

AKADÉMIAI KIADÓ

# Structural brain differences related to compulsive sexual behavior disorder

Journal of Behavioral Addictions

12 (2023) 1, 278–287

DOI:  
10.1556/2006.2023.00008  
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Received: November 13, 2022 • Revised manuscript received: January 16, 2023; February 11, 2023 • Accepted: February 25, 2023

Published online: March 21, 2023

## FULL-LENGTH REPORT



### ABSTRACT

*Background and aims:* Compulsive sexual behavior disorder (CSBD) has been included as an impulse control disorder in the International Classification of Diseases (ICD-11). However, the neurobiological mechanisms underlying CSBD remain largely unknown, and given previous indications of addiction-like mechanisms at play, the aim of the present study was to investigate if CSBD is associated with structural brain differences in regions involved in reward processing. *Methods:* We analyzed structural MRI data of 22 male CSBD patients (mean = 38.7 years, SD = 11.7) and 20 matched healthy controls (HC; mean = 37.6 years, SD = 8.5). Main outcome measures were regional cortical thickness and surface area. We also tested for case-control differences in subcortical structures and the effects of demographic and clinical variables, such as CSBD symptom severity, on neuroimaging outcomes. Moreover, we explored case-control differences in regions outside our hypothesis including white matter. *Results:* CSBD patients had significantly lower cortical surface area in right posterior cingulate cortex than HC. We found negative correlations between right posterior cingulate area and CSBD symptoms scores. There were no group differences in subcortical volume. *Conclusions:* Our findings suggest that CSBD is associated with structural brain differences, which contributes to a better understanding of CSBD and encourages further clarifications of the neurobiological mechanisms underlying the disorder.

### KEYWORDS

compulsive sexual behavior, hypersexual disorder, neuroimaging, surface area, reward, brain structure

## INTRODUCTION

According to the International Statistical Classification of Diseases and Related Health Problems (ICD-11) (World Health Organization, [WHO], 2022), compulsive sexual behavior disorder (CSBD) is characterized by a persistent pattern of failure to control intense sexual impulses or urges often resulting in repetitive sexual behavior. In CSBD, out-of-control sexual behaviors are pursued regardless of adverse consequences. Approximately 3–5% of the general population suffer from CSBD, rendering it a significant health burden (Bóthe et al., 2020; Briken et al., 2022). While promising treatment options are already been offered, their effectiveness can potentially be improved (Antons et al., 2022; Borgogna, Garos, Meyer, Trussell, & Kraus, 2022; Briken, 2020; Turner et al., 2022).

Despite CSBD being classified as an impulse control disorder in ICD-11, the underlying neurobiological mechanisms remain substantially unknown (Derbyshire & Grant, 2015). While similar mechanisms to those playing a role in obsessive compulsive and substance use

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disorders may be involved in CSBD (Coleman, 1991; Gola et al., 2017; Voon et al., 2014), impairments of brain regions regulating sexual arousal and desire have also been proposed (Blum, Badgaiyan, & Gold, 2015; Carnes et al., 2012; Derbyshire & Grant, 2015; Estellon & Mouras, 2012; Kingston & Bradford, 2013; Kor, Fogel, Reid, & Potenza, 2013; Kraus, Voon, & Potenza, 2016; Kuhn & Gallinat, 2016; Weinstein, Katz, Eberhardt, Cohen, & Lejoyeux, 2015). Neuroimaging studies suggested that processing of sexual stimuli is altered in CSBD patients (Stark, Klucken, Potenza, Brand, & Strahler, 2018) and that CSBD is associated with altered functioning in brain areas involved in sensitization, habituation, impulse control, and importantly reward processing (Kowalewska et al., 2018), e.g., fronto-temporal cortices, amygdala, and ventral striatum (Gola & Draps, 2018; Kowalewska et al., 2018; Voon et al., 2014). In summary, previous studies strongly indicate that the brain reward system plays a crucial role in CSBD (Kowalewska et al., 2018; Schmidt et al., 2017; Voon et al., 2014), and that the underlying key mechanisms are similar to those found in substance or behavioral addictions (Gola & Draps, 2018; Kowalewska et al., 2018; Mechelmans et al., 2014). However, structural MRI studies in CSBD are scarce and the brain structural correlates of CSBD remain largely unknown (Balodis & Potenza, 2015).

The aim of the present study was to test for differences in brain structure between CSBD patients and healthy controls, after correcting for age. We focused on predefined regions of interest known to be involved in reward processing. We analyzed structural MRI data from 22 men with CSBD and 20 healthy controls (HC). Main outcome measures were cortical thickness and surface area. We also tested for subcortical differences and correlations between MRI-derived phenotypes and CSBD symptoms. We also explored case-control differences in cortical regions outside our hypothesis and white matter underlying all cortical regions investigated. Moreover, we conducted sensitivity analyses testing for the effects of demographic and clinical variables on our outcomes, such as medication use and other clinical characteristics.

## METHODS

### Participants

The study was conducted at ANOVA, a multidisciplinary clinic for research, assessment, and treatment in the fields of andrology, sexual and transgender medicine at the Karolinska University Hospital, and at Karolinska Institutet, Stockholm, Sweden. Patients seeking treatment for CSBD at ANOVA were recruited through the phone helpline *PreventTell*, designed to prevent sexual offences by targeting persons at risk or with unwanted sexual behaviors. Details on recruitment, eligibility criteria, and procedures can be found in the Supplemental Material and elsewhere (Hallberg et al., 2020; Liberg et al., 2022; Savard et al., 2020). In brief, psychiatric diagnoses were established through interviews with clinical psychologists and psychiatrists, including the Mini-

International Neuropsychiatric Interview (M.I.N.I. v7.0) (Sheehan et al., 1998), and a semi-structured clinical interview focusing on compulsive sexual behavior. Male participants who met criteria for Hypersexual Disorder (HD) as proposed for DSM-5 (Kafka, 2010) and for CSBD according to ICD-11 were included in the present study. Age-matched controls were recruited through online and public advertisements. Controls were physically and psychologically healthy and screened negative for CSBD, e.g. a score below 20 points on the Hypersexual Disorder Screening Inventory (HDSI) (Parsons et al., 2013), and below 53 points on the Hypersexual Behavior Inventory (HBI) (Reid, Garos, & Carpenter, 2011). Patients had not started any psychological treatment.

General inclusion criteria were being male, at least 18 years of age, and fluent in the Swedish language. Detailed exclusion criteria are listed in the Supplemental Material. In short, for both patients and controls, we excluded for MRI contraindications, severe neurological/psychiatric diseases or conditions, current alcohol or substance dependence, pathological gambling or other behavioral addictions, untreated endocrine diseases, and patients using specific medications. Patients with paraphilias (except for pedophilic interests) were included.

We included 23 CSBD males, of which 22 underwent an MRI scan, and 20 HC. While we previously reported functional MRI findings including the same study cohort (Liberg et al., 2022), this study focusses on structural MRI outcomes.

### Measures

**Clinical characteristics and questionnaires.** Clinical characteristics were also assessed through questionnaires; e.g. symptoms of compulsive sexual behavior and neurodevelopmental disorders, depression, drug- and alcohol use and impulsivity. In brief, CSBD related symptoms were assessed using the Hypersexual Behavior Inventory (HBI), a 19-item scale constructed to reflect the proposed criteria of hypersexual disorder (Reid et al., 2011), the Sexual Compulsivity Scale (SCS), a 10-item scale that assesses compulsive sexual behaviors and urges (Kalichman & Rompa, 1995), and the Hypersexual Disorder Screening Inventory (HDSI), a screening questionnaire according to the DSM-5 criteria (Parsons et al., 2013).

We further assessed sexual desire using the Sexual Desire Inventory (SDI, 14 items) (Spector, Carey, & Steinberg, 1996) and severity of depression using the Montgomery Åsberg Depression Rating Scale (MADRS-S, 9 items) (Svanborg & Åsberg, 2001). The full list of questionnaires used in this study can be found in Supplemental Material. Finally, we assessed the frequency of sexual encounters within the last 6 months, as well as weekly pornography consumption.

**Magnetic resonance imaging.** A 3T GE MRI scanner (Discovery MR750) equipped with an eight-channel head coil was used. Sagittal T1-weighted images were acquired using a 3D-BRAVO sequence (TR = 6.40 ms, TE = 2.81 ms, FOV = 24.0 cm, flip angle = 12°, TI = 450 ms, acquisition matrix 240 × 240 × 180, voxel size 1 × 1 × 1 mm<sup>3</sup>).



## Procedure

**Image processing and region of interest selection.** Measures for regional cortical thickness and surface area of 34 regions per hemisphere (Desikan atlas (Desikan et al., 2006)) were obtained from structural T1-weighted images using the semi-automated cortical surface reconstruction and parcellation methods provided by FreeSurfer (Dale, Fischl, & Sereno, 1999; Fischl & Dale, 2000; Fischl, Salat, et al., 2004; Fischl, Sereno, & Dale, 1999; Fischl, van der Kouwe, et al., 2004). The primary analysis was a hypothesis-driven region of interest (ROI) approach and focused on cortical brain areas involved in reward processes. The main selection was done using [neurosynth.org](http://neurosynth.org) (keyword “reward”) which resulted in the selection of following seven cortical structures: medial and lateral orbitofrontal cortex, rostral and caudal cingulate cortex, posterior cingulate cortex, anterior insula, and superior frontal cortex. Three subcortical structures were investigated: nucleus accumbens, amygdala, and caudate. Considering left and right hemispheres, this yielded 14 cortical and six subcortical ROIs. The regions selected align with those reported to be involved in reward-related processes according to recent review articles (Oldham et al., 2018). In addition, the selected ROIs largely overlap with regions related to the processing of visual sexual stimuli and arousal (Georgiadis & Kringelbach, 2012), which is of further interest for CSBD. Regions outside of the main hypothesis were analyzed in exploratory whole brain analyses as well as volume of white matter underlying these regions. Volumetric measures were also obtained using FreeSurfer.

## Statistical analyses

**Group characteristics (demographic and clinical data).** Demographic and clinical variables were compared using *t*-tests or Fisher’s exact  $\chi^2$ .

**Differences in brain structure between CSBD patients and healthy controls.** In main analyses, we tested for the effect of group (CSBD vs HC; independent variable of interest) on cortical thickness (and area), as well as subcortical volumes. Age was used as covariate to account for potential age-related variance in the MRI data given that cortical changes occur along with age. For each imaging phenotype (thickness, area, subcortical volumes) the tests were separately conducted in SPSS v26 using multiple univariate analyses of covariance (ANCOVAs), where brain imaging phenotypes were set as dependent variables, group was entered as fixed factor and age was set as covariate. Since cortical volume is a function of the two genetically and phenotypically independent measures cortical thickness and surface area, these measures were analyzed separately, and cortical volume analyses were omitted (Winkler et al., 2010). We further treated cortical and subcortical analyses as independent studies, which are often reported separately (Hibar et al., 2016, 2018). However, we report cortical and

subcortical findings in the same manuscript. Although we did not have lateral hypotheses, we did not combine measures from left and right hemisphere to total measures. While this increased the number of tests conducted for each phenotype, this approach was chosen to provide lateral information. Hence, within each imaging phenotype, we corrected for multiple testing using Bonferroni’s Dubey Armitage-Parmar/Sidak’s adjustment of  $\alpha$ -level, which considers the number of regions ( $N = 14$  for cortical regions, and  $N = 6$  for the subcortical regions) performed and their inter-correlation ( $r_{\text{thickness}} = 0.41$ ,  $r_{\text{area}} = 0.41$ ,  $r_{\text{subcortical}} = 0.37$ ; leading to adjusted  $\alpha$ -levels of  $\alpha_{\text{thickness}} = \alpha_{\text{area}} = 0.013$ ,  $\alpha_{\text{subcortical}} = 0.016$ ) (Sankoh, Huque, & Dubey, 1997).

Distributions of dependent variables were tested for normality using the one-sample Kolmogorov-Smirnov test. Mean differences between HC and CSBD subjects were quantified with Cohen’s *d*. Results were visualized with ENIGMA Viewer v.2016.07.08, where *p*-values and effect size (Cohen’s *d*) for each brain regions were color-graded and mapped to into the brain space.

## Correlations between structural brain phenotypes and CSBD symptom scores (exploratory).

The primary findings were further complemented with regression analyses testing for correlations between brain regions that showed group differences and CSBD symptoms scores. These analyses were performed in the combined sample of patients and controls to a) widen the range of symptoms and b) increase statistical power. Another rationale was to test if a relationship between symptom scores and structural brain variation can be detected regardless of a categorical diagnostic label. Brain measures were dependent variables, symptoms scores and age were independent variables. We tested for correlations with the following symptoms scores (one at a time): HDSI, HBI, SDI and SCS. Analyses were also run separately within patients and controls, and in the combined cohort while controlling for case-control status, to test if observed correlations were driven by case-control differences. Note, that a correlation driven by case-control differences can still reflect a genuine relationship between CSBD and brain structure, and results obtained when controlling for case-control status should be interpreted with caution.

**Sensitivity analyses.** In sensitivity analyses testing for potential confounders, the main analysis was repeated while adding potential confound variables as covariates. These included demographic or clinical variables (e.g., comorbidities, medication use) as listed in Table 1. The full list of variables tested for in sensitivity analyses can be found the Supplemental Material (Table S1a). Such variables were entered one at a time in the ANCOVA models. In additional tests, when fewer than ten participants had a specific comorbidity or used a specific medication, the sensitivity analyses were repeated one at a time after excluding those individuals. CSBD symptom scores were not controlled for, as they relate to the phenotype of interest and controlling for



Table 1. Demographic and clinical characteristics

Measure	HC ( <i>n</i> = 20)	CSBD ( <i>n</i> = 23)	HC vs. CSBD ( <i>p</i> -value)
Age, mean (SD)	37.6 (8.5)	38.7 (11.7)	0.741
BMI, mean (SD)	23.1 (2.8)	25.8 (4.5)	0.026
Nicotine use (yes/no/ sometimes), <i>n</i>			
Moist snuff	3/16/0*	7/13/0*	0.157
Smoking	0/16/4	0/21/0*	0.048
Handedness (R/L/M), <i>n</i>	16/4/0	16/1/1*	0.822
Sexual orientation Self-identified	1	1	0.919
homosexual, <i>n</i>			
Kinsey scale, mean (SD)	0.6 (1.1)	0.71 (1.3)	0.778
HDSI, mean (SD)	1.9 (2.2)	20.2 (3.8)	<0.001
HBI, mean (SD)	22.5 (4.1)	69.4 (13.4)	<0.001
SDI, mean (SD)	55.2 (12.6)	80.6 (17.1)	<0.001
SCS, mean (SD)	11.2 (0.9)	29.4 (6.3)	<0.001
Pornography consumption times per week, mean (SD)	2.2 (2.3)	13.0 (20.7)	0.033
hours per week, mean (SD)	0.7 (0.7)	9.2 (8.0)	<0.001
age first consumption, mean (SD)	14.2 (3.4)	13.2 (4.9)	0.424
MADRS-S, mean (SD)	3.9 (4.9)	18.3 (7.8)	<0.001
AUDIT, mean (SD)	4.1 (3.8)	6.3 (3.8)	0.059
DUDIT, mean (SD)	2.7 (4.5)	2.1 (3.0)	0.582
RAADS, mean (SD)	6.1 (6.0)	11.1 (7.7)	0.025
ASRS, mean (SD)	14.7 (10.6)	34.2 (11.7)	<0.001
BIS-11, mean (SD)	53.1 (7.3)	66.7 (10.8)	<0.001
BIS/BAS			
BAS drive, mean (SD)	7.4 (2.3)	9.0 (2.7)	0.048
BAS fun seeking, mean (SD)	10.5 (2.5)	11.9 (1.7)	0.037
BAS reward response, mean (SD)	16.3 (2.1)	16.5 (1.6)	0.726
BIS, mean (SD)	17.9 (5.1)	20.7 (3.1)	0.033
STAI-S, mean (SD)	9.3 (2.0)	12.6 (2.5)	<0.001

Notes: Demographic and clinical characteristics mean (SD) or number of participants (*n*) of healthy controls (HC) and males with Compulsive Sexual Behavior Disorder (CSBD) are provided. Results (*p*-values) of group comparisons are presented. Sexual orientation was measured through a 7-point Kinsey scale. Data reported for all patients enrolled. \* indicates variables with missing data. Abbreviations: Hypersexual Disorder Screening Inventory (HDSI), Hypersexual Behavior Inventory (HBI), Sexual Desire Inventory (SDI), Sexual Compulsivity Scale (SCS), Montgomery Asberg Depression Rating Scale (MADRS-S), Alcohol Use Disorders Identification Test (AUDIT), Drug Use Disorders Identification Test (DUDIT), Ritvo Autism Asperger Diagnostic Scale (RAADS), Adult ADHD Self-Report Scale (ASRS), Barratt Impulsiveness Scale (BIS-11), Behavioral Inhibition/Activation System (BIS/BAS), State-Trait Anxiety Inventory - State (STAI-S).

these was assumed to remove variance related to the effects of interest (Hyatt et al., 2020).

## Ethics

The study was carried out in accordance with the Declaration of Helsinki. All participants were informed about the

procedures and provided informed consent. The study was approved by the regional Ethical Review Board in Stockholm, Sweden and was preregistered at [clinicaltrials.gov](https://clinicaltrials.gov) (NCT03495414).

## RESULTS

### Participants

Patient characteristics are shown Table 1. Note that these have previously been reported as part of our functional MRI study (Liberg et al., 2022). CSBD and HC males matched on age, sexual orientation and handedness. HC had lower BMI than CSBD males. With the exception of four occasional smokers in the HC group, both groups contained only non-smoking individuals. While HC had no psychiatric comorbidities, one patient reported panic disorder, and two patients reported depression, generalized anxiety disorder (GAD), and attention-deficit/hyperactivity disorder (ADHD), respectively (Table S1b). In line with this, two patients used stimulants, and four patients used antidepressants (Table S1b). However, groups did not significantly differ in psychiatric comorbidity or medication use. While age was controlled for in main analyses, we conducted further sensitivity tests to test for any potential confounds by demographic and clinical variables (see below).

CSBD patients scored significantly higher than HC on all questionnaires, except on a reward response subscale (BAS), where no group differences were found. CSBD males used pornography more frequently and for longer periods than HC. There were no group differences in age of onset of pornography use, nor for drug and alcohol consumption. We did not find group differences with respect to the frequency of sexual encounters or the number of sexual partners within the last six months (Table S2).

### Main analysis

After multiple comparison correction, we found a significant difference between HC and CSBD patients in right posterior cingulate surface area ( $p = 0.004$ , Cohen's  $d = 0.95$ , Table 2), with CSBD patients showing lower surface area than the control group. Differences in left posterior cingulate and right caudal anterior cingulate (cACC) cortical surface area were statistically significant on the  $p < 0.05$  level but did not survive multiple comparison corrections (Table 2). Similarly, left superior frontal thickness differed between HC and CSBD patients ( $p = 0.022$ , Cohen's  $d = 0.58$ , Table 3) but did not withstand multiple comparison corrections. Effect sizes were moderate to large (Tables 2 and 3). Findings are displayed in Fig. 1 (and Figure S1). There were no group differences in subcortical volume (Table S3).

### Sensitivity and secondary analyses

Group difference in right posterior cingulate surface area remained statistically significant when correcting for any



Table 2. Group differences in cortical surface area

Brain region	HC		CSBD		F (1, 39)	p	Cohens d
	Mean	SD	Mean	SD			
Lh lateral orbito frontal	2,861	276	2780	291	0.846	0.363	0.29
Lh medial orbito frontal	2,098	262	2,034	195	0.834	0.367	0.28
Lh caudal anterior cingulate	666	141	677	111	0.069	0.795	−0.08
Lh rostral anterior cingulate	917	144	876	166	0.688	0.412	0.26
Lh posterior cingulate	1,148	119	1,260	187	5.086	0.030*	−0.71
Lh insula	2,501	181	2,486	247	0.052	0.822	0.07
Lh superior frontal	7,805	719	7,748	665	0.07	0.793	0.08
Rh lateral orbito frontal	2,969	336	2,891	325	0.583	0.450	0.24
Rh medial orbito frontal	2,106	171	2,092	170	0.079	0.780	0.09
Rh caudal anterior cingulate	764	148	664	111	6.166	0.017*	0.76
Rh rostral anterior cingulate	655	111	609	92	2.082	0.157	0.45
Rh posterior cingulate	1,302	181	1,157	117	9.422	0.004**	0.95
Rh insula	2,422	218	2,474	284	0.429	0.516	−0.21
Rh superior frontal	7,583	674	7,480	739	0.215	0.645	0.15

Lh indicates left hemisphere, rh indicates right hemisphere. \* $p < 0.05$  \*\* after multiple comparisons ( $p < 0.011$ ). Surface area measures are given in  $\text{mm}^2$ . Results were mapped into brain space shown in Fig. 1.

Table 3. Group differences in cortical thickness

Brain region	HC		CSBD		F (1, 39)	p	Cohens d
	Mean	SD	Mean	SD			
Lh lateral orbito frontal	2.847	0.137	2.876	0.106	0.605	0.441	0.24
Lh medial orbito frontal	2.644	0.138	2.686	0.116	1.639	0.208	0.33
Lh caudal anterior cingulate	2.691	0.273	2.690	0.226	0.001	0.981	0.00
Lh posterior cingulate	2.580	0.154	2.606	0.169	0.453	0.505	0.16
Lh rostral anterior cingulate	2.944	0.208	2.999	0.177	1.22	0.276	0.28
Lh insula	3.082	0.149	3.111	0.143	0.471	0.497	0.20
Lh superior frontal	2.803	0.154	2.89	0.148	5.726	0.022*	0.58
Rh caudal anterior cingulate	2.524	0.231	2.558	0.255	0.272	0.605	0.14
Rh lateral orbito frontal	2.717	0.113	2.781	0.125	3.344	0.075	0.54
Rh medial orbito frontal	2.637	0.122	2.658	0.161	0.306	0.583	0.15
Rh posterior cingulate	2.580	0.127	2.647	0.154	3.363	0.074	0.48
Rh rostral anterior cingulate	2.944	0.184	2.989	0.264	0.457	0.503	0.20
Rh insula	3.065	0.134	3.09	0.196	0.278	0.601	0.15
Rh superior frontal	2.774	0.135	2.839	0.141	3.986	0.053	0.47

Lh indicates left hemisphere, rh indicates right hemisphere. \* $p < 0.05$ . Thickness measures are given in mm. Results were mapped into brain space shown in Fig. 1.

additional covariate tested, including total brain volume, with one exception: when correcting for depression symptom levels (MADRS-S) scores, the group difference in posterior cingulate surface area was no longer statistically significant ( $p = 0.286$ , Table S1a).

In the combined cohort, we observed a negative correlation between right posterior cingulate area and scales assessing hypersexuality symptoms and sexual compulsivity (HDSI, HBI, SCS). There was no significant correlation with the sexual desire (SDI) score ( $p = 0.061$ , Table S4). These correlations were not significant when correcting for group status or when repeating the correlation analysis within each group separately (CSBD  $p = 0.812$ – $0.989$ ; HC  $p = 0.396$ – $0.972$ , see Table S4 for details).

The exploratory whole brain analysis was indicative for group differences in brain areas not included in the main analysis. These included thickness of the left frontal pole,

right precuneus, and the left superior frontal cortex. Volume of white matter underlying the right posterior cingulate cortex was lower in the patient group than in controls. The full results, including means and SD for each group can be found in the Supplemental material, Tables S5–8.

## DISCUSSION

The neurobiological mechanisms underlying CSBD are unknown, and structural MRI studies that can help provide valuable insights into the neurobiology of CSBD are scarce. This study investigated structural brain differences in reward-related brain areas between 22 males diagnosed with CSBD and 20 healthy controls. Further, we tested for correlation between CSBD symptom severity and the MRI-derived phenotypes.



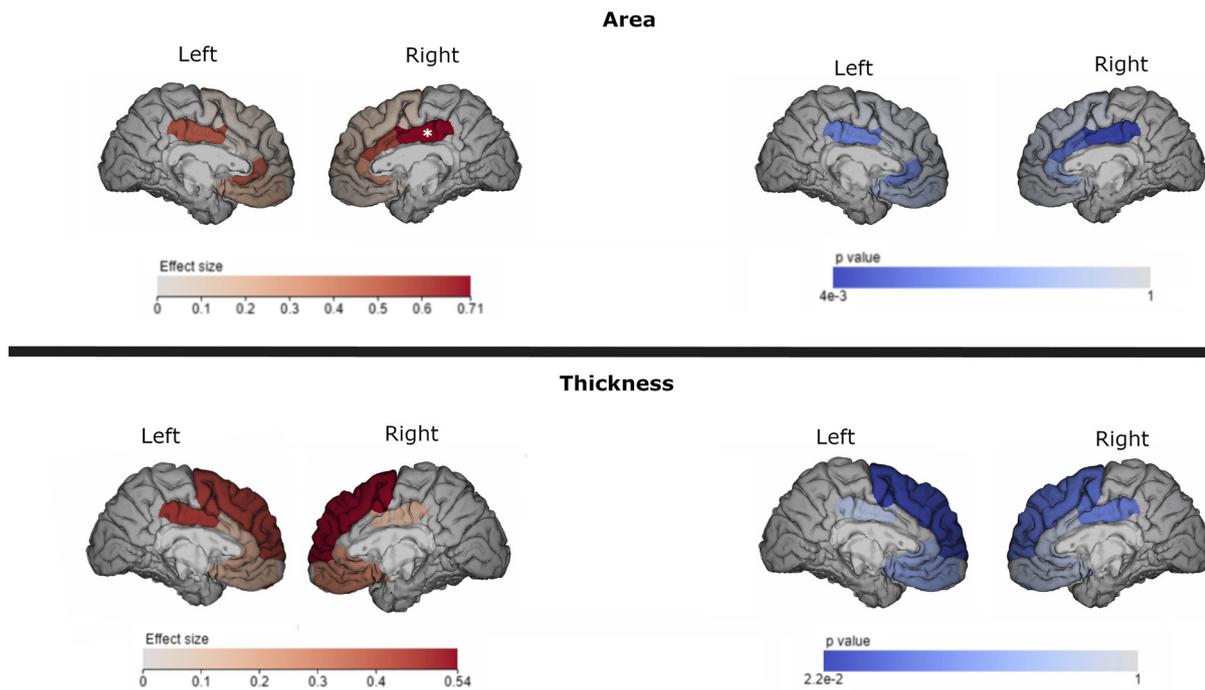


Fig. 1. Case-control differences in cortical thickness and surface area. Results ( $p$ -values) and effect sizes (Cohen's  $d$ ) are indicated (color bars). Numerical results can be found in Tables 2–3 for transparency, significance is thresholded at  $p = 0.05$ . The brain area in which significant case-control differences were observed after multiple comparison correction (right posterior cingulate) is indicated with \* (shown isolated in Figure S1)

As expected, CSBD patients scored significantly higher on CSBD-related symptom measures than HC. The main neuroimaging finding was that, compared to healthy controls, the CSBD group had significantly lower cortical surface area in right posterior cingulate cortex, indicating that CSBD is associated with cortical brain alterations. The results remained robust when controlling for several potential confounding variables.

Intriguingly, in the combined cohort, there were negative correlations between right posterior cingulate surface area and CSBD symptoms scores, indicating that CSBD symptoms were more severe in individuals displaying more pronounced cortical variations. This also suggests that the brain metrics investigated may have functional relevance in CSBD. Furthermore, our results were supported by exploratory analyses which revealed group differences in white matter volume solely in the region underlying the posterior cingulate cortex (see Table S8).

The body of structural neuroimaging studies conducted in compulsive sexual behavior have been increased in the last years, however, is still sparse. For example, Schmidt et al. (Schmidt et al., 2017) reported greater left amygdala gray-matter volume in males with CSBD than males without CSBD. Seok & Sohn (Seok & Sohn, 2018) found lower volume of the left superior and right middle temporal gyrus in males with as compared with males without CSBD. Volume of the left superior temporal gyrus was negatively correlated with the severity of CSBD, suggesting relationships between structural brain phenotypes and CSBD symptoms, as observed in the present study.

However, most previous studies used region of interest approaches that did not include the posterior cingulate cortex and/or have used different cortical metrics (predominantly volume), which does not allow the independent investigation of cortical thickness and surface area. Hence, differences in cortical surface area, as found in the present study, could have been present but remained undetected in previous studies. Moreover, previous studies varied in cohort definition, image processing methodology, and/or MR modality.

The posterior cingulate cortex is highly connected with other brain areas, yet there is no consensus about its function. In a review, Leech et al. (Leech & Sharp, 2014) state that this region is likely to function as an integrative hub, with the ventral part to be involved in memory retrieval and planning, whilst the dorsal part with its prominent connections to the frontal lobes is suggested to be involved in control of attention. Further, it is suggested that the posterior cingulate is a key structure in the network responsible for environmental change detection and provides a signal for behavioral changes when actions result in suboptimal consequences (Pearson, Heilbronner, Barack, Hayden, & Platt, 2011). Traumatic brain injury affecting the posterior cingulate have been reported to result in difficulty with sustained attention and switching from automatic to controlled responses leading to perseverative behavior (Leech & Sharp, 2014). Altered attention has been reported in CSBD (Savard et al., 2021) and one could speculate that impaired modification in behavioral changes in responses to environmental demands contribute to sexual behaviors

being pursued regardless of adverse consequences. Importantly, however, the posterior cingulate cortex is implicated in reward processing (Oldham et al., 2018), which aligns with the theory that reward-related processes are impacted in CSBD (Liberg et al., 2022; Schmidt et al., 2017; Voon et al., 2014). While the exact role of the posterior cingulate in CSBD remains to be clarified, results from previous studies and the present study are in line with the notion that CSBD is associated with brain alterations in areas implicated in sensitization, habituation, impulse control, and reward processing.

### Limitations

The cross-sectional nature of the study does not allow for inference regarding causality. Although structural abnormalities are commonly associated with functional impairments (Burzynska et al., 2012; Joshi et al., 2016), functional MRI studies need to be conducted to enable conclusions about the direct functional involvement of implicated brain areas and their association with CSBD. In addition, in order to detect focal and smaller effects, e.g., in cortical thickness differences, replication in larger samples is needed. Furthermore, the cohort consisted solely of self-referred males. Although female patients seek treatment for CSBD, they are the minority in these patients groups, and future studies should focus on the recruitment of females.

The result that group differences are not significant after correcting for depression symptom levels needs to be interpreted with caution, as depression is highly correlated to CSBD (Antons & Brand, 2021); compulsive sexual behavior has been suggested to be the result of coping mechanism compensating for negative affective states, but distress and depressive symptoms can potentially also be caused by out-of-control sexual fantasies and behaviors (Briken, 2020). While both mechanisms may contribute, they cannot be separated in this study. In addition, we did not evaluate the prevalence of comorbid paraphilic disorder in this study. Although our previous study in an independent sample of CSBD patients, notably from the same catchment area, indicated a paraphilic disorder prevalence of only 8.3% (Hallberg et al., 2020), future studies should investigate the impact of paraphilic disorder, as well as other comorbidities, such as mood disorders, ADHD, and autism, on MRI-based outcomes. Finally, several participants have been recruited during the COVID-19 pandemic. Although studies report no significant changes in pornography use during the COVID-19 pandemic (Grubbs, Perry, Grant Weinandy, & Kraus, 2022; Koós, Demetrovics, Griffiths, & Bóthe, 2022), it remains unknown if sexual behavior changed in our sample, e.g. increased pornography use and reduced number of sexual partners. Thus, it remains to be investigated whether our results are more generalizable to CSBD subgroups with high-frequency pornography use (Antons & Brand, 2021). Nevertheless, our study cohort represented an ecologically valid clinical sample of patients with CSBD.

### CONCLUSIONS

Our findings suggest that CSBD is associated with structural brain differences. This study provides valuable insights into a largely unexplored field of clinical relevance and encourages further clarifications of the neurobiological mechanisms underlying CSBD, which is a prerequisite for improving future treatment outcomes. The findings may also contribute to ongoing discussion around whether the current classification of CSBD as an impulse-control disorder is reasonable.

*Funding sources:* This work was supported by Karolinska Institutet's Research Foundation Grants (2016 and 2017; CA), the Swedish Research Council (2020-01183; JJ, CA), the Swedish Prison and Probation System, and through a regional agreement between Umeå University and Region Västerbotten (ALF) (RV-941086).

*Authors' contribution:* CA was principal investigator and designed the study. CA collected MRI and behavioral data. PG performed the majority of analyses and wrote the first draft of the manuscript. KJÖ, SA, CD, and MI contributed to study design and with clinical advice. KJÖ, JJ, and JS provided important intellectual input and contributed to manuscript writing. JS recruited and screened patients for eligibility and contributed to data collection. All authors had full access to all data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis. All authors have reviewed the manuscript, provided intellectual input, and approved the submission of the manuscript.

*Conflict of interest:* CA is employed by Quantify Research (consultancy work unrelated to the present work). The authors report no financial or other relationship relevant to the subject of this article.

*Acknowledgements:* We thank the study nurses, medical, and administrative staff at ANOVA for their support in data collection and study organization, Christoffer Rahm for discussions during the study design phase, and Christian Mannfolk for his help in the recruitment of HC participants.

### SUPPLEMENTARY MATERIAL

Supplementary data to this article can be found online at <https://doi.org/10.1556/2006.2023.00008>.

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