Effects of whole-body vibrations on neuromuscular fatigue: A study with sets of different durations

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Background. Whole body vibrations have been used as a tool to study neuromuscular integration or as an exercise modality to increase neuromuscular performance and health. There is increasing evidence that longer WBV exposure (up to 10 minutes) induce a decline of neuromuscular parameters. However, the magnitude and origin of WBV induced fatigue are poorly understood.

Purpose. The study aimed to investigate the magnitude and origin of neuromuscular fatigue induced by half-squat long-exposure whole-body vibration intervention (WBV) with sets of different duration and compare it to non-vibration (SHAM) conditions.

Methods. Ten young, recreationally trained adults participated in six fatiguing trials, each consisting of maintaining a squatting position for several sets of the duration of 30, 60 or 180 seconds. The static squatting was superimposed with vibrations (WBV_{30}, WBV_{60}, WBV_{180}) or without vibrations ($SHAM_{30}$, $SHAM_{60}$, $SHAM_{180}$) for a total exercise exposure of 9-minutes in each trial. Maximum voluntary contraction (MVC), level of voluntary activation (%VA), low- (T_{20}) and high-frequency (T_{100}) doublets, low-to-high-frequency fatigue ratio ($T_{20/100}$) and single twitch peak torque (TW_{PT}) were assessed before, immediately after, then 15 and 30 minutes after each fatiguing protocol.

Result. Inferential statistics using RM ANOVA and post hoc tests revealed statistically significant declines from baseline values in MVC, T_{20}, T_{100}, $T_{20/100}$ and TW_{PT} in all trials, but not in %VA. No significant differences were found between WBV and SHAM conditions. Magnitude based inference revealed a *likely small* to *medium* fatiguing effects in favour of WBV_{30} for MVC. *Possibly small* to *likely moderate* fatiguing effect in favour of WBV_{180} were observed for TW_{PT}, T_{20} and $T_{20/100}$.

Conclusion. Our findings suggest that the origin of fatigue induced by WBV is not significantly different compared to control conditions without vibrations. The lack of significant differences in %VA and the significant decline in other assessed parameters suggest that fatiguing protocols used in this study induced peripheral fatigue of a similar magnitude in all trials. However, trials with longer sets duration (WBV_{180}) were likely to induce a possibly larger magnitude of fatigue compared to SHAM condition.

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Introduction

Whole body vibration (WBV) transfers sinusoidal oscillations into the human body, which inspired the use of this physical modality both as a tool to study the sensorimotor integration of the neuromuscular system and as an intervention stimulus with beneficial effects on human health and performance (Rittweger, 2010). Early studies have suggested that short single sessions of WBV of 3 to 5-minutes duration in a squat position immediately increase neuromuscular performance (Bosco et al., 2000; Cardinale & Bosco, 2003), maximal voluntary contraction (MVC), jump performance and myoelectric activity (Alam, Khan & Farooq, 2018). The acute increase in neuromuscular performance after vibration is referred to as 'post-activation potentiation' (PAP) for short-lasting enhancements (less than 1 minute) and as 'post-activation performance enhancement' (PAPE) for more extended performance enhancement periods lasting up to several hours (Blazevich & Babault, 2019). Both phenomena are related to vibration-induced changes in the neuronal control of the affected skeletal muscles that encompass a facilitated central drive (Mileva, Bowtell & Kossev, 2009; Krause et al., 2017) concomitant with modified reflectory activation at the spinal level (Rittweger, Beller & Felsenberg, 2000; Ritzmann et al., 2018) persistent over a period of 15 minutes after vibration exposure (Krause et al., 2016; Ritzmann et al., 2018).

In everyday practice, therapists and practitioners promote longer WBV exposures (10 minutes or longer), although the effects of such exercise modalities are mostly unknown (Torvinen et al., 2002; Zory et al., 2013). By increasing the WBV stimuli duration up to a cumulative total of 10 minutes, it has been suggested that WBV may acutely induce fatigue rather than potentiation (Torvinen et al., 2002; de Ruiter et al., 2003; Erskine et al., 2007; Rittweger, 2010; Zory et al., 2013). For example, Erskine et al. (2007) observed a gradual decrease of MVC after a 10×1-minute WBV intervention. However, no changes in MVC were observed in the control condition without vibrations. Even though various studies have reported a fatigue-induced drop in neuromuscular performance, there have been contradictory findings regarding the underlying mechanisms which favour either a central or peripheral origin. Several authors investigated the effect of WBV on central fatigue (Jordan et al., 2010; Maffiuletti et al., 2013; Zory et al., 2013) and were unable to find any difference in the level of voluntary activation (%VA) between interventions with and without vibrations. In contrast, the protocols of de Ruiter et al. (2003) resulted in a significant drop in %VA after a single session of WBV, and they postulated the involvement of central fatiguing mechanisms after WBV. To the best of our knowledge, the force-frequency fatigue-related mechanisms of WBV-induced peripheral fatigue have not been studied. By comparing the ratio of the electrically induced mechanical responses using low-frequency (below fusion frequency – 20 Hz) and high-frequency (above fusion frequency – 100 Hz) paired supramaximal electrical stimuli, peripheral fatigue can be subdivided into low- and high-frequency (Edwards, 2008; Millet et al., 2011). Analogous exercise-induced fatigue

studies have demonstrated that prolonging exercise stimuli can shift the peripheral fatiguing mechanism towards low-frequency fatigue (Millet & Lepers, 2004; Tomazin et al., 2012).

To better understand the intervention stimuli induced by WBV, it is crucial to establish which fatiguing mechanisms occur after a single session of WBV, and how different vibration parameters affect the magnitude and origin of neuromuscular fatigue. The scientific and practitioner choices for WBV intervention are motivated by achieving high superimposed effects throughout WBV to trigger physiological and neuromuscular adaptations and thus, WBV parameters are combined accordingly (Abercromby et al., 2007; Ritzmann, Gollhofer & Kramer, 2013). Electromyography studies suggest that side-alternating vibration exposure driven by high amplitude and frequency cause the highest activation intensities in distal and proximal leg musculature (Abercromby et al., 2007; Rittweger, 2010; Ritzmann, Gollhofer & Kramer, 2013). In addition to vibration-associated attributes, and in an analogy to strength training, the training load is mainly determined by intensity and volume (Baechle & Earle, 2008). Therefore, volume is subdivided into number of set and repetitions with defined set duration (Campbell et al., 2017). In a similar manner, vibration amplitude and frequency define the training intensity in WBV interventions. However, to the best of our knowledge, there is a lack of studies investigating how WBV intervention volume (set numbers and set duration) affects the occurrence of neuromuscular fatigue.

Therefore, the aim of the present study was to investigate the magnitude and origin of neuromuscular fatigue induced by long-exposure half-squat whole-body vibration intervention (WBV) with sets of different duration and compare it with non-vibration (SHAM) conditions. Thus, and with reference to, previous research involving long-exposure WBV induced fatigue (Erskine et al., 2007; Zory et al., 2013) we selected a long (cumulative exercise time of 9 minutes) static WBV fatiguing intervention divided into sets of different duration (30 s, 60 s or 180 s). In a series of MVC paradigms, we applied different peripheral nerve stimulation techniques, allowing us to distinguish the source of fatigue. We hypothesised that WBV exercise interventions would cause higher magnitudes of fatigue compared to non-vibration intervention (Erskine et al., 2007; Zory et al., 2013). We expected that fatigue magnitude would be dependent on the duration of exposure and would increase with set-duration. We hypothesised that predominantly peripheral, rather than central fatiguing mechanisms, would be causally involved (Jordan et al., 2010; Maffiuletti et al., 2013; Zory et al., 2013).

Materials & Methods

Study design

In a cross-over repeated measures design, each subject performed three different fatiguing exercise interventions with WBV and three exercise interventions in a sham

condition without WBW (SHAM) to determinate the effect of WBV (*Fig.* 1A.). Each intervention comprised a cumulative exercise period with a duration of 9 minutes divided into different sets (either 18 x 30 s or 9 x 60 s or 3 x 180 s) with 120 s rest between sets (Fig 1A). The exercise interventions were performed on an activated vibration platform (WBV_{30}, WBV_{60}, WBV_{180}) and three on an inactive vibration platform ($SHAM_{30}$, $SHAM_{60}$, $SHAM_{180}$). Each intervention was executed on different visits with at least seven days rest in-between. The order was randomised. The subjects were not permitted to undertake explosive strength training or fatiguing workouts for 48 hours before each measuring day, in order to eliminate side-effects. The study design, materials and neuromuscular assessments are available for reference in protocols.io (dx.doi.org/10.17504/protocols.io.beadjaa6)

Neuromuscular assessment in the resting position was performed at t_0 (baseline) prior to exercise intervention. The assessment consisted maximum voluntary contraction (MVC) of the knee extensors, interpolated with a high frequency (T_{MVC}) twitch (10 ms interstimuli interval), followed 3 s later by a 100 Hz doublet (T_{100}), followed 3 s later by a 20 Hz (50 ms interstimuli interval) doublet (T_{20}), and 3 s later by a potentiated single twitch (TW). The assessment procedure was executed according to (Millet et al., 2011) and repeated at 1 minute (t_f), as well as at 15 (t_{f15}) and 30 minutes (t_{f30}) after the final 9-minute intervention. All neuromuscular assessments were performed on the right leg.

Subjects

Ten healthy subjects (6 men and 4 women; age: 21.1 $\pm$ 1.41 years, mass: 77.8 $\pm$ 11.73 kg, BMI: 22.9 $\pm$ 1.25) volunteered to participate in the study. All subjects were recreationally trained athletes, participating in moderate endurance and strength training activities 3 times per week. Exclusion criteria were acute injuries in the upper or lower extremities, locomotor dysfunctions, pregnancy, cardiovascular or neurological conditions. All subjects signed the written informed consent and the study was approved by the Ethics Committee of the Faculty of Sport of the University of Ljubljana 975/2017 and conducted according to the Declaration of Helsinki II.

The sample size was estimated by means of a power analysis based on experimental evidence obtained from Jordan et al. (Jordan et al., 2010)(f = 0.85; alpha = 0.05; power = 0.85).

Intervention

The interventions were performed on a side-alternating vibration platform (Galileo Fit, Novotec Medical, Germany) which was running at a frequency of 26 Hz (Rittweger, Mutschelknauss & Felsenberg, 2003; Cochrane et al., 2010) and off, respectively, for WBV and SHAM conditions. Subjects were instructed to maintain a half-squat position with their knees flexed at an angle of 60° (Ritzmann et al., 2010) for several sets with 2-minute rest between sets. Kinematics were controlled with a goniometer. The subjects

stood with their feet 40 cm apart at a point where the tilting platform reached peak-to-peak displacement amplitude of 5 mm (Ritzmann, Gollhofer & Kramer, 2013). At the beginning of each session, subjects underwent a 6-minute warm-up routine consisting of bench stepping (20 cm high) at a frequency of 0.5 Hz, swapping the leading leg at one minute intervals.

Testing protocols

During the neuromuscular assessment, the subjects remained seated in a custom-built isometric knee extension apparatus equipped with a force transducer (MES, Maribor, Slovenia) (Tomazin, Dolenec & Strojnik, 2008; García-Ramos et al., 2016). The force transducer was calibrated prior to testing sessions. Each subject was seated in an upright position, hip at 100° and trunk leaning against the backrest of the testing apparatus, fixed by straps over the pelvis and a horizontal pad over the distal third of the thigh. The knee joint axis was aligned with the mechanical axis of the dynamometer. The shin pad was placed just superior to the medial malleolus. The right knee joint was fixed at a 60° angle (0° = full extension) (*Fig.* 1C).

Femoral nerve electrical stimulation

The femoral nerve was stimulated by pressing a monopolar cathode (10-mm in diameter, Ag–AgCl, Type 0601000402, Controle Graphique Medical, Brie-Comte-Robert, France) into the femoral triangle of the iliac fossa (*Fig.* 1C). A larger (102mm x 52mm, Compex, SA, Ecublens, Switzerland) self-adhesive electrode placed over the gluteal fold served as the anode. Electrical impulses (single, square wave, 1-ms duration) elicited by a high voltage constant current electrical stimulator (DS7A; Digitimer, Hertfordshire, UK) were used to trigger the muscle response, which was detected as a change in torque of the knee extensors. The stimulation intensity to elicit maximum knee extensors isometric twitch was determined in each subject at the beginning of each trial and maintained for the entire trial. Starting from an intensity of 10 mA, the stimulation intensity was progressively increased by 10 mA until no further increase in torque was observed despite further increment in electrical current. The current at maximal twitch torque was additionally increased by a factor of 1.5 to obtain a supramaximal stimulus (Verges et al., 2009).

Single twitch

The torque change induced by a single supramaximal femoral nerve stimulus (Place et al., 2007) was analysed to obtain the peak torque value (TW_{PT}).

High- and low-frequency doublets

The torque change induced by the paired high-frequency (100 Hz, i.e. 10-ms interstimuli interval) and low-frequency (20 Hz, i.e. 50-ms interstimuli interval) supramaximal

electrical stimuli (Place et al., 2007; Verges et al., 2009) was analysed to obtain the following parameters: peak torque from 100 Hz doublet (T_{100}), peak torque from 20 Hz doublet (T_{20}). In addition, the low- to high-frequency ratio ($T_{20/100}$) was calculated using the following formula:

$$T_{20/100} = \frac{T_{20}}{T_{100}} * 100$$

This ratio was used as a surrogate of low- to high-frequency tetanic stimulation (Verges et al., 2009).

Maximal voluntary contraction with double twitch interpolated techniques

Subjects were asked to perform a 5 s maximal isometric voluntary knee extension (Verges et al., 2009). The signal was smoothed using a 0.5 s window moving average filter and peak torque (MVC) was retained for analysis. The double twitch interpolated technique (Allen, Gandevia & McKenzie, 1995) was performed by superimposing a 100 Hz doublet on the isometric plateau (T_{MVC}). A second analogous stimulation (T_{100}) on the relaxed muscle followed after 3 seconds (*Fig.* 1B). The ratio of the amplitude of the T_{MVC} over T_{100} was then calculated to obtain the level of voluntary activation (%VA):

$$\%VA = \left(1 - \frac{T_{MVC} - MVC}{T_{100}}\right) * 100$$

Statistics

A two-way factorial ANOVA (Type III) was conducted in R(3.5.1) with the afex package (Singmann et al., 2018) to compare the main effects of time (t_0, t_f, t_{f15}, t_{f30}) and trial (WBV_{30}, WBV_{60}, WBV_{180}, $SHAM_{30}$, $SHAM_{60}$, $SHAM_{180}$) and the interaction effect of time $\times$ trial. Generalised eta squared (η_G^2) effect sizes were calculated for the ANOVA main and interaction effects. In the case of statistically significant interactions, post hoc comparisons with Sidak corrections were applied using the emmeans package (Lenth et al., 2018) in order to compare WBV and SHAM condition. Tukey-corrected pairwise post hoc tests were used to calculate differences to baseline within trials.

In addition to inference statistics, standardised changes in the mean of each measure were used to assess the magnitudes of effect (ES) between WBV and SHAM conditions of the same set duration (e.g. $SHAM_{30}$-WBV_{30}, $SHAM_{60}$-WBV_{60}, $SHAM_{180}$-WBV_{180}) and were then calculated using Cohen d. The magnitude of ES was interpreted as follows: trivial = <0.20; small = 0.2–0.59; moderate = 0.60–1.19; large = 1.20–1.99; and very large = >2.0 based on recommendations by Hopkins (Hopkins et al., 2009). Additionally, magnitude-based inferences of observed ES were determined and interpreted qualitatively as: almost certainly not = <0.5%; very unlikely = 0.5%-5%; unlikely = 5%-25%; possible = 25%-75%; likely = 75%-95%; very likely = 95%-99.5%; and almost certain = >99.5% (Hopkins et al., 2009). Statistical significance was set at the level of $p < 0.05$. ES results should be interpreted with caution, since negative

values imply a larger fatiguing effect of WBV compared to SHAM condition and positive values imply a larger fatiguing effect for SHAM condition compared to WBV.

Results

Descriptive statistics for MVC and %VA are displayed in Table 1; descriptive statistics for T_{20}, T_{100} and $T_{20/100}$, TW_{PT} are listed in Table 2.

Maximum voluntary contraction

There was a statistically significant time effect (F (3, 27) = 24.40, p < 0.001, η_G^2 = 0.02), but no significant trial effect (F (5, 45) = 2.13, p = 0.08, η_G^2 = 0.01) nor trial x time interaction effect (F (15, 135) = 0.60, p = 0.87, η_G^2 = 0.002) for MVC. Within-trial post hoc tests showed differences between baseline and post-assessments (*Fig.* 2a). Trials with sets of 30 s duration induced *likely small* to *likely moderate* fatiguing effects lasting up to 30 minutes after the exercise intervention in favour of WBV (*Fig.* 3). *All other trials comparing SHAM and WBV resulted in unclear differences.*

Level of voluntary activation (%VA)

There was a statistically significant time (F (3, 27) = 3.67, p = 0.024, η_G^2 = 0.02) and trial (F (5, 45) = 2.52, p = 0.042, η_G^2 = 0.08) effect, but no trial x time interaction (F (15, 135) = 1,21, p = 0.26, η_G^2 = 0.03) for %VA. Post hoc tests did not reveal significant differences between baseline and post-assessments (*Fig.* 2b). Trials with sets of 30 s duration induced possibly *small* to *likely moderate* fatiguing effects 15 and 30 minutes after exercise intervention in favour of SHAM. Trials with sets of 60 s duration induced *likely small* fatiguing effects 15 minutes after exercise intervention in favour of SHAM. Trials with sets of 180 s duration resulted in *likely small* fatiguing effects 15 minutes after exercise intervention in favour of WBV (*Fig.* 3).

Peripheral fatigue

There was a significant time effect (F (3, 27) = 64.43, p < 0.001, η_G^2 = 0.25) for T_{20}. Trial effects (F (5, 45) = 1.91, p = 0.11, η_G^2 = 0.03) and trial x time interaction effects (F (15, 135) = 0.90, p = 0.56, η_G^2 = 0.007) remained statistically insignificant. Post hoc tests revealed significant differences between baseline and post-assessments for each of the trials (Fig. 4a**,** Table 3). Trials with sets of 30 s duration induced *possibly small* fatiguing effects in favour of WBV 15 and 30 minutes after exercise intervention. Trials with sets of 180 s duration induced *likely small* fatiguing effects in favour of WBV lasting at least 30 minutes after exercise intervention (*Fig.* 5). Trials with sets of 60 s duration resulted in *unclear differences*.

There was a significant *time* effect (*F* (3, 27) = 60.33, p < 0.001, η_G^2 = 0.15) for T_{100}. *Trial* effect (*F* (5, 45) = 2.15, p = 0.07, η_G^2 = 0.03) and *trial x time interaction* effect (*F* (15, 135) = 0.43, p = 0.97, η_G^2 = 0.002) remained statistically insignificant. Post hoc tests revealed significant differences between baseline and post-assessments for each of the trials (*Fig.* 4b, Table 3). All trials *resulted in unclear differences*.
There was a significant time effect (F (3, 27) = 46.33, p < 0.001, η_G^2 = 0.17) for T20/100. Trial effect (F (5, 45) = 1.06, p = 0.40, η_G^2 = 0.02) and trial x time interaction effect (F (15, 135) = 0.97, p = 0.49, η_G^2 = 0.02) remained statistically insignificant. Post hoc tests revealed significant differences between baseline and post-assessments for each of the trials (*Fig.* 4c, Table 3). Trials with sets of 180 s duration resulted in *likely moderate* fatiguing effect in favour of WBV 15 minutes after exercise intervention (*Fig.* 5).

Single twitch

There was a significant time effect (F (3, 27) = 48.80, p < 0.001, η_G^2 = 0.23). Trial effects (F (5, 45) = 0.86, p = 0.52, η_G^2 = 0.006) and trial x time interaction effect (F (15, 135) = 1.05, p = 0.41, η_G^2 = 0.006) remained statistically insignificant for TWPT. Post hoc tests revealed significant differences between baseline and post-assessments for each of the trials (*Fig.* 2c, Table 3). Trials with sets of 180 s duration induced *likely moderate* fatiguing effect in favour of WBV. *Likely small* to *possibly small* fatiguing effects in favour of WBV seemed to last at least 30 minutes after exercise intervention (*Fig.* 3). All other trials comparing SHAM and WBV resulted in unclear differences.

Discussion

The current study aimed to investigate the magnitude and origin of neuromuscular fatigue induced by long-exposure half-squat whole-body vibration intervention (WBV) with sets of different duration and compare it with non-vibration (SHAM) conditions. Our findings revealed a small superimposed effect of WBV compared to control conditions without vibrations.

Maximal voluntary contraction

Knee extensors MVC torque dropped by 7 to 12% after each fatiguing protocol, which is in line with other WBV induced fatigue studies, where MVC torque decreased by approximately 8% (de Ruiter et al., 2003; Erskine et al., 2007; Colson et al., 2009; Zory et al., 2013). Only Maffiuletti et al. (2013) reported a more substantial decline in MVC (-23%), which is likely associated with the application of additional loads coupled with shorter inter-set rest periods compared to other studies and to our specific experimental setting. Although interaction effects remained insignificant, magnitude-based inference showed that with *likely small* to *moderate effect,* WBV_{30} might be more fatiguing compared to $SHAM_{30}$ (*Fig.* 3).
This finding is in contrast with our hypothesis that longer set duration exercises superimposed with vibration (WBV_{180}) would produce greater fatigue compared to $SHAM_{180}$ condition. However, it has been previously suggested that potentiated electrically elicited supramaximal doublets represent a more suitable indicator of peripheral fatigue and contractile impairments compared to MVC torque (Place et al., 2007).

Central fatigue

The level of voluntary activation (%VA) of the knee extensors was not significantly depressed by any intervention utilised in this study, which suggest that mechanisms located in the central nervous system (CNS) were not significantly involved in the decline of MVC. These findings are in line with Colson et al. (2009) and Jordan et al. (2010) but in contrast to de Ruiter et al. (2003) who reported a vibration-induced decline in knee extensors voluntary activation. Being hypothesis-driven, our findings indicate that there are no evident superimposed effects of WBV on central fatiguing mechanisms compared to control conditions without WBV. This should be taken into consideration when designing exercise programs or research studies which intend to induce central fatigue. As such, WBV superimposed exercises are unlikely to be more effective than maintaining a static squat alone.

Peripheral fatigue

(We believe) This is the first study where electrically elicited supramaximal low- and high-frequency doublets were used to assess the origin and magnitude of peripheral fatigue after WBV exposure. For all protocols, T_{20} was more affected than T_{100} leading to a decreased $T_{20/100}$ ratio (Fig. 4**c**). These declines suggest the occurrence of low-frequency fatigue in all trials. It is noteworthy that the $T_{20/100}$ ratio for SHAM interventions returned to baseline values 15 minutes after the intervention, while WBV interventions remained significantly depressed up to 30 minutes after the intervention. This becomes particularly evident in trials using longer set durations (180 s) where the superimposed vibrations (WBV_{180}) induced slightly a larger magnitude of fatigue with *likely moderate effect* (*Fig.* 5) compared to the $SHAM_{180}$ control condition. This suggests that LFF is stronger and more long-lasting when a WBV exercise is executed with an emphasis on exposing to longer sets of vibration. The observation favouring LFF as an underlying mechanism can additionally be supported by the findings obtained from single twitch data. Similar to T_{20} and T_{100}, TW_{PT} progressively decreased as the intervention continued; despite the absence of statistically significant differences between trials, longer set duration (WBV_{180}) induced larger fatigue with *possibly small* to *likely moderate effects* compared to $SHAM_{180}$ (*Fig.* 3).

Underlying mechanisms

The results of this study suggest that there is low evidence of superimposed WBV effect compared to control conditions without WBV. This is particularly true for MVC and the level of voluntary activation. Even though some authors postulated WBV-induced modulation in the neuronal control is manifested as a facilitated central drive (Mileva, Bowtell & Kossev, 2009; Krause et al., 2016) this does not seem to translate into central fatigue. Furthermore, the decline in low- and high-frequency doublets, as well as single twitch torque, suggest that the predominant mechanism leading to the decrease in force production has to be addressed by an impairment in Ca^{2+} handling. This is followed by a gradual recovery of the Ca^{2+} depletion within the 15-30 min following intervention. Underlying cellular fatiguing mechanisms may refer to the three aspects (Westerblad et al., 2000; Allen & Westerblad, 2001; Williams & Ratel, 2009): a) since elicited peak torques progressively dropped at low- and high-frequencies of stimulation during the intervention protocol, there could be direct inhibition of inorganic phosphates (P_i) on Ca^{2+}, thereby producing an impairment in the cross-bridge force generation (Millar & Homsher, 1990); however, this mechanism alone cannot cause low-frequency fatigue (Allen, Lannergren & Westerblad, 1995); b) as a major drop in T_{20} caused a negative $T_{20/100}$ ratio it could indicate that the limiting mechanisms have to be addressed by Ca^{2+}-P_i precipitation in the sarcoplasmic reticulum, thus, leading to a decrease in free Ca^{2+} available for release (Allen & Westerblad, 2001). In addition, c) reduced myofibrillar Ca^{2+} sensitivity can also affect force production (Bruton et al., 2008). Both mechanisms (b and c) have little impact on force production at high frequencies but a large effect on

low frequencies (Westerblad et al., 2000). Therefore, magnitude-based inference statistics indicate that, by prolonging the set duration up to 180 seconds, vibration-induced superimposed spinal neuronal drive throughout reflectory muscle activation induced by WBV (Ritzmann et al., 2010) might have imposed larger peripheral low-frequency fatigue compared to maintaining a half-squat position alone.

Limitations

The study might have some limitations. An important limitation of this study (similar to the majority of other vibration studies) is the lack of WBV load normalisation, as this may have considerable side-effects on the results, as was demonstrated by Di Giminiani et al. (2009). Another limiting aspect deals with different work/rest ratios between long sets (180 s work – 120 s rest) compared to other shorter set durations (30 s – 120 s and 60 s – 120 s).

Conclusions

The outcomes of this study suggest the origin of fatigue induced by half-squat with superimposed vibrations is no different from the control conditions without vibrations. However, longer set durations (WBV_{180}) induced a slightly larger magnitude of neuromuscular fatigue compared to control conditions without vibrations. Due to a lack of significant modulation of voluntary activation, it can be assumed that the superimposed vibrations predominantly affected peripheral mechanisms rather than central ones. The primary vibration-induced peripheral fatiguing mechanism seems to find its origin in low-frequency fatigue which most probably involves Ca^{2+} handling. Based on the outcomes of this investigation, we suggest practitioners and researchers, aiming to induce peripheral fatigue using vibration superimposed exercises, consider using at least 3 sets of 180 seconds of WBV exposure in their fatiguing protocol planning.

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533

Figure 1

Experimental design and settings.

(A) Experimental design comprising the fatiguing protocols for all six visits and the timeline of each visit. Neuromuscular function was assessed before (t_0), immediately after (t_f), 15 (t_{f15}) and 30 (t_{f30}) minutes after vibration intervention. An expanded view of exercise exposure representing the WBV_{60} protocol (nine sets of 60 s of vibration exercise with 120s rest between sets) is presented in detail. (B) Example of a torque signal from the neuromuscular testing procedure. An expanded view of an interpolated twitch is presented in the dotted box. The neuromuscular testing procedure comprised MVC of the quadriceps muscle combined with different electrical stimulation methods to assess the level of voluntary activation - %VA (via the interpolated double twitch technique), quadriceps twitch torques in response to paired electrical stimuli at 100 Hz (T_{100}) and at 20 Hz (T_{20}), as well as single twitch (TW). (C) Schematic of the position of the subject during the neuromuscular assessment. An expanded view of the femoral nerve stimulation point in the popliteal fossa is presented within the box.

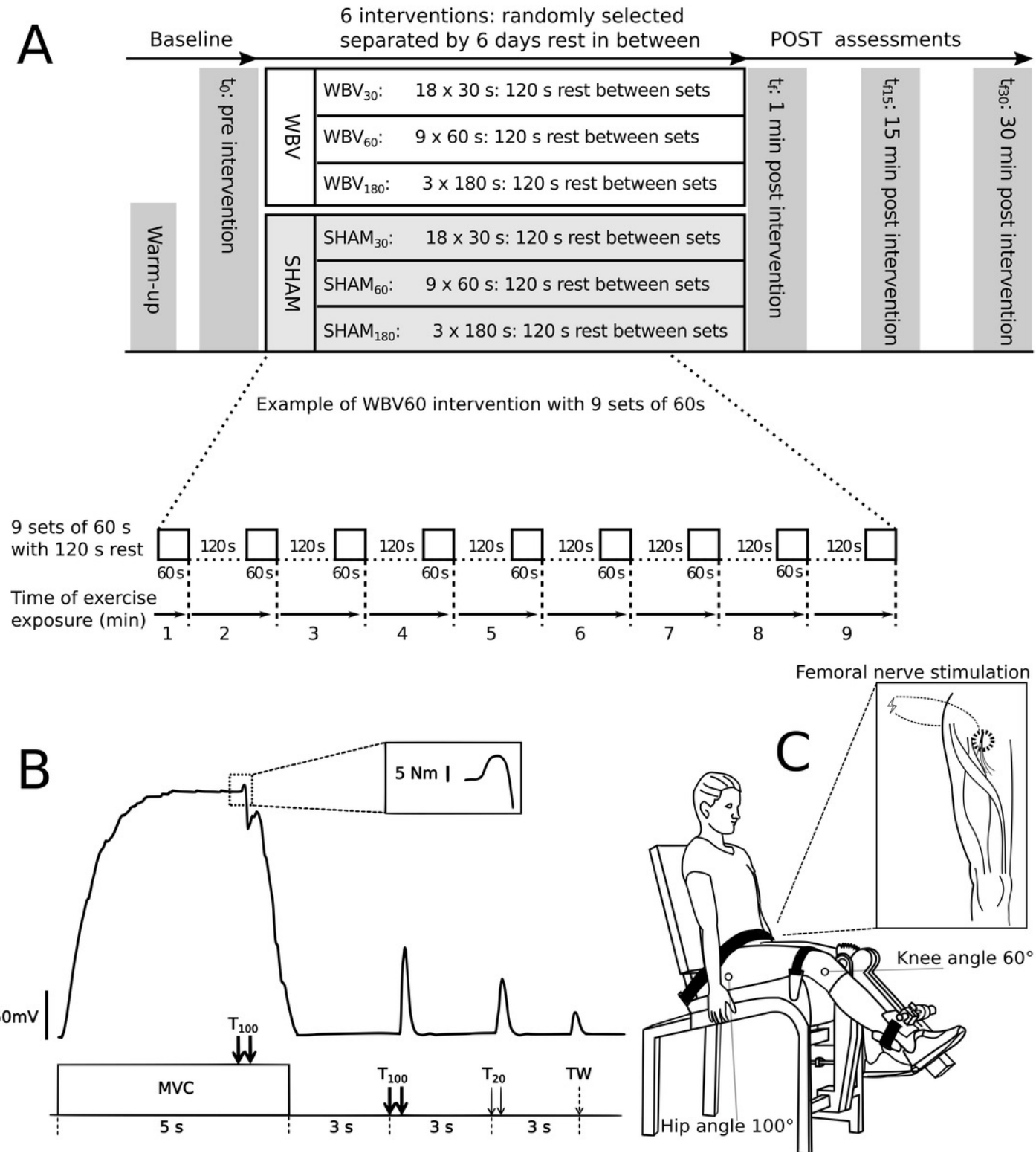

A
Baseline
6 interventions: randomly selected
separated by 6 days rest in between
POST assessments
Warm-up
t0: pre intervention
WBV
WBV30: 18 x 30 s: 120 s rest between sets
WBV60: 9 x 60 s: 120 s rest between sets
WBV180: 3 x 180 s: 120 s rest between sets
SHAM
SHAM30: 18 x 30 s: 120 s rest between sets
SHAM60: 9 x 60 s: 120 s rest between sets
SHAM180: 3 x 180 s: 120 s rest between sets
tf: 1 min post intervention
tf15: 15 min post intervention
tf30: 30 min post intervention
Example of WBV60 intervention with 9 sets of 60s
9 sets of 60 s
with 120 s rest
60s
120s
Time of exercise
exposure (min)
1
2
3
4
5
6
7
8
9
B
5 Nm
50mV
T100
MVC
5 s
3 s
T100
3 s
T20
3 s
TW
C
Femoral nerve stimulation
Knee angle 60°
Hip angle 100°

Figure 2

Relative changes from baseline.

(A) Maximum voluntary contraction (MVC), (B) level of voluntary activation (%VA) and (C) single twitch peak torque (TW_{PT}) for WBV (connected black triangles) and SHAM (connected white circles) for trials with different set durations (30 s, 60 s and 180 s). Values are expressed as mean and standard errors. Black triangles represent statistically significant WBV differences from baseline (▲▲▲ $p < 0.001$; ▲▲ $p < 0.01$; ▲ $p < 0.05$). White circles represent statistically significant SHAM differences from baseline (○○○ $p < 0.001$; ○○ $p < 0.01$; ○ $p < 0.05$).

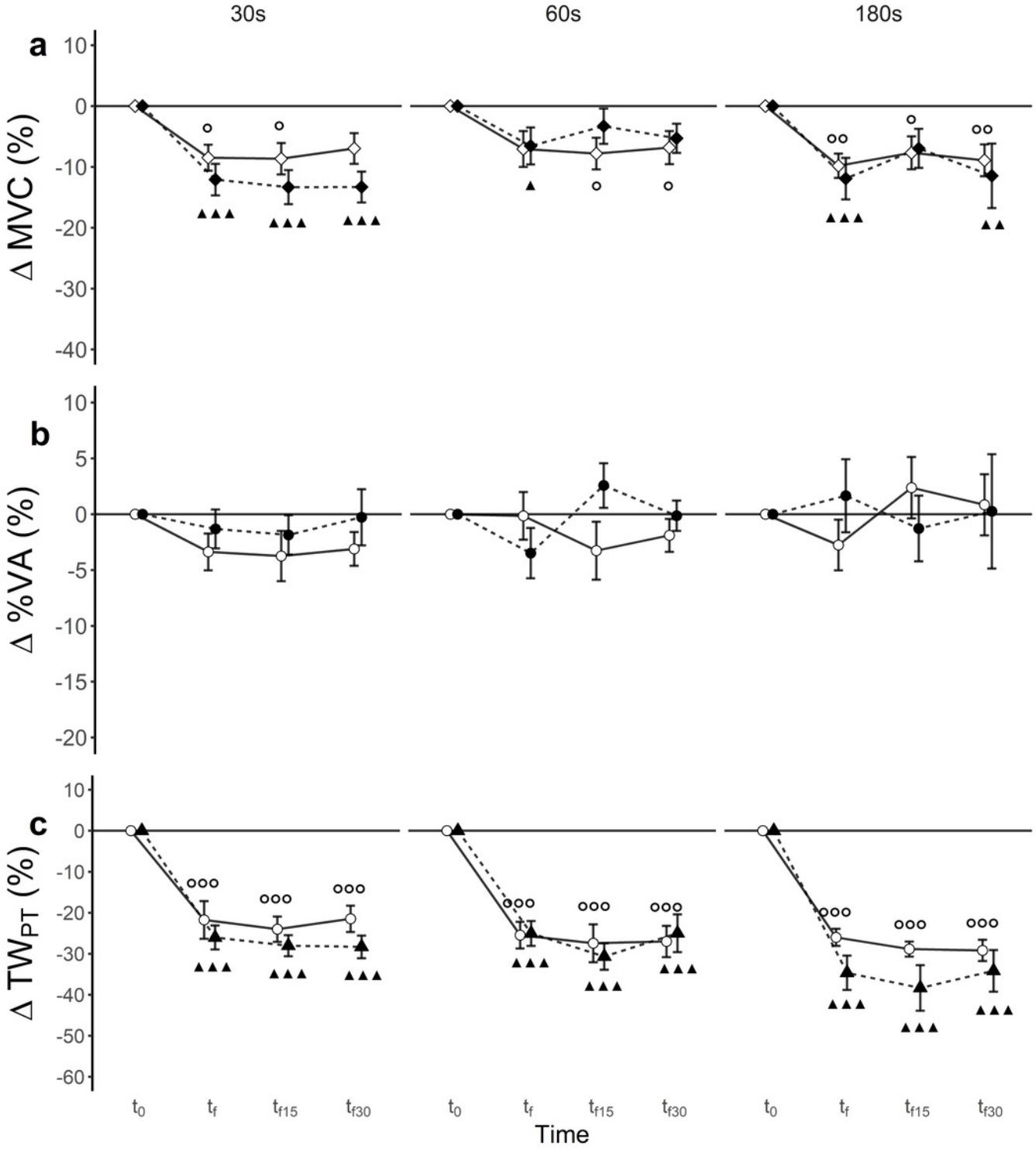
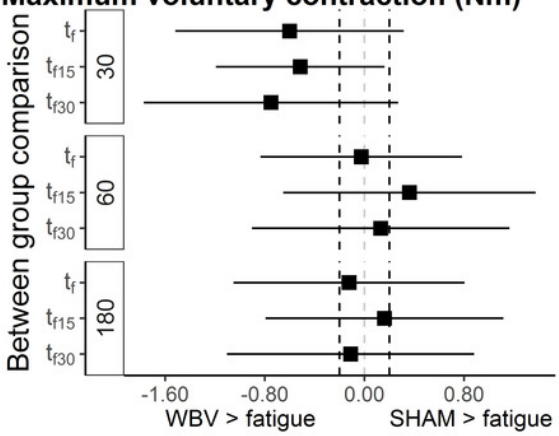

30s
60s
180s
a
Δ MVC (%)
10
0
-10
-20
-30
-40
b
Δ %VA (%)
10
5
0
-5
-10
-15
-20
c
Δ TWPT (%)
10
0
-10
-20
-30
-40
-50
-60
t0
tf
tf15
tf30
Time

Figure 3

WBV and SHAM between-group comparisons for MVC, %VA and TW_{PT}.

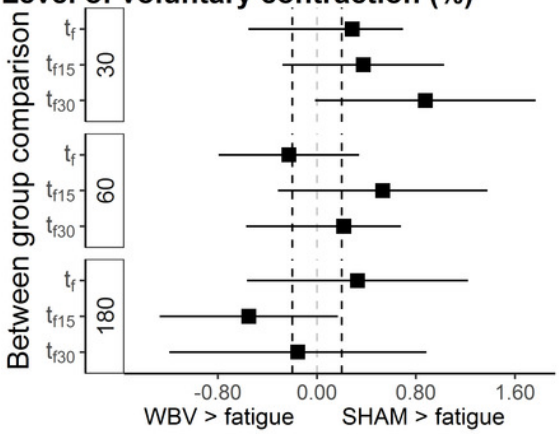
Raw differences (in native units) are presented as mean and 95% confidence intervals. Cohen *d* effect sizes and relative 95% confidence intervals are presented graphically and numerically. The magnitude base inference scores are presented. Negative effect size values represent higher fatigue in favour of the WBV condition, while positive effect size values represent higher fatigue in favour of the SHAM condition.

Maximum voluntary contraction (Nm)



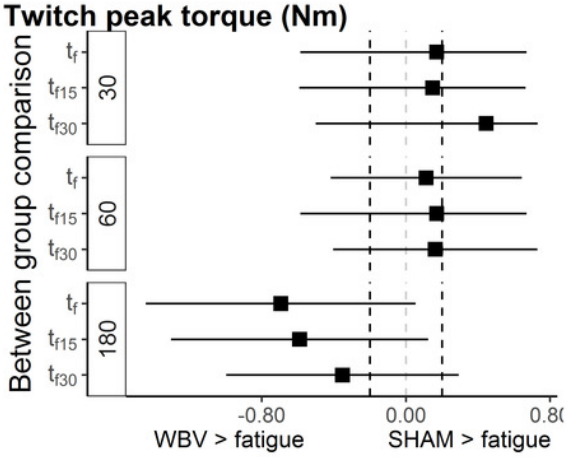
SHAM - WBV raw diff (95% CI)	Cohen *d* [95% CI]	MBI
-11.91 (-30.09, 6.27)	-0.60 [-1.52, 0.32]	*Likely Moderate effect: WBV.*
-9.29 (-21.51, 2.93)	-0.52 [-1.19, 0.16]	*Likely Small effect: WBV.*
-12.33 (-29.11, 4.45)	-0.75 [-1.77, 0.27]	*Likely Moderate effect: WBV.*
-0.61 (-20.03, 18.81)	-0.03 [-0.83, 0.78]	*Unclear Difference.*
7.89 (-14.25, 30.03)	0.36 [-0.65, 1.38]	*Unclear Difference.*
2.97 (-20.44, 26.39)	0.13 [-0.90, 1.17]	*Unclear Difference.*
-3.03 (-25.76, 19.71)	-0.12 [-1.05, 0.80]	*Unclear Difference.*
3.62 (-17.88, 25.12)	0.16 [-0.79, 1.12]	*Unclear Difference.*
-3.23 (-32.15, 25.69)	-0.11 [-1.10, 0.88]	*Unclear Difference.*

Level of voluntary contraction (%)



SHAM - WBV raw diff (95% CI)	Cohen *d* [95% CI]	MBI
1.83 (-1.60, 5.27)	0.28 [-0.55, 0.70]	*Unclear Difference.*
2.23 (-1.66, 6.12)	0.37 [-0.28, 1.03]	*Possibly Small effect: SHAM*
4.04 (-0.08, 8.15)	0.88 [-0.02, 1.77]	*Likely Moderate effect: SHA*
-1.23 (-4.31, 1.85)	-0.23 [-0.79, 0.34]	*Unclear Difference.*
2.27 (-1.35, 5.90)	0.53 [-0.32, 1.38]	*Likely Small effect: SHAM*
1.22 (-2.76, 5.20)	0.22 [-0.57, 0.68]	*Unclear Difference.*
1.97 (-3.42, 7.37)	0.33 [-0.57, 1.22]	*Unclear Difference.*
-3.87 (-8.93, 1.19)	-0.55 [-1.27, 0.17]	*Likely Small effect: WBV.*
-1.16 (-8.88, 6.57)	-0.16 [-1.20, 0.88]	*Unclear Difference.*

Twitch peak torque (Nm)



SHAM - WBV raw diff (95% CI)	Cohen *d* [95% CI]	MBI
-0.93 (-4.13, 2.26)	0.17 [-0.59, 0.67]	*Unclear Difference.*
-0.85 (-3.01, 1.32)	0.15 [-0.59, 0.66]	*Unclear Difference.*
-1.70 (-3.72, 0.33)	0.44 [-0.50, 0.73]	*Unclear Difference.*
0.53 (-1.99, 3.06)	0.11 [-0.42, 0.64]	*Unclear Difference.*
-0.21 (-3.18, 2.75)	0.17 [-0.59, 0.67]	*Unclear Difference.*
0.91 (-2.25, 4.06)	0.16 [-0.40, 0.73]	*Unclear Difference.*
-2.26 (-4.71, 0.18)	-0.70 [-1.44, 0.05]	*Likely Moderate effect: WBV.*
-2.52 (-5.58, 0.53)	-0.59 [-1.30, 0.12]	*Likely Small effect: WBV.*
-1.46 (-4.14, 1.21)	-0.35 [-1.00, 0.29]	*Possibly Small effect: WBV.*

Figure 4

Relative changes from baseline.

(A) low-frequency doublet (T_{20}), (B) high-frequency doublet (T_{100}) and (C) low-high torque frequency ratio ($T_{20/100}$) for WBV (connected black triangles) and SHAM (connected white circles) for trials with different set durations (30 s, 60 s and 180 s). Values are expressed as mean and standard errors. Black triangles represent statistically significant WBV differences from baseline (▲▲▲ $p < 0.001$; ▲▲ $p < 0.01$; ▲○ $p < 0.05$). White circles represent statistically significant SHAM differences from baseline (○○○ $p < 0.001$; ○○ $p < 0.01$; ○ $p < 0.05$).

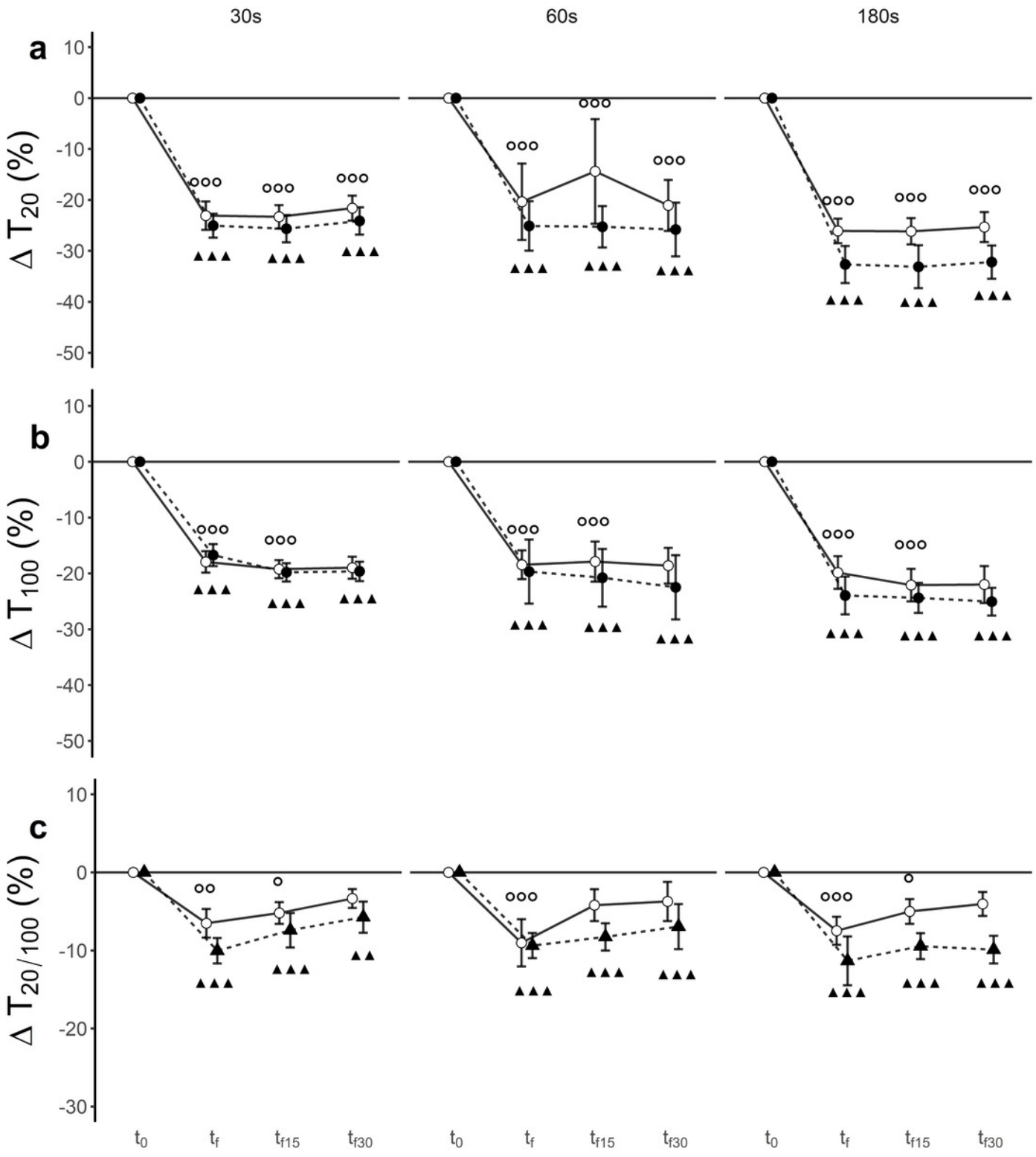
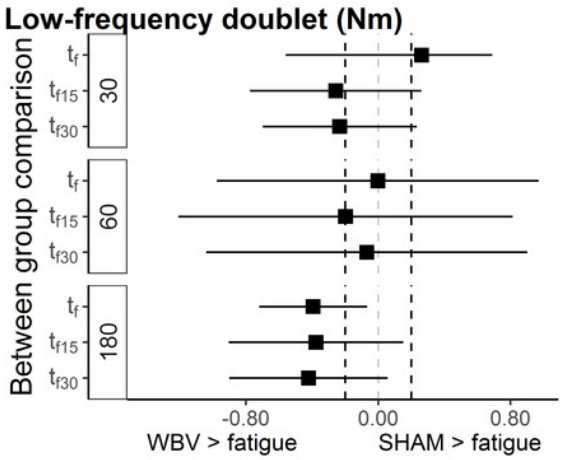

30s
60s
180s
a
b
c
Δ T20 (%)
Δ T100 (%)
Δ T20/100 (%)
10
0
-10
-20
-30
-40
-50
t0
tf
tf15
tf30

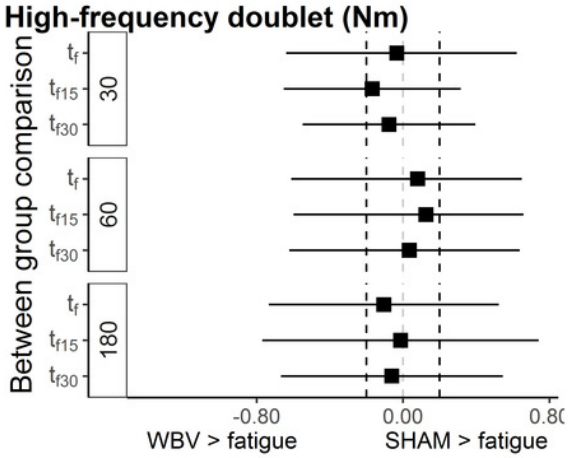
Figure 5

WBV and SHAM between-group comparisons for T_{20}, T_{100}, $T_{20/100}$.

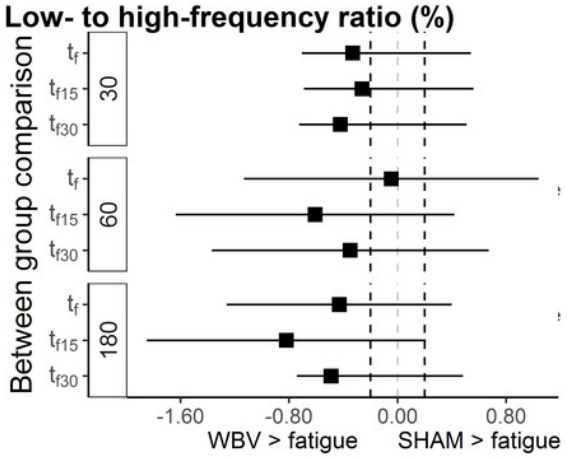
WBV and SHAM between-group comparisons for low-frequency doublet (T_{20}), high-frequency doublet (T_{100}) and low-high torque frequency ratio ($T_{20/100}$). Raw differences (in native units) are presented as mean and 95% confidence intervals. Cohen *d* effect sizes and relative 95% confidence intervals are presented graphically and numerically. The magnitude base inference score is presented as well. Negative effect size values represent higher fatigue in favour of the WBV condition, while positive effect size values represent higher fatigue in favour of the SHAM condition.



	SHAM - WBV raw diff (95% CI)	Cohen *d* [95% CI]	MBI
t_f 30	-2.65 (-8.29, 2.99)	0.26 [-0.56, 0.69]	*Unclear Difference.*
t_{f15} 30	-2.71 (-8.17, 2.75)	-0.26 [-0.78, 0.26]	*Possibly Small effect: WBV.*
t_{f30} 30	-2.41 (-7.23, 2.41)	-0.23 [-0.70, 0.23]	*Possibly Small effect: WBV.*
t_f 60	-0.04 (-16.95, 16.87)	0.00 [-0.98, 0.97]	*Unclear Difference.*
t_{f15} 60	-3.80 (-23.20, 15.60)	-0.20 [-1.21, 0.81]	*Unclear Difference.*
t_{f30} 60	-1.08 (-16.13, 13.97)	-0.07 [-1.04, 0.90]	*Unclear Difference.*
t_f 180	-3.88 (-7.09, -0.66)	-0.39 [-0.72, -0.07]	*Likely Small effect: WBV.*
t_{f15} 180	-4.07 (-9.78, 1.64)	-0.38 [-0.90, 0.15]	*Likely Small effect: WBV.*
t_{f30} 180	-4.55 (-9.72, 0.62)	-0.42 [-0.90, 0.06]	*Likely Small effect: WBV.*



	SHAM - WBV raw diff (95% CI)	Cohen *d* [95% CI]	MBI
t_f 30	0.04 (-4.35, 4.44)	-0.03 [-0.64, 0.62]	*Unclear Difference.*
t_{f15} 30	-1.27 (-4.92, 2.38)	-0.17 [-0.65, 0.32]	*Unclear Difference.*
t_{f30} 30	-0.59 (-4.24, 3.06)	-0.08 [-0.55, 0.40]	*Unclear Difference.*
t_f 60	0.31 (-11.77, 12.40)	0.08 [-0.61, 0.65]	*Unclear Difference.*
t_{f15} 60	-1.28 (-17.57, 15.01)	0.13 [-0.60, 0.66]	*Unclear Difference.*
t_{f30} 60	-2.58 (-20.64, 15.49)	0.03 [-0.62, 0.64]	*Unclear Difference.*
t_f 180	-1.05 (-7.31, 5.22)	-0.10 [-0.73, 0.52]	*Unclear Difference.*
t_{f15} 180	-0.14 (-7.90, 7.62)	-0.01 [-0.77, 0.74]	*Unclear Difference.*
t_{f30} 180	-0.69 (-7.49, 6.12)	-0.06 [-0.67, 0.54]	*Unclear Difference.*



	SHAM - WBV raw diff (95% CI)	Cohen *d* [95% CI]	MBI
t_f 30	-3.53 (-7.79, 0.73)	-0.33 [-0.71, 0.54]	*Unclear Difference.*
t_{f15} 30	-2.34 (-7.67, 2.98)	-0.26 [-0.69, 0.56]	*Unclear Difference.*
t_{f30} 30	-2.47 (-5.83, 0.89)	-0.42 [-0.73, 0.51]	*Unclear Difference.*
t_f 60	-0.38 (-9.31, 8.55)	-0.05 [-1.13, 1.04]	*Unclear Difference.*
t_{f15} 60	-4.08 (-10.98, 2.82)	-0.61 [-1.64, 0.42]	*Unclear Difference.*
t_{f30} 60	-3.32 (-13.01, 6.37)	-0.35 [-1.37, 0.67]	*Unclear Difference.*
t_f 180	-3.57 (-10.45, 3.31)	-0.43 [-1.26, 0.40]	*Unclear Difference.*
t_{f15} 180	-4.54 (-10.25, 1.16)	-0.82 [-1.85, 0.21]	*Likely Moderate effect: WBV.*
t_{f30} 180	-5.59 (-9.92, -1.26)	-0.49 [-0.74, 0.48]	*Unclear Difference.*

Table 1(on next page)

Descriptive statistics (mean and SD), within trial relative change from baseline and Cohen *d* effects size for MVC and %VA.

t_0, baseline; t_f, after intervention; t_{f15}, 15 minutes after intervention; t_{f30}, 30 minutes after intervention.

	t_0	t_f			t_{015}			t_{030}		
	mean (SD)	**mean (SD)**	**Δ %**	**Cohen *d* [95% CI]**	**mean (SD)**	**Δ %**	**Cohen *d* [95% CI]**	**mean (SD)**	**Δ %**	**Cohen *d* [95% CI]**
Maximum voluntary contraction (MVC)										
$SHAM_{30}$	206.04 (67.09)	190.31 (68.15)	-7.63	-0.21 [-0.31, -0.11]	188.93 (68.26)	-8.30	-0.23 [-0.37, -0.09]	191.15 (63.97)	-7.23	-0.21 [-0.35, -0.06]
WBV_{30}	219.46 (64.63)	191.82 (55.25)	-12.59	-0.42 [-0.66, -0.17]	193.06 (67.16)	-12.03	-0.36 [-0.54, -0.18]	192.23 (64.92)	-12.41	-0.38 [-0.54, -0.23]
$SHAM_{60}$	215.61 (64.31)	201.05 (63.94)	-6.75	-0.21 [-0.43, 0.02]	199.34 (61.18)	-7.55	-0.23 [-0.44, -0.03]	200.52 (62.06)	-7.00	-0.22 [-0.45, 0.01]
WBV_{60}	209.75 (63.53)	194.58 (56.78)	-7.23	-0.23 [-0.46, 0.00]	201.37 (57.87)	-3.99	-0.12 [-0.34, 0.09]	197.64 (56.89)	-5.77	-0.18 [-0.38, 0.01]
$SHAM_{180}$	207.81 (62.38)	186.06 (51.73)	-10.47	-0.34 [-0.53, -0.16]	189.68 (52.72)	-8.72	-0.28 [-0.49, -0.08]	188.45 (57.65)	-9.32	-0.29 [-0.47, -0.11]
WBV_{180}	200.36 (62.85)	175.58 (53.91)	-12.37	-0.38 [-0.68, -0.09]	185.85 (61.52)	-7.24	-0.21 [-0.44, 0.02]	177.76 (65.08)	-11.28	-0.32 [-0.66, 0.02]
Level of voluntary activation (%VA)										
$SHAM_{30}$	93.05 (3.00)	89.94 (6.14)	-3.34	-0.58 [-1.22, 0.05]	89.57 (7.14)	-3.74	-0.58 [-1.34, 0.19]	90.12 (4.47)	-3.15	-0.40 [-0.72, 0.52]
WBV_{30}	90.74 (3.98)	89.47 (5.07)	-1.41	-0.25 [-0.97, 0.46]	89.49 (4.58)	-1.38	-0.26 [-0.87, 0.34]	91.85 (4.21)	1.22	0.25 [-0.39, 0.88]
$SHAM_{60}$	89.27 (4.78)	87.71 (6.00)	-1.75	-0.26 [-0.78, 0.26]	88.14 (5.32)	-1.27	-0.20 [-0.74, 0.34]	87.52 (5.39)	-1.96	-0.31 [-0.85, 0.23]
WBV_{60}	90.20 (4.85)	87.41 (5.96)	-3.09	-0.31 [-0.70, 0.55]	91.33 (4.29)	1.26	0.17 [-0.59, 0.67]	89.66 (4.41)	-0.59	-0.03 [-0.64, 0.62]
$SHAM_{180}$	87.40 (6.82)	84.36 (5.47)	-3.47	-0.45 [-1.08, 0.19]	88.73 (5.06)	1.52	0.20 [-0.55, 0.95]	87.45 (6.01)	0.05	0.01 [-0.75, 0.77]
WBV_{180}	88.54 (5.36)	87.48 (4.31)	-1.20	-0.17 [-0.67, 0.59]	86.00 (6.28)	-2.87	-0.40 [-1.03, 0.24]	87.43 (4.33)	-1.25	-0.21 [-0.98, 0.57]

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Table 2 (on next page)

Descriptive statistics (mean and SD), within trial relative change from baseline and Cohen *d* effects size for T_{20}, T_{100}, $T_{20/100}$ and TW_{PT}.

t_0, baseline; t_f, after intervention; t_{f15}, 15 minutes after intervention; t_{f30}, 30 minutes after intervention.

	t_0	t_f				t_{015}			t_{030}	
	mean (SD)	**mean (SD)**	**Δ %**	**Cohen *d* [95% CI]**	mean (SD)	Δ %	Cohen d [95% CI]	mean (SD)	Δ %	Cohen d [95% CI]
Low-frequency doublet (T_{20})										
$SHAM_{30}$	75.26 (18.37)	57.14 (12.56)	-24.07	-1.04 [-1.51, -0.58]	56.97 (11.84)	-24.30	-0.63 [-0.78, 0.42]	57.87 (10.33)	-23.11	-0.63 [-0.78, 0.42]
WBV_{30}	80.47 (20.30)	59.70 (13.91)	-25.80	-1.08 [-1.43, -0.73]	59.47 (15.47)	-26.10	-1.05 [-1.39, -0.72]	60.67 (15.04)	-24.61	-0.63 [-0.78, 0.42]
$SHAM_{60}$	82.03 (22.79)	62.84 (17.27)	-23.39	-0.86 [-1.45, -0.27]	66.78 (19.18)	-18.59	-0.66 [-1.34, 0.03]	62.78 (15.23)	-23.46	-0.90 [-1.39, -0.41]
WBV_{60}	76.11 (18.34)	56.88 (17.20)	-25.26	-0.98 [-1.44, -0.52]	57.06 (17.68)	-25.03	-0.96 [-1.34, -0.58]	55.79 (17.33)	-26.70	-1.03 [-1.52, -0.54]
$SHAM_{180}$	82.24 (17.44)	60.38 (12.87)	-26.58	-0.63 [-0.78, 0.42]	60.35 (13.30)	-26.62	-0.63 [-0.78, 0.42]	60.86 (13.15)	-26.00	-0.63 [-0.78, 0.42]
WBV_{180}	79.59 (18.84)	53.86 (16.14)	-32.33	-1.33 [-1.69, -0.97]	53.62 (17.56)	-32.63	-1.29 [-1.68, -0.90]	53.66 (13.88)	-32.58	-1.42 [-1.82, -1.02]
High-frequency doublet (T_{100})										
$SHAM_{30}$	78.56 (20.46)	63.90 (15.30)	-18.66	-1.04 [-1.51, -0.58]	63.09 (15.68)	-19.69	-0.63 [-0.78, 0.42]	62.73 (13.77)	-20.15	-0.63 [-0.78, 0.42]
WBV_{30}	82.52 (21.65)	67.91 (15.83)	-17.71	-1.08 [-1.43, -0.73]	65.79 (16.42)	-20.28	-1.05 [-1.39, -0.72]	66.10 (16.90)	-19.90	-0.63 [-0.78, 0.42]
$SHAM_{60}$	87.98 (22.21)	71.22 (18.06)	-19.05	-0.86 [-1.45, -0.27]	71.91 (20.34)	-18.27	-0.66 [-1.34, 0.03]	71.07 (18.78)	-19.23	-0.90 [-1.39, -0.41]
WBV_{60}	81.00 (19.98)	64.55 (20.02)	-20.30	-0.98 [-1.44, -0.52]	63.64 (19.06)	-21.43	-0.96 [-1.34, -0.58]	61.50 (18.58)	-24.06	-1.03 [-1.52, -0.54]
$SHAM_{180}$	88.05 (21.36)	69.93 (16.90)	-20.58	-0.63 [-0.78, 0.42]	68.09 (16.89)	-22.68	-0.63 [-0.78, 0.42]	67.69 (15.60)	-23.12	-0.63 [-0.78, 0.42]
WBV_{180}	83.10 (21.07)	63.93 (20.07)	-23.07	-1.33 [-1.69, -0.97]	62.99 (18.42)	-24.20	-1.29 [-1.68, -0.90]	62.06 (16.54)	-25.33	-1.42 [-1.82, -1.02]
Low- to high-frequency doublet ration ($T_{20/100}$)										
$SHAM_{30}$	0.96 (0.06)	0.90 (0.07)	-6.56	-1.04 [-1.51, -0.58]	0.91 (0.08)	-5.15	-0.63 [-0.78, 0.42]	0.93 (0.08)	-3.27	-0.63 [-0.78, 0.42]
WBV_{30}	0.97 (0.05)	0.88 (0.05)	-10.11	-1.08 [-1.43, -0.73]	0.90 (0.07)	-7.50	-1.05 [-1.39, -0.72]	0.92 (0.07)	-5.77	-0.63 [-0.78, 0.42]
$SHAM_{60}$	0.97 (0.07)	0.88 (0.09)	-9.24	-0.86 [-1.45, -0.27]	0.93 (0.08)	-4.27	-0.66 [-1.34, 0.03]	0.94 (0.09)	-3.87	-0.90 [-1.39, -0.41]
WBV_{60}	0.98 (0.06)	0.89 (0.04)	-9.58	-0.98 [-1.44, -0.52]	0.90 (0.06)	-8.42	-0.96 [-1.34, -0.58]	0.91 (0.08)	-7.24	-1.03 [-1.52, -0.54]
$SHAM_{180}$	0.96 (0.08)	0.88 (0.08)	-7.64	-0.63 [-0.78, 0.42]	0.91 (0.09)	-4.98	-0.63 [-0.78, 0.42]	0.92 (0.08)	-4.15	-0.63 [-0.78, 0.42]
WBV_{180}	0.98 (0.06)	0.87 (0.12)	-11.12	-1.33 [-1.69, -0.97]	0.88 (0.07)	-9.53	-1.29 [-1.68, -0.90]	0.88 (0.09)	-9.78	-1.42 [-1.82, -1.02]
Single twitch peak torque (TW_{PT})										
$SHAM_{30}$	26.81 (7.53)	20.62 (6.21)	-23.10	-0.63 [-0.78, 0.42]	20.04 (5.12)	-25.28	-0.63 [-0.78, 0.42]	20.74 (5.39)	-22.65	-0.84 [-1.29, -0.40]
WBV_{30}	26.62 (8.07)	19.50 (5.65)	-26.77	-0.63 [-0.78, 0.42]	19.00 (5.53)	-28.64	-1.00 [-1.36, -0.64]	18.85 (5.34)	-29.19	-1.03 [-1.42, -0.64]
$SHAM_{60}$	27.37 (8.27)	19.83 (4.11)	-27.54	-1.05 [-1.53, -0.56]	19.06 (3.70)	-30.35	-1.18 [-1.81, -0.54]	19.41 (4.35)	-29.10	-1.09 [-1.58, -0.60]
WBV_{60}	26.77 (8.11)	19.76 (5.50)	-26.17	-0.92 [-1.26, -0.57]	18.25 (5.31)	-31.84	-0.63 [-0.78, 0.42]	19.71 (5.78)	-26.37	-0.91 [-1.38, -0.44]
$SHAM_{180}$	26.72 (7.68)	19.85 (6.47)	-25.72	-0.88 [-1.09, -0.67]	18.90 (5.35)	-29.29	-1.07 [-1.36, -0.78]	18.94 (6.39)	-29.15	-0.63 [-0.78, 0.42]
WBV_{180}	27.20 (8.23)	18.06 (7.74)	-33.60	-1.04 [-1.32, -0.75]	16.85 (7.37)	-38.06	-1.20 [-1.58, -0.82]	17.95 (7.35)	-34.01	-1.07 [-1.43, -0.72]

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Table 3(on next page)

Within-trial Tukey corrected t-test comparison with baseline.

Asterisks represent statistically significant differences from baseline (*** $p < 0.001$; ** $p < 0.01$; ** $p < 0.05$). t_0, baseline; t_f, after intervention; t_{f15}, 15 minutes after intervention; t_{f30}, 30 minutes after intervention.

	t_f			t_{f15}			t_{f30}		
visit	t-value	p-value	sig.	t-value	p-value	sig.	t-value	p-value	sig.
Maximum voluntary contraction (MVC)									
$SHAM_{30}$	-2.53	0.04	*	-2.75	0.02	*	-2.40	0.05	
WBV_{30}	-4.45	< 0.001	***	-4.25	< 0.001	***	-4.38	< 0.001	***
$SHAM_{60}$	-2.34	0.06		-2.62	0.03	*	-2.43	0.05	*
WBV_{60}	-2.44	0.05	*	-1.35	0.45		-1.95	0.15	
$SHAM_{180}$	-3.50	0.002	**	-2.92	0.01	*	-3.12	0.006	**
WBV_{180}	-3.99	< 0.001	***	-2.34	0.06		-3.64	0.001	**
Level of voluntary activation (%VA)									
$SHAM_{30}$	-2.02	0.13		-2.26	0.07		-1.90	0.17	
WBV_{30}	-0.83	0.79		-0.81	0.80		0.72	0.85	
$SHAM_{60}$	-1.01	0.68		-0.74	0.84		-1.14	0.59	
WBV_{60}	-1.81	0.20		0.74	0.84		-0.35	0.98	
$SHAM_{180}$	-1.97	0.14		0.86	0.77		0.03	1.00	
WBV_{180}	-0.69	0.87		-1.65	0.27		-0.72	0.85	
Low-frequency doublet (T_{20})									
$SHAM_{30}$	-6.08	< 0.001	***	-6.14	< 0.001	***	-5.84	< 0.001	***
WBV_{30}	-6.97	< 0.001	***	-7.05	< 0.001	***	-6.65	< 0.001	***
$SHAM_{60}$	-6.44	< 0.001	***	-5.12	< 0.001	***	-6.46	< 0.001	***
WBV_{60}	-6.46	< 0.001	***	-6.40	< 0.001	***	-6.82	< 0.001	***
$SHAM_{180}$	-7.34	< 0.001	***	-7.35	< 0.001	***	-7.18	< 0.001	***
WBV_{180}	-8.64	< 0.001	***	-8.72	< 0.001	***	-8.71	< 0.001	***
High-frequency doublet (T_{100})									
$SHAM_{30}$	-5.42	< 0.001	***	-5.71	< 0.001	***	-5.85	< 0.001	***
WBV_{30}	-5.40	< 0.001	***	-6.18	< 0.001	***	-6.07	< 0.001	***
$SHAM_{60}$	-6.19	< 0.001	***	-5.94	< 0.001	***	-6.25	< 0.001	***
WBV_{60}	-6.08	< 0.001	***	-6.41	< 0.001	***	-7.20	< 0.001	***
$SHAM_{180}$	-6.70	< 0.001	***	-7.38	< 0.001	***	-7.52	< 0.001	***
WBV_{180}	-7.08	< 0.001	***	-7.43	< 0.001	***	-7.78	< 0.001	***
Low- to high-frequency doublet ration ($T_{20/100}$)									
$SHAM_{30}$	-3.57	0.001	**	-2.81	0.02	*	-1.78	0.21	
WBV_{30}	-5.57	< 0.001	***	-4.13	< 0.001	***	-3.18	0.005	**
$SHAM_{60}$	-5.09	< 0.001	***	-2.35	0.06		-2.13	0.10	
WBV_{60}	-5.30	< 0.001	***	-4.66	< 0.001	***	-4.01	< 0.001	***
$SHAM_{180}$	-4.12	< 0.001	***	-2.69	0.02	*	-2.24	0.08	
WBV_{180}	-6.14	< 0.001	***	-5.26	< 0.001	***	-5.40	< 0.001	***
Single twitch peak torque (TW_{PT})									
$SHAM_{30}$	-5.70	< 0.001	***	-6.24	< 0.001	***	-5.59	< 0.001	***
WBV_{30}	-6.56	< 0.001	***	-7.01	< 0.001	***	-7.15	< 0.001	***
$SHAM_{60}$	-6.93	< 0.001	***	-7.64	< 0.001	***	-7.33	< 0.001	***
WBV_{60}	-6.44	< 0.001	***	-7.84	< 0.001	***	-6.49	< 0.001	***
$SHAM_{180}$	-6.32	< 0.001	***	-7.20	< 0.001	***	-7.17	< 0.001	***
WBV_{180}	-8.41	< 0.001	***	-9.52	< 0.001	***	-8.51	< 0.001	***

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