

ORIGINAL ARTICLE

Overlapping Functional Representations of Self- and Other-Related Thought are Separable Through Multivoxel Pattern Classification

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Abstract

Self-reflection and thinking about the thoughts and behaviors of others are important skills for humans to function in the social world. These two processes overlap in terms of the component processes involved, and share overlapping functional organizations within the human brain, in particular within the medial prefrontal cortex (MPFC). Several functional models have been proposed to explain these two processes, but none has directly explored the extent to which they are distinctly represented within different parts of the brain. This study used multivoxel pattern classification to quantify the separability of self- and other-related thought in the MPFC and expanded this question to the entire brain. Using a large-scale mega-analytic dataset, spanning three separate studies ($n = 142$), we find that self- and other-related thought can be reliably distinguished above chance within the MPFC, posterior cingulate cortex and temporal lobes. We highlight subcomponents of the ventral MPFC that are particularly important in representing self-related thought, and subcomponents of the orbitofrontal cortex robustly involved in representing other-related thought. Our findings indicate that representations of self- and other-related thought in the human brain are described best by a distributed pattern rather than stark localization or a purely ventral to dorsal linear gradient in the MPFC.

Key words: classification, MPFC, MVPA, self-reference, social

Introduction

A hallmark of human cognition is the ability to reflect upon one's thoughts and behaviors, which inform one's own self-image (James 1890; Gallagher 2000). Similarly, humans are also adept social learners, observing others' behaviors, inferring their mental states, and using this to inform personal impressions (Frith and Frith 2006; Ma et al. 2014). Whether and how judgments about the self and others are distinct is a question that has interested psychologists for decades (Rogers et al. 1977; Bower and Gilligan 1979). Neuroimaging research has more recently contributed to this question, and identified regions of

the brain associated with self-related and other-related thought. The extent to which these neural correlates overlap, however, is still not fully understood. Systematic reviews, individual studies, and meta-analytic research highlight that the medial prefrontal cortex (MPFC), in particular, is involved in both self- and other-related thought (Pfeifer et al. 2007; Jenkins et al. 2008; Bergstrom et al. 2015; Wagner et al. 2019), but that there is likely not a stark sublocalization of these processes (Denny et al. 2012). Instead of a binary distinction between these two processes, it is likely that they are represented grossly similarly in some parts of the brain, but it is not known how distinguishable they are in terms

of more granular representations. Therefore, this research aims to estimate where, and to what extent, self- and other-related thought are distinctly represented in the MPFC, and throughout the brain as a whole.

Neural Correlates of Self-Related Thought

There are several regions of the brain that are associated with self-related processing, including the posterior cingulate cortex, precuneus, and MPFC (Kelley et al. 2002; Ochsner et al. 2004; Mitchell et al. 2005; Johnson et al. 2006; Jenkins and Mitchell 2010; Martial et al. 2018). Whether individuals are assessing emotional content or episodes related to the self (Ochsner et al. 2004; De Pisapia et al. 2019; Verfaellie et al. 2019), the similarity of others' faces to their own (Mitchell et al. 2005) or their own personality traits (Heatherton et al. 2006; D'Argembeau et al. 2007; Beer et al. 2010; Rameson et al. 2010; Martinelli et al. 2013), the MPFC is reliably activated.

The MPFC is a relatively large area of the brain, and both dorsal and ventral portions of the MPFC have been reported in separate research studies of self-related thought (Ochsner et al. 2004; van der Meer et al. 2010). Some research, however, argues that self-related processing is most robustly localized to ventral portions of the MPFC (VMPFC) (van der Meer et al. 2010; Wagner et al. 2012). This position is bolstered by the fact that patients with damage to the VMPFC also show significant deficits in their ability to remember self-relevant information and memories (Wheeler et al. 1997; Philippi et al. 2012). Indeed, individuals with focal damage to the VMPFC have notable difficulty in remembering information about themselves, but do not show similar difficulties in remembering information about other people (Marquine et al. 2016). The partly specialized role for the VMPFC in processing information about the self, relative to others, is further specified by patients with lesions to regions other than the VMPFC (e.g., lateral occipital, temporal, and parietal regions) who do not show similar deficits in self-referential processing (Philippi et al. 2012).

Neural Correlates of Other-Related Thought

The MPFC is also activated when individuals make trait judgments about other people (Mitchell et al. 2002), form impressions of others (Mende-Siedlecki et al. 2013), hold social information in working memory (Meyer and Collier 2020), and when they infer the mental states of others (Saxe 2006; Blakemore 2008; Hampton et al. 2008; Skerry and Saxe 2014). A large body of work has shown that the dorsal MPFC (DMPFC) is more strongly associated with the processing of information relevant to other people than the self (see Van Overwalle 2009 for review). Further, direct contrasts of thinking about others and thinking about oneself have revealed greater activation in the DMPFC, supporting the specificity hypothesis (D'Argembeau et al. 2007). The DMPFC also exhibits increased coherence with other regions of the default mode network (e.g., posterior cingulate cortex (PCC), temporoparietal junction (TPJ)) when assessing the traits of others but not oneself, further linking this subregion of the MPFC to social cognition (Hassabis et al. 2014).

More broadly, thinking about the traits and behaviors of others and inferring their thoughts are also all—to different degrees—associated with activation in the temporal parietal junction, the precuneus, and superior temporal sulcus. (Saxe and Waxler 2005; Saxe 2006; Dodell-Feder et al. 2011). These regions also provide useful information in the prediction of

socially relevant cognition, like assessing the social status of other individuals (Parkinson et al. 2017).

Overlap of Self- and Other-Related Thought

Other neuroimaging literature, however, provides reason to question whether self- and other-related thought can be starkly linked to distinct portions of the MPFC (Saxe et al. 2006). For example, ventral regions of the MPFC are frequently identified in studies of socially oriented thought (Kelley et al. 2002; Lou et al. 2004; D'Argembeau et al. 2005; Koster-Hale et al. 2017) and dorsal regions are similarly activated during self-oriented thought (Craig et al. 1999; Gusnard et al. 2001; Schmitz et al. 2004; Seger et al. 2004). Such double associations have motivated alternative models of organization within the MPFC. More specifically, researchers have suggested that self- and other-related thought may be organized along a ventral to dorsal gradient within the MPFC (Mitchell et al. 2005; Heatherton et al. 2006; Tamir and Mitchell 2010). Perhaps the most influential evidence for this perspective comes from a meta-analysis conducted by Denny et al. (2012). In this study, the authors conducted a review of the literature on self-related thought and other-related thought, and found 47 peak activations for self-thought and 43 peak activations for other-thought across 107 studies within the MPFC. The authors then conducted a logistic regression using the z-coordinate of each activation to model the organization of this body of research. They found a ventral-dorsal gradient, such that peak activations located more ventrally were more likely to be (though not exclusively) reported in a study of self-related thought and activations located more ventrally were more likely to be (though not exclusively) reported in a study of other-related thought (Denny et al. 2012).

Denny et al. (2012) provide a parsimonious probabilistic model for how self- and other-related thought are distributed within the MPFC. What this model does not answer, however, is whether, and to what extent, these two processes can be distinguished within subregions of the MPFC. Indeed, even within the data described by Denny and colleagues, there is substantial overlap in where studies of self- and other-related thought report activations. This suggests that in addition to a dorsal-ventral gradient, there may be varying degrees of overlap in the function of specific regions. In order to directly test the question of separability within subregions, it is necessary to compare direct observations of self-related and other-related thought within the MPFC; a level of analysis which meta-analysis abstracts from, and which requires more direct access to raw data. Denny et al. reveals that, indeed, self- and other-related thought are distributed throughout the MPFC, but do not directly quantify the extent to which these distributed representations are separable.

Quantifying the Separability of Self- and Other-Related Thought

The introduction of multivariate pattern analysis (MVPA) and machine learning in neuroimaging research has made it possible to identify differences between cognitive states through the analysis of their distributed patterns of activation. In one of the foundational MVPA studies, Haxby and colleagues found that multivariate patterns within the ventral temporal (VT) cortex were more sensitive to differences between different object categories compared to univariate analysis (Haxby et al. 2001;

Norman et al. 2006). Activation related to different object categories in the VT overlapped substantially, but the distributed patterns related to each were discernible. Analogously, self- and other-related processing overlap substantially in the MPFC, but it is not yet clear to what extent the patterns of activation related to each are distinguishable.

Recently, there has been a growing interest in applying multivariate approaches to the study of self-related processing and other-related processing (Wagner et al. 2019; Courtney and Meyer 2020; Koski et al. 2020). This research has brought new understanding to the overlap of experienced and vicarious pain, and has found that the pattern of activation elicited by one is notably similar to the other (Corradi-Dell'Acqua et al. 2016; Krishnan et al. 2016). More specific to the aims of the current analysis, Oosterwijk et al. (2017) found that a model trained to classify different forms of self-related processing (e.g., thinking about one's own emotions, actions or physical feelings) is also able to classify these different forms of thinking when they are about other people (Oosterwijk et al. 2017). The evidence thus far indicates that self- and other-related processing are not only overlapping in *where* they activate the cortex but also, at least partially, in *how* they activate the cortex. In the current study, we ask not how well a model trained on different forms of self-related processing generalizes to other-related processing, but instead ask whether, and to what extent self-related processing can be distinguished from other-related processing.

Here, we employ a mega-analytic approach to this question, which integrates raw data from multiple studies. Specifically, in this study, we analyze the data from a large sample of participants from three separate studies ((Cooper et al. 2015; Falk et al. 2015); one unpublished study) to ask how separable self- and other-related thought are within the MPFC and how these processes are functionally organized. We employ machine learning techniques including regularization and crossvalidation to quantify the extent to which these processes are distinct in the MPFC in particular as well as throughout the whole brain. We ask four directed questions regarding the functional organization of self- and other-related processing within the brain. First, we ask whether information from the MPFC is sufficient to reliably distinguish between self- and other-related thought. Second, we ask if the derived organization of regions that distinguish between self- and other-related thought in the MPFC follows a linear pattern along a ventral-dorsal gradient, or whether this organization is more complex. Third, we move beyond the MPFC to explore whether including information from the entire brain significantly improves the characterization and prediction of self- and other-related thought. Finally, we explore the map of voxel weights from the whole brain to ask how self- and other-related thought are organized outside of the MPFC.

Materials and Methods

Participants

One hundred and forty-two ($N = 142$) participants were included in this analysis. Participant data were collected as part of three separate neuroimaging studies ($N_{\text{study1}} = 60$; $N_{\text{study2}} = 39$; $N_{\text{study3}} = 43$). These three neuroimaging experiments included tasks, which probed the neural substrates of self- and other-related thought among other tasks unique to each study. Participants self-identified as 55% women and 45% men. Participants' ages ranged from 18 to 77 with an average

age of 29 years ($SD = 12.61$). Participants self-identified as: 10% Asian, 15% Black, 6% Latino, 5% multiracial, 56% White, 8% other ethnicity. The experimental protocols for each study were approved by the relevant university Institutional Review Boards (University of Pennsylvania and University of Michigan). Informed written consent was obtained from all participants.

Task and Procedure

Participants were recruited for three different neuroimaging experiments, all of which included as part of their procedures a paradigm called the self-localizer task. The self-localizer task is a widely used task to localize areas of the brain associated with self-related thought (Schmitz and Johnson 2006). Participants in the self-localizer task are presented with trait adjectives and are asked to make judgments about each word after its presentation. In all three of the current experiments, participants were asked to make judgments about the relevance of words to themselves, the relevance of words to another person, or whether the words had positive or negative valence. One of the three experiments (study 3) had additional conditions that varied slightly from the other two (described below), but the judgments focused on in the current analysis (relevance to self, relevance to another person) were shared across all three.

In studies 1 (Falk et al. 2015) and 2 (unpublished), participants made trait adjective judgments under three conditions: word describes you, word describes then-president Barack Obama, and word is positive or negative. Participants completed six blocks of each condition, each containing five trait adjective trials, for a total of 30 trials per condition. For each block participants viewed both positive and negative trait adjectives in a pseudorandom order. Each block of words began with a screen that indicated to the participant what type of judgment they were being asked to make. Each word was then presented and the participant made a judgment about the word. The word and the participant's decision remained on the screen until 2.5 s had elapsed, after which the next word was presented. Participants responded "yes" or "no" or "positive" or "negative" depending on whether they were making relevance judgments or valence judgments. Both "yes" and "no" responses in the self and social judgment constitute thinking about oneself or a social target, as either requires participants to reflect on their traits (or the social target's traits) and confirm or deny whether the given adjective matches.

In the third study (Cooper et al. 2015), participants made trait adjective judgments under five conditions. Four conditions required participants to take either their own perspective or the perspective of a friend, and judge whether the word described themselves or their friend. In this way, the participant could either be making judgments about whether they thought a word described themselves, whether they thought a word described a friend, whether they thought their friend would think the word described themselves, or whether they thought their friend would think the word described their friend. In the current analysis, to parallel the first two studies, we only focus on conditions in which participants took their own perspective and judged whether the words described themselves or a friend. Participants also completed a valence judgment condition like those in the studies 1 and 2. All task procedures and timing were the same as in studies 1 and 2 except that in this condition a total of 36 words were viewed and each block contained 6 trait adjectives, three positive and three negative.

Image Acquisition

Data from study 1 were acquired on a 3 Tesla GE Signa MRI scanner. Functional images were acquired using a reverse spiral sequence (TR=2000 ms, TE=30 ms, flip angle=90°, 43 axial slices, FOV=220 mm, slice thickness=3 mm, voxel size=3.44, 3.44, 3.0 mm). In-plane T1-weighted images (43 slices, slice thickness=3 mm, voxel size=0.86, 0.86, 3.0 mm) and high-resolution T1-weighted images (124 slices, slice thickness=1.02, 1.02, 1.2 mm, SPGR) were also acquired for use in coregistration and normalization. Data from study 2 were acquired on a 3 Tesla Siemens Magnetom MRI scanner. Functional images for study 2 were recorded using a multiband sequence (TR=1500 ms, TE=25 ms, flip angle=60°, 54 axial slices, FOV=208 mm, slice thickness=3 mm, voxel size=3.0, 3.0, 3.0 mm). High-resolution T1-weighted images were also acquired (160 slices, voxel size=0.9, 0.9, 1.0 mm) for use in coregistration and normalization. Finally, data from study 3 were acquired using a 3-Tesla GE Signa MRI scanner. Functional images were acquired using a reverse spiral sequence (TR=2000 ms, TE=30 ms, flip angle=90°, 43 axial slices, FOV=220 mm, slice thickness=3 mm, voxel size=3.44, 3.44, 3.0 mm). In-plane T1-weighted images (43 slices, slice thickness=3 mm, voxel size=0.86, 0.86, 3.0 mm) and high-resolution T1-weighted images (124 slices, slice thickness=1.02, 1.02, 1.2 mm, SPGR) were also acquired for use in coregistration and normalization.

fMRI Analysis

Data were preprocessed using Statistical Parametric Mapping (SPM8; Wellcome Department of Cognitive Neurology, Institute of Neurology, London, UK) for all stages except for initial despiking, which was performed using the 3dDespike program implemented in the AFNI toolbox (Cox 1996). Preprocessing steps for all three datasets included slice time correction, realignment, coregistration with both T1-weighted images, segmentation, and normalization to the MNI-152 and resampling to the same voxel size (3 mm).

Task block condition-specific estimates were calculated by running a voxelwise first-level general linear model for every participant, in which the entire block of a specific condition was modeled as its own separate regressor (performed with SPM12). This regression approach (Rissman et al. 2004), results in a separate whole brain map estimate for every self-judgment and other-judgment condition block. For our analysis of the MPFC, all first level block images were then masked using a binary gray matter image of the entire medial wall (3596 voxels). Following Denny et al., the boundaries for this mask were $|x| < 25$, $y > 15$. Importantly, we included the OFC in our mask, unlike Denny and colleagues, who restricted their mask to voxels above $z = -5$ (Denny et al. 2012). For our whole brain analysis, we also masked all first-level block images using a binary gray matter image (Shen et al. 2013). T-maps for the MPFC and whole brain data were flattened to a 1D array using custom code and the python neuroimaging toolbox, Nilearn (Abraham et al. 2014).

MVPA Training and Testing

Participants' data were first shuffled and split into a training and test set using an 80/20 split, and thus containing 111 and 32 participants respectively. Individual participants' data were kept together for this train-test sample split, such that any given participant's data only existed in one of these two samples. For model training, we implemented a Ridge-PCR analysis

method to differentiate self- and other-related processing for the MPFC and whole brain, respectively. This method closely follows Wager et al.'s (2011) LASSO-PCR method, in which PCA is run prior to model tuning and training. The motivation of this approach is twofold: first, PCA is initially performed in order to account for the tendency of L1 and L2 regularization to ignore the natural covariance in fMRI data and select or downweight noncontiguous voxels, and second PCA significantly reduces the size of the feature space during model training, which reduces the computational burden of performing permutation procedures. Our analysis diverged from the LASSO-PCR approach, in that we chose to implement a Ridge regression in order to maintain all features during model training (Hoerl and Kennard 1970). Similar to LASSO-PCR, Ridge-PCR returns linear feature weights, which allowed for easily interpretable brain maps.

PCA was performed with scikit-learn (version 0.22.1), which uses a LAPACK routine to compute the singular value decomposition (<http://www.netlib.org/lapack/faq.html>). Following the procedures of Wager et al., we retain a full-rank set of components for both MPFC and whole-brain analyses. There are 1145 observations in the training data (significantly fewer observations than voxels in both MPFC and whole brain), which resulted in n -observations minus 1 principle components, or 1144. In addition to the findings of Wager et al., classification analysis using a full-rank set of components has been shown to be a powerful method for discriminating various psychological processes from neuroimaging data (Chang et al. 2015; Krishnan et al. 2016; Koban et al. 2021).

Model development for the MPFC and whole brain respectively was performed using the python machine-learning toolboxes, scikit-learn (Pedregosa et al. 2011). To train and tune classifiers, 5-fold randomized crossvalidation was used. A grid-search protocol was implemented to tune the L2 regularization penalty hyperparameter, which tested values between 0.001 and 1 at increments of 0.1. For the classifier, the hyperparameter that resulted in the best average validation accuracy value (within folds) was finally retrained on all training data and carried forward for out-of-sample testing with the independent, held-out test set.

Classification models were tested out-of-sample for their predictive accuracy on the remaining self and other block images in the test sample ($N=384$ images across 32 test-set participants). Confidence intervals for out-of-sample test accuracy were derived using a bootstrap procedure, in which 1000 bootstrap samples were drawn with replacement. Confidence intervals that do not include 50% (chance) are considered significant.

We additionally carried out all of the above procedures using alternative classification models (PLS-DA, SVC-PCA; Platt 1999; Barker 2010) in order to establish that our model results were not dependent on our specific methodological choices. These methods indeed returned very consistent results to our main analysis, and details of these procedures can be found in our [Supplementary Materials](#).

Results

MPFC Classification

To differentiate self- and other- related cognition within the MPFC a Ridge-PCR model was tuned and trained on data extracted from the MPFC. Block images were used for this procedure. To first assess the predictive capacity of these models, the

within sample accuracy for each of five stratified crossvalidation folds was calculated. Overall, the model performed well within the training sample with an average validation accuracy score across folds of 63.7% (min=57%, max=66%). The component weights for the trained model were used to predict whether a not-yet-observed image came from a condition under which a participant was engaging in self- or other-related thought. The MPFC Ridge-PCR model achieved a mean out-of-sample test accuracy of 58.9% (CI=[54%, 64%]).

Whole Brain Classification

Analyses aimed at distinguishing self- versus other-related processing were next repeated using data from the entire brain. A Ridge-PCR classifier was trained and tuned in the same fashion as the MPFC analysis using block-level images and was then tested out of sample for its predictive accuracy. Training prediction accuracy was again calculated for every fold of the 5-fold cross validation and averaged for the model. The whole brain model performed well in the training sample, achieving a training accuracy score of 71.8% (min=67%, max=75%). The whole brain Ridge-PCR classifier had a mean out of sample classification accuracy score of 67.7% (CI=[62%, 73%]). The test accuracy score of the whole brain model was also directionally more accurate than the model using MPFC alone however, the confidence intervals obtained through resampling overlap substantially.

Organization of MPFC Model Weights

To investigate whether there is a ventral to dorsal gradient in the organization of voxel weights for the classification of self- versus other-related thought, we examined model weights from the Ridge-PCR model organized linearly from the most ventral to dorsal portion of the MPFC. These model weights represent the relative contribution and direction of each voxel in discriminating between self- and other-related thought. To obtain this organization, we projected model weights back into their original 3-D voxel space by multiplying the PCA component matrix with the best performing model regression coefficients. We correlated (Pearson r) the ventral to dorsal position (i.e., z -coordinate) of every voxel in the MPFC with the model weight for each voxel. We found initial supporting evidence for a dorsal to ventral gradient in the organization of MPFC weights ($r=0.09$, $P=0.03$); however, this relationship was weak, suggesting the possibility of a more complex organization (Fig. 1).

We obtained a P -value to test the significance of this correlation through a permutation procedure, which involved shuffling the order of the training labels, retraining the model, projecting the trained model weights back into their original 3-D space, and computing the z -coordinate correlation 1000 times. We also tested the reliability of the individual model weights through this same permutation procedure, resulting in a P -value for every model weight (voxel). (see [supplementary materials](#) from [Pereira et al. \(2009\)](#) for review of procedures). We then thresholded our model weights, controlling for multiple comparison (FDR corrected, $\alpha=0.05$). Clusters in the OFC and VMPFC survived this thresholding procedure, as can be seen in [Figure 2](#).

Organization of Whole Brain Model Weights

Voxel weights from the whole brain were also back projected into their original 3D voxel positions. Model weights, that

is, the relative contribution and direction of each voxel in discriminating between self- and other-related thought, were examined across the entire brain. To test the reliability of the model weights, we repeated the permutation procedures reported in the previous section for the whole brain and thresholded our model weights, controlling for multiple comparison (FDR corrected, $\alpha=0.05$). We found that the MPFC was reliably involved in discriminating between self- and other-related processing as were other regions throughout the brain. Peak voxel weights coding for self-related thought were primarily localized to the VMPFC and anterior cingulate cortex (ACC). Voxels coding for other-related thought were more distributed throughout the brain, showing peak voxel weights in the PCC, left angular gyrus, and left temporal lobe (Fig. 3).

Discussion

Can patterns of activation within the brain provide reliable information for discriminating between self- and other-related thought? We first focused on the MPFC, a region frequently and reliably implicated in both processes ([Heatherton 2011](#)), and then moved to examine patterns across the entire brain. Using raw data (i.e., a mega-analysis, as compared to a meta analysis; [Denny et al. 2012](#)) relevant to the question of how these processes are distinctly represented, we provide novel evidence that self- and other-related processing can be reliably distinguished using patterns of activity restricted to voxels within the MPFC. We also found that self- and other-related processing can be reliably distinguished using activity apparent across the whole brain, with differential activation in ACC and VMPFC particularly connected to self-related thought, and PCC, left angular gyrus and left temporal lobe particularly connected to other-related thought. In addition to quantitatively evaluating the explanatory power of the MPFC (and whole brain) for discriminating between self- and other-related thought, we also report data supporting the hypothesis that self- and other-related processing are functionally distributed along a ventral to dorsal gradient within the medial frontal wall ([Denny et al. 2012](#)). However, we also find that a purely linear pattern does not fully explain the effect; rather, self-related thought is most reliably represented within the middle prefrontal cortex and other-related thought within both the OFC and DMPFC, suggesting that curvilinear, rather than linear, gradient might characterize the organization of self- and other-related cognition within the medial wall of the frontal cortex.

In our analysis of the MPFC, we developed a model to discriminate between self- and other-related thought. This model performed well out-of-sample, correctly predicting 59% of our test observations. After training and testing this model, we also investigated how the multivariate pattern of weights were organized within the MPFC, and tested whether model weights followed a linear ventral–dorsal gradient. Here, our analysis of voxel weights and their respective position along the axial plane suggests a more complicated pattern that cannot be described with a simple linear gradient. Our results highlighting a more complex pattern are also consistent with previous reviews of the MPFC that note a significant overlap in how self- and other-related processes are represented in the MPFC ([Wagner et al. 2012](#)), and also help explain how self- and other-related thought can appear overlapping (see [Denny et al. 2012](#)), but still be distinguishable within the MPFC.

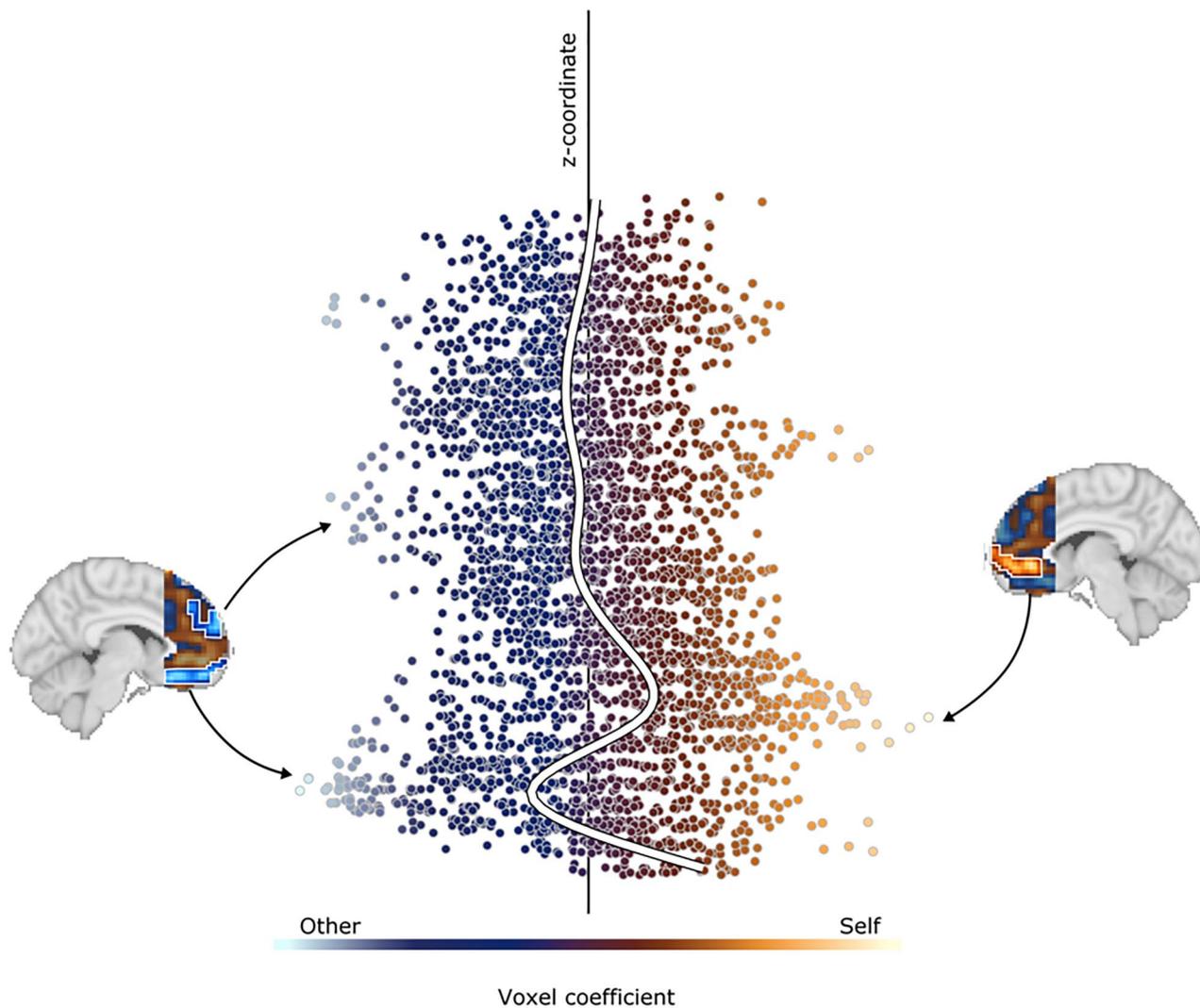


Figure 1. Organization of unthresholded voxel weights from the MPFC along the axial plane. Lighter blue indicates stronger weight and encoding for other-related processing, lighter orange indicates stronger weight and encoding for self-related processing. Z-coordinates for this graph are normalized between 0 and 1, the lowest z-coordinate in this analysis was $z = -30$ and the highest $z = 70$. Highlighted voxels indicate clusters associated with peak weights represented in the distribution. Peak weights for self-related processing were centered around $z = -2$, peak weights for other-related processing were centered around $z = -14$ and $z = 24$. A Loess regression was used to generate the smoothed line in the figure.

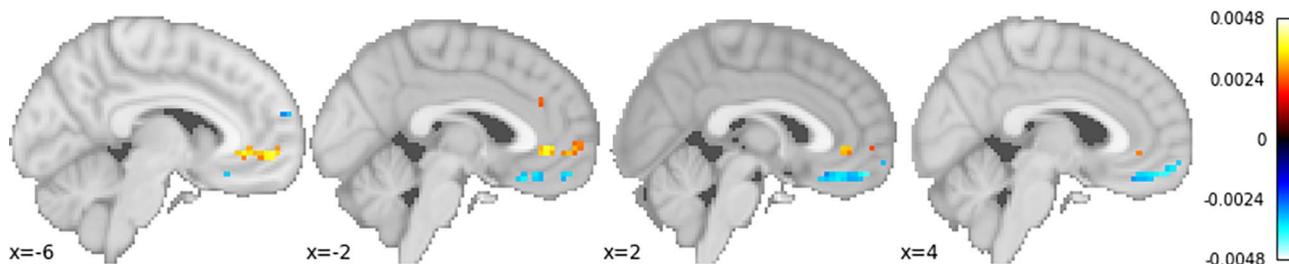


Figure 2. MPFC Ridge-PCR model weights (FDR corrected, $\alpha = 0.05$), back projected onto the whole brain. These results highlight most robust patterns differentiating self-related thought in the VMPFC, and other-related thought in the OFC and a small portion of the DMPFC. Brighter orange indicates greater weight towards encoding self-related thought, brighter blue indicates greater weight towards encoding other-related thought.

We found that voxels that were more strongly and reliably associated with the encoding of self-related (vs. social) processing (controlling for all other activation in the MPFC) were found within the ventral portion of the MPFC and extended

upwards through the ACC. This result falls in line with a wealth of previous research linking middle subregions of the MPFC with self-related thought (Legrand and Ruby 2009; Heatherton 2011). However, as shown in our whole-brain analysis (discussed

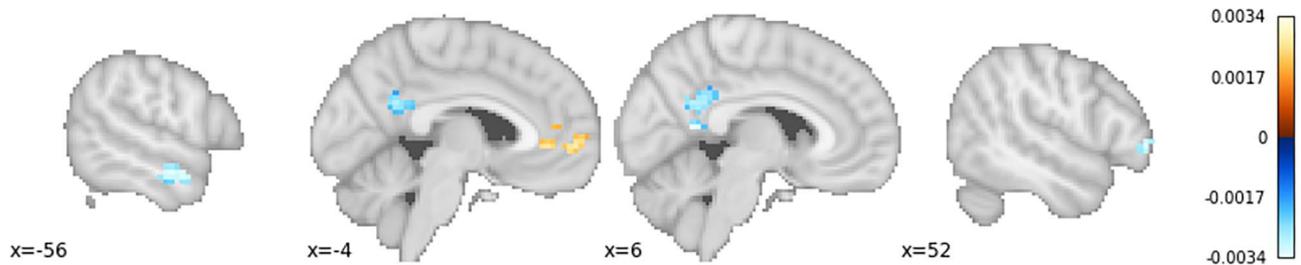


Figure 3. Whole Brain Ridge-PCR model weights (FDR corrected, $\alpha=0.05$), back projected onto whole brain. These results highlight the most robust patterns differentiating self-related thought in the VMPFC, and other-related thought in the PCC, angular gyrus, and temporal lobe. Brighter orange indicates greater weight towards encoding self-related thought, brighter blue indicates greater weight towards encoding other-related thought.

below) and other research (Rose Addis and Tippett 2008; Murray et al. 2012), these areas of the MPFC are not solely responsible for encoding self-related cognition or distinguishing between thinking about oneself from thinking about others. Other regions distributed throughout the human cortex also contribute to this distinct form of cognition. These results specifically indicate that when analysis is restricted to the MPFC, a region that has shown notable functional overlap of self- and other-related thinking, middle subregions stand out as reliably encoding self-related thought.

The most influential voxels coding for other-related thought, however, were found in both the DMPFC and VMPFC, including the orbitofrontal cortex. Dorsal portions of the MPFC are frequently implicated in thinking about the traits and thoughts of other people (Mitchell et al. 2005; Baron et al. 2011), and so our findings complement and extend prior understanding of this subregion. Further, supplemental analyses using the current dataset, including a voxel-wise univariate contrast and classification with average activation in ventral and dorsal portions of the MPFC, bolster our confidence in the current findings. Comparable voxel patterns to our primary findings were found in these additional analyses (Supplementary Materials), indicating that the organization of voxel weights found in our primary analyses are robust to analytic approaches. Lastly, we also found the organization of voxel weights to be robust when we trained the Ridge-PCR model on each of the three studies included in this mega-analysis separately. Studies 1 and 2 were most similar to the full sample model, but study 3 provided a less clear pattern; a result we attribute to its smaller sample size and alternative task design (see Supplementary Materials).

In addition to DMPFC, we also find that the OFC is important in reflecting on information about other people. This aligns with previous findings that the OFC is involved in social cognition (Beer 2006; Völlm et al. 2006; Weaverdyck et al. 2021), and in particular is sensitive to differences in trait judgments between the self and social targets (Beer and Hughes 2010; Hughes and Beer 2012). These findings also extend previous investigations of self- and other-related processing in MPFC (e.g., Denny et al. 2012) by further highlighting the OFC's role in social thinking. Our use of continuous data in a mega-analysis, as opposed to summary statistics focused on peak activation from previously published work, highlights a stronger role of the VMPFC in social cognition than previously emphasized.

As reflected in our findings above, and a substantial body of prior research, MPFC is centrally involved in both self- and other-related processing. However, recent work also highlights the value of incorporating whole brain patterns for characterizing psychological states (Chang et al. 2015; Van Oudenhove

et al. 2020). Thus, in addition to our focus on MPFC, we also developed an additional model, which classified self- and other-related processing using information contained within the entire brain. The inclusion of this information generated a directional improvement in the performance of our model, such that our best performing whole-brain model improved out-of-sample prediction accuracy by nearly 9% relative to our best performing MPFC model, and 18% better than chance. Previous research employing MVPA techniques to classify higher order cognitive processes, such as social perception (Brosch et al. 2013) and different aspects of self-related thought (Oosterwijk et al. 2017), tend to achieve classification accuracy scores of roughly 10–15% greater than chance.

Using data from the entire brain also allowed us to understand the relative contribution of the MPFC in encoding self- and other-related thought. Comparing the voxel weights from our best performing MPFC model and best performing whole-brain model, we saw that many of the most important voxels identified in the MPFC model remained among the most predictive within the whole-brain model. These consistent feature weights lend even stronger support for the notion that the MPFC is a critical brain region in processing both cognition relating to oneself and cognition relating to other people. The whole-brain results also provide additional insights into what other areas of the brain are important for processing information about other people relative to information about the self. Specifically, consistent with past meta-analyses (e.g., Denny et al. 2012; Murray et al. 2012), we found that regions associated with the default mode network (e.g., PCC, temporal lobe and angular gyrus; Raichle 2015) also contained the most strongly weighted voxels encoding for differentiating other-related thought from self-related thought. The PCC and MPFC are associated with both self and social cognition (Saxe and Powell 2006; Qin and Northoff 2011; Brewer et al. 2013; Mahy et al. 2014; Wagner et al. 2019); we also provide evidence from our whole brain analysis that these regions are important for processing information about other people differentially from the self.

Machine learning techniques in neuroimaging are becoming increasingly popular within the field, and can be leveraged to generate externally valid models that can be iteratively refined and tested in additional contexts. Thoughtful interpretation of these models and their weights is necessary though, as they are not a direct measure of change in voxel activation in response to study manipulations. Coefficients from these types of models represent the relative contribution of brain regions in discriminating between conditions and the direction of their associations. The back-projected maps from the models presented here help to describe the relative influence of brain regions

in coding for self against other related thought, and can be further contextualized by comparing them with more traditional contrast maps (see [Supplementary Materials](#)). In addition to careful interpretation of voxel weights, researchers must also be mindful of the bias-variance tradeoff implicit in modeling and prediction procedures, and how this affects meaningful inference. Here, we have made efforts to implement several techniques like stratified k -fold crossvalidation and out-of-sample bootstrapping to bolster confidence in the generalizability of our findings.

Looking forward, future multivariate methods like multivoxel searchlight analysis, will further help to understand the role of the MPFC in processing self- and other-related thought. More specifically, these methods can help to understand *where* in the MPFC these two processes are most reliably distinguishable, a question not answered by the analytic strategy taken here. The current study utilized data from three independent studies, which all asked individuals to explicitly judge the traits of themselves and other people, affording a relatively large overall dataset yielding a complementary high-precision method for identifying mechanisms of self and social cognition when compared to traditional coordinate-based meta-analyses.

Summary and Conclusion

The current study found that self- and other-related thought are represented in partially distinct regions within the MPFC. Self- and other-related cognition were represented in both dorsal and ventral regions of the MPFC, and were weakly organized along the ventral–dorsal gradient previously proposed, with our results clarifying that clusters of VMPFC and OFC are also particularly important for distinguishing social thought. Whole brain analysis further confirmed that the MPFC was primarily responsible for representing these processes, but that the PCC and other regions of the default mode network were also involved in processing thoughts relating to other people. We provide here a key step in applying machine learning techniques to develop neural models of self- and other-related processing.

Positionality Statement

Mindful that our identities can influence our approach to science ([Roberts et al. 2020](#)), the authors wish to provide the reader with potentially relevant information about our backgrounds. With respect to race/ethnicity, one author self-identifies as Asian; five as white. With respect to gender identity, 4 self-identify as men, 2 as women.

Citation Diversity Statement

Recent work in several fields of science has identified a bias in citation practices such that papers from women and other minority scholars are under-cited relative to the number of such papers in the field ([Maliniak et al. 2013](#); [Caplar et al. 2017](#); [Chakravartty et al. 2018](#); [Dion et al. 2018](#); [Thiem et al. 2018](#); [Dworkin et al. 2020](#); [Zhou et al. 2020](#)). Here, we sought to proactively consider choosing references that reflect the diversity of the field in thought, form of contribution, gender, race, ethnicity, and other factors. First, we obtained the predicted gender of the first and last author of each reference by using databases that store the probability of a first name being carried by a woman ([Dworkin et al. 2020](#); [Zhou et al. 2020](#)).

By this measure (and excluding self-citations to the first and last authors of our current paper), our references contain 24% woman(first)/woman(last), 21% man/woman, 20% woman/man, and 35% man/man. This method is limited in that (a) names, pronouns, and social media profiles used to construct the databases may not, in every case, be indicative of gender identity and (b) it cannot account for intersex, nonbinary, or transgender people. We look forward to future work that could help us to better understand how to support equitable practices in science.

Supplementary Material

[Supplementary material](#) can be found at [Cerebral Cortex](#) online.

Notes

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References

- Abraham A, Pedregosa F, Eickenberg M, Gervais P, Mueller A, Kossaifi J, Gramfort A, Thirion B, Varoquaux G. 2014. Machine learning for neuroimaging with scikit-learn. *Front Neuroinform*. 8:14.
- Barker M. 2010. *Partial least squares for discrimination: statistical theory and implementation*. Maude Avenue, Sunnyvale, CA: LAP Lambert Academic Publishing.
- Baron SG, Gobbini MI, Engell AD, Todorov A. 2011. Amygdala and dorsomedial prefrontal cortex responses to appearance-based and behavior-based person impressions. *Soc Cogn Affect Neurosci*. 6(5):572–581.
- Beer JS. 2006. Orbitofrontal cortex and social regulation. In: *Social neuroscience: people thinking about thinking people*, pp. 41–62.

- Beer JS, Hughes BL. 2010. Neural systems of social comparison and the “above-average” effect. *Neuroimage*. 49(3):2671–2679.
- Beer JS, Lombardo MV, Bhanji JP. 2010. Roles of medial prefrontal cortex and orbitofrontal cortex in self-evaluation. *J Cogn Neurosci*. 22(9):2108–2119.
- Bergstrom ZM, Vogelsang DA, Benoit RG, Simons JS. 2015. Reflections of oneself: neurocognitive evidence for dissociable forms of self-referential recollection. *Cereb Cortex*. 25(9):2648–2657.
- Blakemore SJ. 2008. The social brain in adolescence. *Nat Rev Neurosci*. 9(4):267–277.
- Bower GH, Gilligan SG. 1979. Remembering information related to one’s self. *J Res Pers*. 13(4):420–432.
- Brewer J, Garrison K, Whitfield-Gabrieli S. 2013. What about the “self” is processed in the posterior cingulate cortex? *Front Hum Neurosci*. 7:647.
- Brosch T, Bar-David E, Phelps EA. 2013. Implicit race bias decreases the similarity of neural representations of black and white faces. *Psychol Sci*. 24(2):160–166.
- Caplar N, Tacchella S, Birrer S. 2017. Quantitative evaluation of gender bias in astronomical publications from citation counts. *Nat Astron*. 1(6):1–5.
- Chakravartty P, Kuo R, Grubbs V, McIlwain C. 2018. Communication SoWhite. *J Commun*. 68(2):254–266.
- Chang LJ, Gianaros PJ, Manuck SB, Krishnan A, Wager TD. 2015. A sensitive and specific neural signature for picture-induced negative affect. *PLoS Biol*. 13(6):e1002180.
- Cooper N, Tompson S, O’Donnell MB, Falk EB. 2015. Brain activity in self- and value-related regions in response to online anti-smoking messages predicts behavior change. *J Media Psychol*. 27:93–109.
- Corradi-Dell’Acqua C, Tusche A, Vuilleumier P, Singer T. 2016. Cross-modal representations of first-hand and vicarious pain, disgust and fairness in insular and cingulate cortex. *Nat Commun*. 7:10904.
- Courtney AL, Meyer ML. 2020. Self-other representation in the social brain reflects social connection. *J Neurosci Off J Soc Neurosci*. 40(29):5616–5627.
- Cox RW. 1996. AFNI: software for analysis and visualization of functional magnetic resonance neuroimages. *Comput Biomed Res*. 29(3):162–173.
- Craik FIM, Moroz TM, Moscovitch M, Stuss DT, Winocur G, Tulving E, Kapur S. 1999. In search of the self: a positron emission tomography study. *Psychol Sci*. 10(1):26–34.
- D’Argembeau A, Collette F, Van der Linden M, Laureys S, Del Fiore G, Degueldre C, Luxen A, Salmon E. 2005. Self-referential reflective activity and its relationship with rest: a PET study. *Neuroimage*. 25(2):616–624.
- D’Argembeau A, Ruby P, Collette F, Degueldre C, Baeteau E, Luxen A, Maquet P, Salmon E. 2007. Distinct regions of the medial prefrontal cortex are associated with self-referential processing and perspective taking. *J Cogn Neurosci*. 19(6):935–944.
- Denny BT, Kober H, Wager TD, Ochsner KN. 2012. A meta-analysis of functional neuroimaging studies of self- and other judgments reveals a spatial gradient for mentalizing in medial prefrontal cortex. *J Cogn Neurosci*. 24(8):1742–1752.
- De Pisapia N, Barchiesi G, Jovicich J, Cattaneo J. 2019. The role of medial prefrontal cortex in processing emotional self-referential information: a combined TMS/fMRI study. *Brain Imaging Behav*. 13:603–614.
- Dion ML, Sumner JL, Mitchell SM. 2018. Gendered citation patterns across political science and social science methodology fields. *Polit Anal*. 26(3):312–327.
- Dodell-Feder D, Koster-Hale J, Bedny M, Saxe R. 2011. fMRI item analysis in a theory of mind task. *Neuroimage*. 55(2):705–712.
- Dworkin JD, Linn KA, Teich EG, Zurn P, Shinohara RT, Bassett DS. 2020. The extent and drivers of gender imbalance in neuroscience reference lists. *Nat Neurosci*. 23(8):918–926.
- Falk EB, O’Donnell MB, Cascio CN, Tinney F, Kang Y, Lieberman MD, Taylor SE, An L, Resnicow K, Strecher VJ. 2015. Self-affirmation alters the brain’s response to health messages and subsequent behavior change. *Proc Natl Acad Sci USA*. 112(7):1977–1982.
- Frith CD, Frith U. 2006. The neural basis of mentalizing. *Neuron*. 50(4):531–534.
- Gallagher I. I. 2000. Philosophical conceptions of the self: implications for cognitive science. *Trends Cogn Sci*. 4(1):14–21.
- Gusnard DA, Akbudak E, Shulman GL, Raichle ME. 2001. Medial prefrontal cortex and self-referential mental activity: relation to a default mode of brain function. *Proc Natl Acad Sci USA*. 98(7):4259–4264.
- Hampton AN, Bossaerts P, O’Doherty JP. 2008. Neural correlates of mentalizing-related computations during strategic interactions in humans. *Proc Natl Acad Sci USA*. 105(18):6741–6746.
- Hassabis D, Spreng RN, Rusu AA, Robbins CA, Mar RA, Schacter DL. 2014. Imagine all the people: how the brain creates and uses personality models to predict behavior. *Cereb Cortex*. 24(8):1979–1987.
- Haxby JV, Gobbini MI, Furey ML, Ishai A, Schouten JL, Pietrini P. 2001. Distributed and overlapping representations of faces and objects in ventral temporal cortex. *Science*. 293(5539):2425–2430.
- Heatherston TF. 2011. Neuroscience of self and self-regulation. *Annu Rev Psychol*. 62:363–390.
- Heatherston TF, Wyland CL, Macrae CN, Demos KE, Denny BT, Kelley WM. 2006. Medial prefrontal activity differentiates self from close others. *Soc Cogn Affect Neurosci*. 1(1):18–25.
- Hoerl AE, Kennard RW. 1970. Ridge regression: biased estimation for nonorthogonal problems. *Technometrics: a journal of statistics for the physical, chemical, and engineering Sciences*. 12(1):55–67.
- Hughes BL, Beer JS. 2012. Orbitofrontal cortex and anterior cingulate cortex are modulated by motivated social cognition. *Cereb Cortex*. 22(6):1372–1381.
- James W. 1890. *The principles of psychology*, Vol II. New York, NY: Henry Holt and Company, Inc.
- Jenkins AC, Mitchell JP. 2010. Medial prefrontal cortex subserves diverse forms of self-reflection. *Soc Neurosci*. 6(3):211–218.
- Jenkins AC, Macrae CN, Mitchell JP. 2008. Repetition suppression of ventromedial prefrontal activity during judgement of self and others. *Proc Natl Acad Sci USA*. 105(11):4507–4512.
- Johnson MK, Raye CL, Mitchell KJ, Touryan SR, Greene EJ, Nolen-Hoeksema S. 2006. Dissociating medial frontal and posterior cingulate activity during self-reflection. *Soc Cogn Affect Neurosci*. 1(1):56–64.
- Kelley WM, Macrae CN, Wyland CL, Caglar S, Inati S, Heatherston TF. 2002. Finding the self? An event-related fMRI study. *J Cogn Neurosci*. 14(5):785–794.
- Koban L, Lee S, Schelski DS, Simon MC, Lerman C, Weber B, Kable JW, Plassmann H. 2021, March 19, 2021. An fMRI-based brain marker predicts individual differences in delay discounting. *bioRxiv*. doi: 10.1101/2021.03.18.435969.
- Koski JE, McHaney JR, Rigney AE, Beer JS. 2020. Reconsidering longstanding assumptions about the role of medial prefrontal cortex (MPFC) in social evaluation. *Neuroimage*. 214:116752.

- Koster-Hale J, Richardson H, Velez N, Asaba M, Young L, Saxe R. 2017. Mentalizing regions represent distributed, continuous, and abstract dimensions of others' beliefs. *Neuroimage*. 161:9–18.
- Krishnan A, Woo C-W, Chang LJ, Ruzic L, Gu X, López-Solà M, Jackson PL, Pujol J, Fan J, Wager TD. 2016. Somatic and vicarious pain are represented by dissociable multivariate brain patterns. *Elife*. 5:1–42.
- Legrand D, Ruby P. 2009. What is self-specific? Theoretical investigation and critical review of neuroimaging results. *Psychol Rev*. 116(1):252–282.
- Lou HC, Luber B, Crupain M, Keenan JP, Nowak M, Kjaer TW, Sackeim HA, Lisanby SH. 2004. Parietal cortex and representation of the mental self. *Proc Natl Acad Sci USA*. 101(17):6827–6832.
- Ma N, Baetens K, Vandekerckhove M, Kestemont J, Fias W, Van Overwalle F. 2014. Traits are represented in the medial prefrontal cortex: an fMRI adaptation study. *Soc Cogn Affect Neurosci*. 9(8):1185–1192.
- Mahy CEV, Moses LJ, Pfeifer JH. 2014. How and where: theory-of-mind in the brain. *Dev Cogn Neurosci*. 9:68–81.
- Maliniak D, Powers R, Walter BF. 2013. The gender citation gap in international relations. *Int Organ*. 67(4):889–922.
- Marquine MJ, Grilli MD, Rapcsak SZ, Kaszniak AW, Ryan L, Walther K, Glisky EL. 2016. Impaired personal trait knowledge, but spared other-person trait knowledge, in an individual with bilateral damage to the medial prefrontal cortex. *Neuropsychologia*. 89:245–253.
- Martial C, Stawarczyk D, D'Argembeau A. 2018. Neural correlates of context-independent and context-dependent self-knowledge. *Brain Cogn*. 125:23–31.
- Martinelli P, Sperduti M, Piolino P. 2013. Neural substrates of the self-memory system: new insights from a meta-analysis. *Hum Brain Mapp*. 34:1515–1529.
- Mende-Siedlecki P, Cai Y, Todorov A. 2013. The neural dynamics of updating person impressions. *Soc Cogn Affect Neurosci*. 8(6):623–631.
- Meyer ML, Collier E. 2020. Theory of minds: managing mental state inferences in working memory is associated with the dorsomedial subsystem of the default network and social integration. *Soc Cogn Affect Neurosci*. 15(1):63–73.
- Mitchell JP, Banaji MR, Macrae CN. 2005. General and specific contributions of the medial prefrontal cortex to knowledge about mental states. *Neuroimage*. 28(4):757–762.
- Mitchell JP, Heatherton TF, Macrae CN. 2002. Distinct neural systems subserve person and object knowledge. *Proc Natl Acad Sci USA*. 99(23):15238–15243.
- Murray RJ, Schaer M, Debbané M. 2012. Degrees of separation: a quantitative neuroimaging meta-analysis investigating self-specificity and shared neural activation between self- and other-reflection. *Neurosci Biobehav Rev*. 36(3):1043–1059.
- Norman KA, Polyn SM, Detre GJ, Haxby JV. 2006. Beyond mind-reading: multi-voxel pattern analysis of fMRI data. *Trends Cogn Sci*. 10(9):424–430.
- Ochsner KN, Knierim K, Ludlow DH, Hanelin J, Ramachandran T, Glover G, Mackey SC. 2004. Reflecting upon feelings: an fMRI study of neural systems supporting the attribution of emotion to self and other. *J Cogn Neurosci*. 16(10):1746–1772.
- Oosterwijk S, Snoek L, Rotteveel M, Barrett LF, Scholte HS. 2017. Shared states: using MVPA to test neural overlap between self-focused emotion imagery and other-focused emotion understanding. *Soc Cogn Affect Neurosci*. 12(7):1025–1035.
- Parkinson C, Kleinbaum AM, Wheatley T. 2017. Spontaneous neural encoding of social network position. *Nat Hum Behav*. 1(5):1–7.
- Pedregosa F, Varoquaux G, Gramfort A, Michel V, Thirion B, Grisel O, Blondel M, Prettenhofer P, Weiss R, Dubourg V, et al. 2011. Scikit-learn: machine learning in python. *J Mach Learn Res*. 12:2825–2830.
- Pereira F, Mitchell T, Botvinick M. 2009. Machine learning classifiers and fMRI: a tutorial overview. *Neuroimage*. 45(1):S199–S209.
- Pfeifer JH, Lieberman MD, Dapretto M. 2007. “I know you are but what am I!?”: neural bases of self- and social knowledge retrieval in children and adults. *J Cogn Neurosci*. 19(8):1323–1337.
- Philippi CL, Duff MC, Denburg NL, Tranel D, Rudrauf D. 2012. Medial PFC damage abolishes the self-reference effect. *J Cogn Neurosci*. 24(2):475–481.
- Platt J. 1999. Probabilistic outputs for support vector machines and comparisons to regularized likelihood methods. *Advances in Large Margin Classifiers*. 10(3):61–74.
- Qin P, Northoff G. 2011. How is our self related to midline regions and the default-mode network? *Neuroimage*. 57(3):1221–1233.
- Raichle ME. 2015. The brain's default mode network. *Annu Rev Neurosci*. 38:433–447.
- Rameson LT, Satpute AB, Lieberman MD. 2010. The neural correlates of implicit and explicit self-relevant processing. *Neuroimage*. 50(2):701–708.
- Rissman J, Gazzaley A, D'Esposito M. 2004. Measuring functional connectivity during distinct stages of a cognitive task. *Neuroimage*. 23(2):752–763.
- Roberts SO, Bareket-Shayit C, Dollions FA, Goldie PD, Mortenson E. 2020. Racial inequality in psychological research: trends of the past and recommendations for the future. *Perspect Psychol Sci*. 15(6):1295–1309.
- Rogers TB, Kuiper NA, Kirker WS. 1977. Self-reference and the encoding of personal information. *J Pers Soc Psychol*. 35(9):677–688.
- Rose Addis D, Tippett LJ. 2008. The contributions of autobiographical memory to the content and continuity of identity: a social cognitive neuroscience approach. In: *Self continuity individual and collective perspectives*. Psychology Press, p. 71–84.
- Saxe R. 2006. Uniquely human social cognition. *Curr Opin Neurobiol*. 16(2):235–239.
- Saxe R, Powell LJ. 2006. It's the thought that counts: specific brain regions for one component of theory of mind. *Psychol Sci*. 17(8):692–699.
- Saxe R, Moran JM, Scholz J, Gabrieli J. 2006. Overlapping and non-overlapping brain regions for theory of mind and self reflection in individual subjects. *Soc Cogn Affect Neurosci*. 1(3):229–234.
- Saxe R, Waxler A. 2005. Making sense of another mind: the role of the right temporo-parietal junction. *Neuropsychologia*. 43(10):1391–1399.
- Schmitz TW, Johnson SC. 2006. Self-appraisal decisions evoke dissociated dorsal-ventral aMPFC networks. *Neuroimage*. 30(3):1050–1058.
- Schmitz TW, Kawahara-Baccus TN, Johnson SC. 2004. Metacognitive evaluation, self-relevance, and the right prefrontal cortex. *Neuroimage*. 22(2):941–947.
- Seger CA, Stone M, Keenan JP. 2004. Cortical activations during judgments about the self and an other person. *Neuropsychologia*. 42(9):1168–1177.

- Shen X, Tokoglu F, Papademetris X, Constable RT. 2013. Group-wise whole-brain parcellation from resting-state fMRI data for network node identification. *Neuroimage*. 82:403–415.
- Skerry AE, Saxe R. 2014. A common neural code for perceived and inferred emotion. *J Neurosci*. 34(48):15997–16008.
- Tamir DI, Mitchell JP. 2010. Neural correlates of anchoring-and-adjustment during mentalizing. *Proc Natl Acad Sci USA*. 107(24):10827–10832.
- Thiem Y, Sealey KF, Ferrer AE, Trott AM, Kennison R. 2018. *Just ideas? The status and future of publication ethics in philosophy: a white paper*, Technical report.
- van der Meer L, Costafreda S, Aleman A, David AS. 2010. Self-reflection and the brain: a theoretical review and meta-analysis of neuroimaging studies with implications for schizophrenia. *Neurosci Biobehav Rev*. 34(6):935–946.
- Van Oudenhove L, Kragel P, Dupont P, Ly HG, Pazmany E, Enzlin P, Rubio A, Delon-Martin C, Bonaz B, Aziz Q, et al. 2020. Common and distinct neural representations of aversive somatic and visceral stimulation in healthy individuals. *Nat Commun*. 11(1):1–11.
- Van Overwalle F. 2009. Social cognition and the brain: a meta-analysis. *Hum Brain Mapp*. 30(3):829–858.
- Verfaellie M, Wank AA, Reid AG, Race E, Keane MM. 2019. Self-related processing and future thinking: distinct contributions of ventromedial prefrontal cortex and the medial temporal lobes. *Cortex*. 115:159–171.
- Völlm BA, Taylor ANW, Richardson P, Corcoran R, Stirling J, McKie S, Deakin JFW, Elliot R. 2006. Neuronal correlates of theory of mind and empathy: a functional magnetic resonance imaging study in a nonverbal task. *Neuroimage*. 28(1):90–98.
- Wager TD, Atlas LY, Leotti LA, Rilling JK. 2011. Predicting individual differences in placebo analgesia: contributions of brain activity during anticipation and pain experience. *J Neurosci*. 31(2):439–452.
- Wagner DD, Chavez RS, Broom TW. 2019. Decoding the neural representation of self and person knowledge with multivariate pattern analysis and data-driven approaches. *Wiley Interdiscip Rev Cogn Sci*. 10(1):e1482.
- Wagner DD, Haxby JV, Heatherton TF. 2012. The representation of self and person knowledge in the medial prefrontal cortex. *Wiley Interdiscip Rev Cogn Sci*. 3(4):451–470.
- Weaverdyck ME, Thornton MA, Tamir DI. 2021. The representational structure of mental states generalizes across target people and stimulus modalities. *Neuroimage*. 238:118258.
- Wheeler MA, Stuss DT, Tulving E. 1997. Toward a theory of episodic memory: the frontal lobes and autonoetic consciousness. *Psychol Bull*. 121(3):331–354.
- Zhou D, Cornblath EJ, Stiso J, Teich EG, Dworkin JD, Blevins AS, Bassett DS. 2020. *Gender Diversity statement and code note-book v1.0*. Zenodo.