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Schizophrenia Subjects Show Intact Success-Related Neural Activation but Impaired Uncertainty Processing during Decision-Making

Martin P Paulus^{*,1,2,3}, Lawrence Frank^{3,4}, Gregory G Brown^{2,3} and David L Braff²

¹Laboratory of Biological Dynamics and Theoretical Medicine, University of California at San Diego, La Jolla, CA, USA; ²Department of Psychiatry, University of California at San Diego, La Jolla, CA, USA; ³Veterans Affairs San Diego Health Care System, USA; ⁴Department of Radiology, University of California at San Diego, La Jolla, CA, USA

Decision-making is a complex process that is important for everyday life. This study examined the effect of the *degree of success*, and outcome *uncertainty*, on decision-making and associated neural substrate activation in schizophrenia subjects (SZS) and normal comparison subjects (NCS). A total of 15 subjects with a diagnosis of schizophrenia and 15 age- and education-matched NCS participated in this study. These subjects completed the two-choice prediction task during functional magnetic resonance imaging. Decision-making characteristics and activation of neural substrates were obtained at 20, 50, or 80% error rate. Success and uncertainty influenced the behavioral characteristics on the two-choice prediction task, and the task-related activation in SZS and NCS. Neither success nor uncertainty differentially affected the behavioral characteristics of SZS relative to NCS during the two-choice prediction task. Nonetheless, there was a significant interaction between group and error rate in bilateral parietal cortex. The activation in NCS was the highest when the outcome was most uncertain. In contrast, task-related activation in SZS was not modulated by the degree of uncertainty. Thus, SZS failed to utilize the parietal cortex to process decision-making situations with highly uncertain outcomes. *Neuropsychopharmacology* (2003) **28**, 795–806. doi:10.1038/sj.npp.1300108

INTRODUCTION

Decision-making, that is, selecting an action from a number of alternatives when the outcome is uncertain, is a complex process that is important for everyday life (Tversky and Kahneman, 1981). Both cognitive and affective functions are involved in the decision-making process. Among these functions, two key factors influence the response selection during decision-making. First, the degree of success associated with an action strongly influences response selection when the actions are not explicitly related to a reward (Schultz et al, 2000). Second, uncertainty of an outcome, that is the degree to which the subject can predict whether the decision will be associated with success or failure, critically contributes to the ability to form an internal prediction model (Egelman et al, 1998). In the absence of explicit reward, and when the outcome of a decision is uncertain, subjects use the history of successes and failures associated with different response alternatives to form a model and guide their action in order to obtain the highest likelihood of success (Calfee and Atkinson, 1966; Goulet and Barclay, 1967; Ludvigson, 1966).

Decision-making can be dysfunctional in patients with different neuropsychiatric disorders, including schizophrenia (Mogg et al, 1991; Lyon et al, 1986; Garety et al, 1991; Brebion et al, 1997; American Psychiatric Association, 1994; Rahman et al, 1999). Although no single task can probe all aspects of decision-making, we have used the two-choice prediction task to examine the sequential organization of responses during decision-making, and the influence of history of uncertainty or success influence the selection of responses (Paulus et al, 1996, 1999, 2001b). During the twochoice prediction task, subjects are asked to predict repeatedly the location of a stimulus on a computer screen. Within an experimental session, individual schizophrenia patients generate response sequences that are both highly predictable and rigid or highly unpredictable (Paulus et al, 1996) and are not well predicted by the previous stimulus or the previous outcome (Paulus et al, 1999). In contrast, normal comparison subjects (NCS) generate sequences that are moderately unpredictable, and are often related to the previous outcome or the previous stimulus. In schizophrenia patients, as opposed to NCS, the current response can be predicted not only by the immediately preceding response but also by responses that were made many trials before,

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^{*}Correspondence: Dr MP Paulus, Department of Psychiatry, University of California at San Diego, La Jolla, CA 92093-0603, USA, Tel: +1 858 822 1473, Fax: +1 858 642 1429, E-mail: martin@mag.ucsd.edu

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which yields extensive temporal correlations (Paulus *et al*, 1999). These decision-making characteristics in schizophrenia patients are independent of psychopathology and are stable over time (Paulus *et al*, 2001b). In combination, these results support the hypothesis that decision-making dysfunctions in schizophrenia patients are because of abnormal processing of response sequences and altered processing of external stimuli. However, it is unclear whether these dysfunctions are because of changes in the association between response and outcome, or because of the inability to form accurately a prediction model.

Based on behavioral studies, decision-making has been separated into: assessment, that is, relating a stimulus to outcome probabilities; evaluation, that is, optimizing among competing responses; and executive processes, selecting or updating ongoing response strategies (Kahneman and Tversky, 1984). Not surprisingly, functional neuroimaging studies have shown that decision-making is critically dependent on a widely distributed neural network that includes the inferior prefrontal cortex (Paulus et al, 2001a; Ernst et al, 2002), ventromedial and ventrolateral frontal cortex (Elliott et al, 2000, 1999; Rogers et al, 1999), anterior cingulate (Elliott et al, 2000), insula (Critchley et al, 2001), and parietal cortex (Paulus et al, 2001a). Specifically, the inferior prefrontal areas have been linked to the detection and selection of advantageous over disadvantageous responses (Elliott et al, 2000; Rogers et al, 1999; O'Doherty et al, 2001). The dorsolateral prefrontal cortex has been linked to the processes of response selection during decision-making (Paulus et al, 2001a) and the anterior cingulate to the evaluative components, that is, the distinction between hypotheses testing and guessing (Elliott and Dolan, 1998) or reward-based response selection (Bush et al, 2002; Gehring and Willoughby, 2002). The parietal areas have been implicated in the assessment of functions, for example, anticipatory processing during decision-making (Critchley et al, 2001), relative to the utility of behavioral alternatives (Platt and Glimcher, 1999; Shadlen and Newsome, 2001).

Our previous studies of error-rate-related decisionmaking in normal controls showed that inferior prefrontal areas show more activation at low error rates, that is, when response selection was associated with a high success rate, and posterior parietal activation was linked to outcomerelated responses (Paulus *et al*, 2002a). Therefore, if alterations in processing during decision-making in schizophrenia subjects (SZS) were because of changes in successrelated response selection, one would predict that there would be differences in the inferior prefrontal cortical activation. Alternatively, if an alteration in uncertainty processing underlies decision-making dysfunctions in schizophrenia patients, one would predict to see differences predominantly in the posterior parietal cortex.

Functional neuroimaging studies with SZS have shown dysfunctions of the neural substrates critical for decisionmaking, that is, prefrontal cortex, anterior cingulate, and parietal cortex. Specifically, working-memory studies during functional magnetic resonance imaging (fMRI) have shown that both chronic and first-episode never-medicated schizophrenia patients showed deficits in working memory performance associated with an increased or decreased activation (Callicott *et al*, 1998; Barch *et al*, 2001), or an increased spatial heterogeneity of dorsolateral prefrontal cortex activation (Manoach et al, 2000). The dorsolateral prefrontal cortex dysfunction in SZS during a working memory task has been associated with disorganization symptoms (Perlstein et al, 2001). Moreover, anterior cingulate dysfunction, related to the evaluation or detection of response conflict during the Stroop task, has been observed in both PET (Nordahl et al, 2001) and fMRI studies (Carter et al, 2001). Finally, both reduced (Artiges et al, 2000; Fletcher et al, 1998) and increased posterior parietal activation (Callicott et al, 1998; Paulus et al, 2002b) have been observed in response to different tasks in SZS. The heterogeneity of these findings has led some investigators to propose that schizophrenia patients show a disruption of processing across different areas in the brain, rather than a circumscribed abnormality in one area (Friston, 1998). This hypothesis has been supported by some studies (Volz et al, 1999; Meyer-Lindenberg et al, 2001; Lawrie et al, 2002) but not others (Spence et al, 2000; Curtis et al, 1999; Fletcher et al, 1998).

This study examined the effect of the *degree of success* and the uncertainty on decision-making and associated neural substrate activation in SZS, relative to NCS. Based on the response of the subject during the two-choice prediction task, a computer program determined a priori whether the prediction would be 'correct' or 'incorrect'. Specifically, decision-making characteristics and activation of neural substrates were obtained at 20% error rate (2/10 correct predictions), 50% error rate, or 80% error rate. This design enables one to examine the effect of success, that is, the number of correct vs incorrect predictions (high success -20% error rate, low success 80% error rate), and of uncertainty, that is, the degree to which a response predicts an outcome irrespective of success or failure (high unpredictability -50% error rate, low unpredictability -20 and 80% error rate), on decision-making. It was hypothesized that if decision-making dysfunctions in SZS are because of altered processing of failure vs success, neural substrate differences should be most pronounced at high or low error rate, respectively, and should be related to differences in inferior prefrontal cortex processing. Alternatively, if the decision-making dysfunctions are because of altered processing of uncertainty, neural substrate differences should be most pronounced when the outcome is most unpredictable (ie at 50% error rate) and should be related to differences in posterior parietal cortex.

METHODS

This study was approved by the UCSD Human Research Protections Program (000730) and all subjects signed informed consent. Initially, 17 SZS and 16 NCS were enrolled in the study. Two SZS and one NCS were excluded from subsequent analyses because of movement artifact in the echoplanar images. Included were 15 right-handed subjects with the diagnosis of schizophrenia (four females, 11 males), continuous, according to the DSM-IV (1994) with an average age of 41.7 ± 1.6 years (range 30-53), an average education level of 14.4 ± 0.7 years (range 12-23), an average age of onset of 25.9 ± 1.8 years (range 18-39), and an average illness duration of 15.7 ± 2.1 years (range 7-34).

The behavioral and functional neuroimaging data from these subjects were compared to a group of 15 right-handed NCS (four females, 11 males) that were matched on age (mean 41.0 ± 2.1 , range 21-54) and education (mean 15.3 ± 0.56 , range 12-21). All subjects were trained to perform the two-choice prediction and two-choice response task prior to testing during fMRI scanning.

Diagnoses for all subjects were obtained by using a structured clinical interview for DSM-IV diagnosis (SCID-P) (Spitzer *et al*, 1992). Subjects with a major depressive disorder, bipolar, post-traumatic stress, panic, or obsessive-compulsive disorder were excluded from the study. Subjects with nonremovable materials that respond to high magnetic fields, for example, metal fragments, were also excluded. At the time of testing, the SZS were clinically stable, nine subjects were treated with atypical antipsychotic medication, three subjects were treated with typical antipsychotic medications, and three subjects were not treated with any antipsychotic medications at the time of testing.

Task

The two-choice prediction task has been described in detail elsewhere (Paulus, 1997). Briefly, a house flanked by a person to the left and right is shown on a computer screen. The goal for the subject is to match a person on the computer screen with a car that is presented on the far left or right side of the screen. For the two-choice prediction task, the subject's goal is to predict which side the car will be presented and to select the left or right button, respectively. After the subject has made a response, the car is presented for 300 ms on the far left or right side. If the selected response matches the side where the car is presented, the person meets up with the car. For the twochoice response task, the car is presented on the left or right side before the subject is asked to respond, and there is no prediction.

Unbeknown to the subject, on the two-choice prediction task the car is presented according to a predetermined schedule. Specifically, the computer program takes the response of the subject into account, and determines whether a response will be 'correct' or 'incorrect'. The key difference between these two tasks is that during the twochoice prediction task, the subject does not know the correct response in advance, and has to decide in the presence of uncertainty, using the previous responses, stimuli, and outcomes to determine their response. In comparison, during the two-choice response task the subject knows the correct answer before selecting a response, decides in the presence of certainty, and does not need to use the sequences of previous responses, stimuli, or outcomes. The task was presented to the subjects in the MRI using an LCD projector, back-projected onto a screen at the subjects' feet, which could be seen via a mirror attached to the head coil. Subjects requiring corrective lenses were provided with a pair of plastic-framed lenses that approximated their degree of correction. Motor responses were made by the right hand using a button box.

A block design was used for this study. The two-choice prediction task was divided into three trial-blocks, each lasting 90 s. During the first trial-block the computer program assured that 50% of all responses were 'correct',



during the second trial-block 20% of all responses 'correctly predicted' the location of the car, and during the third trialblock 80% of all responses were 'correct' predictions. Therefore, the first trial-block corresponds to 'evenly right or wrong', the second trial-block to being 'mostly wrong', and the third trial-block to being 'mostly right'. The twochoice prediction task was contrasted with the two-choice response task, which was presented between the 50 and 20%, as well as between the 20 and 80% trial blocks for 30 s, to examine task-related activation. The duration of each trial depended on the latency to make a decision, that is, the time between presentation of the initial situation and the selection of the response.

Behavioral Measures

For both the two-choice prediction task and the two-choice response task, the following variables were recorded: (1) the choice selected by the subject (left or right), (2) the response selected by the computer (*left* or *right*), and (3) the subjects' latency of response selection (time from the presentation of the current situation to the selection of the response). Based on these variables, the strategies of decision-making in the presence of uncertainty were assessed using the following three sets of measures: (1) General response biases: the number of left or right responses or stay responses (a left response followed by left response), vs switch (left followed by right response) responses. (2) The degree to which the current response was determined by the previous response, the previous stimulus, or a combination of both, quantified by mutual information measures (Herzel and Grosse, 1995). The mutual information between two events, A and B, expressed in units of bits, is the logarithmic likelihood ratio of the observed occurrence rate over the expected chance occurrence rate of A and B. The win-stay/lose-shift mutual information is computed by coding event A as a win (correct) or lose (incorrect) outcome at trial i-1 and B as the same (stay) or different (shift) response at trial i. (3) The predictability of the response sequence measured by the average entropy and the range of subsequence fluctuations between highly predictable (minimum entropy sequence) and highly unpredictable response sequences (maximum entropy sequence), which operationally defines the degree of response dysregulation during the two-choice prediction task. The dynamical entropy is computed via the determination of unique subsequences of responses during the twochoice prediction task as detailed in Grassberger (1989) and Paulus (1997).

fMRI Protocol and Image Analysis Pathway

Magnetic resonance images were obtained using a 1.5 T whole-body system (Siemens, Erlangen). Anatomical T1-weighted images of the whole brain (MPRAGE, TR = 11.4 ms, TE = 4.4 ms, flip angle = 10°, FOV = 256 × 256, 1 mm³ voxels) were obtained sagitally to identify the anterior/posterior commissure, to coregister the functional image, and to transform the images into Talairach space (Talairach and Tournoux, 1988). A total of 32 slices of T2*-weighted images were obtained in the axial plane using gradient-recalled echo planar imaging (TE = 40 ms, flip angle = 90°, 64×64 pixel FOV = 220 × 220 mm, 3 mm contiguous slice

thickness) every 3000 ms for 112 repetitions, yielding a voxel size of $3.43 \text{ mm} \times 3.43 \text{ mm} \times 3 \text{ mm}$ in order to minimize signal dropout related to magnetic susceptibility variations in the orbitofrontal cortex.

All structural and functional image processing was done using the analysis of functional neuroimages (AFNI) software package (Cox, 1996). Echoplanar images were coregistered using a 3D coregistration algorithm to the echoplanar image that resulted in the smallest amount of image translation and rotation, relative to all other images. The main dependent measure to assess task-related brain activation was the percent signal change during the twochoice prediction task, relative to the two-choice response task across three delay times. Multiple regression analysis was used to quantify the fMRI time series data (Courtney et al, 1997). Three different reference functions were used to measure the degree of echoplanar signal change during the 50, 20, and 80% reinforcement conditions, respectively. The AFNI program 3dDeconvolve was used to calculate the estimated impulse response function between the reference function and the echoplanar time series using a time shift of 1-3 TR (ie 3-9 s) at each voxel. The relative signal change was computed by dividing the regressor coefficients for each time shift by the zeroth-order regressor coefficient, which measures the average signal intensity during the twochoice response task trial-blocks. A Gaussian filter with FWHM 6 mm was applied to voxelwise percent signal change data to account for individual variations of the anatomical landmarks and to yield the highest detection power (Skudlarski et al, 1999). The data of each subject were normalized to Talairach coordinates, and the measure of relative change of signal intensity during a task of interest (two-choice prediction task during different reinforcement conditions), relative to the baseline task (two-choice response task), was entered into a mixed model nested ANOVA. Specifically, task condition (20, 50, 80%) was used as a fixed factor, subjects as a random factor, which together were nested under the group as a fixed factor (SZS, NČS). A threshold adjustment method, based on Monte-Carlo simulations, was used to guard against identifying false positive areas of activation (Forman et al, 1995). Based on these simulations, it was determined that a voxelwise a priori probability of 0.01 would result in a corrected clusterwise activation probability of 0.05 if a minimum volume of $512 \,\mu l$ (or eight contiguous $4 \,mm^3$ voxels) and a connectivity radius of 4 mm was considered. All graphical results are presented as volume-threshold % signal difference or F-maps. Labels for brain activation foci were obtained in Talairach coordinates using the Talairach Demon software (Lancaster et al, 2000).

Statistical Analysis

All analyses for the behavioral data were carried out with SPSS 10.0 (Norusis, 1990). A mixed model ANOVA (fixed factor: task conditions; random factor: subjects) was used to analyze the behavioral measures. The planned comparisons were evaluated using the least significant difference (LSD) *post hoc* analysis. In order to adjust the degrees of freedom for the correlations in within-subject designs (violations of sphericity), Greenhouse–Geisser (GG) corrections were applied. All analyses were conducted using a response

number as a covariate; however, the results did not differ significantly, and the results presented here are based on the mixed model described above.

BEHAVIORAL RESULTS

Error rate significantly affected basic response characteristics during the two-choice prediction task in both groups (Table 2). The error rate effects on response biases, mutual information functions, and the dynamical entropy, however, were similar for both SZS and NCS (Table 2). Specifically, the latency to select a response and the degree to which the current response predicted the next response (mutual information), in both groups, was lowest when the outcome was most unpredictable (Tables 1 and 2). Thus, the degree of response predictability matches the uncertainty of the outcome. In comparison, the probability of switching from the current response to the alternative response was proportional to the error rate for both SZS and NCS (Table 1). Both SZS- and NCS-generated response sequences were most predictable as measured by the average entropy when

Table IAverage Behavioral Measures and SEM for DifferentError Rate Conditions and the Two-Choice Response Task forNormal Comparison Subjects and Schizophrenia Subjects

	Nor compariso	mal n subjects	Schizophrenia subjects	
Measures	Mean	SEM	Mean	SEM
20% error rate				
Response latency (ms) Probability of RIGHT response	750 0.5 I	85 0.03	809 0.53	85 0.03
Probability of	0.38	0.03	0.38	0.04
Mutual information Win-stay mutual information	0.08 0.09	0.03 0.02	0.11 0.19	0.02 0.05
50% error rate				
Response latency (ms) Probability of RIGHT response	904 0.46	4 0.02	978 0.50	4 0.02
Probability of SWITCH response	0.47	0.03	0.44	0.03
Mutual information Win-stay mutual information	0.03 0.11	0.01 0.04	0.04 0.17	0.02 0.04
80% error rate				
Response latency (ms) Probability of	762 0.48	88 0.01	871 0.53	88 0.02
Probability of SWITCH response	0.55	0.03	0.57	0.04
Mutual information Win-stay mutual information	0.05 0.07	0.01 0.02	0.09 0.14	0.02 0.03
Response task				
Response latency (ms) Response errors (% errors)	676 4	72 3	810 17	80 3

 Table 2
 Statistical Analysis of the Behavioral Results for the Two-Choice Prediction Task

	Group		Error rate		Group by error rate	
Measure	F	Þ	F	Þ	F	Þ
Number of responses per block	0.37	NS	5.27	< 0.01	0.454	NS
Probability of RIGHT response	2.0	NS	2.55	< 0.1	0.384	NS
Probability of SWITCH response	0.0	NS	19.72	< 0.01	0.492	NS
Mutual information	2.79	NS	5.08	< 0.05	0.252	NS
Win-stay mutual information	3.12	< 0.1	1.25	NS	0.25	NS
Average entropy	1.89	NS	6.28	< 0.01	0.465	NS
Dysregulation	3.89	< 0.05	2.07	NS	0.74	NS

All F Values are Greenhouse-Geisser Corrected where Appropriate

Dynamical Entropy



Figure I Average and standard error of mean (SEM) of error-rate dependent changes in behavioral measures for the two-choice prediction task in NCS and SZS.

the error rate was low (Figure 1). The dysregulation of response sequences, as measured by the range of response sequence predictabilities, differed significantly across groups (Table 1), but not across error-rate conditions. As shown in Figure 1, SZS showed a slightly but significantly higher degree of dysregulation, when averaged across the different error-rate conditions. Although the latency to select a response during the two-choice response task did not differ between SZS and NCS (Table 1, F(1,28) = 1.7, NS), SZS made significantly more errors than NCS during the two-choice response task (Table 1, F(1,28) = 11.3, p < 0.01).

FMRI RESULTS

The main effect of the task, that is, the difference between two-choice prediction and two-choice response task, as evaluated separately for NCS and SZS, revealed a distributed activation across inferior prefrontal, anterior cingulate, and posterior parietal areas in NCS (Figure 2, top) and across inferior prefrontal and anterior cingulate but not posterior parietal areas in SZS (Figure 2, bottom). When the groups were combined, different error-rate conditions significantly affected distributed areas of the brain including the inferior prefrontal, anterior cingulate, and posterior parietal cortex (Table 3). First, prefrontal areas included bilateral inferior frontal gyrus (BA 44/45) extending to the middle frontal gyrus (BA 9/46) and medially to the medial prefrontal cortex (BA 10/11). Second, both dorsal and rostral anterior cingulate (BA 32) showed error-rate- related activation differences across tasks. Third, several posterior parietal areas, including bilateral precuneus (BA 7/18) and left inferior parietal lobule (BA 40), showed error-rate-related activation changes. When the task \times error-rate analyses was conducted separately for NCS and SZS, significant errorrate-related changes were found in both the inferior prefrontal cortex and the rostral anterior cingulate, but not in the posterior parietal cortex. The average activation across these functional regions of interest showed a Ushaped relation to the error rate (Figure 4). That is, for both NCS and SZS these regions were most active when the subjects were most successful in predicting the outcome, and were least activated when the outcome was most unpredictable.

Several closely related areas revealed a significant task \times group \times error-rate interaction (Figure 3, Table 2). Bilateral precuneus (BA 7) and superior parietal lobule (BA 7) showed different error-rate-related activation in comparison subjects relative to SZS. As shown in Figure 3, whereas most NCS showed a larger task-related activation at 50% error rate, most SZS did not show larger activation at 50% error rates in the precuneus and superior parietal regions.

Normal Comparison Subjects: Two-choice Prediction -Two choice Response Task 10 28 22 34 46 40 52 0.6 0.5 04 0.3 Schizophrenia Subjects: Two-choice Prediction - Two 0.2 choice Response Task 10 16 0.1 0.0 22 28 34 46 40 52

Figure 2 Percent signal difference between two-choice prediction and response task for NCS (top) and schizophrenia subject (bottom); numbers indicate *z* Talairach coordinate.

In order to examine whether the interaction between task and error rate across groups was because of uncertaintyrelated processing in NCS but not in SZS, a contrast vector (activation at 50% error rate $-\frac{1}{2}$ (activation at 20%+activation at 80% error rate)) was created and tested separately for NCS and SZS. As shown in Figure 3, NCS exhibited a significant contrast effect in the posterior parietal cortex (precuneus, BA 7); however, no such area was observed for SZS (figure not shown). Moreover, the average activation from these functional regions of interest showed a significant $contrast \times group$ effect (F(1,28) = 11.45,p < 0.01). A similar contrast analysis with contrasting high success rate with high error rate (20% error rate-80% error rate) did not yield significant differences between NCS and SZS (F(1,28) = 3.27, NS).

To determine whether the difference in the precuneus area was because of the fact that SZS made more errors

during the two-choice response task, an ANCOVA, with response errors as covariate, was conducted between NCS and SZS. The corrected model was highly significant (F(3,27) = 7.9, p < 0.01) and there was a main effect of group (F(1,27) = 5.8, p < 0.05), but not response error (F(1,27) = 3.3, NS), on the contrast vector in the precuneus (BA 7). Whereas the posterior parietal cortex of NCS was relatively more active when the outcome of the decision was most uncertain, no such modulation was found in SZS. Moreover, the lack of modulation was not related to differences in errors made during the two-choice prediction task.

As shown in Figure 3, most SZS showed no change or reduced activation during the 50% error-rate condition. The degree to which this pattern was expressed correlated with duration of illness (r=0.67, p<0.01), but not age (r=0.36, NS) or education (-0.46, NS), in SZS (Figure 3). Thus, the difference in processing uncertainty in the posterior parietal cortex between NCS and SZS increased with increasing duration of illness. There were no significant correlations between the contrast vector in this area and behavioral contrasts (ie difference in response biases, mutual information functions, and dynamical entropy at 50% error rate and 20 or 80% error rate interaction areas.

DISCUSSION

SZS relative to NCS show altered processing of outcome uncertainty, but do not show altered processing of response success or failure. Specifically, there was a significant interaction between group and error rate in several areas of the posterior parietal cortex comprising the precuneus (BA7), bilateral superior parietal lobule (BA 40), and right inferior parietal lobule (BA 40). In each of these areas, NCS, but not SZS, showed the largest activation at 50% error rate and less activation at 20 or 80% error rate. Finally, the degree to which SZS lacked an increase in uncertaintyrelated activation in the precuneus was correlated with illness duration. These results extend previous reports of reduced posterior parietal activation during the two-choice prediction task in SZS (Paulus et al, 2002b) and support the hypothesis of an altered processing of outcome uncertainty during decision-making in SZS that involves the posterior parietal cortex.

In comparison, the behavioral results of this study do not support the hypothesis that the degree of success or the outcome uncertainty differentially affects response selection during decision-making in SZS relative to NCS. As for response switching, the degree to which previous responses or stimuli predicted the current response; the average response entropy and the degree of dysregulation did not reveal an interaction between error rate and group. Thus, both NCS and SZS responded similarly to different error rates.

Success-related changes of activation in neural substrates were similar for both SZS and NCS. At low and high error rates, predominant response strategies develop, for example, lose-shift resulting in repetitive response switching or win-stay in long response repetition. In contrast, when the outcome is highly uncertain, subjects are required to adjust

Volume	x	у	Z	L/R	ВА	Description
Task × error-rate	effect					
2889	-27	-66	38	L	7	Precuneus
43	20	-53	39	R	7	Precuneus
648	10	-43	49	R	7	Precuneus
1350	I	-75	9	R	18	Cuneus
513	-2	-92	15	L	18	Cuneus
5562	-44	-42	40	L	40	Inferior parietal lobule
1161	-55	-28	37	L	40	Inferior parietal lobule
3618	-10	21	-10	L	25	Anterior cingulate
837	51	11	40	R	8/9	Middle frontal gyrus
513	-46	14	44	L	8/9	Middle frontal gyrus
23058	-56	16	3	L	45	Inferior frontal gyrus
9774	57	28	2	R	45	Inferior frontal gyrus
756	25	-36	47	R	3	Postcentral gyrus
702	23	-37	67	R	5	Postcentral gyrus
1863	-19	-17	15	L		Thalamus
513	54	-35	-9	R	20	Middle temporal gyrus
Task × group × e	error-rate effect					
837	-13	-89	36	L	19	Cuneus
1404	0	-48	46	L/R	7	Precuneus
621	5	-51	62	R	7	Precuneus
621	29	-70	58	R	7	Superior parietal lobule
540	-15	-53	61	L	7	Superior parietal lobule
594	-31	-70	37	L	19	Precuneus
972	36	-56	44	R	40	Inferior parietal lobule
864	44	0	-17	R	38	Superior temporal gyrus

Table 3Center of Mass for Volume-Thresholded Mixed ANOVA Clusters of Task-Related (Two-ChoicePrediction—Two-Choice Response Task)Activation for the Group Effect, the Reinforcement Effects Common toBoth Groups and the Group by Reinforcement Interaction, Respectively

Labels are Based on Talairach Demon Software (Lancaster et al, 2000).

strategies. The lack of the inverted U-shaped relation between error rate and task-related activation in posterior parietal cortex in SZS may reflect an altered processing of outcome uncertainty as previously observed (Paulus *et al*, 2002b). The key aspect of high uncertainty, relative to low or high success rate, is the inability to establish repetitive response strategies, for example, repeated win-stay or loseshift responses, as evidenced by the decrease in response predictability. The repeated adjustment of response selection, because of highly unpredictable success or failure, appears to engage posterior parietal in NCS but not in SZS. Thus, the current results are consistent with the general hypothesis that there is a disturbance in the processing of decision-making in SZS when there are no dominant response strategies established.

Both structural (Weinberger *et al*, 1992; Seidman *et al*, 1994; Gur *et al*, 2000) and functional (Weinberger *et al*, 1996; Callicott *et al*, 1998; Manoach *et al*, 1999; Rubia *et al*, 2001a; Volz *et al*, 2001) imaging studies, using a number of different neurocognitive tasks, in SZS have implicated primarily, and some selectively (Barch *et al*, 2001), dorsolateral prefrontal cortex dysfunction in SZS. Others still have correlated evaluative dysfunctions in schizophrenia with impaired performance monitoring in the anterior cingulate (Carter *et al*, 2001, 2000, 1997; Nordahl *et al*, 2001). However, there is increasing evidence of both structural (as reviewed in Shenton *et al*, 2001) and functional parietal cortex dysfunction in SZS. Moreover, blood flow changes in the parietal cortex have been reported

to increase with duration of illness in SZS (Schultz et al, 2002).

The posterior parietal cortex has been implicated in a number of processes important for decision-making. Specifically, attentional processes that involve sustained, and possibly selective, attention (Coull et al, 1996), switching from task-relevant local to global targets (Fink et al, 1996; Lamb et al, 1989), voluntary attentional control (Hopfinger et al, 2000), and the distinction between taskirrelevant and task-relevant events (Downar et al, 2001; Kiehl et al, 2001; McCarthy et al, 1997) support the view that this area is critical for the extraction and selection of task-relevant information. Moreover, this area has been implicated in inhibitory control in a number of different paradigms (Garavan et al, 1999; Menon et al, 2001; Rubia et al, 2001b; Steel et al, 2001; Doricchi et al, 1997), that is, the allocation of resources to a response that has to compete with a highly overlearned and potentially habitual behavior. Several studies using decision-making paradigms have implicated the right posterior parietal cortex in autonomic arousal processes (Tranel and Damasio, 1994; Critchley et al, 2000), risk-taking decision-making (Ernst et al, 2002), and guessing (Elliott et al, 1999).

Dissociation between frontoparietal processing has been reported in other neuroimaging studies with SZS. For example, whereas SZS showed performance-related activation deficits during a working-memory task in the dorsolateral prefrontal cortex, these subjects also showed a working-memory load-independent reduction in the



Figure 3 Task × group × error rate interaction (upper left) and contrast effect (upper right) in NCS. The contrast was inversely correlated with illness durationin SZS (lower right). Activation patterns in individual subjects show an uncertainty-related pattern in NCS (bottom left) but not SZS (bottom right).

parietal cortex (Fletcher *et al*, 1998). Moreover, workingmemory-related activation in the posterior parietal cortex has been found to be correlated with somatic hallucinations (Shergill *et al*, 2001; Lennox *et al*, 2000). Others have suggested that parietal cortex dysfunction in SZS may contribute to the difference in semantic fluency vis-à-vis phonologic fluency (Feinstein *et al*, 1998). SZS showed significantly less activation in posterior parietal areas during a randomization task, which are though to contribute to the perseverative tendencies in these subjects (Artiges *et al*, 2000). Electrophysiological studies have shown reduced parietal P300 amplitude, which has been interpreted to signify a dysfunction in the continuous memory updating of current events (Nieman *et al*, 2002).

The lack of uncertainty-related activation in posterior parietal areas in SZS would suggest that SZS engage less processing resources than NCS. How is this compatible with the previously reported increase in outcome-related strategy (Paulus *et al*, 2002b)? The win-stay/lose-shift strategy consists of two steps. First, the subject needs to remember where the stimulus was presented during the previous trial. Second, the subject needs to determine whether the





Task By Error-Rate Effect



Figure 4 Task-related activation averaged across all functional regions of interest that showed a significant effect of error rate (Table 1). Significant increase in activation at low error rates was common in both SZS and NCS.

prediction was correct or incorrect. If the current choice was simply based on the presentation of the previous stimulus, irrespective of the previous correct or incorrect prediction, one would expect increased win-stay/lose-shift consistent responses across all error-rate conditions. The current behavioral results are consistent with this view, showing that SZS tend to select the response based on the location of the previous stimulus and irrespective of whether the prediction was 'correct' or 'incorrect'. In NCS, increased activation in the posterior parietal cortex was associated with a decreased frequency of the dominant response strategy (win-stay/lose-shift) (Paulus et al, 2001a). Processing of prospective gains during decision-making is most difficult when the outcome of the decision is most uncertain and the outcomes of previous responses are most variable at the 50% error-rate condition. Thus, outcome uncertainty and variable history of 'correct' and 'incorrect' responses may force NCS to increase assessment processing during decision-making using the posterior parietal cortex, a process that is missing in SZS.

SZS, relative to NCS, adjusted their response characteristics similarly for different error rates. Specifically, both groups changed the switching rate, proportional to the error rate, and exhibited the lowest response predictability when the outcome associated with a response was most uncertain (ie at 50% error rate). Similarly, both groups showed a network of neural substrates that showed error-rate-related changes in activation. Bilateral inferior frontal gyrus (BA 44/45) and middle frontal gyrus (BA 9/46), ventromedial prefrontal cortex (BA 10/11), and anterior cingulate (BA 32) showed error-rate-related activation changes. These areas have been implicated in spatial attentional processes (Mesulam, 1999), conflict and error monitoring (Carter *et* *al*, 2000; Botvinick *et al*, 1999), inhibitory processes (Menon *et al*, 2001; Liddle *et al*, 2001; Rubia *et al*, 2001b), and control of eye movements (Luna *et al*, 1998). Thus, both groups allocate similar error-rate-related processing resources during this decision-making task. This finding is consistent with the observation that SZS respond similar to NCS concerning reinforcement contingencies (for a review, see Stieper *et al*, 1972).

There are several caveats to the conclusions and limitations of the current study. First, the functional imaging differences between NCS and SZS were not corroborated by behavioral differences that correlated with activation differences in the posterior parietal cortex. The divergence between behavioral measures and functional imaging results may signify that the current measures do not adequately quantify the degree of assessment prior to selecting a response. Event-related fMRI designs or electrophysiological methods that provide high temporal resolution may need to be employed to clarify the relation between behavioral differences and brain processes during the assessment phase of decision-making. Second, the degree of dysregulation in SZS, although significant across error rates, was small and did not reach significance for 50% error rates. It has been pointed out by others (Carter et al, 2001) that fMRI studies may select higher-functioning SZS and, thus, may create a selection bias that reduces performance differences between these groups (Resnick, 1992). Third, the presentation of the error-rate conditions during the fMRI experiment does not control for order effects. Although both behavioral and neuroimaging effects showed order-unrelated changes, future designs may need to employ longer functional sessions with multiple errorrate conditions. Fourth, symptom status was not evaluated in close proximity to testing. Thus, it is unclear if the relation between (1) the lack of uncertainty-related activation in the posterior parietal cortex and (2) illness duration is mediated by the severity of illness at the time of testing. Fifth, SZS committed significantly more errors during the two-choice response task. Although the effect of uncertainty was still significant when the number of errors was entered into the analysis as a covariate, there is still a possibility that activation differences between the two-choice prediction task and the two-choice response task may relate to the number of errors during the two-choice response task.

In conclusion, SZS and NCS show similar task-related activation in response outcome success, which encompasses a distributed network comprising inferior prefrontal, posterior parietal, and cingulate cortex. Whereas NCS showed uncertainty-related activation in the posterior parietal cortex, SZS did not show error-rate-related differences in these areas. Thus, inadequate processing of uncertainty in the posterior parietal cortex may be key substrates in the error-rate-related decision-making dysfunctions in schizophrenia.

ABBREVIATIONS

NCS, normal comparison subjects; SZS, schizophrenia subjects; BA, Brodmann area; fMRI, functional magnetic resonance imaging; DSM, Diagnostic and Statistical Manual of Mental Disorders; SCID, Structured Clinical Interview for DSM-IV diagnoses; MPRAGE, magnetization-prepared rapid acquisition of gradient echo; FOV, field of view; AFNI, analysis of functional neuroimages; FWHM, full width half maximum; ANOVA, analysis of variance; ANCOVA, analysis of covariance.

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806