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Atypical Dorsolateral Prefrontal Activity in Female Adolescents With Conduct Disorder During Effortful Emotion Regulation

Nora Maria Raschle, Lynn Valérie Fehlbaum, Willeke Martine Menks, Anne Martinelli, Martin Prätzlich, Anka Bernhard, Katharina Ackermann, Christine Freitag, Stephane De Brito, Graeme Fairchild, and Christina Stadler

ABSTRACT

BACKGROUND: Conduct disorder (CD), which is characterized by severe aggressive and antisocial behavior, is linked to emotion processing and regulation deficits. However, the neural correlates of emotion regulation are yet to be investigated in adolescents with CD. Furthermore, it remains unclear whether CD is associated with deficits in emotional reactivity, emotion regulation, or both.

METHODS: We used functional magnetic resonance imaging to study effortful emotion regulation by cognitive reappraisal in 59 female adolescents 15 to 18 years of age (30 with a CD diagnosis and 29 typically developing (TD) control adolescents).

RESULTS: Behaviorally, in-scanner self-report ratings confirmed successful emotion regulation within each group individually but significant group differences in emotional reactivity and reappraisal success when comparing the groups (CD < TD). Functional magnetic resonance imaging results revealed significantly lower activation in left dorsolateral prefrontal cortex and angular gyrus in CD compared with TD adolescents during emotion regulation, but no group differences for emotional reactivity. Furthermore, connectivity between left dorsolateral prefrontal cortex and the bilateral putamen, right prefrontal cortex, and amygdala was reduced in CD compared with TD adolescents during reappraisal. Callous-unemotional traits were unrelated to neural activation, but these traits correlated negatively with behavioral reports of emotional reactivity.

CONCLUSIONS: Our results demonstrate reduced prefrontal brain activity and functional connectivity during effortful emotion regulation in female adolescents with CD. This sheds light on the neural basis of the behavioral deficits that have been reported previously. Future studies should investigate whether cognitive interventions are effective in enhancing emotion-regulation abilities and/or normalizing prefrontal and temporoparietal activity in female adolescents with CD.

Keywords: Conduct disorder, Dorsolateral prefrontal cortex, Emotion processing, Emotion regulation, Female adolescents, fMRI

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Adolescence is a time of profound changes, marked by an increase in emotionality and a growing need to acquire socioaffective skills, such as the ability to effectively regulate one's emotions (1,2). Emotion regulation skills provide a means of controlling the intensity, duration, or extent of the experience evoked by an emotional stimulus or situation (3,4). Proficient emotion regulation is typically attained in late adolescence or early adulthood (5) and has been linked to better social functioning and psychological and physical health (4,6). Deficient emotion regulation, in contrast, has been suggested to be associated with childhood psychopathologies, including conduct disorder (CD) (7–9).

CD is a psychiatric disorder of childhood and adolescence that is characterized by repetitive and persistent aggressive

and nonaggressive behaviors that violate others' basic rights or major age-appropriate societal norms (10). Overall, the lifetime prevalence of CD is estimated at 9.5% (11) but differs between the sexes [i.e., 12.0% for male and 7.1% for female adolescents (11)]. The origin of behaviors characteristic of CD [e.g., irritability, anger outbursts, or intense emotional responses (12)] may be underpinned by variations in emotional reactivity (13,14) and/or emotion-regulation difficulties (8,15). Considerable heterogeneity observed within CD adolescents has led to attempts to subtype these individuals according to specific features. Most prominently, variations in callous-unemotional (CU) traits, psychopathy, or levels of anxiety have been suggested to impact the behavioral and neural characteristics associated with CD (15–19).

The neural substrates of emotion regulation have been studied in healthy (20,21) and clinical (22–24) populations. Within the neuroimaging literature, effortful emotion-regulation strategies, such as cognitive reappraisal, have been the most commonly studied and effective strategies (3,21), and these are core targets of various intervention approaches (25–27). While different forms of emotion regulation exist, this paper focuses on effortful emotion regulation via cognitive reappraisal. Effortful emotion regulation by cognitive reappraisal follows an initial reaction to an emotional stimulus that activates affect-related brain regions, such as amygdala, insula, and striatum. Based on data from healthy adolescents and adults (27–29), two models of emotion regulation have been proposed (30). The mediation hypothesis of emotion regulation suggests that emotion regulation is achieved by prefrontal-subcortical mediation effects (28,30). According to this hypothesis, emotion regulation may include both 1) the activation of brain regions associated with cognitive control mechanisms, attention, and response inhibition (e.g., bilateral ventrolateral and dorsolateral prefrontal cortex [vlPFC/dlPFC]) or temporoparietal brain regions (e.g., angular/middle temporal gyrus) and 2) the modulation, or downregulation, of affective regions [e.g., amygdala, insula, or ventral striatum (27–29)]. Alternatively, the direct pathway hypothesis describes emotion regulation primarily through activity in prefrontal and cortical brain regions, without further involvement of subcortical systems (28,30).

While neuroimaging studies have examined the neural correlates of effortful emotion regulation in psychiatric populations or those who experienced childhood adversity (23,24,31), no data exist yet on CD. Meta-analytic studies of adolescents with CD or disruptive behavior have revealed functional (32–34) and structural (33–35) alterations in brain regions implicated within the emotion processing and regulation network, including vlPFC/dlPFC, anterior cingulate cortex or temporal gyrus, limbic brain regions (e.g., amygdala), insula, and striatum. In line with previous work, it was further demonstrated that the neural phenotype of CD varies depending on the level of CU traits or presence of comorbid disorders (35–37). For example, CD adolescents with high CU traits show hyporeactivity to emotional stimuli as compared to CD adolescents with low CU traits (15,32,37,38). Most research on CD to date has been conducted in male adolescents only (9,32); substantially less is known about CD in female adolescents (39).

Here we aimed to bridge this gap in the literature by investigating the neural basis of emotion regulation in female adolescents with CD. In line with the mediation hypothesis of cognitive emotion regulation derived from normative data in typically developing (TD) control adolescents (27–29,40), the known neural phenotype of CD (32–34), and reports of emotion processing and regulation deficits in CD (9), we expected to observe atypical neural activation during effortful emotion regulation in female adolescents with CD compared with in TD individuals. This would be reflected by the following hypotheses: 1) hypoactivation of prefrontal and cognitive control regions (e.g., dlPFC/vlPFC, anterior middle cingulate, superior temporal gyrus, and angular gyrus) in individuals with CD compared with in TD individuals; 2) deficient modulation of affect-related regions (amygdala, ventral striatum, insula), which would be demonstrated by continuing heightened

activation of these regions despite attempts to regulate emotions; and/or 3) no need to initiate emotion regulation because of initial differences in emotional reactivity and/or processing (i.e., reduced reactivity in female adolescents with CD and high CU traits).

METHODS AND MATERIALS

Participants and Measures

All participants included were recruited as part of the Neurobiology and Treatment of Female Adolescent Conduct Disorder study (9) and were tested at 2 sites (Universities of Basel and Frankfurt). A total of 59 female adolescents 15 to 18 years of age with either a clinical diagnosis of CD ($n = 30$; average age = 16.28 years) or without a current clinical diagnosis (TD: $n = 29$; average age = 16.74 years) were included. We initially scanned 65 female participants 15 to 18 years of age. In a first step, we excluded 5 participants to match groups for age, sex, site, and pubertal status, which may all impact the neural correlates of reappraisal (40–42) [pubertal status was measured through the Pubertal Development Scale (43)]. One TD participant was additionally excluded owing to neuroanatomical abnormalities. CD and common psychiatric disorders were assessed using the Kiddie Schedule for Affective Disorders and Schizophrenia–Present and Lifetime Version (44). Of the participants in the CD group, 26 met the diagnostic criteria for current CD (≥ 3 CD symptoms; 14 of them had comorbid oppositional defiant disorder [ODD]), 3 had 2 current CD symptoms while meeting full ODD criteria, and 1 participant had 1 CD symptom plus current ODD. Analyses were repeated including just those participants who met full CD criteria ($n = 26$) and including only the larger of the 2 sites (24 CD, 17 TD from Basel) to exclude diagnosis-dependent or nonlinear effects of site on our functional magnetic resonance imaging (fMRI) results (Supplemental Figure S2). For all participants, exclusion criteria included IQ < 70 [Wechsler Intelligence Scale for Children-IV/Wechsler Adult Intelligence Scale-IV (45)], autism spectrum disorder, psychosis, any neurological or genetic disorder, and standard MRI exclusion criteria. Finally, all participants were asked to complete the self-report Youth Psychopathic Traits Inventory (46) to assess psychopathic and CU traits and the parent-report Child Behavior Checklist (47) (owing to time constraints, Child Behavior Checklist data were only available for 17 participants with CD and 27 TD participants). Local ethics committees reviewed and approved the study at each site (Ethics Committees of Northwest and Central Switzerland in Basel and the Medical Faculty of Goethe University Frankfurt). Adolescents and their parents or caregivers provided informed assent and consent, respectively.

fMRI Task

All participants completed an age-appropriate adaptation of an fMRI emotion-regulation task (3,20,21,48) (see the task design and training protocol in the Supplement). Each experimental trial lasted 20,000 ms and started with a 2500-ms instruction cue, indicating whether the subjects will have to look at a neutral or negative picture or regulate emotions evoked by a negative emotional picture (i.e., decrease the emotions). Cognitive emotion regulation required the participants to

actively reappraise the experienced negative emotion to reduce the intensity of their emotional state. More specifically, they were asked to reinterpret the situations and scenes observed to experience more positive feelings [see also (3,20,21,48)]. The instruction cue was followed by a negative or neutral image presented for 10,000 ms. Subsequently, in-scanner self-report ratings of the strength of the negative affect (on a Likert scale of 1 = lowest to 4 = highest; presented for 5000 ms) were collected, before a relaxation period of 2500 ms preceded the next trial. This design allows modeling of 3 experimental conditions: 1) look neutral (nonemotional), 2) look negative (no regulation), and 3) decrease negative (emotion regulation). Participants completed 48 trials in total (16 in each of the 3 conditions). The experiment was divided across 2 runs (~8 minutes each).

Image Acquisition

Prior to data collection, both sites underwent site-qualification procedures and preassessments to ensure comparability of fMRI data [see protocols in the Supplement and in (49,50)]. Per run, 196 images were acquired with a 41-slice echo planar imaging interleaved sequence on a Siemens Prisma 3T scanner (Berlin, Germany) at the Basel site and a Siemens Trio 3T MR at the Frankfurt site. Image acquisition at both sites included the following specifications: repetition time = 2500 ms; echo time = 30 ms; flip angle = 83°; field of view = 192 mm; voxel size = 3 × 3 × 2 mm. The first 4 functional volumes were discarded from later analysis to account for T1-weighted equilibration effects.

fMRI Analysis

All fMRI data were analyzed using SPM12 (University College London, London, UK; <http://www.fil.ion.ucl.ac.uk/spm/doc/manual.pdf>) running under MATLAB2018b (The MathWorks, Inc., Natick, MA). To account for movement artifacts, all images were first realigned and unwarped with reference to the first image. Next, each participant's structural scan was used for the coregistration and segmentation functions prior to normalization into standard space (McConnell Brain Imaging Center ICBM 152 template). Finally, all images were smoothed using a 6-mm full width at half maximum isotropic kernel. Using the ART imaging toolbox (NeuroImaging Tools and Resources Collaboratory; https://www.nitrc.org/projects/artifact_detect/), 7 separate regressors accounting for motion and variations in mean signal intensity were added to the first-level model. Additional regressors omitting images with visible motion artifacts (51) were used to remove artifactual time points. Regressors of interest were created using a boxcar function for each experimental condition (look neutral, look negative, decrease negative), and contrasts of interest included emotional reactivity (look negative > look neutral) and emotion regulation (activation by emotion regulation: decrease negative > look negative; modulation by emotion regulation: decrease negative < look negative). Affect rating and relaxation periods were not included in the neuroimaging analyses.

At the second level, differences in blood oxygen level-dependent signal intensity change in response to contrasts of interest were assessed using random-effects group analyses. General linear models were constructed using site and

IQ as covariates of no interest. Contrasts for emotional reactivity and emotion regulation were entered in 1-sample *t* tests assessing within-group activation as well as 2-sample *t* tests to compare CD and TD groups. Statistical significance is reported based on a cluster-building threshold of $p < .001$ and a small-volume familywise error correction for a priori-defined regions of interest (ROIs) ($p < .05$; peak-level inference). This small-volume correction was achieved by creating an individual mask including all affective and prefrontal ROIs previously implicated in emotion regulation [affective: automatic anatomical labeling-based bilateral insula/amygdala and a 10-mm-spherical ROI for ventral striatum; cognitive: 10-mm-spherical ROIs for bilateral dlPFC/vlPFC, anterior cingulate, and angular and left middle temporal gyrus; according to (27–29); further details in the Supplement]. To inform about areas not previously implicated in emotion regulation, regions surviving whole-brain familywise error correction ($p < .05$, cluster-building threshold $p < .001$; cluster-level inference) are additionally reported.

Post Hoc Analyses: ROI and Functional Connectivity

For post hoc assessments and displaying purposes, we further extracted mean parameter estimates in a priori-defined affective and cognitive ROIs (as described above and in the Supplement) for emotion regulation (decrease negative vs. look negative). To assess emotional reactivity, mean parameter estimates for emotional reactivity (look negative – look neutral) were extracted for affective ROIs. Two-sample *t* tests implemented in SPSS, version 25.0 (IBM Corp., Armonk, NY) and corrected for multiple comparisons were used to assess post hoc assessments of group differences within a priori-defined ROIs.

Modulation of subcortical areas by cortical regions during cognitive emotion regulation has been demonstrated by some, but not all previous neuroimaging studies (21,30,52). Thereby, prefrontal-amygdala connections are considered particularly important (52,53). We followed up on this evidence by first directly testing correlations between activity in the amygdala and dlPFC (defined as independent anatomical ROIs) during emotion regulation. Second, we tested for additional regions that showed task-dependent functional correlations (coactivations) with neural activation in left angular gyrus and left dlPFC during emotion regulation using a weighted task-dependent and seed-based connectivity approach through the CONN toolbox [NeuroImaging Tools and Resources Collaboratory; <https://web.conn-toolbox.org/> (54)]. The condition time series convolved with a canonical hemodynamic response function served as weights within the analysis. Temporal correlations between activation in left angular gyrus and left dlPFC seeds to all other brain voxels were computed using a general linear model approach (further details in the Supplement).

Self-reported Emotion Intensity and Emotion Regulation Success

Self-reports of in-scanner affect ratings on a 1 to 4 Likert scale directly after each trial were obtained via button press to assess group differences on a behavioral level: ratings for each condition (emotional reactivity [look negative – look neutral])

Table 1. Group Characteristics

	CD, Mean \pm SD or <i>n</i>	TD, Mean \pm SD or <i>n</i>	CD/TD, <i>n</i>	<i>p</i> , Sig. 2-tailed
Age, Years	16.28 \pm 0.85	16.74 \pm 0.99	30/29	.061
Handedness (Right/Left/Ambidextrous)	22/2/3	21/8/0	27/29	.893
Pubertal Status ^a	4.2	4.4	29/29	.133
Age of Onset (Childhood/Adolescence), Years	10/19	—	29/—	
IQ				
Performance IQ	103.67 \pm 13.64	105.34 \pm 11.25	30/29	.609
Verbal IQ	95.67 \pm 16.70	106.72 \pm 12.91	30/29	.006
Total IQ	99.90 \pm 12.95	106.24 \pm 9.64	30/29	.038
Comorbidities (DSM-5)				
Attention-deficit/hyperactivity disorder	7	0	29/26	
Major depressive disorder	5	0	28/26	
Generalized anxiety disorder	0	0	29/26	
Posttraumatic stress disorder	2	0	30/26	
Alcohol dependence	1	0	28/26	
Substance dependence	5	0	28/26	
YPI				
Grandiose-manipulative	39.24 \pm 10.43	32.97 \pm 8.90	29/29	.017
Callous-unemotional	30.10 \pm 6.97	24.52 \pm 5.16	29/29	.001
Impulsive-irresponsible	39.93 \pm 7.71	30.93 \pm 6.48	29/29	<.001
Total score	109.28 \pm 20.31	88.41 \pm 15.30	29/29	<.001
CBCL				
Internalizing subscale	65.47 \pm 10.28	49.3 \pm 10.62	17/27	<.001
Externalizing subscale	71.12 \pm 5.85	47.0 \pm 7.84	17/27	<.001
Total score	69.65 \pm 6.97	48.00 \pm 10.16	17/27	<.001

Unless otherwise indicated, *t* tests nonsignificant at threshold *p* = .05. For IQ, standard scores are reported; for Youth Psychopathic Traits Inventory (YPI) and Child Behavior Checklist (CBCL), mean scores are reported.

CD, conduct disorder; Sig., significance; TD, typically developing.

^aPubertal status was measured using the Pubertal Development Scale.

and emotion-regulation success (look negative – decrease negative condition). Differences between conditions within each group were evaluated using paired-samples *t* tests, whereas independent-samples *t* tests were used to compare the CD and TD groups.

RESULTS

Descriptive Statistics

Female adolescents with CD had significantly lower total and verbal IQ than TD female adolescents, but there were no group differences in age, performance IQ, or pubertal status. Adolescents with CD, compared with TD adolescents, scored significantly higher in psychopathic and CU traits and externalizing and/or internalizing symptoms (Table 1; see Supplemental Table S1 for site distributions and effects).

Self-report of Emotional Reactivity and Emotion Regulation Success

Relative to TD adolescents, adolescents with CD reported significantly lower scores for in-scanner emotional reactivity and emotion-regulation success (both *p* \leq .001) (Figure 1). However, both adolescents with CD and TD adolescents displayed significant within-group emotional reactivity and regulation success (mean scores and detailed group statistics in Table 2).

fMRI Results

Within-Group (1-Sample *t* Tests). The neural correlates of emotion regulation (activation by emotion regulation: decrease negative > look negative) in TD adolescents corresponded to a network of brain regions previously linked to cognitive reappraisal (27–29,40) and were reflected in significant activation of left angular gyrus and dlPFC (superior/middle frontal gyrus). No modulation by emotion regulation (decrease negative < look negative) of affect-associated regions was observed in the TD group. Adolescents with CD showed no significant activation increases during effortful emotion regulation, but they did show a significant decrease in activation in inferior orbitofrontal cortex, hippocampus/thalamus, and occipital cortex (Figure 2, Supplemental Table S2).

Between-Group (2-Sample *t* Tests). Group comparisons revealed significantly lower activation during emotion regulation (decrease negative > look negative) in adolescents with CD, compared with TD adolescents, in left dlPFC (inferior/middle frontal gyrus) and angular gyrus (Figure 2, Table 3). There were no significant differences in activation between adolescents with CD and TD adolescents for emotional reactivity (look negative > look neutral). Our findings remained unchanged when analyses were repeated in an IQ-matched subsample or when restricted to include 1 site per diagnosis only (see Supplemental Figure S2 and the Supplement).

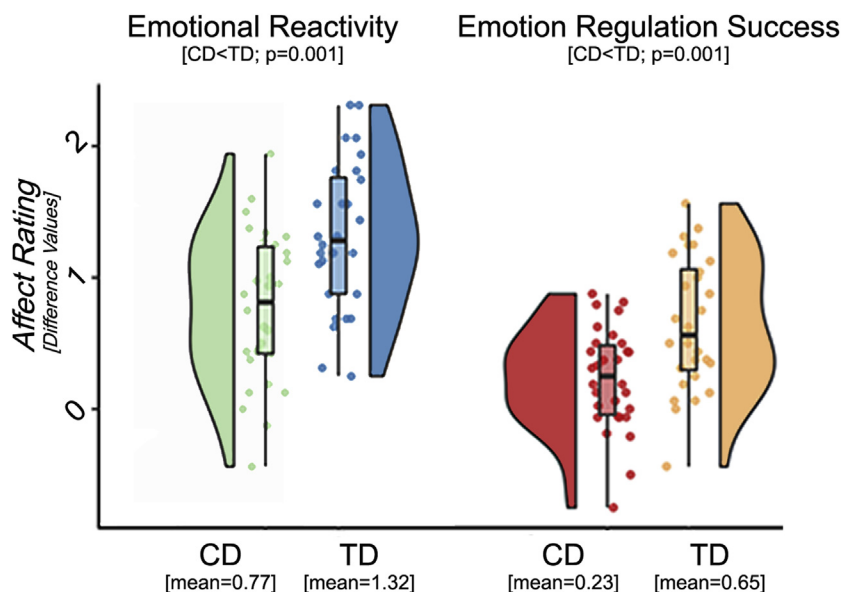


Figure 1. In-scanner affect ratings. Behavioral in-scanner reports as displayed using “raincloud” plots, which combine boxplots, raw jittered data, and a split-half “violin” [see (74)], revealed significantly lower ratings for female adolescents with conduct disorder (CD) compared with typically developing (TD) control adolescents in respect to emotional reactivity (difference for look negative – look neutral) and emotion regulation success (difference for look negative – decrease negative).

Post Hoc ROI Results. To assess putative models of cognitive reappraisal as previously suggested in healthy adolescents and adults (27–29) and for consequent follow-up analyses (e.g., assessment for CU traits-related effects), we extracted mean parameter estimates from the relevant ROIs. We expected emotional reactivity-related activation in amygdala, striatum, and insula regions, which in turn may be altered in CD. No significant group differences in emotional reactivity were observed in the ROIs. We expected the neural correlates of emotion regulation to differ in adolescents with CD compared with TD individuals. In line with the mediation hypothesis of emotion regulation (28,30), amygdala, striatum, and insula ROIs were hypothesized to decrease in activation (reflecting modulation), while cognitive ROIs were expected to be activated in TD adolescents, but less so in CD adolescents. Between-group findings were assessed using 2-sample *t* tests adjusted for multiple comparisons (Bonferroni correction). Familywise adjusted α levels were achieved by dividing $p < .05$ by the number of tests conducted for modulation by emotion regulation (adjusted $p < .008$) or activated activation by emotion regulation (adjusted $p < .006$). We found that adolescents with CD showed lower left angular and middle

temporal gyrus activation than did TD adolescents during emotion regulation, following correction for multiple comparisons (Figure 3, Supplemental Table S2).

Post Hoc Analyses Assessing Functional Connectivity During Emotion Regulation. Correlational analyses (controlling for site and IQ) between extracted neural activation parameters from left dIPFC and amygdala ROIs during emotion regulation (decrease > look negative) were significant in TD individuals ($p < .001$; $r = .747$), but not in individuals with CD ($p = .170$; $r = .257$), giving rise to significant group differences in left dIPFC–amygdala connectivity ($p < .010$). Additionally, task-dependent and seed-based functional connectivity analyses revealed that areas of coactivation based on the left dIPFC differed between TD adolescents and adolescents with CD during reappraisal (2-tailed significance, cluster-building $p < .001$ and clusterwise false discovery rate correction of $p < .05$) (Figure 4). Follow-up analyses of this group effect revealed that individuals with CD, compared with TD individuals, showed lower connectivity between the left dIPFC seed and bilateral putamen, right orbitofrontal cortex/vIPFC, and amygdala. No group differences were observed for left angular gyrus.

Table 2. In-Scanner Affect Ratings

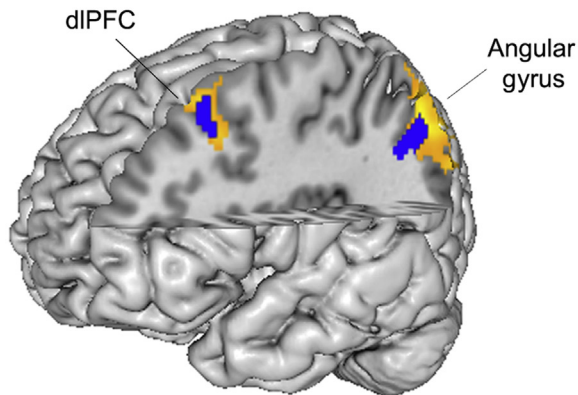
	Look Neutral	Look Negative	Decrease Negative	Emotion Intensity (Within Group; p Paired t / r)	Emotion-Regulation Success (Within Group; p Paired t / r)
CD	1.454	2.224	1.993	0.770 (<0.001 / .5961)	0.231 (0.003 / .1839)
TD	1.333	2.649	2.003	1.316 (<0.001 / .8036)	0.646 (<0.001 / .5052)
CD vs. TD, p 2-Tailed	.221 ($r = .1602$)	.010 ($r = -.3301$)	.944 ($r = -.0092$)	.001 ($r = -.4337$)	.001 ($r = -.4197$)

CD, conduct disorder; TD, typically developing.

Activation by emotion regulation (decrease negative > look negative)

Two-sample *t*-test: CD < TD

One-sample *t*-test: TD



Modulation by emotion regulation (decrease negative < look negative)

Two-sample *t*-test: not significant

One-sample *t*-test: CD

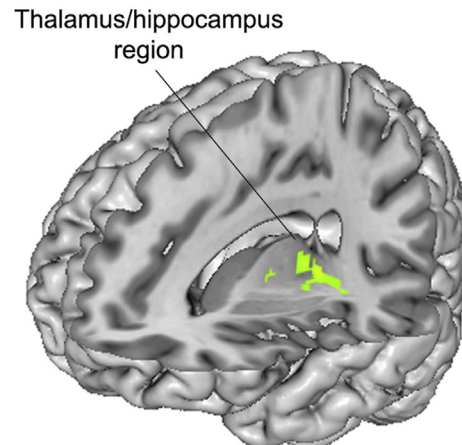


Figure 2. Neural correlates of emotion regulation in and group differences between adolescents with conduct disorder (CD) and typically developing (TD) adolescents. Statistical parametric maps reflecting the neural correlates of emotion regulation. Activation by emotion regulation: (Left) Clusters that were significantly activated in the TD group are shown in orange-yellow; regions that were less active in participants with CD than in TD participants are shown in blue (based on a cluster-building $p < .001$ and small-volume familywise error correction of $p < .05$). (Right) Regions that were modulated during emotion regulation in the CD group are shown in green (based on an exploratory whole-brain search with a cluster-building $p < .001$ and familywise error-corrected $p < .05$). dlPFC, dorsolateral prefrontal cortex.

Influence of CU Traits on Behavior and Neural Activation. We assessed whether CU traits or the presence of comorbidities may have impacted the observed neural and behavioral group differences reported.

To assess CU traits and neural activity, the scaled mean parameter estimates from independent ROIs for areas of interest (i.e., left dlPFC and angular gyrus) were entered as individual dependent variables into 2 stepwise multiple regression models assessing changes in R^2 , with group status (TD, CD), covariates (site and IQ), CU traits, and comorbidities (attention-deficit/hyperactivity disorder, major depressive disorder, posttraumatic stress disorder, alcohol dependence, substance dependence) as independent variables. Our findings revealed that group differences in left angular gyrus and dlPFC (CD < TD) were unrelated to CU traits. Likewise, covariates and comorbidities were not significant factors in the model.

To assess CU traits and in-scanner affect ratings, we used stepwise multiple regression models to determine whether the variance in behavioral findings (emotional reactivity and emotion-regulation success as predictors) was explained by the independent variables—group status (TD, CD), covariates (site, IQ), CU traits, and comorbidities. The only significant predictor of variance in emotion-regulation success was group status ($p = .002$; 18.2%), while variance in emotional reactivity was explained by group status ($p < .001$; 21.5%), with an additional 9.8% of the variance explained by CU traits ($p = .009$). Covariates and comorbidities were not significant factors in the model. Follow-up partial correlational analysis,

accounting for site and IQ, within the CD and TD groups individually demonstrated a negative relationship between emotional reactivity and CU traits in adolescents with CD ($r = -.390$, $p = .044$), but not in TD adolescents ($r = -.269$, $p = .184$) (Supplemental Figure S4).

DISCUSSION

This is the first study to examine neural activity during effortful emotion regulation in female adolescents with CD. In line with our main hypothesis, our results indicate that female adolescents with CD, compared with TD adolescents, displayed atypical neural activation during emotion regulation, as indicated by reduced left dlPFC and angular gyrus activation. This was supported by behavioral findings of deficient emotion regulation in female adolescents with CD. Additionally, post hoc functional connectivity analyses revealed weaker connectivity between the left dlPFC and the right amygdala, vlPFC/orbitofrontal cortex, and bilateral putamen in the CD group versus the TD group. CU traits were negatively correlated with emotional reactivity as measured behaviorally, but unlike in previous studies (18,19,55), they were unrelated to the neural correlates of emotion regulation or reactivity.

Corresponding to hypothesis 1 and in line with prior evidence (3,21,28), our data indicate an involvement of left hemisphere angular gyrus and dlPFC during emotion regulation in TD adolescents, which is reduced in adolescents with CD. Functional neuroimaging studies in healthy control adolescents have suggested that the dlPFC is central to

Table 3. Peak Activation Reports for Activation and Modulation by Emotion Regulation

Brain Region		Side	<i>t</i>	<i>P</i> _{FWE}	Cluster Size (<i>k</i>)	MNI Coordinates		
Lobe	Area					x	y	z
Small-Volume Correction								
TD: Activation by ER (Decrease Neg > Look Neg)								
Parietal	Angular gyrus, superior/inferior parietal lobe	L	5.57	.006	327	-46	-56	40
Frontal	Superior/middle frontal gyrus (dlPFC)	L	4.71	.038	231	-30	18	42
TD: Modulation by ER (Look Neg > Decrease Neg)								
CD: Activation by ER (Decrease Neg > Look Neg)								
CD: Modulation by ER (Look Neg > Decrease Neg)								
TD > CD: Activation by ER (Decrease Neg > Look Neg)								
Parietal	Angular gyrus, mid occipital/inferior parietal lobe	L	4.85	.009	129	-30	18	40
Frontal	Inferior/middle frontal gyrus (dlPFC)	L	4.55	.023	118	-40	-58	36
TD > CD: Modulation by ER (Look Neg > Decrease Neg)								
Whole-Brain Correction								
TD: Activation by ER (Decrease Neg > Look Neg)								
Parietal	Angular gyrus, superior/inferior parietal, middle occipital lobe	L	5.57	<.001	972	-46	-56	40
Frontal	Middle/superior frontal lobe	L	4.71	.028	385	-30	18	42
TD: Modulation by ER (Look Neg > Decrease Neg)								
CD: Activation by ER (Decrease Neg > Look Neg)								
CD: Modulation by ER (Look Neg > Decrease Neg)								
Frontal	Olfactory, anterior cingulate, caudate, putamen, OFC	L/R	6.57	.012	180	-18	30	-6
Temporal	Thalamus, hippocampus, lingual, precuneus, parahippocampus	L/R	6.56	<.001	412	-6	-24	6
Occipital	Fusiform gyrus, inferior/middle occipital gyrus, lingual, calcarine	R	4.96	.003	238	30	-84	-10
Occipital	Middle, inferior, superior occipital lobe	L	4.57	.008	195	-22	-98	8
TD > CD: Activation by ER (Decrease Neg > Look Neg)								
Parietal	Angular, inferior parietal, mid occipital lobe	L	4.64	.046	231	-38	-58	34
TD > CD: Modulation by ER (Look Neg > Decrease Neg)								

CD, conduct disorder; dlPFC, dorsolateral prefrontal cortex; ER, emotion regulation; FWE, familywise error; L, left; MNI, Montreal Neurological Institute; Neg, negative; NS, not significant; OFC, orbitofrontal cortex; R, right; TD, typically developing.

emotion regulation (28,30,52) and is associated with tasks requiring cognitive control [e.g., working memory, dual-task performance, or response inhibition (56,57)]. The mediation hypothesis of emotion regulation suggests that the dlPFC achieves negative affect reduction by modulating subcortical regions such as the amygdala (30). Successful coupling between prefrontal and subcortical brain regions has been associated with emotion-regulation success (53), while reduced prefrontal-subcortical connectivity has been associated with increased symptoms in psychiatric populations [e.g., posttraumatic stress disorder/anxiety (58)]. However, depending on the strategy used, PFC activity can be related to both positive and negative appraisal processes and amygdala decreases and increases, respectively (30). Overall, functional and structural alterations of the dlPFC are common in adolescents with CD/ODD (34,39,59). Our findings of reduced dlPFC activation during emotion regulation in CD may thus reflect a more generic disruption in PFC functioning in CD. Some of the behavioral characteristics seen in individuals with CD may be explained by deficient recruitment of cognitive

control regions (i.e., dlPFC and angular gyrus). Reduced prefrontal activity in CD has previously also been associated with implicit forms of emotion regulation (60,61).

Contrary to hypothesis 2 and prior evidence (14,20,28,39,62), no significant neuronal evidence of down-regulation/modulation of affect-related areas was observed during reappraisal. On a first view, our data may therefore be less supportive of the mediation hypothesis of emotion regulation because emotion regulation was only associated with prefrontal and angular gyrus activation, without accompanying modulation of subcortical structures. However, the developmental and clinical sample investigated renders interpretation challenging. Developmental research has indicated that a reduced amygdala response following emotion regulation may become apparent only in late adolescence (40). The expected inverse prefrontal-subcortical coupling could still be developing with age (40,63). Furthermore, differences in brain age (i.e., delayed developmental trajectories) are characteristic of psychiatric disorders (64), complicating interpretations. Finally, potentially coexisting positive and negative dlPFC-amygdala

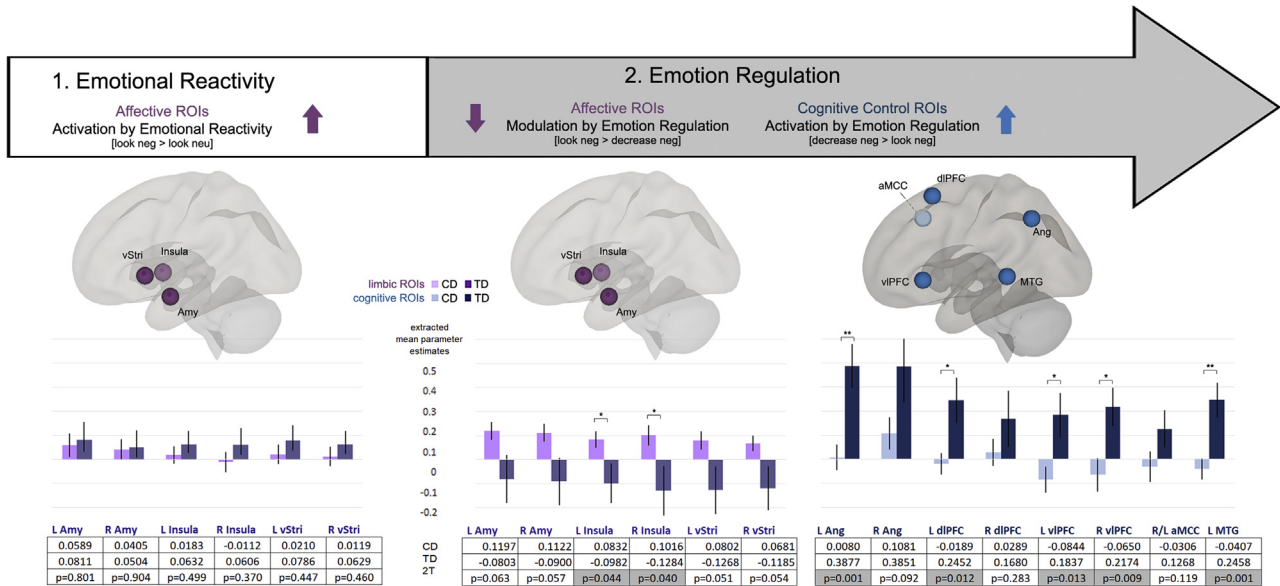


Figure 3. Post hoc display of neural activation in regions of interest (ROIs) and group differences in activation. Graphical display reflecting processes preceding emotion regulation (emotional reactivity) and actual emotion regulation (activation of cognitive ROIs [blue] or modulation of affective ROIs [lilac] during emotion regulation). Extracted mean parameter estimates for ROIs are reported for adolescents with conduct disorder (CD) (lilac/blue: light colors) and typically developing (TD) adolescents (lilac/blue: dark colors). Two-sample *t* tests corrected for multiple comparisons indicate hypoactivation in left angular and middle temporal gyrus in adolescents with CD compared with TD adolescents. *Uncorrected *p* < .05; ***p* corrected for multiple comparisons. Amy, amygdala; aMCC, anterior middle cingulate gyrus; Ang, angular gyrus; dIPFC, dorsolateral prefrontal cortex; L, left; MTG, middle temporal gyrus; neg, negative; neu, neutral; R, right; vIPFC, ventrolateral prefrontal cortex; vStri, ventral striatum.

correlations could cancel each other out (30). Transient subcortical modulatory effects within the amygdala may have been missed in the present paradigm using a relatively long blocked time window for the modeled regressors of interest. It remains to be investigated whether shorter, event-related designs enable detection of more transient subcortical signals (65). Overall, the neural differences identified here could reflect fundamental differences in emotion regulation (consistent across the life span) between individuals with CD or TD individuals and/or transient effects resulting from delayed brain development in CD that may eventually normalize.

While some previous evidence supports a prefrontal-subcortical mediation theory [e.g., (20,21,28,66)], the precise nature of such relations, particularly in clinical populations, warrants further investigation. Using post hoc assessments of functional connectivity, we observed positive connectivity between the left dIPFC and bilateral putamen, amygdala, and vIPFC during reappraisal in TD individuals, which was reduced in the CD group. Putamen and nucleus accumbens conjointly constitute the dorsal striatum, which is in close anatomical and functional association with the insula (67). Functionally, the putamen has been implicated during tasks of emotion and reward processing, learning, or working memory (68). Overall, coactivation findings of dIPFC with vIPFC/prefrontal brain regions, amygdala, and striatum are in line with meta-analytic evidence on emotion regulation (28).

The within-group analyses of behavioral affect ratings collected in the scanner suggest that both TD adolescents and adolescents with CD showed intact emotional reactivity (their affect ratings were higher to negative than neutral images) and were able to successfully apply emotion-regulation strategies

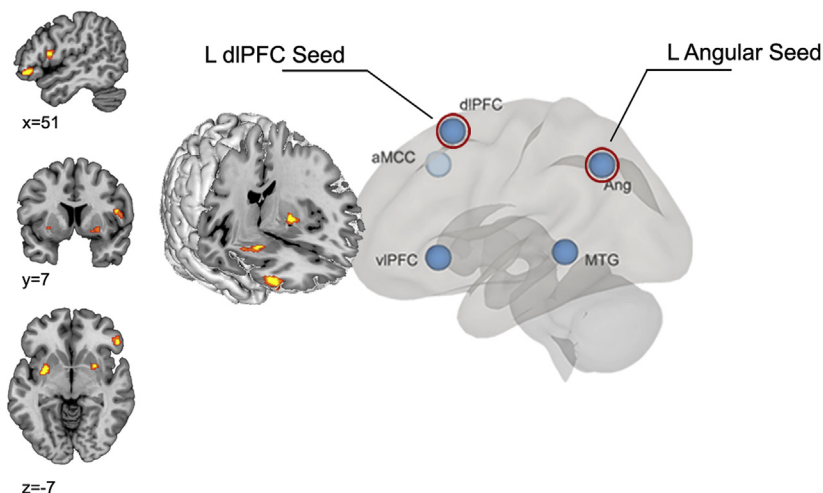
(affect ratings of negative images were lower in the decrease condition than in the look condition). However, adolescents with CD were significantly less successful in emotion regulation and reported lower emotional reactivity overall, which corresponds to behavioral deficits reported for CD (8,9,15,69,70). Hypothesis 3 was therefore only partly supported: there were behavioral differences for both emotion regulation and emotional reactivity (CD < TD), but there were only neural differences between the groups in respect to reappraisal and not emotional reactivity, contrary to previous reports (14,39,71).

Because of the intricate nature of the task design and population studied, there are several limitations that should be noted. Behavioral evidence of reduced emotional reactivity could suggest that despite reported reappraisal success, there was less need for or potential to show emotion regulation in the CD group. While prior studies suggest stability, reliability, and predictive value of in-scanner self-reports (30,72), and while we carefully trained all participants, we cannot fully exclude the possibility that participants with CD were less motivated to engage in effortful reappraisal or that their reports were influenced by reporting biases. Finally, it is possible that differences in the choice of emotion-regulation strategies (e.g., increased use of suppression in CD) impacted the present findings. The neural correlates of emotion regulation are known to depend on the strategy used [e.g., PFC activation during suppression and reappraisal, but decreased amygdala and insula response for reappraisal vs. increased response in the same areas during suppression (73)]. Additional noteworthy limitations are the presence of comorbid disorders, particularly substance and alcohol abuse (information on potential

Seed-based functional connectivity during emotion regulation

(TD>CD; decrease negative > look negative)

ANCOVA (controlling for IQ and site)
p<0.05, FDR-corr



H	Region	MNI			Cluster size	p-FDR cluster-size	Effect size	T
		x	y	z				
<i>L dIPFC Seed</i>								
R	vIPFC, frontal pole, inferior frontal gyrus, OFC	54	32	-6	66	0.009	0.47	5.43
L	putamen	-30	-2	-8	49	0.021	0.29	5.01
R	putamen, amygdala	20	4	-10	43	0.025	0.30	4.83
R	inferior frontal gyrus, precentral gyrus	48	8	12	39	0.027	0.36	4.53

Figure 4. Reduced functional connectivity in adolescents with conduct disorder (CD) compared with typically developing (TD) adolescents during emotion regulation. Connectivity differences (CD < TD) were demonstrated by group assessments in coactivation patterns for left (L) dorsolateral prefrontal cortex (dIPFC) with bilateral putamen, right (R) orbitofrontal cortex (OFC)/ventrolateral prefrontal cortex (vIPFC) and amygdala during cognitive reappraisal (decrease negative > look negative). aMCC, anterior middle cingulate gyrus; ANCOVA, analysis of covariance; Ang, angular gyrus; FDR-corr, false discovery rate-corrected; H, hemisphere; MNI, Montreal Neurological Institute; MTG, middle temporal gyrus.

substance consumption in the days prior to testing was incomplete) and the absence of a male comparison group. Accumulating evidence suggests that the neural correlates of CD may be sex specific (49,59). It remains to be investigated whether male adolescents with CD show similar distinct neural alterations (if any) during emotion regulation.

The present findings of atypical left angular gyrus and dIPFC activation during emotion regulation in female adolescents with CD and reduced functional connectivity between the left dIPFC and cortical (vIPFC/orbitofrontal cortex, putamen), and subcortical (amygdala) regions emphasize the importance of addressing emotion regulation when treating CD. Future studies may investigate whether targeted interventions, such as cognitive behavioral therapy, enhance emotion-regulation skills in adolescents with CD and/or ameliorate the alterations in prefrontal and parietal activation observed here.

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CF receives royalties from books on autism spectrum disorder, attention-deficit/hyperactivity disorder, and major depressive disorder and is a consultant for Roche and Desitin. GF is an editor of the journal *European Child and Adolescent Psychiatry*. All other authors report no biomedical financial interests or potential conflicts of interest.

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