromyographic (EMG) signals from the plantar-flexor muscles. This suggested that  
72 muscle activation was reduced to achieve lesser biological ankle joint stiffness when assistance was  
73 being provided by the device. Whilst the studies of Ferris et al. (18) and Chang et al. (8) both suggest  
74 that spring-loaded ankle exoskeletons might unload biological tissues, it was not the aim of these  
75 studies to link the mechanics of hopping in ankle exoskeletons to the associated metabolic cost.  
76 Therefore, they did not determine the mechanical demands and metabolic power requirements of  
77 the task concurrently. Doing so would help to reveal if assisting only at the ankle joint could reduce  
78 the mechanical demands on leg muscles sufficiently to reduce the metabolic cost of a bouncing gait.

79 The aim of this study was to investigate the effects of spring-loaded ankle exoskeletons on the  
80 metabolic cost and mechanics of bilateral human hopping. It was hypothesised that when hopping  
81 with the spring-loaded ankle exoskeletons, participants would reduce plantar-flexor muscle activity  
82 and that this would reduce the mechanical power output by biological tissues (muscle-tendon units)

83 at the ankle. Because of reduced mechanical demand placed on muscles, it was also hypothesised  
84 that hopping with spring-loaded exoskeletons would reduce metabolic power. Furthermore, it was  
85 expected that the ankle joint would dominate overall mechanical input to the task and thus,  
86 metabolic reductions would be proportionally less than those previously achieved with a full leg  
87 exoskeleton (21) (i.e., if the ankle joint provides 60% of the mechanical input, the observed  
88 reductions will be 60% of that with full leg assistance).

## 89 **Materials and Methods**

90 Seven male participants (mean  $\pm$  sd, age =  $28 \pm 7$ , height =  $1.8 \pm 0.06$  m, mass =  $80 \pm 10$  kg) gave  
91 written informed consent to participate in this study. All participants were in good health and had no  
92 recent history of lower limb musculo-skeletal injury. All procedures were approved by an  
93 institutional review board and complied with the guidelines for research involving human  
94 participants as set out in the Declaration of Helsinki.

95 Because metabolic power required for hopping and the effect of exoskeletons is frequency  
96 dependant (21), participants hopped in time with a metronome at four different frequencies (2.2  
97 Hz, 2.5 Hz, 2.8 Hz, 3.2 Hz). There were three different conditions: (1) Without an exoskeleton (NE);  
98 (2) With an exoskeleton but with no spring (NS); (3) With a spring-loaded exoskeleton to assist  
99 plantar-flexion (S). Each trial lasted 4 minutes. Because of the physically challenging nature of the  
100 task, each condition was performed on a separate day to minimise fatigue effects. The order of  
101 conditions and frequencies was randomized. The intent of this experiment was to test the effects of  
102 the exoskeletons on naïve users, so participants were given sufficient familiarisation with hopping in  
103 the exoskeletons to feel comfortable but not for long enough to introduce any training effects.

104 An example of the exoskeletons used is shown in Figure 1. The exoskeleton consisted of a carbon  
105 fibre cuff around the upper shank which was connected to a carbon fibre foot section via two  
106 aluminium bars which had a freely rotating joint aligned with the participants' malleoli. The foot

107 section was embedded in a training shoe, through the sole and around the heel. An extension spring  
108 could be attached to a bracket on the posterior aspect of the cuff and a bolt on the heel of the foot  
109 segment via a number of metal links. The number of links was adjusted for each participant such  
110 that the resting length of the spring coincided with an ankle angle of  $127^\circ$  which has been  
111 determined as the typical angle at ground contact in hopping (18). This same approach was used by  
112 Ferris et al. (18) for a similar exoskeleton. A compression load cell (Omegadyne Inc., OH,USA) was  
113 placed on the inferior side of the bolt at the heel of the foot segment and attached to the links in  
114 series with the spring. This was used to measure the forces in the spring. The stiffness of the spring  
115 in tension was  $5 \text{ kNm}^{-1}$  and its moment arm about the joints was 0.135 m. This gave a rotational  
116 stiffness of  $1.59 \text{ Nm}/^\circ$  ( $91 \text{ Nm/rad}$ ) which is approximately 40% of ankle stiffness during unassisted  
117 hopping at preferred frequency (18).

118 An eight camera motion analysis system (Vicon, Oxford, UK) was used to capture the three-  
119 dimensional positions of 22 reflective markers attached to the pelvis and right leg. Raw marker  
120 positions were filtered using a second order low-pass butterworth filter with a cut-off of 10 Hz. A  
121 static standing trial was captured and the positions of markers on segment end points were used to  
122 calibrate a four segment (pelvis, thigh, shank and foot) model for each subject using established  
123 inertial parameters (10). Clusters of three or four markers on rigid plates were attached to the pelvis,  
124 thigh and shank segments to track segment motion during hopping. For the foot, a cluster of three  
125 markers was attached directly to the shoe. Joint angles for the hip knee and ankle were computed in  
126 three dimensions as the orientation of the distal segment with reference to the proximal segment  
127 and differentiated to calculate joint velocities.

128 Three-dimensional (3D) ground reaction forces applied to the left and right legs were computed  
129 during hopping using a split belt instrumented treadmill (Bertec, OH, USA). Participants hopped such  
130 that each foot was on a separate half of the treadmill and thus, the two 3D force vectors could be  
131 attributed separately to the left and right legs. Raw analogue force platform signals were filtered

132 using a low-pass Butterworth digital filter with the cut-off set to 35 Hz. Inverse dynamic analyses  
 133 (Winter, 1990) were then used to compute net joint moments which were multiplied with joint  
 134 velocities to calculate joint powers at the hip, knee and ankle. Kinematics and kinetics were  
 135 calculated for the right leg only and it was assumed that the left leg behaved symmetrically. Inverse  
 136 dynamics procedures were performed with Visual 3D software (C-motion Inc., Germantown, MD,  
 137 USA).

138 The mechanical demands of hopping under different conditions was partly assessed by the average  
 139 positive power ( $\bar{P}_{mech}^+$ ) of the task. More details of the computation of  $\bar{P}_{mech}^+$  can be found  
 140 elsewhere (17). Briefly, it was calculated for each joint [ankle ( $\bar{P}_{ank}^+$ ), knee ( $\bar{P}_{knee}^+$ ) and hip ( $\bar{P}_{hip}^+$ )] as  
 141 the integral of all periods of instantaneous positive power for that joint during a hop, divided by the  
 142 time per hop.  $\bar{P}_{ank}^+$ ,  $\bar{P}_{knee}^+$  and  $\bar{P}_{hip}^+$  were summed to compute total average positive power ( $\bar{P}_{tot}^+$ ).  
 143 When the spring-loaded exoskeletons were being worn, only a portion of  $\bar{P}_{ank}^+$  was provided by  
 144 biological tissues and the rest came from the exoskeleton. To separate out these contributions, the  
 145 power output of the exoskeleton was determined as follows. First, the force in the spring (measured  
 146 by the load cell, Figure 1) was multiplied by the spring's moment arm about the joint of the  
 147 exoskeleton, to give the moment provided by the spring. This moment was multiplied by ankle  
 148 angular velocity to compute the instantaneous power provided by the exoskeleton. The average  
 149 positive power of the exoskeleton ( $\bar{P}_{exo}^+$ ) was calculate as it was for joints. The contribution of  
 150 biological tissues at the ankle ( $\bar{P}_{bio}^+$ ) was calculated as  $\bar{P}_{ank}^+$  minus  $\bar{P}_{exo}^+$ . All average powers were  
 151 normalised to body mass ( $W \cdot kg^{-1}$ ).

152 The rate at which muscles produce force may also be an important mechanical determinant of  
 153 metabolic cost (27). Therefore, a rate of force metric was also computed. This metric was the  
 154 average rate of joint moment production ( $\bar{M}^+$ ). Somewhat analogous to average positive power, this  
 155 value was calculated for each joint [ankle ( $\bar{M}_{ank}^+$ ), knee ( $\bar{M}_{knee}^+$ ) and hip ( $\bar{M}_{hip}^+$ )] by integrating the  
 156 first derivative of each joint moment during periods of increasing moment production (i.e. when the

157 derivative was positive), summing each of the integral values during each hop and dividing the result  
158 by the time taken for an entire hop cycle. As for average positive power,  $\bar{M}_{ank}^+$  was reduced to  
159 separate exoskeleton and biological contributions when a spring-loaded exoskeleton was being  
160 used.

161 Surface electromyography (EMG) was used to record muscle activity from Medial Gastrocnemius  
162 (MG), Lateral Gastrocnemius (LG), Soleus (SO) and Tibialis Anterior (TA). All four channels were  
163 recorded using wired electrodes (Biometrics Ltd, UK) that were carefully placed over muscle bellies  
164 after the skin surface was prepared by light abrasion and cleaned with an alcohol swab. D.C. offsets  
165 were removed from raw signals which were then band-pass filtered (20-300 Hz). The data were then  
166 smoothed by calculating the root-mean-squared (RMS) value of the signals over a rolling window of  
167 20 ms. The RMS of each muscle's signal was also calculated over the period of ground contact and  
168 the aerial phase of each hop as a metric of total activity over these two phases of the hop. For each  
169 session of data collection, the participant performed a set of 20 hops at 2.2 Hz while wearing  
170 exoskeletons without a spring. The maximum of the processed signals from each muscle for this  
171 condition was used to normalise EMG signals from the experimental conditions on that day. This was  
172 to avoid any effects of day-to-day variation in electrode placement on signal magnitudes.

173 Rates of oxygen consumption and carbon dioxide production during hopping trials were recorded  
174 using a portable metabolic system (OXYCON MOBILE, Viasys Healthcare, CA, USA). Prior to hopping,  
175 measurements were made during five minutes of quiet standing and values from the last two  
176 minutes were averaged and used to calculate rates of metabolic energy consumption whilst  
177 standing. For the hopping trials, data from the last two of the four minutes were averaged for the  
178 calculation of metabolic rate. Visual inspection of rates of oxygen consumption with time (averaged  
179 over 30 s intervals) confirmed that participants were at steady-state during this period and the  
180 respiratory exchange ratio was never greater than one. Rates of oxygen consumption and carbon  
181 dioxide production were converted to metabolic powers using standard equations detailed by



182 Brockway (5). Net metabolic powers during hopping were calculated by subtracting metabolic power  
183 during standing from metabolic power during hopping and these values were normalized to  
184 individual body mass ( $W \cdot kg^{-1}$ ). Metabolic data were presented as the normalised net value, unless  
185 otherwise stated.

186 All the kinematic and kinetic data for individual participants was reduced to the mean of at least 10  
187 hops for each experimental condition. Unless otherwise stated, the values presented in this paper  
188 are the mean  $\pm$  standard error for the whole participant group. To test for statistical differences in  
189 dependent variables between conditions a two-way ANOVA with repeated measures was employed  
190 using SPSS software (IBM,USA). The two independent variables for the ANOVA were: 1. Spring  
191 condition (3 levels – NE, NS, S) and 2. Hopping Frequency (4 levels – 2.2, 2.5, 2.8, 3.2 Hz). F-ratios for  
192 main and interaction effects were considered significant for  $P < 0.05$ . If a significant main effect was  
193 found, paired t-tests were used to make pair-wise comparisons between levels of independent  
194 variables.

## 195 **Results**

196 The S condition was described by similar hip and knee joint kinematics to the other NS and NE  
197 conditions but it did cause a shift in ankle angle to a generally more plantar-flexed position (Figure  
198 2). This shift was systematic, allowing the ankle to move through a similar range of motion ( $15-30^\circ$ ,  
199 dependant on frequency) in all conditions but with a more plantar-flexed touchdown angle for S.  
200 Similarly, joint kinetics (moments and powers) were similar across the NE, NS and S conditions for  
201 the hip and knee (Figures 3 and 4). Total ankle power had generally greater peaks in the S condition  
202 (Figure 3) indicating that the sum of exoskeleton power and biological ankle power for S was greater  
203 than the biological power in NE and NS. The ankle joint moments showed that the exoskeleton  
204 provided between 30 and 50% of the total moment (Figure 3) and this significant contribution was  
205 confirmed in the  $\bar{P}_{mech}^+$  data which showed that  $\bar{P}_{exo}^+$  accounted for 36-38% of  $\bar{P}_{tot}^+$  while  $\bar{P}_{bio}^+$   
206 accounted for 23-38% (Figure 5).

207 Net metabolic power typically followed a U-shaped relationship with frequency for NE, NS and S  
208 (Figure 6). This meant that net metabolic power requirements were least for the intermediate  
209 frequencies (2.5 & 2.8 Hz). Notably, there was a relatively large increase in net metabolic power  
210 from 2.8 Hz to 3.2 Hz in the S condition (Figure 6). This caused the mean value for net metabolic  
211 power for S to be greater than for NE at 3.2 Hz when it had been consistently less than NE and NS at  
212 all other frequencies. Net metabolic power was significantly less for S than NS at 2.2 (-16%,  $P=0.009$ ),  
213 2.5 (-19%,  $P = 0.006$ ) and 2.8 Hz (-20%,  $P = 0.047$ ) but only at 2.5 Hz (-13%,  $P = 0.016$ ) compared to  
214 NE (Figure 6).

215 The biological contribution to  $\bar{P}_{tot}^+$  was significantly less for S than for NS ( $\approx 30$ -40%) and NE ( $\approx 20$ -  
216 30%) at all frequencies of hopping (Figure 6b, ANOVA  $P < 0.001$ ). As can be observed in Figure 6b,  
217 this difference was greater at low frequencies than at high frequencies with the linear fits starting to  
218 converge at high frequencies. Biological  $\bar{P}_{mech}^+$  was greater at low frequencies than at high  
219 frequencies for S, NS and NE. This was in contrast to the biological contribution to  $\bar{M}_{ank}^+$  which  
220 increased with hopping frequency. There was a significant difference in  $\bar{M}_{ank}^+$  between the S than in  
221 NS conditions at 2.2 (21%,  $P < 0.001$ ), 2.5 (24%,  $P = 0.003$ ) and 2.8 Hz (20%,  $P = 0.001$ ) (Figure 6b).  
222  $\bar{M}_{ank}^+$  was only significantly less for S than NS at 2.2 (18%,  $P = 0.007$ ) and 2.5 Hz (19%,  $P = 0.05$ )  
223 (Figure 6c).

224 Time histories of the smoothed and normalised EMG signals are shown in Figure 7. MG, LG and SO  
225 all typically had lower activations toward the end of the aerial phase for S than for NE and NS. This  
226 was apparent in the RMS values for these muscles during the aerial phase which were typically  
227 significantly less for S than for NS and NE (Figure 8). During ground contact only SO had significantly  
228 lower activation for S (Figure 8). Contrary to the plantar-flexor activations, TA exhibited significantly  
229 increased activation during ground contact and the aerial phase for all frequencies in the S condition  
230 compared to NE and NS (Figure 8).

231

## 232 Discussion

233 This study aimed to investigate the effects of spring-loaded ankle exoskeletons on the metabolic cost  
234 and mechanics of bilateral human hopping. First, it was hypothesised that the use of bilateral  
235 spring-loaded ankle exoskeletons during two-legged hopping would reduce ankle plantar-flexor  
236 muscle activity. This prediction was supported for all the plantar-flexors tested during the aerial  
237 phase but only soleus during ground contact. It was also hypothesised that the reduced activity  
238 would reduce the contribution of plantar-flexors to mechanical power output. This was found to be  
239 the case across all frequencies. The final prediction made was that using spring-loaded ankle  
240 exoskeletons would lower the metabolic power requirement of hopping when compared to hopping  
241 without an exoskeleton or without a spring. This prediction was supported for some conditions at  
242 certain frequencies but not all. These findings raise interesting questions regarding the mechanisms  
243 by which spring-loaded ankle exoskeletons may be able to lower metabolic cost and also highlights  
244 some potential obstacles to this goal.

245 The results showed that the use of spring-loaded ankle exoskeletons resulted in significant  
246 reductions in the  $\bar{P}_{mech}^+$  production by biological tissues (muscle-tendon units) when hopping at all  
247 frequencies compared to the NE and NS conditions (Figure 6). This was due to a reduction in the  
248 biological plantar flexion moment at the ankle (Figure 3). This result was not surprising as it has been  
249 previously shown that humans reduce biological ankle and limb stiffness to maintain overall system  
250 stiffness when hopping with spring-loaded ankle or full leg exoskeletons (8, 18, 21). The reduction in  
251 biological contribution to the plantar-flexion moment was primarily achieved by a reduction in  
252 soleus muscle activity during the ground contact phase of hops (Figure 8). MG and LG RMS EMG  
253 values were only less during the aerial phase and this was primarily due to a reduction in activity late  
254 in this phase (a similar trend was apparent in soleus for the aerial phase). Therefore it seems that  
255 pre-activation of plantar-flexors was also less when using spring-loaded exoskeletons.

256 The reduction in biological  $\bar{P}_{mech}^+$  seen at all frequencies in the S condition was one indication that  
257 the mechanical demands on biological tissues was less with assistance. However, significant  
258 reductions in metabolic power only occurred at 2.5 Hz ( $-0.9 \text{ W}\cdot\text{kg}^{-1}$ ) compared to the NE condition  
259 and at 2.2 ( $-1.3 \text{ W}\cdot\text{kg}^{-1}$ ), 2.5 ( $-1.5 \text{ W}\cdot\text{kg}^{-1}$ ) and 2.8 Hz ( $-1.6 \text{ W}\cdot\text{kg}^{-1}$ ) compared to the NS condition.  
260 Furthermore, where there were reductions in metabolic power, they were not proportional with  
261 reductions in biological  $\bar{P}_{mech}^+$  (Figure 6). Therefore, whilst  $\bar{P}_{mech}^+$  provides some indication of the  
262 mechanical demands of the task, it alone cannot fully explain the trends in metabolic cost that were  
263 observed. This is particularly notable at 3.2 Hz where the metabolic power requirement was greater  
264 for the S condition than for the NE condition despite a reduction in  $\bar{P}_{mech}^+$ .

265 Another factor that may help to explain the observed trends in metabolic power is the metabolic  
266 cost associated with producing muscular force. An increasing number of studies of human (and  
267 animal) movement have indicated that a significant portion (up to 50%) of the metabolic cost of  
268 locomotion is related to the metabolic energy used by muscles to produce force (22-23, 26-27, 32).  
269 This cost has been proposed to increase proportionally with the rate at which force must be  
270 produced (27). The cost of producing force may be particularly important in muscle groups such as  
271 the plantar-flexors that rely on the series compliance of tendons to cycle energy while the muscle  
272 itself contracts relatively isometrically during stance (25, 28). In such contractions, the contractile  
273 elements of muscle do only small amounts of work but must produce forces sufficient to 'anchor'  
274 the tendon at the myotendinous junction. In a study of human bouncing tasks, Dean and Kuo (9)  
275 actually showed that the rate of force cycling in ankle plantar-flexors increased with bouncing  
276 frequency and that this could explain why metabolic cost began to increase at frequencies above 3  
277 Hz, despite muscular work decreasing. If the rate of force production becomes a significant factor in  
278 determining metabolic cost at high frequencies ( $>3 \text{ Hz}$ ), it might explain why our spring-loaded  
279 exoskeletons were unable to reduce metabolic power at 3.2 Hz despite reducing biological  
280 mechanical power (assuming similarities between bouncing and hopping). In the present study  $\bar{M}_{ank}^+$

281 was considered as a metric for rate of force production by the muscles acting at the ankle joint.  
282 Figure 6 shows that  $\bar{M}_{ank}^+$  increased with frequency lending some support to the notion that it  
283 becomes more influential at higher frequencies. In fact, at the two highest frequencies in this study,  
284  $\bar{M}_{ank}^+$  could explain the trends in metabolic power. At 3.2 Hz no significant differences in metabolic  
285 power were observed and there were also no significant reductions in  $\bar{M}_{ank}^+$  (Figure 6). For hopping  
286 at 2.8 Hz there was a significant reduction in metabolic power for S compared to NS that coincided  
287 with a reduction in  $\bar{M}_{ank}^+$  but neither variable was reduced compared to NE (Figure 6). Therefore, at  
288 higher frequencies metabolic power seems more dependent on  $\bar{M}_{ank}^+$  than  $\bar{P}_{mech}^+$  and at 3.2 Hz the  
289 exoskeletons did not reduce  $\bar{M}_{ank}^+$  sufficiently.

290 However, at lower frequencies this may not be the case. For both 2.2 Hz and 2.5 Hz there were  
291 significant reductions in both  $\bar{P}_{mech}^+$  and  $\bar{M}_{ank}^+$  for S compared to NE and NS but metabolic cost was  
292 not reduced at 2.2 Hz compared to NE (Figure 6). A possible explanation for this might be related to  
293 the energy consumed by more proximal muscle groups. Although the muscles acting at the ankle  
294 joint were responsible for the majority of power output, a non-trivial contribution was provided by  
295 knee extensors (18-46%) and this was greatest at 2.2 Hz (Figure 5). Work provided by more proximal  
296 muscle-tendon units has been postulated to be done less efficiently (33). This is owing to their lesser  
297 ability to make use of series elastic compliance than muscle groups such as the plantar-flexors.  
298 Therefore, although the contributions of more proximal muscle groups to power output was less  
299 than at the ankle, they probably provided this power less efficiently. Thus, they may be responsible  
300 for a larger proportion of the total energy consumed than mechanical power output suggests.  
301 Because the muscles acting at the knee were providing a greater proportion of the total positive  
302 power at 2.2 Hz, the effectiveness of the exoskeletons at reducing overall mechanical demands may  
303 have been diminished. This may have contributed to the lack of significant reductions in metabolic  
304 power compared to NE.

305 The influence of more proximal muscle groups was highlighted by the lesser reductions in metabolic  
306 power that were achieved in this study than by a full leg exoskeleton in a previous study (21). With a  
307 full leg exoskeleton, reductions were in the region of  $2.0\text{-}2.5\text{ W}\cdot\text{kg}^{-1}$  ( $\approx 30\%$ ) (21). By comparison, in  
308 the present study the greatest reductions observed were approximately  $1.5\text{ W}\cdot\text{kg}^{-1}$  ( $\approx 20\%$ ).  
309 Therefore, it seems that it would be beneficial to assist at more proximal joints as well as the ankle.  
310 As can be seen from figure 5, the contribution of the hip to overall mechanical power output was  
311 minimal. Therefore it might be that assisting at the ankle and knee joints would provide the desired  
312 reductions in metabolic cost without requiring a device that assists at the hip too. However, this may  
313 only apply to hopping in place which requires no forward swing of the leg.

314 Another further explanation for the lesser metabolic power reductions observed here may be  
315 apparent in the RMS EMG data for TA. There was a significant increase in TA RMS EMG during both  
316 the aerial and ground contact phases of hopping across all frequencies when hopping with the  
317 spring-loaded exoskeletons. Particularly notable was the 2-3 fold increase in normalised EMG signal  
318 magnitude during the aerial phase (Figures 7 and 8). In the NE and NS conditions there was apparent  
319 pre-activation of the plantar-flexors prior to ground contact that was opposed by some co-activation  
320 of TA (Figure 7). In this case, TA may be serving to stiffen the ankle joint to allow pre-activation of  
321 the plantar-flexors without generating rotation at the ankle joint (there was  $<5^\circ$  of rotation at the  
322 ankle during the aerial phase despite considerable plantar-flexor activity). When hopping with  
323 spring-loaded exoskeletons, the pre-activation of the plantar-flexors was less (Figures 7 and 8) and  
324 this resulted in significantly smaller RMS EMG values for the plantar-flexors during the aerial phase  
325 (Figure 8). This might lead one to think that TA should have exhibited lesser activation at the same  
326 time to maintain equilibrium of the force couple at the ankle joint and consistent kinematics.  
327 However, as shown in Figure 2, the spring was actually exerting force to generate a plantar-flexion  
328 moment during the aerial phase. Therefore, TA had to be active to oppose this moment and was  
329 effectively co-activating antagonistically with the spring. It was inferred from the raised activation of  
330 TA in the spring-loaded condition that the moment generated by the spring during the aerial phase

331 was greater than that generated by the plantar-flexors in the NS and NE conditions. A consequence  
332 of raised TA co-activation could be increased metabolic energy consumption that partially cancelled  
333 out the benefits of reduced plantar-flexor activation. Elevated levels of antagonistic co-activation has  
334 been cited as a factor contributing to increased energy consumption for locomotion in older  
335 populations (24, 30). Therefore, the elevated activation of TA to work against the spring during the  
336 aerial phase may have contributed to there being lesser metabolic power reductions than were  
337 expected.

338 The apparent engagement of the spring during the aerial phase of hops likely represents a design  
339 flaw of the exoskeletons. The spring was attached such that its slack length would be reached when  
340 the ankle was at  $127^\circ$  of flexion. This angle was chosen from a previous study using similar devices as  
341 the typical angle at touchdown (18). Thus, it was intended that the spring would remain slack (i.e.  
342 exerting no force) until contact was made with the ground and it could be used to store energy  
343 transferred from kinetic and potential energy of the body. However, Figure 2 shows that the  
344 touchdown angle for the NS and NE conditions was less (more dorsi-flexed) than  $127^\circ$  ( $\approx 116-125^\circ$ ).  
345 Extension of the spring would have been required to reach these angles when wearing the spring-  
346 loaded exoskeletons. As can be seen in Figure 2, the touchdown angle of the ankle was greater  
347 (more plantar-flexed) in the spring-loaded condition and at  $\approx 132^\circ$  should have placed the spring  
348 below its slack length. However, the spring was still exerting force prior to touchdown and thus must  
349 have been longer than its slack length. The results indicate that the angle at which the spring  
350 engages is an important factor and should be optimised (possibly on an individual basis) to maximise  
351 energy storage but minimise aerial co-activation of TA. Alternatively, a clutching mechanism that  
352 selectively engages the spring after landing and disengages it after take-off might solve this issue.  
353 Such mechanisms are useful for walking (34) and running (11) with spring-loaded assistive devices,  
354 to prevent wearers from working against springs during the swing-phases of gait.

355 Other aspects of the exoskeleton that should be considered as ‘tuneable’ are its mass and spring  
356 stiffness. Mass is a relatively simple concept in that it is a matter of making the exoskeletons as light  
357 as possible. The exoskeletons used here were prototypes and thus were not fine-tuned to be as light  
358 as would be desirable. The added mass of the exoskeletons adds mechanical demands and  
359 associated metabolic cost to the task. In fact, it can be seen from the size of the pies in Figure 5 that  
360 overall mechanical power output was greater in the NS and the S conditions than for NE. When  
361 drawing comparisons between the S and NS conditions, significant reductions in metabolic cost were  
362 observed at 2.2, 2.5 and 2.8 Hz. This indicated that the assistance provided by the spring was  
363 typically sufficient to counteract the added mass of the exoskeleton, but only improved upon energy  
364 consumption of normal hopping at 2.5 Hz. Reductions in metabolic cost relative to the NE condition  
365 would be expected to reach similar values to reductions compared to the NS condition as the mass  
366 of the device approaches zero.

367 With regard to spring stiffness, it may be necessary to design experiments to try and find an optimal  
368 stiffness for metabolic reductions. The spring chosen for this study was selected as the stiffest spring  
369 in which a pilot subject could complete the task. It was considered that this was the spring that  
370 would provide maximal unloading of biological tissues. However, maximal unloading does not  
371 necessarily correspond to the greatest reductions in metabolic cost (4). In fact, Grabowski and Herr  
372 (21) showed with their full-leg exoskeleton that a stiffer spring can increase the net metabolic  
373 power required for hopping compared to a less stiff spring. They proposed that this was because the  
374 stiffer exoskeleton may have compromised stability and/or sensory feedback. As a proxy for stability  
375 Grabowski and Herr (21) calculated the distance moved by the centre of pressure between  
376 consecutive hops and showed that this distance was typically larger for the stiffer exoskeleton. In  
377 this study the same metric was calculated (dHop) and was typically greater for the spring-loaded  
378 condition, suggesting that participants were less stable in that condition. Therefore, although  
379 intuition might suggest that a greater stiffness (up to the normal ankle stiffness) would be more  
380 beneficial, it might actually impair the person’s ability to hop. It seems plausible that there would be



381 some optimal stiffness that maximises energy storage and return without significantly impairing  
382 stability and sensory feedback. Furthermore, this optimal stiffness might vary with hopping  
383 frequency and between individuals depending on their body mass and stature. Careful studies  
384 designed to explore these factors are still needed to fine-tune the exoskeleton design.

385 This study showed that spring-loaded ankle exoskeletons can be used to reduce the metabolic cost  
386 of bilateral human hopping at certain frequencies. It has highlighted the potential for such devices to  
387 assist with bouncing gaits and some of the important design problems that must be addressed.  
388 Future studies should aim to investigate optimal spring stiffness and engagement angle. They should  
389 also consider how the exoskeletons interact with the underlying muscle-tendon units and how they  
390 might be re-designed to be used in running where the kinematics become more complex than for  
391 hopping.

#### 392 **Acknowledgements**

393 The authors would like to thank Ben Robertson (North Carolina State University) for assistance with  
394 data collection.

#### 395 **Disclosures**

396 The authors declare no conflicts of interest in relation to this study

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#### 398 **References**

- 399 1. **Alexander R.** *Elastic Mechanisms in Animal Movement*. Cambridge: Cambridge University  
400 Press, 1988.
- 401 2. **Biewener AA, and Roberts TJ.** Muscle and tendon contributions to force, work, and elastic  
402 energy savings: a comparative perspective. *Exercise and Sport Sciences Reviews* 28: 99-107, 2000.
- 403 3. **Blickhan R.** THE SPRING MASS MODEL FOR RUNNING AND HOPPING. *Journal of*  
404 *Biomechanics* 22: 1217-1227, 1989.

- 405 4. **Bregman DJJ, van der Krogt MM, de Groot V, Harlaar J, Wisse M, and Collins SH.** The effect  
406 of ankle foot orthosis stiffness on the energy cost of walking: A simulation study. *Clinical*  
407 *Biomechanics* 26: 955-961, 2011.
- 408 5. **Brockway JM.** DERIVATION OF FORMULAS USED TO CALCULATE ENERGY-EXPENDITURE IN  
409 MAN. *Human Nutrition-Clinical Nutrition* 41C: 463-471, 1987.
- 410 6. **Cavagna GA.** Storage and utilization of elastic energy in skeletal muscle. *Exercise and Sport*  
411 *Sciences Reviews* 5: 89-129, 1977.
- 412 7. **Cavagna GA, and Kaneko M.** Mechanical work and efficiency in level walking and running.  
413 *Journal of Physiology* 268: 467-481, 1977.
- 414 8. **Chang YH, Roiz RA, and Auyang AG.** Intralimb compensation strategy depends on the nature  
415 of joint perturbation in human hopping. *Journal of Biomechanics* 41: 1832-1839, 2008.
- 416 9. **Dean JC, and Kuo AD.** Energetic costs of producing muscle work and force in a cyclical  
417 human bouncing task. *Journal of Applied Physiology* 110: 873-880, 2011.
- 418 10. **Dempster AD.** Space requirements of the seated operator Ohio: Wright-Patterson Air Force  
419 Base, 1955.
- 420 11. **Dollar AM, and Herr H.** Design of a quasi-passive knee exoskeleton to assist running. In: *IEEE*  
421 *international conference on intelligent robots and systems*. Nice, France: IEEE, 2008, p. 747-754.
- 422 12. **Farley CT, Blickhan R, McMahon TA, and Taylor CR.** MECHANICS OF HUMAN HOPPING.  
423 *Journal of Biomechanics* 20: 896-896, 1987.
- 424 13. **Farley CT, Blickhan R, Saito J, and Taylor CR.** HOPPING FREQUENCY IN HUMANS - A TEST OF  
425 HOW SPRINGS SET STRIDE FREQUENCY IN BOUNCING GAITS. *Journal of Applied Physiology* 71: 2127-  
426 2132, 1991.
- 427 14. **Farley CT, Glasheen J, and McMahon TA.** RUNNING SPRINGS - SPEED AND ANIMAL SIZE.  
428 *Journal of Experimental Biology* 185: 71-86, 1993.
- 429 15. **Farley CT, and Morgenroth DC.** Leg stiffness primarily depends on ankle stiffness during  
430 human hopping. *Journal of Biomechanics* 32: 267-273, 1999.

- 431 16. **Farris DJ, and Sawicki GS.** Human medial gastrocnemius force–velocity behavior shifts with  
432 locomotion speed and gait. *Proceedings of the National Academy of Sciences* 109: 977-982, 2012.
- 433 17. **Farris DJ, and Sawicki GS.** The mechanics and energetics of human walking and running: a  
434 joint level perspective. *Journal of The Royal Society Interface* 9: 110-118, 2012.
- 435 18. **Ferris DP, Bohra ZA, Lukos JR, and Kinnaird CR.** Neuromechanical adaptation to hopping  
436 with an elastic ankle-foot orthosis. *Journal of Applied Physiology* 100: 163-170, 2006.
- 437 19. **Ferris DP, and Farley CT.** Interaction of leg stiffness and surface stiffness during human  
438 hopping. *Journal of Applied Physiology* 82: 15-22, 1997.
- 439 20. **Ferris DP, Louie M, and Farley CT.** Running in the real world: adjusting leg stiffness for  
440 different surfaces. *Proceedings of the Royal Society of London Series B-Biological Sciences* 265: 989-  
441 994, 1998.
- 442 21. **Grabowski AM, and Herr HM.** Leg exoskeleton reduces the metabolic cost of human  
443 hopping. *Journal of Applied Physiology* 107: 670-678, 2009.
- 444 22. **Grabowski AM, and Kram R.** Effects of Velocity and Weight Support on Ground Reaction  
445 Forces and Metabolic Power During Running. *Journal of Applied Biomechanics* 24: 288-297, 2008.
- 446 23. **Griffin TM, Roberts TJ, and Kram R.** Metabolic cost of generating muscular force in human  
447 walking: insights from load-carrying and speed experiments. *Journal of Applied Physiology* 95: 172-  
448 183, 2003.
- 449 24. **Hortobagyi T, Finch A, Solnik S, Rider P, and DeVita P.** Association Between Muscle  
450 Activation and Metabolic Cost of Walking in Young and Old Adults. *Journals of Gerontology Series a-  
451 Biological Sciences and Medical Sciences* 66: 541-547, 2011.
- 452 25. **Ishikawa M, Pakaslahti J, and Komi PV.** Medial gastrocnemius muscle behaviour during  
453 human running and walking. *Gait and Posture* 25: 380-384, 2007.
- 454 26. **Kram R.** Muscular force or work: what determines the metabolic energy cost of running?  
455 *Exercise and Sport Sciences Reviews* 28: 138-143, 2000.

- 456 27. **Kram R, and Taylor CR.** ENERGETICS OF RUNNING - A NEW PERSPECTIVE. *Nature* 346: 265-  
457 267, 1990.
- 458 28. **Lichtwark GA, Bougoulias K, and Wilson AM.** Muscle fascicle and series elastic element  
459 length changes along the length of the human gastrocnemius during walking and running. *Journal of*  
460 *Biomechanics* 40: 157 - 164, 2007.
- 461 29. **McMahon TA, and Cheng GC.** THE MECHANICS OF RUNNING - HOW DOES STIFFNESS  
462 COUPLE WITH SPEED. *Journal of Biomechanics* 23: 65-78, 1990.
- 463 30. **Peterson DS, and Martin PE.** Effects of age and walking speed on coactivation and cost of  
464 walking in healthy adults. *Gait & Posture* 31: 355-359, 2010.
- 465 31. **Roberts TJ.** The integrated function of muscles and tendons during locomotion. *Comparitive*  
466 *Biochemistry and Physiology part A* 133: 1087-1099, 2002.
- 467 32. **Roberts TJ, Kram R, Weyand PG, and Taylor CR.** Energetics of bipedal running I. Metabolic  
468 cost of generating force. *Journal of Experimental Biology* 201: 2745-2751, 1998.
- 469 33. **Sawicki GS, Lewis CL, and Ferris DP.** It Pays to Have a Spring in Your Step. *Exercise and Sport*  
470 *Sciences Reviews* 37: 130-138, 2009.
- 471 34. **Wiggin MB, Collins SH, and Sawicki GS.** An exoskeleton using controlled energy storage and  
472 release to aid ankle propulsion. In: *IEEE international conference on rehabilitation robotics*. Zurich,  
473 Switzerland: IEEE, 2011, p. 1-5.
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**480 Figure Captions**

481 Figure 1. Sketch of the spring-loaded exoskeleton and load-cell

482 Figure 2. Group mean ankle (A), knee (B) and hip (C) angles for each frequency, plotted over an  
483 average hop (normalised over 101 points). The left column is the no exoskeleton (NE) condition,  
484 middle is the no spring (NS) condition and right is the spring-loaded (S) condition. Vertical lines  
485 represent the range of normalised take-off times across frequencies.

486 Figure 3. Group mean ankle (A), knee (B) and hip (C) moments plotted over an average hop  
487 (normalised over 101 points). The additional data series of smaller magnitude on the S ankle plot are  
488 the exoskeleton moments for the corresponding frequencies and the hatched area highlights the  
489 contribution from biological tissues to the total moment for 2.2 Hz. The left column is the No  
490 Exoskeleton (NE) condition, middle is the no spring (NS) condition and right is for the spring-loaded  
491 (S) condition. Vertical lines represent the range of normalised take-off times across frequencies.

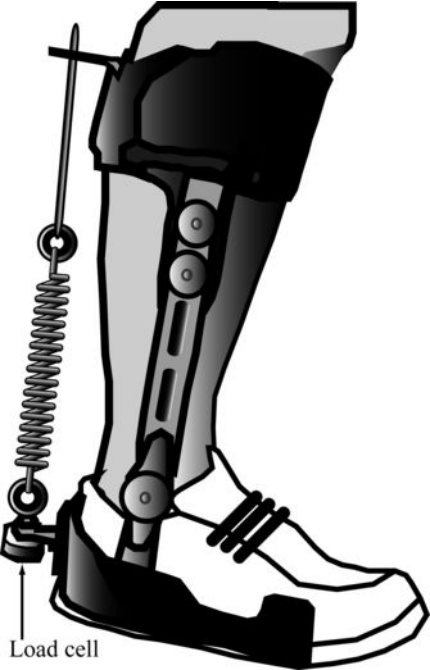
492 Figure 4. Group mean ankle (A), knee (B) and hip (C) powers plotted over an average hop  
493 (normalised over 101 points). The additional data series of smaller magnitude on the S ankle plot are  
494 the exoskeleton powers for the corresponding frequencies and the hatched area highlights the  
495 contribution from biological tissues to the total ankle power for 2.2 Hz. The left column is the no  
496 exoskeleton (NE) condition, middle is the spring (S) condition and right is for the no spring (NS)  
497 condition. Vertical lines represent the range of normalised take-off times across frequencies.

498 Figure 5. Pie charts showing the percentage of total average positive power contributed by the  
499 biological tissues at the ankle (white), knee (thick hatching), hip (black) and by the exoskeleton  
500 (narrow hatching) for the spring-loaded (A), no spring (B) and no exoskeleton (C) conditions. The  
501 total area of the pies represents the total average positive power relative to all other pies.

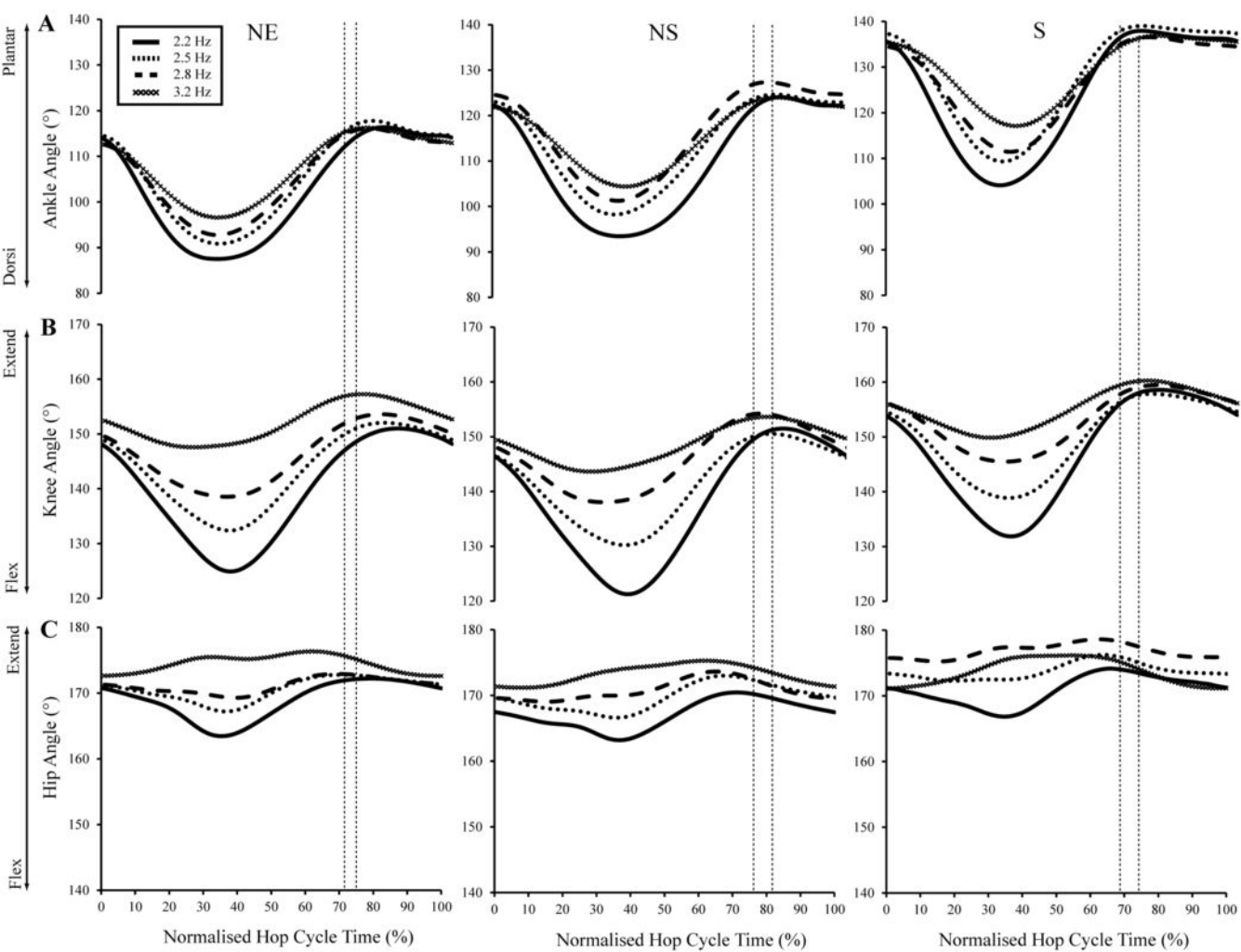
502 Figure 6. Metabolic power (A), biological contribution to  $\bar{P}_{tot}^+$  (B) and Biological  $\bar{M}_{ank}^+$  (C) across  
503 hopping frequency for S (filled diamonds), NS (filled circles) and NE (open triangles). \* denotes  
504 significant difference from S at that frequency and † denotes a significant main effect of frequency

505 Figure 7. Group mean normalised and smoothed time histories of EMG activity for MG (A), LG (B), SO  
506 (C) and TA (D) at each frequency for NE, NS and S. Vertical lines indicate the range of normalised  
507 take-off times across frequencies.

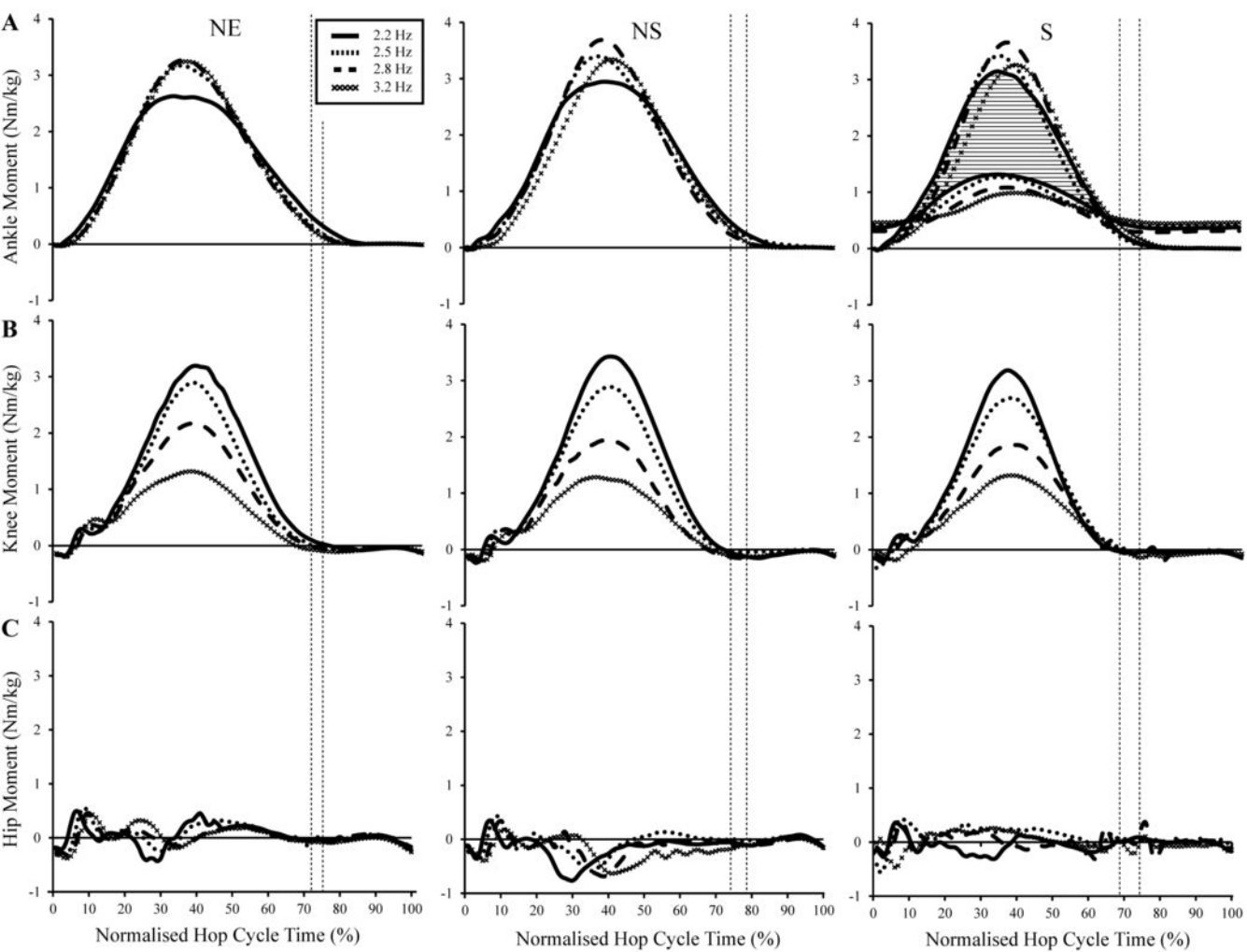
508 Figure 8. Group mean  $\pm$  sem RMS EMG values for MG, LG, SO and TA during the ground contact (A)  
509 and aerial (B) phases of hops. \* denotes statistical significantly different from the S condition and †  
510 denotes significant effect of frequency

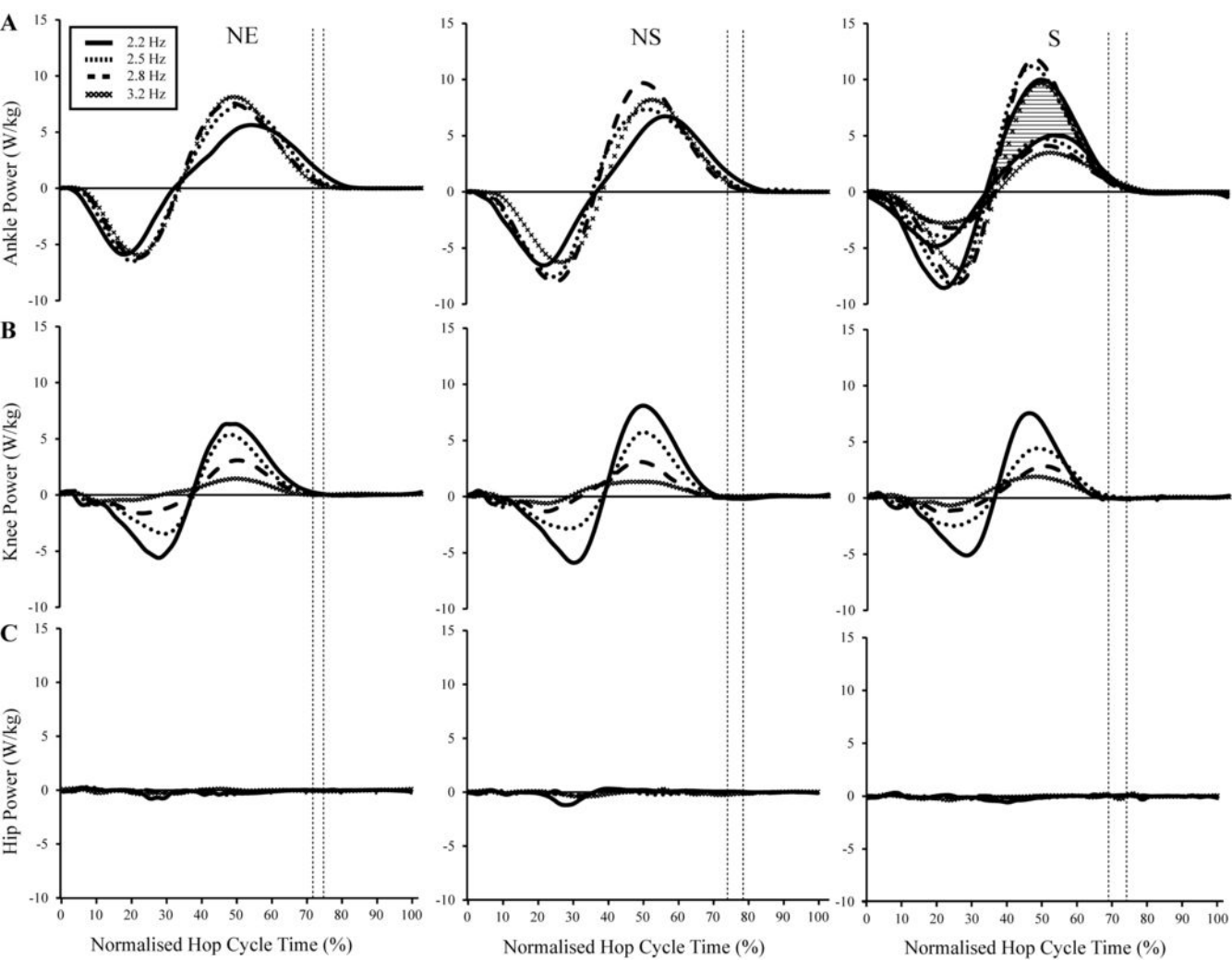


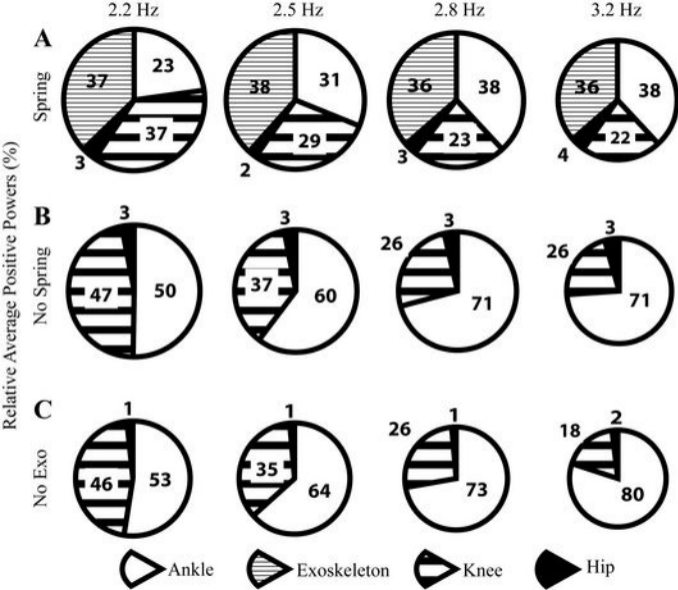
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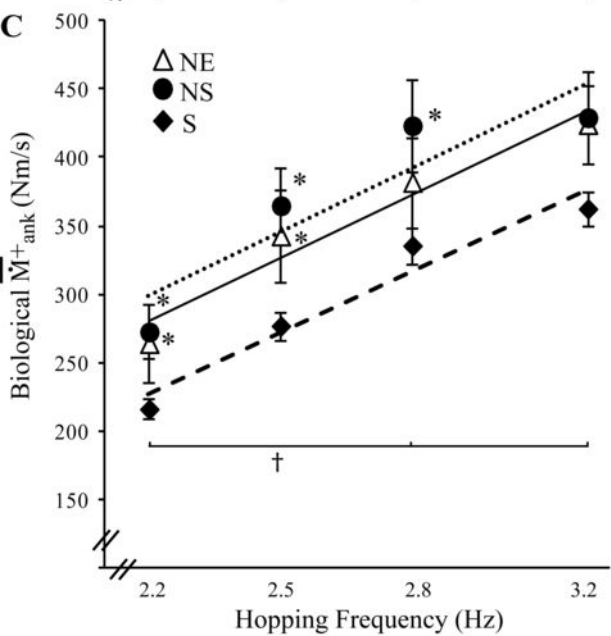
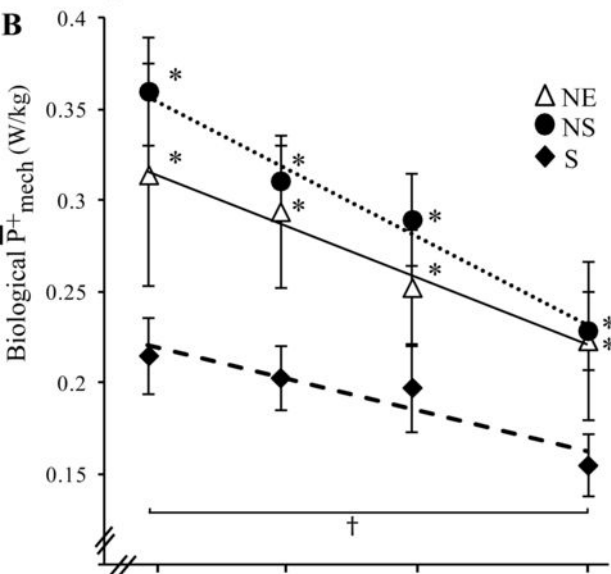
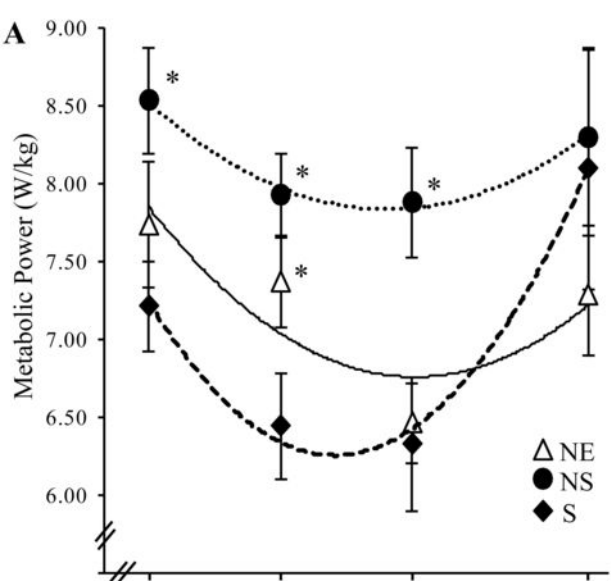


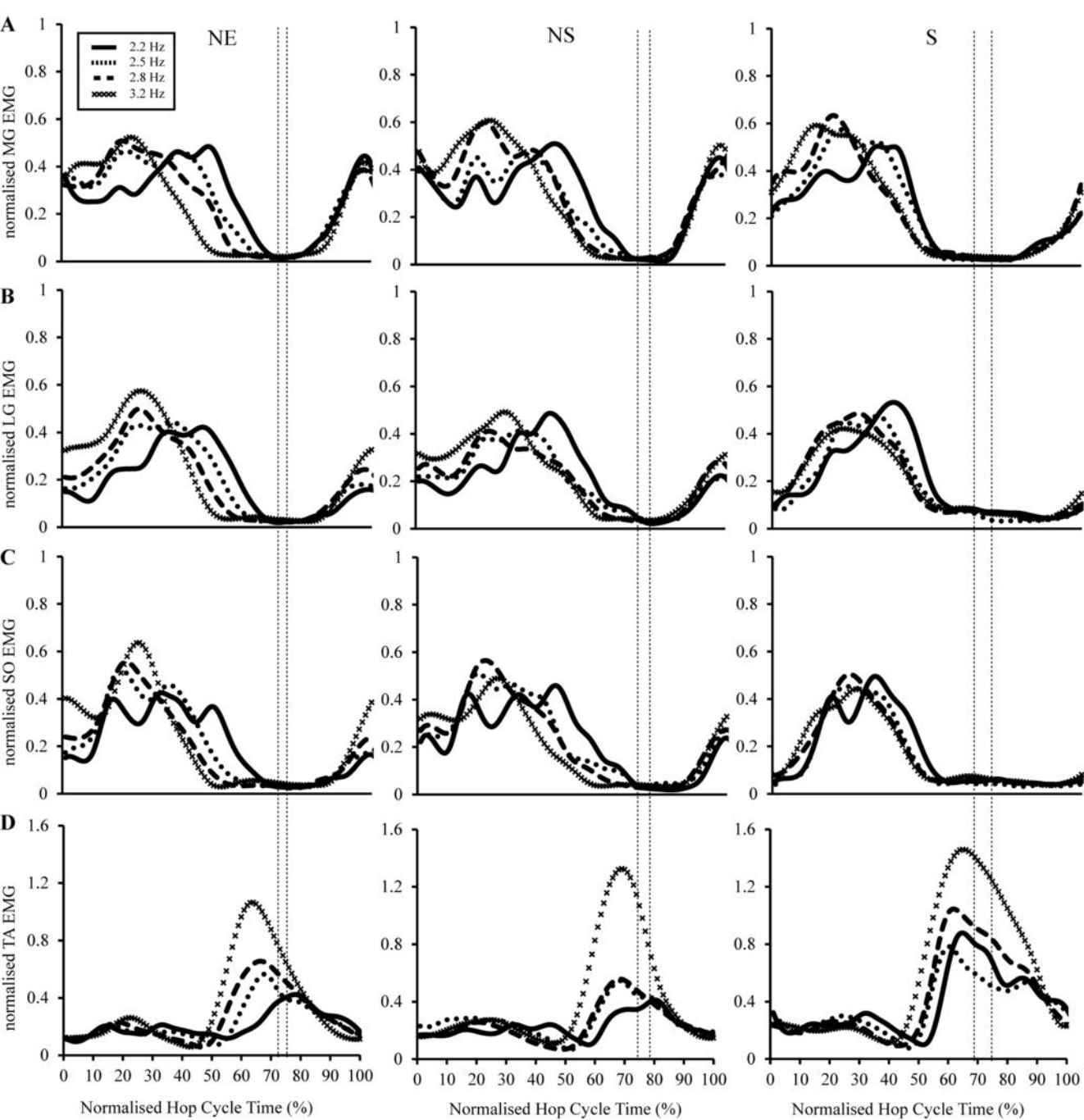












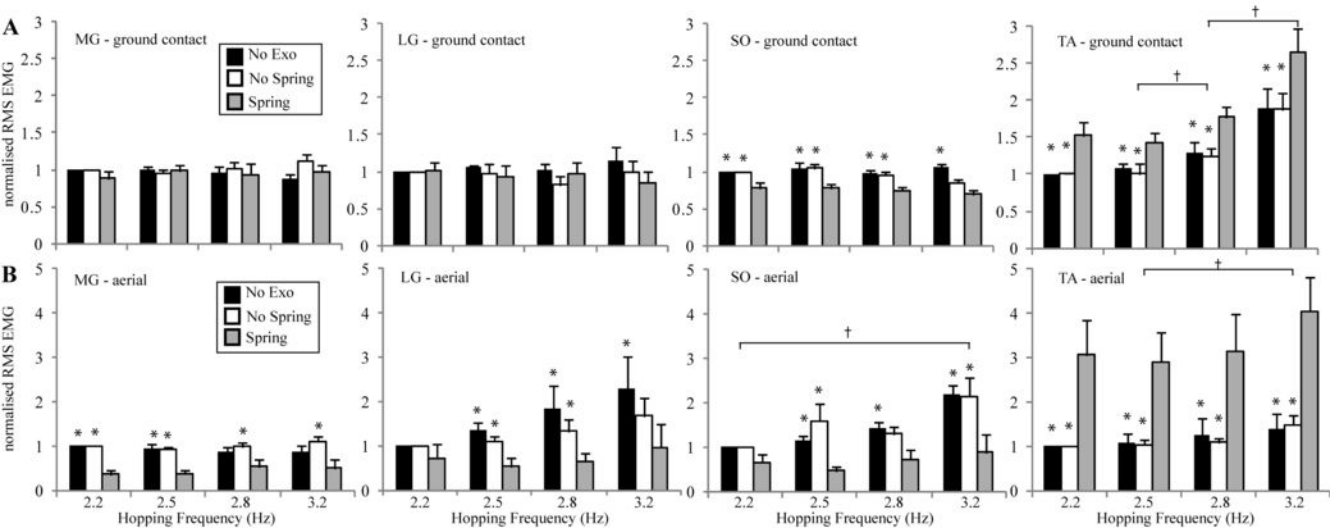


Table 1. Group mean  $\pm$  sd hopping metrics at each frequency for each spring condition. ANOVA P values are for main effects of the spring condition (S) and frequency ( $\omega$ ) and the interaction effect between the two ( $S*\omega$ ).

	$\omega$ (Hz)	Hop Height (mm)	Ground Contact Time (ms)	Actual Frequency (Hz)	Duty Factor	dHop (mm)
No Exo	2.2	22 $\pm$ 8	337 $\pm$ 41	2.2 $\pm$ 0.1	0.75 $\pm$ 0.09	18 $\pm$ 5
	2.5	20 $\pm$ 8	285 $\pm$ 35	2.5 $\pm$ 0.1	0.72 $\pm$ 0.08	20 $\pm$ 4
	2.8	13 $\pm$ 5	256 $\pm$ 29	2.8 $\pm$ 0.0	0.72 $\pm$ 0.07	17 $\pm$ 5
	3.2	9 $\pm$ 4	225 $\pm$ 19	3.2 $\pm$ 0.1	0.73 $\pm$ 0.06	17 $\pm$ 5
No Spring	2.2	13 $\pm$ 6	352 $\pm$ 28	2.2 $\pm$ 0.1	0.78 $\pm$ 0.06	20 $\pm$ 5
	2.5	13 $\pm$ 6	300 $\pm$ 23	2.5 $\pm$ 0.0	0.75 $\pm$ 0.06	22 $\pm$ 10
	2.8	11 $\pm$ 4	264 $\pm$ 19	2.8 $\pm$ 0.0	0.74 $\pm$ 0.05	20 $\pm$ 7
	3.2	5 $\pm$ 2	264 $\pm$ 12	3.2 $\pm$ 0.1	0.79 $\pm$ 0.04	25 $\pm$ 7
Spring	2.2	17 $\pm$ 14	312 $\pm$ 35	2.3 $\pm$ 0.2	0.70 $\pm$ 0.05	25 $\pm$ 12
	2.5	15 $\pm$ 9	279 $\pm$ 27	2.5 $\pm$ 0.0	0.69 $\pm$ 0.07	22 $\pm$ 5
	2.8	13 $\pm$ 7	259 $\pm$ 23	2.8 $\pm$ 0.2	0.72 $\pm$ 0.04	28 $\pm$ 9
	3.2	9 $\pm$ 4	236 $\pm$ 22	3.1 $\pm$ 0.1	0.74 $\pm$ 0.05	29 $\pm$ 8
ANOVA P (S)		0.19	0.08	0.100	0.171	0.038*
ANOVA P ( $\omega$ )		<0.000*	<0.000*	<0.000*	0.014	0.424
ANOVA P (S* $\omega$ )		0.259	0.003*	0.263	0.071	0.432

\*Indicates statistically significant effect.