

Sensory gating and suppression of subjective peripheral sensations during voluntary muscle contraction

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Research article

Keywords: subjective peripheral sensation, F-wave, somatosensory evoked potential, muscle contraction

Posted Date: August 6th, 2020

DOI: <https://doi.org/10.21203/rs.3.rs-26883/v2>

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Version of Record: A version of this preprint was published on October 1st, 2020. See the published version at <https://doi.org/10.1186/s12868-020-00592-2>.

Abstract

Background: During voluntary muscle contraction, sensory information induced by electrostimulation of the nerves supplying the contracting muscle is inhibited and the somatosensory evoked potentials (SEPs) amplitude decreases. This depression of sensory input during voluntary muscle contraction has been demonstrated by many studies using electrophysiological methods. However, the association between the electrophysiological response of the sensory system during sustained muscle contraction and subjective peripheral sensation (SPS) is still unclear. The aim of this study was to investigate changes in spinal excitability, SEPs, and SPS during voluntary muscle contraction.

Results: The appearance rate of the F-wave was significantly higher during muscle contraction than during rest, whereas no significant difference was observed in F-wave latency between muscle contraction and rest. Furthermore, the P25 amplitude of SEPs was significantly lower during muscle contraction than during rest, whereas the N20 amplitude of SEPs exhibited no significant differences. The SPS was significantly lower during muscle contraction than during rest.

Conclusions: We conclude that sensory gating, which is found in the P25 component of SEPs during muscle contraction, is one of the neurophysiological mechanisms underlying the suppression of SPS.

Background

Human motor control results from integration of the sensory and motor systems, which are regulated by the relationship between the upper and lower centers, i.e., the brain and spinal cord, respectively. When peripheral nerves are electrically stimulated, a direct motor response (M-wave) occurs in the muscle due to anterograde conduction from the stimulated area. At the same time, the ascending afferent input is projected to the somatosensory cerebral cortex via the spinal cord, and somatosensory evoked potentials (SEPs) in the cortex can be recorded. During voluntary muscle contraction, sensory information induced by electrostimulation of the nerves supplying the contracting muscle is inhibited and the SEP amplitude decreases [1-4]. The suppression of the sensory potential is known as "gating". With respect to gating during voluntary movement, previous studies reported that the amount of gating depends on the difficulty of the movement [5], and that gating is observed not only during motion but also pre-motion [6]. Furthermore, suppression of sensory input has already occurred when the spinal cord (a midpoint between the cerebral cortex and periphery) receives input [7]. Therefore, it seems that the main functional role of gating is to eliminate unnecessary sensory information during the execution of purposeful voluntary movements.

F-waves are muscle action potentials recorded when electrostimulation to a peripheral nerve causes retrograde conduction to the axon of an α -motoneuron, and anterograde conduction occurs once again through the automatic firing of a motoneuron in the anterior horn of the spinal cord. The F-wave appearance rate indicates the number of motor units participating in the waveform [8] and naturally varies at a certain width even in the resting state. This natural variation suggests that sensory input

influences the effect the central nervous system has on α -motoneurons. H-reflex is also a non-invasive technique that is used to assess responses of motor units activated by the afferent pathway [9]. Previous study using the H-reflex have reported that the excitability of spinal α -motoneurons that innervate active muscles increases during voluntary contraction in the muscle groups of the upper and lower limbs [10]. Furthermore, it is known that the excitability of spinal α -motoneurons increases during remote [11] and contralateral muscle contraction [12]. Therefore, the facilitation of spinal α -motoneurons during muscle contraction may occur not only locally but also systemically.

The depression of sensory input during voluntary muscle contraction has been demonstrated by many studies using electrophysiological methods for animals and humans. It has been reported that the P25 amplitude was attenuated, although the N20 amplitude remained stable during voluntary muscle contraction tasks [1, 13] and reaction tasks [14]. Furthermore, these responses were observed after the tasks were completed [15, 16]. Thus, on the basis of the results of a previous report on the change in each component of SEPs under various conditions, the functional significance of attenuation of sensory input during motion is certain. However, the association between the electrophysiological response of the sensory system during exercise and subjective peripheral sensation (SPS) is still unclear. For example, it is often observed that athletes who have pain in the motor system (e.g., mild ankle sprains or muscle soreness) hardly recognize the pain sensation after a few minutes of exercise. However, the neurophysiological aspect of this phenomenon has not been studied to date. Therefore, we hypothesized that the change in electrophysiological response in sensory input affects the subjective peripheral sensation during motor output. The aim of this study was to investigate the changes in spinal excitability, SEPs, and SPS during voluntary muscle contraction.

Results

The appearance rate of the F-wave was significantly higher during muscle contraction ($83.4 \pm 12.3\%$) than during rest ($34.2 \pm 16.0\%$; $t(12) = 9.638$, $P < 0.001$; Fig. 1B). No significant differences in F-wave latency were observed between muscle contraction (26.7 ± 1.7 ms) and rest (27.2 ± 1.8 ms; $t(12) = 1.257$, $P = 0.23$; Fig. 1C). The F-wave amplitude was significantly higher during muscle contraction ($6.6 \pm 4.0\%$) than during rest ($2.7 \pm 1.3\%$; $t(12) = 3.661$, $P = 0.003$; Fig. 1D). For the N20 amplitude of SEPs, no significant differences were observed between muscle contraction (2.5 ± 1.1 μ V) and rest (3.0 ± 1.3 μ V; $t(12) = 2.083$, $P = 0.06$; Fig. 2B). The P25 amplitude of SEPs was significantly lower during muscle contraction (4.23 ± 1.70 μ V) than during rest (5.35 ± 2.10 μ V; $t(12) = 2.982$, $P = 0.001$; Fig. 2C). The accuracy rate of 20 filament stimulations was significantly lower during muscle contraction ($57.7 \pm 22.9\%$) than during rest ($78.5 \pm 14.2\%$; $t(12) = 4.158$, $P = 0.001$; Fig. 3).

Discussion

In this study, we aimed to investigate changes in spinal excitability, SEPs, and SPS during voluntary muscle contraction. We found that the F-wave appearance rate increased during the isometric pinching task and that this natural variation increases during the pinching task. It is possible that the increase in

the F-wave appearance rate during the pinching task is caused by suppression of the corticospinal tract, which converges on the spinal anterior motor nerve, and inhibitory systems such as the higher control system.

The potential source of N20 is considered to be the 3b area, which is interpreted simply as the stage at which sensory stimulation has reached the primary sensory cortex via the thalamus [17]. Furthermore, P25 is considered a component derived from a higher level than the 3b area [18]. This suggests that the submaximal isometric pinching force used in our current study suppresses the somatosensory input of a higher level than the 3b area. Previous studies on SEP gating during voluntary movement have reported that gating does not occur in the components corresponding to N20 [19, 20]. For these reasons, although the electrophysiological input that is projected to the primary somatosensory area is the same for a given amount of physical stimulation (regardless of the presence or absence of the motor task), this electrophysiological input is suppressed during the subsequent more complex process of information processing.

In this study, the accuracy rate for cutaneous stimulation at the sensory threshold was used as an index of SPS, and this parameter was reduced by submaximal isometric muscle contraction. This is considered the result of presynaptic inhibition of peripheral sensation during motor output. During muscle contraction, some sensory inputs derived from skin sensory receptors are already suppressed within peripheral nerves, i.e., before reaching the neural circuits of the brain and spinal cord [21]. The numerous neural circuits in the cerebral cortex and spinal cord play various important roles during motor behavior. The results of this study suggest that there is a mechanism that facilitates the necessary circuits by inhibiting unnecessary circuits that are related to motion during motor output.

Transcranial focused ultrasound of the primary sensory cortex (S1), which enhances performance in sensory discrimination tasks [22], decreased short latency (P27-N20 and N33-P27) SEP components. Considering these results, we hypothesized that the pulsed acoustic pressure wave to S1 locally shifts the balance of excitation and inhibition by acting on mechanically sensitive components of the brain (cell membranes, ion channels, and synaptic vesicle cycles). Based on this hypothesis, we determined that the pulsed acoustic pressure waves dampen excitation or increase local interneuron firing, leading to attenuation of SEP amplitudes and resulting in an improvement in the cortical representation of tactile stimuli. Furthermore, S1 is already activated by the input from primary motor cortex (M1) before it receives the input from sensory receptors during voluntary movement [23]. In the present study as well, it can be predicted that S1 was activated with the execution of the muscle contraction task. However, in this study, the input from the peripheral sensory receptor to S1 is hardly necessary in the isometric muscle contraction task at the submaximal strength. Therefore, we considered that the inhibitory adjustments for S1 activated by the input from M1 had occurred. The inhibitory effect on S1 seems to contribute to the improvement in the precision of motor system adjustments. This hypothesis helps explain the decrease in the P25 amplitude in the present study.

Furthermore, it is possible that the threshold of cutaneous surface sensation was increased by the motor output. Most of the decrease in the accuracy rate observed in this study was due to the increased inability to recognize the sensory stimulation. This indicates the need for a test in which participants could recognize sensory information correctly. There was a certain degree of freedom regarding the stimulation site and filament stimulation intensity, as the sensory stimulation was applied manually in this study. This implies that the sensory threshold during motor output does not increase uniformly at all sites but varies according to each site or receptor.

This study demonstrates the neurophysiological aspect of changes in subjective sensation during muscle contraction and that subjective sensations in humans are closely related to the attenuation of sensory potential.

Conclusions

The present study has demonstrated that increasing spinal excitability, which innervates active muscle groups, causes depression of P25 components in the operating limb area that governs muscle group actions. As such, we have clarified the mechanisms underlying SPS of operating limbs. We conclude that sensory gating, which is found in the P25 component of SEP during muscle contraction, is one of the neurophysiological mechanisms responsible for suppressing SPS.

Methods

Participants

Participants comprised 13 healthy adult males with no history of neurophysiological diseases. Mean (\pm SD) age was 27 ± 11 years, height was 167 ± 2 cm, and weight was 66 ± 6 kg). This study was conducted in accordance with the Declaration of Helsinki. Informed consent to participate in the study was obtained from the participants verbally and in writing. This study was approved by the Ethical Review Board of the Kibi International University (No. 18-37).

Electrical stimulation

A square wave pulse lasting 0.2 ms was generated in the median nerve in the carpal area on the right side (i.e., ipsilateral to the pinching hand). Using a surface electrostimulation apparatus (NM-420S, Nihon Kohden, Japan), the stimulation electrodes were placed 20 mm apart, with the cathode proximal and the anode distal. The stimulation electrodes were secured using a fixation band to regulate their pressure. The minimum intensity of electrostimulation to induce an M-wave was confirmed, and the stimulation region was set appropriately.

F-wave recording and analysis

A surface electromyogram (EMG) during electrostimulation was recorded in the right abductor pollicis brevis using Ag/AgCl bipolar electrodes (5 mm diameter, 20 mm interelectrode distance; Nihon Kohden, Japan) on the same side as the electrostimulation. The recording electrode was attached onto the muscle belly of the abductor pollicis brevis, and the reference electrode onto the first proximal phalanx; both electrodes were fixed with surgical tape. The attachment site for the recording electrode was wiped with alcohol and preprocessed using sandpaper to create an interelectrode resistance < 5 k Ω . An EMG/evoked potential testing device (Neuropack MEB-9404, Nihon Kohden, Japan) was used, and F-wave waveforms were recorded using a band-pass filter of 1.5–3 kHz and a sampling frequency of 10 kHz. The electrostimulation intensity was set to 1.2x, which obtained a maximum M-wave. The stimulation frequency was set to 1 Hz and was carried out for approximately 1 min. The F-wave analysis used the three factors appearance rate, latency, and amplitude F/M ratio during the second half (30 s) of electrostimulation. To determine the percentage of the appearance rate, all identifiable resulting waveforms (total n = 30) on the monitor (500 μ V/D) were targeted. For the F-wave amplitude, the average value of the peak-to-peak amplitude for the F-waves was expressed as a ratio compared to the maximum M-wave amplitude. Latency was the average time from electrostimulation until F-wave initiation.

SEP recordings and analysis

Based on the international 10-20 system, SEPs were recorded from the somatosensory area of the right upper arm (C3': 2 cm posterior from C3) on the stimulation side. The reference electrode was placed at the point Fz. Electrodes were attached onto the skin surface using conductive paste. Interelectrode resistance was set to < 5 k Ω . The electrostimulation intensity was fixed at just above the motor threshold with a repetition rate of 3 Hz. An EMG/evoked potential testing device (Neuropack MEB-9404, Nihon Kohden, Japan) was used, and SEP waveforms were recorded using a band-pass filter of 20 Hz–10 kHz and a sampling frequency of 10 kHz, and 200 responses were averaged. SEP waveforms were evaluated for 100 ms, and evaluations were performed at the time of electrostimulation and 100 ms after stimulation. Epochs with artifacts owing to eye movement or blinking ($>\pm 6 \mu$ V from baseline) were automatically excluded before averaging. A plate electrode was used to record the evoked electroencephalogram (Ag/AgCl electrode, NE-132B (Φ 10 mm), Nihon Kohden, Japan). The amplitudes from baseline to N20 and the N20-P25 peak-to-peak amplitude, which are early components after electrostimulation, were analyzed for SEPs. The amplitude of each component was measured from preceding peaks.

Subjective peripheral sensation (SPS)

Prior to the experiment, the SPS threshold on the dorsal surface of the right hand was measured using a Semmes-Weinstein monofilament test (SAKAI Medical Co., Ltd., Tokyo, Japan). After establishing their peripheral sensory thresholds, participants reported the presence or absence of peripheral cutaneous stimulation during the SPS measurements. Monofilament stimulation was carried out using gradually thicker filaments, starting from thin filaments of 0.008 g. Confirmation tests were repeated approximately five times for each intensity, and the filament thickness that could be sensed correctly at a rate of

approximately 100% was defined as the SPS threshold. The experimenter lowered the filament vertically onto the hand, removed it, and returned it to its original position within 1 s. The stimulation interval of the filament was random and carried out a total of 20 times. The participants were instructed to give verbal cues when they sensed filament stimulation; afterwards, their accuracy rate was calculated. All measurements were performed by the same experimenter. The filament stimulation site was marked using a marker, so that deviation of a large stimulation site owing to experimenter error did not occur.

Testing procedure

The participants were seated in a chair with both arms placed on armrests with the eyes open to perform a pinching task with the right hand while keeping the arm in a neutral position. Before the task, maximum voluntary isometric force (MVIF) was measured, from which the target tension level was calculated. The participants held the pinch force meter with the thumb and index finger and squeezed with maximal force for 5 seconds in the experimental position. The maximum value of force exertion at 5 seconds was defined as MVIF. The target value was displayed visually on a computer placed 1 m in front of the participant. After sufficient rest of more than 15 minutes, participants maintained 30% MVIF for approximately 2 min. Tension was not exerted in the rest condition. Each measurement item and each condition were randomly set and measured for each subject.

Statistics

All values are shown as the mean \pm standard deviation. Comparisons were analyzed using paired t-tests; statistical significance levels were set at $< 5\%$ ($P < 0.05$) with GraphPad Prism Ver 8.3.1 for Macintosh (GraphPad Software, San Diego, CA, USA).

Abbreviations

SEP, somatosensory evoked potential; SPS, subjective peripheral sensation; EMG, electromyogram; MVIF, maximum voluntary isometric force

Declarations

Ethics approval and consent to participate

This study was conducted in accordance with the Declaration of Helsinki. Informed consent to participate in the study was obtained from the participants verbally and in writing. This study was approved by the Ethical Review Board of the Kibi International University (No. 18–37).

Consent for publication

Not applicable

Availability of data and materials

The datasets supporting the conclusions of this article are included within the article (and its additional files).

Competing interests

The authors declare that they have no conflicts of interest.

Funding

This work was supported by the JSPS KAKENHI Grant Number JP 18K17909.

Authors' contributions

TT, HY, and SO conceived and designed the research. KS and MM conducted the experiments. TT and KS analyzed the data. TT wrote the manuscript. All authors read and approved the final manuscript.

Acknowledgments

Not applicable

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Figures

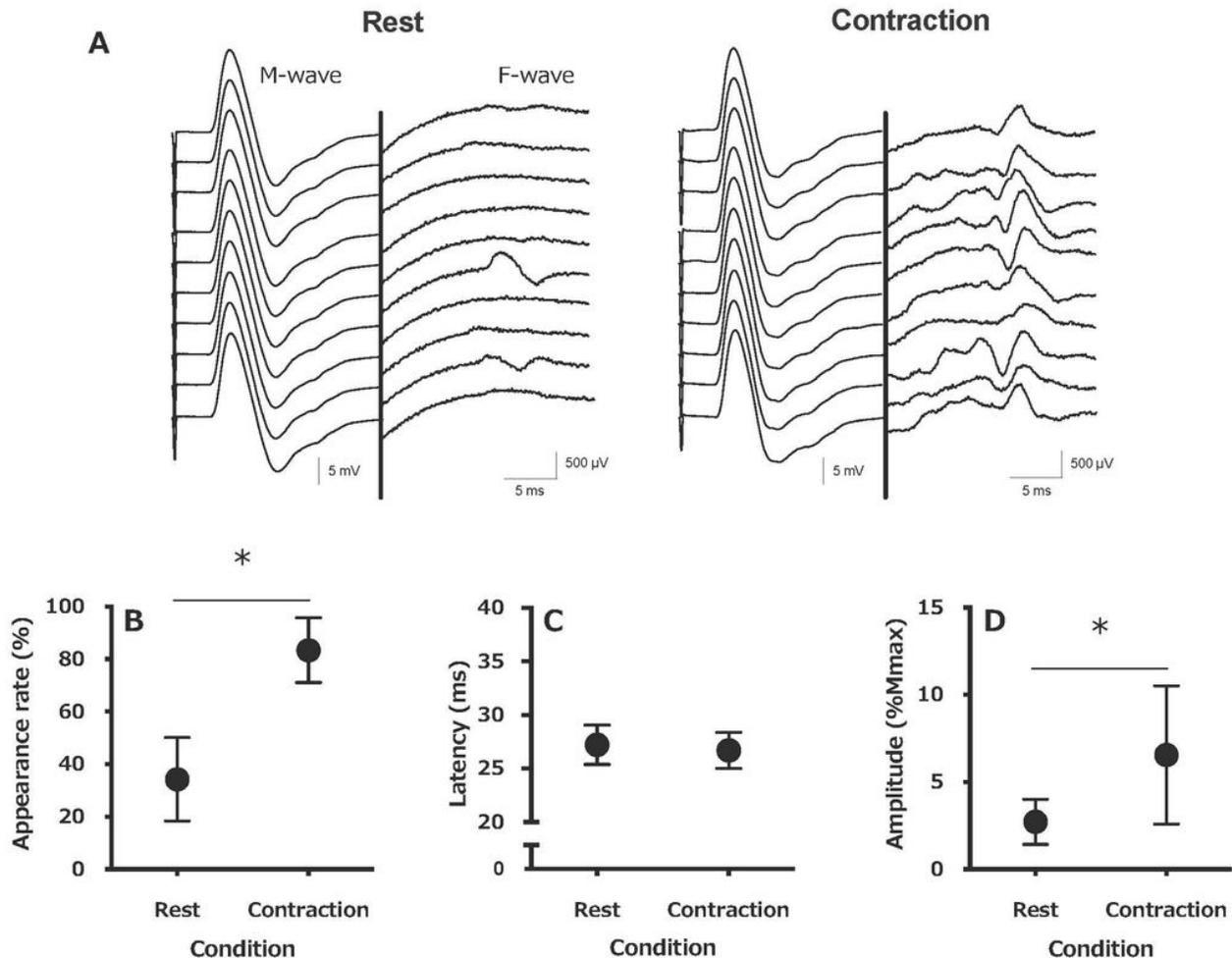


Figure 1

A) A typical example of an induced electromyogram waveform from a single participant in both conditions (for 10 stimulations). The waveforms of both conditions are M-waves on the left side and F-waves on the right side of the central thick line. The display range of the F-wave is 10x that of the M-wave. The changes in average F-wave appearance rate (B), latency (C), and amplitude (D) of all participants in both conditions are displayed. * P < 0.05

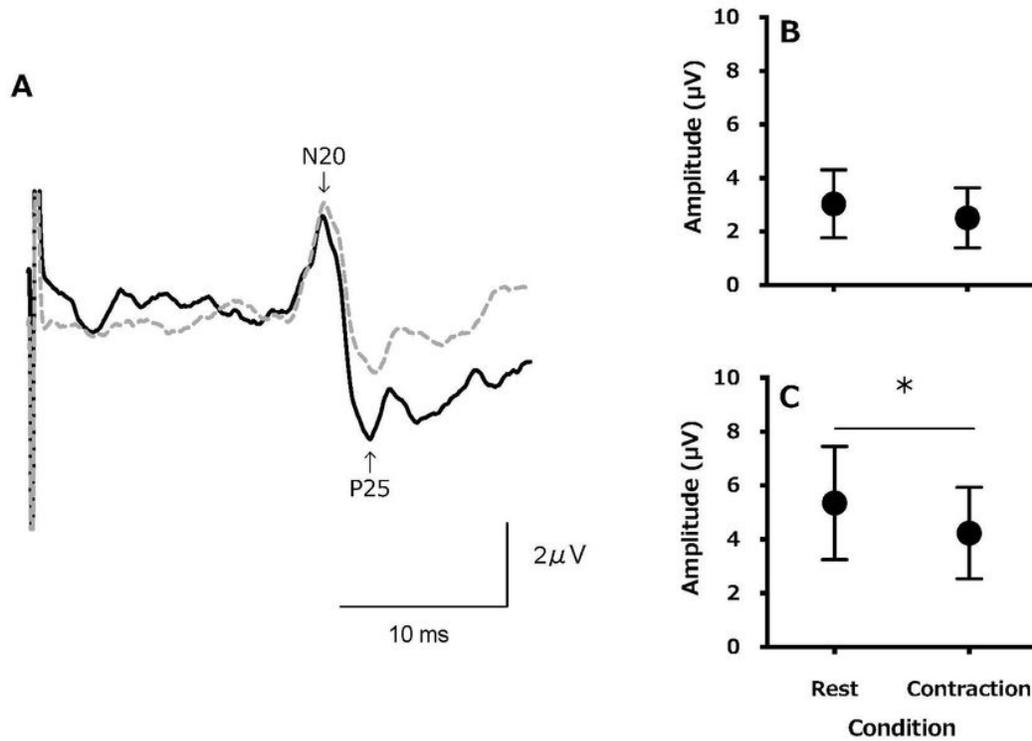


Figure 2

A) A typical example of a somatosensory evoked potential (SEP) waveform from a single participant (for 10 stimulations) during rest (solid black line) and contraction (gray dashed line). B-C) The changes in average SEP amplitude for N20 (B) and P25 (C) of all participants in both conditions. * $P < 0.05$

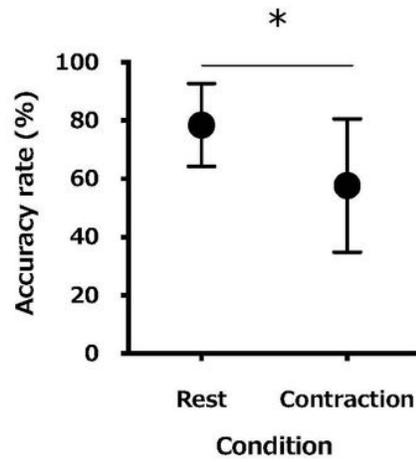


Figure 3

The changes in the average accuracy rate of all participants in both conditions. * P < 0.05