Comparing electromyography, accelerometry, and visual inspection to assess the resting motor threshold for transcranial magnetic stimulation

Gautier Hamoline[1]*, Antoine E Lunardi [1,] Marcos Moreno-Verdú [1], Baptiste M Waltzing [1], Elise E Van Caenegem [1], Siobhan MacAteer[1], Robert M Hardwick [1],

[1] Brain, Action, and Skill Laboratory, Institute of Neurosciences, UCLouvain, Belgium

*Address for Correspondence: Gautier Hamoline, UCLouvain, Institute of NeuroScience, Cognition & Systems Division, Avenue Mounier 54, 1200 Bruxelles, Belgium. gautier.hamoline@uclouvain.be

Abstract

Introduction: Electromyography (EMG) remains the gold standard for estimating the

Resting Motor Threshold (RMT) Transcranial Magnetic Stimulation (TMS) studies,

but its cost and limited accessibility often lead researchers to use visual inspection

(VIS). However, VIS may introduce variability and systematic bias. Accelerometry

(ACC) offers a cost-effective, objective alternative to capture TMS-evoked responses.

Objective: To compare the RMT as estimated using EMG, ACC, and VIS.

Methods: Five participants underwent TMS while EMG, ACC, and video recordings

were collected. Separately, 64 observers judged hand movement in videos to estimate

RMT via VIS. RMTs were compared across the three methods using Bayesian model

comparison, Bland-Altman analyses, and Intraclass Correlation Coefficients (ICCs).

Results: RMTs estimated via EMG were lower than those obtained using either ACC

or VIS. Compared to EMG, VIS tended to overestimate RMT (mean bias = 5.23%,

95%CI = [1.00–11.00]), while ACC and VIS estimates were more closely aligned (mean

bias = 0.43%, 95%CI = [–4.00 – 5.00]). ICC (2,1) values indicated moderate reliability

for VIS vs EMG (mean = 0.580, 95%CI = [0.389 – 0.748]), and good-to-excellent

reliability for VIS vs ACC (mean = 0.845). However, bootstrapped 95% confidence

intervals identified significant variability in the estimates provided by visual

inspection, ranging from +1 to +11 for VIS vs EMG, but as low as -4 to +5 for VIS vs

ACC.

Conclusions: EMG remains the most sensitive technique for estimating the RMT, but

when EMG is not feasible, accelerometery provides a quantifiable, more objective, and

less variable alternative than visual inspection.

Keywords

Transcranial Magnetic Stimulation, Electromyography, Accelerometry, Visual Inspection

Highlights

• Visual inspection introduces observer-related variability in RMT estimation

• Visual inspection & accelerometry systematically overestimate RMT

compared to EMG

• EMG remains the most sensitive method for estimating RMT

• Accelerometry is an objective alternative to visual inspection if EMG is

unavailable

1. Introduction

Research on the effects of Transcranial Magnetic Stimulation (TMS) has largely focused on the primary motor cortex of the human brain (1). When an area of the primary motor cortex linked to a target muscle is stimulated, it can activate the corticospinal pathway leading to muscle contractions, and potentially induce movements in that muscle. By employing electromyography (EMG), the resulting muscle activity from this stimulation can be recorded as a 'Motor Evoked Potential' or MEP (2). Among the features of the MEP, the most commonly analysed is its amplitude, often measured from peak to peak in microvolts (μV). Measuring MEP amplitudes is central to defining the intensity of TMS that is delivered in most TMS studies, which is typically defined based on the Resting Motor Threshold (RMT) — the minimal stimulator output necessary to produce MEPs with an amplitude of ≥50μV in at least 5 out of 10 trials.

Although much of the early research on TMS has focused on fundamental motor responses, over time, TMS has garnered significant attention for its wide-ranging applications across diverse fields. In neurology, TMS has been used to examine conditions such as multiple sclerosis (3–7), stroke (8,9), epilepsy (10,11), Parkinson's disease (12–15), dystonia (16–18) and myopathy (19–21). The application of repetitive TMS has demonstrated substantial promise in treating psychiatric disorders, leveraging its ability to modulate neural activity in targeted brain regions. Conditions such as major depressive disorder (MDD) (22–24), obsessive-compulsive disorder (OCD) (23,25–27), anxiety(28), post-traumatic stress disorder (PTSD)(29,30) and schizophrenia (31,32) have all benefited from TMS interventions.

Additionally, its efficacy extends to behavioral disorders, including addiction (33),

attention-deficit/hyperactivity disorder (ADHD) (34), and eating disorders (35).

Beyond these possible therapeutic applications, TMS has also proven to be a valuable

tool in cognitive research, to improve our understanding of cognitive processes such

as mental imagery, decision making and action preparation (36). Thus, while TMS

was initially developed to probe motor system function through the measurement of

MEPs, it has since evolved into a versatile tool with broad applications.

Regardless of the exact field of study, EMG remains the standard for assessing TMS

responses. However, EMG presents several drawbacks. The initial cost of EMG

equipment such as apparatus and amplifiers, combined with recurring expenses for

consumables like electrodes, electrode gel, and cleaning alcohol, makes it a relatively

expensive technique (37). Additionally, precise electrode placement requires

understanding of muscle anatomy and physiology (38). Preparatory steps, including

skin cleaning, hair removal, and verifying the clarity of the signal, all add to the time

required to create optimal recordings using this method (39). Despite its widespread

use among research teams investigating motor control and neurophysiology, many

laboratories employ EMG solely to evaluate the RMT (40–42). However, due to the

costs and preparation time associated with EMG, some groups instead measure the

"visually determined" RMT. This approach defines RMT as the lowest stimulator

output that generates a visible movement (usually of the hand) in at least 5 out of 10

trials (43). While more accessible and less expensive than using EMG, the visual

inspection method is inherently subjective (43) and therefore subject to inter- and

intra-individual variability; moreover, it focuses on identifying the presence/absence

of movement without providing other quantifiable measures (e.g. the speed or

magnitude of the induced movement). Furthermore, previous reports provide

conflicting accounts of the relationship between measures collected using visual

inspection and EMG, with research indicating the visually defined threshold can differ

from that identified using EMG by between –10 to +2% of maximal stimulator output

(43,44). This is notable as under- or over-estimating the RMT could in turn lead to the

application of stimulation intensities that are too low to be effective, or that may

exceed TMS safety guidelines (45). Together, these limitations underscore the need

to investigate alternative methods to EMG that are both cost-effective and efficient

but also allow precise quantification of TMS evoked responses.

Accelerometry provides a promising alternative to visual inspection and a potentially

more accessible method than EMG in assessing TMS responses. Specifically,

accelerometry is more cost-effective than EMG, requiring smaller initial outlays and

incurring lower recurring expenses. Accelerometry also enables researchers to

objectively quantify TMS responses, and is therefore likely to be more reliable than

the subjective assessments provided by visual inspection. Past studies have used

accelerometry to examine how TMS-evoked movement direction and magnitude

change with training (46–48). Recently, our laboratory has explored using

accelerometry as a tool to measure different characteristics of TMS evoked responses;

although accelerometry tends to overestimate the RMT, comparisons of

measurements of MEP amplitudes and accelerometer jerk readings were highly

correlated, demonstrating this measurement tool shows promising potential (49).

In this paper, the aim is to investigate and compare the measurement of the Resting

Motor Threshold (RMT) using three different methods: electromyography (EMG),

accelerometry (ACC), and visual inspection (VIS). By evaluating the accuracy,

reliability, and practical application of each technique, the present study will

determine their respective strengths and limitations in capturing the RMT.

2. Materials and Methods

2.1. General Design

The study leveraged a mixed-model design to examine data from a total sample of 105 participants. The final sample after application of exclusion criteria (see below) included 69 participants. This final sample comprised two groups of participants.

Data from Group 1 provided direct recordings of responses to TMS. Data collection involved recording data from five participants using three methods: electromyography, accelerometry, and video recordings of the hand during TMS (i.e. recordings demonstrating whether the intensity of TMS was sufficient to produce movement or not).

Data from Group 2 then focused on the visual determination of the resting motor threshold by individual observers. Participants in this second group observed video recordings taken from the first group, providing responses to indicate whether they observed a movement in each trial in response to TMS.

2.2. Group 1: Recording of responses to TMS (EMG, Accelerometry, and Video)

2.2.1. Participants

Five participants (2 females, 3 males), all right-handed, with an average age of 24.2 years (ranging from 21 to 29 years, SD = ± 2.31) took part in this part of the study. The safety screening checklist recommended by Lefaucheur et al. (50) was administered prior to stimulation to confirm participant eligibility and ensure safe

application for the use of TMS. The study was approved by the Ethics Committee

Saint-Luc Hospital, UCLouvain (2020/30JUL/389 - DNMT-TMS – amendment 2), and

each participant provided written informed consent prior to the experiment.

Participants also consented to video recordings of their hands to be presented to

other participants (see below).

2.2.2. Transcranial Magnetic Stimulation (TMS)

TMS was administered using a monophasic Magstim 200^2 stimulator with a figure-of-

eight coil (2 x Ø70 mm) directed posteriorly and toward the ipsilateral side with a

random interval of 4 to 6 seconds between each stimulation. To ensure precise

targeting, participants wore a head tracker (NT-103) and the Visor2 neuronavigation

system (ANT Neuro, Netherlands) was used, which provided real-time guidance for

accurate coil positioning.

2.2.3. Electromyography (EMG)

EMG was utilized to record Motor Evoked Potentials (MEPs) from the First Dorsal

Interosseus (FDI) muscle of the index finger on the dominant hand. Before placing the

electrodes, the skin was cleaned with an alcohol solution and shaved if necessary to

remove hair and enhance conductivity. Two self-adhesive, pre-gelled bipolar surface

electrodes (Blue Sensor N, Ambu®, Denmark) were positioned on the muscle body

and its distal insertion. A reference electrode was placed on the styloid process of the

ulna. EMG signals were sampled at 2 kHz with an online digital notch filter (50Hz),

bandpass filtered at 20–450Hz, and amplified using a D360 8-Channel Patient

Amplifier (Digitimer®, England).

2.2.4. Accelerometer Recordings

A 8791A250 K-Shear® Miniature Triaxial Accelerometer (Kistler, USA) was secured

to the nail of the index finger using micropore adhesive tape (Fig. 1). The

accelerometer captured movement in the x, y, and z axes, enabling a complete

reconstruction of the motion. The data were collected using a 4-Channel

PiezoSmart® (TEDS) Power Supply/Signal Conditioner (Kistler, USA) and recorded

as the finger's acceleration (m/s^2). Prior to analysis, the data were low-pass filtered

offline using a 20 Hz 4th order Butterworth filter.

2.2.5. Video recordings

Video recordings of the experiment were captured using a GoPro Hero 8 Black, set to

record at 1080p resolution and 120 frames per second. The camera was mounted on

a tripod, positioned to film the hand from above and from the contralateral side (see

Fig. 1). This setup was chosen to simulate the perspective of an experimenter during

a typical experimental session and offered a clear view of the hand.

2.2.6. Protocol Design

Participants sat on a chair with the palmar side of the dominant hand resting face

down on the table, and were able to make adjustments until they found a comfortable

position that they could maintain throughout testing. The corticomotor area over the

left hemisphere of the brain corresponding to the right first dorsal interosseus muscle

was identified, and the motor hotspot was determined. RMT was assessed using EMG,

defined as the lowest intensity of maximal stimulator output (MSO) to produce a MEP

of ≥50μV in at least 5 out of 10 trials. Stimulation was randomly applied at intensities

ranging from -5%MSO to +20%MSO of the EMG-defined RMT. An external

experimenter (A.L.) visually monitored the participant's hand and signaled to the

experimenter delivering TMS (G.H.) if an intensity of TMS that produced a movement

on each trial (i.e. 10/10 trials) was observed, and future intensities of stimulation

were applied only below this intensity. This procedure determined the highest

stimulation intensity around RMT that elicited visible movements (range: +9% to

+19% MSO).

2.2.7.   Data Processing

EMG and accelerometer data were recorded using a Power1401–3A system and

Signal software (both from Cambridge Electronic Design, England). The data were

processed using custom MATLAB scripts.

EMG signal analysis focused on identifying the 'peak-to-peak' amplitude of the Motor

Evoked Potential (MEP), which was calculated as the difference between the highest

positive and negative peaks of the response in a post-stimulation window of 200ms.

Accelerometer data, capturing movements in the x, y, and z axes, were combined

using the Cartesian formula (absolute acceleration = $\sqrt{(x^2 + y^2 + z^2)}$) to generate a

single value representing the total acceleration of the sensor (i.e. a measure sensitive

to any movement of the finger in response to TMS, analogous to visual inspection).

The derivative of the acceleration was then calculated to determine the 'jerk' (m/s^3),

with the absolute peak jerk being the primary signal of interest. Jerk is particularly

useful for analysing rapid or transient movements, which are common in TMS-

induced responses.

The processed accelerometer data were also used to identify the RMT as determined

by accelerometry. The RMT was defined as the intensity of the maximal stimulator

output (MSO) at which at least 5 out of 10 trials exhibited a peak jerk (m/s^3)

exceeding the 95% confidence interval calculated from the 200 ms preceding the TMS

stimulation (49)(i.e., a baseline, motionless period for each participant).

2.3. Group 2: Video observations to determine the visual RMT

2.3.1. Participants

Initially, 100 participants were recruited through a combination of in-person sessions (20 participants) and the Prolific online platform (80 participants). However, after data verification and attention checks were considered (see below), a total of 64 participants were retained for analysis. These 64 participants (33 females, 31 males) had an average age of 34.1 years (range: 18–62 years, SD = ±9.4 - for full descriptive data for the final sample see Supplementary Table 1). The study was approved by the Ethics Committee Saint-Luc Hospital, UCLouvain (2020/30JUL/389 - DNMT-TMS – amendment 3), and participants gave informed consent though a computerized form. Participants received compensation of up to 8€ for their time.

2.3.2. Procedure and stimuli

The experimental protocol was implemented using PsychoPy version 2024.4, running on Python 3.10 (51) and transformed to PsychoJS experiment to be hosted online on Pavlovia servers (52).

The experiment began with initial screens that provided instructions to the observer. Each trial of the experiment then began with a preparation screen (see Figure 1), displaying a fixation cross in the center of the screen for 1 second. The main stimuli in this experiment consisted of short video clips, each lasting 2 seconds, with a single pulse of TMS being delivered at a specific intensity 1 second into the video. Following the video, participants were given a 10-second response window in which they

answered "yes" (right arrow) "no" (left arrow) to indicate whether they observed any

movement in the hand. In the case of a technical issue with the stimulus presentation,

participants could select the option "problem" (down arrow), which would mark the

trial for removal from any subsequent analysis.

After the initial instructions were provided, participants observed two example

videos: one without any movement in response to TMS, and one with a pronounced

movement in response to TMS. Participants then completed blocks in which they

observed 10 trials where TMS was delivered at a fixed intensity of MSO. At the end of

each block (i.e., after viewing all 10 videos for a given intensity level), participants

encountered a pause screen and were able to rest as long as they required before

beginning the next block of trials.

Each participant in Group 2 was randomly assigned to view videos taken from one of

the participants in Group 1 (from this point onwards we refer to participants in Group

1 as "observers" to help distinguish them from participants in Group 1). Notably, the

number of trials required to produce a clear visual response in all trials at a fixed

intensity varied from participant to participant; as such, the total number of trials that

observers in Group 2 needed to complete varied. Observers saw trials delivered

across at least 15 intensities, and a maximum 20 intensities (average 17.8 intensities

– see also table 1 for summary information). Every 5 blocks, an attention task

appeared and asked the observer to press the "up" arrow; this attention check was

designed to be able to catch any participants who were not following the instructions

given (e.g. simply repeatedly pressing the same response button across all trials, or

randomly selecting one of the three buttons that were normally used to provide a

response).

Figure 1 : Trial structure for the visual inspection task.

2.3.3. Data processing

For each observer, the number of "movement" and "non-movement" responses were

recorded. These responses were then collapsed across each intensity level to

determine the RMT, defined as the intensity at which the proportion of trials with a

"movement" response reached or exceeded 50%. This approach is equivalent to the

standard EMG/visual method but accounts for any trials that were removed due to

technical errors. In total 34 trials were removed (i.e. 0.3% of all trials presented).

2.4. Statistical analyses

All statistical analyses were performed using RStudio (version 4.4.1).

2.4.1. Bayesian ANOVA

A Bayesian ANOVA mixed-effects model was implemented to analyze the effect of the conditions on the dependent variable RMT. This approach allows for the estimation for the effects of conditions (EMG, ACC, and VIS) while accounting for model uncertainty. Random effects, including video series (i.e. different participants from Group 1) and observers (i.e. different participants from Group 2), were incorporated to control inter-individual variability. To interpret the Bayes factors (BFs), statistical analysis followed the classification proposed by Lee & Wagenmakers (53). A grouped scatter plot was created to visualize the data distribution and inter-individual variability in RMT measurements. To further refine the analysis, a Model-Averaged Posterior Summary was conducted, providing weighted mean estimates of condition effects while integrating model selection uncertainty. The posterior mean, standard deviation, and 95% credible intervals were extracted for each condition (EMG, ACC, VIS) to offer a more robust evaluation of their impact on RMT.

Bland-Altman Plots

Separate Bland-Altman Plots were generated to examine the agreement between EMG

(i.e. the gold standard) and ACC, as well as EMG and VIS, by analyzing systematic biases

and limits of agreement. The Bland-Altman plot takes the difference between two

measures plotted against their mean, and is therefore useful to observe systematic bias

from one method to the other. This visualization was essential for identifying potential

measurement discrepancies between the different methods, and assessing their level of

agreement. For data examining VIS conditions, we calculated 95% confidence intervals

based on a bootstrapping procedure. For each of the video series presented (i.e. data

from a single participant in Group 1), 10,000 resamples of the corresponding VIS data

(i.e. observers from Group 2 who saw videos from that specific participant in Group 1)

were generated with replacement. The mean for each resample was then taken,

allowing us to identify a 95% confidence interval (i.e. the 2.5th and 97.5th percentiles).

This approach thus ensured that the analyses were not biased by unequal observer

contributions or systematic rating sequences, allowing us to obtain more robust

estimations of the effects across conditions.

2.4.2. Intraclass Correlation Coefficient (ICC) analyses

Several reliability and agreement analyses were performed to model condition

effects. First, Intraclass Correlation Coefficient (ICC) with a two-way random effect

model of absolute agreement and single measures (i.e. ICC(2,1), used to evaluate the

agreement across raters/measurements) was computed to assess the absolute

agreement between EMG (gold standard) and VIS, considering differences across

observers(54). Given that the number of observers varied between participants and

video series, we implemented a bootstrap-based resampling procedure. We

resampled the data 10,000 times, such that each of the 5 different video series

(depicting data from Group 1) was assessed by one randomly selected observer (i.e.

a single participant from Group 2 was selected from all those available who had

watched the corresponding video series). The ICC(2,1) was then calculated for each

resample. This approach allowed us to calculate both the mean value, and a

bootstrapped 95% confidence interval (taking the 2.5% and the 97.5% percentile ICC

values), allowing us to generate a more detailed estimation of the distribution of ICC

values. To verify that this resampling did not bias the results, an ICC analysis based

on the mean ratings per condition (i.e. ICC (2,k)) analysis with a two-way random

effect model of absolute agreement and multiple measures was also calculated and

compared with the ICC (2,1) confidence interval.

Additionally, an ICC (2,1) was computed to assess agreement between ACC and VIS

conditions, following the same bootstrapping resampling to ensure robustness

despite the variability in observer numbers. The ICC (2,k) was computed following

the same methodology as described above.

Finally, the ICC (2,1) was computed to assess agreement between ACC and EMG

conditions following without bootstrapping due to the number of data acquired (no

ICC (2,k) analysis was performed as only one measure was available per participant

in Group 1).

To establish reliability thresholds, analysis referred to the classification proposed in

Koo & Li (2016) (54) as values below 0.5 indicate poor reliability, those between 0.5

and 0.75 reflect moderate reliability, values ranging from 0.75 to 0.9 suggest good

reliability, and values exceeding 0.90 denote excellent reliability. ▯

3. Results

3.1. Bayesian Model Comparisons

Bayesian Model Comparison of a null model (which included video series and observers as random effects), and a 'condition' model (which included the conditions EMG, ACC, and VIS as fixed effects) indicated there was "extreme" evidence in favour of the model including the factor of condition (BF_{10}= 11381). A grouped scatter plot demonstrates the effect of the different conditions (Fig. 2).

Figure 2. Individual Resting Motor Threshold (RMT) estimates by method and participant. Each panel shows data for one of the five participants from Group 1. Grey dots represent RMT estimates based on visual inspection responses from one of the

The Model Averaged Posterior Summary provides estimates for key variables (Table. 1). Notably, the RMT as estimated by EMG was lower than that for either ACC or VIS, with non-overlapping confidence intervals for these comparisons indicating there was a statistically meaningful difference when estimating the RMT using EMG compared to other techniques. By contrast, the overlapping 95% confidence intervals for ACC and VIS indicates their estimates are statistically indistinguishable within the model, suggesting no meaningful difference in estimates of the RMT exists between these two techniques. Furthermore, the positive estimates for ACC and VIS and the fact that the 95% credible intervals remain entirely above zero, suggesting a tendency to overestimate RMT when compared to EMG.

	Level	Mean	SD	95% Credible Interval	
				Lower	Upper
Intercept		50.1	2.1	45.6	54.4
	ACC	1.3	0.6	0.1	2.4
Condition	**EMG**	-3.0	0.6	-4.3	-1.8
	VIS	1.7	0.5	0.6	2.8

Table 1: Model-averaged posterior summary statistics for the effects of condition on

Resting Motor Threshold (RMT). The intercept reflects the grand mean baseline RMT (in

% of maximum stimulator output), while condition estimates indicate the deviation

from the intercept for each technique: accelerometry (ACC), electromyography (EMG),

and visual inspection (VIS). Positive values represent an overestimation relative to the

baseline, and negative values represent an underestimation. Posterior means, standard

deviations (SD), and 95% credible intervals are reported.

3.2. Bland-Altman analysis

When comparing EMG to VIS (Fig 3A), the analysis revealed a systematic bias, as the

95% confidence interval of the mean difference did not include zero, indicating that

VIS consistently overestimated RMT relative to EMG rather than fluctuating randomly

around it (mean=5.30, 95%CI=2.30-7.22) Additionally, the wide confidence interval

suggested substantial variability in the differences between VIS and EMG data.

By contrast, when comparing ACC to VIS (Fig 3B), the analysis indicated no systematic

bias, as the 95% confidence interval of the mean difference included zero

(mean=0.44, 95%CI=-1.25-1.90). However, despite this absence of bias, the broad

confidence interval suggested high variability across observers, indicating that while

VIS did not consistently over- or underestimate ACC, individual assessments varied

considerably between observers.

Finally, comparing EMG to ACC (Fig 3C), the analyses showed systematic bias, as the

95% confidence interval of the mean difference is higher than zero (mean=4.8, 95%CI

=1.20-7.00).

Figure 3: Bland-Altman plots comparing Visual Inspection with Electromyography (A),

Visual Inspection with Accelerometry (B) and Accelerometry with Electromyography

(C) as reference methods for resting motor threshold. The x-axis represents the mean

RMT (%) between the two methods, while the y-axis shows the difference in RMT (%)

between visual inspection and the reference method. Each dot represents an individual

measurement pair (though note that the data presented here are representative of a

bootstrap analysis with 10,000 resamples; due to the high density of data, numerous

data points overlap). The solid horizontal line indicates the mean bias, representing the

systematic difference between the methods, while the dashed lines denote the 95%

confidence interval. Each plot includes 10,000 data points from the resamples. Due to

the high density of data, numerous points overlap in regions, leading to visual clustering.

3.3. Intraclass Correlation Coefficient Analyses

Summary statistics for the bootstrapped analysis are provided in Table 2.

For the bootstrap based ICC (2,1) analyses, when comparing VIS to EMG, the mean ICC value identified only 'acceptable' reliability, and the 95% confidence interval included responses ranging from 'unacceptable-to-good' (mean ICC=0.58, 95%CI=0.39-0.74), with the majority of resamples indicating only 'acceptable' reliability. By contrast, when comparing VIS to ACC, the mean ICC value identified 'good' reliability, with a 95% confidence interval including a range of responses spanning from 'acceptable-to-excellent' (mean ICC = 0.84, 95%CI=0.62-1.00) (Fig. 4).

To complement these results, ICC (2,k) values based on averaged ratings were also calculated. These showed "excellent" reliability for VIS vs ACC (point estimate = 0.980, 95% CI = 0.815–0.997) and "good" reliability for VIS vs EMG, though the 95% confidence interval indicated considerable variability, including responses ranging from 'unacceptable-to-excellent' (point estimate = 0.753, 95% CI = 0.132–0.973).

Finally, the ICC (2,1) comparison between EMG and ACC showed "acceptable" reliability, again with a wide 95% confidence interval, which included responses ranging from 'unacceptable-to-excellent' (point estimate= 0.587, 95%CI=0.108-0.943). No ICC (2,k) was calculated for this comparison due to the limited number of observations available for EMG and ACC.

	ICC (2,1)			ICC (2,k)		
	Mean Point	**95% CI**		**Point**	**95% CI**	
	Estimate	**Lower**	**Upper**	**Estimate**	**Lower**	**Upper**
VIS vs ACC	0.845	0.615	1.000	0.980	0.815	0.997
VIS vs EMG	0.580	0.389	0.748	0.753	0.132	0.973
ACC vs EMG	0.587	-0.108	0.943	N/A	N/A	N/A

Table 2: Intraclass Correlation Coefficient (ICC) values comparing Resting Motor

Threshold (RMT) point estimates across methods. ICC (2,1) values reflect agreement

between individual observer responses, while ICC (2,k) values reflect agreement based

on average ratings per condition. Comparisons include visual inspection (VIS),

accelerometry (ACC), and electromyography (EMG). Values are presented with their

corresponding 95% confidence intervals. No ICC (2,k) was calculated for the ACC–EMG

comparison as only one comparison was available per due to limited sample size.

Figure 3 : Results of the bootstrap analysis demonstrating the distribution of Intraclass

Correlation Coefficients (ICC 2,1) for visual inspection compared to accelerometry

(Yellow) and electromyography (Blue). Violin plots represent the full distribution of ICC

values with the dot representing the mean and vertical line the 95%CI, while box plots

indicate the median, interquartile range, and overall dispersion. Each dashed line

represents the reliability threshold based on Koo and Li (2016).

4. Discussion

The present study compared the resting motor threshold as estimated using

electromyography, accelerometry, and visual inspection. A series of analyses

indicated that while estimates of the resting motor threshold produced by

accelerometry and visual inspection are generally similar, these techniques typically

over-estimate the threshold as determined using electromyography. However,

further analysis supported the view that accelerometry provides a more objective

and quantifiable approach to determining the resting motor threshold than visual

inspection, which varied considerably between observers. These findings support the

view that electromyography remains the gold-standard for estimating the resting

motor threshold, but in cases where the use of electromyography is not feasible, the

objective and less variable nature of accelerometry provides a preferable alternative

to visual inspection.

The present manuscript supports the view that estimation of the resting motor

threshold using electromyography represents the 'gold standard'. Bayesian Model

comparison and Bland-Altman analyses both indicated that visual inspection over-

estimated the RMT when compared to the use of EMG. This overestimation is easily

explained by the fundamental differences in what each method measures. EMG

captures an intrinsic characteristic of the muscle—its electrical activity—providing a

direct assessment of neuromuscular activation. In contrast, accelerometry and visual

inspection rely on an extrinsic characteristic (i.e., overt movement of the finger).

Muscle contraction (detectable by EMG) does not always result in displacement of the

finger (as measured using ACC and VIS), and movement requires overcoming factors

such as inertia and frictional forces. These differences would explain why ACC and

VIS tend to overestimate RMT, as they may fail to detect subtle contractions that do

not produce sufficient movement.

These results are partially in line with the previous research that has compared the

use of visual inspection to EMG when estimating the resting motor threshold (43,44).

However, the protocols and findings of these previous studies remain subject to

discussion. Pridmore et al. (1998) (42) reported that motor thresholds determined

using EMG compared to visual inspection showed a difference of less than 10% in

percent of maximum stimulator output, suggesting a general consistency between

these approaches. However, they also found that in most cases, the threshold detected

through visual inspection was reached at *lower* intensities than with EMG. This

discrepancy raises questions about the comparability and sensitivity of these

methods in assessing cortical excitability, particularly regarding the potential

influence of methodological differences on threshold determination. Meanwhile,

Balslev et al. (2007) (45) employed a visual inspection protocol assessing the effects

of eight different coil orientations on the resting motor threshold as estimated using

EMG and visual inspection. While their results indicated that visual inspection

typically over-estimated the RMT as determined by EMG by approximately 2% of

MSO, as the comparison between these two methods was not the main focus of their

study, only data from four participants using a primarily repeated-measures design

(i.e. 8 different coil orientations were assessed for each participant) was collected.

This procedure may therefore have led to an under-estimation of differences between

the RMT as estimated using EMG compared to visual inspection. We also note that

the results of the present study are generally in line with the findings of our previous

research (49), in which we found that the RMT as estimated via EMG and ACC were

highly correlated, but that RMTs identified by EMG were generally lower than those

identified using accelerometry. An important consideration when interpreting our

findings is the pronounced measurement error inherent to VIS, which distinguishes

it from techniques such as EMG and accelerometry. When the same motor event is

recorded and analyzed using EMG or accelerometry, and the same detection

algorithm is applied, the outcome is reproducible, yielding a single, objective value

per trial. In contrast, VIS introduces considerable variability, as it relies on subjective

human judgment.

Further analyses considered whether accelerometry provided any benefits compared

to visual inspection. Here we note that while accelerometry provides a single

measurement that can be quantified objectively, visual inspection, by comparison,

remains a more subjective approach to identifying the Resting Motor Threshold.

Bland-Altmann analysis indicated that while the average RMT provided by visual

inspection matched well to that of estimates provided by EMG, there was

considerable variability around the mean (95%CI ranging from −4 to +5 MSO).

Similarly, our ICC bootstrapping analysis indicated that while on average VIS has

"good" reliability when compared to ACC, there was again a large disparity in the

95%CI, indicating that, in the best-case scenario, an observer (VIS) may provide a

value close to ACC, but in other cases, their estimate of the RMT could be far from it.

Given these findings, we propose that researchers who are not able to use EMG to

determine the RMT should prefer the use of accelerometry, rather than visual

inspection, wherever possible. Unlike VIS, which depends on individual observers

and introduces variability, accelerometry provides a single, consistent measurement

per trial, reducing subjectivity and improving measurement reliability. Nonetheless,

a limitation of the accelerometry is that it requires real-time (online) data processing

through a dedicated script.

However, several limitations should be acknowledged. First, the sample size for the

EMG and the accelerometry conditions was limited to five participants, which

constrains the generalizability of these findings and increases the risk of sampling

bias. We note, however, that the results of the comparisons between EMG and ACC in

the present study are in agreement with our previous work which has demonstrated

the same effects in a larger sample size (49). Second, a large proportion of the data

for visual inspection were collected in a remote, online experiment, which prevented

close monitoring of participant engagement. The inclusion of attention checks

therefore helped to maintain high quality data (i.e. approximately 1/3 of participants

were rejected based on this measure). Third, the visual inspection condition relied on

a participant reviewing recorded videos, rather than observing the stimulation in real

time. This may have limited the accuracy of our visual inspection measurements, as

subtle movements or contextual cues available in a live setting may not be as easily

detectable on video. Nevertheless, a major strength of the present study is a much

larger sample size when examining visual inspection compared to previous studies.

Another important limitation of both visual inspection and accelerometry is their

inability to assess background muscle activity prior to TMS. EMG, in contrast, allows

for the quantification of baseline muscle activation, which can significantly influence

MEP responses and RMT estimates. Failing to detect such pre-activation may lead to

an underestimation of the RMT and contribute to the variability of non-EMG methods.

A further limitation is that the observers in the present study can be considered

'naive' or untrained, when compared to an experimenter who may have more

experience with observing responses in relation to TMS. While we would agree that

training could have reduced the variability in the present experiment, individual

differences in perception and judgment would remain, meaning that variability in

visual inspection measurements would be reduced, but not fully eliminated; thus,

visual inspection would still introduce variability to the measurement of the resting

motor threshold that accelerometry would remove. Additionally, the convergence of

our findings with previous research (e.g., our prior work on EMG and ACC

correspondence) further supports the reliability of the present data. In particular, the

strong agreement observed between EMG and ACC measurements confirms the

replicability of our previous results, and highlights the potential of using an equation

to convert the RMT as calculated by accelerometry to an estimated value of the RMT

as would be determined by EMG. Accelerometry therefore remains a viable

alternative when EMG is not available.

In conclusion, the present study indicates that EMG provides the most sensitive

approach to identifying the RMT, most likely due to its ability to measure the intrinsic

excitability of the corticospinal system. In situations where access to

electromyography is limited, researchers should consider using accelerometry in the

place of visual inspection, as accelerometry provides a more objective and reliable

approach.

5. Research data

The research data supporting the findings of this study are stored in an OSF (Open Science Framework) repository and are available upon reasonable request from the corresponding author at gautier.hamoline@uclouvain.be. Open public access is restricted to protect participant confidentiality in accordance with the guidelines of the approving ethics committee.

6. Acknowledgements

Gautier Hamoline is supported by a PhD fellowship from the Faculty of Motricity Sciences, UCLouvain, and the Belgian Fund for Scientific Research (FNRS J.0084.21). Robert Hardwick is supported by grants from the UCLouvain Special Research Funds Seedfunds (FSR 1C21300057) and the Belgian Fund for Scientific Research (FNRS F.4523.23 and FNRS J.0084.21). Marcos Moreno-Verdu is supported by a grant from the Belgian Fund for Scientific Research (FNRS 1.B359.25). Elise Van Caenegem is supported by a grant from the Belgian Fund for Scientific Research (FNRS 1.AB19.24). Baptiste Waltzing and Siobhan McAteer are supported by the Belgian Fund for Scientific Research (FNRS F.4523.23).

7. Author Roles (CRediT)

Author	Conceptualization	Data curation	Formal analysis	Funding acquisition	Investigation	Methodology	Project administration	Resources	Software	Supervision	Validation	Visualization	Writing – original draft	Writing - review and editing
Hamoline Gautier	x	x	x		x	x	x	x	x	x	x	x	x	x
Antoine Lunardi		x			x								x	
Marcos Moreno-Verdu			x						x			x		x
Baptiste M Waltzing		x												x
Elise E Van Caenegem		x												x
Siobhan MacAteer									x					x
Robert M Hardwick	x			x	x	x			x	x	x		x	x

8. Bibliography

1.   Hallett M. Transcranial magnetic stimulation and the human brain. Nature.

juill 2000;406(6792):147-50.

2.   Barker AT, Jalinous R, Freeston IL. Non-invasive magnetic stimulation of

human motor cortex. Lancet. 11 mai 1985;1(8437):1106-7.

3.   Aloizou AM, Pateraki G, Anargyros K, Siokas V, Bakirtzis C, Liampas I, et al.

Transcranial magnetic stimulation (TMS) and repetitive TMS in multiple sclerosis.

Reviews in the Neurosciences. 1 nov 2021;32(7):723-36.

4.   Beer S, Rösler KM, Hess CW. Diagnostic value of paraclinical tests in multiple

sclerosis: relative sensitivities and specificities for reclassification according to the

Poser committee criteria. J Neurol Neurosurg Psychiatry. août 1995;59(2):152-9.

5.   Hess CW, Mills KR, Murray NM. Measurement of central motor conduction in

multiple   sclerosis   by   magnetic   brain   stimulation.   Lancet.   16   août

1986;2(8503):355-8.

6.   Rossini PM, Zarola F, Floris R, Bernardi G, Perretti A, Pelosi L, et al. Sensory

(VEP, BAEP, SEP) and motor-evoked potentials, liquoral and magnetic resonance

findings in multiple sclerosis. Eur Neurol. 1989;29(1):41-7.

7.   Šoda J, Pavelin S, Vujović I, Rogić Vidaković M. Assessment of Motor Evoked

Potentials in Multiple Sclerosis. Sensors. janv 2023;23(1):497.

8.   McDonnell MN, Stinear CM. TMS measures of motor cortex function after

stroke: A meta-analysis. Brain Stimulation. 1 juill 2017;10(4):721-34.

9.   Richards LG, Stewart KC, Woodbury ML, Senesac C, Cauraugh JH. Movement-

dependent stroke recovery: A systematic review and meta-analysis of TMS and fMRI

evidence. Neuropsychologia. 1 janv 2008;46(1):3‑11.

10.   Badawy RAB, Strigaro G, Cantello R. TMS, cortical excitability and epilepsy:

The clinical impact. Epilepsy Research. 1 févr 2014;108(2):153‑61.

11.   Theodore WH. Transcranial Magnetic Stimulation in Epilepsy. Epilepsy Curr.

1 nov 2003;3(6):191‑7.

12.   Brys M, Fox MD, Agarwal S, Biagioni M, Dacpano G, Kumar P, et al. Multifocal

repetitive TMS for motor and mood symptoms of Parkinson disease. Neurology. nov

2016;87(18):1907‑15.

13.   Cohen OS, Rigbi A, Yahalom G, Warman-Alaluf N, Nitsan Z, Zangen A, et al.

Repetitive Deep TMS for Parkinson Disease: A 3‑Month Double-Blind, Randomized

Sham-Controlled Study. Journal of Clinical Neurophysiology. mars 2018;35(2):159.

14.   Elahi B, Elahi B, Chen R. Effect of transcranial magnetic stimulation on

Parkinson motor function—Systematic review of controlled clinical trials. Movement

Disorders. 2009;24(3):357‑63.

15.   Ollanova SS, Ugli TSZ. ANALYSIS OF THE EFFICACY OF TMS IN RESPECT OF

NONMOTOR DISORDERS IN PARKINSON'S DISEASE. World Bulletin of Public Health.

24 mars 2023;20:71‑5.

16.   Brighina F, Romano M, Giglia G, Saia V, Puma A, Giglia F, et al. Effects of

cerebellar TMS on motor cortex of patients with focal dystonia: a preliminary report.

Exp Brain Res. 1 févr 2009;192(4):651-6.

17.   Lozeron P, Poujois A, Richard A, Masmoudi S, Meppiel E, Woimant F, et al.

Contribution of TMS and rTMS in the Understanding of the Pathophysiology and in

the Treatment of Dystonia. Front Neural Circuits [Internet]. 10 nov 2016 [cité 6 janv

2025];10.   Disponible   sur: https://www.frontiersin.org/journals/neural-

circuits/articles/10.3389/fncir.2016.00090/full

18.   Quartarone A. Chapter 43 - Transcranial magnetic stimulation in dystonia. In:

Lozano AM, Hallett M, éditeurs. Handbook of Clinical Neurology [Internet]. Elsevier;

2013 [cité 6 janv 2025]. p. 543-53. (Brain Stimulation; vol. 116). Disponible sur:

https://www.sciencedirect.com/science/article/pii/B9780444534972000437

19.   Liepert J, Schoser BGH, Weiller C. Motor excitability in myopathy. Clinical

Neurophysiology. 1 janv 2004;115(1):85-9.

20.   Molenaar JP, van Kleef E, van Zandvoort E, van Alfen N, van Engelen BG,

Voermans NC, et al. Detecting impaired muscle relaxation in myopathies with the use

of motor cortical stimulation. Neuromuscular Disorders. 1 mai 2023;33(5):396-404.

21.   Nardone R, Versace V, Sebastianelli L, Brigo F, Golaszewski S, Christova M, et

al. Cortical involvement in myopathies: Insights from transcranial magnetic

stimulation. Clinical Neurophysiology. 1 oct 2017;128(10):1971-7.

22.   Pridmore S, Turnier-Shea Y, Rybak M, Naguy A. Transcranial Magnetic

Stimulation (TMS) for Major Depressive Disorder—Modus Operandi!

Psychopharmacol Bull. 11 août 2023;53(3):55-60.

23.   Tendler A, Roth Y, Harmelech T. Deep repetitive TMS with the H7 coil is

sufficient to treat comorbid MDD and OCD. Brain Stimulation: Basic, Translational,

and Clinical Research in Neuromodulation. 1 mai 2021;14(3):658-61.

24.   van Rooij SJH, Arulpragasam AR, McDonald WM, Philip NS. Accelerated TMS -

moving quickly into the future of depression treatment. Neuropsychopharmacol. janv

2024;49(1):128-37.

25.   Carmi L, Yagon UA, Dar R, Zohar J, Zangen A. Deep Transcranial Magnetic

Stimulation (Tms) in Obsessive Compulsive Disorder (Ocd) Patients. European

Psychiatry. mars 2015;30(S1):1-1.

26.   Carmi L, Alyagon U, Barnea-Ygael N, Zohar J, Dar R, Zangen A. Clinical and

electrophysiological outcomes of deep TMS over the medial prefrontal and anterior

cingulate cortices in OCD patients. Brain Stimulation. 1 janv 2018;11(1):158-65.

27.   Harmelech T, Tendler A, Arikan MK, Çetin HL, Esmeray MT, Ilhan R, et al. Long-

term outcomes of a course of deep TMS for treatment-resistant OCD. Brain

Stimulation: Basic, Translational, and Clinical Research in Neuromodulation. 1 janv

2022;15(1):226-8.

28.　Cirillo P, Gold AK, Nardi AE, Ornelas AC, Nierenberg AA, Camprodon J, et al.

Transcranial magnetic stimulation in anxiety and trauma-related disorders: A

systematic review and meta-analysis. Brain and Behavior. 2019;9(6):e01284.

29.　Karsen EF, Watts BV, Holtzheimer PE. Review of the Effectiveness of

Transcranial Magnetic Stimulation for Post-traumatic Stress Disorder. Brain

Stimulation. 1 mars 2014;7(2):151-7.

30.　Yan T, Xie Q, Zheng Z, Zou K, Wang L. Different frequency repetitive

transcranial magnetic stimulation (rTMS) for posttraumatic stress disorder (PTSD):

A systematic review and meta-analysis. Journal of Psychiatric Research. 1 juin

2017;89:125-35.

31.　Cole JC, Green Bernacki C, Helmer A, Pinninti N, O'reardon JP. Efficacy of

Transcranial Magnetic Stimulation (TMS) in the Treatment of Schizophrenia: A

Review of the Literature to Date. Innov Clin Neurosci. 2015;12(7-8):12-9.

32.　Hou M di, Santoro V, Biondi A, Shergill SS, Premoli I. A systematic review of

TMS and neurophysiological biometrics in patients with schizophrenia. Journal of

Psychiatry and Neuroscience. 21 déc 2021;46(6):E675-701.

33.　Gersner R, Barnea-Ygael N, Tendler A. Moderators of the response to deep TMS

for smoking addiction. Front Psychiatry [Internet]. 9 janv 2023 [cité 6 janv 2025];13.

Disponible　　　　　　　　　　　　　　　　　　　　　　　　　　　　　　sur:

https://www.frontiersin.org/journals/psychiatry/articles/10.3389/fpsyt.2022.107

9138/full

34.   Hoegl T, Bender S, Buchmann J, Kratz O, Moll GH, Heinrich H. [Transcranial

magnetic stimulation (TMS), inhibition processes and attention deficit/hyperactivity

disorder (ADHD) - an overview]. Z Kinder Jugendpsychiatr Psychother. nov

2014;42(6):415-28; quiz 428-9.

35.   Jáuregui-Lobera I, Martínez-Quiñones JV. Neuromodulation in eating

disorders and obesity: a promising way of treatment? Neuropsychiatric Disease and

Treatment. 26 oct 2018;14:2817-35.

36.   Chepurova A, Hramov A, Kurkin S. Motor Imagery: How to Assess, Improve Its

Performance, and Apply It for Psychosis Diagnostics. Diagnostics (Basel). 11 avr

2022;12(4):949.

37.   Ambrosini E, Ferrante S, van de Ruit M, Biguzzi S, Colombo V, Monticone M, et

al. StimTrack: An open-source software for manual transcranial magnetic stimulation

coil positioning. Journal of Neuroscience Methods. 1 janv 2018;293:97-104.

38.   Micera S, Carpaneto J, Raspopovic S. Control of Hand Prostheses Using

Peripheral Information. IEEE Reviews in Biomedical Engineering. 2010;3:48-68.

39.   Peri E, Ambrosini E, Colombo VM, Ruit M van de, Grey MJ, Monticone M, et al.

Intra and inter-session reliability of rapid Transcranial Magnetic Stimulation

stimulus-response curves of tibialis anterior muscle in healthy older adults. PLOS

ONE. 14 sept 2017;12(9):e0184828.

40.   Cantello R, Rossi S, Varrasi C, Ulivelli M, Civardi C, Bartalini S, et al. Slow

Repetitive TMS for Drug-resistant Epilepsy: Clinical and EEG Findings of a Placebo-

controlled Trial. Epilepsia. 2007;48(2):366-74.

41.   Hoogendam JM, Ramakers GMJ, Di Lazzaro V. Physiology of repetitive

transcranial magnetic stimulation of the human brain. Brain Stimulation. 1 avr

2010;3(2):95-118.

42.   Ikeguchi M, Touge T, Kaji R, Deguchi K, Sasaki I, Tsukaguchi M, et al. Durable

effect of very low-frequency repetitive transcranial magnetic stimulation for

modulating cortico-spinal neuron excitability. International Congress Series. 1 mars

2005;1278:272-5.

43.   Pridmore S, Fernandes Filho JA, Nahas Z, Liberatos C, George MS. Motor

Threshold   in   Transcranial   Magnetic   Stimulation: A   Comparison   of   a

Neurophysiological Method and a Visualization of Movement Method. The Journal of

ECT. mars 1998;14(1):25.

44.   Balslev D, Braet W, McAllister C, Miall RC. Inter-individual variability in

optimal current direction for transcranial magnetic stimulation of the motor cortex.

Journal of Neuroscience Methods. 15 mai 2007;162(1):309-13.

45.   Rossi S, Hallett M, Rossini PM, Pascual-Leone A. Safety, ethical considerations,

and application guidelines for the use of transcranial magnetic stimulation in clinical

practice and research. Clinical Neurophysiology. 1 déc 2009;120(12):2008-39.

46.   Classen J, Liepert J, Wise SP, Hallett M, Cohen LG. Rapid Plasticity of Human
Cortical Movement Representation Induced by Practice. Journal of Neurophysiology.
févr 1998;79(2):1117-23.

47.   Duque J, Mazzocchio R, Stefan K, Hummel F, Olivier E, Cohen LG. Memory
Formation in the Motor Cortex Ipsilateral to a Training Hand. Cerebral Cortex. 1 juin
2008;18(6):1395-406.

48.   Mawase F, Uehara S, Bastian AJ, Celnik P. Motor Learning Enhances Use-
Dependent Plasticity. J Neurosci. 8 mars 2017;37(10):2673-85.

49.   Hamoline G, Van Caenegem EE, Waltzing BM, Vassiliadis P, Derosiere G, Duque
J, et al. Accelerometry as a tool for measuring the effects of transcranial magnetic
stimulation. Journal of Neuroscience Methods. 1 mai 2024;405:110107.

50.   Lefaucheur JP, André-Obadia N, Poulet E, Devanne H, Haffen E, Londero A, et
al. Recommandations françaises sur l'utilisation de la stimulation magnétique
transcrânienne répétitive (rTMS) : règles de sécurité et indications thérapeutiques.
Neurophysiologie Clinique/Clinical Neurophysiology. 1 déc 2011;41(5):221-95.

51.   Peirce J, Gray JR, Simpson S, MacAskill M, Höchenberger R, Sogo H, et al.
PsychoPy2: Experiments in behavior made easy. Behav Res. 1 févr
2019;51(1):195-203.

52.   Bridges D, Pitiot A, MacAskill MR, Peirce JW. The timing mega-study:
comparing a range of experiment generators, both lab-based and online. PeerJ. 20 juill
2020;8:e9414.

53.   Lee MD, Wagenmakers EJ. Bayesian statistical inference in psychology:

Comment on Trafimow (2003). Psychological Review. 2005;112(3):662-8.

54.   Koo TK, Li MY. A Guideline of Selecting and Reporting Intraclass Correlation

Coefficients for Reliability Research. Journal of Chiropractic Medicine. 1 juin

2016;15(2):155-63.

Supplementary materials

Group 2 (Observers)						
Video Series Source (i.e. Group 1 Participant ID)	**1**	**2**	**3**	**4**	**5**	**Total**
Valid	10	17	13	9	15	64
Rejected	10	3	7	11	5	36
Mean age (years)	35.3	34,2	35.3	35.5	30.2	34.1
Min age (years)	20	20	18	19	18	18
Max age (years)	54	54	53	59	62	62
Intensities presented	20	15	19	20	15	86

Descriptive Statistics

	RMT				
	1	2	3	4	5
Valid	11	17	14	10	17
Missing	0	0	0	0	0
Mean	57.9	49.6	52.3	52.0	47.3
Std. Deviation	1.6	1.6	2.8	2.7	1.3
Coefficient of variation	0.027	0.033	0.053	0.053	0.027
Range	4	5	10	7	6
Minimum	57	47	47	50	45
Maximum	61	52	57	57	51