Reduced Orbitofrontal Gray Matter Concentration as a Marker of Premorbid Childhood Trauma in Cocaine Use Disorder

Keren Bachi, Muhammad A. Parvaz, Scott J. Moeller, Gabriela Gan, Anna Zilverstand, Rita Z. Goldstein and Nelly Alia-Klein*

Departments of Psychiatry and Neuroscience, Icahn School of Medicine at Mount Sinai, New York, NY, United States, Department of Psychiatry, Stony Brook University School of Medicine, Stony Brook, NY, United States, Department of Psychiatry and Psychotherapy, Central Institute of Mental Health, Medical Faculty Mannheim, Heidelberg University, Mannheim, Germany

Background: Childhood trauma affects neurodevelopment and promotes vulnerability to impaired constraint, depression, and addiction. Reduced gray matter concentration (GMC) in the mesocorticolumbic regions implicated in reward processing and cognitive control may be an underlying substrate, as documented separately in addiction and for childhood trauma. The purpose of this study was to understand the contribution of childhood maltreatment to GMC effects in individuals with cocaine use disorder.

Methods: Individuals with cocaine use disorder were partitioned into groups of low vs. high childhood trauma based on median split of the total score of the Childhood Trauma Questionnaire (CTQ; CUD-L, N = 23; CUD-H, N = 24) and compared with age, race, and gender matched healthy controls with low trauma (N = 29). GMC was obtained using voxel-based morphometry applied to T1-weighted MRI scans. Drug use, depression and constraint were assessed with standardized instruments.

Results: Whole-brain group comparisons showed reduced GMC in the right lateral orbitofrontal cortex (OFC) in CUD-H as compared with controls (cluster-level $p_{FWE-corr} < 0.001$) and CUD-L (cluster-level $p_{FWE-corr} = 0.035$); there were no significant differences between CUD-L and controls. A hierarchical regression analysis across both CUD groups revealed that childhood trauma, but not demographics and drug use, and beyond constraint and depression, accounted for 37.7% of the variance in the GMC in the right lateral OFC ($p < 0.001$).

Conclusions: Beyond other contributing factors, childhood trauma predicted GMC reductions in the OFC in individuals with cocaine use disorder. These findings underscore a link between premorbid environmental stress and morphological integrity of a brain region central for behaviors underlying drug addiction. These results further highlight the importance of accounting for childhood trauma, potentially as a factor predisposing to addiction, when examining and interpreting neural alterations in cocaine addicted individuals.

Keywords: cocaine dependence, childhood maltreatment, gray matter, voxel-based morphometry, addiction
INTRODUCTION

The child maltreatment-morbidity connection makes up for a large fraction of the global burden of all disease and premature mortality (Shalev et al., 2013, 2016; Chen et al., 2016). For example, childhood maltreatment (and associated adversity) accounts for 64% of the population-attributed risk for addiction to illicit drugs (Dube et al., 2003), potentially predisposing vulnerable individuals to substance use disorders and comorbid psychopathology. Specifically, childhood maltreatment increased the likelihood of lifetime substance use disorders, including the initiation of substance use especially during early adolescence (Dube et al., 2003; Shin et al., 2010). Childhood maltreatment has also been implicated in greater substance-cue reactivity and enhanced substance-use symptom severity inclusive of interpersonal and substance-related financial problems and risk of relapse (Westermeyer et al., 2001; Schumacher et al., 2006; Van Dam et al., 2014).

The potential impact of childhood maltreatment on substance use are elucidated via examination of how early exposures to social-relational stress may affect neurodevelopment in prefrontal cortical regions (De Bellis et al., 2002; Hanson et al., 2010; Hart and Rubia, 2012; De Bellis and Zisk, 2014; Paquola et al., 2016; Teicher et al., 2016) crucial for the development of highest-order cognitive abilities (Arnsten, 2009). Exposure to even brief periods of intense stress causes significant structural remodeling within the rodent prefrontal cortex (see Holmes and Wellman, 2009 for review). This structural remodeling involves impairments in orbitofrontal cortex (OFC)-dependent cognitive functions via excessive stimulation of dopaminergic and noradrenergic receptors (see Arnsten, 2009 for review), thereby enhancing drug self-administration (see Miczek et al., 2004; Sinha, 2008 for reviews). Children who have experienced early maternal deprivation exhibit an accelerated development in prefrontal cortical-amygnda connectivity as measured with an fMRI task of viewing emotional faces and analyzed with a psychophysiological interaction analysis (Gee et al., 2013). Previous reports suggest that this accelerated brain development in humans parallels that in rodents (McEwen, 2008; Callaghan and Richardson, 2011) and that it is interpreted to reflect an ontogenetic adaptation to adversity before adulthood (Anisman et al., 1998; de Kloet and Oitzl, 2003; Callaghan and Richardson, 2011). Such neurodevelopmental structural remodeling in mesocorticostral regions and specifically in the OFC, may be key in development of substance use disorders since these regions mediate behaviors of cognitive and impulse control and reward processing (e.g., enhanced drug over natural reward salience concomitant with a decrease in inhibitory control), as highlighted by the Impaired Response Inhibition and Salience Attribution (iRISA) model (Goldstein and Volkow, 2002, 2011).

Studies of adolescents and adults without psychopathology reported widespread reductions in gray matter (GM) volumes in several brain regions in childhood trauma victims (e.g., Tomoda et al., 2009: visual association cortices; Edmiston et al., 2011: corticostral-limbic regions; Cohen et al., 2006: anterior cingulate cortex and caudate nuclei; Thomaes et al., 2010: anterior cingulated cortex, OFC and hippocampus; Rinne-Albers et al., 2017: anterior cingulate and superior temporal cortices). There have been previous efforts to disentangle effects of childhood trauma and psychiatric disorders on brain structure. Overall GM volume, and volumes of regions in the prefrontal cortex, including the anterior cingulate cortex and inferior frontal gyrus, were negatively correlated with sexual abuse severity in individuals with psychotic disorders (Sheffield et al., 2013). Similarly, hippocampal volume was reduced in women with a history of prepubertal physical and/or sexual abuse and current unipolar major depressive disorder (Vytilingam et al., 2002; Frodl et al., 2010; Opel et al., 2014). The amygdala has also been implicated in childhood trauma and mood and anxiety disorders (Nemeroff, 2016). Functional MRI studies found that amygdala responsiveness to threat-related facial expressions was strongly associated with Childhood Trauma Questionnaire (CTQ) scores, an index of childhood adversity, an effect not attributed to recent life stressors, current depression or anxiety or sociodemographic factors (Dannlowski et al., 2012). A meta-analysis of neuroanatomy in children/adolescents and adults with a history of childhood maltreatment and comorbid psychiatric disorders (e.g., anxiety, depression) reporting the prefrontal cortex, hippocampus, parahippocampus, striatum and OFC as regions showing structural alterations associated with childhood trauma (Lim et al., 2014). The most consistent brain structural abnormalities in this meta-analysis were decreases in GM volumes in the right OFC-temporo-limbic and left inferior frontal regions, as attributed to the typically observed deficits in cognitive control in this population (Lim et al., 2014).

Drug addiction is associated with reductions in GM volumes across several of the same regions affected by childhood maltreatment (Hall et al., 2015). Decreased GM density in mesocorticoloimbic regions in adults with substance use disorders has been attributed to genetics (Alia-Klein et al., 2011) and factors including the severity of drug use (Sim et al., 2007; Moeller et al., 2016), craving (Wrase et al., 2008; Morales et al., 2015), abstinence (Connolly et al., 2013), or co-morbid depression (Goodkind et al., 2015).

The underlying physiology of decreased GM in substance use disorders may be associated with the acute impact of drugs (as seen with psychostimulants) on inducing large and fast increases in extracellular dopamine, which mimic yet surpass those induced by physiological dopamine cell firing, with marked decreases in dopamine D2 receptor availability and dopamine release with chronic drug abuse (Volkow et al., 2009). This decrease in dopamine function is associated with reduced regional activity in the OFC (Volkow et al., 2009). In addition, in the OFC psychostimulants have been associated with long-lasting decreases in dendritic density (Crombag et al., 2005) likely reflecting decreases in neuronal plasticity (Schoenbaum and Shaham, 2008), which is consistent with decreased GM in the OFC in human substance use disorders (Franklin et al., 2002; Ersche et al., 2013; Hall et al., 2015). Childhood trauma, however, may also be a contributing factor impacting structural and functional deficits in this region in ways that heighten vulnerability to impulsivity, depression symptoms, and substance use disorders in adulthood (Jaffee et al., 2004;
Goldstein and Volkow, 2011; McCrory et al., 2011; Puetz and McCrory, 2015). Indeed, childhood adversity enhances the development of antisocial personality traits including reduced constraint, and may lead to chronic depression in adulthood (Beach et al., 2010; Hengartner et al., 2015; Khan et al., 2015; Negele et al., 2015; Thibodeau et al., 2015; Nelson et al., 2017). Low constraint (as a measure of impulsivity) and depression have been identified as risk factors for substance use and are central to the substance use disorders phenotype (Crews and Boettiger, 2009; de Wit, 2009; Green et al., 2012).

Understanding the neural mechanisms that may underlie the impact of childhood maltreatment on substance use disorders could contribute to pre-empting the adverse consequences of exposure and informing neural investigations of addictions, individualized addiction interventions, and to address the long-lasting consequences of childhood trauma exposure (Heilig et al., 2016; Teicher et al., 2016).

The purpose of this study is to examine the potential association between childhood maltreatment and gray matter concentration (GMC) effects in individuals with cocaine use disorder. We further explored, as a post hoc analysis, the contribution of the demographic and clinical variables that could contribute to the extent of the observed structural deficits. Since a high incidence of early trauma has been observed in individuals with cocaine use disorders (iCUD; Back et al., 2000; Khoury et al., 2010), and because both substance use disorders and maltreatment are associated with OFC GM deficits, and considering the impact of psychostimulants on OFC neuroplasticity and psychiatric comorbidity, we hypothesized that: (1) iCUD who have history of childhood trauma will exhibit reduced GMC of the OFC; and (2) childhood trauma will explain OFC GM variability in iCUD above and beyond the contribution of demographics, drug effects, and constraint and depression symptoms.

MATERIALS AND METHODS

Participants
Seventy six right-handed individuals were recruited by advertisements in local newspapers and partitioned into three groups: 29 healthy controls (CON), 23 iCUD with low childhood trauma (CUD-L), and 24 iCUD with high childhood trauma (CUD-H). All participants provided written informed consent in accordance and approval of the Institutional Review Board of the Icahn School of Medicine at Mount Sinai. All participants were healthy, were not taking any medications, and were excluded if they had contraindications to the magnetic resonance (MR) imaging environment (e.g., metal in the body or claustrophobia), history of head trauma or loss of consciousness (>30 min), other neurological disease, history of major medical conditions (cardiovascular, endocrinologic, oncologic, or autoimmune diseases), major psychiatric disorders (other than cocaine dependence and alcohol and past cannabis abuse for the CUD group and/or nicotine dependence for both groups), and urine positive (Biopsych; Biopsych Triage, San Diego, CA, USA) for psychoactive drugs or their metabolites (phencyclidine, benzodiazepines, amphetamines, cannabis, opiates, barbiturates and inhalants) except for cocaine in iCUD. With the exception of cocaine, alcohol, cannabis and nicotine, current use or dependence on other drugs was denied. See Table 1 for comparison of demographic, drug use and clinical characteristics between the groups.

Measures
Substance Use and Psychiatric Assessment
The Structured Clinical Interview for DSM-IV Axis I Disorders (research version; Ventura et al., 1998), the Addiction Severity Index (McLellan et al., 1992), Cocaine Selective Severity Assessment Scale (assessing withdrawal symptoms; Kampman et al., 1998), Severity of Dependence Scale (Gossop et al., 1992), and Cocaine Craving Questionnaire (Tiffany et al., 1993) were used to assess substance use and psychiatric history.

Childhood Trauma Questionnaire
Childhood maltreatment was assessed in all 76 participants with the brief screening version of the CTQ (Bernstein et al., 2003). Due to the high prevalence of co-occurring maltreatment types (Scher et al., 2004), and the association of exposure to multiple types of trauma with poor health outcomes (Huang et al., 2012; Chen et al., 2016; Shalev et al., 2016), we used the CTQ total score, calculated as the sum of scores from each maltreatment subtype, as a continuous measure of individual history of maltreatment (Elton et al., 2014). Following Elton et al. (2014), low vs. high childhood trauma was determined based on a median split of the total CTQ score (= 43; range: 25–125) among all iCUD. Consistent with the median split, the current results showed higher rates of childhood trauma among the CUD-H as compared with the other two study groups and with normative data from a community sample (Scher et al., 2004), where median split of CTQ total scores were = 32 for both sexes combined (Normative data < CUD-H, t = 11.4, p < 0.001; Normative data = CUD-L, t = 1.08, p = 0.29; Normative data < CON, t = 2.4, p = 0.025).

Depression Symptoms
Beck Depression Inventory-II (BDI; Beck et al., 1996) was used to assess the severity of depressive symptoms occurring over the 2 weeks preceding the study. The BDI-II is based on the DSM-IV criteria (American Psychiatric Association, 2000), and is considered a valid and reliable instrument for depression screening in the general population (Beck et al., 1996; Vanheule et al., 2008). The items are summed to give a total score (range 0–63). A higher score on the BDI-II denotes more severe depression.

Personality
The Multidimensional Personality Questionnaire (MPQ) Brief Form (Patrick et al., 2002; Tellegen and Waller, 2008) was used as a trait measure of constraint. Its scales represent three higher order factors and 10 primary personality dimensions. The higher order factor of constraint (reverse-scored impulsivity and behavioral restraint) embodies both affect and temperament.
TABLE 1 | Demographic, drug use and clinical variables.

| Characteristics | Controls N = 29 | Cocaine users low childhood trauma N = 23 | Cocaine users high childhood trauma N = 24 |
|-----------------|----------------|------------------------------------------|------------------------------------------|
| **Demographics** |                |                                          |                                          |
| Age             | 41.9 ± 7.9     | 47.0 ± 7.9                                | 45.8 ± 7.8                               |
| Education*      | 14.6 ± 1.9     | 13.1 ± 1.8                                | 12.7 ± 1.6                               |
| Gender          | 20 men (69.0%) | 20 men (87.0%)                            | 18 men (76.0%)                           |
| Race            | 22 African American (75.9%) | 19 African American (82.6%) | 16 African American (66.7%)             |
| **Drug use lifetime years/age of initiation/years of heavy use** |                |                                          |                                          |
| Nicotine        | 16.7 ± 15.5/16.3 ± 2.2/NA | 21.8 ± 11.2/15.4 ± 4.1/NA | 22.3 ± 11.9/17.4 ± 6.7/NA               |
| Alcohol         | 9.3 ± 10.1/17.3 ± 5.8/1.8 ± 1.2 | 14.8 ± 12.1/15.3 ± 3.6/3.9 ± 2.9 | 17.3 ± 10.1/13.9 ± 2.8/4.1 ± 3.6         |
| Cocaine         | NA             | 18.5 ± 8.8/20.9 ± 5.3/3.8 ± 3.8            | 16.8 ± 9.5/20.7 ± 6.3/6.9 ± 7.8         |
| Cannabis**      | 0.6 ± 2.1/19.0 ± 2.9/0.6 ± 0.3 | 5.9 ± 7.5/17.2 ± 1.4/1.2 ± 0.5 | 10.3 ± 12.2/20.1 ± 5.3/1.5 ± 1.7       |
| Severity of cocaine dependence | NA            | 4.0 ± 5.1                                 | 4.8 ± 5.2                                |
| Cocaine craving | NA             | 13.3 ± 10.7                               | 12.8 ± 14.0                              |
| Cocaine withdrawal | NA          | 15.9 ± 13.0                              | 16.1 ± 12.8                              |
| Cocaine: days since last use* | NA            | 4.0 ± 4.8                                | 1.7 ± 1.2                                |
| Current alcohol | 0              | 2 abuse                                   | 11 dependence                            |
| **Clinical**    |                |                                          |                                          |
| Childhood trauma* | 34.2 ± 5.1    | 33.2 ± 5.4                                | 57.9 ± 11.1                              |
| Constraint*     | 82.3 ± 8.2     | 82.6 ± 9.4                                | 57.1 ± 14.1                              |
| Depression*     | 2.9 ± 4.2      | 5.8 ± 6.8                                 | 10.1 ± 8.5                               |

Notes: *p < 0.001 (Controls < Cocaine users low < Cocaine users high); **p < 0.001 (Controls < Cocaine users low < Cocaine users high); 1 <p < 0.001, 1 p = 0.055 (Controls = Cocaine users low < Cocaine users high; for constraint Controls = Cocaine users low < Cocaine users high); NA: not applicable; 1 N = 10; 2 N = 21; 3 N = 23; 4Reported number of days since last cocaine use prior to urine test (N = 7 for each CUD group); 5 N = 23; 6The instruments used to derive drugs use/clinical measures: the Structured Clinical Interview for DSM-IV Axis I Disorders (research version; Ventura et al., 1998), the Addiction Severity Index (McLellan et al., 1992), Cocaine Selective Severity Assessment Scale (assessing withdrawal symptoms; Kaplan et al., 1998), Severity of Dependence Scale (Gossop et al., 1992), and Cocaine Craving Questionnaire (Tiffany et al., 1993). Childhood Trauma Questionnaire (Bernstein et al., 2003) was used to assess childhood trauma. Beck Depression Inventory-II (BDI; Beck et al., 1996) used to assess the severity of depressive symptoms occurring over the 2 weeks preceding the study. The Multidimensional Personality Questionnaire (MPQ) Brief Form (Patrick et al., 2002; Tellegen and Waller, 2008) was used as a trait measure of constraint.

Voxel-Based Morphometry

Voxel based morphometry (VBM), which enables an unbiased voxel-by-voxel comparison of cortical volumes (Whitwell, 2009), was used to quantify regional GMC in study participants.

M1 Acquisition and Processing

T1-weighted anatomical images were acquired on a 3T Skyra (Siemens, Erlangen, Germany) using vendor provided 32 channel head coil with a 3D MPRAGE sequence [FOV 256 × 256 × 179 mm³, 0.8 mm isotropic resolution, TR/TE/T1 = 2400/2.07/1000 ms, flip angle 8° with binomial (1, −1) fat saturation, bandwidth 240 Hz/pixel, echo spacing 7.6 ms, inplane acceleration (GRAPPA) factor of 2, total acquisition time ~7 min]. Structural T1 images were preprocessed using the “HCP PreFreeSurfer structural pipeline” (based on FSL 5.0.6 and FreeSurfer 5.3.0-HCP) to align the origin to the anterior−posterior commissure line, correct image distortions (bias-field inhomogeneities), and to normalize T1-weighted images to the MNI space using a FLIRT affine linear and then a FNIRT nonlinear registration. FSL’s image registration tools were applied to register the native structural scans to the MNI template. FLIRT (FMRIB’s linear image registration tool) was used for linear (affine; translate, rotate, zoom and shear) registration of images. For instances where the linear transform did not achieve adequate registration, FNIRT (FMRIB’s non-linear image registration tool) was used for non-linear registration.

Voxel-Based Morphometry Analysis

All VBM procedures were conducted using Statistical Parametric Mapping (SPM12 and SPM8; Wellcome Trust Centre for Neuroimaging, UK), a program running through Mathworks (Matlab 7.5; MathWorks, Natick, MA, USA) with VBM8 toolbox (Gaser, C, University of Jena, Department of Neurology and Psychiatry, Germany). Total cortical GMC measures were obtained by unified segmentation (Ashburner and Friston, 2005) of each T1 image into GM, white matter and cerebrospinal fluid. A hidden Markov random field was used to maximize segmentation accuracy (Cuadra et al., 2005). The segmentation was then visually inspected for accuracy. Unmodulated normalized GM images (which are GMC, and thereby differ from GM volume that consists of modulated cerebrospinal fluid. A hidden Markov random field was used to maximize segmentation accuracy (Cuadra et al., 2005). The segmentation was then visually inspected for accuracy. Unmodulated normalized GM images were then smoothed using an 8-mm full-width at half-maximum Gaussian kernel.

Although we had a region specific hypothesis for the OFC, we conducted whole-brain analyses, recognizing that other regions may also be involved (e.g., limbic). We performed a one-way analysis of covariance (ANCOVA) at the whole-brain level for between group comparisons to assess the effect

---

1http://fsl.fmrib.ox.ac.uk/fsl/flirt/

2https://fsl.fmrib.ox.ac.uk/fsl/fslwiki/FNIRT

3www.fil.ion.ucl.ac.uk/spm

4http://www.neuro.uni-jena.de/vbm/
of childhood trauma on GMC with education, age, and total intracranial volume (TIV) included in the models as covariates. We applied a search threshold of \( p < 0.01 \), adjusted for cluster size, with an extent threshold of 100 contiguous voxels. Only voxels which survived correction for multiple comparisons in the entire volume using the false discovery rate (FDR) method \( (p < 0.05) \) were considered as significant (Genovese et al., 2002). To further examine the main effect and investigate the contribution of childhood trauma, we conducted a planned follow-up comparison between high and low trauma among CUD (CUD-L vs. CUD-H) using whole-brain statistical parametric maps. Here we controlled for multiple comparisons by using the more stringent voxel-level family wise error (FWE) corrected method \( (p < 0.05) \) at the cluster level. The contribution of childhood trauma to OFC GMC within the entire CUD group (the only region that differed between CUD-H and CUD-L) above and beyond that of the other variables (i.e., demographics and cocaine use, that have not been included in the previous steps of analysis) was evaluated by a multiple regression analysis, conducted in SPSS version 23.0 (SPSS Inc., Chicago, IL, USA). This post hoc analysis was an additional step to verify that the reported whole brain effects were not driven by other variables that had significant differences between the groups of interest. The first block of this hierarchical regression model included age, education, TIV, and lifetime years of drug use (cocaine, alcohol and cannabis). The second block added constraint and depression, and the third block added the CTQ total scores. To further characterize the potential psychological and behavioral implications of GMC reductions across the whole sample, we examined correlations between the GMC variance in the selected region of interest with constraint and depression.

**RESULTS**

For descriptive characteristics of the sample see Table 1. The three groups were matched on gender, race and age, but differed in education (CON > CUD-L = CUD-H, \( p < 0.001 \); used as a covariate in all analyses). Lifetime drug use (in years) of nicotine, alcohol, cocaine, and cannabis, and age of drug use initiation, and years of heavy use (besides nicotine), as well as severity of cocaine dependence, cocaine craving, withdrawal symptoms, time since last cocaine use and current alcohol dependence did not differ between the CUD groups. Yet, iCUD with increased childhood trauma severity reported reduced constraint (CON = CUD-L > CUD-H, \( p < 0.001 \)) and a trend for increased depression symptoms (CON = CUD-L < CUD-H, \( p = 0.055 \)).

The whole-brain SPM analysis showed a significant group main effect in the right lateral OFC, the bilateral middle temporal gyri, and the dorsal anterior cingulate gyrus (Table 2 and Figure 1). Follow-up pairwise whole-brain comparisons showed significantly reduced GMC in the right lateral OFC in the CUD-H as compared to CON and even as compared to CUD-L. In contrast, CUD-L and CON groups did not differ (Table 2 and Figure 2). Additionally, reduced GMC in the bilateral middle temporal gyri were observed in CUD-H as compared to CON. Although there was a main effect in the dorsal anterior cingulate gyrus, the more stringent post hoc between group comparisons were not significant. An alcohol dependence diagnosis status was not associated with the GMC results, nor did correlations between each drug use variable in Table 1 (lifetime years, age initiation and years of heavy use) with GMC in our main effect regions reached significance.

Across all CUD, the post hoc analysis using hierarchical regression demonstrated that the continuous measure of childhood trauma accounted for 37.7% of variance in GMC in the right lateral OFC (third block; \( \Delta R^2 = 0.24, p < 0.001 \); Table 3), while constraint and depression explained 10.2% (second block; \( \Delta R^2 = 0.20, p = 0.012 \), above and beyond the effects of age, education, TIV and lifetime drug use variables (first block \( p = 0.853 \)). Thus, beyond demographics and drug use, and beyond constraint and depression, the experience of childhood trauma related to OFC GMC reduction (Figure 3).

**TABLE 2** | Main effect and between group comparisons of whole-brain gray matter concentration.

| Region                        | BA | Side | Voxel | Peak Z | Cluster P Value | x    | y    | z    |
|------------------------------|----|------|-------|--------|----------------|------|------|------|
| **Group main effect**        |    |      |       |        |                |      |      |      |
| Lateral orbitofrontal gyrus  | 11 | R    | 883   | 4.05   | 0.020          | 29   | 47   | −14  |
| Middle temporal gyrus        | 21 | R    | 1069  | 3.64   | 0.014          | 59   | 3    | −27  |
| Middle temporal gyrus        | 22 | L    | 1010  | 3.86   | 0.014          | −63  | −15  | −8   |
| Dorsal anterior cingulate gyrus | 32 | L    | 1108  | 3.55   | 0.014          | −11  | 33   | 19   |
| **Between-groups comparisons** |    |      |       |        |                |      |      |      |
| CUD-L > CUD-H                |    |      |       |        |                |      |      |      |
| Lateral orbitofrontal gyrus  | 11 | R    | 1537  | 3.49   | 0.035          | 30   | 48   | −12  |
| CON > CUD-H                  |    |      |       |        |                |      |      |      |
| Lateral orbitofrontal gyrus  | 47 | R    | 15,097| 4.53   | <0.001         | 39   | 41   | −15  |
| Middle temporal gyrus        | 21 | R    | 420   | 4.20   | 0.014          | 59   | 3    | −27  |
| Middle temporal gyrus        | 22 | L    | 4802  | 4.31   | <0.001         | −63  | −15  | −8   |

**Note:** \(^1\)Analyses are one-way ANCOVAs (controlling for age, education and total intracranial volume), thresholded at \( p < 0.01 \), uncorrected, and an extent threshold of 100 contiguous voxels. Only voxels which survived correction for multiple comparisons in the entire volume using the false discovery rate (FDR) method, \( p < 0.05 \), were considered significant; \(^2\)Analyses are pairwise between-groups comparisons of gray matter concentration (controlling for education, age, and total intracranial volume), corrected for multiple comparisons using the voxel-level family wise error (FWE) corrected method, \( p < 0.05 \); CON, controls; CUD-L, Individuals with cocaine use disorder with low childhood trauma; CUD-H, Individuals with cocaine use disorder with high childhood trauma; BA, Brodmann Area; R, right; L, left.
Correlations of GMC across the whole sample revealed that the decreased GMC in the right lateral OFC was also significantly associated with increased depression ($r = -0.382, p = 0.001$) and decreased constraint ($r = 0.329, p = 0.004$; Figure 4).

DISCUSSION

This study was conducted to discern the morphological and comorbid symptom properties of childhood maltreatment in iCUD. Consistent with our hypotheses, the main results of this study are that: (1) GMC of the right lateral OFC is reduced with greater childhood trauma in iCUD such that the most abnormality was observed in iCUD with high trauma, followed by iCUD with low trauma, and then by controls; (2) above and beyond the effects of demographics, lifetime drug use, constraint and depression, childhood trauma has a noticeable and significant effect on OFC GMC reduction in iCUD.

Our main results are consistent with previous findings demonstrating brain’s GM decreases as related to childhood trauma in individuals with a history of childhood maltreatment and comorbid psychiatric disorders. For example, an extensive meta-analysis (comprising 331 individuals with a history of childhood maltreatment and psychiatric comorbidities and 362 comparison subjects) showed that deficits in the right OFC/superior temporal gyrus were the most robust GM abnormalities in non-addicted individuals (Lim et al., 2014). Reduced GMC in the middle temporal gyrus was observed in CUD-H as compared to CON (but not CUD-L) as well as in subjects with childhood verbal aggression trauma measured in early adulthood (Tomoda et al., 2011). Our main results are also consistent with a morphometric analysis of 148 healthy individuals (without any history of psychiatric disorders) where high CTQ scores were associated with reduced GM volumes in the OFC and anterior cingulate gyrus (Dannlowski et al., 2012). In contrast to this latter study, we did not report significant results for the hippocampus, insula, and caudate. Of note is that in a previous study of addicted individuals as compared to controls, childhood trauma was similarly associated with GMC reductions in the hippocampus (and also in the parahippocampus and anterior fusiform gyrus) in individuals with alcohol (95.8% of the sample), cocaine (79.2%), and/or cannabis (77.8%) use disorders (Van Dam et al., 2014). These inconsistencies in results may be attributed to differences in the samples and methods used in these studies. Although smaller in size (our total sample size was 76 vs. 148 and 177 in the previous studies), we only included current actively drug using iCUD with nearly double lifetime years of cocaine use for our participants as compared to the...
It is possible that prefrontal cortical GM volumes, which mature later during childhood and adolescence (Sowell et al., 2003) at a vulnerable developmental timing that coincides with drug experimentation, predispose individuals to substance abuse risk (Weiland et al., 2014). It is also possible that prenatal drug exposure (to alcohol and/or cigarette smoking), which are associated with reduced OFC cortical thickness, increases substance use during adolescence (Kühn et al., 2016). Both factors may be more pronounced with childhood maltreatment.

Previous research showed that childhood maltreatment contributes to a high prevalence of co-morbid personality...
disorders in addicted populations (Bernstein et al., 1998). Indeed, in our study CUD-H reported less constraint and higher depression than CUD-L. Such greater symptomatology in CUD-H may confer a greater severity of substance use disorders and an increased risk for relapse, as associated with lower GMC in the OFC. Thus, iCUD with histories of childhood trauma may represent a clinically and biologically distinct subtype. Recognition of this distinction may be essential in determining the biological bases of this subtype and guide individualized treatment (Nanni et al., 2012; Teicher and Samson, 2013; Teicher et al., 2016).

These results should be considered in light of the study’s limitations. A major limitation is in absence of a control group who experienced childhood trauma (although note that our healthy control group reported higher levels of childhood trauma than the normative CTQ data). Our three group study design is similar to previous studies that compared brain’s GM following early life adversity in two groups with psychiatric comorbidity (e.g., major depression and psychotic disorder patients) with and without early trauma and a healthy control group without trauma (e.g., Vythilingam et al., 2002; Sheffield et al., 2013). Nevertheless, future studies including a control group with high childhood trauma and no substance use disorder would be critical to attribute the observed low OFC GMC results specifically to the trauma or to an interaction of trauma with substance use. As well, it remains to be studied whether individuals with childhood trauma with comorbid substance use disorders and major depression represent a qualitatively different subtype than those subjects included in the current study. In addition, given that strong statistical effects were not anticipated (since we were measuring effects of events that probably have occurred decades ago), we used a search threshold of 0.01 uncorrected together with different multiple correction methods (FDR and FWE). Although these analytical strategies enabled us to capture the effects of maltreatment that contributed evident variance beyond demographics and drug use, and comorbid symptomatology, recent reports point to inaccuracy of FDR correction methods for voxel-wise analyses as compared with topological FWE or FDR control (Chumbley et al., 2010); therefore, these results need to be replicated in a larger sample and with more stringent statistical thresholds. Lastly, childhood trauma was retrospectively assessed and self-reported. This approach is broadly accepted as reliable and valid (Dill et al., 1991; Hardt and Rutter, 2004; Anda et al., 2006); nevertheless, future prospective and well documented assessments need to account for type, timing and duration of childhood trauma, since specific neurobiological consequences of extent of childhood trauma have been associated with unique
windows of vulnerability in various brain regions (Teicher et al., 2006; Andersen et al., 2008; Teicher and Samson, 2013; Pechtel et al., 2014; Nemeroff, 2016).

CONCLUSION

Above and beyond effects of demographics, lifetime drug use and constraint and depression, our results point to a noticeable and significant contribution of adverse childhood experiences, predating drug use, to GMC reduction of the OFC in cocaine use disorders. These findings suggest a link between premorbid environmental stress and OFC morphology in addicted individuals, and highlight the importance of accounting for childhood trauma in future examination of neuroanatomy in substance use. Early trauma may result in a different addiction-related phenotype, due to the potency of the effects of maltreatment on the developing brain. Although MRI scans are not routinely used in treatment, these findings underscore individual variance within the population of substance users and highlight the importance of individualized interventions to ameliorate the possible consequences of childhood trauma exposure for addicted individuals.

ETHICS STATEMENT

This study was carried out in accordance with the recommendations of the Institutional Review Board of the Icahn School of Medicine at Mount Sinai with written informed consent from all subjects. All subjects gave written informed consent in accordance with the Declaration of Helsinki. The protocol was approved by the Institutional Review Board of the Icahn School of Medicine at Mount Sinai.

AUTHOR CONTRIBUTIONS

KB, MAP, SJM, GG, AZ, RZG and NA-K (all authors) fulfilled the following four criteria: (1) substantial contributions to the conception or design of the work; or the acquisition, analysis, or interpretation of data for the work; (2) drafting the work or revising it critically for important intellectual content; (3) final approval of the version to be published; and (4) agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

FUNDING

The support of the following sources is gratefully acknowledged: National Institute on Drug Abuse (NIDA): T32-DA007135-31 (KB), K01DA037452 (SJM), R01DA041528 and U01DA041174 (RZG); National Institute of Mental Health (NIMH): R01MH0990134 (NA-K); Icahn School of Medicine at Mount Sinai: KL2TR001435 (MAP), Department of Environmental Medicine and Public Health support (KB); NWO Netherlands: Rubicon 446-14-015 (AZ).

REFERENCES

Alia-Klein, N., Parvaz, M. A., Woicik, P. A., Konova, A. B., Maloney, T., Shumay, E., et al. (2011). Gene disease interaction on orbitofrontal gray matter in cocaine addiction. Arch. Gen. Psychiatry 68, 283–294. doi: 10.1001/archgenpsychiatry.2011.10

American Psychiatric Association. (2000). Diagnostic and Statistical Manual of Mental Disorders. 4th Edn. Washington, DC: American Psychiatric Association.

Anda, R. F., Felitti, V. J., Bremner, J. D., Walker, J. D., Whitfield, C., Perry, B. D., et al. (2006). The enduring effects of abuse and related adverse experiences in childhood. A convergence of evidence from neurobiology and epidemiology. Eur. Arch. Psychiatry Clin. Neurosci. 256, 174–186. doi: 10.1007/s00406-005-0624-4

Andersen, S. L., Tomada, A., Vincow, E. S., Valente, E., Polcari, A., and Teicher, M. H. (2008). Preliminary evidence for sensitive periods in the effect of childhood sexual abuse on regional brain development. J. Neuropsychiatry Clin. Neurosci. 20, 292–301. doi: 10.1176/appi.neuropsych.20.3.292

Anisman, H., Zaharia, M. D., Meaney, M. J., and Merali, Z. (1998). Do early-life experiences, predating drug use, to GMC reduction of the OFC

Ashburner, J., and Friston, K. J. (2005). Unified segmentation. NeuroImage 26, 839–851. doi: 10.1016/j.neuroimage.2005.02.018

Back, S., Dansky, B. S., Coffey, S. F., Saladin, M. E., Sonne, S., and Brady, K. T. (2000). Cocaine dependence with and without posttraumatic stress disorder: a comparison of substance use, trauma history and psychiatric comorbidity. Am. J. Addict. 9, 51–62. doi: 10.1080/10550490050172227

Beach, S. R. H., Brody, G. H., Gunter, T. D., Packer, H., Wernert, P., and Philibert, R. A. (2010). Child maltreatment moderates the association of MAOA with symptoms of depression and antisocial personality disorder. J. Fam. Psychol. 24, 12–20. doi: 10.1037/a0018074

Beck, A. T., Steer, R. A., and Brown, G. K. (1996). Beck Depression Inventory-Second Edition Manual. San Antonio, TX: The Psychological Corporation.

Bernstein, D. P., Stein, J. A., and Handelmann, L. (1998). Predicting personality pathology among adult patients with substance use disorders: effects of childhood maltreatment. Addict. Behav. 23, 855–868. doi: 10.1016/s0306-4603(98)00072-0

Bernstein, D. P., Stein, J. A., Newcomb, M. D., Walker, E., Bogge, D., Ahluvalia, T., et al. (2003). Development and validation of a brief screening version of the Childhood Trauma Questionnaire. Child Abuse Negl. 27, 169–190. doi: 10.1016/s0145-2134(02)00541-0

Callaghan, B. L., and Richardson, R. (2011). Maternal separation results in early emergence of adult-like fear and extinction learning in infant rats. Behav. Neurosci. 125, 20–28. doi: 10.1037/a0022008

Chen, E., Turiano, N. A., Mroczek, D. K., and Miller, G. E. (2016). Association of reports of childhood abuse and all-cause mortality rates in women. JAMA Psychiatry 73, 920–927. doi: 10.1001/jamapsychiatry.2016.1786

Chumbley, J., Worsley, K., Flandin, G., and Friston, K. (2010). Topological FDR for neuroimaging. NeuroImage 49, 3057–3064. doi: 10.1016/j.neuroimage.2009.10.090

Cohen, R. A., Grieve, S., Hoth, K. F., Paul, R. H., Sweet, L., Tate, D., et al. (2006). Early life stress and morphometry of the adult anterior cingulate cortex and caudate nuclei. Biol. Psychiatry 59, 975–982. doi: 10.1016/j.biopsych.2005.12.016

Connolly, C. G., Bell, R. P., Foxe, J. J., and Garavan, H. (2013). Dissociated grey matter changes with prolonged addiction and extended abstinence in cocaine users. PLoS One 8:e59645. doi: 10.1371/journal.pone.0059645

Crews, F. T., and Boettiger, C. A. (2009). Impulsivity, frontal lobes and risk for addiction. Pharmacol. Biochem. Behav. 93, 237–247. doi: 10.1016/j.pbb.2009.04.018
Schoenbaum, G., and Shaham, Y. (2008). The role of orbitofrontal cortex in drug addiction: a review of preclinical studies. *Neuropsychopharmacology* 32, 2229–2237. doi: 10.1038/sj.np.1704316

Sinha, R. (2008). Chronic stress, drug use, and vulnerability to addiction. *Ann. N Y Acad. Sci.* 1141, 105–130. doi: 10.1196/annals.1415.030

Sowell, E. R., Peterson, B. S., Thompson, P. M., Welcome, S. E., Henkenius, A. L., and Toga, A. W. (2003). Mapping cortical change across the human life span. *Nat. Neurosci.* 6, 309–315. doi: 10.1038/nn1008

Teicher, M. H., and Samson, J. A. (2013). Childhood maltreatment and psychopathology: a case for ecopeptotypic variants as clinically and neurobiologically distinct subtypes. *Am. J. Psychiatry* 170, 1114–1133. doi: 10.1176/appi.ajp.2013.12070957

Teicher, M. H., Samson, J. A., Anderson, C. M., and Ohashi, K. (2016). The effects of childhood maltreatment on brain structure, function and connectivity. *Nat. Rev. Neurosci.* 17, 652–666. doi: 10.1038/nrn.2016.111

Teicher, M. H., Samson, J. A., Polcari, A., and McGreenery, C. E. (2006). Sticks, stones, and hurtful words: relative effects of various forms of childhood maltreatment. *Am. J. Psychiatry* 163, 993–1000. doi: 10.1176/appi.ajp.163.6.993

Tellegen, A., and Waller, N. G. (2008). “Exploring personality through test construction: development of the multidimensional personality questionnaire,” in *The SAGE Handbook of Personality Theory and Assessment: Volume 2—Personality Measurement and Testing: Personality Measurement and Assessment*, eds G. J. Boyle, G. Matthews and D. H. Saklofske (London: SAGE Publications Ltd), 261–292.

Thibodeau, E. L., Cicchetti, D., and Rogosch, F. A. (2015). Child maltreatment, impulsivity, and antisocial behavior in african-american children: moderation effects from a cumulative dopaminergic gene index. *Dev. Psychopathol.* 27, 1621–1636. doi: 10.1016/j.devps.2015.09.003

Thomaes, K., Dorrepaal, E., Draijer, N., de Ruiter, M. B., van Balkom, A. J., Smit, J. H., et al. (2010). Reduced anterior cingulate and orbitofrontal volumes in child abuse-related complex PTSD. *J. Clin. Psychiatry* 71, 1636–1644. doi: 10.4088/JCP.09m04754blu

Tiffany, S. T., Singleton, E., Haeretzen, C. A., and Henningsfield, J. E. (1993). The development of a cocaine craving questionnaire. *Drug Alcohol Depend.* 34, 19–28. doi: 10.1016/0376-8716(93)90042-o

Tomoda, A., Navaleta, C. P., Polcari, A., Sadato, N., and Teicher, M. H. (2009). Childhood sexual abuse is associated with reduced gray matter volume in visual cortex of young women. *Biol. Psychiatry* 66, 642–648. doi: 10.1016/j.biopsych.2009.04.021

Tomoda, A., Sheu, Y. S., Rabi, K., Suzuki, H., Navaleta, C. P., Polcari, A., et al. (2011). Exposure to parental verbal abuse is associated with increased gray matter volume in superior temporal gyrus. *Neuroimage* 54, S280–S286. doi: 10.1016/j.neuroimage.2010.05.027

Van Dam, N. T., Rando, K., Potenza, M. N., Tuit, K., and Sinha, R. (2014). Childhood maltreatment, altered limbic neurobiology, and substance use relapse severity via trauma-specific reductions in limbic gray matter volume. *JAMA Psychiatry* 71, 917–925. doi: 10.1001/jamapsychiatry.2014.0441

Vanheule, S., Desmet, M., Groenwynck, H., Rosseel, Y., and Fontaine, J. (2008). The factor structure of the beck depression inventory-II: an evaluation. *Assessment* 15, 177–187. doi: 10.1177/1073191107312161

Ventura, J., Liberman, R. P., Green, M. F., Shaner, A., and Mintz, J. (1998). Training and quality assurance with the Structured Clinical Interview for DSM-IV (SCID-I/P). *Psychiatry Res.* 79, 163–173. doi: 10.1016/S0165-1781(98)00038-9

Volkow, N. D., Fowler, J. S., Wang, G. J., Balser, R., and Telang, F. (2009). Imaging dopamine’s role in drug abuse and addiction. *Neuropharmacology* 56, 3–8. doi: 10.1016/j.neuropharm.2008.05.022
Vythilingam, M., Heim, C., Newport, J., Miller, A. H., Anderson, E., Bronen, R., et al. (2002). Childhood trauma associated with smaller hippocampal volume in women with major depression. *Am. J. Psychiatry* 159, 2072–2080. doi: 10.1176/appi.ajp.159.12.2072

Weiland, B. J., Korycinski, S. T., Soules, M., Zubieta, J.-K., Zucker, R. A., and Heitzeg, M. M. (2014). Substance abuse risk in emerging adults associated with smaller frontal gray matter volumes and higher externalizing behaviors. *Drug Alcohol Depend.* 137, 68–75. doi: 10.1016/j.drugalcdep.2014.01.005

Westermeyer, J., Wahmanholm, K., and Thuras, P. (2001). Effects of childhood physical abuse on course and severity of substance abuse. *Am. J. Addict.* 10, 101–110. doi: 10.1080/105504901750227769

Whitwell, J. L. (2009). Voxel- based morphometry: an automated technique for assessing structural changes in the brain. *J. Neurosci.* 29, 9661–9664. doi: 10.1523/JNEUROSCI.2160-09.2009

Wrase, J., Makris, N., Braus, D. F., Mann, K., Smolka, M. N., Kennedy, D. N., et al. (2008). Amygdala volume associated with alcohol abuse relapse and craving. *Am. J. Psychiatry* 165, 1179–1184. doi: 10.1176/appi.ajp.2008.07121877

**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2018 Bachi, Parvaz, Moeller, Gan, Zilverstand, Goldstein and Alia-Klein. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.