Made available by Hasselt University Library in https://documentserver.uhasselt.be

Direct eye contact enhances mirroring of others' movements: A transcranial magnetic stimulation study Peer-reviewed author version

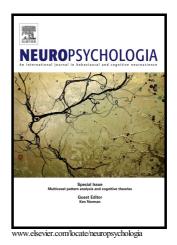
Prinsen, Jellina; Bernaerts, Sylvi; Wang, Yin; de Beukelaar, Toon; CUYPERS, Koen; Swinnen, Stephan & Alaerts, Kaat (2016) Direct eye contact enhances mirroring of others' movements: A transcranial magnetic stimulation study. In: NEUROPSYCHOLOGIA, 95, p. 111-118.

DOI: 10.1016/j.neuropsychologia.2016.12.011 Handle: http://hdl.handle.net/1942/22904

Author's Accepted Manuscript

Direct eye contact enhances mirroring of others' movements: A transcranial magnetic stimulation study

Jellina Prinsen, Sylvie Bernaerts, Yin Wang, Toon T. de Beukelaar, Koen Cuypers, Stephan P. Swinnen, Kaat Alaerts



 PII:
 S0028-3932(16)30449-3

 DOI:
 http://dx.doi.org/10.1016/j.neuropsychologia.2016.12.011

 Reference:
 NSY6202

To appear in: Neuropsychologia

Received date: 29 July 2016 Revised date: 11 November 2016 Accepted date: 7 December 2016

Cite this article as: Jellina Prinsen, Sylvie Bernaerts, Yin Wang, Toon T. de Beukelaar, Koen Cuypers, Stephan P. Swinnen and Kaat Alaerts, Direct eye contact enhances mirroring of others' movements: A transcranial magnetic stimulation study, *Neuropsychologia* http://dx.doi.org/10.1016/j.neuropsychologia.2016.12.011

This is a PDF file of an unedited manuscript that has been accepted fo publication. As a service to our customers we are providing this early version o the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting galley proof before it is published in its final citable form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain

Direct eye contact enhances mirroring of others' movements: A transcranial magnetic stimulation study Jellina Prinsen^{a*}, Sylvie Bernaerts^a, Yin Wang^b, Toon T. de Beukelaar^c, Koen

Cuypers^{c,d}, Stephan P. Swinnen^c, Kaat Alaerts^a

 ^aResearch Group for Neuromotor Rehabilitation, Department of Rehabilitation Sciences, KU Leuven, Tervuursevest 101, 3001 Leuven, Belgium
 ^bDepartment of Psychology, New York University, New York, United States
 ^cMovement Control and Neuroplasticity Research Group, Department of Kinesiology, KU Leuven, Tervuursevest 101, 3001 Leuven, Belgium
 ^dREVAL Rehabilitation Research Centre, Biomedical Research Institute, Faculty of Medicine and Life Sciences, Hasselt University, Agoralaan, 3590 Diepenbeek, Belgium.
 ^{*}Correspondence: jellina.prinsen@kuleuven.be

Abstract

Direct eye contact is a powerful social cue to regulate interpersonal interactions. Previous behavioral studies showed a link between eye contact and motor mimicry, indicating that the automatic mimicry of observed hand movements is significantly enhanced when direct eye contact exists between the observer and the observed model. In the present study, we aim to investigate the neurophysiological basis of the previously reported behavioral enhancements. Here, transcranial magnetic stimulation (TMS) was applied to assess changes in corticomotor excitability at the level of the primary motor cortex (M1) to explore whether and how the motor system is facilitated from observing others' hand movements and, in particular, how this process is modulated by eye contact. To do so, motor evoked potentials (MEPs) were collected from two hand muscles while participants received single-pulse TMS and naturally observed video clips of an actor showing hand opening movements or static hands. During the observation, either direct or averted eye gaze was established between the subject and the observed actor. Our findings show a clear effect of eye gaze on observationinduced motor facilitation. This indicates that the mapping or 'mirroring' of others' movements is significantly enhanced when movement observation is accompanied by direct eye gaze compared to averted eye gaze. Our results support the notion that eye contact is a powerful social signal with the ability to direct human non-verbal social behavior. Furthermore, our findings are important for understanding the role of the mirror motor system in the mapping of socially relevant actions.

Abbreviations

APB, abductor pollicis brevis; FC, fixation count; FDI, first dorsal interosseous; IFG, inferior frontal gyrus; IPL, inferior parietal lobule; M1, primary motor cortex; MEP, motor-evoked potential; mPFC, medial prefrontal cortex; RMSE, root mean square error; rMT, resting motor threshold; STS, superior temporal sulcus; TFD, total fixation duration; TMS, transcranial magnetic stimulation.

Keywords: Movement observation; gaze processing; eye contact; mirror-motor system; transcranial magnetic stimulation

1 Introduction

Human social interaction is a complex behavior between two or more individuals to communicate thoughts, intentions, emotional states and actions to one another. Ever since their discovery, 'mirror neurons' have been suggested to form an integral part of the neural circuitry that mediates our capacity to understand the meaning of the actions and behaviors of others (Gallese, 2009).

Neurons with mirror properties were first discovered using single-cell recordings in the ventral premotor cortex of macaque monkeys (Rizzolatti et al., 1988), and were shown to have the ability to fire not only when the monkey executes a certain motor action, but also when the monkey observes another individual performing the motor action (di Pellegrino et al., 1992; Gallese et al., 1996; Rizzolatti et al., 1996). Using movement observation paradigms in combination with functional neuroimaging techniques such as fMRI (Buccino et al., 2001; Iacoboni et al., 1999) and PET (Grafton et al., 1996; Rizzolatti et al., 1996) a homologous action observation—execution matching system or 'mirror system' has been localized in the human brain. Particularly, both frontal (inferior frontal gyrus; IFG) and parietal (inferior parietal lobule; IPL) areas have been shown to become increasingly activated during the mere observation of others' actions (Chong et al., 2008; Kilner et al., 2009).

Overall, and according to the notion of 'embodied cognition', this process of 'mapping' observed actions onto the corresponding sensorimotor representations has been hypothesized to form the core neural mechanism by which others' actions and emotional states can be simulated, recognized and understood (lacoboni, 2009; lacoboni et al., 2005; Rizzolatti and Craighero, 2004; Rizzolatti and Fabbri-Destro, 2008). However, note that also weaker accounts of 'embodied cognition' have been put forward, arguing that conceptual 'understanding of actions' may not be represented exclusively in terms of sensorimotor

processes, but may additionally involve an abstract or modality-independent representation (Caramazza et al., 2014; Mahon, 2015).

In the past decade, the non-invasive brain stimulation technique transcranial magnetic stimulation (TMS) has been used extensively to measure resonant 'mirror motor' activity in the observer's motor system. By applying TMS over the primary motor cortex (M1), a motor evoked potential (MEP) can be elicited from the contralateral muscles to obtain a measure of cortico-motor excitability (Fadiga et al., 1995). Interestingly, a number of studies (for a review, see Fadiga, Craighero and Olivier, 2005) have shown that during the mere observation of others' actions, cortico-motor excitability within parts of M1 becomes increasingly facilitated, as indicated by significant enhancements in MEP amplitudes. Furthermore, this process has been shown to be highly muscle-specific, such that modulations in M1 cortico-motor excitability are predominantly observed in the muscles that are used in the observed action (Alaerts, Heremans, Swinnen, & Wenderoth, 2009; Alaerts, Swinnen, & Wenderoth, 2009; Strafella & Paus, 2000). Besides muscular involvement, a number of studies used the TMS technique to explore how different kinematic features of the observed actions are encoded by the observer's motor system, such as temporal dynamics (Gangitano, Mottaghy, & Pascual-Leone, 2001), grip force (Alaerts, de Beukelaar, Swinnen, & Wenderoth, 2011; Alaerts et al., 2010; Alaerts, Swinnen, & Wenderoth, 2010), orientation (Maeda, Kleiner-Fisman, & Pascual-Leone, 2002) and predictability (Maeda, Chang, Mazziotta, & Iacoboni, 2001).

The mapping mechanism for conveying information from others' behaviors is not only affected by kinematic features, but may also be influenced by the processing of socially relevant cues from the observed environment (Wang and Hamilton, 2012). One such powerful social cue is perceived eye contact. The role of eye gaze in social behavior has been investigated extensively, with several neuroimaging studies showing that observed eye contact is a strong modulator of activity in regions of the 'social brain', a network of structures that is specialized to process social information such as faces, theory of mind and empathy, but also biological motion, action and goal direction (for a review, see Senju and Johnson, 2009). Particularly within the superior temporal sulcus (STS), brain activity has been shown to be specifically enhanced when direct eye contact is perceived (Pageler et al., 2003; Pelphrey, Viola and McCarthy, 2004). Furthermore, the STS region has also been hypothesized to form an integral part of the 'extended' mirror system network by providing the main visual input to upstream fronto-parietal mirror-motor regions (Grèzes et al., 2001; Grossman and Blake, 2002; Grossman et al., 2000).

To date however, only a handful of studies have explored the effect of perceived eye contact on the processing of the actions and movements of others. In terms of movement mimicry, a recent behavioral study by Wang, Newport and Hamilton (2011) provided first indications that the tendency of an observer to mimic others' actions is enhanced when eye contact exists between the observer and the model. As a form of unconscious imitation, mimicry is strongly associated with the mirror neuron system (Iacoboni, 2009). In particular, reaction times for mimicking a hand closing or opening movement were shown to be faster when direct eye contact was established, rather than when eye gaze was averted (Wang, Newport, et al., 2011). Also a magnetoencephalographic (MEG) study by Kilner, Marchant and Frith (2006) provided evidence that the social relevance of a stimulus (modulated in terms of the observer's viewpoint) can enhance putative mirror neuron activity. Together, these observations provide first indications that activity within the human mirror system can be influenced by distinct socially relevant cues from the observed environment.

To the best of our knowledge, no studies to date have directly investigated the neurophysiological basis of the effect of eye contact on motor resonance, as research has mainly focused on mimicry (Wang, Newport, et al., 2011; Wang, Ramsey, et al., 2011) or the influence of higher-order cognitive processes such as social relevance observation (Kilner et al., 2006). However, since eye contact is a powerful social cue, it would be interesting to directly explore whether direct eye gaze can modulate the mapping of others' actions in the observer's motor system. In the present study, the TMS technique was used to assess the effect of eye gaze on motor facilitation of M1 during movement observation. In particular, single-pulse TMS was applied over left M1 to measure the level of cortico-motor excitability of two hand muscles (right abductor pollicis brevis (APB) and first dorsal interossei (FDI)) during the observation of an actor performing simple hand movements involving those muscles. During the movement observation trials, the actor looked either directly towards or away from the observing participant to assess the effect of direct versus averted eye gaze on observation-induced motor facilitation at the level of M1. If eye gaze forms a salient social cue for modulating the process of mirror-motor mapping at the level of M1, we expected TMS-evoked MEPs to be higher when accompanied by direct gaze compared to averted gaze.

2 Material and Methods

2.1 Main experiment: Measurements of cortico-motor excitability during movement observation

2.1.1 Participants

Thirty-three right-handed individuals (16 males and 17 females) aged between 19 and 26 years old (mean \pm SD: 22;7 \pm 1;8 years;months) participated in this study. Handedness was assessed with the Edinburgh Handedness Questionnaire (EHQ; Oldfield, 1971). All participants provided signed written informed consents prior to the experiment, reported no history of neurological/psychiatric illness or motor dysfunctions of the hands/arms and met safety criteria for TMS. Ethical approval for the experiment was granted by the local Ethics Committee for Biomedical Research at the Katholieke Universiteit Leuven and conformed to the Code of Ethics of the World Medical Association (Helsinki, 1964). One female subject was excluded due to technical problems during the experiment.

To explore whether modulating effects of eye gaze were related to inter-individual differences in social responsiveness, subjects completed the Dutch self-report version of the Social Responsiveness Scale for adults (SRS-A; Constantino and Todd, 2005). The SRS-A (64 items) is a widely used screening tool to identify the presence and extent of any social impairments in the typical population using a four-point Likert-scale. It encompasses four subscales, including social awareness (19 items; $\alpha = .80$), social communication (22 items; $\alpha = .88$), social motivation (11 items; $\alpha = .83$) and rigidity/repetitiveness (12 items; $\alpha = .79$). Lower scores indicate higher social responsiveness. For raw SRS scores, a cut-off point of 54 is suggested for signaling impairments in social responsiveness (Noens et al., 2012).

2.1.2 General procedure

Participants were seated in a comfortable chair approximately 80 cm in front of a widescreen DELL monitor (resolution: 1920×1080 pixels, refresh frequency: 60 Hz) on which video stimuli of hand movements were displayed with a frame rate of 29 Hz. The right hand was placed palm-down on a soft cushion on their lap and participants were asked to relax their hand muscles while spontaneously viewing the presented video clips. During the experiment subjects' vision of their own hands was obstructed by another cushion placed on top of their hands.

2.1.3 Electromyography recordings and TMS

Dependent measures of cortico-motor excitability, i.e. motor-evoked potentials (MEPs), were recorded via electromyography (EMG). To do so, disposable self-adhesive electrodes were

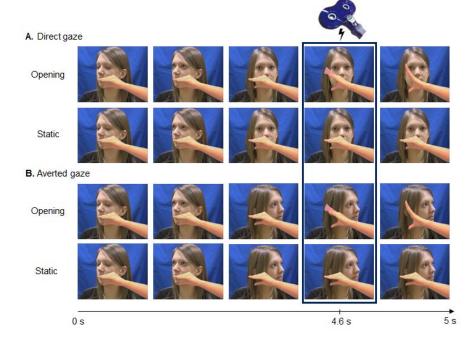
attached to the muscle bellies of the right hand abductor pollicis brevis (APB) and first dorsal interossei (FDI), with two referential electrodes attached at the wrist. Both muscles were shown to be involved in the to-be-observed hand movement (hand opening), although activations were more pronounced for the APB compared to the FDI muscle (see supplementary methods). Single-pulse transcranial magnetic stimulation (TMS), using a Magstim 200 stimulator (Magstim Company Ltd, UK) with a hand-held 70 mm figure-of-eight coil, was administered to locate the optimal scalp site for stimulating the primary motor cortex (M1) ("hotspotting"). The coil was positioned over the left hemisphere, tangentially to the scalp and 45° away from the midsagittal line, such that the induced current flow was in a posterior anterior direction, i.e. approximately perpendicular to the central sulcus. Optimal coil location for the experimental TMS-stimulation of M1 was determined as the site that produced maximal responses in the contralateral APB muscle while at rest. Although parameter setting procedures were prioritized for the APB, MEPs were simultaneously obtained from the APB and FDI muscles. Due to the overlap of hand muscle representations in M1, stimulation parameter settings are assumed to be satisfactorily effective for assessing condition-specific modulations simultaneously from both muscles (Facchini et al., 2002; Gertner and Classen, 2006; Krings et al., 1998; Scheiber, 1990). Next, the resting motor threshold (rMT) was defined for each participant as the lowest stimulation intensity that produced a peak-to-peak MEP of at least 50 µV in five out of ten consecutive trials (Rossini et al., 1994). During the experimental procedure, stimulation intensity was set at a suprathreshold of 130% of the subject's rMT (Alaerts, Swinnen and Wenderoth, 2009). Signal Software (version 2.02, Cambridge Electronic Design, UK) was used for EMG-recordings and triggering of the TMS-stimulator. EMG recordings were sampled at 2000 Hz via a CED Power 1401 unit (Cambridge Electronic Design, UK), amplified, band-pass filtered (5-1000 Hz) and stored on a PC for offline analysis.

2.1.4 Video stimuli

During TMS, video clips were presented to the observing participants. Video stimuli were identical to those used in a previous study by Wang, Newport, et al. (2011) and Wang, Ramsey and Hamilton (2011) in which an actor performed a head movement followed by a simple intransitive (i.e. not directed towards an object) hand movement (**figure 1**). At the onset of each clip, the actor was facing away from the camera with her eyes closed and her left hand static in front of her face. Then, the actor opened her eyes and turned her head either towards the camera, which resulted in direct gaze towards the observer, or away from the camera, providing averted gaze. Her hand remained static during the duration of the head movement. Subsequently, the actor performed a hand movement (i.e. opening of the

hand) or the hand remained static. This resulted in a 2 × 2 factorial design with the factors 'observed hand movement' (opening or static) and 'gaze direction' (direct or averted). An illustration of the different conditions of the factorial design is provided in **figure 1**. Each of the four conditions was presented five times in blocks of four five-second video clips (i.e. total of 20 trials per condition). Block presentation order was randomized across subjects and experimental blocks were randomly interleaved with four 'baseline' blocks in which only a blue background was shown. During movement observation, TMS pulses were delivered approximately 4.6 seconds after the start of the video clip which corresponded to the execution phase of the observed hand opening movement (see **figure 1**). Subjects' attention to the presented videos was randomly assessed between blocks by asking the subject to report the type of hand movement and gaze direction that was previously observed. In 92.5% of the assessments, subjects gave a correct response, ensuring attention to the presented videos. Video presentation timing was controlled by LabVIEW software (version 14.0, National Instruments, UK) and was triggered by the Signal Software for TMS-stimulation and EMG-recording.

Figure 1. Illustration of the experimental video clips. Participants were presented with a series of video clips of an actor performing a head movement to establish direct or averted gaze towards the observer, followed by a hand opening movement or no movement (static hand). This resulted in a 2×2 factorial design with the factors 'observed hand movement' (opening or static hand) and 'gaze direction' (direct or averted gaze).



2.1.5 Data analysis and statistics

Based on the recorded EMG data, peak-to-peak amplitudes of the TMS-evoked MEPs were determined. Additionally, background EMG was quantified by calculating the root mean square error (RMSE) across the 110 to 10 millisecond interval prior to TMS-stimulation. Since background EMG is known to modulate the size of MEP amplitudes (Devanne et al., 1997; Hess et al., 1987), peak-to-peak MEP amplitudes from trials with excessive background EMG (exceeding 2.5 standard deviations away from the mean) were discarded (2.42% of the trials for the APB, and 1.94% of the trials for the FDI). Further, MEP peak-to-peak amplitudes were considered as outliers and were removed from the analysis when they exceeded Q3 ± 1.5*(Q3-Q1), with Q1 and Q3 denoting the first and third quartile computed for each condition in each subject (Electronic Statistics Textbook, 2008, StatSoft). This resulted in an additional omission of 8.48% of the trials for the APB and 8.45% of the trials for the FDI. Note that the total number of discarded trials was similar across conditions (*F*(4,124) = 1.67, *p* = .16) and muscles (*F*(1,31) = 0.16, *p* = .70).

MEP peak-to-peak amplitudes and RMSE-scores were averaged separately for each condition. Due to high inter-individual differences in raw MEP responses, MEPs recorded during the four experimental conditions were normalized relative to baseline MEP responses separately for each subject (Halaki and Ginn, 2012). Shapiro-Wilk's W tests ensured a normal distribution of the MEP data for each condition. Repeated measures analyses of variance (ANOVAs) with the within-subject factors 'observed hand movement' (opening hand, static hand) and 'gaze direction' (direct gaze, averted gaze) were performed for each muscle (APB, FDI) separately on the normalized MEP amplitudes to explore whether corticomotor excitability of M1 is modulated by movement observation and/or eye contact. Normalized MEP amplitudes were entered in two repeated measures analyses of variance (ANOVAs), one for each muscle separately, with the within-subjects factors 'observed hand movement' (opening hand, static hand) and 'gaze direction' (direct gaze, averted gaze) to explore whether cortico-motor excitability of M1 is modulated by movement observation and/or eye contact. Fisher's least significant difference (LSD) contrasts were used for posthoc between-condition analyses. All statistics were calculated with Statistica 10 (StatSoft, USA) and results were considered significant with a *p*-value lower than .05.

2.2 Control experiment: Gaze behavior during movement observation

To explore whether gaze behavior and/or attention towards the presented hand movement was similar for the direct and averted eye gaze conditions, an additional eye tracking experiment was conducted while participants observed the four video clips of the main

experiment ('observed hand movement' (opening or static) × 'gaze direction' (direct or averted)).

2.2.1 Participants

Twenty-eight new subjects (20 males and 8 females), between the ages of 18 and 29 years participated in the additional eye tracking experiment to measure gaze behavior during observation of the video clips adopted in the main experiment. All participants provided signed written informed consents prior to the experiment and had normal or corrected-to-normal eye vision. Participants of the eye tracking experiment were not the same as those participating in the main movement observation TMS experiment.

2.2.2 Procedure

During the eye tracking session, the four video clips as described above were presented on a Tobii T120 binocular eye tracking device. The Tobii eye tracking system consists of a high-resolution camera embedded in a 17 inch TFT monitor (resolution: 1280×1024 pixels, sampling rate: 120 Hz, average precision: 0.5° of visual angle). Subjects were seated approximately 60 cm from the device. After a five-point calibration procedure, participants were instructed to naturally view the videos that were shown on the screen. Each of the four conditions was presented in one block of four five-second video clips (i.e. total of four trials per condition). An inter-block interval consisting of a black screen was shown for one second between blocks. The order of block presentation was randomized across subjects.

2.2.3 Data analysis and statistics

Two areas of interest (AOI) were defined for each video: the hand region and the eye region. These AOIs were defined using rectangular definition tools to mark the corresponding regions. Dependent measures included (i) the total fixation duration (TFD), which measures the sum of the duration for all fixations within an AOI; and (ii) fixation count (FC), which was calculated as the number of times the participant fixates the AOI (i.e. the number of times the participant's eye gaze enters and leaves the AOI). The gaze data was checked for outliers (none) and normality was assessed by means of Shapiro-Wilk's *W* tests (gaze data was sufficiently normally distributed). For each dependent variable (TFD, FC) and AOI (hand region, eye region), a repeated measures ANOVA with the within-subject factors 'observed hand movement' (opening hand, static hand) and 'gaze direction' (direct gaze, averted gaze) was conducted to examine gaze behavior for each condition.

3 Results

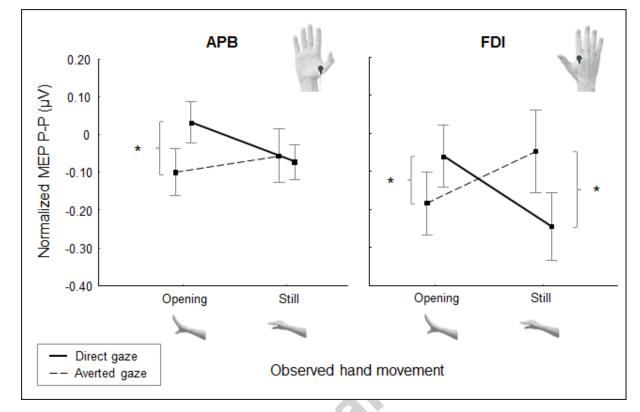
3.1 M1 facilitation during movement observation

A repeated measures ANOVA with the within-subject factors 'observed hand movement' (opening hand, static hand) and 'gaze direction' (direct gaze, averted gaze) was conducted on the normalized MEP data separately for each muscle (APB, FDI) to explore the effect of eye gaze on observation-induced facilitation of the primary motor cortex (M1). **Figure 2** displays the MEP amplitude data separately for each muscle and condition.

In both muscles, a two-way interaction between 'observed hand movement' and 'gaze direction' was revealed (tentatively in the APB: F(1,31) = 2.89; p = 0.06; $\eta^2 = .09$; *power* = .38; significantly in the FDI: F(1,31) = 7.07; p < .05, $\eta^2 = .19$; *power* = .73), indicating a differential impact of eye gaze on observation-induced M1 facilitation. Direct exploration of the difference in MEP response between direct and averted eye gaze showed that for observing the hand opening movement, MEP responses of the APB and FDI were significantly higher for the direct eye gaze condition compared to the averted eye gaze condition (Fisher LSD: both p < .05; **figure 2**, left panel). During observation of the static hand condition, MEP responses in the APB muscle were not significantly different between the direct and averted eye gaze conditions (p = .75), whereas in the FDI muscle, M1 facilitation was reversibly modulated, indicating significantly *lower* MEP responses when the static hand observation was accompanied by direct versus averted eye gaze (p < .001; **figure 2**, right panel). No main effects of 'gaze direction' (all p > .29) or 'observed hand movement' (all p > .48) were revealed in either muscle.

MEP scores were not confounded by modulations in background EMG scores. This was tested by conducting similar ANOVAs to the corresponding background EMG data (i.e. normalized RMSE-scores). Background EMG was generally small and condition-specific modulations were minimal, as no significant main or interaction effects were revealed. For all conditions and muscles, background EMG scores are listed in **supplementary table 1**.

Figure 2. Normalized MEP peak-to-peak amplitude data for the APB and FDI during action observation, separately for each condition. Vertical bars denote standard errors of the mean. * Denotes a significant difference between conditions (p < .05).



3.2 Link with social responsiveness

Participants reported a mean total SRS-A score of 36.15 (SD = 19.74), which, as a group, is well below the cut-off score of 54 signaling impairments in social responsiveness. However, for five out of the 32 participants a total score higher than the cut-off point was reported, indicative of impairments in terms of social responsiveness. To explore whether interindividual differences in social responsiveness were related to the extent by which direct gaze elicited higher MEP responses compared to averted gaze during movement observation, a regression analysis was conducted with 'gaze effect' (difference in MEP response between direct and averted eye gaze conditions) as dependent variable and 'social responsiveness (sub)score' as predictor (across muscles). Overall, *beta*-values were generally small and none of the relationships reached significance (all p > .25, see **supplementary table 2**) indicating that social responsiveness was not predictive for the extent of M1 facilitation.

3.3 Gaze behavior during movement observation

An additional eye tracking experiment was conducted to explore whether the observer's gaze behavior was different when the movement observation conditions (opening hand, static

hand) were accompanied by direct versus averted gaze. To do so, eye tracking was performed and the total fixation duration (TFD) and fixation count (FC) were determined in an area-of-interest (AOI) centered over the hand and eye region of the presented video clip.

For the TFD, a repeated-measures ANOVA with the within-subject factors 'observed hand movement' (opening hand, static hand) and 'gaze direction' (direct, averted) revealed no significant main effect of 'gaze direction', indicating that across movement observation conditions, participants fixated an equal amount of time towards the hand region during direct as during averted gaze conditions (F(1,27) = 0.79, p = .38, $\eta^2 = .03$, *power* = .14). Similarly, no significant effect of 'gaze direction' was revealed for the FC data, indicating that participants made a comparable amount of saccades towards the hand region for the direct and averted gaze conditions (F(1,27) = 0.75, p = .39, $\eta^2 = .03$, *power* = .13). Note however, that the ANOVA revealed significant main effects of 'observed hand movement', indicating that irrespective of gaze direction (direct, averted), subjects looked significantly more (FC: F(1,27) = 21.24, p < .0001, $\eta^2 = .44$, *power* = .99) and longer (TFD: F(1,27) = 26.62, p < .0001, $\eta^2 = .50$; *power* = .99) towards the hand region when the actor performed the opening hand movement, compared to when the actor's hand remained static.

The eye tracking data were additionally used to explore whether direct versus averted eye gaze differentially modulated the observer's gaze behavior towards the eye region of the actor. Not surprisingly, for the eye region AOI, a significant main effect of 'gaze direction' was revealed, indicating that across hand movements, subjects looked longer towards the eye region during direct gaze than during averted gaze (i.e. indicative of the establishment of eye contact between the actor and the observer during direct gaze conditions) (TFD: F(1,27) = 8.73, p < .01, $\eta^2 = .24$; *power* = .81). In terms of FC, participants made a comparable amount of saccades toward the eye region during direct gaze as during averted gaze (F(1,27) = 0.06, p = .81, $\eta^2 = .002$, *power* = .06). Mean TFD and FC values are displayed separately for each AOI and condition in **supplementary table 3**.

4 Discussion

In the present study, transcranial magnetic stimulation (TMS) was used to assess corticomotor excitability at the level of the primary motor cortex (M1) during movement observation, and, in particular, whether observation-induced facilitation of M1 is altered when accompanied by direct or averted eye gaze. Overall, our results show that observationinduced M1 facilitation was most pronounced when direct eye gaze was observed, indicating that eye gaze forms a salient social cue that can modulate the extent by which others' actions are 'mapped' onto the observer's motor system.

As such, the present TMS study extends previous findings from behavioral studies by Wang, Newport, et al. (2011) and Wang, Ramsey, et al. (2011) studying the effect of eye gaze on automatic motor mimicry using similar video clips as those adopted in the present study. In these studies, a stimulus-response compatibility paradigm was adopted where participants were asked to perform the same movement or the opposite movement as viewed in the video clip, and a clear congruency effect was found indicating that responses were significantly faster when the same movement was performed (e.g. hand opening observed hand opening performed), compared to trials in which the opposite hand movement was performed (e.g. hand closing observed - hand opening performed). Interestingly, Wang, Newport, et al. (2011) demonstrated that this mimicry congruency effect was even more enlarged when direct eye contact was established between the observer and the observed actor, indicating a rapid modulation of mimicry by eye contact. Our study provides insights into the neurophysiological mechanism underlying this modulating effect of eye gaze on automatic motor simulation, by showing that direct eye gaze can significantly enhance the extent by which the observed movement is mirrored onto the observer's motor system. Our findings are also in agreement with previous results from an MEG study by Kilner et al. (2006). In this study, MEG was used to record cortico-motor activity whilst participants observed upper limb movements of an actor that was facing away or towards them. Results from this study showed that cortico-motor responses to movement observation (oscillatory activity in the 7-12 Hz frequency range) are dependent on the relative perspective of the observed model towards the observer, such that cortico-motor modulations to movement observation were only present when the actor was facing towards the observer, not when the actor was facing away from the observer. The authors suggested that distinct 'social' signals accompanying the observed movements and actions of other people (e.g. the perspective relative to the observer) can modulate visuospatial attention, such that only specific visual information of the most salient and most socially relevant actions is allowed to enter the mirror system for further processing. Our study extends these findings by showing that not the perspective per se, but the establishment of direct eye gaze may be a highly salient cue

in determining the extent by which an observed action will be mapped onto the observer's motor system. In addition to perceived eye contact, also other social signals may sophistically direct motor resonance (Wang and Hamilton, 2012). Indeed, previous studies have shown that also social cues such as self-construal (Obhi et al., 2011), social interaction (Hogeveen and Obhi, 2012) and power (Hogeveen et al., 2014) can influence motor resonance. It has therefore been argued that the control of motor resonance may involve a 'social top-down response modulation' (STORM) that is dependent on the social context in which others' actions are observed (Wang and Hamilton, 2012).

Previous fMRI studies in humans (Kampe et al., 2003; Nummenmaa and Calder, 2009; Senju and Johnson, 2009b) and single-cell recordings in monkeys (Emery, 2000; Perrett et al., 1992) consistently showed that the medial prefrontal cortex (mPFC) and the superior temporal sulcus (STS) are increasingly activated during direct versus averted gaze, highlighting the importance of these two regions in gaze processing. Wang, Ramsey, et al. (2011) replicated these findings and additionally suggested that the mPFC may well be the originator of the effect of eye gaze on motor mimicry by modulating functional connectivity with the STS, i.e. the main visual input region to the fronto-parietal mirror system. In other words, the model by Wang, Ramsey, et al. (2011) suggested that gaze-related activations at the level of mPFC may impose a top-down control over the processing of visuo-motor information at the level of the STS, which in turn may impact the extent by which observed actions are processed in down-stream mirror regions in the inferior frontal gyrus, ventral premotor regions and inferior parietal lobule. In this view, M1 may be conceived as the endstate region of a chain of cortico-cortical connections signaling on whether or not the corticospinal tract and the corresponding peripheral muscles are to be recruited for initiating overt motor simulation. Correspondingly, by receiving direct input from upstream premotor and mirror regions, modulations in cortico-motor excitability at the level of M1 may reflect an endstate cortical measure of how the brain 'evaluated' the social relevance or saliency of the observed visual scene. Our results therefore provide additional support to the notion that direct eye gaze from the actor forms a strong mediator for evaluating whether or not visuomotor information of the observed action is sufficiently relevant to be processed up to the level of M1. Instead of simulating all possible movement-related information perceived in a visual scene, eye contact may direct the motor system to give preference to processing visuo-motor input originating from the most socially relevant person. Note that in the FDI muscle, but not in the APB, we found an inverse effect of eye gaze on M1 facilitation during the observation of the static hand, indicating increased M1 excitability for the averted compared to the direct gaze condition. One potential interpretation could be that during the trials in which no actual hand opening movement was observed (only a static hand), direct

eye gaze might have induced an increased inhibitory effect on M1 excitability to encode more efficiently that *no* movement is observed during these trials. From this perspective, it can be hypothesized that direct eye primes the observation-to-execution mapping system by increasing the signal-to-noise ratio when perceiving motion stimuli (i.e., by effectively heightening M1 excitability during actual movement observation, and inhibiting M1 excitability when no movement is observed). This interpretation remains speculative however as it is unclear why this effect was then only significantly observed in the FDI muscle and not in the APB muscle.

Nummenmaa and Calder (2009) showed that observing another person's gaze can automatically induce gaze following, thereby shifting spatial attention toward the scene observed by the model. In this view, an alternative explanation for the observed gaze effect of the present study can be put forward, suggesting that the averted gaze conditions induced a shift of spatial attention away from the observed hand movement, thereby reducing the observation-induced M1 facilitation. We explicitly addressed this alternative explanation in a control eye-tracking experiment, in which the same video clips were displayed while the participants' gaze behavior was recorded. Overall, we found no indications that participants spend less time fixating on the to-be-observed hand movement in video clips with averted gaze compared to direct gaze, which makes it implausible that differences in visual spatial attention are responsible for the encountered gaze-related modulations in observationinduced M1 excitability. Also Wang, Newport, et al. (2011) explicitly addressed this issue in their control "flashbox" experiment in which distracting stimuli (i.e. flashing white squares) were displayed in the periphery during the movie clip to draw participant's attention. These manipulations did however not alter the effects of eye gaze on motor mimicry, indicating the robustness of the eye gaze effect relative to the presence of distracters competing for attention. Furthermore, previous studies have shown that eye gaze can have a direct effect on several physiological measures such as skin conductance, indicating a heightened response of the observer's autonomic nervous system during direct versus averted eye gaze (Hietanen et al., 2008; Pönkänen et al., 2011). While direct eye gaze conditions may have induced a similar heightening of arousal in the present study, it is unlikely that these arousal effects directly affected the modulation of MEP responses, since enhancements in M1 facilitation were specifically observed during the movement observation condition (i.e. hand opening) and not during the observation of the static hand. Also background EMG scores which were measured before the TMS pulse - did not show any condition specific modulations.

Nevertheless, future research on the effect of eye gaze on motor facilitation would benefit from the inclusion of parallel assessments of skin conductance to explore the possibility of arousal-related effects further. Also in terms of the assessment of viewing behavior and attention to the presented stimuli, future studies would benefit from the inclusion of online eye tracking. In the present study, assessments of viewing behavior were only assessed in a separate eye tracking experiment in which the participants were different from those participating in the main experiment. While this additional eye tracking experiment already provided relevant information related to viewing behavior towards the presented stimuli, online eye tracking would have allowed a direct assessment of viewing behavior and attention on a trial-by-trial basis and its potential relationship to the evoked MEP responses. In the present TMS experiment, attention to the presented stimuli was only assessed randomly, by asking the participants to verbally report the type of hand movement and gaze direction that was observed in the previous video clip. Since this assessment might have affected the subjects' explicit awareness of the presented experimental manipulations, it should be beneficial for future experiments to adopt other strategies to assess the subjects' online attention to the presented stimuli, such as the inclusion of eye tracking.

Overall, mirror motor mapping is hypothesized to form the basic neural mechanism by which others' actions and emotional states can be simulated, recognized and imitated. Considering the hypothesized link between motor simulation and these high-level social skills, the 'broken mirror theory of autism' has been put forward, postulating that behavioral deficits in action understanding, imitation and empathy seen in Autism Spectrum Disorders (ASD) may result from aberrant functioning of the fronto-parietal mirror motor mapping (Ramachandran and Oberman, 2016). While a number of neurophysiological studies provided support for aberrant 'mirroring' in ASD (Dapretto et al., 2006; Enticott et al., 2012; Hadjikhani et al., 2006; Oberman et al., 2005), also several studies found no evidence to support this account (for a review, see Hamilton, 2013). Related to these controversies, and considering that eye contact may form a highly salient cue and perhaps even a prerequisite for the initiation of motor simulation, it would be interesting for future research to evaluate whether and how gaze-related effects on motor mirroring are affected in patients with ASD, who are wellknown to display particular difficulties with engaging mutual eye contact (Kaartinen et al., 2012; Kylliäinen and Hietanen, 2006; Kylliäinen et al., 2012; Senju and Johnson, 2009a). Note that in the present study, we found no significant relationships between scores on the social responsiveness scale (SRS) and the extent of the 'gaze effect'. However, considering that inter-individual differences in SRS scores were generally small in our rather homogenous sample of neurotypical students, it should be interesting for future studies to explore the relationship between the eye gaze effect and social responsiveness in more

heterogeneous samples including individuals with particular implications in the social domain, such as ASD or social anxiety disorders (Myllyneva et al., 2015; Wieser et al., 2009).

Further, considering that based on the present sample, uncertainty exists with regard to the behavioral correlates of the observed gaze effect on motor mirroring, it should be interesting for future studies to explore whether - in addition to social competence - potentially also other personality traits may be informative in predicting inter-individual variations in the observed gaze effect. For example, based on the work by Hietanen et al. (2008), several links have been suggested between seeing direct or averted eye gaze and inter-individual differences in the motivational system towards approach and avoidance. In particular, neuroticism and scales assessing social phobia have been suggested to form important predictors of behavioral direct gaze avoidance and subjective averted gaze preference (Myllyneva et al., 2015; Uusberg et al., 2015). In this view, it should also be interesting for future research to discern whether inter-individual differences in these motivational preferences towards eye contact may be important in determining the facilitative effect of eye gaze on motor mirroring.

To sum up, the present results provide evidence that the mapping of others' movements onto the observer's motor system is enhanced when direct compared to averted eye gaze is established between the observer and the observed model. These findings support the notion that eye contact is a powerful and highly salient social signal with the ability to modify activity in the human mirror-motor system, thereby directing human social interactions.

5 Funding

This research was supported by grants from the Flanders Fund for Scientific Research (FWO [KAN 1506716N, KAN 1521313N and G.0401.12] and the Branco Weiss fellowship of the Society in Science - ETH Zurich granted to K.A. J.P. is supported by an internal fund of the KU Leuven [STG/14/001].

6 Acknowledgments

The authors would like to thank Jaana van Overwalle, Michaël Doumen and Stephanie Brams for their assistance in data collection and analysis. We would also like to thank Prof. dr. Antonia Hamilton for her contribution in inspiring the design of the present study.

References

- Alaerts, K., de Beukelaar, T. T., Swinnen, S. P., & Wenderoth, N. (2012). Observing how others lift light or heavy objects: Time-dependent encoding of grip force in the primary motor cortex. *Psychological Research*, *76*(4), 503-531. doi:10.1007/s00426-011-0380-1
- Alaerts, K., Heremans, E., Swinnen, S. P., & Wenderoth, N. (2009). How are observed actions mapped to the observer's motor system? Influence of posture and perspective. *Neuropsychologia*, 47(2), 415–422. doi:10.1016/j.neuropsychologia.2008.09.012
- Alaerts, K., Senot, P., Swinnen, S. P., Craighero, L., Wenderoth, N., & Fadiga, L. (2010). Force requirements of observed object lifting are encoded by the observer's motor system: A TMS study. *European Journal of Neuroscience*, *31*(6), 1144–1153. doi:10.1111/j.1460-9568.2010.07124.x
- Alaerts, K., Swinnen, S. P., & Wenderoth, N. (2009). Is the human primary motor cortex activated by muscular or direction-dependent features of observed movements? *Cortex*, *45*(10), 1148–1155. doi:10.1016/j.cortex.2008.10.005
- Alaerts, K., Swinnen, S. P., & Wenderoth, N. (2010). Observing how others lift light or heavy objects: Which visual cues mediate the encoding of muscular force in the primary motor cortex? *Neuropsychologia*, 48(7), 2082-2090. doi:10.1016/j.neuropsychologia.2010.03.029
- Buccino, G., Binkofski, F., Fink, G. R., Fadiga, L., Fogassi, L., Gallese, V., ... Freund, H.-J. (2001). Action observation activates premotor and parietal areas in a somatotopic manner: An fMRI study. *European Journal of Neuroscience*, *13*(2), 400–404. doi:10.1111/j.1460-9568.2001.01385.x
- Caramazza, A., Anzellotti, S., Strnad, L., & Lingau, A. (2014). Embodied cognition and mirror neurons: A critical assessment. *Annual Review of Neuroscience*, 37, 1-15. doi: 10.1146/annurev-neuro-071013-013950.
- Chong, T. T.-J., Cunnington, R., Williams, M. A., Kanwisher, N., & Mattingley, J. B. (2008). fMRI adaptation reveals mirror neurons in human inferior parietal cortex. *Current Biology*, *18*(20), 1576–1580. doi:10.1016/j.cub.2008.08.068
- Constantino, J. N., & Todd, R. D. (2005). Intergenerational transmission of subthreshold autistic traits in the general population. *Biological Psychiatry*, 57(6), 655–660. doi:10.1016/j.biopsych.2004.12.014
- Dapretto, M., Davies, M. S., Pfeifer, J. H., Scott, A. A., Sigman, M., Bookheimer, S. Y., &

Iacoboni, M. (2006). Understanding emotions in others: mirror neuron dysfunction in children with autism spectrum disorders. *Nature Neuroscience, 9,* 28–30. doi:10.1038/nn1611

- de Beukelaar, T. T., Alaerts, K., Swinnen, S. P., & Wenderoth, N. (2015). Motor facilitation during action observation: The role of M1 and PMv in grasp predictions. *Cortex*, 75, 180–192. doi:10.1016/j.cortex.2015.11.009
- Devanne, H., Lavoie, B. A., & Capaday, C. (1997). Input–output properties and gain changes in the human corticospinal pathway. *Experimental Brain Research*, *114*(2), 329–338.
- di Pellegrino, G., Fadiga, L., Fogassi, L., Gallese, V., & Rizzolatti, G. (1992). Understanding motor events: A neurophysiological study. *Experimental Brain Research*, *91*(1), 176– 180. doi:10.1007/BF00230027
- Emery, N. J. (2000). The eyes have it: the neuroethology, function and evolution of social gaze. *Neuroscience and Biobehavioral Reviews*, *24*(6), 581–604. doi:10.1016/S0149-7634(00)00025-7
- Enticott, P. G., Kennedy, H. A., Rinehart, N. J., Tonge, B. J., Bradshaw, J. L., Taffe, J. R., Daskalakis, Z. J., & Fitzgerald, P. B. (2012). Mirror neuron activity associated with social impairments but not age in autism spectrum disorder. *Biological Psychiatry*, *71*, 427–33. doi:10.1016/j.biopsych.2011.09.001
- Facchini, S., Romani, M., Tinazzi, M., & Agliotti, S. M. (2002). Time-related changes of excitability of the human motor system contingent upon immobilisation of the ring and little fingers. *Clinical Neurophysiology*, *113*(3), 367-375.
- Fadiga, L., Craighero, L., & Olivier, E. (2005). Human motor cortex excitability during the perception of others' action. *Current Opinion in Neurobiology*, 15(2), 213–218. doi:10.1016/j.conb.2005.03.013
- Fadiga, L., Fogassi, L., Pavesi, G., & Rizzolatti, G. (1995). Motor facilitation during action observation: a magnetic stimulation study. *Journal of Neurophysiology*, 73(6), 2608– 2611.
- Gallese, V. (2009). Motor abstraction: A neuroscientific account of how action goals and intentions are mapped and understood. *Psychological Research*, *73*(4), 486-498.
- Gallese, V., Fadiga, L., Fogassi, L., & Rizzolatti, G. (1996). Action recognition in the premotor cortex. *Brain*, *119*(2), 593–609. doi:10.1093/brain/119.2.593
- Gangitano, M., Mottaghy, F. M., & Pascual-Leone, A. (2001). Phase-specific modulation of cortical motor output during movement observation. *NeuroReport*, *12*(7), 1489–1492.
- Gertner, R., & Classen, J. (2006). Modular organization of finger movements by the human central nervous system. *Neuron*, *52*(4), 731-742.
- Grafton, S. T., Arbib, M. A., Fadiga, L., & Rizzolatti, G. (1996). Localization of grasp representations in humans by positron emission tomography. *Experimental Brain*

Research, 112(1), 103–111. doi:10.1007/BF00227183

- Grèzes, J., Fonlupt, P., Bertenthal, B., Delon-Martin, C., Segebarth, C., & Decety, J. (2001).
 Does perception of biological motion rely on specific brain regions? *NeuroImage*, *13*(5), 775–785. doi:10.1006/nimg.2000.0740
- Grossman, E. D., & Blake, R. (2002). Brain areas active during visual perception of biological motion. *Neuron*, *35*(6), 1167–1175. doi:10.1016/S0896-6273(02)00897-8
- Grossman, E., Donnelly, M., Price, R., Pickens, D., Morgan, V., Neighbor, G., & Blake, R. (2000). Brain areas involved in perception of biological motion. *Journal of Cognitive Neuroscience*, *12*(5), 711–720. doi:10.1162/089892900562417
- Hadjikhani, N., Joseph, R. M., Snyder, J., & Tager-Flusberg, H. (2006). Anatomical differences in the mirror neuron system and social cognition network in autism. *Cerebral Cortex, 16*, 1276–1282. doi:10.1093/cercor/bhj069
- Halaki, M., & Ginn, K. (2012). Normalization of EMG signals: To normalize or not to normalize and what to normalize to? In G.R. Naik (Ed.), *Computational Intelligence in Electromyography Analysis – A Perspective on Current Applications and Future Challenges* (pp. 175-194). InTech. doi: 10.5772/49957
- Hamilton, A.F. de C. (2013). Reflecting on the mirror neuron system in autism: A systematic review of current theories. *Developmental Cognitive Neuroscience*, 3, 91–105. doi:10.1016/j.dcn.2012.09.008
- Hess, C. W., Mills, K. R., & Murray, N. M. (1987). Responses in small hand muscles from magnetic stimulation of the human brain. *Journal of Physiology, 388*, 397–419.
- Hietanen, J. K., Leppänen, J. M., Peltola, M. J., Linna-Aho, K., & Ruuhiala, H. J. (2008).
 Seeing direct and averted gaze activates the approach-avoidance motivational brain systems. *Neuropsychologia*, 46(9), 2423–2430. doi:10.1016/j.neuropsychologia.2008.02.029
- Hogeveen, J., Inzlicht, M., & Obhi, S. S. (2014). Power changes how the brain responds to others. *Journal of Experimental Psychology*, *143*, 755–762.
- Hogeveen, J., & Obhi, S. S. (2012). Social interaction enhances motor resonance for observed human actions. *The Journal of Neuroscience*, 32(17), 5984–5989. doi:10.1523/JNEUROSCI.5938-11.2012
- Iacoboni, M. (2009). Imitation, empathy, and mirror neurons. *Annual Review of Psychology*, *60*, 653–70. doi:10.1146/annurev.psych.60.110707.163604
- Iacoboni, M., Molnar-Szakacs, I., Gallese, V., Buccino, G., Mazziotta, J. C., & Rizzolatti, G. (2005). Grasping the intentions of others with one's own mirror neuron system. *PLoS Biology*, *3*(3), e79. doi:10.1371/journal.pbio.0030079
- Iacoboni, M., Woods, R. P., Brass, M., Bekkering, H., Mazziotta, J. C., & Rizzolatti, G. (1999). Cortical mechanisms of human imitation. *Science*, *286*(5449), 2526–2528.

doi:10.1126/science.286.5449.2526

- Kaartinen, M., Puura, K., Mäkelä, T., Rannisto, M., Lemponen, R., Helminen, M., ... Hietanen, J. K. (2012). Autonomic arousal to direct gaze correlates with social impairments among children with ASD. *Journal of Autism and Developmental Disorders*, 42(9), 1917–1927. doi:10.1007/s10803-011-1435-2
- Kampe, K. K. W., Frith, C. D., & Frith, U. (2003). "Hey John": Signals conveying communicative intention toward the self activate brain regions associated with "mentalizing", regardless of modality. *Journal of Neuroscience*, 23(12), 5258–5263.
- Kilner, J. M., Marchant, J. L., & Frith, C. D. (2006). Modulation of the mirror system by social relevance. Social Cognitive and Affective Neuroscience, 1(2), 143–148. doi:10.1093/scan/nsl017
- Kilner, J. M., Neal, A., Weiskopf, N., Friston, K. J., & Frith, C. D. (2009). Evidence of mirror neurons in human inferior frontal gyrus. *The Journal of Neuroscience*, 29(32), 10153– 10159. doi:10.1523/JNEUROSCI.2668-09.2009
- Krings, T., Naujokat, C., Graf v. Keyserlingk, D. (1998). Representation of cortical motor function as revealed by stereotactic transcranial magnetic stimulation. *Electroencephalography and Clinical Neurophysiology*, *102*(2), 85-93.
- Kylliäinen, A., & Hietanen, J. K. (2006). Skin conductance responses to another person's gaze in children with autism. *Journal of Autism and Developmental Disorders*, *36*(4), 517–525. doi:10.1007/s10803-006-0091-4
- Kylliäinen, A., Wallace, S., Coutanche, M. N., Leppänen, J. M., Cusack, J., Bailey, A. J., & Hietanen, J. K. (2012). Affective-motivational brain responses to direct gaze in children with autism spectrum disorder. *Journal of Child Psychology and Psychiatry*, *53*(7), 790– 797. doi:10.1111/j.1469-7610.2011.02522.x
- Maeda, F., Chang, V. Y., Mazziotta, J., & Iacoboni, M. (2001). Experience-dependent modulation of motor corticospinal excitability during action observation. *Experimental Brain Research*, 140(2), 241–244. doi:10.1007/s002210100827
- Maeda, F., Kleiner-Fisman, G., & Pascual-Leone, A. (2002). Motor facilitation while observing hand actions: Specificity of the effect and role of observer's orientation. *Journal of Neurophysiology*, 87(3), 1329–1335.
- Mahon, B. Z. (2015). What is embodied about cognition? *Language, Cognition and Neuroscience, 30*(4), 420-429. doi: 10.1080/23273798.2014.987791
- Myllyneva, A., Ranta, K., & Hietanen, J. K. (2015). Psychophysiological responses to eye contact in adolescents with social anxiety disorder. *Biological Psychology*, 109, 151– 158. doi:10.1016/j.biopsycho.2015.05.005
- Noens, I., De la Marche, W., & Scholte, E. (2012). SRS-A Screeningslijst voor autismespectrumstoornissen bij volwassenen. Handleiding [SRS-A Screening tool for

autism spectrum disorders in adults. A manual]. Amsterdam, The Netherlands: Hogrefe Uitgevers.

- Nummenmaa, L., & Calder, A. J. (2009). Neural mechanisms of social attention. *Trends in Cognitive Sciences*, *13*(3), 135–143. doi:10.1016/j.tics.2008.12.006
- Oberman, L. M., Hubbard, E. M., McCleery, J. P., Altschuler, E. L., Ramachandran, V. S., & Pineda, J. A. (2005). EEG evidence for mirror neuron dysfunction in autism spectrum disorders. *Cognitive Brain Research, 24*(2), 190–198. doi:10.1016/j.cogbrainres.2005.01.014
- Obhi, S. S., Hogeveen, J., & Pascual-Leone, A. (2011). Resonating with others: the effects of self-construal type on motor cortical output. *The Journal of Neuroscience*, 31(41), 14531–14535. doi:10.1523/JNEUROSCI.3186-11.2011
- Oldfield, R. C. (1971). The assessment and analysis of handedness: The Edinburgh inventory. *Neuropsychologia*, *9*(1), 97–113. doi:10.1016/0028-3932(71)90067-4
- Pageler, N. M., Menon, V., Merin, N. M., Eliez, S., Brown, W. E., & Reiss, A. L. (2003). Effect of head orientation on gaze processing in fusiform gyrus and superior temporal sulcus. *NeuroImage*, 20(1), 318–329. doi:10.1016/S1053-8119(03)00229-5
- Pelphrey, K. A., Viola, R. J., & McCarthy, G. (2004). When strangers pass: Processing of mutual and averted social gaze in the superior temporal sulcus. *Psychological Science*, *15*(9), 598–603. doi:10.1111/j.0956-7976.2004.00726.x
- Perrett, D. I., Hietanen, J. K., Oram, M. W., & Benson, P. J. (1992). Organization and functions of cells responsive to faces in the temporal cortex. *Philosophical Transactions* of the Royal Society of London. Series B, Biological Sciences, 335(1273), 23–30. doi:10.1098/rstb.1992.0003
- Pönkänen, L. M., Peltola, M. J., & Hietanen, J. K. (2011). The observer observed: frontal EEG asymmetry and autonomic responses differentiate between another person's direct and averted gaze when the face is seen live. *International Journal of Psychophysiology*, 82(2), 180–7. doi:10.1016/j.ijpsycho.2011.08.006
- Ramachandran, V. S., & Oberman, L. M. (2006). Broken mirrors: A theory of autism. *Scientific American*, 295(5), 62–69. doi:10.1038/scientificamerican1106-62
- Rizzolatti, G., Camarda, R., Fogassi, L., Gentilucci, M., Luppino, G., & Matelli, M. (1988). Functional organization of inferior area 6 in the macaque monkey. *Experimental Brain Research*, 71(3), 491–507. doi:10.1007/BF00248742
- Rizzolatti, G., & Craighero, L. (2004). The mirror neuron system. *Annual Review of Neuroscience*, 27, 169–192. doi:10.1146/annurev.neuro.27.070203.144230
- Rizzolatti, G., & Fabbri-Destro, M. (2008). The mirror system and its role in social cognition. *Current Opinion in Neurobiology*, *18*(2), 179–184. doi:10.1016/j.conb.2008.08.001
- Rizzolatti, G., Fadiga, L., Gallese, V., & Fogassi, L. (1996). Premotor cortex and the

recognition of motor actions. *Cognitive Brain Research*, *3*(2), 131–141. doi:10.1016/0926-6410(95)00038-0

- Rizzolatti, G., Fadiga, L., Matelli, M., Bettinardi, V., Paulesu, E., Perani, D., & Fazio, F. (1996). Localization of grasp representations in humans by PET: 1. Observation versus execution. *Experimental Brain Research*, *111*(2), 246–252. doi:10.1007/BF00227301
- Rossini, P. M., Barker, A. T., Berardelli, A., Caramia, M. D., Caruso, G., Cracco, R. Q., ...
 Tomberg, C. (1994). Non-invasive electrical and magnetic stimulation of the brain, spinal cord and roots: basic principles and procedures for routine clinical application.
 Report of an IFCN committee. *Electroencephalography and Clinical Neurophysiology*, *91*(2), 79–92. doi:10.1016/0013-4694(94)90029-9
- Scheiber, M. H. (1990). How might the motor cortex individuate movements. *Trends in Neurosciences*, *13*, 440–445.
- Senju, A., & Johnson, M. H. (2009a). Atypical eye contact in autism: Models, mechanisms and development. *Neuroscience and Biobehavioral Reviews*, 33(8), 1204–1214. doi:10.1016/j.neubiorev.2009.06.001
- Senju, A., & Johnson, M. H. (2009b). The eye contact effect: Mechanisms and development. *Trends in Cognitive Sciences*, *13*(3), 127–134. doi:10.1016/j.tics.2008.11.009
- Strafella, A. P., & Paus, T. (2000). Modulation of cortical excitability during action observation: A transcranial magnetic stimulation study. *NeuroReport*, *11*(10), 2289– 2292. doi:doi.org/10.1097/00001756-200007140-00044
- Uusberg, H., Allik, J., & Hietanen, J. (2015). Eye contact reveals a relationship between Neuroticism and anterior EEG asymmetry. *Neuropsychologia*, *76*, 161-168.
- Wang, Y., & Hamilton, A. F. de C. (2012). Social top-down response modulation (STORM): a model of the control of mimicry in social interaction. *Frontiers in Human Neuroscience*, 6, 1–10. doi:10.3389/fnhum.2012.00153
- Wang, Y., Newport, R., & Hamilton, A. F. de C. (2011). Eye contact enhances mimicry of intransitive hand movements. *Biology Letters*, *7*(1), 7–10. doi:10.1098/rsbl.2010.0279
- Wang, Y., Ramsey, R., & Hamilton, A. F. de C. (2011). The control of mimicry by eye contact is mediated by medial prefrontal cortex. *Journal of Neuroscience*, *31*(33), 12001–12010. doi:10.1523/JNEUROSCI.0845-11.2011
- Wieser, M. J., Pauli, P., Alpers, G. W., & Mühlberger, A. (2009). Is eye to eye contact really threatening and avoided in social anxiety? An eye-tracking and psychophysiology study. *Journal of Anxiety Disorders*, 23(1), 93–103. doi:10.1016/j.janxdis.2008.04.004

Highlights

- TMS was used to assess gaze-induced modulations in M1 excitability during movement • observation.
- M1 facilitation occurred mostly when movements in combination with direct eye gaze • were observed.
- Direct eye gaze can modulate excitability in the observer's motor system. ٠

.....