

# The Extended Language Network: A Meta-Analysis of Neuroimaging Studies on Text Comprehension

Evelyn C. Ferstl,<sup>1,2\*</sup> Jane Neumann,<sup>1</sup> Carsten Bogler,<sup>1</sup>  
and D. Yves von Cramon<sup>1,2</sup>

<sup>1</sup>Max-Planck-Institute of Cognitive and Brain Sciences, Leipzig, Germany

<sup>2</sup>Day Clinic of Cognitive Neurology, University of Leipzig, Leipzig, Germany

---

**Abstract:** Language processing in context requires more than merely comprehending words and sentences. Important subprocesses are inferences for bridging successive utterances, the use of background knowledge and discourse context, and pragmatic interpretations. The functional neuroanatomy of these text comprehension processes has only recently been investigated. Although there is evidence for right-hemisphere contributions, reviews have implicated the left lateral prefrontal cortex, left temporal regions beyond Wernicke's area, and the left dorso-medial prefrontal cortex (dmPFC) for text comprehension. To objectively confirm this extended language network and to evaluate the respective contribution of right hemisphere regions, meta-analyses of 23 neuroimaging studies are reported here. The analyses used replicator dynamics based on activation likelihood estimates. Independent of the baseline, the anterior temporal lobes (aTL) were active bilaterally. In addition, processing of coherent compared with incoherent text engaged the dmPFC and the posterior cingulate cortex. Right hemisphere activations were seen most notably in the analysis of contrasts testing specific subprocesses, such as metaphor comprehension. These results suggest task dependent contributions for the lateral PFC and the right hemisphere. Most importantly, they confirm the role of the aTL and the fronto-medial cortex for language processing in context. *Hum Brain Mapp* 29:581–593, 2008. © 2007 Wiley-Liss, Inc.

**Key words:** text comprehension; theory-of-mind; pragmatics; language interpretation; anterior temporal lobe; fronto-medial cortex; right hemisphere; metaanalysis; neuroimaging

---

## INTRODUCTION

Language comprehension requires more than just understanding words and sentences. Over and above these linguistic processes, realized in left-sided perisylvian brain regions, a variety of additional cognitive processes are recruited. For creating a coherent representation of a story or a dialogue, it is necessary to bring in general world knowledge, to integrate the current utterance with the

prior context, or to check the consistency of the resulting interpretation with the communicative situation. Because of these requirements, it is not surprising that neuroimaging studies of text comprehension have unveiled an extended language network<sup>1</sup> (ELN) of brain regions to be involved during inferencing and interpretation.

Two single studies illustrating this network are the seminal PET study by Mazoyer et al. [1993], as well as a recent experiment using fMRI [Xu et al., 2005]. The activation patterns include the anterior temporal lobes (aTL) bilaterally,

---

\*Correspondence to: Evelyn C. Ferstl, Department of Psychology, University of Sussex, Pevensey 1, Falmer, Brighton BN1 9QH, United Kingdom. E-mail: e.c.ferstl@sussex.ac.uk

Received for publication 16 January 2006; Revised 13 April 2007; Accepted 16 April 2007

DOI: 10.1002/hbm.20422

Published online 7 June 2007 in Wiley InterScience (www.interscience.wiley.com).

---

<sup>1</sup>For lack of a better term, and consistent with conventional usage, the term "network" is used throughout the manuscript to refer to a set of cortical regions contributing to the same functional tasks. We do not claim that the meta-analysis method provides insight into whether these regions are anatomically or even functionally connected.

extending into the temporal poles, as well as activation all along the superior temporal sulcus (STS) on the left side. Many studies have found inferior frontal gyrus (IFG) activation. Importantly, fronto- and parieto-medial activations are reported when coherent language is read or heard. Based on reviews of imaging studies, Ferstl [in press] and Mar [2004] have shown that these results are rather stable across studies.

There are several open questions, however. The first question concerns the overlap of the ELN with the network identified to be crucial for Theory-of-Mind processes [ToM; Frith and Frith, 2003; Gallagher and Frith, 2003]. ToM or mentalizing refers to the ability to attribute other people's motivations, emotions, actions, and thoughts to their mental states and to their beliefs about the world. During communication, ToM processes help to understand the speaker's or writer's intentions, and some pragmatic theories in linguistics equate ToM and language processing in context [cf. Frith and Frith, 2003; Sperber and Wilson, 1995]. The brain regions consistently reported in ToM studies are the aTL, the temporo-parietal junction (TPJ), and most importantly, the dorso-medial prefrontal cortex (dmPFC). We have found this latter region to be involved during inference processes in text comprehension [Ferstl and von Cramon, 2001, 2002; Ferstl et al., 2005]. When inspecting the available neuroimaging literature on text comprehension, it can be seen that several studies were actually designed to test ToM processes rather than language comprehension [e.g., Fletcher et al., 1995; Vogeley et al., 2001]. Thus, it is an open question whether the apparent overlap between the ToM network and the ELN might be an artefact of including ToM studies in reviews of text comprehension research [e.g., Mar, 2004], or whether it is because of an overlap of the processes recruited in the two domains.

The second question concerns the role of the lateral prefrontal cortex for language comprehension in context. Although there is neuropsychological evidence for text comprehension deficits after left PFC lesions [Ferstl et al., 2002], many imaging studies show clear dominance of temporal regions for text comprehension [e.g., Crinion et al., 2002]. Moreover, the lateral PFC encompasses a number of subregions with dissociable functions [Bookheimer, 2002; Brass et al., 2005]. The more ventral<sup>2</sup> PFC including the IFG is important for phonological, syntactic, and semantic processes. More dorsal PFC regions in the middle frontal gyrus and inferior frontal sulcus are important for executive functions and working memory. Although the linguistic processes are expected to be comparable to those on the word and sentence level, it is likely that the dorso-lateral PFC contribution, related to executive

functions, increases with the task demands. Whether it is crucial for text comprehension however, or merely a by-product of memory or executive functions is an open question.

The third question concerns the contribution of the right hemisphere (RH). Neuropsychological theories and clinical observations have long focussed on the importance of the RH for text comprehension and pragmatic interpretation [e.g., Beeman, 2005; Beeman and Chiarello, 1998; Brownell et al., 2000]. However, the evidence from both patient studies and imaging studies is mixed [Ferstl, in press; Lehmann and Tompkins, 2000; Mar, 2004]. For many of the subprocesses of text comprehension, there are studies showing clear RH activations and others that do not [metaphors: Rapp et al., 2004 vs. Bottini et al., 1994, situation model building: Maguire et al., 1999 vs. George et al., 1999, etc.; inferences: Mason and Just, 2004; Ferstl and von Cramon, 2001]. Depending on the particular focus, the interpretations of these studies, as well as overarching reviews of this work, stress the role of the RH [e.g., Bookheimer, 2002; Mar, 2004] or the functions of the left fronto-medial and lateral PFC regions [Ferstl, in press].

To further investigate these issues, we present a quantitative review of 23 relevant text comprehension studies. The analysis was conducted using a meta-analysis method developed by Neumann et al. [2005], which combines activation likelihood estimation [ALE; Turkeltaub et al., 2002] with replicator dynamics [Schuster and Sigmund, 1983; see also Lohmann and Bohn, 2002, for application in functional imaging]. In ALE, single activation peaks are considered evidence for underlying probability distributions. Based on these distributions, cortical regions or clusters are determined in which nonrandom activation can be assumed. The ALE result consists of a number of cortical areas that play a central role in processing the cognitive task of interest. In a second step, functionally related subnetworks are identified. The goal is to group ALE clusters together, based on their co-occurrence in the investigated studies. In order to do so, activation peaks falling within the ALE-clusters are subjected to a replicator process.

The contrasts included in the meta-analysis were collected from studies on higher level language comprehension. Studies were included in which either connected text was presented (e.g. stories) or in which pragmatic interpretations were required (e.g. metaphor comprehension). Studies on the comprehension of single sentences, in particular those concerned with syntactic or semantic processes, were excluded. Because of the small number of relevant articles, studies using fMRI as well as PET, auditory as well as visual presentation, and all languages were considered (e.g. German, Italian, French, Japanese, and English). Care was taken to exclude any contrasts in which ToM processes were targeted explicitly.

Four separate analyses were conducted based on a classification of the contrasts reported in the studies. First, all contrasts in which connected language was compared to a resting baseline were collected (Rest). To subtract percep-

<sup>2</sup>Following the neurological convention, the terms, "dorsal" (high) and "ventral" (low) are used when referring to the inferior–superior dimension of brain regions, corresponding to the z-axis in stereotactic space.

tual processes contrasts using a nonlanguage baseline, such as speech played backward, were considered (Language). The third analysis used contrasts in which word or sentence level language processing was subtracted by comparing coherent to incoherent text (Coherence). Finally, despite the heterogeneity of the relevant studies, an exploratory analysis was conducted using coordinates reflecting specific subprocesses of pragmatic language comprehension, in particular situation model building and metaphor comprehension; Special).

The first goal of the study was to replicate the results obtained in prior reviews for language comprehension compared with a resting or nonlanguage baseline [Ferstl, in press; Mar, 2004]. In particular, the importance of the aTLs, the fronto-medial cortex, and the TPJ were to be confirmed. Second, the role of the right hemisphere was of interest. By assessing the coactivations of regions using replicator dynamics, we attempted to gain insight into the interplay between regions, and possibly between homolog regions in the two hemispheres. A comparison of the four analyses was expected to shed light on the specificity of the regions for language processing in context. Of particular interest was whether the fronto-medial cortex could be associated with coherence building.

## METHODS

### Identification of Coordinates

Twenty three available neuroimaging studies were included in which connected text was presented. Studies using single sentences were included only if the task explicitly focussed on pragmatic or interpretative features (e.g. metaphor comprehension).

To identify relevant articles, a literature search using the database Current Contents<sup>®</sup> (all editions) was conducted in the fall of 2005. Current Contents is provided by the Institute for Scientific Information-Thomson Scientific<sup>®</sup> (<http://scientific.thomson.com/>) and contains bibliographic references of over 7,500 journals, including medical and psychology journals from 1993 onward. The results of a search using the key words discourse, text, context, and comprehension were screened for imaging studies meeting the requirements. Because not all authors use these keywords, in particular if other issues were in the focus of the study [e.g. Giraud et al., 2000], the search results were supplemented with other relevant studies that had come to our attention. Studies were excluded that used an anatomically based region of interest analysis without providing coordinates in a standard stereotactic space. Contrasts using special populations were excluded as well (e.g., left handers, brain injured patients).

All reported contrasts were categorized according to the baseline used. First, nine contrasts from seven studies were identified in which text comprehension was compared with a resting baseline without stimulus presentation (Rest). Second, 16 contrasts from 11 studies were

chosen that used a perceptual, nonlanguage baseline, such as speech played backwards or nonletter strings (Language). Third, 13 contrasts from 10 studies were identified in which coherent, comprehensible text was compared with an incoherent language baseline, such as word lists or unrelated sentences (Coherence). Finally, nine contrasts from six studies were considered in which specific processes such as metaphor comprehension or topic continuity were targeted by considering direct comparisons of qualitatively different language comprehension tasks (Special). Three of these studies concerned metaphor comprehension and moral interpretation, whereas the others were concerned with situation model building, i.e., with setting up a globally coherent representation of connected text [Kintsch, 1998]. It is important to keep in mind that this last category combines heterogeneous subprocesses that might not yield consistent, overlapping results.

Care was taken to exclude contrasts specifically probing ToM processes. For instance, from the study by Fletcher et al. [1995] only the contrast comparing physical stories to unrelated sentences was entered into the analyses, but not the contrast comparing ToM stories or all stories to unrelated sentences. The complete list of contrasts is given in Tables I-IV.

For each of the four categories, all coordinates listed in the articles were entered. If necessary, MNI coordinates were transformed into Talairach coordinates using the formula proposed by Matthew Brett ([www.mrc-cbu.cam.ac.uk](http://www.mrc-cbu.cam.ac.uk)). The total number of coordinates entering the analyses was 93 for Rest, 156 for Language, 76 for Coherence, and 48 for Special.

### Meta-Analysis

The analysis was performed as proposed by Neumann et al. [2005]. First, activation likelihood estimation (ALE) [Derrfuss et al., 2005; Lancaster et al., 2005; Turkeltaub et al., 2002] was applied to the list of peak coordinates of the activations reported in the original articles. Single activation peaks were represented by three-dimensional Gaussian probability distributions with a standard deviation of 4 mm (FWHM = 9.4 mm). This standard deviation was chosen to approximately match filter sizes commonly used in fMRI studies. The union of the distributions yielded empirical activation likelihood estimates for all voxels, including voxels that were not in the collection of reported peaks. A voxel in close vicinity to one or several peak coordinates thereby received a higher ALE compared with voxels far removed.

Using Monte Carlo simulations, an ALE map for randomly distributed activation peaks was computed, containing as many activation peaks as reported. Specifically for 1,000 times, voxels of a brain volume mask were randomly chosen to represent activation peaks according to a uniform distribution, and ALE values were calculated for all voxels in the brain volume. The entire volume of the

**TABLE I. Contrasts entering into the meta-analysis for the comprehension of coherent language compared with a resting baseline**

Authors	Year	Method	Modality	Contrast	No. of peaks
Caplan, Dapretto	2001	fMRI	Auditive	Reasoning vs. rest (make sense judgment on question-answer pairs)	12
Caplan, Dapretto	2001	fMRI	Auditive	Topic vs. rest (topic judgment on question-answer pairs)	16
Kansaku et al.	2000	fMRI	Auditive	Story—no voice females	4
Kansaku et al.	2000	fMRI	Auditive	Story—no voice males	2
Miura et al.	2005	fMRI	Visual	Modern Japanese text vs. rest	4
Papathanassiou et al.	2000	PET	Auditive	Story–Rest	16
Perani et al.	1996	PET	Auditive	Italian story—attentive silence	10
Rapp et al.	2004	fMRI	Visual	Literal sentences vs. baseline (evaluation of connotation)	12
Tzourio et al.	1998	PET	Auditive	Right handers: text–Rest	17

The number of peak coordinates from each contrast is given in the rightmost column. The total number of peak coordinates entering into this analysis was 93.

brain normalized to the standard size as provided by the software package LIPSIA [Lohmann et al., 2001] served as mask volume, but the distribution of random activation peaks was restricted to the area spanned by the minimum and maximum Talairach coordinates of the empirical activation peaks. The resulting 1,000 ALE maps were averaged to yield the map that served as null hypothesis against which the significance of the empirical ALE values was tested [Turkeltaub et al., 2002]. Voxels were considered significantly activated if they exceeded an ALE threshold corresponding to an  $\alpha$ -level of 0.05% (uncorrected). Topologically connected areas of at least three voxels exceeding this threshold were considered as clusters of activation.

From the coordinates falling within the determined clusters, a co-occurrence matrix was formed, recording for

each pair of clusters the number of co-occurrences in the individual contrasts. This matrix was then subjected to a replicator process. Based on the principles of natural selection, this process determines subnetworks of ALE clusters with the property that every cluster included in the network co-occurs relatively often with every other network member [Lohmann and Bohn, 2002; Neumann et al., 2005; Schuster and Sigmund, 1983]. Specifically, at the beginning of the replicator process, each ALE cluster was assigned a so-called membership value. After convergence of the replicator process, a cluster was considered a member of the so-called dominant network, if its membership value exceeded the average membership value  $1/n$ , where  $n$  is the number of ALE clusters included in the analysis. This way, a network of cortical areas that are likely to be most

**TABLE II. Contrasts entering into the meta-analysis reflecting language processing compared with a nonlanguage perceptual baseline**

Authors	Year	Method	Modality	Contrast	No. of peaks
Alho et al.	2003	PET	Auditive	L-C (attending to story presented in left ear–visual random text)	8
Alho et al.	2003	PET	Auditive	R-C (attending to story presented in right ear–visual random text)	9
Crinion et al.	2003	PET	Auditive	Stories–reversed stories (T–) (no attention to voice identity)	9
Crinion et al.	2003	PET	Auditive	Stories–reversed stories (T+) (attention to voice identity)	10
Ferstl, von Cramon	2001	fMRI	Visual	Sentence pairs vs. letter strings (coherence judgment)	10
Ferstl, von Cramon	2002	fMRI	Auditive	Sentence pairs vs. pronounceable pseudo-language—part A: logic	8
Giraud et al.	2000	PET	Auditive	Story–vowels	7
Kansaku et al.	2000	fMRI	Auditive	Story–reverse story (female speaker)	2
Kansaku et al.	2000	fMRI	Auditive	Story–reverse story (male speaker)	2
Perani et al.	1996	PET	Auditive	Italian—story played backwards	5
Perani et al.	1998	PET	Auditive	Italian vs. backwards Japanese	4
Perani et al.	1998	PET	Auditive	Native language—native language backwards (Spanish or Catalan)	17
Robertson et al.	2000	fMRI	Visual	Sentences with indefinite articles vs. nonletter character strings	8
Robertson et al.	2000	fMRI	Visual	Sentences with definite article vs. nonletter strings	8
Vorobyev et al.	2004	PET	Auditive	V-C (reading story–visual random text)	15
Xu et al.	2005	fMRI	Visual	Narrative–letter strings	34

Sixteen contrasts from 11 studies were considered relevant. The total number of peak coordinates entering into the analysis was 156.

**TABLE III. Contrasts entering into the meta-analysis for the comprehension of coherent compared with incoherent language**

Authors	Year	Method	Modality	Contrast	No. of peaks
Bottini et al.	1994	PET	Visual	Sentence processing vs. lexical decision	16
Ferstl, von Cramon	2001	fMRI	Visual	Coherent vs. incoherent sentence pairs	2
Ferstl, von Cramon	2002	fMRI	Auditive	Coherent vs. incoherent—part A: logic	12
Fletcher et al.	1995	PET	Visual	Physical stories vs. unlinked sentences	4
Giraud et al.	2000	PET	Auditive	Story-sentences	5
Kuperberg et al.	2000	fMRI	Auditive	Normal sentences vs. pragmatic violations	10
Maguire et al.	1999	PET	Visual	Standard story vs. nonstandard story, without picture	2
Maguire et al.	1999	PET	Visual	Nonstandard stories, with relevant picture vs. without, 1st presentation	1
Maguire et al.	1999	PET	Visual	With relevant picture vs. no picture, 2nd presentation	1
Robertson et al.	2000	fMRI	Visual	Definite vs. indefinite articles	2
Vogele et al.	2001	fMRI	Visual	Stories (no ToM: T-) vs. sentences	4
Xu et al.	2005	fMRI	Visual	Narrative-sentences	17

Studies were included in which coherent texts were compared with an incoherent or less coherent language baseline. 76 peak coordinates entered into this analysis.

relevant for the cognitive task was chosen. After identifying this dominant network, its clusters were removed from the co-occurrence matrix and the replicator process was repeated to yield a second network from the remaining clusters [Lancaster et al., 2005; Lohmann and Bohn, 2002; Neumann et al., 2006]. For each of the resulting networks, the contributing contrasts were inspected. To uncover findings that have been replicated at least once, only those replicator networks are reported which co-occurred in at least two studies.

## RESULTS

### Rest

The contrasts in which language was compared with a resting baseline yielded a bilateral, fronto-temporal network of regions listed in Table V and displayed in Figure 1. In addition to large mid superior temporal gyrus activations, there were significant homolog regions in the lateral

aTLs, reaching from the STS into the temporal pole. Taken together, the sizes of the temporal activations were comparable in the two hemispheres. Lateral prefrontal activations included the frontal operculum in both hemispheres, a left-sided premotor region in the precentral gyrus and a right-sided area in the middle frontal gyrus close to the junction of the precentral and inferior frontal sulci. The only medial region showing a significant contribution was the presupplementary motor area (pre-SMA; BA 6/8).

The results of the replicator dynamics yielded a dominant network consisting of the two anterior temporal foci (R1, R2) and the left STG region (R3). These regions were concurrently activated in three of the seven studies.

### Language

The language contrasts, compared with a perceptual baseline, uncovered a large network of fronto-temporal regions, remarkably similar to that found for the Rest contrasts. These results are shown in Table VI and Figure 2.

**TABLE IV. Contrasts entering into the meta-analysis reflecting special text comprehension processes**

Authors	Year	Method	Modality	Contrast	No. of peaks
Bottini et al.	1994	PET	Visual	Metaphors task vs. sentence task	6
Caplan, Dapretto	2001	fMRI	Auditive	Topic vs. reasoning	14
Ferstl et al.	2005	fMRI	Auditive	Inconsistent stories vs. consistent; event related	1
Ferstl et al.	2005	fMRI	Auditive	Chronological stories: inconsistent vs. consistent; epoch-related	2
Ferstl et al.	2005	fMRI	Auditive	Emotional stories: inconsistent vs. consistent, epoch-related	3
Nichelli et al.	1995	PET	Visual	Semantic-font	3
Nichelli et al.	1995	PET	Visual	Moral-semantic	7
Rapp et al.	2004	fMRI	Visual	Metaphor vs. literal	3
Xu et al.	2005	fMRI	Visual	Outcome-setting	9

This analysis is considered exploratory because of the diversity of processes included, such as metaphor comprehension or the integration of inconsistent information. The total number of peak coordinates entering into this analysis was 48.

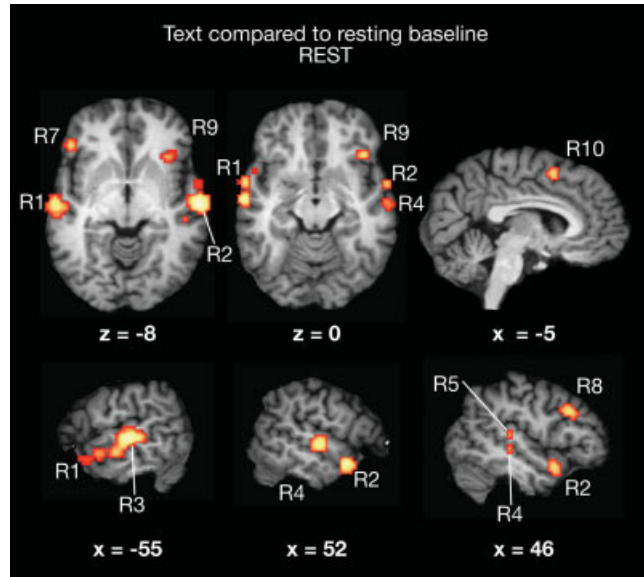
**TABLE V. Clusters in the ALE solution for all contrasts in which language processing was compared with a resting baseline**

ID	Size	Side	x	y	z	Anatomical location	RD
R1	1107	L	-59	0	-9	aTL	1
R2	2133	R	52	6	-15	aTL	1
R3	4401	L	-59	-24	6	STG/STS	1
R4	2295	R	58	-18	0	STG/STS	
R5	216	R	49	-33	6	STG/STS	
R6	405	L	-47	0	48	GPrec	
R7	405	L	-47	30	0	IFG, triangular part (BA 47)	
R8	810	R	46	18	30	MFG/IFS	
R9	675	R	37	21	-6	IFG, opercular part	
R10	405	L	-5	6	51	pre-SMA	

Peak coordinates and sizes of the regions displayed in Figure 1 are listed. The replicator dynamics yielded one network of coactivated clusters, marked in the right-most column.

aTL, anterior temporal lobe; STG, superior temporal gyrus; STS, superior temporal sulcus; GPrec, precentral gyrus; IFG, inferior frontal gyrus; IFS, inferior frontal sulcus; MFG, middle frontal gyrus; SMA, supplementary motor area; RD, label for the replicator dynamics membership of the cluster.

The most obvious difference is that the language network is more clearly left-lateralized. Once more, there were bilateral contributions of the aTLs, and left-dominant mid-



**Figure 1.**

Results of the meta-analysis of language processing in context against a resting baseline, based on the contrasts listed in Table I. The labels of the regions correspond to those provided in Table V.

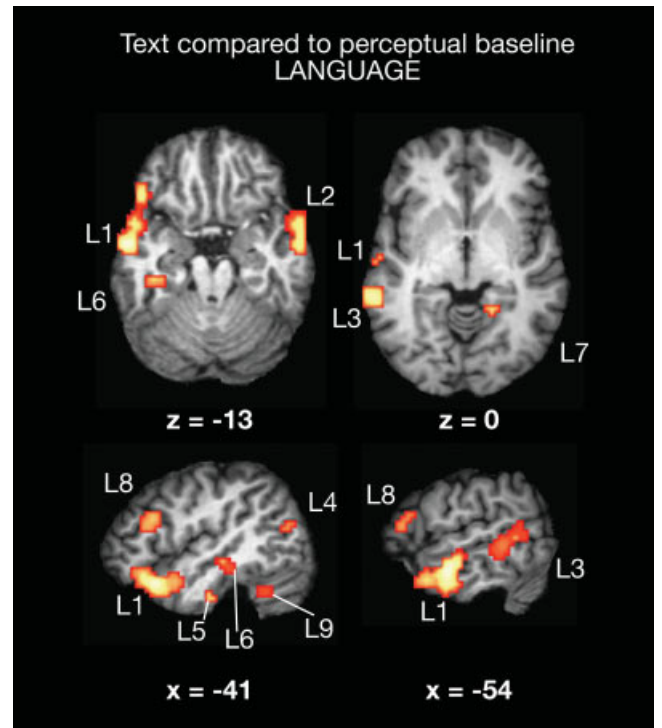
**TABLE VI. Clusters in the ALE solution for all contrasts in which language processing was compared with a perceptual baseline**

ID	Size	Side	x	y	z	Anatomical location	RD
L1	7317	L	-44	15	-21	aTL	1
L2	3159	R	55	9	-18	aTL	1
L3	2457	L	-59	-42	3	MTG/posterior STS	2
L4	162	L	-44	-69	15	MTG/posterior STS	
L5	162	L	-44	-18	-30	ITG/OTS	
L6	675	L	-41	-30	-9	ITG/fusiform gyrus	
L7	162	R	16	-48	0	SColl	
L8	1431	L	-47	21	18	IFG/triangular part	2
L9	216	L	-41	-54	-24	Cerebellum	

Peak coordinates and sizes of the regions displayed in Figure 2 are listed. The replicator dynamics yielded two networks of coactivated clusters, marked in the right-most column with the same number.

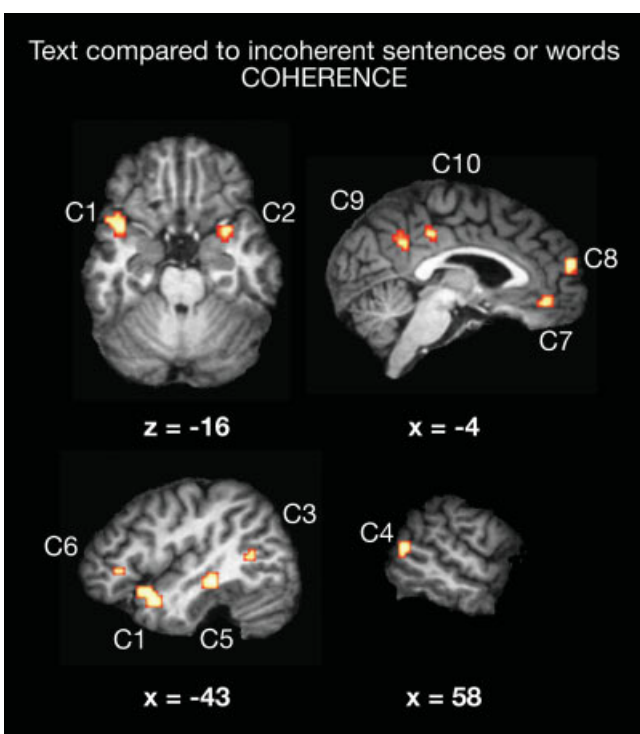
ITG, inferior temporal gyrus; SColl, collateral sulcus; OTS, occipitotemporal sulcus; all other abbreviations as in Table 5.

dle and posterior temporal regions. In addition to these regions, which were only slightly more ventral than the STG regions reported in the Rest analysis, a focus in the inferior temporal lobe and a contribution of the right col-



**Figure 2.**

Results of the meta-analysis of language processing in context against a nonlanguage baseline, based on the contrasts listed in Table II. The labels of the regions correspond to those provided in Table VI.



**Figure 3.**

Results of the meta-analysis of comprehending coherent language compared with incoherent language. This analysis is based on the contrasts listed in Table III. The labels of the regions correspond to those provided in Table VII.

lateral sulcus emerged. Finally, there were prefrontal activations in the triangular part of the left IFG and in the right precentral sulcus, as well as a focus in the left lateral cerebellum.

The replicator dynamics identified the bilateral anterior temporal regions to be dominant (L1, L2). Seven contrasts from five studies contributed to this result. When removing this first replicator solution, the left IFG (L8) and the left posterior STS (L3) proved to form a subnetwork. These regions might be considered part of the perisylvian language cortex and thus to reflect word and sentence level processes.

### Coherence

The contrasts reflecting the comprehension of coherent, comprehensible text showed bilateral aTL activation. The foci are more medial than those in the two previous analyses. All other regions were in the left hemisphere. As before, the mid and posterior middle temporal gyrus, as well as the left IFG were part of the significant network. In addition, four left-medial areas proved to be important for coherence building: two regions lay in the posterior cingulate cortex and inferior precuneus (PCC/prec, BA23/31) one in the dmPFC (BA 10), and one in the ventro-medial

prefrontal cortex [BA 11]. In the recently introduced terminology by Ramnani and Owen [2004], these areas are the polar and rostral sections of BA 10.

Based on two studies, the replicator dynamics identified the left aTL (C1) and the mid MTG (C5) as coactivated. The second subnetwork consisted of the dmPFC (C8) and the more posterior of the two PCC/prec regions (C9). Although this latter result resembles the findings from the coherence judgment task [Ferstl and von Cramon, 2001, 2002], two studies from other laboratories gave rise to this finding [Bottini et al., 1994; Xu et al., 2005] (Fig. 3, Table VII).

### Special

The contrasts reflecting special text interpretation processes such as metaphor comprehension or the understanding of the moral of a story are to be taken with caution. Only nine studies entered into this analysis and only 48 coordinates were used. Nevertheless, the results once more yield bilateral anterior temporal regions. However, the RH area is now larger, and the LH peak lies more ventrally, at the anterior end of the inferior temporal gyrus. In addition, there were three regions in the left IFG (BA45/47/11). Right-sided foci included two dorso-lateral PFC regions, close to the inferior frontal and precentral sulci (BA9/46), and the TPJ.

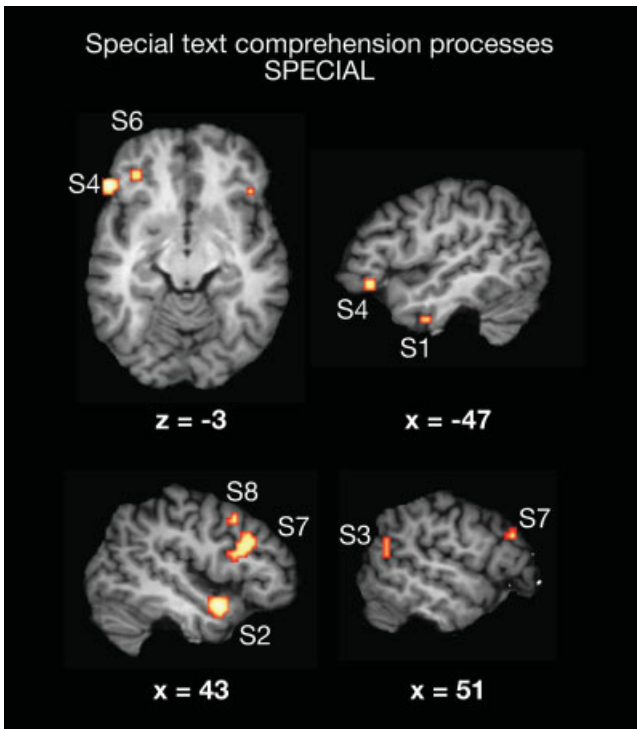
Because of the heterogeneity of the contrasts, only the two largest regions, the right aTL and the right IFS, were based on coordinates from more than one study. Similarly, none of the replicator dynamics solutions reflected co-occurrence in more than one study (Fig. 4, Table VIII).

**TABLE VII. Peak coordinates and sizes of the regions found significant for the ALE analysis of all contrasts reflecting coherence building**

ID	Size	Side	x	y	z	Anatomical location	RD
C1	999	L	-38	6	-21	aTL	1
C2	324	R	28	3	-18	aTL	
C3	189	L	-41	-57	9	posterior STS	
C4	405	R	58	-57	12	posterior STS	
C5	513	L	-41	-30	-6	mid MTG	1
C6	162	L	-41	30	0	IFG, opercular part, BA 47	
C7	162	R	1	45	-6	vmPFC (BA 11)	
C8	432	L	-5	57	15	dmPFC (BA 9/10)	2
C9	432	L	-8	-51	33	IPrec	2
C10	108	L	-5	-30	36	IPrec	

The regions are displayed in Figure 3. The replicator dynamics yielded two networks of coactivated clusters, marked in the right-most column with the same number.

vmPFC, ventro-medial prefrontal cortex; dmPFC, dorso-medial PFC; IPrec, inferior precuneus; RD, label for the replicator dynamics membership of the cluster.



**Figure 4.**

Results of the meta-analysis for special processes of text comprehension, based on the contrasts listed in Table IV. The labels of the regions correspond to those provided in Table VIII.

**Overlap**

To visually illustrate the stability of the anterior temporal contribution, an overlap map of the ALE maps was calculated. There was no overlapping result for all four analyses. Excluding the Special network, only one voxel was common to the Rest, Language, and Coherence comparison.

**TABLE VIII. Significant clusters from the ALE of all contrasts in which special issues on contextual language integration were investigated**

ID	Size	Side	x	y	z	Anatomical location
S1	108	L	-47	-9	-30	aTL/ITG
S2	945	R	43	3	-15	aTL
S3	108	R	52	-57	24	TPJ
S4	540	L	-50	27	-6	IFG/BA47
S5	108	L	-38	33	0	IFG/GOP
S6	108	L	-32	33	-6	IFG/GOP
S7	864	R	43	18	21	IFS/SPrec
S8	108	R	43	12	39	GPreC

Peak coordinates and sizes of the regions are listed. The regions are displayed in Figure 4. The replicator dynamics did not yield subnetworks coactivated in more than one study. IFS, inferior frontal sulcus; GOP, posterior orbital gyrus; SPrec, precentral sulcus; other abbreviations as before.

sons. The location of this voxel in the left aTL is shown in Figure 5.

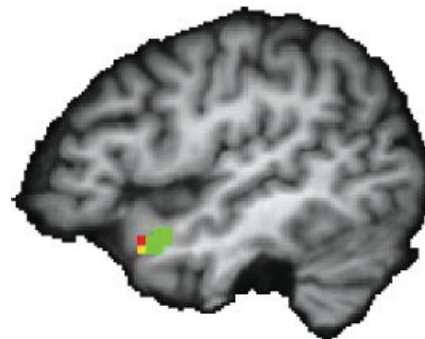
**DISCUSSION**

The meta-analysis of language processing in context took advantage of a two-step procedure. In the first step, significant clusters of activation were determined based on activation likelihood estimation. In the second step, replicator dynamics were used to identify subnetworks of coactivated regions. Previous studies using similar numbers of peak coordinates as used here have established the method as reliable [Lancaster et al., 2005; Neumann et al., 2005]. Moreover, the analyses are based on a larger number of studies than prior qualitative reviews [e.g., Gernsbacher and Kaschak, 2003; Mar, 2004] so that we can be confident about their results.

**The Anterior Temporal Lobes**

The most striking result of the meta-analysis is the stability of activation in the aTLs. In all four analyses, which were in part based on different studies from different laboratories using different scanning methods, modalities, and materials, bilateral involvement of this region emerged. An illustration of the overlap between the results uncovered a small region in the left aTL that contributed to three of the four results. The main difference between the analyses was that for the coherence analysis, the activation was more medial than for the Rest, Language, and Special analyses.

In all four analyses, the coordinates of the homolog regions in the left and right hemispheres were in remarkably close vicinity. In the Rest and Language analyses, the replicator dynamics solution identified the left and right aTL regions as the dominant subnetwork. This result sug-



**Figure 5.**

The spatial overlap of the ALE clusters from the three comparisons: Rest, Language, and Coherence. Only one voxel in the left aTL was contained in all three networks (marked in red:  $x = -47, y = -12, z = -15$ ). The neighboring regions were common to two comparisons (green: Rest and Language, yellow: Language and Coherence).

gests a bilateral function of the aTL in which the homolog regions are closely interrelated.

The meta-analyses corroborate the role of the aTLs for text comprehension. Although their specific function is still under debate, a crucial contribution to sentence comprehension has been appreciated recently [for review see Stowe et al., 2005]. The theoretical proposals for aTL functionality include memory processes, in particular for autobiographical and emotional, episodic memory, and semantic processes, in particular category specific retrieval of proper names or animate entities [e.g., Damasio et al., 1996; Leveroni et al., 2000; Maratos et al., 2001; Martin and Chao, 2001]. The finding of sensitivity to syntactic violations [Friederici et al., 2003] might either be explained by resulting difficulties with semantic integration or by slight shifts in the anatomical location. In studies on language interpretation, the activations tend to be at the anteriormost end of the STS, reaching into the temporal pole, whereas activations related to syntactic features lay more dorsally, in the anterior temporal plane. Attempts to dissociate these anatomical regions [Humphries et al., 2005] are often difficult because lack or violations of syntactic structure immediately affect the ease of semantic integration [Ferstl, in press].

Because the temporal lobes are multimodal association areas, it is likely that syntactic, semantic, and episodic information sources are indeed integrated to transform the language input into a meaning based representation. The most parsimonious account, in the framework of text comprehension, is that the aTL implements propositionalization, the process required for combining words into semantically based content units [Kintsch, 1998]. This process, similar to the concept of semantic encoding [Stowe et al., 2005], clearly utilizes prosodic, syntactic, and lexical information to derive a semantic representation. It is an open question whether the more medial anterior temporal regions uncovered in the coherence contrasts have a different functionality, for example related to autobiographical memory [Fink et al., 1996].

### The Middle and Posterior Temporal Lobe

In all four analyses, middle and posterior temporal activations were significant. These activations were in the mid portion of the superior temporal gyrus in the Rest analysis, reflecting the fact that the majority of contrasts entering into this analysis used auditory presentation. Turkeltaub [2002], in their meta-analysis of word reading, reported bilateral peaks very close to ours. These middle temporal activations are slightly more ventral, in the STS, when a perceptual baseline is subtracted [cf. Dehaene et al., 1997]. In the language analysis, there were three foci along the left STS and the adjacent part of the middle temporal gyrus. For more specific contrasts, the activation peaks appear to lay more posteriorly and dorsally. In particular, the Coherence analysis and the Special analysis yielded foci in the left posterior middle temporal gyrus and the right TPJ, respectively. Bavelier et al. [1997], in an early

study on sentence processing, analyzed the time course of temporal activation. They found a spread of activation from mid superior temporal lobe both anteriorly into the temporal pole and posteriorly into the TPJ. Thus, the mid portion is likely to implement basic language perception, whereas the anterior and posterior temporal lobes are concerned with integration and interpretation.

### The Lateral Prefrontal Cortex

The activations in the lateral PFC lay predominantly in the IFG. One activation fell into the dorsal triangular part close to the inferior frontal sulcus (BA 44/45), traditionally named Broca's area. This region proved important for language processing compared with a perceptual baseline. Because this region co-occurred with activation in superior temporal gyrus (or Wernicke's area) this result is likely to reflect language processing on the word and sentence level. This activation can be considered to reflect language processing on the word and sentence level.

Most of the IFG activations were in the opercular part or reaching into the orbito-frontal cortex (BA47/12). Three of these activations were found in the Special analysis, and come from one study only [Ferstl et al., 2005]. More readily interpretable are the bilateral activations in the frontal operculum for the Rest analysis. This finding has often been reported for auditory language processing, independent of context or comprehensibility [Meyer et al., 2000]. However, the left-sided opercular region was also significant in the analysis of Coherence. Thus, it is likely that its function goes beyond language perception. Proposals include the processing of sentence level context [Baumgärtner et al., 2002] or the need to make decisions based on recently encountered stimuli [Petrides et al., 2002; Ferstl et al., 2005].

Interestingly, the meta-analyses reported here reflect the lack of large dorso-lateral prefrontal activations in many of the studies. There were two right-sided foci. In the Rest analysis, the right middle frontal gyrus proved active, whereas in the Special analysis a slightly more ventral region in the inferior frontal sulcus emerged.

### The Medial Wall of the Left Hemisphere

Medial activations proved reliable in two of the four analyses. An area in the presupplementary motor area (pre-SMA) appeared when language processing was compared with a resting baseline. The peak of this activation is almost identical to that found in the meta-analysis of word reading [Turkeltaub et al., 2002]. Thus, it is likely that the activation is related to inner speech used for encoding both auditory and written language.

Consistent with previous reports on inferencing, we also found medial activations in the meta-analysis of coherence building. There was a pair of activations in the fronto- and parieto-medial cortices, which replicator dynamics suggested to be coactivated. Because of cortico-cortical connec-

tions of these two regions [Barbas, 1992], coactivation of fronto- and parieto-medial areas has been reported in numerous studies [e.g., Zysset et al., 2002, 2003], and in particular in studies on inferencing [Ferstl and von Cramon, 2001, 2002]. The replicator solution reported here was based on coordinates from two studies from other laboratories, though [Bottini et al., 1994; Xu et al., 2005]. This finding shows that the results obtained with the coherence judgment task [Ferstl and von Cramon, 2001, 2002] generalize to other materials and tasks.

The small ventro-medial region in BA 11 (or, more specifically, in rostral BA 10 in the terminology recently suggested by Ramnani and Owen, 2004) was also significant in the Coherence analysis. A number of recent results have implicated this region for emotion processing in the context of verbal tasks and on verbal humor [Ferstl et al., 2005; Goel and Dolan, 2001, 2004; Siebörger et al., 2004]. And indeed, among the studies contributing to this result is an experiment in which funny cartoons provide the background knowledge for loosely connected sentences [Maguire et al., 1999]. The activation uncovered by the meta-analysis might thus reflect concurrent affective reactions elicited by the language stimuli.

### The Right Hemisphere

There were right hemisphere activations in all analyses. Besides the aforementioned regions in the right temporal lobe, right prefrontal activations were reliable in the Rest and Special analyses. For auditory presentation, the finding of bilateral frontal operculum activation replicates results from sentence processing studies [e.g., Meyer et al., 2000]. The more dorsal right PFC activation found for Special contrasts was based mostly on the study by Caplan and Dapretto [2001]. However, these authors reported bilateral patterns, including left PFC activations.

The prediction of the RH being important for inference processes [Beeman, 2005; Beeman and Chiarello, 1998; Mason and Just, 2004] was not supported. The analysis of coherence building yielded a clearly left dominant network, with the right aTL as the only significant RH region. It is important to note, however, that some imaging studies reporting RH activation for inference processes had to be excluded because of the lack of reported coordinates [George et al., 1999; Mason and Just, 2004]. As argued elsewhere [Ferstl, in press], it might be that a region of interest analysis, as conducted in these studies, compensates for the higher anatomical variability of the RH, and is thus more powerful to detect RH contributions.

### Theory-of-Mind

It would be desirable to conduct a meta-analysis of ToM studies using the mathematical methods presented here for an objective comparison. However, the results presented strongly suggest an overlap between the ELN and the regions implicated for ToM processes in qualitative reviews [Frith and Frith, 2003]. Although in contrast to

other reviews, specific contrasts testing for ToM using verbal materials were excluded, the aTL, TPJ, and dmPFC regions were clearly significant in several analyses. The most striking result was the network obtained in the Coherence analysis. In addition to the aTLs, the dmPFC, and the posterior STS, this network included the IFG (BA 45/47) and the lower precuneus, regions closely connected to the dmPFC [Ramnani and Owen, 2004]. In particular, the Language and the Coherence analyses yielded posterior STS activations bilaterally, slightly more ventral than the TPJ region identified by Saxe and Kanwisher [2003] for ToM-story comprehension. Thus, the dmPFC seems to be specific for the successful interpretation of coherent language, rather than have a role in language comprehension in general.

Despite this overlap between the networks, there is no obvious causal direction. Although any communication might entail a ToM component [cf. Frith and Frith, 2003; Sperber and Wilson, 1995], many ToM tasks are explicitly verbal. For instance, the ToM-stories used by Saxe and Kanwisher [2003], but not their control texts, require elaborate inference processes. Even nonverbal ToM tasks [e.g., Castelli et al., 2000] are likely to elicit verbalization or narrativization [Bruner and Feldman, 1993]. In the only imaging study specifically designed to dissociate verbal inference from mentalizing processes [Ferstl and von Cramon, 2002], we found both coherence and ToM to be sufficient, but neither of them necessary, for engaging the dmPFC. Thus, we have argued for a domain-independent general process encompassing inferencing, evaluation, and ToM [Ferstl and von Cramon, 2002; Zysset et al., 2002, 2003], going beyond the processing of self-relevant stimulus materials [cf. Northoff and Bermpohl, 2004].

### Limitations of the Method

One serious limitation of the method is that only those coordinates can be included that the authors choose to report. Studies using anatomically defined regions of interest rather than standard stereotactic coordinates have to be excluded altogether. Some studies include only one peak coordinate for each connected region, even if it is rather large, and others report submaxima within these regions. Some authors use a masking procedure and some do not use whole-head measurement but only scan a predefined section of the brain. Thus, the reported results might be a subset of the relevant regions only.

Note further that the included studies have not been weighted for the number of subjects or scans, or the statistical threshold reported. Although the possible influence of such weighting has been discussed in the literature for some time [e.g., Lancaster et al., 2005; Neumann et al., 2005; Nielsen et al., 2004; Turkeltaub et al., 2002], to our knowledge no agreed-upon solution exists to date. This is because of the complex relationship between the relevant factors [see Lancaster et al., 2005 for a discussion of this issue]. Thus, changing or altering the data as little as possible seemed to be the most objective choice.

A second problem, specific to the issue at hand, is that there still is a lack of relevant empirical data. Particularly for the Special analysis, the large variety of tasks, methods, and presentation modalities does not allow us to attribute the resulting network to specific subprocesses of text comprehension. For example, an evaluation of lateral prefrontal activation can only be thoroughly investigated when the number of available studies renders feasible a comparison across tasks with differing complexity. Further experimentation is needed so that the suggestions reported here become more definite. In particular, it is necessary to conduct well-designed studies to evaluate specific subprocesses of comprehension and pragmatic interpretation.

## CONCLUSIONS

The application of recently developed meta-analysis methods to the functional neuroanatomy of text comprehension has identified a network of fronto-temporal brain regions, extending beyond the perisylvian language cortex. In addition to left inferior frontal and posterior superior temporal regions, there was an extremely stable contribution of the aTLs. Importantly, the specific role of fronto- as well as parieto-medial regions for inference processes was strengthened. In contrast to previous reviews of text comprehension studies [e.g., Gernsbacher and Kaschak, 2003; Mar, 2004], the meta-analysis did not provide evidence for a special role of the right hemisphere for inference processes. An attribution of right prefrontal and right temporal functions to special comprehension tasks [e.g., Caplan and Dapretto, 2001; Ferstl et al., 2005] requires further empirical replications.

## REFERENCES

- Alho K, Vorobyev VA, Medvedev SV, Pakhomov SV, Roudas MS, Tervaniemi M, van Zuijen T, Näätänen R (2003): Hemispheric lateralization of cerebral blood-flow changes during selective listening to dichotically presented continuous speech. *Brain Res Cogn Brain Res* 17:201–211.
- Barbas H (1992): Architecture and cortical connections of the prefrontal cortex in the rhesus monkey. *Adv Neurol* 57:91–115.
- Baumgärtner A, Weiller C, Büchel C (2002): Event-related fMRI reveals cortical sites involved in contextual sentence integration. *Neuroimage* 16:736–745.
- Bavelier D, Corina D, Jezzard P, Padmanabhan S, Clark VP, Karni A, Prinster A, Braun A, Lalwani A, Rauschecker JP, Turner R, Neville H (1997): Sentence reading: A functional MRI study at 4 Tesla. *J Cogn Neurosci* 9:664–686.
- Beeman M (2005): Bilateral brain processes for comprehending natural language. *Trends Cogn Sci* 9:512–518.
- Beeman M, Chiarello C (1998): Complementary right- and left-hemisphere language comprehension. *Psychol Sci* 7:2–8.
- Bookheimer S (2002): Functional MRI of language: New approaches to understanding the cortical organization of semantic processing. *Annu Rev Neurosci* 25:151–188.
- Bottini G, Corcoran R, Sterzi R, Paulesu E, Schenone P, Scarpa P, Frackowiak RSJ, Frith CD (1994): The role of the right hemisphere in the interpretation of figurative aspects of language: A positron emission tomography activation study. *Brain* 117:1241–1253.
- Brass M, Derrfuss J, Forstmann B, von Cramon DY (2005): The role of the inferior frontal junction area in cognitive control. *Trends Cogn Sci* 9:314–316.
- Brownell HH, Griffin R, Winner E, Friedmann O, Happe F (2000): Cerebral lateralization and theory of mind. In: Baron-Cohen, S, Tager-Flusberg H, Cohen DJ, editors. *Understanding Other Minds: Perspectives from Developmental Cognitive Neuroscience*. Oxford: Oxford University Press. pp 306–333.
- Bruner J, Feldman C (1993): Theories of mind and the problem of autism. In: Baron-Cohen S, Tager-Flusberg H, Cohen DJ, editors. *Understanding Other Minds: Perspectives from Autism*. Oxford: Oxford University Press. pp 267–291.
- Caplan R, Dapretto M (2001): Making sense during conversation: An fMRI study. *Neuroreport* 12:3625–3632.
- Castelli F, Happé F, Frith U, Frith C (2000): Movement and mind: A functional imaging study of perception and interpretation of complex intentional movement patterns. *Neuroimage* 12:314–325.
- Crinion JT, Lambon-Ralph A, Warburton EA, Howard D, Wise RJS (2003): Temporal lobe regions engaged during normal speech comprehension. *Brain* 126:1193–1201.
- Damasio H, Grabowski TJ, Tranel D, Hichwa RD, Damasio AR (1996): A neural basis for lexical retrieval. *Nature* 380:499–505.
- Dehaene S, Dupoux E, Mehler J, Cohen L, Paulesu E, Perani D, et al. (1997): Anatomical variability in the cortical representation of first and second language. *Neuroreport* 8:3809–3815.
- Derrfuss J, Brass M, Neumann J, von Cramon DY (2005): Involvement of the inferior frontal junction in cognitive control: Meta-analyses of switching and Stroop studies. *Hum Brain Mapp* 25:22–34.
- Ferstl EC: The functional neuroanatomy of text comprehension: What's the story so far? In: Schmalhofer F, Perfetti CA, editors. *Higher Level Language Processes in the Brain: Inference and Comprehension Processes*. Mahwah, NJ: Lawrence Erlbaum (in press).
- Ferstl EC, von Cramon DY (2001): The role of coherence and cohesion in text comprehension: An event-related fMRI study. *Brain Res Cogn Brain Res* 11:325–340.
- Ferstl EC, von Cramon DY (2002): What does the fronto-medial cortex contribute to language processing: Coherence or theory of mind? *Neuroimage* 17:1599–1612.
- Ferstl EC, Guthke T, von Cramon DY (2002): Text comprehension after brain injury: Left prefrontal lesions affect inference processes. *Neuropsychology* 16:292–308.
- Ferstl EC, Rinck M, von Cramon DY (2005): Emotional and temporal aspects of situation model processing during text comprehension: An event-related fMRI study. *J Cogn Neurosci* 17:724–739.
- Fink GR, Markowitsch HJ, Reinkemeier M, Bruckbauer T, Kessler J, Heiss WD (1996): Cerebral representation of one's own past: Neural networks involved in autobiographical memory. *J Neurosci* 16:4275–4282.
- Fletcher PC, Happé F, Frith U, Baker SC, Dolan RJ, Frackowiak RSJ, Frith CD (1995): Other minds in the brain: A functional imaging study of "theory of mind" in story comprehension. *Cognition* 57:109–128.
- Friederici AD, Rüschemeyer SA, Hahne A, Fiebach C (2003): The role of left inferior frontal and superior temporal cortex in sentence comprehension: Localizing syntactic and semantic processes. *Cereb Cortex* 13:170–177.
- Frith CD, Frith U (2003): Development and neurophysiology of mentalizing. *Philos Trans R Soc Lond B Biol Sci* 358:459–473.

- Gallagher HL, Frith CD (2003): Functional imaging of "theory of mind". *Trends Cogn Sci* 7:77–83.
- Gernsbacher MA, Kaschak MP (2003): Neuroimaging studies of language production and comprehension. *Annu Rev Psychol* 54:16.1–16.24.
- Giraud AL, Truy E, Frackowiak RSJ, Gregoire MC, Pujol JF, Collet L (2000): Differential recruitment of the speech processing system in healthy subjects and rehabilitated cochlear implant patients. *Brain* 123:1391–1402.
- Goel V, Dolan RJ (2001): The functional anatomy of humor: Segregating cognitive and affective components. *Nat Neurosci* 4:237–261.
- Goel V, Dolan RJ (2004): Reciprocal neural response within lateral and ventral medial prefrontal cortex during hot and cold reasoning. *Neuroimage* 20:2314–2321.
- Humphries C, Love T, Swinney D, Hickok G (2005): Response of anterior temporal cortex to syntactic and prosodic manipulations during sentence processing. *Hum Brain Mapp* 26:128–138.
- Kansaku K, Yamaura A, Kitazawa S (2000): Sex differences in lateralization revealed in the posterior language areas. *Cereb Cortex* 10:866–872.
- Kintsch W (1998): *Comprehension: A paradigm for cognition*. Cambridge: Cambridge University Press. 461 p.
- Kuperberg GR, McGuire PK, Bullmore ET, Brammer MJ, Rabe-Hesketh S, Wright IC, Lythgoe DJ, Williams SCR, David AS (2000): Common and distinct neural substrates for pragmatic, semantic, and syntactic processing of spoken sentences: An fMRI study. *J Cogn Neurosci* 12:321–341.
- Lancaster J, Laird, AR, Fox, PM, Glahn DE, Fox, PT (2005): Automated analysis of meta-analysis networks. *Hum Brain Mapp* 25:174–184.
- Lehmann MT, Tompkins CA (2000): Inferencing in adults with right hemisphere brain damage: An analysis of conflicting results. *Aphasiology* 14:485–499.
- Leveroni CL, Seidenberg M, Mayer AR, Mead LA, Binder JR, Rao SM (2000): Neural systems underlying the recognition of familiar and newly learned faces. *J Neurosci* 20:878–886.
- Lohmann G, Bohn S (2002): Using replicator dynamics for analyzing fMRI data of the human brain. *IEEE Trans Med Imaging* 21:485–492.
- Lohmann G, Muller K, Bosch V, Mentzel H, Hessler S, Chen L, Zysset S, von Cramon DY (2001): LIPSIA - a new software system for the evaluation of functional magnetic resonance images of the human brain. *Computerized Medical Imaging and Graphics* 25:449–457.
- Maguire EA, Frith CD, Morris RGM (1999): The functional neuroanatomy of comprehension and memory: The importance of prior knowledge. *Brain* 122:1839–1850.
- Mar RA (2004): The neuropsychology of narrative: Story comprehension, story production and their interrelation. *Neuropsychologia* 42:1414–1434.
- Maratos EJ, Dolan RJ, Morris JS, Henson RNA, Rugg MD (2001): Neural activity associated with episodic memory for emotional context. *Neuropsychologia* 29:910–920.
- Martin A, Chao LL (2001): Semantic memory and the brain: Structure and processes. *Curr Opin Neurobiol* 11:194–201.
- Mason RA, Just MA (2004): How the brain processes causal inferences in text: A multiple process theory of the function of the language network in both hemispheres. *Psychol Sci* 15:1–7.
- Mazoyer BM, Tzourio N, Frak V, Syrota A, Murayama N, Levyrier O, Salamon G, Dehaene S, Cohen L, Mehler J (1993): The cortical representation of speech. *J Cogn Neurosci* 5:467–479.
- Meyer M, Friederici AD, von Cramon DY (2000): Neurocognition of auditory sentence comprehension: Event related fMRI reveals sensitivity to syntactic violations and task demands. *Brain Res Cogn Brain Res* 9:19–33.
- Miura N, Watanabe J, Iwata K, Sassa Y, Riera J, Tsuchiya H, Sato S, Horie K, Takahashi M, Kitamura M, Kawashima R (2005): Cortical activation during reading of ancient versus modern Japanese texts: An fMRI study. *Neuroimage* 26:426–431.
- Neumann J, Lohmann G, Derrfuss J, von Cramon DY (2005): Meta-analysis of functional imaging data using replicator dynamics. *Hum Brain Mapp* 25:165–173.
- Neumann J, von Cramon DY, Forstmann BU, Zysset S, Lohmann G (2006): The parcellation of cortical areas using replicator dynamics in fMRI. *Neuroimage* 32:208–219.
- Nichelli P, Grafman J, Pietrini P, Clark K, Lee KY, Miletich R (1995): Where the brain appreciates the moral of a story. *Neuroreport* 6:2309–2313.
- Nielsen FA, Hansen LK, Balslev D (2004): Mining for associations between text and brain activation in a functional neuroimaging database. *Neuroinformatics* 2:369–380.
- Northoff G, Bermpohl F (2004): Cortical midline structures and the self. *Trends Cogn Sci* 8:102–107.
- Papathanassiou D, Etard O, Mellet E, Zago L, Mazoyer B, Tzourio-Mazoyer N (2000): A common language network for comprehension and production: A contribution to the definition of language epicenters with PET. *Neuroimage* 11:347–357.
- Perani D, Dehaene S, Grassi F, Cohen L, Cappa SF, Dupoux E, Fazio F, Mehler J (1996): Brain processing of native and foreign languages. *Neuroreport* 7:2439–2444.
- Perani D, Paulesu E, Galles NS, Dupoux E, Dehaene S, Bettinardi V, Cappa SF, Fazio F, Mehler J (1998): The bilingual brain. Proficiency and age of acquisition of the second language. *Brain* 121:1841–1852.
- Petrides M, Alivisatos B, Frey, S (2002): Differential activation of the human orbital, mid-ventrolateral, and mid-dorsolateral prefrontal cortex during the processing of visual stimuli. *Proc Natl Acad Sci USA* 99:5649–5654.
- Ramnani N, Owen AM (2004): Anterior prefrontal cortex: Insights into function from anatomy and neuroimaging. *Nat Rev Neurosci* 5:184–194.
- Rapp AM, Leube DT, Erb M, Grodd W, Kircher TJ (2004): Neural correlates of metaphor processing. *Brain Res Cogn Brain Res* 20:395–402.
- Robertson DA, Gernsbacher MA, Guidotti SJ, Robertson RRW, Irwin W, Mock BJ, Campana ME (2000): Functional neuroanatomy of the cognitive process of mapping during discourse comprehension. *Psychol Sci* 11:255–260.
- Saxe R, Kanwisher N (2003): People thinking about thinking people. The role of the temporo-parietal junction in "theory of mind". *Neuroimage* 19:1835–1842.
- Schuster P, Sigmund K (1983): Replicator dynamics. *J Theor Biol* 100:533–538.
- Sieböcker FT, Ferstl EC, Volkmann B, von Cramon DY (2004): Spass beiseite! Eine fMRI-Studie und eine behaviorale Patientstudie zu verbalem Humor und sprachlicher Revision [All joking aside! An fMRI study and a behavioral patient study on verbal humour and linguistic revision]. *Zeitschrift für Neuropsychologie* 15.
- Sperber D, Wilson D (1995): *Relevance: Communication and cognition*, 2nd ed. Oxford: Blackwell. 326 p.

- St. George M, Kutas M, Martinez A, Sereno MI (1999): Semantic integration in reading: Engagement of the right hemisphere during discourse processing. *Brain* 122:1317–1325.
- Stowe LA, Haverkort M, Zwarts F (2005): Rethinking the neurological basis of language. *Lingua* 115:997–1042.
- Turkeltaub PE, Eden GF, Jones KM, Zeffiro TA (2002): Meta-analysis of the functional neuroanatomy of single-word reading: Method and validation. *Neuroimage* 16:765–780.
- Tzourio N, Crivello F, Mellet E, Nkanga-Ngila B, Mazoyer B (1998): Functional anatomy of dominance for speech comprehension in left handers vs. right handers. *Neuroimage* 8:1–16.
- Vogele K, Bussfeld P, Newen A, Herrmann S, Happe F, Falkai P, Maier W, Shah NJ, Fink GR, Zilles K (2001): Mind Reading: Neural mechanisms of theory of mind and self-perspective. *Neuroimage* 14:170–181.
- Vorobyev VA, Alho K, Medvedev SV, Pakhomov SV, Roudas MS, Rutkovskaya JM, Tervaniemi M, van Zuijen TL, Näätänen, R (2004): Linguistic processing in visual and modality-nonspecific brain areas: PET recordings during selective attention. *Brain Res Cogn Brain Res* 20:309–322.
- Xu J, Kemeny S, Park G, Frattali C, Braun A (2005): Language in context: Emergent features of word, sentence, and narrative comprehension. *Neuroimage* 25:1002–1015.
- Zysset S, Huber O, Ferstl EC, von Cramon DY (2002): The anterior frontomedian cortex and evaluative judgment: An fMRI study. *Neuroimage* 15:983–991.
- Zysset S, Huber O, Samson A, Ferstl EC, von Cramon DY (2003): Functional specialization within the anterior medial prefrontal cortex: A functional magnetic resonance imaging study with human subjects. *Neurosci Lett* 335:183–186.