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José Mira, Thomas Lapole, Robin Souron, Laurent Messonnier, Guillaume Y. Millet, et al.. Cortical voluntary activation testing methodology impacts central fatigue. *European Journal of Applied Physiology*, 2017, 117 (9), pp.1845-1857. <10.1007/s00421-017-3678-x>. <hal-02409531>

HAL Id: hal-02409531

<https://hal.science/hal-02409531v1>

Submitted on 1 Sep 2023

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Cortical voluntary activation testing methodology impacts central fatigue

José Mira^{1,2} · Thomas Lapole³ · Robin Souron³ · Laurent Messonnier¹ · Guillaume Y. Millet² · Thomas Rupp¹

Abstract

Purpose Currently, cortical voluntary activation (VA_{TMS}) is assessed by superimposing transcranial magnetic stimulation (TMS) on a maximal voluntary contraction (MVC), 75% MVC and 50% MVC, each contraction being interspersed with 5–10 s of relaxation. Here, we assessed whether this traditional approach (TRADI) underestimates central fatigue due to this short recovery compared to a continuous method (CONTI).

Methods VA_{TMS} , motor-evoked potential (MEP), and cortical silent period (CSP) of the *vastus lateralis* were determined in 12 young healthy adults before and after a 2-min sustained MVC of knee extensors in two randomly assigned sessions. In TRADI, evaluations comprised a 7-s rest between the three contractions (100, 75, and 50% MVC) and evaluation following the 2-min sustained MVC started after a minimal rest (3–4 s). In CONTI, evaluations were performed with no rest allowed between the three levels of contraction, and evaluation after the 2-min sustained MVC commenced without any rest.

Results MVC was equally depressed at the end of the 2 min in both conditions. Post 2-min sustained MVC,

VA_{TMS} change was greater in CONTI than in TRADI (−29 (15)% [−42, −17] vs. −9 (4)% [−13, −5], respectively, $P < 0.001$). Differences were also observed between TRADI and CONTI for MEP and CSP immediately after the fatiguing exercise. All differences between the two methods disappeared after 2 min of recovery.

Conclusion After a 2-min sustained MVC, a few seconds of recovery change the amount of measured VA_{TMS} and associated parameters of central fatigue. The continuous method should be preferred to determine deficits in voluntary activation.

Keywords Central fatigue · Corticospinal excitability · Intracortical inhibition · Maximal muscle contraction · Recovery · Transcranial magnetic stimulation

Abbreviations

BF	Biceps femoris
CI	Confidence interval
CONTI	Continuous method
CSP_{50} – CSP_{75} – CSP_{100}	Cortical silent period at 50–75–100% MVC
EMG	Electromyography
ERT	Estimated resting twitch
FNES	Femoral nerve electrical stimulation
MEP_{50} – MEP_{75} – MEP_{100}	Motor-evoked potential at 50–75–100% MVC
M-wave	Muscle compound action potential on the relaxed muscle
Msup	Muscle compound action potential measured during voluntary contraction (50% MVC)
MVC	Maximal voluntary contraction
NMF	Neuromuscular function

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POST	At the end of the 2-min sustained MVC
PRE	Before the 2-min sustained MVC
Pt	Peak twitch
R2–R4–R6	2–4–6 min after the end of the 2-min sustained MVC
RF	Rectus femoris
RMS	Root-mean square
SIT	Superimposed twitch
VA	Voluntary activation
VA _{TMS}	Cortical voluntary activation
VL	Vastus lateralis
TMS	Transcranial magnetic stimulation
TRADI	Traditional method

Introduction

Neuromuscular fatigue has been defined as any exercise-induced decline in maximal force or power produced by a muscle (Bigland-Ritchie and Woods 1984; Gandevia 2001). Fatigue is generally quantified as a decrement in maximal isometric strength that develops soon after exercise onset. The underlying mechanisms may originate at any level of the motor pathway and are usually divided into central and peripheral sites of failure. The former refers to the decline in voluntary activation or neural drive of the central nervous system (Goodall et al. 2012a), while the latter concerns alterations at or distal to the neuromuscular junction.

Peripheral fatigue is classically identified as a decrease in twitch or tetanic force evoked by peripheral nerve stimulation on relaxed muscles (Millet et al. 2011). The most common method to characterize central fatigue is maximal voluntary activation (VA) calculated by twitch interpolation (Merton 1953). This technique consists of superimposing peripheral electrical stimulation during a maximal voluntary contraction (MVC), which results in a superimposed twitch (SIT). The greater the SIT relative to a reference stimulation on a relaxed muscle, the lower the VA. This observed activation deficit may be due to the inability of the motor units to fire quickly enough or to a failure in their spatial recruitment (Gandevia 2001). Any decrease in VA with exercise characterizes the presence of central fatigue, which may originate anywhere at or above the site of the stimulation. Consequently, it is not possible to distinguish the changes happening within the central nervous system. The use of other techniques is, therefore, helpful to identify the different failure sites proximal to the neuromuscular junction, which may be spinal and/or supraspinal.

Transcranial magnetic stimulation (TMS) has grown in popularity due to its clinical significance in the diagnosis

[e.g., idiopathic generalized epilepsy (Reutens et al. 1993)] and possible treatment of diseases [e.g., depression (Triggs et al. 1999) and Parkinson's disease (Siebner et al. 1999)]. In addition, single and double pulse TMS (e.g., motor-evoked potential, MEP; cortical silent period, CSP; short and long intracortical inhibitions) is extensively used to understand and characterize fatigue etiology in sport (Goodall et al. 2012b) and medical conditions such as multiple sclerosis (Thickbroom et al. 2006), cancer-related fatigue (Cai et al. 2014), and chronic fatigue syndrome (Sacco et al. 1999). In the voluntary activation calculation through TMS (VA_{TMS}), the superimposed stimulus is evoked by magnetic stimulation applied directly over the motor cortex. If a SIT is observed during a MVC, it means that a reserve of force, which was not used voluntarily, was still available. Thus, motor cortical output was unable to drive all motor units maximally before stimulation (Taylor et al. 2005), which may suggest that part of the central failure could be supraspinal (Gandevia et al. 1996). Nonetheless, a decline in VA_{TMS} after exercise does not exclusively indicate the presence of supraspinal fatigue, as fatigue at the spinal level may still occur.

Since TMS is a relatively recent technique, most of the experiments used peripheral stimulation to assess VA. Nonetheless, VA_{TMS} and its changes with fatigue have been shown to be reliable for the upper (Todd et al. 2004) and lower (Goodall et al. 2009; Sidhu et al. 2009a) limbs. Since corticospinal excitability is enhanced during voluntary contraction (Todd et al. 2003), the reference twitch needs to be estimated. To do so, a linear regression between SIT and voluntary force is drawn to obtain an estimated resting twitch (ERT) by extrapolation (Todd et al. 2003). For a linear relationship between TMS and voluntary force to reliably estimate the resting twitch, TMS should be evoked at contractions above 50% MVC (Goodall et al. 2009, 2012b). Hence, the most currently used method encompasses a superimposed TMS during an MVC and two submaximal contractions (75% MVC and 50% MVC), interspersed with a short period (5–10 s) between contractions.

When investigated for a few seconds and up to several minutes after repeated or sustained contractions, VA_{TMS} has been observed to decline in upper (Gandevia et al. 1996; Hunter et al. 2008; Szubski et al. 2007) and lower (Bachasson et al. 2016; Goodall et al. 2009; Gruet et al. 2014; Mileva et al. 2012) limbs. However, considering the rapid recovery of neuromuscular function that occurs within the first 2 min after exercise (Froyd et al. 2013), the rate of fatigue, and notably central fatigue, which appears to recover more quickly than peripheral fatigue, may have been underestimated in these studies. For instance, Gruet et al. (2014) reported a full recovery of knee extensors VA_{TMS} 96 s after high-intensity quadriceps contractions to task-failure, whereas peak twitch (Pt) only recovered

partially 276 s after exercise. As a consequence, existing evidence may not reflect the true central fatigue induced by exercise.

Therefore, the purpose of the current study was to compare the traditional method (TRADI) of VA_{TMS} assessment with a continuous method (CONTI), where neuromuscular function was evaluated with no rest between contractions and immediately at the end of exercise. We hypothesized that VA_{TMS} deficit would be smaller in TRADI due to the rapid recovery of central fatigue, confirming the need to use a CONTI method to assess the actual VA depression induced by exercise. Since peripheral fatigue recovery is slower, we also hypothesized that the methodology would either have a minimal or non-existent effect on it.

Materials and methods

Subjects

Twelve healthy active men [mean (SD), age 23 (3) years, height 181 (6) cm, and mass 74 (8) kg] participated in the study. Written informed consent was obtained from all subjects prior to their participation. This study was approved by the ethics board at the University of Calgary (REB 14-1625) and conformed to the standards from the latest edition of the *Declaration of Helsinki*. All subjects were in good physical condition with no signs or symptoms of neurological, cardiovascular, circulatory or orthopedic disorders, or contraindications to TMS. Subjects refrained from any type of physical exercise at least 2 days before testing and did not drink any beverages with caffeine on test days.

Experimental protocol

To compare the rate of central fatigue using the CONTI and TRADI methods, VA_{TMS} was determined before and after a 2-min sustained MVC of knee extensors in two randomly assigned sessions, performed on the same day, with a wash-out period of 1.5 h. Corticospinal excitability (through MEP) and intracortical inhibition (through CSP) were also measured with TMS. Femoral nerve electrical stimulation (FNES) was applied to assess neuromuscular transmission (through muscle compound action potential, M-wave) and contractile properties (through Pt). Electromyography activity (EMG) of vastus lateralis (VL), rectus femoris (RF), and biceps femoris (BF) muscles was monitored throughout the sessions.

Figure 1a illustrates the experimental protocol, which consisted, after the baseline measurements, in two times

the following with a 1.5-h wash-out period: neuromuscular function (NMF) tests before (PRE), immediately after (POST) and 2–4–6 min after (R2–R4–R6, respectively) a 2-min sustained MVC (fatiguing task). The baseline measurements (Fig. 1b) comprised (a) the determination of stimulation intensity to elicit the highest M-wave and Pt response in the unfatigued state (this intensity was further used for stimulation after the fatiguing protocol); (b) a warm-up composed of 20 submaximal contractions with progressive increases in force followed by two 3–4-s MVCs interspersed with 1 min to set the percentage MVC used for optimal TMS site and intensity determination; (c) the determination of optimal TMS site; and (d) the determination of optimal TMS intensity. The NMF tests performed before the fatiguing task (PRE) were performed twice before each 2-min sustained MVC. Figure 1c shows the NMF tests methodology to assess central (VA_{TMS} , MEP, and CSP) and peripheral (Pt, M-wave) fatigue. The CONTI method comprised a single voluntary contraction including an initial ~3–4-s MVC followed by a voluntary reduction in force output to match 75% MVC and then 50% MVC with no rest period in-between. To allow for performance of this continuous voluntary contraction without any rest between the 3 levels of force (i.e., MVC, 75 and 50% MVC), one experimenter monitored the instantaneous values of force produced during MVC (LabChart software was continuously providing the highest value of force achieved). Then, after the TMS had been delivered during the MVC, the scale of the force channel of LabChart was reset to the maximum value of force produced before TMS delivery. Since before the start of the experiment, two red twines (target line) were affixed on the screen corresponding to 75 and 50% of the maximum force value displayed in the scale of the force channel, resetting this maximum value ensured that the upper twine corresponded to 75% and the lower twine to 50% of the maximum value represented in the channel. This reset was displayed in less than 2 s after TMS, while the subject was either relaxing between contractions (TRADI) or still contracting at the previous torque level (CONTI). This procedure enabled the performance of the NMF test right after the 2-min sustained MVC without any rest in CONTI.

The TRADI method consisted of three ~3–4-s voluntary contractions: one MVC, one 75% MVC, and one 50% MVC, each contraction being interspersed by ~7-s relaxation periods. In TRADI, a 3–4-s rest was also allowed between exercise termination and fatigue evaluation. This was not the case in CONTI. During voluntary contractions, TMS was only delivered when the subjects attained and maintained the desired force level. Immediately after TMS delivery, subjects were asked to re-contract at the same force level as fast as possible.

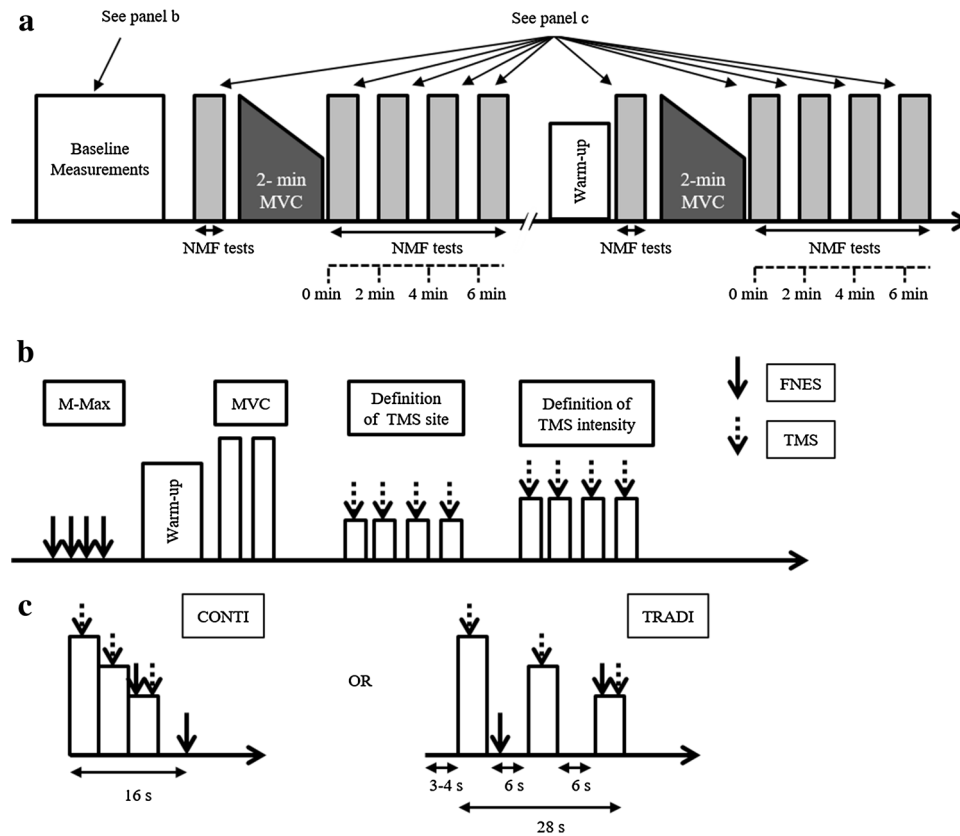


Fig. 1 **a** Experimental protocol, which consisted of baseline measurements and two times the following testing (interspersed by a 1.5-h wash-out period): neuromuscular function (NMF) tests before (PRE), immediately after (POST) and 2–4–6 min (R2–R4–R6) after a 2-min sustained MVC (fatiguing task). **b** Baseline measurements, which comprised: (a) electrically evoked responses to elicit the highest M-wave and twitch response in the unfatigued state; (b) warm-up followed by $2 \times 3\text{--}4\text{-s}$ MVCs; (c) determination of optimal TMS site;

and (d) determination of optimal TMS intensity. The *black arrow* indicates femoral nerve electrical stimulation (FNES), whereas the *dotted arrow* represents the transcranial magnetic stimulation (TMS). **c** The NMF tests methodology to assess central and peripheral fatigue. The continuous method (CONTI) comprised one unique voluntary contraction, while the traditional method (TRADI) consisted of three $\sim 3\text{--}4\text{-s}$ voluntary contractions (MVC, 75% MVC, and 50% MVC) interspersed by $\sim 7\text{-s}$ relaxation periods

Dependent variables

Force measurements

Subjects sat on a custom-built quadriceps chair with the right hip angle set at 90° and knee joint angle set at 100° of flexion (full knee extension = 180°). A noncompliant strap connected to a force transducer was attached 3–5 cm above the subject's lateral malleoli. Force data were acquired using a PowerLab data acquisition system (16/30—ML880/P, ADInstruments, Bella Vista, Australia) at a sampling rate of 2 kHz. The subjects were firmly secured to the chair with noncompliant straps to minimize body movement. The target force to meet during each submaximal contraction of the TMS testing procedure and real-time visual feedback of the force produced was provided to the subject via custom software and macros (LabChart 7, ADInstruments) on a wide computer screen throughout the experiment. Strong verbal

encouragement was given during brief and prolonged MVCs.

Electromyographic recordings

The EMG signals of the right VL, RF and BF were recorded using bipolar silver chloride surface electrodes of 10-mm diameter (Type 0601000402, Contrôle Graphique Medical) during the voluntary contractions and electrical/magnetic stimuli. The recording electrodes were secured lengthwise to the skin over the muscle belly, with an inter-electrode distance of 25 mm. The positions of the electrodes were marked on the skin in case replacement was necessary. The reference electrode was attached to the patella. Low impedance ($Z < 5\text{ k}\Omega$) at the skin–electrode surface was obtained by abrading the skin with fine sand paper and cleaning with alcohol. EMG data were recorded with a PowerLab system (16/30—ML880/P, ADInstruments) with a sampling frequency of 2 kHz. The EMG signal was amplified

with octal bio-amplifier (Octal Bioamp, ML138, ADInstruments) with a bandwidth frequency ranging from 5 to 500 Hz (input impedance = 200 M Ω , common mode rejection ratio = 85 dB, gain = 1000), transmitted to the PC and analyzed with the LabChart 7 software (ADInstruments).

Femoral nerve electrical stimulation

Electrical stimulation was delivered percutaneously to the femoral nerve via a self-adhesive electrode (10-mm diameter, Ag–AgCl, Type 0601000402, Contrôle Graphique Medical, Brie-Comte-Robert, France). The anode, a 10 \times 5 cm self-adhesive stimulation electrode (Medicomplex SA, Ecublens, Switzerland), was located in the gluteal fold. A constant current stimulator (Digitimer DS7A, Hertfordshire, United Kingdom) was used to deliver a square-wave stimulus of 1-ms duration with maximal voltage of 400 V. The optimal stimulus intensity was determined from knee extensors response producing both maximal Pt and maximal M-wave amplitude. The stimulating intensity [151 (67) mA] was supramaximal (i.e., 120% of optimal intensity) and held constant throughout the protocol. Supramaximal FNES was delivered: (1) 2 s after the MVC in relaxed muscle in TRADI or 2 s after last 50% MVC in relaxed muscle in CONTI and (2) during the 50% MVC (see Fig. 1c).

Transcranial magnetic stimulation

A magnetic stimulator (Magstim 200², The Magstim Company, Dyfed, United Kingdom) was used to stimulate the motor cortex area of the right leg. Single TMS pulses of 1-ms duration were delivered via a concave double-cone coil (diameter 110 mm, maximum output 1.4 T) positioned over the vertex of the scalp and held tangentially to the skull. To determine optimal TMS site, the coil was positioned to preferentially activate the left motor cortex (contralateral to the right leg) and elicit the largest MEP in the VL with only a small MEP in the BF (<10% of VL MEP) during isometric knee extension at 10% MVC with a stimulus intensity of 50% of maximal stimulator power output (Sidhu et al. 2009a). The optimal stimulus site was marked on a breathable swimming cap, which was worn on the head throughout the protocol to ensure reproducibility of the stimulus conditions for each subject. To determine optimal stimulation intensity, TMS was elicited during brief (~5 s) submaximal (20% MVC) isometric knee extensions at 30, 40, 50, 60, 70, and 80% of maximal stimulator power output in random order. Four consecutive contractions were performed at each stimulus intensity with 10 s of rest between contractions. The stimulus intensity [68 (10)% of maximal stimulator power output] that elicited the largest right VL MEPs and SIT with small MEP of the right BF

(<10% of VL MEP) was considered optimal and employed throughout the protocol, as previously suggested (Rupp et al. 2012). If a plateau was not reached, higher intensities (i.e., 90 and 100%) were investigated.

Data analysis

Mechanical responses

Peak forces measured during MVC were calculated as the maximal values obtained before the stimuli. At PRE, the highest of the two MVCs was chosen for further analysis. To calculate global force quantity (intensity \times duration) produced throughout the 2-min sustained MVC (Rozand et al. 2014), force time integral (N s) was calculated and summed over eight periods of 15 s along the 2 min. Pt was also determined.

M-wave

M-wave peak-to-peak amplitudes, durations, and areas were calculated from VL EMG responses to single FNES in the relaxed muscle (M-wave) and at 50% MVC (Msup, Fig. 1c).

TMS parameters

Motor-evoked potential areas of quadriceps muscles during submaximal (MEP₅₀ and MEP₇₅ for 50 and 75% MVC, respectively) and maximal contractions (MEP₁₀₀) were normalized to Msup (Sidhu et al. 2012). The duration of the CSP during submaximal (CSP₅₀ and CSP₇₅ for 50 and 75% MVC, respectively) and maximal contractions (CSP₁₀₀) was determined manually as the interval from the stimulus to the return of continuous EMG activity (Sidhu et al. 2009b). VA_{TMS} was quantified by measurement of the superimposed force responses to TMS. At PRE, only the highest VA_{TMS} value from the two NMF tests was used for further analysis. The SIT amplitude evoked during contractions at 100, 75, and 50% MVC was calculated, and the y-intercept of the linear regression between the mean SITs and voluntary force was used to quantify ERT (Todd et al. 2003). For both TRADI and CONTI, VA_{TMS} (%) was then calculated using the following equation:

$$VA_{TMS} = [1 - (SIT/ERT)] \times 100.$$

The reliability of this method for the determination of VA_{TMS} has been previously described for the knee extensors (Goodall et al. 2009; Sidhu et al. 2009a). 6 VA_{TMS} out of a total of 120 assessments had to be ruled out, as 5 showed a poor linear regression between SIT and voluntary force and 1 POST NMF test (in TRADI condition) was not performed by a subject.

EMG during 2-min sustained MVC

During the 2-min sustained MVC, the EMG activity of VL, RF, and BF was calculated as the average root-mean-square (RMS) over the same 15-s periods, as for force analysis (see above).

Statistical analyses

Data are presented in the text as mean (SD) and 95% confidence interval [CI]. Data are shown in the figures and tables as mean (SD). The statistical analyses were performed with Statistica (version 8, Tulsa, USA). Normality of distribution and homogeneity of variances of the main variables were checked using a Shapiro–Wilk normality test and the Levene’s test, respectively. A two-way ANOVA (condition \times time) with repeated measures was performed for the two conditions (CONTI and TRADI) and five timepoints of each condition (PRE, POST, R2, R4, and R6) or nine timepoints when investigating torque/EMG kinetics during the 2-min MVCs (MVC PRE, 0–15, 15–30, 30–45, 45–60, 60–75, 75–90, 90–105, and 105–120 s). Because VA_{TMS} was not normally distributed at PRE, this parameter was analyzed with a two-way ANOVA with repeated measures performed on the change from baseline measurements, i.e., condition (CONTI, TRADI) \times time (Δ PRE-POST, Δ PRE-R2, Δ PRE-R4, and Δ PRE-R6). Where the ANOVA revealed significant main effects or interactions, the data were further explored using pair-wise comparisons with a Bonferroni correction. For all statistical analyses, an alpha level of 0.05 was used as the cutoff for significance.

Results

Fatiguing protocol

Figure 2 shows the typical force trace and EMG VL during the TRADI and CONTI evaluation of one subject. Force and EMG VL values were not different between TRADI and CONTI during the 2-min sustained MVC (Fig. 3). Force was significantly depressed at the end of the 2-min MVC ($P < 0.001$) to a similar extent in both conditions (-75 (5)% [$-78, -73$] vs. -74 (7)% [$-77, -70$] in TRADI and CONTI, respectively). VL and RF EMG were depressed at the end of the 2-min MVC, with no differences between conditions. In addition, force quantity produced during the 2-min MVC showed no differences between TRADI and CONTI (28.7 (4.5) kN s [$26.1, 31.2$] vs. 28.0 (4.6) kN s [$25.4, 30.6$], respectively, NS).

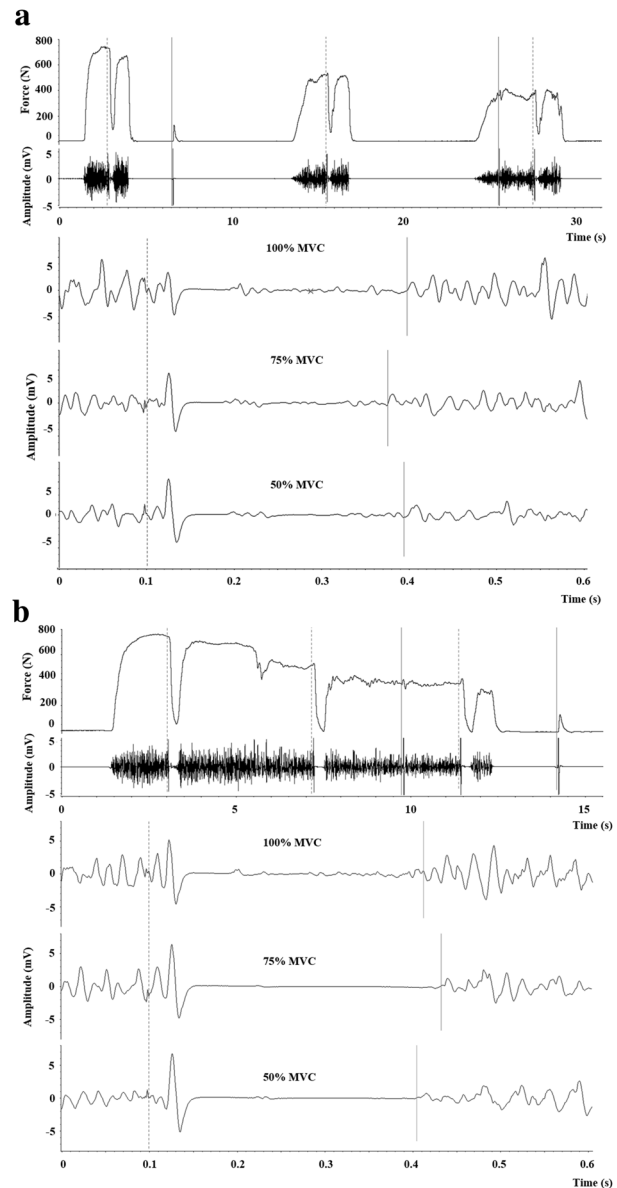
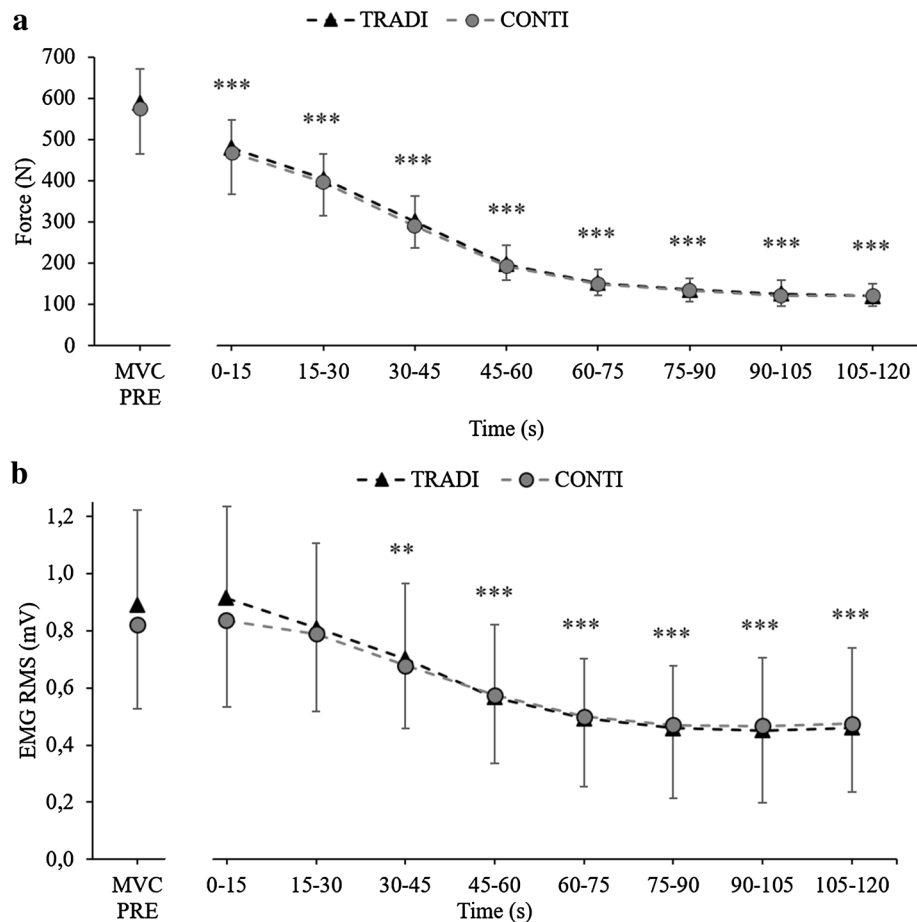


Fig. 2 Typical force trace and VL EMG during TRADI (a) and CONTI (b) evaluation before the fatiguing task. The bottom part of the panel shows the MEP and CSP at MVC, 75% MVC and 50% MVC. Dotted lines represent TMS. The solid lines at the top represent FNES, whereas at the bottom they represent the end of the CSP

Maximal voluntary force

Maximal voluntary contraction values are reported in Fig. 4a. At PRE, MVCs were similar in both conditions. MVC changes from PRE to POST were significantly greater in CONTI compared to TRADI (-79 (6)% [$-83, -76$] vs. -45 (10)% [$-51, -40$], respectively, $P < 0.001$). At R2, R4, and R6, the MVCs were not different between conditions. Despite a partial recovery, MVC at R6 was still lower than at baseline both in TRADI (-15 (8)%

Fig. 3 **a** Force during MVC PRE and during 15-s periods of the 2-min sustained MVC. **b** EMG VL during MVC PRE and during 15-s periods of the 2-min sustained MVC. Significantly different from MVC PRE: ** $P < 0.01$, *** $P < 0.001$



[-19, -10], $P < 0.001$) and in CONTI (-15 (7)% [-19, -11], $P < 0.001$).

Peripheral fatigue

Mechanical and EMG responses to electrical stimulation are shown in Table 1. Pt did not differ between conditions at PRE. At POST, Pt was significantly depressed ($P < 0.001$) to a similar extent in both conditions (-62 (15)% [-71, -53] vs. -63 (18)% [-73, -52] in TRADI and CONTI, respectively). Values at R6 indicated a partial and similar recovery in TRADI and CONTI. The M-wave area was not different between conditions at PRE. There was a significant effect of time ($P < 0.001$), i.e., M-wave area increased similarly in both TRADI and CONTI at POST (17 (9)% [11, 22] vs. 17 (11)% [10, 24], respectively, $P < 0.001$), R2 (20 (6)% [16, 24] vs. 11 (11)% [5, 18], respectively, $P < 0.001$), R4 (8 (21)% [-4, 21] vs. 7 (8)% [3, 12], respectively, $P = 0.001$), and R6 (10 (8)% [5, 15] vs. 5 (9)% [0, 10], respectively, $P = 0.009$).

Voluntary activation

VA_{TMS} at PRE was not different between conditions (Fig. 4b). At POST, VA_{TMS} decreased significantly in both conditions, but the decline was significantly greater in CONTI compared to TRADI (-29 (15)% [-40, -19] vs. -9 (4)% [-12, -6], respectively, $P < 0.001$). At R2, VA_{TMS} was not significantly different compared to PRE anymore for both conditions.

Corticospinal excitability and intracortical inhibition

Figure 5 shows the changes in MEP during fatigue and recovery. At PRE, MEP_{100} , MEP_{75} , and MEP_{50} did not differ between CONTI and TRADI. At POST, MEP_{100} showed a significant increase in CONTI only (115 (63)% [76, 154], $P < 0.001$) and this parameter was significantly greater in CONTI than in TRADI at this timepoint ($P = 0.004$). From R2, MEP_{100} values were not different from PRE, whatever the conditions. MEP_{75} remained unchanged throughout the protocol for both conditions. There was a significant effect of time ($P < 0.001$) for MEP_{50} , as it decreased similarly

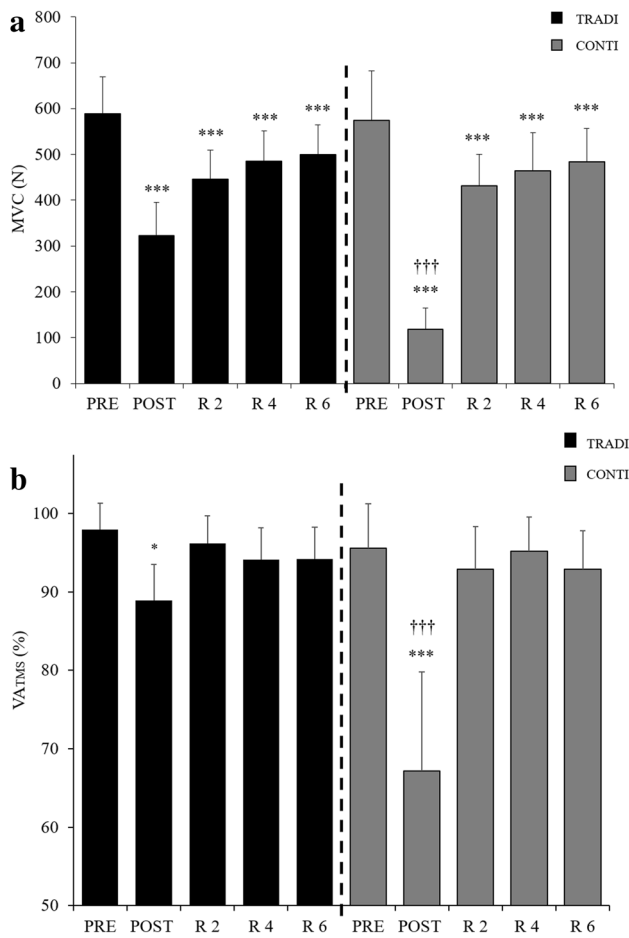


Fig. 4 a Maximal voluntary contraction force (MVC) before (PRE), at the end of the 2-min sustained MVC (POST), and 2 min (R2), 4 min (R4), and 6 min (R6) after exercise. $N = 12$. **b** Maximal voluntary activation by TMS (VA_{TMS}) before (PRE), at the end of the 2-min sustained MVC (POST), and 2 min (R2), 4 min (R4), and 6 min (R6) after exercise. $N = 8$. Dashed line indicates wash-out period of 1.5 h. Data are mean (SD) in TRADI and CONTI conditions. Significantly different from PRE of the same condition: * $P < 0.05$, *** $P < 0.001$; significantly different from the same time-point in TRADI: ††† $P < 0.001$

in both conditions at POST (-10 (24)% [-25 , 4] vs. -33 (19)% [-45 , -21] in TRADI and CONTI, respectively). At R2, MEP_{50} had completely recovered in both conditions.

Changes in CSP are shown in Table 1. At PRE, CSP_{100} , CSP_{75} , and CSP_{50} did not differ between CONTI and TRADI. CSP_{100} was significantly greater at POST in CONTI only (22 (18)% [11, 32], $P = 0.003$). CSP_{75} analysis showed an effect of time. Although no significant changes were observed compared to PRE, CSP_{75} was significantly longer at POST compared to R4 (3 (13)% [-5 , 11] vs. 8 (17)% [-2 , 18] in TRADI and CONTI, respectively, $P = 0.017$) and R6 (8 (11)% [2, 15] vs. 7 (13)% [0, 15] in TRADI and CONTI, respectively, $P = 0.002$). There was a significant time effect for CSP_{50} , as it was significantly longer at POST compared

Table 1 Peak twitch force (Pt), EMG responses to single-pulse stimuli on the relaxed vastus lateralis muscle (M-wave), and vastus lateralis cortical silent period (CSP) during contractions at 50, 75, and 100% of MVC before (PRE), at the end of the 2-min sustained MVC (POST), and 2 min (R2), 4 min (R4), and 6 min (R6) after exercise

	TRADI					CONTI				
	Pre	Post	R2	R4	R6	Pre	Post	R2	R4	R6
Pt (N)	136 (33)	51 (22)†††	94 (30)†††	96 (34)†††	94 (33)†††	132 (26)	47 (19)†††	94 (28)†††	100 (23)†††	103 (17)†††
M-wave amplitude (mV)	15.9 (5.4)	16.7 (5.2)	16.7 (5.3)	16.0 (6.2)	15.0 (7.0)	16.1 (5.1)	16.2 (4.9)	16.6 (5.1)	16.9 (5.3)	16.8 (5.2)
Duration (ms)	10.7 (2.2)	11.7 (2.6)	11.2 (2.5)	11.4 (3.0)	10.4 (2.4)	10.1 (1.8)	10.8 (2.7)	10.5 (2.2)	10.0 (2.0)	9.9 (2.0)
Area ($\mu V \cdot s$)	106 (38)	122 (42)†††	127 (47)†††	119 (50)††	117 (43)††	104 (40)	121 (42)†††	116 (41)††	113 (43)††	111 (43)††
CSP_{50} (ms)	303 (71)	322 (82)††	287 (74)	305 (98)	284 (56)	319 (78)	360 (71)††	325 (74)	314 (71)	313 (77)
CSP_{75} (ms)	297 (79)	304 (80)	296 (67)	292 (68)	277 (73)	316 (76)	344 (80)	314 (72)	311 (71)	316 (72)
CSP_{100} (ms)	302 (82)	314 (92)	297 (83)	319 (100)	301 (72)	298 (81)	357 (84)**	299 (70)	296 (84)	295 (77)

$N = 11$. Data are mean (SD) for TRADI and CONTI conditions

Significantly different from PRE (main effect of time): ** $P < 0.01$, *** $P < 0.001$; significantly different from PRE of the same condition: ** $P < 0.01$

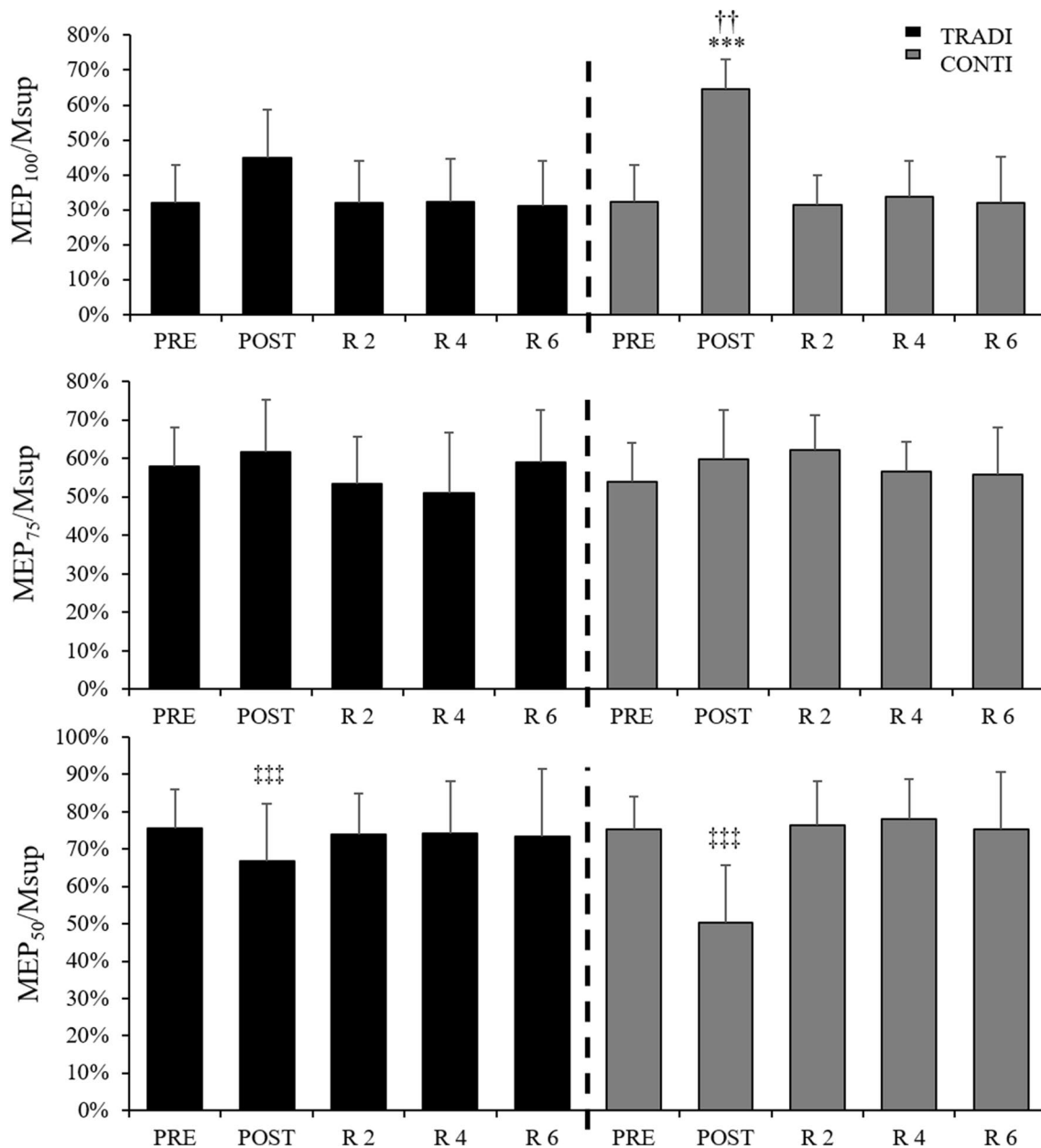


Fig. 5 Vastus lateralis motor-evoked potential areas (MEP) assessed by TMS and normalized to Msup areas. Data are presented before (PRE), at the end of the 2-min sustained MVC (POST), and 2 min (R2), 4 min (R4), and 6 min (R6) after exercise. **a–c** MEPs obtained during MVC (MEP₁₀₀) and during submaximal contraction at 75% MVC (MEP₇₅) and 50% MVC (MEP₅₀), respectively. $N = 10$. Dashed

line indicates wash-out period of 1.5 h. Data are mean (SD) for TRADI and CONTI conditions. Significantly different from PRE of the same condition: $***P < 0.001$; significantly different from the same timepoint in TRADI: $††P < 0.01$; significantly different from PRE (main effect of time): $†††P < 0.001$

to PRE in both conditions (7 (10)% [1, 12] vs. 15 (10)% [9, 21] in TRADI and CONTI, respectively, $P = 0.002$).

Discussion

The objective of this paper was to determine whether the methodology to evaluate VA_{TMS} would influence the rate

of central fatigue and corticospinal properties. Our study design is the first to permit the comparison between the conventional method, which starts the neuromuscular evaluation a few seconds after exercise termination and allows a short relaxation between contractions, with a continuous method, which comprised fatigue assessment at exercise completion and with no recovery period between contractions. The main results are that 3–4 s between the end of

exercise and neuromuscular evaluation, together with 7 s of rest between contractions, leads to a 20% underestimation of the extent of central fatigue assessed by VA_{TMS} . Furthermore, intracortical inhibition and corticospinal excitability changes were also underestimated when short relaxation times were allowed. Therefore, this study highlights the need for measuring central parameters immediately after exercise completion to adequately appreciate fatigue amplitude and etiology. The proposed method to evaluate central fatigue has implications not only in the field of sport but also in the clinical setting, where quantifying and characterizing fatigue is relevant (Cai et al. 2014; Sacco et al. 1999; Thickbroom et al. 2006).

Maximal voluntary force changes and peripheral fatigue

The present study showed a ~75% decline in force at the end of the 2-min sustained MVC, regardless of the condition. These findings are in accordance with previous literature, as 100-s and 2-min sustained MVCs of the knee extensors have been shown to induce force decrements of about 63–79% (Doix et al. 2013; Goodall et al. 2009; Kennedy et al. 2014; Lloyd et al. 2015; Martin and Rattey 2007; Place et al. 2007; Rattey et al. 2006; Siegler and Marshall 2015).

Although there was a similar force drop during the 2-min sustained MVC, the fatiguing protocol induced a decline in MVC at POST of about 79% in CONTI and 45% in TRADI. Previous literature has documented a decrement in knee extensors MVC ranging from 10 to 33% at the POST 2-min measurement (Doix et al. 2013; Goodall et al. 2009; Kennedy et al. 2014; Lloyd et al. 2015; Martin and Rattey 2007; Place et al. 2007; Rattey et al. 2006). The lower decline in these studies is likely due to a delay in neuromuscular assessment after fatigue, which ranged from a few seconds up to 30 s (Martin and Rattey 2007). Our study corroborates this idea, as a 3–4-s rest period led to a 34% recovery in force. Neuromuscular function evaluation should, therefore, be performed as close to exercise completion as possible to avoid fatigue underestimation and fatigue etiology misinterpretations, as discussed below.

As opposed to MVC, Pt was reduced after exercise to a similar extent in both conditions. The 63% decline observed in the present study was greater compared to the decrease of 26% in Pt reported previously (Goodall et al. 2009). At the end of the fatiguing protocol, Goodall et al. (2009) performed 3 MVCs with peak twitch assessment 2 s after each MVC. The rest (even if minimal) between the end of the fatigue protocol and the first MVC, between each MVC and peripheral stimulation, and between peripheral stimulation and the following MVC might have led to some additional

recovery. In addition, the authors averaged the three peak twitch values. Due to the rest between assessments, the second and third evoked responses already benefited from some recovery (probably more than 15–20 s), which might have significantly affected the average obtained from the three assessments. Therefore, the rest between assessments and the use of the average of the three peak twitch values may help explain the lower peak twitch depression in Goodall et al. (2009) compared to our study.

The lack of difference in Pt depression between TRADI and CONTI suggests that it may take longer to recover from peripheral fatigue. In fact, Pt was still depressed 6 min after exercise in both conditions.

Cortical voluntary activation and corticospinal excitability changes

Since the linear association between SIT and voluntary force was preserved even after fatigue, we were able to extrapolate the ERT (Todd et al. 2003) in both methods. Goodall et al. (2009) documented a 17% VA_{TMS} decline after the same knee extensor fatiguing protocol. To our knowledge, only one study (Sidhu et al. 2009b) used a CONTI method to assess neuromuscular fatigue. However, in this study, the continuous method was used after a 30-s MVC, which was itself conducted after cycling exercise. Thus, it is not possible to differentiate the role of methodology vs. the role of the type of fatiguing exercise in VA_{TMS} reduction. The present study is the first one to demonstrate that a few seconds of relaxation were enough to observe a pronounced difference in VA_{TMS} between CONTI and TRADI.

Differences between conditions were not limited to VA_{TMS} . Indeed, MEP_{100} was increased at POST in CONTI only, with no differences anymore at R2. The increased MEP_{100} in CONTI is in agreement with previous literature in the upper body for sustained MVC, since MEP_{100} in these experiments was performed without rest (Gandevia et al. 1996; Kennedy et al. 2014; Szubski et al. 2007). Our findings are also consistent with previous works on the knee extensors, as enhanced MEP_{100} was reported at task failure after a repeated isometric submaximal contraction protocol (Gruet et al. 2014). Gruet et al. (2014) also reported a very rapid recovery, as MEP_{100} was not different from baseline 96 s after task failure. This quick recovery is further supported by our study, as the TRADI condition did not show any significant MEP_{100} depression at POST. Thus, it seems that after sustained maximal contractions, only a few seconds are sufficient for TMS-induced changes in corticospinal excitability during MVC to be reversed.

This is the first study documenting changes in MEP at various force levels after a 2-min sustained MVC of the knee extensors. Strikingly, MEP_{50} was depressed at POST

in CONTI and TRADI. To our knowledge, only Todd et al. (2003) assessed MEP changes at the same relative forces after a sustained contraction of the elbow flexors. In their study, however, the sustained contraction terminated when the subjects were no longer able to maintain 60% of the initial MVC. Although MEP_{50} was significantly smaller with fatigue, MEP_{100} and MEP_{75} remained unaltered. These findings are, at least partly, in agreement with the changes observed in our study. Indeed, it seems that a longer recovery time is required for MEP_{50} to return back to baseline, as there was a similar depression in both TRADI and CONTI. Todd et al. (2003) further support this idea, as the depression in MEP_{50} was still evident 8 s after exercise termination. In our study, MEP_{100} did not show any differences at POST in TRADI, suggesting a quicker recovery than MEP_{50} . Thus, Todd et al. (2003) may have failed to report any changes at MEP_{100} due to the 8-s rest allowed after exercise completion.

Before exercise, MEP amplitude usually reaches its peak near 50% MVC, decreasing thereafter in stronger contractions up to 100% MVC. Since motoneuron discharge rate increases from 50% up to 100% MVC, it is more likely that upon TMS delivery more motoneurons are in the refractory period at 100% MVC than at 50% MVC. Thus, motoneurons are less responsive to TMS at maximal voluntary contractions, which explain why MEP is smaller at 100% MVC than at 50% MVC (Todd et al. 2016).

As a result of the slowing of motoneuron discharge rate during a 2-min MVC (Bigland-Ritchie et al. 1983), it is plausible to admit that, after exercise, discharge frequency during an MVC is closer to that of a 50% MVC before exercise. Therefore, after exercise, TMS is likely to elicit a greater response at 100% MVC, because less motoneurons are in the refractory period, which may explain the exercise-induced increase in MEP_{100} observed in CONTI.

Motoneuron discharge rate at 50% MVC after a 2-min MVC is likely to be similar to the discharge rate of contractions <50% MVC before exercise. The lower corticospinal excitability at these discharge rates with lower spatial recruitment at these torque levels may explain the decrease in MEP_{50} after exercise.

Although the pronounced exercise-induced changes in absolute force might have affected motor unit recruitment (both spatial and temporal) at submaximal and maximal force intensities, which in turn might have induced different responses in MEP_{100} and MEP_{50} , this remains speculative and needs to be addressed in future research.

Intracortical inhibitions

The 2-min sustained MVC induced a lengthening in CSP_{50} in both conditions, while CSP_{100} was increased solely in CONTI. At the end of a sustained contraction, intracortical

inhibitions during MVC have been shown to be enhanced both in the upper (Gandevia et al. 1996; Szubski et al. 2007; Taylor et al. 1996) and lower limbs (Gruet et al. 2014; Mileva et al. 2012; Sidhu et al. 2009b). Since CSP_{100} returned to baseline at POST in TRADI, it appears that 3–4 s are sufficient to see a full recovery in CSP_{100} . Conversely, the lengthened CSP_{100} 6 s after exercise in Gruet et al. (2014) would suggest a more long-lasting CSP_{100} increase. The use of a repeated isometric submaximal contraction protocol by Gruet et al. (2014) differed from the sustained MVC used in our study, which may explain the contrasting results in CSP_{100} .

Most studies assessed intracortical inhibition during maximal contractions only. Nonetheless, longer CSP_{50} have been previously reported in knee extensors (Gruet et al. 2014; Sidhu et al. 2009b) and ankle flexors (Mileva et al. 2012) after fatiguing tasks. Since in Gruet et al. (2014), these changes were only noticeable right at the end of the fatiguing protocol (i.e., the first post measurement revealed a total recovery of intracortical inhibition), a 6-s relaxation appeared to be enough to observe a full recovery of intracortical inhibitions in that study. However, we observed a CSP_{50} depression both in CONTI and in TRADI. Further research is thus required to investigate if fatigue-induced changes in CSP depend on the intensity of contraction (maximal vs. submaximal) as well as on motor task intensity and/or modality (i.e., sustained vs. intermittent).

The concomitant analysis of MEPs and CSPs helps to explain the etiology of VA_{TMS} deficit. In the CONTI session, VA_{TMS} , MEP, and CSP changes occurred at POST, with a complete recovery of all parameters in R2. However, the TRADI condition revealed an attenuated but still significant VA_{TMS} decline at POST, with changes only in MEP_{50} and CSP_{50} , which confirms that central fatigue does not necessarily imply modified corticospinal excitability and inhibition (Gandevia 2001). Consequently, central mechanisms other than changes in intracortical inhibitions and excitability of motor cortical cells with corticospinal projections might explain central fatigue. Such a hypothesis has been previously suggested, as when the elbow flexors were held ischemic following a sustained MVC, MEPs recovered, while VA_{TMS} did not (Gandevia et al. 1996; Taylor et al. 2000).

In addition to the impact of the methodology used on central fatigue quantification, the different timings of NMF assessment at POST (i.e., 3–4 s after 2-min MVC in TRADI vs. no rest in CONTI) might have also contributed to the differences observed in central fatigue. The fact that the MVC was significantly different at POST between TRADI and CONTI provides a good indication that neuromuscular function can significantly recover within 3–4 s, reinforcing the need to reduce any period of recovery when measuring fatigue.

Conclusion

The method widely used to quantify neuromuscular changes following a fatiguing protocol comprises electrical and magnetic stimulation at submaximal and maximal intensities, interspersed with a few seconds of recovery. In addition, most literature that reported central and peripheral fatigue allowed a period of rest between exercise completion and fatigue evaluation, which may range from a few seconds up to 5 min. The present study showed that the methodology greatly influences fatigue quantification, as the traditional method led to fatigue underestimation of maximal voluntary activation decline and associated parameters. Thus, the results from previous research that used this methodology might not reflect the actual central fatigue caused by the fatiguing protocols and prevent the right analysis of fatigue etiology. This is especially important in protocols that use maximal sustained contractions, as in this case, the recovery of the neuromuscular system is very quick. The differences between the two methods are not expected to be as high for other types of fatiguing protocols. The continuous method used in the present study should be preferred to evaluate central fatigue induced by exercise in sport and in the clinical setting.

Acknowledgements We would like to thank Dr. John Temesi for his valuable comments and revision of the manuscript, and Douglas Doyle-Baker for the English revision.

Compliance with ethical standards

Funding This work was supported by a doctoral research grant from the University Savoie Mont Blanc.

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

Ethical approval All procedures were performed in accordance with the ethical standards of the institutional research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

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