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Punishment and psychopathy: a case-control functional MRI investigation of reinforcement learning in violent antisocial personality disordered men



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Summary

Background Men with antisocial personality disorder show lifelong abnormalities in adaptive decision making guided by the weighing up of reward and punishment information. Among men with antisocial personality disorder, modification of the behaviour of those with additional diagnoses of psychopathy seems particularly resistant to punishment.

Methods We did a case-control functional MRI (fMRI) study in 50 men, of whom 12 were violent offenders with antisocial personality disorder and psychopathy, 20 were violent offenders with antisocial personality disorder but not psychopathy, and 18 were healthy non-offenders. We used fMRI to measure brain activation associated with the representation of punishment or reward information during an event-related probabilistic response-reversal task, assessed with standard general linear-model-based analysis.

Findings Offenders with antisocial personality disorder and psychopathy displayed discrete regions of increased activation in the posterior cingulate cortex and anterior insula in response to punished errors during the task reversal phase, and decreased activation to all correct rewarded responses in the superior temporal cortex. This finding was in contrast to results for offenders without psychopathy and healthy non-offenders.

Interpretation Punishment prediction error signalling in offenders with antisocial personality disorder and psychopathy was highly atypical. This finding challenges the widely held view that such men are simply characterised by diminished neural sensitivity to punishment. Instead, this finding indicates altered organisation of the information-processing system responsible for reinforcement learning and appropriate decision making. This difference between violent offenders with antisocial personality disorder with and without psychopathy has implications for the causes of these disorders and for treatment approaches.

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Introduction

Most violent crimes are committed by a small number of men.¹ They display a pattern of antisocial and aggressive behaviour that begins in childhood and remains stable throughout the lifespan. They meet diagnostic criteria for conduct disorder in childhood and for antisocial personality disorder in adulthood. Lifelong patterns of poor decision making, impulsivity, and risk-taking behaviours characterise such men, and persist despite repeated punishments enacted by parents, teachers, and the criminal justice system.

Within this population, a subgroup presents with antisocial personality disorder and psychopathy, defined by the psychopathy checklist-revised (PCL-R)² as including callousness, lack of empathy, an interpersonal style involving grandiosity and manipulation of others, and persistent reactive and instrumental aggression. Such individuals show reduced tonic skin conductance and cortisol concentrations, and have difficulty recognising fear and sadness in the faces of others.³ By contrast, men who have antisocial personality disorder without psychopathy are characterised by emotional lability, mood and

anxiety disorders, and reactive aggression.⁴ Although the two phenotypes emerge early in life, those who develop antisocial personality disorder with psychopathy begin offending at an earlier age, engage in a broader range and higher frequency of offending behaviours,⁴ and respond less well to treatment in childhood⁵ and adulthood⁶ than those without psychopathy. The two groups show distinct differences in brain structure⁷ and functional responses to empathy-eliciting scenarios⁸ and emotional stimuli when engaged in goal-directed behaviour.⁹ Diagnostic classification systems, however, do not distinguish between antisocial personality disorder with and without psychopathy. Rehabilitation programmes typically exclude offenders with psychopathy.

Many decisions, including whether or not to engage in an antisocial act, involve the ability to assess consequences. Selecting an action is dependent on reinforcement learning, whereby possible rewards are weighed against possible punishments on the basis of past experience.¹⁰ Men with antisocial personality disorder show impairments in adaptive decision making, characterised by abnormal processing of reinforcement information.^{11,12} Their

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behaviour seems to be driven more by potential rewards than potential punishments (reward dominance). Impaired learning about the consequences of actions might result in overly optimistic and inaccurate predictions of favourable outcomes, resulting in frustration, which could predispose to reactive aggression. Impaired self-control lowers the threshold for aggression in response to such frustration.¹³ In behavioural studies offenders with antisocial personality disorder and psychopathy have shown notable impairment in using reinforcement information when choosing between punished and rewarded objects in passive avoidance,¹⁴ extinction,¹⁵ and reversal learning tasks.^{11,16,17} The failure to use punishment information to signal inappropriate behaviour is thought to be the primary deficit in psychopathy, and emerges early in childhood.¹⁸

Reversal learning tasks examine the ability to adjust behaviour to changes in reinforcement contingency. Studies of healthy adults and adults with brain lesions have identified a neural network recruited during reversal learning.¹⁹ The dorsomedial prefrontal cortex, ventrolateral prefrontal cortex, and dorsolateral prefrontal cortex serve, respectively, to allocate attentional resources, assess conflicting responses, and initiate response modulation. The ventromedial prefrontal cortex, posterior cingulate cortex, and dorsal caudate track changes in reinforcement information, reducing activation in response to punishment, thus signalling behavioural error, change in reinforcement contingency (when reward is expected but punishment received), and the need to adapt behaviour.^{20,21}

Little is known about the neural systems serving reinforcement learning anomalies in offenders with antisocial personality disorder with and without psychopathy. Offline versions of probabilistic response reversal tasks are self-paced, which means that intervals between the stimulus presentation, response, and feedback vary across trials.¹¹ A variant of this task has been developed for the functional MRI (fMRI) scanning environment that is designed specifically to minimise behavioural differences between antisocial individuals and normal controls. Importantly, the scanner version of the task provides reinforcement more often and more consistently than the offline task. This change lessens task difficulty and the likelihood of confounding introduced by large differences in performance, such as differential contamination by error events.²² The scanner version of the task has been used to assess adolescents with conduct disorder and callous unemotional traits,²¹ which are the childhood antecedents of antisocial personality disorder with psychopathy.⁵ Compared with healthy teenagers, adolescents who have conduct disorder with callous unemotional traits showed intact recruitment of the dorsomedial prefrontal cortex, dorsolateral prefrontal cortex, and inferior frontal cortices when changing their behaviour in response to punishment signals, but did not appropriately reduce activation of the ventromedial prefrontal cortex. This finding was interpreted as intact recruitment of regions involved in response change immediately after

punishment, but impairment in regions that alter long-term value associated with responses, thereby increasing the likelihood of future incorrect responding.

We used the scanning environment probabilistic response reversal task to do an fMRI study of reinforcement processing and decision making in violent adult offenders with antisocial personality disorder with and without psychopathy. Focusing on responses to punished reversal errors relative to rewarded correct responses, we tested the hypothesis that violent offenders with antisocial personality disorder and psychopathy would show increased activation within the ventromedial prefrontal cortex, caudate, and posterior cingulate cortex to punished reversal errors, as identification of anomalous responding to reinforcement information within such individuals could be useful as a diagnostic biomarker.

Materials and methods

Participants and study design

Between January, 2007, and January, 2011, we enrolled 50 men, aged 20–50 years, with reading age higher than 10 years as defined by the Schonell instrument. Eligible participants had no history of major mental disorders (bipolar 1, bipolar 2, major depression, or psychotic disorders), as defined by the structured clinical interview for Diagnostic and Statistical Manual of Mental Disorder, fourth edition (DSM-IV), axis I disorders (SCID I), or self-reported neurological disorders, head injury resulting in loss of consciousness for 1 h or longer, severe visual or hearing impairments, or contraindications to MRI. Offenders with convictions for violent crimes (murder, rape, attempted murder, and grievous bodily harm) who met DSM-IV criteria for antisocial personality disorder were recruited via the National Probation Service of England and Wales. Healthy non-offenders were recruited via community websites and from unemployment offices. All participants completed diagnostic and PCL-R interviews and authorised access to their criminal records. A cross-cultural validation study²³ of the PCL-R demonstrated that cutoff scores for psychopathy in men vary between North America (30 of a possible 40 points) and Europe (25 of a possible 40 points). We used a score of 25 as the threshold for psychopathy. We calculated total PCL-R, factor 1 (interpersonal and affective traits) and factor 2 (impulsivity and antisocial lifestyle) scores for all participants.⁷

This study was approved by the Joint South London and Maudsley and the Institute of Psychiatry NHS Research and Ethics Committee. Participants were paid the UK minimum hourly wage for their time and were encouraged to desist from using substances, including alcohol and illicit drugs (appendix), from 2 weeks before and during the study, and use was assessed in urine and saliva samples collected at each session.

Measures

Forensic psychiatrists did diagnostic interviews in which they assessed participants with SCID I and SCID for axis

See Online for appendix

II disorders (SCID II) in video-taped interviews. Trained forensic psychiatrists and psychologists administered the PCL-R in interviews, which were also video-taped. A random sample of 25% of participants was rerated by a second psychologist to test the intraclass correlation for total scores. All participants completed the Wechsler adult intelligence scale, third edition, and the reactive-proactive aggression questionnaire.

Probabilistic response reversal task

The scanning environment event-related probabilistic response reversal task assesses the ability to learn (acquisition phase) and alter (reversal phase) stimulus-response associations as a function of contingency change. Participants were presented with pairs of images (line drawings of animals or furniture) on a screen while in the fMRI scanner and asked to select one. Each choice received positive or negative feedback (gain or loss of points;

figure 1). To increase task difficulty, stimulus pairs had reinforcement probabilities of 100:0 (positive or negative feedback was given for 100% of right or wrong selections, respectively) or 80:20 (positive or negative feedback was given for 80% of right or wrong selections, respectively, but swapped for 20% of trials). In the acquisition phase, pairs of images were presented 20 times for the 100:0 condition or 25 times for the 80:20 condition, with each individual image assigned rewarding reinforcement. In the reversal phase, the reinforcement contingency associated with the pair of images then reversed and they were presented a further 20 or 25 times. Running point totals were displayed for each acquisition or reversal trial (appendix).

Image acquisition and processing

Participants were scanned in a 1.5 T Excite MRI scanner (GE Medical Systems, Hatfield, UK), with use of an eight-channel head coil. Four dummy acquisition scans were

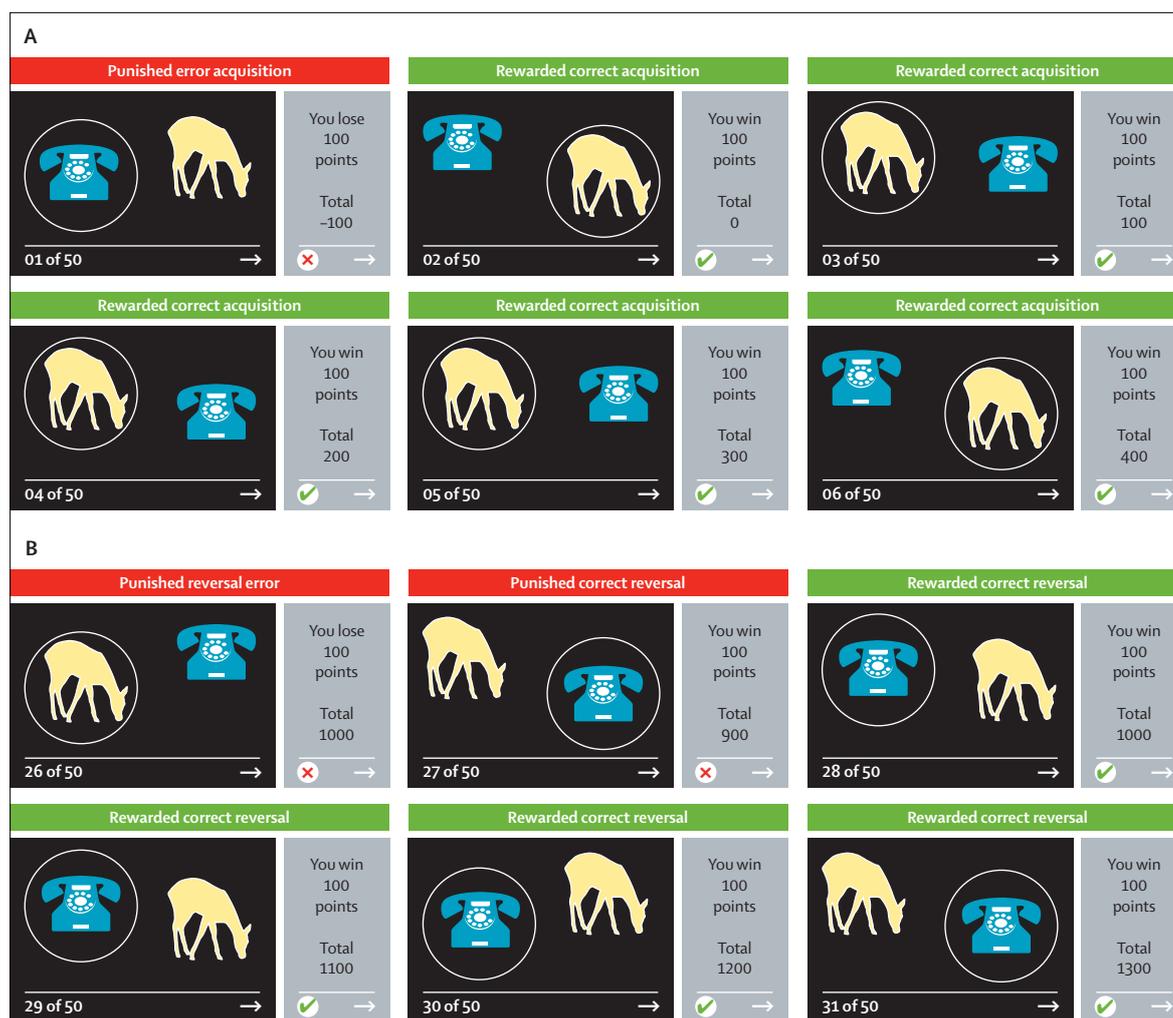


Figure 1: Schematic of the probabilistic response-reversal task

(A) The acquisition phase. (B) The reversal phase. 80:20 reinforcement probability is illustrated, where positive or negative feedback was given for 80% of right or wrong selections, respectively, but was swapped for 20% of trials. Participants selected one of the pair of images and received rewarding or punishing feedback for 25 acquisition trials, after which the contingency was reversed for 25 trials so that previously rewarded images resulted in punishment.

used to establish steady-state longitudinal magnetisation. For each participant, 192 T_2^* -weighted images were acquired with a gradient-echo, planar, imaging sequence with repetition time 2500 ms, echo time 40 ms, flip angle 90°, and in-plane resolution 3.75 mm². 25 axial slices of 5 mm thickness with 0.5 mm gaps between were collected. A high-resolution T1-weighted spoiled gradient re-echo image was also acquired, which was used for coregistration and normalisation of the data. 124 slices of 1.6 mm thickness were collected with repetition time 34 ms, echo time 9 ms, flip angle 30°, field of view 20 cm, and an acquisition matrix of 256 × 192.

Statistical analysis

Demographic and clinical variables were compared with two-sample *t* tests, ANOVA, or χ^2 and Fisher's exact tests. To test for differences in performance between groups, the percentage of errors and response latency were each assessed with 3 × 2 (group × phase) repeated-measures ANOVA. In the general linear model, we applied seven conditions modelled at the first level for each participant: baseline; rewarded correct acquisition; rewarded correct reversal; punished correct acquisition; punished correct reversal; punished errors acquisition; punished errors reversal. The numbers of trials within each of these conditions (excluding the motor baseline which was consistent for each block) for each group were also assessed with one-way ANOVA. All analyses were done with SPSS (version 15).

Image preprocessing and analyses were done with SPM5 software (running under Matlab 7.0.1 on a UNIX platform). All fMRI time series were realigned,

coregistered to the spoiled gradient re-echo image, spatially normalised to the International Consortium for Brain Mapping template image and smoothed with an 8 mm full width at half maximum Gaussian kernel. Movement parameters were included as regressors. At the group level we assessed the contrast in blood-oxygenation-level-dependent (BOLD) responses for punished reversal errors versus rewarded correct responses (rewarded correct acquisition + rewarded correct reversal).

One-sample *t* tests were done to identify punished reversal error and rewarded correct response task systems for each group, with an initial significance threshold of $p=0.001$ and a familywise error cluster corrected significance threshold of 0.05. The *t* test values, which were modelled within the SPM factorial design, were calculated to investigate potential differences between groups to the punished reversal error and rewarded correct response trials, with an initial significance threshold of $p=0.005$ and familywise error cluster corrected. We extracted β values from regions of differential activation from men with antisocial personality disorder with and without psychopathy by use of the SPM toolbox for illustrative purposes. Finally, a secondary analysis was done to explore whether the observed group differences in BOLD contrast between offenders with antisocial personality disorder with and without psychopathy for punished reversal errors versus rewarded correct responses were specifically related to the syndrome of psychopathy or could alternatively be explained by antisocial personality disorder symptom severity. Thus, independent correlations between activity during punished reversal errors compared

For the SPM toolbox see <http://marsbar.sourceforge.net/>

	Group			Group comparison		Post-hoc test		
	NO (n=18)	ASPD-P (n=20)	ASPD+P (n=12)	Statistic	p value	NO vs ASPD-P	NO vs ASPD+P	ASPD-P vs ASPD+P
Age (years)	34.8 (8.8)	36.8 (7.6)	40.1 (8.9)	$F_{2,47}=1.39$	0.26	1	0.31	0.88
FSIQ score	97.2 (11.0)	91.7 (12.2)	89.8 (11.2)	$F_{2,47}=1.71$	0.19	0.46	0.3	1
Duration of education (years)	12.2 (1.5)	10.5 (0.8)	10.4 (1.2)	$F_{2,47}=11.43$	0.001	0.001	0.001	1
Personality disorder other than ASPD (%)								
Cluster A	0	10	8.3	1.92	0.45
Cluster B	0	10	8.3	1.92	0.45
Cluster C	0	5	8.3	1.65	0.71
PCL-R score (mean [range])								
Total (0–40)	3.4 (0–8)	15.9 (10–24)	28.2 (26–30)	$F_{2,45}=246.79$	0.001	0.001	0.001	0.001
Factor 1 (0–20)	0.6 (0–4)	4.5 (0–11)	9.6 (6–13)	$F_{2,45}=55.36$	0.001	0.001	0.001	0.001
Factor 2 (0–20)	2.3 (0–6)	10.3 (5–15)	15.8 (12–19)	$F_{2,45}=110.70$	0.001	0.001	0.001	0.001
Age at first violent conviction (years)	N/A	22.3 (9.9)	17.1 (3.8)	$t=2.08, df=27$	0.05
Number of violent convictions	N/A	4.5 (3.2)	7.6 (5.9)	$t=-1.68, df=15$	0.11
Aggression score								
Total	7.1 (3.6)	15.7 (7.3)	22.9 (12.0)	$F_{2,42}=13.44$	0.001	0.001	0.002	0.22
Proactive	2.0 (2.3)	7.4 (3.9)	13.8 (6.6)	$F_{2,42}=24.03$	0.012	0.001	0.001	0.02
Reactive	5.1 (3.4)	8.3 (5.7)	11.7 (7.0)	$F_{2,42}=4.91$	0.001	0.151	0.027	0.44

Group data are mean (SD) unless stated otherwise. NO=non-offenders. ASPD-P=violent offenders with antisocial personality disorder but not psychopathy. ASPD+P=violent offenders with antisocial personality disorder and psychopathy. FSIQ=full-scale intelligence quotient. PCL-R=psychopathy checklist-revised. N/A=not applicable.

Table 1: Sociodemographic, clinical, and behavioural characteristics of participants

with rewarded correct responses for PCL-R scores and total number of SCID II antisocial personality disorder symptoms in the entire group of offenders with antisocial personality disorder were modelled.

Role of the funding source

The funder of the study had no role in the study design, data collection, data analysis, data interpretation, or writing of the report. The corresponding author had full access to all the data in the study and had final responsibility for the decision to submit for publication.

Results

The three groups did not differ significantly except for education and PCL-R score (table 1). The offenders with antisocial personality disorder and psychopathy were significantly younger at first conviction than those without psychopathy. They also had higher numbers of convictions for violent crime, but this difference was not significant. Significantly more offenders had lifetime diagnoses of alcohol and cocaine dependence than non-offenders (appendix). Of note, though, the proportions of offenders with and without psychopathy and with lifetime substance use disorders did not differ.

Intraclass correlation for PCL-R total scores was 0.83. Groups did not differ significantly for percentage errors ($F_{2,47}=1.22$, $p=0.31$) or response latencies ($F_{2,47}=0.87$, $p=0.43$) in the acquisition and reversal phases of the task (appendix). For percentage errors, there was an effect of phase ($F_{1,47}=107.4$, $p=0.001$), with more errors being committed during the reversal phase, but no significant phase-group interaction ($F_{2,47}=0.511$, $p=0.60$). For response latencies, an effect of phase was seen but was not quite significant ($F_{1,47}=3.51$, $p=0.07$), with responses being slower during the reversal phase, but no significant interaction was seen for phase and group ($F_{2,47}=0.15$, $p=0.86$). The number of trials for each condition (appendix) and the proportion of trials completed were similar in all groups ($p=0.25$).

Offenders with antisocial personality disorder and psychopathy, compared with those without psychopathy and non-offenders, displayed increased BOLD responses to punished reversal error trials within the bilateral posterior cingulate cortex and precuneus (Brodmann areas 23 and 31). Increased activation was also seen in offenders with psychopathy during punished reversal errors trials in the right anterior insula when compared with offenders without psychopathy, but not non-offenders. A difference between offenders with antisocial personality disorder and psychopathy and non-offenders, however, did become evident when a lower cluster-defining threshold of $p=0.01$ was used. Conversely, the antisocial offenders without psychopathy and the non-offenders showed significantly increased BOLD responses during rewarded correct response trials in the right superior temporal gyrus (Brodmann area 22), extending to the anterior middle temporal gyrus (Brodmann area 21) compared with

	Brodmann area	MNI coordinates			Cluster size	Z score	p value*
		x	y	z			
ASPD+P>ASPD-P, punished reversal errors>rewarded correct responses							
Left posterior cingulate cortex							
Posterior cingulate cortex	23	-3	-24	30	818	3.84	0.001
Posterior cingulate cortex	31	0	-39	27
Precuneus	7	-3	-69	42
Right posterior cingulate cortex							
Posterior cingulate cortex	31	6	-45	36
Precuneus	7	0	-63	36
Posterior cingulate cortex	23	9	-57	12
Right anterior insula							
Inferior frontal gyrus	44	51	3	18	204	3.16	0.044
Insula	..	36	3	0
Insula	..	33	9	-3
ASPD+P>non-offenders, punished reversal errors>rewarded correct responses							
Left posterior cingulate cortex							
Posterior cingulate cortex	23	-3	-39	27	281	3.47	0.011
Posterior cingulate cortex	31	-3	-63	15
Posterior cingulate cortex	23	-6	-54	9
Right posterior cingulate cortex							
Posterior cingulate cortex	23	3	-54	9
Precuneus	7	3	-66	33
Precuneus	7	3	-54	39
ASPD-P>ASPD+P, rewarded correct responses>punished reversal errors							
Right superior temporal gyrus							
Middle temporal gyrus	21	54	-36	-6	211	3.68	0.039
Superior temporal gyrus	21	60	-18	-3
Superior temporal gyrus	22	45	-21	3
Non-offenders>ASPD+P, rewarded correct responses>punished reversal errors							
Right superior temporal gyrus							
Superior temporal gyrus	22	45	-18	3	251	3.76	0.019
Superior temporal gyrus	..	48	-6	0
Superior temporal gyrus	..	39	-15	18

BOLD=blood-oxygenation-level-dependent. ASPD-P=violent offenders with antisocial personality disorder but not psychopathy. ASPD+P=violent offenders with antisocial personality disorder and psychopathy. *Corrected for familywise error.

Table 2: Areas with significantly different BOLD responses for the contrast punished reversal error trials versus rewarded correct response trials

offenders with antisocial personality disorder and psychopathy. No significant differences in BOLD activity in either direction were seen between the offenders with antisocial personality disorder but not psychopathy and non-offenders (table 2, figures 2, 3).

All three groups showed significantly greater activation during punished reversal error trials than during rewarded correct response trials within the ventrolateral prefrontal cortex, dorsolateral prefrontal cortex, and dorsomedial prefrontal cortex. Offenders with antisocial personality disorder with psychopathy and non-offenders also showed significantly greater activity within the inferior parietal lobe during punished reversal error trials than during rewarded correct response trials. Only offenders with antisocial personality disorder and psychopathy showed significantly

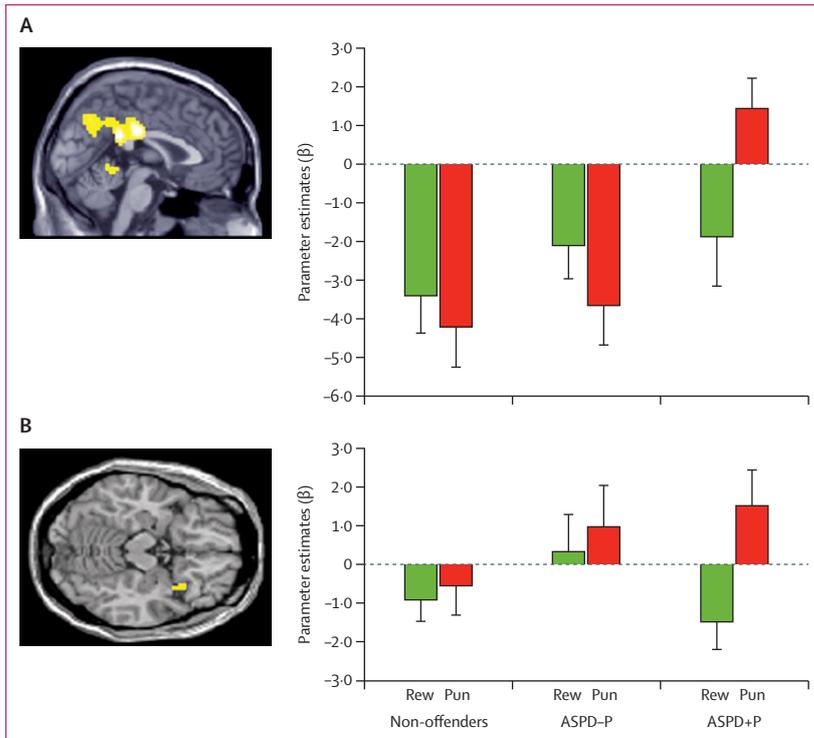


Figure 2: Mean values for the contrast punished reversal errors > rewarded correct responses
 (A) Bilateral posterior cingulate cortex. (B) Right anterior insula. In these two regions the ASPD+P group showed significantly increased activation to punished reversal errors relative to the two comparison groups. Areas of significantly increased activation in ASPD+P group compared with the ASPD-P group are overlaid on the MNI T1 template. Error bars represent SE. Rew=rewarded correct responses. Pun=punished reversal errors. ASPD-P=antisocial personality disorder without psychopathy. ASPD+P=antisocial personality disorder with psychopathy.

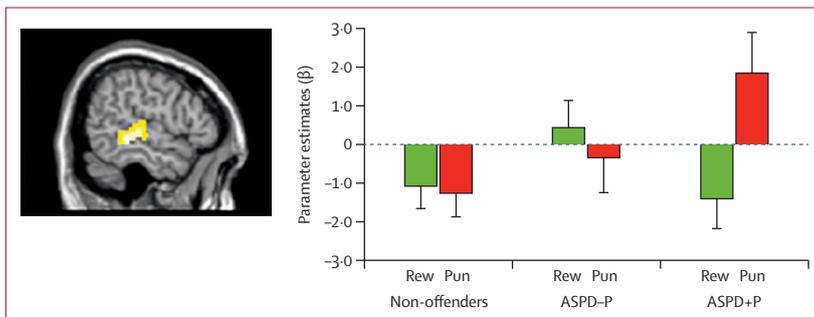


Figure 3: Mean values for the contrast rewarded correct responses > punished reversal errors in the right superior temporal gyrus
 In this region the ASPD-P and the non-offender groups showed significantly increased activation to rewarded correct responses relative to the ASPD+P group. Error bars represent SE. Rew=rewarded correct responses. Pun=punished reversal errors. ASPD-P=antisocial personality disorder without psychopathy. ASPD+P=antisocial personality disorder with psychopathy.

greater BOLD responses in the posterior cingulate cortex during punished reversal error trials than during rewarded correct response trials. By contrast, those without psychopathy and non-offenders showed significantly greater activity to rewarded correct response trials than to punished reversal error trials within this region and in the superior temporal gyrus. In offenders with antisocial personality disorder and psychopathy, no regions showed

significantly greater activity to rewarded correct response trials than to punished reversal error trials. The results remained unchanged when group comparisons were repeated excluding offenders who had positive urinary drug screens on the day of scanning (appendix).

The secondary correlation analyses in offenders showed a positive relation between PCL-R scores and activity within the posterior cingulate cortex region (MNI coordinates $x=-3$, $y=-66$, and $z=42$, cluster size 323, $Z=3.52$, $p=0.005$ corrected) during punished reversal error trials versus rewarded correct response trials. However, no significant relation was seen between the number of SCID II antisocial personality disorder symptoms and activation in this area.

Discussion

We investigated the neural basis of reversal learning in violent offenders with antisocial personality disorder with and without psychopathy. Offenders with antisocial personality disorder and psychopathy displayed abnormal responding to punishment signals within the posterior cingulate and insula, illustrated by significantly greater BOLD activation to punished reversal errors than to rewarded correct responses in both regions. These differences were not related to the number of antisocial personality disorder symptoms. Offenders with antisocial personality disorder but not psychopathy and non-offenders showed significant reductions in activation in the posterior cingulate cortex, and the offenders without psychopathy had significantly reduced activation in the anterior insula.

Reversal learning impairments have been robustly identified in children with conduct disorder and callous unemotional traits and in adult men with psychopathy.^{12,24} In this study, however, we noted no behavioural deficits, which is consistent with results of the fMRI investigation with the same task in adolescents with conduct disorder and callous unemotional traits.²¹ In these two fMRI studies, abnormal activations were seen in brain regions signalling prediction errors. Adolescents with conduct disorder and callous unemotional traits showed greater activity within the ventromedial prefrontal cortex to punished reversal errors than did healthy adolescent controls.²¹ Offenders with antisocial personality disorder and psychopathy, unlike those without psychopathy and non-offenders, showed significantly greater activation within the posterior cingulate cortex to punished reversal errors. Consistent with previous findings in healthy adults, reduced BOLD responses within the posterior cingulate to punished reversal errors compared with those for rewarded correct responses were seen in the offenders with antisocial personality disorder but not psychopathy and in the non-offenders.^{19,20}

The posterior cingulate cortex plays an important part in altering behaviour in response to an unexpected change.²⁵ Animal and human studies have shown the relevance of this cortex in representing subjective value, particularly in relation to rewards,²⁶ and in error signalling after the omission of an expected reward.²⁷ The posterior cingulate

typically responds to prediction error in conjunction with the ventromedial prefrontal cortex.²⁷ In previous studies, structural posterior cingulate cortex abnormalities were seen in children with callous unemotional traits²⁸ and some adults with psychopathy,²⁹ but these were not seen in the sample in this study.⁷ Reduced posterior cingulate cortex responsivity to emotional stimuli,³⁰ including personal moral dilemmas,³¹ has also been seen in investigations of adults with psychopathy. A white-matter diffusion tensor imaging study of the offenders with antisocial personality disorder and psychopathy from this study revealed reduced fractional anisotropy suggestive of reduced axonal integrity and organisation in the dorsal cingulum, which links the posterior cingulate cortex to the medial prefrontal cortex.³² The findings from this study support the proposition that the representation of reinforcement value is profoundly disturbed in adult men with psychopathy (panel).³³

Thus, in adolescents²¹ and adults, rather than psychopathic traits being associated with decreases in activity within reinforcement-sensitive regions after unexpected punishment, significantly greater activation within these regions was seen in response to punished trials. This result challenges the notion that individuals with psychopathy are simply insensitive to subjective value, prediction errors, or both.³³ Men with psychopathy were sensitive to this information and used it to inform appropriate behavioural change, but processed it in a highly atypical way. This atypical processing was associated with psychopathy but not with the number of antisocial personality disorder symptoms. Replications and further elucidation of this anomalous responding are urgently needed, not least because treatments for psychopathy that increase sensitivity to reinforcement information might be misdirected. Instead, interventions aimed at modifying the subjective value accorded to punishment information, perhaps by focusing attention appropriately, might need to be developed.¹⁵

Among the offenders with antisocial personality disorder and psychopathy, a similar abnormality was seen within the anterior insula. The anterior insula is connected to limbic regions, such as the ventromedial prefrontal cortex and amygdala, and is innervated by dopaminergic neurons. This area is involved in motivation and represents context-dependent aversive value and reward and tracks the salience of outcomes, including recognition of errors.³⁴ Insular damage in human beings promotes risky decision-making because of impaired signalling of the probability of aversive outcomes.³⁵ Reductions in grey-matter volume in the insula have been reported in adults with psychopathy,^{7,29} as have reductions in insula activity during aversive conditioning³⁰ and atypically increased activation in response to empathy-eliciting scenarios.⁸

Regions involved in representing subjective value, such as the ventromedial prefrontal cortex and posterior cingulate cortex, have been proposed to integrate input from the superior temporal gyrus.³⁶ In this study, offenders

Panel: Research in context

Systematic review

We searched PubMed for reports published in English up to Aug 8, 2014, with the search terms "punishment AND psychopathy", "fMRI AND psychopathy", "response reversal AND psychopathy", "reinforcement learning AND psychopathy", and "fMRI AND antisocial". We also checked the reference lists of identified reports for relevant publications. We found no studies that had assessed the neural underpinnings of reversal learning in men with antisocial personality disorder with and without psychopathy.

Interpretation

We found distinctive neural mechanisms related to the severe impairment in learning from punishment that characterises violent offenders with antisocial personality disorder and psychopathy. Important neurobiological distinctions between men with antisocial personality disorder with and without psychopathy have been theorised. The lack of evidence for such differences, however, has led to diagnostic classification schemes that favour a single diagnosis of antisocial personality disorder, which has greatly hindered progress in understanding the causes of subtypes and their optimum treatment. Child diagnostic classification systems specify the developmental precursor of psychopathy, limited prosocial emotions, as an important subgroup within those with conduct disorder. Our findings add to the weight of evidence encouraging similar specifications in adult diagnostic schemes to distinguish the syndrome of psychopathy from antisocial personality disorder and to take the subgroups into account when planning rehabilitation programmes. As most violent crimes are committed by men with this early-onset stable pattern of antisocial and aggressive behaviour, interventions that target the specific underlying brain mechanisms and effect change in the behaviour have the potential to significantly reduce the rate of violent crime.

with antisocial personality disorder without psychopathy and non-offenders showed significantly greater activity within this region to rewarded correct responses than to punished reversal errors, whereas offenders with antisocial personality disorder and psychopathy showed significantly less activity. These results augment those from previous studies that have suggested structural²⁹ and functional³⁷ abnormalities within the superior temporal gyrus in individuals with psychopathy. Moreover, they suggest that this dysfunction contributes to deficits in decision making.³⁶

This study has several limitations. The violent offenders with antisocial personality disorder who participated in the study, like most men with this disorder, had additional personality disorders and current or previous substance use disorders. The proportions of such disorders that could affect reversal learning, however, were similar in our two offender groups. The observed functional differences, therefore, cannot simply be attributed to these factors. The use of the contrast of punished reversal errors versus rewarded correct responses cannot distinguish between value representations (contingency tracking) and prediction errors (the amount and valence of surprise associated with feedback on a given trial). The strengths of the study included diagnoses and PCL-R ratings made by trained clinicians, the use of official criminal records to classify participants, and measurement of substance use before the scan.

We identified neural dysfunctions in violent offenders with antisocial personality disorder and psychopathy that

did not characterise antisocial personality disorder in violent offenders without psychopathy or non-offenders. The important areas of differential activity in offenders with antisocial personality disorder and psychopathy were the posterior cingulate cortex and anterior insula, where activity was increased in response to punished reversal errors, which is indicative of dysfunctional prediction error signalling. Additionally, offenders in this group were hypo-responsive to reward information in the superior temporal gyrus, which suggests a failure to consolidate reward information. We have provided further evidence of distinctive neural anomalies that distinguish between individuals with antisocial personality disorder with and without psychopathy. Diagnostic classification schemes, offender rehabilitation programmes,³⁸ and childhood prevention programmes would benefit from taking account of this mounting evidence.

Contributors

RJB, Dff, VK, SH, and NB conceived and designed the study and Dff, VK, SH, and NB obtained funding. SG and NB were responsible for acquisition of data. SG, RJB, Dff, and NB analysed and interpreted the data. SG, SH, and NB did the statistical analyses. AS, SH, and NB provided administrative, technical, or material support. SH and NB supervised the study. SG, SH, and NB drafted the paper, and all authors contributed to critical revision for intellectual content.

Declaration of interests

We declare no competing interests.

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