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TMS demonstrates that both right and left superior temporal sulci are important for facial expression recognition

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Prior studies demonstrate that a face-responsive region in the posterior superior temporal sulcus (pSTS) is involved in facial expression recognition. Although this region can be identified in both hemispheres, studies more commonly report it in the right hemisphere. However, the extent to which expression recognition is lateralised in pSTS remains unclear. In the current study, we used transcranial magnetic stimulation (TMS) to systematically compare the causal contribution of the right pSTS (rpSTS) with the left pSTS (lpSTS) during facial expression recognition. TMS was delivered over the functionally localised rpSTS, lpSTS and the control vertex site while participants (N = 30) performed an expression matching task and a control object matching task. TMS delivered over the rpSTS impaired expression recognition more than TMS delivered over the lpSTS. Crucially, TMS delivered over the rpSTS and lpSTS impaired task performance more than TMS delivered over the control site. TMS had no effect on the control task. This causally demonstrates that while task disruption was greater in the rpSTS, both the rpSTS and the lpSTS were engaged in facial expression recognition. Our results indicate that cognitive functions that are seemingly lateralised in neuroimaging studies, still rely on computations performed in both hemispheres for optimum task performance.

1. Introduction

Recognising facial expressions is an important aspect of human social interaction. Expressions provide us with one of the richest sources of information about another person's emotional state. Models of face processing propose that expressions are computed in an anatomically distributed, and highly interacting network in the human brain. Two components in this network are located in the bilateral posterior superior temporal sulcus (pSTS) (Haxby et al., 2000, 2002; Palermo and Rhodes, 2007; Vuilleumier et al., 2004). While the functional contribution of the right pSTS (rpSTS) to expression recognition has been extensively investigated (Allison et al., 2000; Andrews and Ewbank, 2004; Phillips et al., 1998; Pitcher, 2014; Winston, O'Doherty and Dolan, 2003), the functional involvement of the lpSTS in expression recognition is less clear. In addition, the extent to which expression recognition is lateralised in the pSTS is unknown. In the current study, we used functional magnetic resonance imaging (fMRI) and transcranial magnetic stimulation (TMS) to systematically investigate the causal contribution of the right and left pSTS to expression recognition.

Models of face processing propose that the human STS is a core node of a distributed face network (Adolphs, 2002; Calder and Young, 2005;

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Haxby et al., 2000). Evidence from a range of experimental techniques demonstrates that the posterior region of the right STS is involved in expression recognition. For example, single-cell recordings in patients undergoing craniotomy showed that expression identification evoked changes in neuronal activity in the rpSTS (Ojemann et al., 1992). Direct electrical stimulation to the rpSTS also impaired patients' ability to identify facial expressions, demonstrating that the region is causally engaged in expression recognition (Fried et al., 1982). Similarly, expression recognition impairments have been reported in patients with lesions to the rpSTS (Fox et al., 2011; Rapcsak et al., 1989) and when TMS was delivered over the rpSTS of healthy participants (Pitcher, 2014). Many neuroimaging studies have also demonstrated an increased response in the pSTS during expression recognition tasks (e.g., Andrews and Ewbank, 2004; Engell and Haxby, 2007; Gur et al., 1994; Narumoto et al., 2001; Phillips et al., 1998; Streit et al., 1999; Winston et al., 2004; Winston et al., 2003). It is important to note that in the majority of these studies, the rpSTS was more robustly identified across participants than the lpSTS (but see Engell and Haxby, 2007).

This functional asymmetry of the face-responsive pSTS suggests that expression recognition may be preferentially processed in the right hemisphere (Narumoto et al., 2001). Such a hypothesis is consistent with



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ABSTRACT

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an earlier neuropsychological study in which patients with lesions in the right hemisphere were significantly worse in performing tasks using emotional faces than patients with left-hemispheric lesions (DeKosky et al., 1980). A right hemisphere advantage was also demonstrated by the left visual field superiority for the recognition of emotional facial expressions in healthy individuals (Sackeim et al., 1978). The neuropsychological and behavioural evidence demonstrates that faces are preferentially processed in the right hemisphere but does not address whether this asymmetry is due to the asymmetry of the face-responsive area in the pSTS. As stated above, neuroimaging studies report that the rpSTS is more commonly identified than the lpSTS, but the neuroimaging methods do not causally address whether only the rpSTS, or the bilateral pSTS, is engaged in expression recognition.

In the current study, we systematically investigated the causal contributions of the right and left pSTS to facial expression recognition using TMS. Functional magnetic resonance imaging (fMRI) was used to individually localise the face-responsive area in the right and left pSTS of every participant. TMS was then delivered over the rpSTS, lpSTS and the vertex while participants performed facial expression and object recognition tasks. The vertex acted as a control site and the object task acted as a control task for the non-specific effects of TMS. Our aim was to provide a fuller picture of the functional properties of the pSTS in the extended face processing network, and to further contribute to the debate about hemispheric specialisation of expression recognition.

2. Materials and methods

2.1. Participants

Thirty-one right-handed volunteers participated in this study. One participant found TMS uncomfortable; he withdrew from the study and his data were discarded. All remaining participants (15 women and 15 men; aged between 18 and 44 years, mean: 23, SD: 6) were neurologically healthy with normal or corrected-to-normal vision. Informed consent was obtained after the experimental procedures were explained. A *post hoc* power analysis in GPower (Erdfelder et al., 1996) indicated that with the present sample, power of 97% was achieved with alpha set at 0.05. All participants were paid for their participation. The study was approved by the York Neuroimaging Centre (YNiC) Research Ethics Committee at the University of York.

2.2. Experimental procedures

Each participant completed two sessions, performed on different days. During the first session, participants were scanned using fMRI to functionally localise face-responsive regions in the right and left pSTS. These regions were then used as stimulation targets in the TMS study that was performed in the subsequent session. The fMRI session lasted approximately 1 h while the TMS session lasted approximately 1.5 h.

2.3. FMRI functional localisation

2.3.1. Procedure

Functional data were acquired over 6 block-design runs, lasting 234 sec each. During those runs, participants were instructed to watch videos of faces, bodies, scenes, objects, or scrambled objects, without performing any overt task. Each run contained two sets of five consecutive stimulus blocks to form two blocks per stimulus category per run. Each block lasted 18 sec and contained stimuli from one of the five stimulus categories. Each functional run also contained three 18 sec rest blocks, which occurred at the beginning, middle, and end of the run. During the rest blocks, a series of six uniform color fields were presented for 3 sec each. The order of stimulus category blocks in each run was palindromic (e.g., rest, faces, objects, scenes, bodies, scrambled objects, rest, scrambled objects, bodies, scenes, objects, faces, rest) and randomised across runs. At the end of the session a structural brain scan was

collected to anatomically localise the functional data for each participant.

2.3.2. Stimuli

The stimuli consisted of 3 sec movie clips of faces, bodies, scenes, objects and scrambled objects. Movies of bodies and scenes were not relevant to this study hence their data are not presented. The main motivation for using dynamic faces in the localisation procedure was to maximise chances of finding face-responsive areas in pSTS. This region was shown to respond stronger to dynamic stimuli than to the static stimuli, while activations for both types of stimuli spatially overlapped (Pitcher et al., 2011). These stimuli have also been used in prior fMRI and TMS studies of the pSTS (Pitcher et al., 2014; Pitcher et al., 2017). There were 60 movie clips for each category in which distinct exemplars appeared multiple times. Movies of faces and bodies were filmed on a black background and framed close-up to reveal only the faces or bodies of 7 children as they danced or played with toys or adults (who were out of frame). Movies of scenes included fifteen different locations which were mostly pastoral scenes filmed from a car window while driving slowly through leafy suburbs, along with some other films taken while flying through canyons or walking through tunnels that were included for variety. Movies of objects used 15 different moving objects that were selected in a way that minimizes any suggestion of animacy of the object itself or of a hidden actor pushing the object. Those included mobiles, windup toys, toy planes and tractors, and balls rolling down sloped inclines. Movies of scrambled objects were constructed by dividing each object movie clip into a 15×15 box grid and spatially rearranging the location of each of the resulting movie frames. Within each block, stimuli were randomly selected from within the entire set for that stimulus category. This meant that the same movie clip could appear within the same block but given the number of stimuli this did not occur frequently.

2.3.3. Data collection

Imaging data were collected using a 3T GE HDx Excite MRI scanner at YNiC. Functional images were acquired with an 8-channel phased array head coil (GE) tuned to 127.4 MHz and a gradient-echo EPI sequence (38 interleaved slices, repetition time (TR) = 3 sec, echo time (TE) = minimum full, flip angle = 90°; voxel size = $3 \times 3 \times 3$ mm; matrix size = 128×128) providing whole brain coverage. Slices were aligned with the anterior to posterior commissure line. Structural images were acquired using the same head coil and a high-resolution T-1 weighted 3D fast spoilt gradient (SPGR) sequence (176 interleaved slices, repetition time (TR) = 7.8 sec, echo time (TE) = minimum full, flip angle = 20° ; voxel size = $1 \times 1 \times 1$ mm; matrix size = 256×256).

2.3.4. Data analysis

Data were analysed using fMRI Expert Analysis Tool (FEAT) included in the FMRIB (v6.0) Software Library (www.fmrib.ox.ac.uk/fsl). In the first-level analysis, as part of the pre-statistical processing, singleparticipant functional images underwent extraction of non-brain structures performed with the Brain Extraction Tool (BET). In addition, interleaved slice timing correction, MCFLIRT motion correction, spatial smoothing using a 5 mm full-width half-maximum Gaussian kernel, highpass temporal filtering, and pre-whitening were applied to the data. The pre-processed functional images were entered into a general linear model (GLM) with five independent predictors, corresponding to the five categories of visual stimuli (i.e., Faces, Bodies, Scenes, Objects, Scrambled Objects), to compute participant-specific patterns of activation. The model was convolved using a double-gamma hemodynamic response function (HRF) to generate the main regressors. Temporal derivatives for each condition were included.

Face-responsive areas in the right and left pSTS were identified using a contrast of faces greater than objects. First-level functional results for each participant were registered to their anatomical scan using a 12 degree-of-freedom affine registration. All analyses were conducted at the whole-brain level and differences between conditions were considered significant at Z = 3.1 and cluster p = 0.05, using a cluster-wise

significance test.

To examine the hemispheric laterality of facial expression recognition, region-of-interest (ROI) analyses were performed. For each participant, regional masks were created using a sphere with 5 mm radius centred at the strongest peaks within the right and left pSTS activation clusters (i.e., peaks targeted with TMS), defined by the individual contrast of (Faces > Objects). The mean intensity of blood-oxygen-level dependent (BOLD) signal was then extracted from the (Faces > Objects) contrast for the right and left pSTS, and compared using a paired two-tailed *t*-test.

In order to report coordinates of TMS target sites in standard space, each participant's structural scan was registered to the Montreal Neurological Institute (MNI)-152 template. Note that all stimulation was done in native anatomical space and the standard space coordinates were computed solely for reporting purposes. In addition, the MNI coordinates from each participant were presented on the MNI brain in order to demonstrate the anatomical variability of the pSTS hot spots across individuals.

2.4. TMS experiment

2.4.1. Procedure

The TMS session involved the acquisition of behavioural data while participants performed computer-based visual facial expression and visual object matching tasks. The object recognition task acted as a control task. Each task was performed under four different stimulation conditions: i) TMS delivered over the rpSTS; ii) TMS delivered over the lpSTS; iii) TMS delivered over the vertex (control site); and iv) no TMS (behavioural baseline). Both tasks were taken and adapted from previous studies ran by Pitcher and colleagues (Pitcher et al., 2009; Pitcher et al., 2008). The facial expression matching task required participants to judge whether two faces of different people had the same expression. In the object matching task, participants were asked to judge whether two objects were the same. This task was used to control for non-specific to TMS effects that could result from differences in somatosensory sensation of stimulation that varied between the sites. Particularly, TMS over both pSTS sites produced mild peripheral jaw muscle twitching while the vertex stimulation did not produce any muscular responses. Participants sat 57 cm away from the monitor and used their right index or middle finger to respond "yes" or "no", respectively, by pressing appropriate keys on a keyboard. They were instructed to respond as accurately and quickly as possible.

Participant completed four runs (one run per TMS condition) of the facial expression matching task followed by four runs of the object matching task, or vice versa. Task order was counterbalanced across participants. The order of the TMS conditions was randomised across participants, but kept the same for both tasks in each participant. Each task run consisted of 72 trials, with half 'same' and half 'different' trials. The trial design is shown in Fig. 1. Each trial commenced with a fixation cross displayed for 2000 msec, followed by the target image displayed for another 250 msec, then a fixation cross displayed for another 1000 msec, and the match image displayed for another 250 msec. A trial ended with a blank white screen that was displayed until the participant responded. All stimuli were presented in the centre of a white screen on a Mitsubishi Diamond Pro 2070SB 22-inch CRT monitor, set at 1024×768 resolution and refresh rate of 85 Hz.

2.4.2. Stimuli

The stimuli in the expression matching task were taken from (Ekman and Friesen, 1976), and consisted of grayscale pictures of six female models (C, MF, MO, NR, SW) expressing six emotions: happy, sad, surprise, fear, disgust, and anger. Each picture was cropped with the same contour to cover the hair and neck of the models. Identity of the two faces within each trial was always different and the six expressions were presented an equal number of times within each run.

For the object matching task, pictures of novel, abstract objects were downloaded from Michael Tarr's website (http://wiki.cnbc.cmu.edu/ TarrLab). The "different" trials comprised of two objects that were morphed so that the objects were seen from the same viewing angle and had the same overall shape but varied in local details to different degrees. The percentage difference between the two images was either 20%, 30%, 50%, 80%, or 100%. A number of each morph type was equal across runs.

All stimuli were static in contrast to the stimuli used in the functional localisation. However, we did not expect any significant spatial differences in localisation of the static and dynamic stimuli based on our



Fig. 1. The experimental trial procedure. The expression matching task (A) required the participant to judge whether two faces belonging to different people had the same expression. The object matching task (B) required the participant to judge whether two objects were identical. TMS was applied at the onset of the match image at a frequency of 10 Hz for 500 msec.

previous work (Pitcher et al., 2011) that showed similar responses to both types of stimuli in pSTS. Also, the spatial resolution of TMS, measured in tens of millimetres (Brasil-Neto, McShane, Fuhr, Hallett and Cohen, 1992), provided a degree of spatial tolerance in case of spatial differences and intra-subject variability in the functional location at different time points (Duncan et al., 2009).

2.4.3. Data collection

TMS was delivered using a Magstim Rapid2 stimulator and a Magstim coated Alpha Flat 50 mm diameter figure-of-eight coil (Magstim, Carmarthenshire, UK). The stimulation intensity was set to 60% of the maximum stimulator output for all participants. A single intensity was used for all participants on the basis of our previous studies (Pitcher et al., 2008, 2009). During TMS runs, a train of five pulses at a frequency of 10 Hz was delivered for 500 msec on each trial. The onset of the TMS coincided with the onset of the match image. The TMS frequency, intensity, and duration were within established international safety limits (Rossi et al., 2009; Wassermann, 1998). The TMS coil was held against the participant's head by the experimenter who manually controlled its position throughout testing.

TMS target sites in the right and left pSTS were marked as stimulation targets on each participant's MRI scan using the Brainsight frameless stereotaxy system (Rogue Research, Montreal, Canada). During testing, a Polaris Vicra infrared camera (Northern Digital, Waterloo, ON, Canada) was used in conjunction with the Brainsight to register the participant's head to their MRI scan for accurate stimulation targeting throughout the experiment. All participants wore earplugs in both ears to attenuate the sound of the coil discharge and avoid damage to the ear (Counter et al., 1991). In some participants, stimulation affected the peripheral jaw muscle and produced a small jaw twitch. One participant described stimulation to his rpSTS as uncomfortable and was excluded from the study. The remaining participants tolerated TMS well.

2.4.4. Data analysis

A TMS control baseline was calculated as the mean accuracy score for vertex and no TMS condition in each participant. This type of TMS baseline was used as it constitutes a more representative measure of the control condition for stimulation than each of the two measures in isolation. It was possible to average vertex and no TMS conditions (both equal 82% in the expression matching task and 77% in the object matching task) as there was no significant difference in task performance between them (both paired two-tailed *t*-tests: t(29) < 0.79; p > 0.44; Cohen's effect size d < 0.001). Analyses of the main data with any of those conditions in separation showed the same pattern of results as when the conditions were averaged together.

Performance accuracy and reaction times (RTs) were analysed using IBM SPSS Statistics (v24.0) in a 2×3 repeated measures ANOVA, with Task (Facial Expression Matching Task and Object Matching Task) and Stimulation (TMS to rpSTS, TMS to lpSTS, and TMS Control Baseline) as independent factors. *Post hoc* paired two-tailed *t*-tests (with Bonferroni correction for multiple comparisons) were used to further characterize significant main effects and interactions from the ANOVA.

3. Results

3.1. FMRI functional localisation

We were able to successfully identify a face-responsive area in the right and left pSTS in all participants. The coordinates and strength of the peak activation varied across individuals (see Table 1 for information on individual peak coordinates in the standard MNI space and Fig. 2 for their illustration). The group mean peak coordinates in the standard space [rpSTS: x = 55, y = -39, z = 9; lpSTS: x = -56, y = -45, z = 9] were consistent with previous studies (e.g., Engell and Haxby, 2007).

The ROI analyses of the average mean voxel intensity showed significantly stronger response in the rpSTS (7.90 a.u.) than in the lpSTS

Table 1

Information about the rpSTS and lpSTS, including individual peak coordinates in the standard MNI space; BOLD intensity for ROIs and TMS effects on accuracy during the expression matching task. A TMS effect is a difference between the TMS and baseline conditions where negative values indicate successful disruption of processing.

Participant (G)	rpSTS					lpSTS				
	x	у	Z	BOLD intensity (a.u.)	TMS effect (%)	x	у	z	BOLD intensity (a.u.)	TMS effect (%)
1 (F)	59	-37	8	6.24	-9	-60	-45	10	6.68	-6
2 (F)	50	-38	16	12.34	-2	-47	-61	12	6.37	-2
3 (F)	46	-24	$^{-1}$	4.83	-3	-66	-32	2	3.45	-6
4 (F)	56	-39	4	11.53	-2	-49	-45	1	9.81	8
5 (F)	53	-39	0	7.30	$^{-1}$	-57	-43	0	2.94	-2
6 (F)	58	-42	10	7.07	1	-52	-44	13	4.19	-3
7 (M)	54	-43	15	11.75	-3	-52	-40	12	9.89	0
8 (F)	54	-33	8	5.54	-3	-49	-30	1	4.36	-1
9 (F)	50	-43	12	8.27	$^{-1}$	-51	-42	10	5.09	-2
10 (M)	58	-49	10	5.55	-6	-64	-47	9	5.62	-2
11 (M)	50	-38	-3	4.04	$^{-1}$	-65	-33	6	3.26	1
12 (F)	51	-38	5	11.00	-3	-48	-45	7	13.42	1
13 (M)	46	-42	17	7.43	2	-48	-45	4	3.55	-3
14 (M)	43	-30	3	12.76	-4	-53	-43	11	7.35	-3
15 (M)	51	-37	4	14.99	$^{-1}$	-61	-38	0	7.60	2
16 (F)	55	-44	16	9.40	-3	-45	-43	17	10.63	2
17 (M)	50	-43	9	6.61	-6	-49	-52	16	8.54	1
18 (M)	51	-38	10	8.54	-6	-62	-52	11	11.48	-4
19 (F)	65	-29	4	7.12	-6	-47	-58	15	4.70	-4
20 (M)	61	-48	8	5.39	-13	-58	-35	10	3.87	-8
21 (M)	60	-33	5	6.31	-3	-65	-35	13	2.59	-1
22 (M)	64	-52	8	7.81	-10	-56	-62	7	5.22	-3
23 (M)	59	-39	12	13.52	-6	-66	-27	8	6.81	1
24 (M)	58	-36	12	4.12	-9	-62	-41	6	2.56	-10
25 (F)	55	-29	6	12.36	-11	-65	-46	14	3.89	-6
26 (F)	68	-36	10	4.94	-4	-61	-43	6	4.78	2
27 (M)	64	-43	27	5.34	$^{-1}$	-60	-44	9	7.55	-2
28 (M)	52	-48	0	7.11	1	-51	-50	3	6.09	-3
29 (F)	56	-43	3	2.20	-15	-54	-68	14	2.82	-6
30 (F)	53	-46	33	5.53	-5	-66	-52	8	4.78	-12
Mean	55	-39	9	7.90	-4	-56	-45	9	6.00	-2

(6.00 a.u.; t(29) = 3.56; p = 0.001; Cohen's effect size d = 0.68). In 22 participants, activation intensity was greater in the rpSTS than in the lpSTS while in the remaining 8 participants the opposite pattern was observed (see Table 1).

3.1.1. TMS study

The mean accuracy results are shown in Fig. 3. The main effects of Task (F(1, 29) = 2.54; p = 0.12; partial $\eta^2 = 0.08$) and Stimulation (F(2, 58) = 2.22; p = 0.12; partial $\eta^2 = 0.07$) were not significant. However, there was a significant two-way interaction between Task and Stimulation (F(2, 58) = 11.20; p < 0.001; partial $\eta^2 = 0.28$). *Post hoc t*-tests showed that during the expression matching task, TMS delivered over the rpSTS (78%; t(29) = 6.21; p < 0.001; Cohen's effect size d = 1.05) and lpSTS (80%; t(29) = 3.31; p = 0.003; Cohen's effect size d = 0.49) significantly impaired accuracy in relation to the TMS control baseline condition (82%). TMS delivered over the rpSTS (78%; t(29) = 0.50; p = 0.62; Cohen's effect size d = 0.19) had no effect on performance of the object matching control task in relation to the baseline condition (77%).

In addition, the difference between the accuracy in the rpSTS and lpSTS in the expression matching task was significantly different (t(29) = 2.75; p = 0.01; Cohen's effect size d = 0.49). In the facial expression task, TMS delivered over the rpSTS impaired accuracy in 26 participants while TMS delivered over the lpSTS impaired accuracy in 21 participants. 18 participants showed effect of TMS in both hemispheres while in the remaining 9 and 3 participants TMS had an effect only in the rpSTS and the lpSTS, respectively.

RTs showed no significant two-way interaction between Task and Stimulation (F(2, 58) = 1.51; p = 0.23; partial $\eta^2 = 0.05$) or main effects of Task (F(1, 29) = 1.89; p = 0.18; partial $\eta^2 = 0.06$) and Stimulation (F(2, 58) = 0.62; p = 0.54; partial $\eta^2 = 0.02$).

4. Discussion

In the present study we used fMRI and TMS to systematically investigate the causal contributions of the right and left pSTS to facial expression recognition. fMRI results showed that the rpSTS response to faces was greater than the response of the lpSTS. The TMS results



Fig. 2. Top panel illustrates the individual peak coordinates for all 30 participants (black circles) and the mean group coordinates (red circles) in the left and right pSTS, presented in the standard MNI-152 space. Bottom panel shows bar plots for the mean group BOLD signal intensity extracted from the (Faces > Objects) contrast for the left and right pSTS. Error bars represent SEM. *p < 0.05.



Fig. 3. Group mean accuracy scores during the expression and object matching tasks for the three stimulation conditions: i) TMS over the rpSTS (light grey) ii) TMS over the left pSTS (dark grey), and iii) TMS control baseline (black). Error bars represent SEM. *p < 0.05.

revealed that both the right and left pSTS are causally involved in recognising facial expressions. However, consistent with the fMRI results, TMS delivered over the rpSTS produced a greater disruption in task performance than TMS delivered over the lpSTS. Our results demonstrate that the rpSTS was more engaged in task performance than its left homologue, but optimal facial expression recognition requires computations performed in both right and left pSTS.

Our neuroimaging results are consistent with the previous neuroimaging study (Engell and Haxby, 2007) that revealed increased activation in the right and left pSTS during expression recognition tasks. It is also in line with the face processing model proposed by Haxby and colleagues (Haxby et al., 2000) that include both the right and left pSTS in the face network. These results are, however, seemingly inconsistent with other neuroimaging studies (e.g., Narumoto et al., 2001; Streit et al., 1999; Winston et al., 2004; Winston et al., 2003) that demonstrated a neural response to expressions in the rpSTS only, and consequently focused solely on the role of the rpSTS in processing expressions. Such a discrepancy in detecting responses in both hemispheres may result from differences in the experimental design. Our functional localiser used dynamic, rather than static, face stimuli as those have been demonstrated to be more suitable for eliciting responses in the pSTS, a brain region preferentially engaged in moving faces (Pitcher et al., 2011; Polosecki et al., 2013). Also, using static face stimuli in the current TMS study supported our previous findings (Pitcher et al., 2011) which showed that static faces and dynamic faces activate the same face-responsive areas in the pSTS. Based on our previous findings of a stronger response to dynamic faces than static faces in pSTS, we would predict that TMS effects in the expression matching task would be greater if dynamic stimuli was used.

Our findings are inconsistent with a recent fMRI study (De Winter et al., 2015) that specifically measured lateralisation in face processing areas while humans and monkeys watched dynamic expressions. In humans, the rpSTS was found to be the only region that showed a clear lateralisation for visual non-linguistic facial expressions, with no involvement of the left hemisphere. In contrast, visual linguistic expressions were found to fully engage lpSTS, but not its right homologue. No consistent pattern of lateralisation was found in monkeys. The authors argued that their findings support the verbal versus visuospatial model (Corballis et al., 2000), which proposes that the right hemispheric dominance for visual stimuli evolved as a cost of language specialisation in the opposite hemisphere. While our results are still consistent with the idea of the hemispheric specialisation for visual and language functions, they do not support the absolute segregation of these functions into different hemispheres.

The extent to which cognitive functions are functionally lateralised in the human brain remains an active debate in the cognitive neuroscience. One of the issues is the extent to which we investigate the functional contribution of the less dominant hemisphere to our chosen cognitive operation. The current study addresses this debate by looking specifically at facial expression recognition in the right and left pSTS. By using a robust functional localiser, we were able to identify the face-responsive pSTS in both hemispheres for each participant, and then demonstrate with TMS that although both regions contribute to accurate expression identification, the rpSTS contribution dominates over the lpSTS. Although our study focused on lateralisation of face processing in the pSTS, this region is also implicated in other cognitive functions, some related to language (for review see Price, 2012). Left-hemispheric dominance for language processing was proposed in the first lesion-based neurological model of language (Geschwind, 1970), and later supported by neuroimaging studies (Petersen et al., 1989; Pujol et al., 1999; Scott et al., 2000; Springer et al., 1999; Zatorre et al., 1992). Based on our findings, we predict that both the right and the left pSTS are important for optimum performance of language functions, but lpSTS involvement would dominate over the rpSTS.

A discrepancy in evidence for functional lateralisation between brain stimulation studies and neuroimaging studies has also been reported in brain areas engaged in other cognitive tasks. For example, Santiesteban et al. (2015) examined the lateralisation of socio-cognitive abilities in the temporoparietal junction (TPJ) using transcranial direct current stimulation (tDCS). Their results demonstrated that the right and left TPJ is causally involved in tasks requiring imitation control and visual perspective-taking. This contrasts with the evidence from the neuroimaging literature suggesting unilateral activation of the right TPJ during imitation control (Brass et al., 2005; Spengler et al., 2009) or left TPJ during visual perspective-taking (Schurz et al., 2013). As the authors argue, those differences can result from the application of the over-conservative statistical thresholds in neuroimaging studies to avoid Type I errors while potentially creating Type II errors. It is also possible that those differences are caused by the propagation of the effects of the unilateral stimulation to the opposite hemisphere. However, the reports of selective effects on cognitive tasks following unilateral stimulation over the right and left TPJ (Heinisch et al., 2010; Uddin et al., 2007) and in our case pSTS (e.g., Oliveri et al., 2004; Pobric et al., 2008) do not support this conclusion.

It is also worth noting that our study demonstrated that the strength of activation and size of TMS effect varied across individuals. Some participants exhibited a greater activation, or a greater TMS effect in the left rather than the right pSTS (see Table 1). Similar inter-individual variability in the magnitude of lateralisation in pSTS was found by De Winter et al. (2015). In their study, a number of participants showed lateralisation of dynamic facial expressions to the lpSTS. These differences could be explained by inter-individual differences in i) the development of hemispheric lateralisation for language and faces (Dundas et al., 2012); ii) strategies used for face recognition; iii) functional location between the static and dynamic faces; or lack of individualised stimulation parameters.

The current TMS results replicate and extend the results from our earlier TMS study in which TMS delivered over the rpSTS impaired expression recognition (Pitcher, 2014). The present study, additionally demonstrates that the lpSTS is causally important for recognising expressions, albeit to a lesser extent. In another preceding study (Dzhelyova et al., 2011), TMS delivered over the right and left pSTS impaired judgments of facial trustworthiness. However, this study did not systematically explore the differences between the left and right pSTS. In contrast, our results are not consistent with another set of findings in which TMS delivered over the rpSTS impaired eye gaze discrimination, but did not impair expression recognition (Pourtois et al., 2004). This may be due to methodological differences in how TMS target sites were selected. Pourtois and colleagues identified the pSTSbased on the EEG electrode system while we used functional fMRI localisers to identify target sites individually in each participant. It is apparent from our localisation results (Fig. 2) that the location of the target sites varied greatly across the two hemispheres in each individual and across individuals, indicating the need for using a more precise localisation method when studying the pSTS.

Although our study provided strong evidence for the importance of the right and left pSTS in recognising expressions, the precise role of this region requires further investigation. It has been suggested that the pSTS may be involved in extracting information about the eye gaze in order to interpret expressions, and may be homologous to a region in the superior bank of the monkey's STS where cells respond preferentially to gaze direction (Hasselmo et al., 1989; Perrett et al., 1985). Also in humans, a number of studies demonstrated that the pSTS is involved in eye gaze discrimination (Engell and Haxby, 2007; Pourtois et al., 2004; Puce et al., 1998). It is not possible to determine from our fMRI data whether the pSTS responded to changes in eve gaze because we did not systematically manipulate this factor. There is evidence in the neuroimaging literature (Engell and Haxby, 2007) suggesting that gaze-direction and expressions are represented by distinct but overlapping regions in the rpSTS. Therefore, the specifics of the functional contributions of various sub-regions of the pSTS to facial expression processing still require further investigation, especially with the causal methods like TMS. It is also worth noting that the pSTS is engaged in a range of other cognitive tasks including recognition of intentional actions (Saxe et al., 2004) or body perception (Basil et al., 2017), and future TMS studies could assess how these functions are lateralised.

Our study demonstrated that both the right and left pSTS make a functional contribution to accurate facial expression recognition. Nevertheless, the engagement of the rpSTS was greater than the lpSTS, suggesting the functional domination of this process in the rpSTS. Although this study supports the concept of right-hemispheric specialisation of face processing, it also shows that the face regions in the left hemisphere play a crucial role in this process and their contribution should not be neglected.

Declarations of interest

None.

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