doi:10.1093/scan/nsp001 SCAN (2012) 7, 521–534

Common and distinct brain networks underlying explicit emotional evaluation: a meta-analytic study

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Brain mechanisms underlying explicit evaluation of emotion have been explored using different tasks including 'stimulus-focused evaluation', 'evaluation of one's own emotion' and 'evaluation of others' emotions'. Yet the extent to which similar brain mechanisms underlie different evaluation tasks is unclear. A meta-analysis of published neuroimaging studies of explicit emotional evaluation was conducted to examine common and distinct regions underlying these different evaluation tasks. This study revealed regions common to all three tasks: The amygdala and LPFC as common regions may be involved in emotion—cognition interactions, and the DMPFC may possibly play integrative roles in explicit emotional evaluation. Distinct regions were also identified: (i) the sensory cortex and VLPFC were specifically associated with 'stimulus evaluation', possibly involved in perceptual and conceptual processing; (ii) the insula and rACC were specifically associated with 'evaluation of one's own emotion', potentially associated with interoceptive and experiential processing; and (iii) the STS and TPJ were specifically associated with 'evaluation of others' emotions', potentially reflecting their roles in TOM and empathy. These findings suggest that different types of explicit emotional evaluation may involve common and distinct networks and provide new insights on multiple mechanisms underlying explicit emotional evaluation.

Keywords: explicit evaluation; emotion; neuroimaging; meta-analysis

INTRODUCTION

This study examines brain mechanisms associated with different types of explicit emotional evaluation. We define explicit evaluation of emotion as the processes of how people interpret, evaluate and judge emotional meanings and value (either valence or discrete emotions) of objects (Cunningham et al., 2003; Barrett, 2005; Nielsen and Kaszniak, 2007). During explicit evaluation of emotion, people evaluate emotional value in conscious and deliberate manners and verbalize their responses. Explicit evaluation of emotion allows people to label their emotions, and to express or report current or past emotions, so that they can explain their emotions and share them with other people. However, due to limitations of an imaging environment, explicit emotional evaluation should be simplified and restricted; for example, subjects are asked to evaluate a specific emotional value (e.g. valence, happy and sad, or emotional intensity) under specific task instructions and respond by pressing a button in the scanner, a task termed 'on-line' in this article.

Neuroimaging studies have identified brain regions that are activated more by explicit than implicit/automatic

Received 1 February 2008; Accepted 31 December 2008 Advance Access publication 6 March 2009

We thank Walter Schneider, Julie Fiez and William Klein, for comments on previous versions of this manuscript. This work was supported by N00014-05-1-0881 and MH074807.

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emotional processing (for a review, see Phan *et al.*, 2002; Hutcherson *et al.*, 2005). However, results are inconsistent; for example, (i) increased activation (Liberzon *et al.*, 2000; Gorno-Tempini *et al.*, 2001; Lee *et al.*, 2004; Vollm *et al.*, 2006), decreased activation (e.g. Critchley *et al.*, 2000; Hariri *et al.*, 2000; Taylor, Phan, Decker *et al.*, 2003), or no different activation in the amygdala (e.g. Hutcherson *et al.*, 2005) in explicit emotional evaluation compared to control conditions; and (ii) increased activation in the medial prefrontal cortex (MPFC; Taylor *et al.*, 2003) and in the lateral PFC (e.g. Gorno-Tempini *et al.*, 2001; Lee *et al.*, 2004), or both PFC regions (e.g. Hutcherson *et al.*, 2005).

One potential reason for such discrepancies is that previous studies have used different explicit evaluation tasks, which may depend on diverse brain networks, to examine underlying brain mechanisms of explicit evaluation of emotion. The extent to which similar brain mechanisms underlie such different evaluation tasks is unclear. Perhaps there are central shared mechanisms for explicit evaluation of emotion regardless of evaluation tasks or distinct mechanisms depending on emotional evaluation tasks.

In particular, previous studies have used tasks that ask participants to explicitly evaluate (i) the emotionality of stimuli/situations (e.g. pushing a button for whether a picture or word is positive, negative, or neutral in tone; for example, Critchley *et al.*, 2000; Hariri *et al.*, 2000; Narumoto *et al.*, 2000), (ii) one's own emotional states (e.g. 'How do

you feel?' Gusnard *et al.*, 2001; Ochsner *et al.*, 2004a; Hutcherson *et al.*, 2005) and (iii) other people's emotional states (e.g. 'How does the person of this picture feel?') (e.g. Mitchell *et al.*, 2005; Jackson *et al.*, 2006; Vollm *et al.*, 2006). Answers to these questions may differ. For example, a picture of an evil-doer being apprehended may simultaneously yield a positive feeling for the observer, but an assertion that the evil-doer is feeling upset.

Two lines of evidence suggest the specific importance of these three domains. Developmental psychologists emphasize that emotional knowledge acquired by engaging with physical stimuli/situations, one's own body and other people is important for normal emotional development (Trevarthen and Aitken, 2001). Self-report studies also demonstrate that people describe specific situations, their own internal states and other people's internal states when they talk about emotion (Davitz, 1969; Stein and Levine, 1999; Barrett *et al.*, 2001). Thus, the following sections will concentrate on these three domains.

Emotion-cognition interactions are hypothesized to be common to all three types of emotional evaluation (Frijda and Zeelenberg, 2001; Roseman and Smith, 2001; Scherer, 2001; Lewis, 2005). Thus, we hypothesized that neural substrates of interactions between cognition and emotion would be common to all three tasks. Such interactions, from emotional evaluation to emotion regulation to emotional interference with basic cognitive processes are subserved by several brain regions which were used as a priori structures in the present investigation. These included several prefrontal cortex (PFC) regions including the caudal anterior cingulate cortex (cACC), ventromedial, dorsomedial and dorsolateral prefrontal cortices (VMPFC, DMPFC, DLPFC), and limbic regions such as amygdala, thalamus, hippocampus and orbitofrontal cortex (OFC; see Table 1; for a review, see Lewis, 2005).

In addition, the three specific evaluation tasks may also each recruit distinct mechanisms.

For example, stimulus-focused evaluation may focus on physical features and meaning of stimuli/situations (e.g. Arnold, 1960; Roseman, 1984) and include interpretation of perceived information in terms of conceptual knowledge of stimuli in the representation and memory system (e.g. Leventhal and Scherer, 1987; Smith and Kirby, 2001). Perceptual processing may rely on modality of a stimulus or types of stimuli, indicating that this processing activates sensory cortex regions relevant to the modality such as visual stimuli (i.e. faces and words) and visual cortex (i.e. face fusiform area: FFA; for example, Adolphs, 2002a, b). Higher level of conceptual processing is associated with the ventrolateral PFC (VLPFC: BA47), a core brain region involved in explicit evaluation of a stimulus (Cunningham et al., 2004) and evaluation of object knowledge (e.g. Mitchell et al., 2002). Thus, this evidence suggests the sensory cortices and VLPFC (BA47) may play a key role in evaluating emotionality of stimuli/situations.

Table 1 Summary of hypothesized brain regions associated with theoretical accounts

Theoretical accounts	Possible brain networks (BA)
General mechanisms	Common brain networks
Emotion—cognition interactive processing	PFC regions (e.g. LPFC, MPFC) Subcortical-Limbic regions (e.g. amygdala)
Evaluative and regulatory processing	DMPFC (BA10/32) and PCC
Task-dependent mechanisms	Distinct brain networks
Stimulus-focused evaluation	
Perceptual processing	Sensory-related brain regions (e.g. FFA)
High-level of conceptual processing	VLPFC (BA47)
Evaluation of one's own emotion	
Interoceptive processing	Insula (BA13) and rACC (BA24/32)
Experiential processing	rACC (BA24/32)
Evaluation of other's emotions	
Inferential processing (Theory of Mind)	STS (BA22/39) and TPJ (BA39/40)
Empathetic processing	STS and TPJ
(Perspective-taking)	

BA: Brodmann Area; LPFC: Lateral Prefronal Cortex (PFC); MPFC: Medial PFC; DMPFC: Dorsomedial PFC; PCC: Posterior Cingulate Cortex; FFA: Fusiform Face Area; VLPFC: Ventrolateral PFC; rACC: rostral Anterior Cingulate; STS: Superior Temporal Sulcus; TPJ: Temporo-Parietal Junction.

In contrast, evaluation of one's own emotion may involve inferring emotional experience from inner states and conscious thoughts about current and past experience. Evaluative processing of one's own emotion may thus include evaluation of bodily experience (interoception), conscious thoughts and a felt action tendency (for a review, see Lane, 2000; Lambie and Marcel, 2002). The insula is a key structure associated with subjective awareness of inner states (Craig, 2002, 2004) and has been implicated in subjective interoceptive and emotional states (Critchley et al., 2004). The rostral ACC (rACC) has also been implicated in the representation of conscious emotional experience (Lane, 2000; Lane and McRae, 2004; Barrett et al., 2007). Therefore, the insula and rACC are hypothesized to play specific roles in evaluating one's own emotional experience.

Evaluation of others' emotions involves understanding others' psychological properties including beliefs, intention and emotion. Theory of Mind (TOM) suggests that people have mental state concepts such as belief and intention, so they use this explicit knowledge or rules to infer others' mental states (for a review, see Gallese and Goldman, 1998). Another theoretical account, simulation theory (ST), proposed that people put themselves 'in the other person's shoes' to infer other people's mental states (Gallese and Goldman, 1998; Goldman and Sripada, 2005). Similarly, empathy requires perspective taking and simulation of others' emotions (e.g. Decety and Jackson, 2004). Neuroimaging studies have revealed brain regions associated with TOM and ST. The superior temporal sulcus (STS) is implicated in understanding of others' intentionality (Frith and Frith, 1999; Gallagher and Frith, 2003) and the temporal

poles and temporo-parietal junction (TPJ) are involved in reasoning about others' mental states (Blakemore *et al.*, 2004; Saxe, 2006). Thus, these brain regions may be particularly associated with evaluation of others' emotions.

To summarize, well-documented theoretical and empirical evidence allows us to hypothesize brain networks underlying explicit emotional evaluation. Therefore, our meta-analytic study focused on mainly these hypothesized brain regions. Table 1 presents theoretical accounts and possible brain networks associated with general/shared and task-dependent mechanisms underlying explicit evaluation of emotion which served as our hypotheses.

To understand the extent to which such shared and distinct regions are associated with different types of emotional evaluation, we conducted a quantitative meta-analytic review of published neuroimaging studies. Neuroimaging studies were, therefore, categorized into three different tasks to evaluate emotion consciously including (i) evaluating the emotionality of a stimulus (stimulus-focused emotion), (ii) evaluating one's own emotion and (iii) evaluating others' emotions. Overlapping brain regions among neuroimaging studies using different evaluation tasks were considered as shared/common brain mechanisms. Distinct regions among these studies were interpreted as task-dependent mechanisms.

METHODS

Studies were identified primarily by searching the PUBMED database. Keywords such as emotion, evaluation, cognition and imaging (fMRI and PET) were used to find neuro-imaging studies related to emotion and evaluation, which yielded over 100 studies available until 2006. Another search was performed with more specific keywords such as identification, recognition, judgment, self-reports, experience and empathy. A final search reviewed reference lists in the identified papers was performed. The title and abstract of each study were checked to decide whether the study used online evaluation tasks to assess brain mechanisms of explicit emotional evaluation.

Papers were included if they met the following criteria: (i) They involved healthy participants; (ii) They used on-line tasks to evaluate emotions which subjects evaluated explicitly emotion in the scanner; (iii) They used trial-based emotional evaluation (e.g. evaluation per trial/stimulus) in either block or event-related designs; (iv) They collected imaging data from the whole brain and used either wholebrain analysis or both ROI-based and whole-brain analysis; (v) They reported standard Talairach (Talairach and Tournoux, 1988) or Montreal Neurologic Institute (MNI) coordinates. The 37 studies identified were assigned into three groups. Studies were categorized as stimulus-focused evaluation (STIM) if they used tasks to focus on evaluating emotionality of stimuli including explicit identification, judgment or labelling of emotional stimuli. Studies were categorized as evaluation of one's own emotion (SELF) if they used an evaluation task which required subjects to pay attention to their own emotional states. Studies were categorized as *evaluation of others' emotions* (OTHER) if they used any tasks that give instructions about evaluating others' emotion and pain or about judging empathy of other people.

Table 2 quantitatively summarizes methodological information including the sample size, evaluation tasks, types and modalities of stimuli, emotions and contrasts in the identified studies. Specific tasks of emotional evaluation were broadly assigned to four categories including 'identification', 'labeling', 'matching' and 'intensity' tasks. The identification task was defined as any tasks demanding subjects to assess stimuli, their own emotion or others' emotions based on emotional valence (e.g. positive and negative), discrete emotions (e.g. happy, sad and fear) or social emotions (e.g. shame). The labelling task was involved in judging stimuli by two linguistic labels that were presented on the screen (Hariri et al., 2000). In the matching task, subjects matched sampled faces/words with one of two faces (Narumoto et al., 2000). The intensity task included any tasks if the subjects evaluated emotional intensity of stimuli, their own emotion (arousal) or others' emotions. As summarized in Table 2, most studies have used either identification or intensity tasks, but the STIM more frequently used identification than intensity task, whereas two other groups used both tasks equally. The STIM group was more heterogeneous in stimulus modality including auditory and olfactory compared to the two other groups. In addition, the STIM and OTHER groups were more diverse in emotions such as discrete emotions and social emotions than the SELF group. Most studies contrasted explicit conditions with implicit conditions (e.g. passive viewing or gender task) whereas some studies contrasted emotion conditions with neutral conditions. Some studies also used both contrast methods.

To determine whether hypothesized brain regions were activated in each study, reported coordinates were checked manually. The coordinates were recorded separately in the left and right side of a priori ROIs for this study (see Table 1). The ROIs were defined based on the Brodmann Areas (BA) and previous literature (e.g. Bush et al., 2000; Amodio and Frith, 2006). For example, rACC (BA24/32) and cACC (BA24) were defined based on the boundaries used in Bush and colleagues (2000). The coordinates in each study were checked for possible location on one of the ROIs. If several coordinates were reported in the same brain regions, one coordinate with the highest statistical values (e.g. z scores or t values) was selected as a representative to control the relative contributions by each study. Relative distance among these coordinates within the same regions was not considered to avoid some possible biases by a coder in picking the coordinates. Figure 1 shows all the regions of interests and coordinates (peaks) in these regions. Coordinates from neuroimaging studies that reported

Table 2 List of neuroimaging studies in this meta-analytic study

First author)	Imaging studies (First author)		naging N Tech	Speci	ific Task	(Sti	mulu	IS	Emotion																Contrast		
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Nakamura PET 7 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0				lden	Label	Mat	Int	Fa	W	Pic	Film	Sent	W	Sent	Snd		Pos	Neg	Ar			Н	S	D	F	Ang	Sur	Ex vs I	E vs N
Hariff Milk 18 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	STIM (n = 20)																												
Namumoto MRII 8				0													0	0											
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Gompi Fight 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0						0																	0	0	0		0		
Lange MRIN 9																						-				0			
Winston MRI 12				0			0															0		0	٥			0	0
Pandlso PET 17 0																						٥	٥	٥				0	U
Liberon PET 10				0			U	U		٥							0	0				U	U	U	U			U	0
Taylor PET 10				U			0										U											0	0
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Maddock MARI 24 0				-			0		0	-							-											-	0
Imaizumi	Maddock	fMRI	8	0													0	0											0
Imaizumi	Cunningham	fMRI	24	0					0								0	0										0	
Wildgrider MRI 10 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0		PET	6	0									0									0		0		0	0	0	
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Royer PET 12 2 0 0 0 0 0 0 0 0	Wildgruber	fMRI	10	0										0								0	0	0	0	0		0	
September MRI	Royet	PET								0					0	0	0	0										0	0
Subtotals	Royet						0									0	0	0										0	
SELF (n = 9) Causard MRI 24 0	Royet	fMRI	28	0												0	0	0										0	
Lane PET 10 0 0 0 0 0 0 0 0 0	Subtotals			12	1	1	7	7	3	5	0	0	1	2	1	3	8	11	1	0	0	6	3	5	5	5	3	16	7
Gusnard fMRI 13 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0																													
Ochsner* MRI 13 0																													
Lee fMRI 10																													
Phan MRI 12 0				0																								0	
Garett fMRI 9							0																					_	0
Jackson a MRI 18				0			•										0											0	
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Rubysh PET 10 0										0	•									0		^	0						
OTHER (n = 8) Mitchell				0			U				U	0									0	U	U						
Mitchell fMRI 18	Subtotals			5	0	0	4	0	0	7	1	1	0	0	0	0	5	6	0	1	1	1	1	0	0	0	0	7	2
Mitchell fMRI 18 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	OTHER (n = 8)																												
Ochsnera fMRI 13 0 0 0 0 0 0 0 0 0		fMRI	18				0	0									0											0	
Jackson FMRI 15				0				·		0								0											
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Lawrence fMRI 12 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0																					0								
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Subtotals 4 0 0 4 1 0 4 1 2 0 0 0 0 2 1 0 2 3 1 0 0 0 1 0 7 Specific Task Stimulus Type Emotion Pain Soc Discrete Uis Vis Aud Olf/Tas Dimension Pain Soc Discrete Iden Label Mat Int Fa W Pic Film Sent W Sent Snd Pos Neg Ar H S D F Ang Sur Ex vs I E	Hynes	fMRI	20	0								0									0							0	
Specific Task Stimulus Type Emotion Olf/Tas Dimension Pain Soc Discrete H S D F Ang Sur Ex vs I E	Ruby ^a	PET	10	0								0									0							0	
Vis Aud Olf/Tas Dimension Pain Soc Discrete Iden Label Mat Int Fa W Pic Film Sent W Sent Snd Pos Neg Ar H S D F Ang Sur Ex vs I E	Subtotals			4	0	0	4	1	0	4	1	2	0	0	0	0	2	1	0	2	3	1	0	0	0	1	0	7	1
Iden Label Mat Int Fa W Pic Film Sent W Sent Snd Pos Neg Ar H S D F Ang Sur Ex vs I E				Speci	ific Task	(Stimulus Type							Emotion		tion				Contras	t							
								Vis	5				Au	d		Olf/Tas	Dime	ension		Pain	Soc	Dis	crete	;					
				Iden	Label	Mat	Int	Fa	W	Pic	Film	Sent	W	Sent	Snd		Pos	Neg	Ar			Н	S	D	F	Ang	Sur	Ex vs I	E vs N
Totals 21 1 1 15 8 3 16 2 3 1 2 1 3 15 18 1 3 4 8 4 5 5 6 3 30 1	Totals			21	1	1	15	8	3	16	2	3	1	2	1	3	15	18	1	3	4	8	4	5	5	6	3	30	10

Imaging Tech = Imaging Technique, Specific Task: Iden. = Identification Task, Label = Labelling Task, Mat = Matching Task, Int = Intensity Task, Stimulus: Vis = Visual, Aud = Auditory, Olf/Tas = Olfactory/Taste, Fa = Face, W = Word, Pic = Picture, Sent = Sentence, Snd = Sound, Emotion: Pos = Positive, Neg = Negative, Ar = Arousal, Soc = Social, H = Happy, S = Sad, D = Disgust, F = Fear, Ang = Anger, Sur = Surprise, Contrast: Ex = Explicit Condition, I = Implicit/other control (e.g., rest) Condition, E = Emotional Condition, N = Neutral Condition. (i) Lists of studies in STIM (Imaizumi et al., 1997; Nakamura et al., 1999; Paradiso et al., 1999; Critchley et al., 2000; Hariri et al., 2000; Liberzon et al., 2000; Narumoto et al., 2000; Royet et al., 2000; Gorno-Tempini et al., 2001; Royet et al., 2001; Tabert et al., 2001; Lange et al., 2003; Maddock et al., 2003; Royet et al., 2003; Taylor et al., 2003; Winston et al., 2003; Cunningham et al., 2004; Wildgruber et al., 2004; Wildgruber et al., 2005; Grimm et al., 2006); (ii) Lists of studies in SELF (Gusnard et al., 2001; Lee et al., 2004; Ochsner et al., 2004; Phan et al., 2004; Ruby and Decety, 2004; Garrett and Maddock, 2006; Jackson et al., 2006; Lawrence et al., 2006; Jackson et al., 2006; Vollm et al., 2006; Clawrence et al., 2006; Jackson et al., 2006; Vollm et al., 2006; Clawrence et al., 2006; Jackson et al., 2006; Vollm et al., 2006)

^aThe studies that examine two evaluation tasks and directly compare their effects.

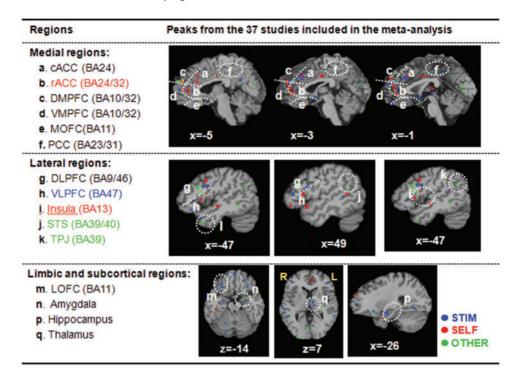


Fig. 1 The boundaries of a priori ROIs in this study and activation peaks colour-coded based on three different groups of emotional evaluation.

Montreal Neurological Institute (MNI) brain were transformed to Talairach coordinate system (http://imaging.mrc-cbu.cam.ac.uk/imaging/MniTalairach) and were double checked using the AFNI program (Analysis of Functional NeuroImages; Cox, 1996).

A meta-analysis was performed using the Activation Likelihood Estimate (ALE) method (Turkeltaub *et al.*, 2002), implemented in BrainMap (Laird *et al.*, 2005) using the coordinates in regions listed in Table 1. The ALE values for each voxel were computed using a 3D Gaussian probability density function with FWHM = 10 mm, as recommended by Laird *et al.* The voxel-wise significance of the ALE values was determined by a permutation test using 5000 permutations (Laird *et al.*, 2005). The images of P values for each voxel were corrected for multiple comparisons using false discovery rates (FDR), at P < 0.05 corrected. The final step was to identify clusters of voxels exceeding 100 mm³ in volume, as commonly used in the GingerALE software (http://brainmap.org).

Separate meta-analyses were conducted with the STIM, SELF and OTHER groups. Overlapping brain maps were drawn via conjunction analysis on the three maps derived from the separate meta-analyses. To identify distinct brain regions associated with individual evaluation tasks, meta-analytic contrast maps were created using a subtraction meta-analysis (Laird *et al.*, 2005). The contrast maps showed regions in which the two groups of foci are significantly different. Conjunction maps between two contrast maps (e.g. SELF *vs* STIM and SELF *vs* OTHER) were also created to identify unique distinct regions of each specific group

(e.g. SELF) that are commonly different from the two other groups (STIM and OTHER).

RESULTS

Common networks associated with explicit evaluation of emotion

To elucidate common brain regions associated with explicit emotional evaluation, we found brain regions associated with each emotional evaluation task separately and then identified overlapping area commonly activated by three different types of explicit emotional evaluation.

The results from the separate meta-analysis of the 141 coordinates identified in 20 studies (32 contrasts) of 'STIM' showed activation in several prefrontal regions (VLPFC [BA47], DMPFC [BA32], DLPFC [BA9] and IFG [BA46]), visual cortices (fusiform and ligual gyrus, BA19) and subcortical-limbic regions (PCC [BA31], thalamus, amygdala and amygdala extending to the parahippocampal gyrus; Table 3). The separate map was created using 61 coordinates in nine studies (14 contrasts) of SELF including rACC (BA32), DMPFC (BA10), insula (BA13), LPFC (BA45/47), cACC and amygdala (Table 4). The map from the separate meta-analysis of 59 coordinates in eight studies (12 contrasts) of OTHER group showed activation in the STS (BA39), TPJ (BA39), DMPFC (BA10), precuneus/PCC (BA31), LPFC (BA44/47), VLPFC (BA47) and amygdala (Table 5).

In short, amygdala, LPFC and DMPFC seemed to activate commonly in all three conditions of explicit emotional

Table 3 Reported foci resulting from meta-analysis in *stimulus-focused evaluation* (voxel-wise P < 0.05; FDR corrected)

Region	Side	ВА	Х	у	Z	Volume (mm ³)	Maximum ALE Value
Cinqulate gyrus	R	32	2	28	30	6128	0.0170
Inferior frontal gyrus	L	47	-28	26	—10	5072	0.0090
Inferior frontal gyrus	R	47	44	28	-10	4480	0.0111
Amygdala	L		-24	-6	-14	2672	0.0150
Thalamus (ventral lateral nucleus)	L		-10	-10	6	2264	0.0092
Amygdala extending to parahippocampal gyrus	R		24	-6	-12	2224	0.0124
Inferior frontal gyrus	L	9	-48	10	28	1720	0.0115
Posterior cingulate	L	31	-6	-54	24	1696	0.0089
Middle frontal gyrus	R	9	50	20	28	840	0.0065
Superior temporal gyrus	L	22	-62	-44	14	816	0.0071
Lingual gyrus	L	19	-16	-44	-2	624	0.0067
Inferior occipital gyrus	R	19	44	-78	-4	592	0.0076
Lingual gyrus	R	19	18	-58	-2	528	0.0064
Middle frontal gyrus	R	11	24	50	-10	440	0.0057
Medial frontal gyrus	L	32	-6	10	44	424	0.0073
Posterior cingulate	R	30	4	60	8	328	0.0059
Lingual gyrus	L	19	-22	-62	-2	256	0.0060
Fusiform gyrus	L	19	-46	-66	-14	200	0.0056
Inferior frontal gyrus	R	46	50	30	12	160	0.0051
Anterior cingulate	L	24	-6	24	-6	144	0.0054
Medial frontal gyrus	R	11	4	58	-14	120	0.0047

Table 4 Reported foci resulting from meta-analysis in *evaluation of one's own emotion* (voxelwise P < 0.05; FDR corrected)

Region	Side	ВА	Х	у	Z	Volume (mm ³)	Maximum ALE
Medial frontal gyrus	L	10	-2	50	12	8312	0.0104
Insula	L	13	-46	11	2	4560	0.0080
Inferior frontal gyrus	R	45	48	20	10	4120	0.0057
Amygdala	L		-20	-6	—12	1184	0.0069
Cingulate gyrus	R	32	8	14	40	1056	0.0071
Thalamus (medial dorsal nucleus)	R		4	-18	12	888	0.0061
Cingulate gyrus	L	24	0	-2	34	520	0.0045
Middle frontal gyrus	R	47	48	36	-8	480	0.0050
Cingulate gyrus	L	32	-4	26	26	448	0.0043
Thalamus (pulvinar)	R		4	-28	2	112	0.0040
Medial frontal gyrus	R	8	4	48	42	104	0.0039

Table 5 Reported foci resulting from meta-analysis in *evaluation of others' emotions* (voxelwise P < 0.05; FDR corrected)

Region	Side	ВА	Х	у	Z	Volume (mm ³)	Maximum ALE
Precuneus	L	31	-10	-56	34	3312	0.0080
Superior frontal gyrus	L	9	-8	54	28	3136	0.0081
Medial frontal gyrus	R	10	2	56	10	2080	0.0066
Superior temporal gyrus	R	39	48	-52	24	1592	0.0078
Medial frontal gyrus	R	11	4	50	-18	1560	0.0052
Fusiform gyrus	L	20	-58	-4	-26	1496	0.0083
Superior temporal gyrus	L	39	-50	60	18	1480	0.0069
Amygdala extending to parahippocampal gyrus	L		-22	-2	-18	592	0.0051
Inferior frontal gyrus	L	47	-54	28	-4	552	0.0062
Cuneus	L	18	0	-78	14	504	0.0043
Middle frontal gyrus	L	46	-46	18	24	120	0.0040
Inferior frontal gyrus	L	44	-46	12	16	112	0.0042

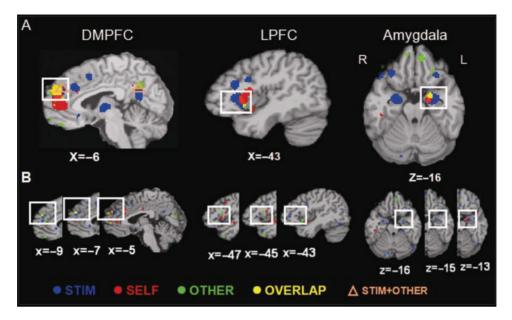


Fig. 2 Overlapping areas representing common brain networks underlying explicit evaluation of emotion. (**A**) The overlapping areas included the DMPFC, LPFC and amygdala. Different colours represent brain regions associated with different evaluation tasks (Blue: brain regions activated by *stimulus-focused evaluation* (STIM), Red: brain regions associated with *evaluation of one's own emotion* (SELF), Green: brain areas associated with *evaluation of others' emotions* (OTHER), Yellow: Overlapping areas commonly associated with three tasks (OVERLAP) and Triangle: Overlapping areas between STIM and OTHER. (**B**) All reported peaks near the overlapping areas. (Abbreviations: DMPFC = Dorsomedial Prefrontal Cortex, LPFC = Lateral Prefrontal Cortex).

evaluation although slightly different sizes and locations of activation were identified within these regions. Overlapping brain regions were created using three maps generated by separate meta-analyses. This confirmed that some parts of these brain regions overlapped across different evaluation conditions, providing evidence that some activation peaks in each group were located near the overlapping area. Thus, these overlapping areas represent higher probability of common involvement in different types of emotional evaluation. However, activation peaks in broad regions (e.g. LPFC) were more distributed and spatially distinguishable compared to peaks in small regions (e.g. amygdala), so activation peaks in broad regions may be separable. To understand whether similar numbers of peaks from each group were located in the regions (suggesting common functionality), Figure 2 displays overlapping regions (A) and individual peaks (B) in the DMPFC, left LPFC and left amygdala. Because observed peak distributions largely overlapped across tasks, the amygdala, LPFC and DMPFC were considered common brain networks involved in explicit emotional evaluation irrespective of the different types of explicit evaluation tasks.

Distinct networks depending on different evaluation task

To examine which brain regions were specifically involved in individual evaluation tasks, three contrast meta-analytic maps (STIM *vs* SELF, STIM *vs* OTHER and SELF *vs* OTHER) were created using the same procedures, yielding brain regions which showed significant differences between

two comparison conditions. Conjunction maps between two contrast maps represented distinct regions commonly more activated by one specific group than two other groups. Table 6 summarizes identified foci in each conjunction map showing significant differences between one group and two other groups.

In the conjunction map of STIM vs SELF and STIM vs OTHER, studies of stimulus-focused evaluation showed more activation in prefrontal regions (e.g. DMPFC [BA32], VLPFC [BA47], DLPFC [BA9]), sensory regions (e.g. visual and temporal cortices) and subcortical regions (e.g. thalamus and parahippocampus gyrus) compared to both studies of evaluation of one's own emotion and of others' emotions, suggesting that these regions are differentially involved in evaluation of emotional stimuli (Table 6). Figure 3 shows activation clusters in the right VLPFC (BA47) and visual cortex hypothesized regions specifically associated with evaluation of stimuli. Consistent with our hypothesis, sensory cortices and VLPFC (BA47) were revealed as distinct brain regions specifically associated with evaluation of emotional stimuli.

The conjunction map of SELF *vs* STIM and SELF *vs* OTHER demonstrated that brain regions such as the rACC (BA24/32) and insula (BA13) near the IFG are more likely associated with *evaluation of one's own emotion* compared to other groups (Table 6). Identified clusters in the rACC and left insula extending to the IFG are displayed in Figure 3. Consistent with our hypotheses, the rACC and insula were specifically involved in evaluation of one's own emotional states.

Regions	Side	BA	X	у	Z	Volume (mm ³)	Maximum ALE
Conjunction map of stimulus-focused: 'STIM' greater	r than 'SELF' and	d 'STIM' greater t	han 'OTHER'				
Cingulate gyrus	R	32	2	30	30	2048	0.014
Inferior frontal gyrus	R	47	46	26	-12	2000	0.009
Amygdala	R		24	-6	-12	1744	0.012
Amygdala	L		-26	-6	-14	656	0.009
Ventral lateral nucleus	L		-10	-10	6	1160	0.009
Posterior cingulate gyrus	L	31	-6	-54	22	1072	0.008
Inferior frontal gyrus	L	9	-48	10	28	816	0.010
Inferior frontal gyrus	L	47	-28	24	-10	664	0.008
Superior temporal gyrus	L	22	-62	-44	14	656	0.007
Parahippocampal gyrus	L	19	-16	-44	-4	368	0.006
Lingual gyrus	R	19	18	-58	-2	312	0.006
Inferior frontal gyrus	R	9	52	8	24	224	0.006
Middle frontal gyrus	R	11	24	50	-10	192	0.006
Medial frontal gyrus	L	32	-6	10	46	192	0.007
Anterior cingulate gyrus	L	32	-18	40	-10	176	0.006
Posterior cingulate gyrus	R	30	4	-60	8	168	0.006
Inferior frontal gyrus	L	13	-42	24	6	168	0.006
Lingual gyrus	L	19	-22	-62	-2	136	0.006
Inferior occipital gyrus	R	19	44	-78	-4	112	0.006
Conjunction map of one's own emotion: 'SELF' great	ter than 'STIM'	and 'SELF' greate	r than 'OTHER'				
Anterior gingulate gyrus	L	24/32	-2	46	8	296	0.007
Cingulate gyrus	R	32	8	14	40	216	0.007
Insula extending to the Inferior Frontal gyrus	L	13/47	-50	14	0	208	0.006
Conjunction map of others' emotions: 'OTHER' great	ter than 'STIM' d	ınd 'OTHER' great	er than 'SELF'				
Superior temporal gyrus	R	39	48	-52	24	488	0.008
Temporo-parietal junction	L	39	-50	-60	18	256	0.007

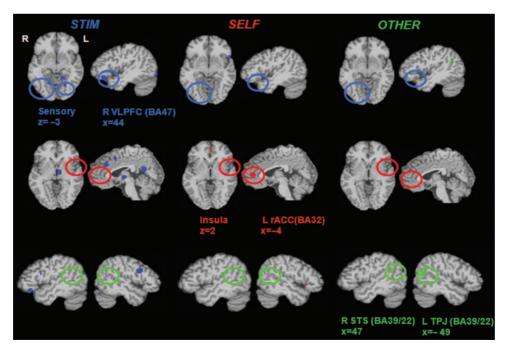


Fig. 3 Contrast meta-maps derived from the comparisons among three different tasks to assess brain mechanisms associated with explicit evaluation of emotion (first column: STIM, second column: SELF and third column: OTHER). Explicit evaluation of emotion engages distinct brain networks depending on different evaluation tasks. First row shows the results in the sensory (visual) cortex and VLPFC. Activation in these regions was identified in only *stimulus-focused evaluation* condition. Second row displays the results in the insula and rACC. Activation in both regions was revealed in only *evaluation of one's own emotion* condition. Third row presents the findings in the STS and TPJ. Both regions were activated by only *evaluation of others' emotions* condition. These results suggest that some brain regions are involved in specific types of emotional evaluation. (Abbreviations: L = left, R = right, VLPFC = ventrolateral prefrontal cortex, rACC = rostral anterior cingulate cortex, STS = superior temporal sulcus, TPJ = temporo-parietal junction).

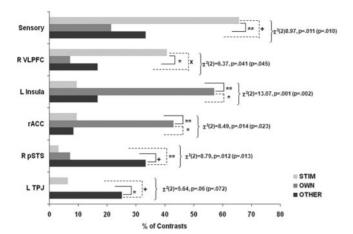


Fig. 4 Comparison of activation frequencies across evaluation task groups (STIM, SELF and OTHER). χ^2 results in each region were displayed (Fisher's exact tests were reported in the parenthesis). Follow-up 2 \times 2 comparisons were reported (x = non significant, + P < 0.10, *P < 0.05 and **P < 0.01).

Brain regions including the STS and TPJ were hypothesized to specifically be associated with evaluation of other people's emotions. The STS (BA39) and TPJ (BA39) were commonly activated by studies of evaluation of others' emotions compared to studies of two other tasks (Table 6). Figure 3 presents activation clusters in the right STS and left TPJ. This result also supports the hypothesis that the STS and TPJ are involved in evaluation of others' emotions via inferring emotional states in others.

Volume-wise differences for regions of interest. The ALE method provides relative peak concentration in a given region for each group which allows for the possibility that differences between tasks are apparent when two tasks activate different parts of the same ROI. To examine whether activation frequencies within the entire volume of each region of interest were significantly different across the three groups of studies, χ^2 and Fisher's exact tests were conducted. Data were coded as 1 or 0 if the distinct regions were activated or not in each group, respectively. Figure 4 displays percentages of activation peaks in each group and χ^2 results. Activation frequencies in the sensory, VLPFC, insula, rACC and STS were significantly different among three task groups, but activation frequencies in the TPJ were marginally different among groups. These χ^2 results were consistent with ALE results. Follow-up 2×2 comparisons were conducted. Except for the comparison between STIM vs OTHER in the VLPFC, most 2×2 comparisons showed significant or marginally significant differences in activation frequencies.

Potential confounds. Finally, potential confounds were examined using a logistic regression analysis. One group of emotional evaluation was more diverse than the other groups in stimulus type (e.g. face, word, film and picture) and emotions (e.g. valence, discrete emotions, social emotions and pain). A logistic regression was performed to

examine whether these factors would predict activation in the distinct regions. Each study was coded based on whether the evaluation task was performed according to valence, discrete emotions, social emotions and pain and further based on whether the task was performed with presentation of specific stimulus types such as face, word, film and picture. The emotion factor did not predict activation in the VLPFC (Wald $\chi^2 = 0.05$, P = 0.975), sensory (Wald $\chi^2 = 0.51$, P = 0.774), insula (Wald $\chi^2 = 4.17$, P = 0.124), rACC (Wald $\chi^2 = 1.38$, P = 0.50) except for the STS (Wald $\chi^2 = 7.17$, P < 0.05) and TPJ (Wald $\chi^2 = 6.08$, P < 0.05). Emotion predicted activation in the STS and TPJ because studies which showed activation in the STS and TPJ used particularly pain and social emotions. However, only three studies reported activation in the STS and TPJ, so this factor did not have sufficient variance to account for activation. Stimulus type did not significantly predict activation in any distinct regions (VLPFC: Wald $\chi^2 = 2.99$, P = 0.392; sensory: Wald $\chi^2 = 7.24$, P = 0.065; insula: Wald $\chi^2 = 2.60$, P = 0.785; rACC: Wald $\chi^2 = 1.07$, P = 0.785; STS: Wald $\chi^2 = 4.51$, P = 0.211; TPJ: Wald $\chi^2 = 0.987$, P = 0.804). Thus, these potential factors did not confound our results. However, one should interpret such findings with caution because this method to control for confound variables is useful when many studies reported activation in brain regions (Wager et al., 2007).

DISCUSSION

A meta-analysis was conducted to elucidate common and distinct brain networks associated with general and task-specific mechanisms of explicit emotional evaluation. Consistent with our hypotheses, results suggest that the amygdala, LPFC (BA47) and DMPFC (BA10/32) are involved in general mechanisms underlying the explicit evaluation of emotion, whereas some brain regions are involved in task-specific mechanisms associated with specific evaluation tasks. Task instructions for cognitive evaluation of emotional stimuli might recruit sensory and VLPFC (BA47) regions. Conscious evaluation of one's own emotion might activate the insula (BA13) and rACC (BA24), whereas evaluation of other people's emotions might be associated with activation in the STS (BA39) and TPJ (BA39).

General brain mechanisms of explicit emotional evaluation

We hypothesized that cognition—emotion interactions involving evaluative and regulatory processing, subserved by prefrontal and limbic regions, would be shared mechanisms associated with many types of explicit evaluation of emotion. In support of this idea, three hypothesized regions, the amygdala, LPFC and DMPFC were observed as common to all three emotional evaluation tasks.

The amygdala is involved in a broad range of emotional processing including perceptual processing (e.g. recognition of facial expressions; Adolphs, 2002a), encoding emotional

arousal (e.g. Anderson *et al.*, 2003) and evaluation of emotional valence (Paradiso *et al.*, 1999; Liberzon *et al.*, 2000). The amygdala appears to participate in fast and automatic processing of emotion (LeDoux, 1996; Adolphs, 2002b). The amygdala may thus be involved in the initial stages in the process of explicit emotional evaluation such as encoding of emotional information and initial emotional reactivity to emotional information.

The LPFC is associated with memory retrieval (e.g. Wagner *et al.*, 2001; Bunge *et al.*, 2004) and cognitive control including selective attention (Bishop *et al.*, 2004; Cardillo *et al.*, 2004; Bedwell *et al.*, 2005). Similarly, the LPFC may guide selective attention to the given task of explicit emotional evaluation and may maintain the task instruction information to accomplish the desired consequences of conscious evaluation of emotion.

The DMPFC (BA10/32) is connected with other brain regions associated with emotional and cognitive processing (e.g. amygdala, LPFC and cACC), possibly reflecting a role in emotion—cognition integration (Paradiso *et al.*, 1999; Taylor *et al.*, 2003; Lieberman *et al.*, 2007). Explicit emotional evaluation also includes evaluative processing depending on individuals' internal value systems, associated with DMPFC activity (Cunningham *et al.*, 2004). The DMPFC has also been implicated in emotion regulation that may occur after conscious evaluation of emotion (Ochsner *et al.*, 2004b; Banks *et al.*, 2007).

To summarize, the amygdala, LPFC and DMPFC (BA10/32) appear commonly associated with explicit emotional evaluation. Divergent experimental designs across studies may prevent identification of other overlapping regions. Potentially, the amygdala is involved in initial emotional processing of explicit evaluation, whereas the LPFC more specifically subserves cognitive aspects of evaluation. These regions could interact via the DMPFC, a convergent region involved in integrative and evaluative processing.

Task-dependent brain mechanisms of explicit emotional evaluation

We examined underlying brain mechanisms distinctively associated with three explicit emotional tasks that required *stimulus-focused evaluation*, and focusing on *one's own emotion* and *other people's emotions*.

Sensory cortex and VLPFC (BA47) were associated with evaluation of stimuli/situations in accordance with our hypothesis. Presumably, such *stimulus-focused evaluation* requires perceptual processing of physical features, yielding recruitment of sensory cortex (visual and temporal cortices) in the evaluation of emotional stimuli (Paradiso *et al.*, 1999; Critchley *et al.*, 2000; Gorno-Tempini *et al.*, 2001; Taylor *et al.*, 2003). The VLPFC (BA47) is associated with evaluative judgment (Cunningham *et al.*, 2003), appraisal of object knowledge (Mitchell *et al.*, 2002) and declarative knowledge about emotional stimuli (Schaefer *et al.*, 2003), possibly

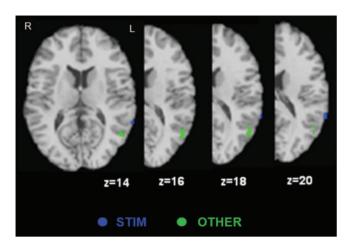
responsible for higher-level of conceptual processing of emotional stimuli.

Also consistent with our hypothesis, the insula (BA13) and rACC were specifically associated with *evaluation of one's own emotion*. The insula has been associated with perception of inner state changes such as interoceptive processing, mediating conscious emotional experience (Craig, 2002; Critchley *et al.*, 2004). The rACC (BA24/32) is also involved in subjective emotional experience (e.g. Lane, 2000).

Finally, consistent with our hypotheses, the STS and TPJ were involved in evaluation of other people's emotions. Both regions are associated with social affective judgment including simulation and empathetic processing as well as social cognitive judgment including understanding of and reasoning others' intention and thoughts (Frith and Frith, 1999; Ochsner *et al.*, 2004a; Ruby and Decety, 2004; Vollm *et al.*, 2006)

Although we found distinct regions associated with three different tasks, it is possible that there are common mechanisms associated with both stimulus-focused evaluation and evaluation of others' emotions using faces. Potentially, evaluating the emotionality (e.g. angry or sad) for face stimuli could make people evaluate emotional states of the subjects whose faces are being viewed. But it is also possible that this task is more dependent on perception and can be accomplished without inferring others' emotional states. In contrast, explicitly demanding people evaluate emotional states of others by presenting only faces is likely to recruit mechanisms associated with TOM. Consistently, past research on empathy has attempted to distinguish evaluating emotions based on more perceptual cues (e.g. stimulus) from evaluating emotions based on inferences about others' emotions (Eisenberg et al., 1997; Graham and Ickes, 1997). Our distinct regions also supported the idea that stimulus-focused evaluation is specifically associated with the visual cortex; however, evaluation of others' emotions is specifically associated with the STS and TPJ, although the precuneus/PCC and DMPFC are overlapped between two tasks (Figure 2).

The ALE and χ^2 results broadly agreed, but two-way χ^2 and ALE were slightly different. It is possible that ALE and χ^2 results are different because the ALE method provides relative peak concentration in a given region for each group whereas χ^2 results provide absolute differences in activation frequencies across groups (Wager *et al.*, 2007). Thus, a region-wide analysis using χ^2 does not detect significant differences in activation frequencies among groups, whereas a finer spatial resolution analysis using ALE does detect different activation clusters if groups differ in the subregions of a single structure they activate. For example, χ^2 did not show significant differences in the TPJ among groups, but ALE detected TPJ as a distinct region associated with evaluation of others' emotions. As shown in Figure 5, peaks of OTHER group were placed on different



Common and distinct brain networks underlying emotional evaluation

Fig. 5 Peaks activated by STIM and OTHER in the TPJ. Peaks in two groups were distinguishable in the TPJ.

subregions of the TPJ compared to peaks of STIM group. Therefore, activation frequencies in this region might not be significantly different among the groups whereas the cluster of subregions might be detected as a distinct region associated with evaluation of others' emotions.

In summary, different tasks to assess underlying processing of explicit emotional evaluation recruited distinct brain networks suggested by theoretical accounts. Distinct brain networks may be specialized for specific emotional evaluation, so these networks are necessary to accomplish specific evaluation tasks.

Multiple processes in explicit evaluation of emotion

Findings from our meta-analysis supported the notion that explicit emotional evaluation is associated with shared mechanisms common to all types of emotional evaluation such as PFC-subcortical interactions, as well as different mechanisms specifically involved in different types of emotional evaluation. PFC-subcortical interactions are commonly acknowledged in imaging research on explicit emotional evaluation. Adding distinct brain mechanisms associated with specific tasks, in particular self vs other evaluation, sheds additional light on understanding the process of explicit emotional evaluation.

There are several reasons why the distinct mechanisms associated with evaluation of three objects should be considered in explicit emotional evaluation. First, as mentioned earlier, developmental psychologists suggest that engagements with people, other physical objects (e.g. stimuli and situations) and self are critical in normal emotional development (Trevarthen and Aitken, 2001). Abnormal development of one distinct mechanism can cause specific affective developmental disorders such as relationships between other-specific mechanisms and autistic spectrum disorder (ASD). Second, different types of explicit emotional evaluation can be used to detect affective disorders such as ASD, alexithymia and psychopathy (for a review, see Blair, 2008a, b; Decety and Moriguchi, 2007). Individuals with ASD exhibit problems with distinct mechanisms associated with evaluating others' emotions whereas individuals with alexithymia exhibit abnormality in distinct mechanisms associated with evaluating one's own emotional states. Thus, distinct mechanisms associated with different types of evaluation contribute to the understanding of mechanisms in affective disorder.

Third, distinct mechanisms of different evaluation based on objects to be evaluated may be differentially associated with other emotional processes such as emotional regulation (Ochsner et al., 2004b). Consistent with our findings, Ochsner and colleague reported distinct functions in the VLPFC and VMPFC (rACC), associated with situationfocused and self-focused regulation, respectively. Specific distinct mechanisms may be associated with particular types of emotion regulation such as reappraisal and suppression. For example, reappraisal (e.g. reinterpreting emotional situations) may be more likely associated with stimulusfocused evaluation whereas suppression (e.g. inhibiting one's own emotional behaviour) may be more likely associated with self. Thus, distinct mechanisms of emotional evaluation contribute to the understanding of other emotional processing.

Possible interactive model

Both common and distinct networks associated with explicit evaluation of emotion were observed. Based on theoretical framework and these results, we suggest the interactive model in Figure 6, which depicts salient interactions between regions subserving common and more distinct functions for different types of explicit emotional evaluation. Of course, the model could include many more connections and reciprocal interactions; rather, our attempt here is to illustrate some of the most conceptually salient connections which tell a story suggested by the meta-analytic data and well-known functions in the regions.

In our model and in all types of explicit emotional evaluation, the amygdala is hypothesized to be involved in encoding emotional information and initial responses to the information, and the LPFC is hypothesized to manage selective attention to guide a desired task performance. The DMPFC possibly plays a role in bridging between the common input regions (amygdala and LPFC; Lieberman et al., 2007). Integrative information in the DMPFC could further be fed back to the amygdala and LPFC subserving cognition-emotion interactions. Task-dependent mechanisms would be required to accomplish additional processing demanded by specific task instructions. For example, in the case of stimulus-focused evaluation, the sensory cortices and VLPFC would be recruited for perceptual and conceptual processing of stimuli. We assume that these task-dependent mechanisms would operate for specific processes related to particular task demand, possibly after early common mechanism but before late common mechanism.

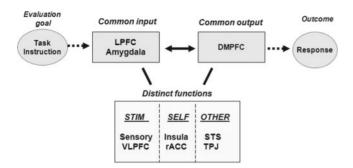


Fig. 6 Hypothetical interactive model of explicit emotional evaluation. Dotted lines represent hypothesized precedence based on the task design (i.e. task instructions are presented before stimuli and responses are generated after some amount of processing). The bi-directional arrow between common input and output regions reflects the extensive literature documenting functional and anatomical connectivity between these regions. Connections to regions associated with distinct functions are less well specified in the literature, and thus future research is necessary to establish directionality of relationships.

Future research could test this hypothesized interactive model using different types of emotional evaluation tasks within the same fMRI design, controlling for experimental factors such as stimulus characteristics. Our model's dynamic interactions and directionality among hypothesized common and distinct regions could specifically be examined via analyses of functional and effective connectivity. Reciprocal connectivity between the amygdala, LPFC and DMPFC is predicted to show a good fit to the fMRI data independent of particular evaluation tasks. Modulation by specific evaluation tasks would be supported if (i) a specific task is associated with increased relationships between early common regions and specific task-dependent regions; and (ii) enhanced activity in the specific networks alters connectivity among the common networks, ideally by feed-forward signals to the DMPFC, which consequently interacting with amygdala and LPFC.

In general, the reciprocal and causal connectivity suggests that underlying mechanisms of explicit emotional evaluation can be better understood within more distributed neural networks. This concept is consistent with recent suggestions that broader and more distributed networks would be more appropriate for elucidating emotion—cognition interactive mechanisms (Taylor and Liberzon, 2007; Pessoa, 2008). This model also demonstrates one example of how subtle differences in task instructions could contribute to alteration of neural circuits associated with psychological mechanisms. Therefore, careful selection of task instructions is suggested.

Limitations and concluding remarks

There are several limitations of this review. We included coordinates in only hypothesized brain regions for this meta-analysis, so we did not examine how other brain regions are associated with different types of conscious emotional evaluation. There is also a difference in the

number of studies in three assigned groups. The group of stimulus-focused evaluation included more studies (20 studies) than two other groups (nine and eight studies in evaluation of one's own emotion and others' emotions, respectively). This may cause additional brain regions including the DLPFC and thalamus associated with stimulus-focused evaluation. The ALE technique does not account for the size and shape of activity in each study. Thus, differential roles among adjacent regions such as the insula and LPFC may not adequately be separated by this technique. In addition, nearly many of the examined tasks used stimuli for which the nominal interpretation of the stimulus would be consistent among any of the three tasks which were required. For example, a picture of a crying person would be evaluated as negative and might make the observer feel sad in addition to provoking an empathetic response. Thus, regions labelled as common may reflect processes that occurred regardless of the nominal task as a function of the stimulus, but are not actually common to these tasks. Experiments using stimuli that are evaluated differently from different perspectives (e.g. stimuli in which the observer would take pleasure in the subject's misfortune) could be helpful in this regard.

Despite these limitations, the proposed framework integrates evidence across a variety of theoretical accounts and neuroimaging studies of explicit emotional evaluation. We suggest that explicit evaluation of emotion is not a unitary process but instead multiple processes mediated by shared/common and distinct mechanisms. The review has important implications for the investigation of understanding mechanisms underlying explicit evaluation of emotion. Given the different brain responses associated with different tasks, instructions for emotional evaluation tasks should be specific in terms of particular research interests in specific mechanisms underlying explicit evaluation. This framework may also have clinical implications for understanding mechanisms of affective disorders such as alexithymia and autism. These disorders are often considered to reflect general deficits in emotional evaluation. Potentially, by carefully assessing evaluation of different domains (e.g. self vs other) pockets of preserved competence could be identified.

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