Title: Gymnasts' ability for general motor imagery evaluated by bioelectric sensorimotor

rhythms

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Abstract

Introduction: Previous psychological studies using questionnaires have consistently reported that athletes have superior motor imagery ability, both for sports-specific movements and general movements. However, regarding general motor imagery, no physiological studies have demonstrated differences in neural activity between athletes and non-athletes. Gymnasts may be a suitable population for investigating this issue because they are likely to have particularly superior motor imagery ability due to their frequent usage of motor imagery as part of daily practice. **Purpose**: The purpose of the present study was to examine differences in bioelectric sensorimotor rhythms during kinesthetic motor imagery of general movements between gymnasts and non-gymnasts. **Methods**: Healthy young participants (16 gymnasts and 16 non-gymnasts) performed repeated motor execution and motor imagery of general movements (wrist dorsiflexion and shoulder abduction of the dominant hand). Scalp electroencephalogram (EEG) was recorded over the contralateral sensorimotor cortex. During motor execution and motor imagery, sensorimotor EEG power is known to decrease in the α- (8–15 Hz) and β-bands (16–35 Hz), referred to as event-related desynchronization (ERD). We calculated the maximal peak of ERD both in the α- (αERDmax) and β-bands (βERDmax) as a measure of changes in corticospinal excitability. **Results**: ERD magnitude during motor imagery was significantly greater in gymnasts, who subjectively evaluated their motor imagery as being more vivid. In particular, αERDmax was greater in gymnasts

compared with non-gymnasts for both motor imagery tasks, whereas βERDmax was greater

in gymnasts only for shoulder abduction imagery. **Conclusion:** We observed gymnasts'

signature of flexibly modulating sensorimotor rhythm with no movement, which may be the

basis of their superior general motor imagery ability.

Keywords: athletes, kinesthetic motor imagery (KMI), electroencephalogram (EEG), event-

related desynchronization (ERD), the Kinesthetic and Visual Imagery Questionnaire (KVIQ)

Introduction

Motor imagery is regularly used by athletes to improve performance (1,2). Previous studies have shown that performing motor imagery in training improves performance in various tasks, including sequence learning (3,4), jump height (5) and free-throw shooting (6). To explain these performance gains, several neuroscience studies have provided evidence that motor imagery activates some neural substrates in common with actual movement, including the primary motor cortex, supplementary motor area, and inferior parietal lobe (7–10), as well as inducing neural plasticity in these areas (11–13).

In the field of sports psychology, several cross-sectional questionnaire studies have consistently reported that athletes have superior motor imagery ability compared with non-athletes (14–16). Furthermore, previous studies have indicated that athletes can perform motor imagery more vividly than non-athletes, not only for specialized movements in their own sports but also for general movements such as raising the arm and jumping (15,17). The findings of these studies suggest that the neural activity underlying motor imagery ability may differ between athletes and non-athletes, not only for sports-specific imagery but also for general motor imagery.

However, in the field of applied physiology, to the best of our knowledge, no studies have demonstrated differences in neural activity during general motor imagery between athletes and non-athletes, although some studies using electroencephalogram (EEG)

(17), magnetoencephalography (MEG) (18), functional magnetic resonance imaging (fMRI)

(19) and transcranial magnetic stimulation (TMS) (20) have reported signature neural activity

patterns during sports-specific motor imagery in athletes. For instance, when tennis players

imagined movements specifically related to tennis, their corticospinal excitability became

higher than that of non-athletes, whereas such a difference between athletes and non-athletes

was not observed when they imagined other general movements (20). Thus, there is currently

a gap in findings between psychological and physiological studies regarding differences in

motor imagery ability, particularly for general movements between athletes and non-athletes.

To clarify this issue, the present study investigated differences in general motor

imagery ability between gymnasts and healthy adults (i.e., non-gymnasts) from both

psychological and physiological points of view. As a psychological indicator, we evaluated

the subjective vividness of motor imagery using The Kinesthetic and Visual Imagery

Questionnaire (KVIQ-20) (21). As a physiological indicator, we evaluated sensorimotor

rhythms using EEG. Event-related desynchronization (ERD) is a measure of decreases in

power of the EEG sensorimotor rhythm within the α- and β-bands from the resting-state to

the motor execution or motor imagery state (22,23), which is known to reflect increased

neuronal excitability in the corticospinal system (24). We chose gymnasts as a population of

athletes for the following reasons: 1) gymnasts perform motor imagery frequently as a part of

their daily practices because of the high risk of serious injury in their performance; 2)

gymnasts were assumed to have higher motor imagery abilities than athletes engaged in other

sports, because it has been reported that motor imagery is more vivid in athletes engaged in

individual and/or non-contact sports compared with athletes engaged in team and/or contact

sports (25); 3) to the best of our knowledge, no previous studies have measured neural

activity during motor imagery in gymnasts.

Methods

Ethical Approval

This study was conducted in accordance with the Declaration of Helsinki. All

experimental protocols and procedures were approved by the Research Ethics Committee in

Shonan Fujisawa Campus, Keio University (Approval Number 167). The examiners provided

a detailed explanation of the purpose, experimental procedures, potential benefits, and risks

involved. After receiving all of the relevant information, participants provided written

informed consent before participating in the experiment.

Participants

We recruited 16 gymnasts (11 men, 5 women, aged 18–24 years) and 16 healthy

adults (8 men, 8 women, aged 19–22 years) as a non-gymnast group. All participants were

right-handed. All gymnasts had been practicing at least for 7 years (range: 7–16 years) and

had participated in an all-Japan intercollegiate gymnastic championship at least once. Note

that two gymnasts were members of the Japanese national gymnastics team. The non-

gymnasts group had no experience of gymnastic training. None of the participants had

experienced any neurological and musculoskeletal disorders.

Psychological assessments

Procedures

Motor imagery ability was tested using a psychological questionnaire translated into

Japanese: The Kinesthetic and Visual Imagery Questionnaire (KVIQ-20) (21,26). Briefly, the

KVIQ-20 tests how vividly a person is able to imagine their own movements subjectively,

using two types of motor imagery: kinesthetic motor imagery (KMI) and visual motor

imagery (VMI). KMI involves imagining the feeling when we perform actual motor tasks.

Conversely, VMI involves imagining to see ourselves or the field of vision where we perform

the tasks (21,27). Participants sat comfortably in a chair next to the examiner and watched the

examiner's example once. Then, they actually performed the exercise, followed by VMI or

KMI of the exercise they had just performed. Participants were then asked to evaluate the

vividness of the motor imagery on a five-point ordinal scale (the more vivid the motor

imagery, the higher the scale score). This procedure was repeated for 10 different simple

exercises: neck flexion/extension, shoulder elevation, forward shoulder flexion, elbow

flexion/extension, thumb-fingers opposition, forward trunk flexion, knee extension, hip

abduction, foot tapping, and foot external rotation.

Analyses

We evaluated the vividness of motor imagery by summing all KVIQ scores for KMI

and VMI, respectively. If a participant could imagine their movements perfectly, the score

was 50 points.

Physiological assessments

Recordings

Scalp EEG signals were recorded with eight passive Ag/AgCl electrodes around the

sensorimotor area related to right upper limbs (Cz, C1, C3, C5, FC1, FC3, CP1, and CP3) in

accord with the extended international 10-20 system. Electrodes with a diameter of 18 mm

were mounted on an electrode cap (g.GAMMAcap 1027; Guger Technologies, Graz,

Austria). Reference and ground electrodes were placed on the right and left earlobes,

respectively. Surface electromyogram (EMG) signals were recorded from the right deltoid

muscle (DEL) and the right extensor carpi radials muscle (ECR). Two passive Ag/AgCl

electrodes with a diameter of 10 mm were placed over each muscle belly with inter-electrode

distances of 20 mm. All EEG and EMG signals were amplified and bandpass-filtered (EEG,

0.5–200 Hz; EMG, 5–1000 Hz) using a linked biosignal recording system (g.BSamp 0201a;

Guger Technologies, Graz, Austria). All analog EEG and EMG signals were converted to

digital signals at a sample rate of 1000 Hz using an AD converter with 16-bit resolution (NI

USB-6259, National Instruments, Austin, TX, United States) that was controlled by data-

logger software originally designed using MATLAB software (The MathWorks, Inc., Antic,

MA, United States).

Procedures

Following the psychological assessment, we performed physiological EEG and

EMG measurements. The participants sat comfortably in the seat of the dynamometer device.

A computer monitor for visual feedback was placed 2 m in front of participants' eyes. First,

resting-state EEG was recorded over 60 s. Participants relaxed and fixated their eyes on a

cross (+) displayed at the center of the monitor during recording. Participants then performed

several practice trials of maximal voluntary contractions (MVCs) of wrist dorsiflexion and

shoulder abduction. After these movements were practiced, participants performed MVC

once each for wrist dorsiflexion and shoulder abduction. When performing each MVC, EMG

activity of the contracting muscle was recorded. The EMG signals were full-wave rectified.

We found a 0.5-s period of stable force exertion during MVC and calculated the integrated

EMG value (iEMGmax) in this period. In the following experiment, 20% of this iEMGmax

value was used as a target value for visual feedback.

The physiological data recordings during motor execution and KMI were performed

after several practice trials. Visual feedback was presented on the screen, with a red cursor to

represent muscle contraction level as a relative value in %iEMGmax and a vertical blue line

to represent a target value. Furthermore, instructions for each phase, including "Rest",

"Relax", "Ready", "Contraction", or "Imagery", were displayed on the monitor. In the wrist

dorsiflexion task, participants' dominant hand was positioned on the armrest and fixed by a

belt with the palm down. In the shoulder abduction task, the dominant upper limb was

lowered to the side of the body while bending the elbow lightly, and the arm was fixed by a

belt. In both the wrist dorsiflexion and shoulder abduction tasks, participants performed

repeated motor execution and KMI according to the procedure used in our previous study

(28).

The experimental paradigm is shown in Figure 1. In detail, each trial was started

from the rest phase and the word "Rest" was displayed on the monitor for 7 s. During the rest

phase, participants were able to adjust their posture freely and/or blink their eyes strongly.

After the rest phase, the word "Relax" was displayed on the monitor for 3 s. During the relax

phase, participants were instructed to relax as much as possible, without performing any

movement. The word "Ready" was then displayed for 3 s, accompanied by a short sound

presented every second. During the ready phase, participants prepared for the next

instruction. After the ready phase, the word "Contraction" was displayed for 5 s. During the

contraction phase, participants performed isometric voluntary contraction (wrist dorsiflexion

or shoulder abduction) at 20% of iEMGmax by their dominant hand. In the contraction phase,

participants were instructed to contract their muscle so that the cursor could follow the target

line as accurately as possible. After the contraction phase, the word "Relax" was displayed

for 2 s. In this relax phase, participants were instructed to relax as much as possible, without

any movement. After the rest, relax, and ready phases, the imagery phase was started, and

participants performed KMI of the preceding contraction for 5 s with their eyes open, and

without any movement. During the imagery phase, we checked that no EMG activity

occurred. When the imagery phase finished, the relax phase was presented again for 2 s. This

flow was conducted for each trial, including motor execution and KMI, and five trials were

repeated within each set. Six sets were performed for each task. Thus, a total 30 trials were

performed for both wrist flexion and shoulder abduction tasks. We set the wrist dorsiflexion

and the shoulder abduction task in a randomized order across participants. The duration of the

set interval was longer than 2 min, to provide sufficient rest for participants.

Analyses

To remove noise arising from the electric power, the EEG and EMG signals were

notch-filtered at 50 Hz. The EEG signals over C1 and C3 were derived with a four-neighbor

Laplacian spatial filter. For example, in the case of C3, the EEG signal over C3 was

subtracted by an average of C1, C5, FC3 and CP3. The Laplacian derivation method is

known to strongly emphasize cortical activity originating below the electrode of interest (29).

If Laplacian-derived EEG included potentials exceeded 50 μV, we considered the trial to

contain an artifact, and excluded the data from future analyses. Additionally, visual

inspection was performed to reject additional artifacts missed by the automatic inspection.

ERDs during motor execution and KMI of each task were calculated as follows.

After separating the data into motor execution and KMI periods, we extracted the 30 1 s data

windows in the same period from the data for each trial. Then, fast Fourier transformation

was performed using Welch's method for the data (window length, 1 s; window function,

Hanning-window; overlap, 0), and the power spectrum densities (PSDs) of the EEG signal

were calculated. This process was repeated by sliding the 1 s data window in 50 ms steps.

The ERDs were calculated using the following equation:

$$ERD(f,t) = \frac{R(f) - A(f,t)}{R(f)} \times 100\%, \qquad (\text{eq. 1})$$

where A is the EEG PSDs at time t, frequency f, and R is the mean PSDs of the baseline

period (last 1 s in the relax phase). This equation indicates that the positively greater the ERD

value, the larger the decrease in EEG PSD during motor execution or KMI compared with the

relax phase. Because the most reactive frequency band of ERD was slightly different across

participants (30), we determined an electrode and the 3 Hz-frequency width showing the

largest ERD in each of the α-band (8–15 Hz) and β-band (16–35 Hz) during motor execution.

Because it has been suggested that the functional roles played by ERD differ between the α-

and β-bands (31,32), we analyzed ERDs from these two frequency bands separately. The

magnitude of ERD in the α-band (αERDmax) and β-band (βERDmax) were measured by

calculating the peak value of ERD for motor execution and KMI of each task, respectively

(28,33).

To compare the features of EEG during the relax phase in the task with continuous

resting-state for a prolonged period, we also analyzed the α-band or β-band PSDs for both

data sets. For the relax phase EEG, the final 1 s periods in the relax phase, which were used

as the baseline periods for ERD analyses, were extracted from all trials, and combined to

create a 60 s relax phase EEG signal. For the resting-state EEG, a continuous 60 s period

with few artefacts was extracted. We then calculated the ratio of the sum of EEG power

within the α-band or β-band PSD to that of the entire frequency range (4–50 Hz) (named,

EEGα-PSD and EEGβ-PSD) for both data sets, and compared these values between relax

phase EEG during tasks and resting-state EEG.

Statistical analyses

Two-side unpaired t-tests were performed on VMI and KMI scores of KVIQ

between groups (non-gymnasts vs. gymnasts), to confirm differences in the subjective

vividness of motor imagery between them. To test differences in αERDmax and βERDmax

during motor execution or KMI, we performed two-way mixed-model analysis of variance

(ANOVA) between participant groups (gymnasts and non-gymnasts) and tasks (wrist

dorsiflexion and shoulder abduction). If the interaction was significant, we performed a two-

sided unpaired t-test for groups (gymnasts vs. non-gymnasts) and a two-sided paired t-test for

tasks (wrist dorsiflexion vs. shoulder abduction). To test differences in the EEGα-PSD or

EEGβ-PSD between resting-state EEG and relax phase EEG during tasks, we also performed

two-way ANOVA between participant groups (gymnasts and non-gymnasts) and conditions

(resting-sate and relax phase). The p-values of 0.05 were used to indicate statistical

significance. All statistical analyses were performed using SPSS statistics software (IBM

SPSS Statistics 25, IBM developerWorks, Tokyo, Japan).

Results

KVIQ

Figure 2 shows group data (mean ± S.D.) for KVIQ scores obtained from VMI and

KMI tasks between gymnasts and non-gymnasts. The KVIQ scores were significantly greater

in gymnasts, both in VMI (gymnasts, 42.68 ± 6.22; non-gymnasts, 35.44 ± 8.73, $p = 0.011$)

(Figure 2A) and KMI (gymnasts, 43.50 ± 6.78; non-gymnasts, 35.94 ± 8.24, $p = 0.008$)

(Figure 2B). The KVIQ results indicate that gymnasts subjectively evaluated how vividly

they could imagine their own movements.

ERD magnitude

259   Typical examples of EEG signals, EEG time-frequency maps and ERD time courses

260   during wrist dorsiflexion ME from a non-gymnast and a gymnast are shown in Figure 3A and

261   3B, respectively. From these time-frequency maps, a decrease of EEG power can be observed

262   around 12 Hz and 22 Hz in the contraction phase (0 to 5 s) compared with the relax phase (−6

263   to −3 s) in both participants when performing motor execution.

264   Figure 3C shows group data for αERDmax during wrist dorsiflexion and shoulder

abduction motor execution. An ANOVA on αERDmax during motor execution revealed no

significant effects of group ($F_{1,30} = 0.209$, $p = 0.651$) and task ($F_{1,30} = 0.831$, $p = 0.369$),

while a significant interaction was obtained ($F_{1,30} = 4.654$, $p = 0.0391$). An unpaired t-test for

group revealed no significant difference in αERDmax between gymnasts and non-gymnasts

for wrist dorsiflexion (gymnasts, 64.41 ± 20.96; non-gymnasts, 65.58 ± 13.97, $p = 0.855$) and

shoulder abduction task (gymnasts, 70.13 ± 21.07; non-gymnasts, 63.26 ± 16.77, $p = 0.316$).

A paired t-test for task revealed no significant difference in αERDmax between wrist

dorsiflexion and shoulder abduction execution both for non-gymnasts (wrist dorsiflexion,

65.58 ± 13.97; shoulder abduction, 63.26 ± 16.77, p = 0.303) and gymnasts (wrist

dorsiflexion, 64.41 ± 20.96; shoulder abduction, 70.13 ± 21.07, p = 0.078). Figure 3D shows

group data for βERDmax during wrist dorsiflexion and shoulder abduction motor execution.

An ANOVA on the βERDmax during motor execution with groups and task revealed no

significant effects of group ($F_{1,30}$ = 0.571, p = 0.456) and task ($F_{1,30}$ = 1.249, p = 0.273), and

interaction ($F_{1,30}$ = 1.815, p = 0.188). The results revealed no effects of sports experience and

body part on ERD magnitude during motor execution.

280   Typical examples of EEG signals, EEG time-frequency maps and ERD time courses

281   during wrist dorsiflexion KMI from a non-gymnast and a gymnast are shown in Figure 4A

282   and 4B, respectively. From the time-frequency maps for a non-gymnast participant, we did

283   not observe clear ERD in the imagery phase (0 to 5 s) compared with the relax phase (−6 to

284   −3 s) in both the α-band and β-band. (Figure 4A). Conversely, clear ERD can be observed in

285   the time-frequency-map for the gymnast participant around 12 Hz and 22 Hz (Figure 4B).

286   Figure 4C shows group data for the αERDmax during wrist dorsiflexion and

shoulder abduction KMI. An ANOVA on αERDmax during KMI showed significant effects

of group ($F_{1,30}$ = 5.437, p = 0.027) and task ($F_{1,30}$ = 10.975, p = 0.002). No significant

interaction effect ($F_{1,30}$ = 0.266, p = 0.610) was observed. Figure 4C shows group data for

βERDmax during wrist dorsiflexion and shoulder abduction KMI. An ANOVA on

βERDmax during KMI showed no significant effects of group ($F_{1,30}$ = 0.876, p = 0.357), but

significant effects of task ($F_{1,30}$ = 8.019, p = 0.008) and a significant interaction ($F_{1,30}$ =

7.421, p = 0.010). An unpaired t-test for group revealed a significant difference in βERDmax

between gymnasts and non-gymnasts for the shoulder abduction task (gymnasts, 46.08 ±

18.09; non-gymnasts, 32.27 ± 17.84, p = 0.021) but not for the wrist dorsiflexion task

(gymnasts, 46.42 ± 22.76; non-gymnasts, 49.56 ± 17.84, p = 0.666). A paired t-test for task

revealed a significant difference in βERDmax between the wrist dorsiflexion and shoulder

abduction imagery conditions for non-gymnasts (wrist dorsiflexion, 49.56 ± 17.84; shoulder

abduction, 32.27 ± 17.84, p = 0.002) but not for gymnasts (wrist dorsiflexion, 46.42 ± 22.76;

shoulder abduction, 46.08 ± 18.09, p = 0.938). These results indicated that sports experience

affected ERD magnitude during KMI.

Comparison of EEGα-PSD and EEGβ-PSD between resting-state EEG and relax phase

EEG during tasks

Figure 5A shows group data for the EEGα-PSD in resting-state EEG and relax phase

EEG during the tasks. An ANOVA examining EEGα-PSD data revealed no significant

effects of group ($F_{1,30}$ = 0.486, p = 0.491) or condition ($F_{1,30}$ = 4.135, p = 0.051); however, a

significant interaction ($F_{1,30}$ = 6.382, p = 0.017) was observed. An unpaired t-test for group

revealed no significant difference in EEGα-PSD during resting-state EEG (gymnasts, 0.473 ±

0.155; non-gymnasts, 0.482 ± 0.166, p = 0.878) and that during the relax phase (gymnasts,

0.482 ± 0.155; non-gymnasts, 0.403 ± 0.125, p = 0.878; p = 0.122) between gymnasts and

non-gymnasts. In non-gymnasts, a paired t-test revealed significant differences in the EEGα-

PSD between conditions (resting-state, 0.482 ± 0.166; relax phase, 0.403 ± 0.125, p = 0.008).

However, in gymnasts, no significant differences in EEGα-PSD were observed between

conditions (resting-state, 0.473 ± 0.155; relax phase, 0.481 ± 0.155, p = 0.715). Figure 5B

shows group data for EEGβ-PSD. An ANOVA on EEGβ-PSD showed no significant effects

of group ($F_{1,30}$ = 0.106, p = 0.747) and condition ($F_{1,30}$ = 1.787, p = 0.191), and no significant

interaction ($F_{1,30}$ = 0.063, p = 0.804). These results indicate that, in non-gymnasts, the EEGα-

PSD was smaller in the relax phase EEG than during resting-state EEG, while such a

difference was not observed in gymnasts. Conversely, EEGβ-PSD did not differ between

resting-state EEG and relax phase EEG in both gymnasts and non-gymnasts.

Discussion

The purpose of the present study was to clarify differences in bioelectric

sensorimotor rhythm during general motor imagery (i.e., non-sports specific motor imagery)

between gymnasts and non-gymnasts. The results revealed that, when required to repeatedly

switch between relaxing and motor execution or KMI, the ERD magnitude during general

KMI was significantly greater in gymnasts, who subjectively evaluated their motor imagery

as more vivid, while no difference between groups was observed during motor execution. In

particular, the ERD magnitude in the α-band was greater in gymnasts compared with non-

gymnasts, both in wrist dorsiflexion and shoulder abduction KMIs, whereas the ERD

magnitude in the β-band was greater in gymnasts only in shoulder abduction KMI.

Difference in general KMI ability between gymnasts and non-gymnasts

We evaluated ERD as a physiological indicator of motor imagery ability in the

present study, because it is considered to reflect changes in corticospinal excitability (24) and

is associated with subjective vividness of motor imagery measured by KVIQ (28). It should

be noted that the present results revealed greater ERD magnitude during general KMI in

gymnasts than in non-gymnasts, although differences in neural activity between athletes and

non-athletes have not been reported in general motor imagery in previous studies using EEG

(17), MEG (18), TMS (20) or fMRI (19). This may be related to the fact that gymnasts

perform motor imagery frequently as a part of their daily practice to reduce the risk of serious

injury in their practice. Furthermore, a previous psychological study showed that the

vividness of general motor imagery was higher in athletes engaged in individual and/or non-

contact sports compared with athletes engaged in team and/or contact sports (25). Thus, as

gymnasts have superior motor imagery ability among athletes, they may provide a

particularly suitable population for highlighting differences in neural activity during general

motor imagery compared with non-athletes.

It is possible that the present task protocol, in which participants performed KMI

following motor execution repeatedly in the order of seconds, led to the current finding of

greater ERD magnitude in gymnasts. In psychological questionnaires, the conventional

procedure for measuring motor imagery ability is to examine participants while they perform

motor execution then motor imagery in one trial, and subjectively evaluate the vividness of

the motor imagery of the preceding movement (21). In physiological experiments, however,

the conventional procedure involves evaluating neural activity while participants perform

only motor imagery (17–20). Thus, there has been a methodological gap in the approach for

examining of motor imagery between psychological questionnaire studies and experimental

physiological studies. In response to direct questioning in the current study, gymnasts

reported that they usually perform actual movements and KMIs alternately in their daily

practice. Therefore, the method for measuring motor imagery in the present physiological

experiment was designed in accord with the procedure of psychological questionnaire

measurement. The current physiological findings may have been due to differences between

gymnasts and non-gymnasts in the ability to flexibly modulate corticospinal excitability when

imagining their own movements, by referring to the actual movement.

As shown in equation (1), we were able to confirm that the ERD was determined by

both the degree of synchronization during the relax phase ($R(f)$) and the degree of

desynchronization during KMI ($A(f, t)$). As shown in Fig. 5, first, gymnasts could return their

sensorimotor α-rhythm during the relax phase in the task to the same power level as during

the resting-state for 60 s, whereas non-gymnasts could not. Thus, gymnasts appeared to be

good at relaxing deeply by making their sensorimotor rhythms more synchronized within a

short period of time. However, higher EEGα-PSD during the relax phase does not appear to

be the only factor involved in gymnasts' greater ERD magnitude in the α-band. As shown in

Fig. 2, second, differences in αERDmax between groups were not observed in motor

execution but were found in KMI. Thus, gymnasts also appeared to be good at increasing

corticospinal excitability by making their sensorimotor rhythms more desynchronized, even

in KMI. Overall, the present results demonstrate that gymnasts have the ability to generate a

clear contrast in the state of the sensorimotor cortex, when required to repeatedly switch

across relaxing, motor execution and motor imagery conditions. On the basis of the current

findings, we believe that the ability to modulate the brain state without any movement is a

core aspect of superior motor imagery ability in gymnasts.

Difference in functional role of ERD between α- and β-band

383   Interestingly, the present study demonstrated different results between ERD

magnitude in the α-band and β-band. Several previous studies reported that functional roles

played by the sensorimotor rhythm are different between frequency bands. During actual

muscle contraction with weak-to-moderate intensity, the sensorimotor rhythm is known to be

coherent with EMG activity only in the β-band, with no significant coherence in the α-band

(34–36). When focusing on oscillatory power itself, the EEG spectral power in the

sensorimotor area contralateral to the contracted/imagined limb was decreased in both the α-

and β-bands, while that in task-irrelevant cortical regions was increased in the α-band

(22,37), but not in the β-band (31). During KMI, the ERD magnitude was increased by

increasing task demand in the β-band, but not in the α-band (32,38). Taken together, these

findings suggest that the functional roles of sensorimotor rhythms for movement/imagery

should be distinguished between the α-band and β-band.

First, αERDmax was larger in gymnasts than non-gymnasts during both KMI tasks

(i.e., wrist dorsiflexion and shoulder abduction). In task-relevant cortical regions, neural

populations are assumed to be disinhibited by the ERD of the sensorimotor area in the α-

band, which would allow reallocation of computational resources (32). However, task-

irrelevant cortical regions are assumed to be inhibited by enhancing their α-oscillations (23).

As gymnasts are required to perform skilled movements successively, they are trained to

quickly switch their attention across their body parts by facilitating task-relevant regions and

inhibiting task-irrelevant regions. The present results regarding ERD in the α-band would

reflect such an ability of gymnasts.

Second, βERDmax was larger in gymnasts only during shoulder abduction KMI, but

not during wrist dorsiflexion KMI. This task-specificity in βERDmax may be caused by ERD

in the β-band playing a role in the calculation of specific motor commands. In general, wrist

movement is used frequently in daily life, which makes it easy for most people to imagine

wrist dorsiflexion. Thus, βERDmax would not be differed between gymnasts and non-

gymnasts in the wrist dorsiflexion task. However, as isometric shoulder abduction is a

movement rarely used in daily life, it may be difficult for most people to imagine this

movement. Conversely, gymnasts are well-trained to move their upper limbs, including the

shoulder joints, both dynamically (i.e., giant swing) and statically (i.e., handstand and rings).

Therefore, it would be easy for gymnasts to imagine shoulder abduction because they are

skilled at adjusting the movement parameters of their shoulder joints. We assume that ERD in

the β-band is an indicator for how precisely a person can imagine their own movement

kinesthetically.

Limitations

In the present study, only upper limb movements (i.e., wrist dorsiflexion and

shoulder abduction) were examined. Gymnasts use their upper limb muscles specifically as

anti-gravity muscles for postural non-gymnastic movements, such as handstand and pommel

horse. This usage of the upper limbs is unique relative to the movements of non-gymnasts.

The uniqueness of gymnasts' upper limb usage may lead to superior motor imagery ability

regarding upper limb movements. Thus, we cannot clearly predict whether similar results

would be obtained when performing similar experiments for other body parts. However, the

KVIQ results demonstrated that gymnasts tended to show higher scores for all movements. In

addition, we found that most gymnasts performed motor imagery of various body parts in

their daily practice. On the basis of these findings, we speculate that gymnasts have superior

motor imagery ability irrespective of body parts, although confirming this possibility will

require further investigation.

The present study is the first to observe differences in physiological indices between

athletes and non-athletes. This means that the present study can bridge the gap between

psychology and physiology studies regarding differences in general motor imagery ability

between athletes and non-athletes. However, because only gymnasts participated in this

study, it is unclear whether the present results are specific to gymnasts or apply generally to

athletes performing any sports. Because differences in general motor imagery ability would

be expected among athletes, further investigation is needed to elucidate sports-specific

differences in ability for general movement imagery. In any case, the present study indicated

the importance of comparing corticospinal excitability measured by ERD for evaluating

motor imagery ability. In future studies, imagery training using bioelectrical signals may

provide a useful tool for improving the motor imagery ability of athletes.

Conclusion

The present study demonstrated that, during general KMI, the corticospinal

excitability measured by ERD magnitude was significantly greater in gymnasts compared

with non-gymnasts. These results are consistent with higher subjective vividness in gymnasts

measured using the KVIQ psychological questionnaire. The observed signature of flexibly

modulating sensorimotor rhythm with no movement would be the basis of their superior

general motor imagery ability in gymnasts.

Author Contributions

H.S. and J.U. conceptualized and designed the study, interpreted data, wrote the manuscript,

and acquired funding. H.S. acquired and analyzed data. J.U. supervised the study.

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We declare that the results of the study are presented clearly, honestly, and without

fabrication, falsification, or inappropriate data manipulation. The results of the present study

do not constitute endorsement by American College of Sports Medicine (ACSM).

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References

1.   Cumming J, Hall C. Deliberate imagery practice: the development of imagery skills in

competitive athletes. *J Sports Sci*. 2002;20(2):137–45.

2.   Mizuguchi N, Nakata H, Uchida Y, Kanosue K. Motor imagery and sport

performance. *J Phys Fit Sport Med*. 2012;1(1):103–11.

3.   Sobierajewicz J, Przekoracka-Krawczyk A, Jaśkowski W, Verwey WB, van der Lubbe

R. The influence of motor imagery on the learning of a fine hand motor skill. *Exp*

Brain Res. 2017;235(1):305–20.

4.   Lebon F, Horn U, Domin M, Lotze M. Motor imagery training: Kinesthetic imagery

strategy and inferior parietal fMRI activation. *Hum Brain Mapp*. 2018;39(4):1805–13.

5.   Battaglia C, D'Artibale E, Fiorilli G, et al. Use of video observation and motor

imagery on jumping performance in national rhythmic gymnastics athletes. *Hum Mov*

Sci. 2014;38:225–34.

6.   Peynircioğlu ZF, Thompson JLW, Tanielian TB, Tanielian TB. Improvement

strategies in free-throw shooting and grip-strength tasks. *J Gen Psychol*.

2000;127(2):145–56.

7.   Decety J. Neural mechanisms subserving the perception of human actions. *Trends*

Cogn Sci. 1999;3(5):172–8.

8.   Guillot A, Collet C, Nguyen VA, Malouin F, Richards C, Doyon J. Brain activity

489   during visual versus kinesthetic imagery: An fMRI study. *Hum Brain Mapp*.

490   2009;30(7):2157–72.

491   9.   Hanakawa T. Functional Properties of Brain Areas Associated With Motor Execution

492   and Imagery. *J Neurophysiol*. 2002;89(2):989–1002.

493   10.   Zabicki A, De Haas B, Zentgraf K, Stark R, Munzert J, Krüger B. Imagined and

494   executed actions in the human motor system: Testing neural similarity between

495   execution and imagery of actions with a multivariate approach. *Cereb Cortex*.

496   2017;27(9):4523–36.

497   11.   Ietswaart M, Johnston M, Dijkerman HC, et al. Mental practice with motor imagery in

498   stroke recovery: Randomized controlled trial of efficacy. *Brain*. 2011;134(5):1373–86.

499   12.   Pascual-Leone A, Dang N, Cohen LG, Brasil-Neto JP, Cammarota A, Hallett M.

500   Modulation of muscle responses evoked by transcranial magnetic stimulation during

501   the acquisition of new fine motor skills. *J Neurophysiol*. 1995;74(3):1037–45.

502   13.   Ruffino C, Papaxanthis C, Lebon F. Neural plasticity during motor learning with

503   motor imagery practice: Review and perspectives. *Neuroscience*. 2017;341:61–78.

504   14.   Di Corrado D, Guarnera M, Quartiroli A. Vividness and transformation of mental

505   images in karate and ballet. *Percept Mot Skills*. 2014;119(3):764–73.

15.   Isaac AR, Marks DF. Individual differences in mental imagery experience:

Developmental changes and specialization. *Br J Psychol*. 1994;85(4):479–500.

16.   Jansen P, Lehmann J. Mental rotation performance in soccer players and gymnasts in

an object-based mental rotation task. *Adv Cogn Psychol*. 2013;9(2):92–8.

17.   Di Nota PM, Chartrand JM, Levkov GR, Montefusco-Siegmund R, DeSouza JFX.

Experience-dependent modulation of alpha and beta during action observation and

motor imagery. *BMC Neurosci*. 2017;18(1):28.

18.   Kraeutner SN, McWhinney SR, Solomon JP, Dithurbide L, Boe SG. Experience

modulates motor imagery-based brain activity. *Eur J Neurosci*. 2018;47(10):1221–9.

19.   Wei G, Luo J. Sport expert's motor imagery: Functional imaging of professional motor

skills and simple motor skills. *Brain Res*. 2010;1341:52–62.

20.   Fourkas AD, Bonavolont V, Avenanti A, Aglioti SM. Kinesthetic imagery and tool-

specific modulation of corticospinal representations in expert tennis players. *Cereb*

Cortex. 2008;18(10):2382–90.

21.   Malouin F, Richards CL, Jackson PL, Lafleur MF, Durand A, Julien D. The

Kinesthetic and Visual Imagery Questionnaire (KVIQ) for Assessing Motor Imagery

in Persons with Physical Disabilities: A Reliability and Construct Validity Study. *J*

Neurol Phys Ther. 2007;31(1):20–9.

22. Pfurtscheller G, Lopes da Silva FH. Event-related EEG/MEG synchronization and

desynchronization: basic principles. *Clin Neurophysiol*. 1999;110(11):1842–57.

23. Pfurtscheller G, Neuper C, Andrew C, Edlinger G. Foot and hand area mu rhythms. *Int

J Psychophysiol*. 1997;26(1–3):121–35.

24. Takemi M, Masakado Y, Liu M, Ushiba J. Event-related desynchronization reflects

downregulation of intracortical inhibition in human primary motor cortex. *J

Neurophysiol*. 2013;110(5):1158–66.

25. Di Corrado D, Guarnera M, Vitali F, Quartiroli A, Coco M. Imagery ability of elite

level athletes from individual vs. Team and contact vs. No-contact sports. *PeerJ*.

2019;7:1–14.

26. Nakano H, Kodama T, Ukai K, Kawahara S, Horikawa S, Murata S. Reliability and

validity of the Japanese version of the kinesthetic and visual imagery questionnaire

(KVIQ). *Brain Sci*. 2018;8(5).

27. Hall C, Pongrac J, Buckholz E. The measurement of imagery ability. *Hum Mov Sci*.

1985;4(2):107–18.

28. Toriyama H, Ushiba J, Ushiyama J. Subjective Vividness of Kinesthetic Motor

Imagery Is Associated With the Similarity in Magnitude of Sensorimotor Event-

Related Desynchronization Between Motor Execution and Motor Imagery. *Front Hum

Neurosci*. 2018;12(July):1–10.

29.   McFarland DJ, McCane LM, David S V., Wolpaw JR. Spatial filter selection for EEG-

based communication. *Electroencephalogr Clin Neurophysiol*. 1997;103(3):386–94.

30.   Pfurtscheller G, Neuper C. Future prospects of ERD/ERS in the context of brain–

computer interface (BCI) developments. *Prog Brain Res*. 2006;159:433–7.

31.   Stolk A, Brinkman L, Vansteensel MJ, et al. Electrocorticographic dissociation of

alpha and beta rhythmic activity in the human sensorimotor system. *Elife*. 2019;8:1–

24.

32.   Brinkman L, Stolk A, Dijkerman HC, de Lange FP, Toni I. Distinct Roles for Alpha-

and Beta-Band Oscillations during Mental Simulation of Goal-Directed Actions. *J*

Neurosci. 2014;34(44):14783–92.

33.   Takemi M, Masakado Y, Liu M, Ushiba J. Is event-related desynchronization a

biomarker representing corticospinal excitability? *Proc Annu Int Conf IEEE Eng Med*

Biol Soc EMBS. 2013;281–4.

34.   Ushiyama J, Takahashi Y, Ushiba J. Muscle dependency of corticomuscular coherence

in upper and lower limb muscles and training-related alterations in ballet dancers and

weightlifters. *J Appl Physiol*. 2010;109(4):1086–95.

35.   Ushiyama J, Yamada J, Liu M, Ushiba J. Individual difference in β-band

corticomuscular coherence and its relation to force steadiness during isometric

voluntary ankle dorsiflexion in healthy humans. *Clin Neurophysiol*. 2017;128(2):303–

11.

36.   Suzuki R, Ushiyama J. Context-Dependent Modulation of Corticomuscular Coherence

in a Series of Motor Initiation and Maintenance of Voluntary Contractions. *Cereb*

Cortex Commun. 2020;1(1):1–14.

37.   Pfurtscheller G. Event-related synchronization (ERS): an electrophysiological

correlate of cortical areas at rest. *Electroencephalogr Clin Neurophysiol*.

1992;83(1):62–9.

38.   van Elk. Semantics in the motor system: motor-cortical beta oscillations reflect

semantic knowledge of end-postures for object use. *Front Hum Neurosci*.

2010;4(February):1–12.

Figures

Figure 1. Experimental paradigm of physiological experiment. Participants performed

isometric contraction in the contraction phase and performed motor imagery of the same

movement in the imagery phase. The diagram shows the flow in each trial, which was

repeated five times within each set. Six sets were performed for each of the wrist dorsiflexion

and shoulder abduction tasks.

Figure 2. Results of psychological experiment. Group data (Mean ± S.D.) for visual motor

imagery (VMI) (A) and kinesthetic motor imagery (KMI) scores (B) obtained from the

Kinesthetic and Visual Imagery Questionnaire (KVIQ) are shown for both groups. The gray

bar represents data for non-gymnasts, while the black bar represents data for gymnasts. *P <

0.05.

Figure 3. Results from physiological experiments for motor execution. Typical time courses

of single trial electroencephalogram (EEG), time-frequency map, α-band event-related

desynchronization (ERD), and β-band in wrist dorsiflexion motor execution are shown for

non-gymnast (A) and gymnast (B) participants. Note that participants performed wrist

dorsiflexion motor execution from 0 s to 5 s. Group data (Mean ± S.D.) for the maximal peak

of ERD both in the α- (αERDmax) (C) and β-bands (βERDmax) (D) during motor execution

are shown across groups and tasks. The gray bar represents data for non-gymnasts, while the

black bar represents data for gymnasts. No significant differences were observed across

groups and tasks.

Figure 4. Results from physiological experiments for KMI. Typical time courses of single

trial EEG, time-frequency map, α-band ERD, and β-band in wrist dorsiflexion KMI are

shown for non-gymnast (A) and gymnast (B) participants. Note that participants performed

wrist dorsiflexion KMI from 0 s to 5 s. Group data (Mean ± S.D.) for αERDmax (C) and

βERDmax (D) during KMI are shown across groups and tasks. The gray bar represents data

for non-gymnasts, while the black bar represents data for gymnasts. *P < 0.05.

608   Figure 5. Results of the ratio of the α-band and β-band power spectrum densities. Group data

609   (Mean ± S.D.) for the ratio of the α-band (A) and β-band power spectrum densities (PSD)

610   (B) are shown for both groups. The gray bar represents data for non-gymnasts, while the

611   black bar represents data for gymnasts. *P < 0.05.