

## RESEARCH ARTICLE

# Attention and Executive Function in Children Diagnosed with Attention Deficit Hyperactivity Disorder and Comorbid Disorders

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### Abstract

**Objective:** The goal of this study was to examine the relationship between comorbid disorders and executive function (EF) in children diagnosed with Attention Deficit/Hyperactivity Disorder (ADHD). **Methods:** Three hundred and fifty-five, 6-12 year old children clinically diagnosed with ADHD were included in the study. Comorbid anxiety disorders, Oppositional Defiant Disorder (ODD) and Conduct Disorder (CD) were examined. The EF domains were assessed using the Conners' Continuous Performance Test (CPT), Wisconsin Card Sorting Test (WCST), Tower of London (ToL), Finger Windows (FW) and Self Ordered Pointing Test (SOPT). **Results:** The findings indicate that children with comorbid anxiety disorders performed worse in domains measured by CPT and prior to controlling for age and sex, by FW. However, once sex was controlled for the results for FW were no longer significant. Children with CD obtained lower scores on WCST. Furthermore, a significant sex by CD interaction was observed. **Conclusion:** These results indicate that comorbid disorders should be carefully examined as they play a significant role in EF performance and subsequently in day-to-day functioning of children with ADHD.

**Key Words:** ADHD, Executive Function, comorbid disorders, conduct disorder

### Résumé

**Objectif:** Cette étude visait à examiner la relation entre les troubles comorbides et la fonction exécutive (FE) chez les enfants ayant reçu un diagnostic de trouble de déficit de l'attention avec hyperactivité (TDAH). **Méthodes:** Trois cent cinquante-cinq enfants de 6 à 12 ans chez qui le TDAH a été cliniquement diagnostiqué ont été inclus dans l'étude. Les troubles anxieux, le trouble oppositionnel avec provocation (TOP) et le trouble des conduites (TC) comorbides ont été examinés. Les domaines de la FE ont été évalués à l'aide du test de performance continu de Conners (CPT), du test de tri de cartes du Wisconsin (WCST), du test de la Tour de Londres (ToL), du test Finger Windows (FW) et du test de pointage auto-imposé (SOPT). **Résultats:** Les résultats indiquent que les enfants souffrant de troubles anxieux comorbides avaient un plus mauvais rendement dans les domaines mesurés par le CPT et avant le contrôle pour l'âge et le sexe, dans FW. Cependant, une fois le sexe contrôlé, les résultats de FW n'étaient plus significatifs. Les enfants souffrant du TC obtenaient

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Submitted: October 5, 2015; Accepted: October 10, 2016

des scores plus faibles au WCST. En outre, une interaction significative du sexe dans le TC a été observée. **Conclusion:** Ces résultats indiquent que les troubles comorbides devraient être examinés avec soin car ils jouent un rôle significatif dans le rendement de la FE et subséquemment, dans le fonctionnement quotidien des enfants souffrant du TDAH.

**Mots clés:** TDAH, fonction exécutive, troubles comorbides, trouble des conduites

## Introduction

Attention Deficit/Hyperactivity Disorder (ADHD) is a neurodevelopmental disorder, more commonly diagnosed in boys (Rucklidge, 2010). Children diagnosed with ADHD are at higher risk for educational failure, social difficulties, as well as high risk behaviour (Biederman et al., 2006). ADHD rarely occurs in isolation and there is now a well-established body of literature that has identified a range of frequently occurring comorbid psychiatric disorders that include oppositional defiant disorder (ODD), conduct disorder (CD) and anxiety disorders (Biederman et al., 2008; Schatz & Rostain, 2006). Rates of comorbid disorders range from 24% to 71%, varying across studies and across disorders (Jensen et al., 2001; Kadesjo & Gillberg, 2001; Robison, Sclar, Skaer & Galin, 1999). With about 50% of children meeting criteria for ODD or CD, and 25 to 33% for anxiety disorders, the presence of comorbidities plays a significant negative role in the degree of impairment and on the course of the disorder. This presents an additional challenge for the diagnosis and treatment of children with ADHD (Ollendick, Jarrett, Grills-Taquechel, Hovey & Wolff, 2008; Cuffe et al., 2015; Pliszka, Sherman, Barrow, & Irick, 2000).

In addition to comorbid disorders, children with ADHD often exhibit Executive Function (EF) deficits. EF represents a set of cognitive processes that integrate information from working memory with information about context in order to select optimal action (Willcutt, Doyle, Nigg, Faraone & Pennington, 2005). Among children diagnosed with ADHD, deficits have been reported in the EF domains of response inhibition and execution, vigilance, working memory, set and task-switching/cognitive flexibility and planning (Toplak, Bucciarelli, Jain & Tannock, 2009; Willcutt et al., 2005).

EF deficits are not specific to ADHD, but have also been associated with other psychiatric conditions such as symptoms of anxiety/depression (Emerson, Mollet & Harrison, 2005), obsessive compulsive disorder (Chamberlain et al., 2007) and CD (Toupin, Dery, Pauze, Mercier & Fortin, 2000; Pajer et al., 2008). Subsequently, several studies have examined EF deficit among children with a diagnosis of ADHD and comorbid disorders, however, these are limited in number and have shown inconsistent results (Doyle, 2006). For example, the presence of comorbid ODD or CD was reported not to affect EF performance as presence of ADHD accounted for EF deficit in children diagnosed

with ADHD and ODD/CD (Kalff et al., 2002; Oosterlaan, Scheres and Sergeant, 2005). Similarly, in teens with disruptive behavioural disorders the presence of comorbid ADHD determined worse performance on EF tasks (Hummer et al., 2011). Rhodes et al. (Rhodes, Park, Seth, & Coghill, 2012) on the other hand found that compared to typically developing children, children with either ADHD or ODD or both performed worse on EF tasks. In adults, those with dual diagnosis of ADHD and CD showed poorer EF performance compared to ADHD only (Fischer, Barkley, Smallish & Fletcher, 2005).

Among children diagnosed with ADHD, those who reported anxiety have been shown to have/demonstrate better behavioural inhibition, while parent reported anxiety was not associated with neurocognitive dysfunction (Bloemsma et al., 2013). In a study examining working memory in four subgroups (children with anxiety, with ADHD, with ADHD and anxiety (N = 108) and a control group), children with dual diagnosis were found to display a similar impairment to children with only ADHD diagnosis, compared to both the control group and anxiety only group (Manassis, Tannock, Young & Francis-John, 2007).

In addition to EF differences seen among children presenting with various psychopathologies, sex differences have also been reported on some measures of EF (De Luca et al., 2003; Seidman et al., 2005), attention (Newcorn et al., 2001), and reaction time (Brocki & Bohlin, 2004). In general, girls have been found to be less impaired in such domains as compared to boys (Hasson & Fine, 2012). Moreover, sex differences have been well documented in the phenotypic expression of the ADHD, with girls demonstrating lower levels of disruptive behaviour and higher levels of inattentive symptoms (Stefanatos & Baron, 2007).

Given the frequently observed EF deficit and the high rate of comorbidities in ADHD population, the main aim of the current study is to examine the role comorbid disorders play on EF task performance in a large sample of children clinically diagnosed with ADHD. We examined the three most common comorbidities reported among elementary school age children. We hypothesised that the presence of anxiety disorders, CD or ODD will have an additive negative effect on EF, resulting in poorer performance compared to children without comorbid disorders. In addition, due to reported sex differences in ADHD and in EF we expected to find sex differences in EF performance among children diagnosed with ADHD.

This study will contribute to identifying the impact anxiety disorders, ODD and CD have on EF, and will allow clinicians to refine individual intervention by paying closer attention to neurocognitive delays that may interfere with the child's functioning. In addition, this study addresses several of the limitations of previous studies, particularly with respect to sample size and the use of a thorough diagnostic and standardised assessment procedure.

## Methods

### Participants

Children ages six to 12 diagnosed with ADHD and referred to the ADHD clinic were recruited sequentially for this study. Children were excluded from the study if they met the DSM-IV criteria for psychosis, had a chronic medical condition, autism spectrum disorder, Gilles de la Tourette's syndrome, or had an IQ less than 70. The final sample consisted of 355 children, 267 boys and 88 girls. The mean age for the group was 9.40 years (SD = 1.66), with no statistical difference in age between boys and girls.

### Design and setting

The study was conducted at a university affiliated mental health institute. Internal Research Ethics Board approval was obtained for the study protocol. All participants signed the informed consent and children gave verbal assent to participate in the study. Primary caregivers completed standardised interviews and questionnaires while children were assessed using EF measures. All medications were stopped prior to assessment and children underwent a minimum five-day medication washout period prior to completing the EF assessment. Two hundred thirty-nine children were not taking any medication when enrolled in the study, 105 children were taking medication at the time of participation. Of these children, 82 children were taking methylphenidate, 15 children were taking other commonly used stimulant or non-stimulant medication, and 12 children were taking risperidone (n=10), seroquel (n=1), clonidine (n=1) or haloperidole (n=1) alone or in combination. The washout period was determined by the team psychiatrist according to the type of medication the child was prescribed. Children taking neuroleptic medication had a washout period of two weeks. All EF measures were administered in the morning, to minimize the possible effects of fatigue on task performance.

### Clinical Assessment

All children were diagnosed with ADHD and/or comorbidities by child psychiatrists according to the fourth edition of the DSM-IV interview based on school reports, observation of the child, and clinical interview with the family. To confirm the diagnosis as well as assess comorbidities a trained research assistant interviewed a parent, using *The National Institutes of Mental Health Diagnostic Interview Schedule*

**Table 1. Distribution of comorbid disorders**

No comorbid disorders	31% (N=110)
ANX only	13.5% (N=48)
CD only	5.1% (N=18)
ODD only	19.4% (N=69)
ANX+CD	7.6% (N=27)
ANX+ODD	23.4% (N=83)
Note: ANX = anxiety disorders; CD = Conduct Disorder; ODD = Oppositional Defiant Disorder.	

*for Children 4<sup>th</sup> edition* (DISC-IV: Shaffer, Fisher, Lucas, Dulcan & Schwab-Stone, 2000). DISC – IV is a highly structured, DSM-IV diagnostic interview which assesses most common mental disorders in children. The hierarchical nature of the interview allows for thorough assessment of each symptom, including duration, the frequency of occurrence, and interference with functioning. The following comorbid disorders were examined in this study: Oppositional Defiant Disorder (ODD), Conduct Disorder (CD), and anxiety disorders. DISC-IV was used to report on positive ADHD symptoms.

### Child measures

Wechsler Intelligence Scale for Children – 3<sup>rd</sup> (WISC-III) or 4<sup>th</sup> (WISC-IV) editions was used to assess the children's general cognitive ability (Wechsler, 1991). Children were assessed using the following measures: Conners Continuous Performance Test (CPT: Conners, Epstein, Angold & Klaric, 2003), Wisconsin Card Sorting Test (WCST: Heaton, Chelune, Talley, Kay, & Curtiss, 1993), Tower of London (ToL: Anderson, Anderson & Lajoie, 1996), Self-Ordered Pointing Test (SOPT: Petrides & Milner, 1982), and Finger Windows (FW: Sheslow & Adams, 2001).

### Statistical Analysis

Statistical analyses were conducted using the SPSS software, Chicago, IL, USA. Continuous variables were expressed as means  $\pm$  standard deviation (SD). Categorical variables were expressed as proportions (%). Chi Square tests were used to compare proportions between groups. SPSS General Linear Model procedure (GLM) multivariate analysis of covariance (MANCOVA) or univariate analysis of covariance (ANCOVA) were used to examine the effect of multiple fixed factors and control for confounding variables (sex and age). Separate MANCOVA analyses were conducted grouping variables according to domains they examine. CPT variables were examined together, WCST variables were examined together, and working memory measures (SOPT and FW) were examined together. As only

Table 2a. Demographic and clinical characteristics of children with and without anxiety disorders					
	With Anxiety n=158		Without Anxiety n=197		p
	Mean	SD	Mean	SD	
Age	9.43	1.68	9.37	1.65	n.s
Percent boys	74.1% (n=117)		76.1% (n=150)		n.s
FSIQ	96.99	12.32	97.4	13.49	n.s
DISC-IV Inattentive symptoms	7.29	2.16	6.56	2.28	.002
DISC-IV Hyperactive symptoms	5.47	2.64	4.79	2.71	.017

SD = Standard Deviation; FSIQ = Full Scale IQ.

Table 2b. Demographic and clinical characteristics of children with and without ODD					
	With ODD n=152		Without ODD n=203		p
	Mean	SD	Mean	SD	
Age	9.40	1.74	9.39	1.60	n.s
Percent boys	73.7% (n=112)		76.4% (n=155)		n.s
FSIQ	99.07	12.18	95.91	13.39	.030
DISC-IV Inattentive symptoms	7.35	2.04	6.54	2.35	.001
DISC-IV Hyperactive symptoms	5.93	2.43	4.47	2.72	<.001

SD = Standard Deviation; FSIQ = Full Scale IQ; ODD=Oppositional Defiant Disorder

Table 2c. Demographic and clinical characteristics of children with and without CD					
	With CD n=45		Without CD n=310		p
	Mean	SD	Mean	SD	
Age	9.08	1.55	9.37	1.65	n.s
Percent boys	80% (n=36)		74.5% (n=231)		n.s
FSIQ	95.93	13.35	97.41	12.93	n.s
DISC-IV Inattentive symptoms	7.07	2.39	6.86	2.24	n.s
DISC-IV Hyperactive symptoms	6.78	1.92	4.85	2.71	<.001

SD = Standard Deviation; FSIQ = Full Scale IQ; CD = Conduct Disorder

one measure was available to assess planning (ToL total score) ANCOVA was used for this measure.

## Results

### Clinical Profile

Only 31% of children did not meet the criteria for any of the three examined comorbidities. Almost half of the group (44.5%) met the criteria for one or more anxiety disorders, 42.8% for ODD and 12.7% for CD. As many as 110 children were diagnosed with more than one comorbid disorder; 13.5% met the criteria for anxiety disorders only, 7.6%

for anxiety disorders and CD and 23.5% for anxiety disorders and ODD (Table 1).

No age ( $F(1,353) = .054, p = .816$ ) or sex differences ( $\chi^2(1, N = 355) = .527, p = .468$ ) were found between children with and without comorbidities (Tables 2a, 2b and 2c). Children with anxiety disorders and ODD presented with more inattentive symptoms (anxiety:  $F(1,353) = 9.34, p = .002$ ; ODD:  $F(1,353) = 11.45, p = .001$ ) and hyperactive symptoms (anxiety:  $F(1,353) = 5.71, p = .017$ ; ODD:  $F(1,353) = 27.22, p < .001$ ), while children with CD more hyperactive symptoms ( $F(1,353) = 21.24, p < .001$ ) compared to those without the comorbidities.

**Table 3. CPT and comorbid disorders**

	Anx		ODD		CD		Total
	Yes	No	Yes	No	Yes	No	
	M (SD)	M (SD)	M (SD)	M (SD)	M (SD)	M (SD)	
Omission errors T score	56.63 (14.81)	56.74 (15.57)	55.61 (14.49)	57.49 (15.71)	59.82 (15.82)	56.28 (15.1)	56.69 (15.21)
Commission errors T scores	54.74* (8.01)	53.4 (8.05)	54.55 (8.52)	53.59 (7.68)	53.86 (7.1)	54.02 (8.19)	54 (8.05)
RT T scores	52.31 (10.38)	51.52 (11.48)	51.06 (11.2)	52.48 (10.83)	53.26 (10.53)	51.67 (11.06)	51.87 (10.99)
RT Standard Error T scores	58.66* (10.7)	56.41 (10.3)	57.23 (10.66)	57.57 (10.45)	60.17 (9.52)	57.02 (10.62)	57.42 (10.53)
RT Block Change T scores	52.65 (12.62)	51.82 (11.5)	51.52 (11.75)	52.69 (12.2)	55.92 (11.43)	51.65 (12.01)	52.19 (12.01)
RT – ISI T scores	56.44* (14.65)	54.05 (12.07)	55.29 (13.82)	55.01 (12.98)	55.77 (12.83)	55.03 (13.42)	55.13 (13.33)

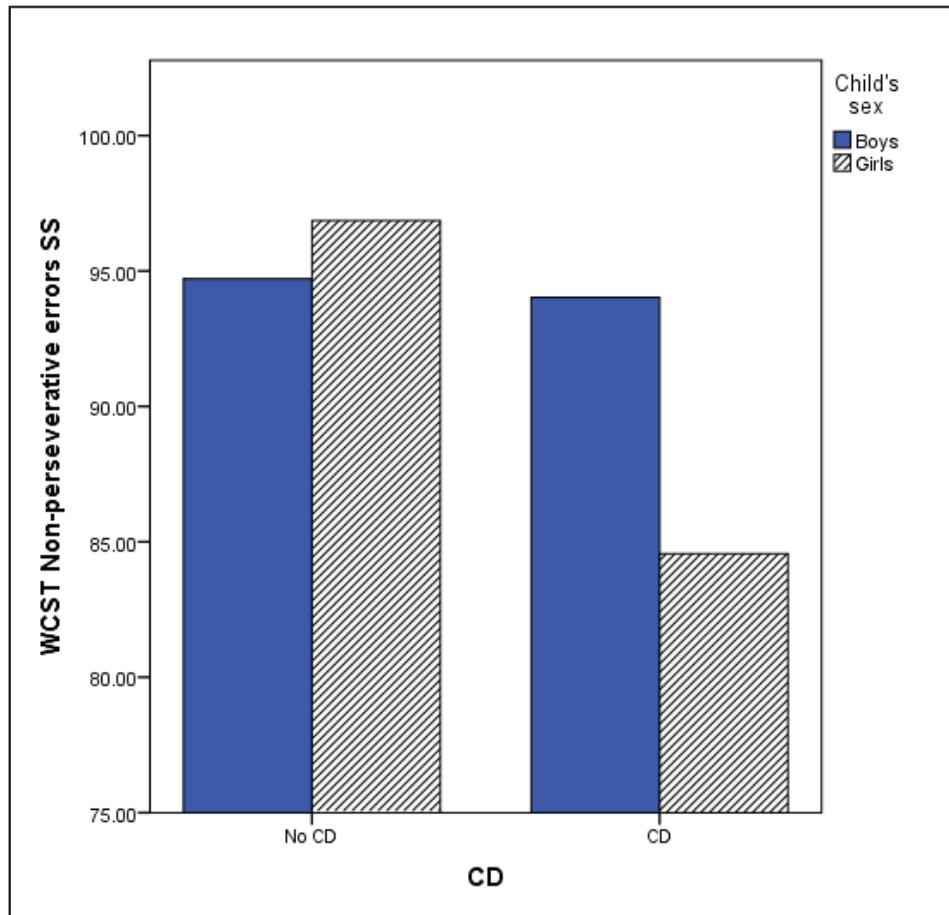
M = Mean; SD = Standard Deviation; RT = Reaction Time; ANX = anxiety disorders; CD = Conduct Disorder; ODD = Oppositional Defiant Disorder; No Com = no comorbid disorders; ISI = Inter –Stimuli – Interval.  
\* = p<.05

**Table 4. WCST and comorbid disorders**

	Anx		ODD		CD		Total
	Yes	No	Yes	No	Yes	No	
	M (SD)						
WCST Perseverative responses SS	97.2	98.99	98.6	97.9	94.9	98.66	98.2
	-13.54	-13.7	-15.26	-12.32	-13.08	-13.67	-13.64
WCST Perseverative Errors SS <sup>a</sup>	96.92	99.18	99.18	97.43	94.51*	98.71	98.18
	-13.43	-12.4	-13.46	-12.43	-12.98	-12.82	-12.89
WCST Non Perseverative Error SS <sup>a</sup>	94.13	95.45	95.41	94.45	92.13*	95.26	94.86
	-14.58	-14.92	-15.22	-14.43	-15.63	-14.62	-14.76
WCST number of categories completed SS <sup>a</sup>	4.28	4.6	4.48	4.44	3.89*	4.54	4.46
	-1.82	-1.66	-1.83	-1.67	-1.97	-1.69	-1.74
WCST Trials to complete first category SS <sup>b</sup>	24.73	21.32	22.68	22.96	32.13*	21.49	22.84
	-28.53	-21.97	-25.16	-25.15	-37.45	-22.56	-25.12
WCST Failure to maintain set SS	1.55	1.56	1.53	1.58	1.51	1.57	1.56
	-1.46	-1.38	-1.27	-1.52	-1.51	-1.4	-1.42

ANX = anxiety disorders; CD = Conduct Disorder; ODD = Oppositional Defiant Disorder; WCST = Wisconsin Card Sorting Test; SS = Standard Score; M = Mean; SD = Standard Deviation;  
\* = p < .05; <sup>a</sup>lower scores on this measure indicates poor performance; <sup>b</sup>higher score on this measure indicates poor performance.

Figure 1. WCST Non-persistent errors



Higher scores correspond to better performance; WCST = Wisconsin Card Sorting Test; CD = Conduct Disorder

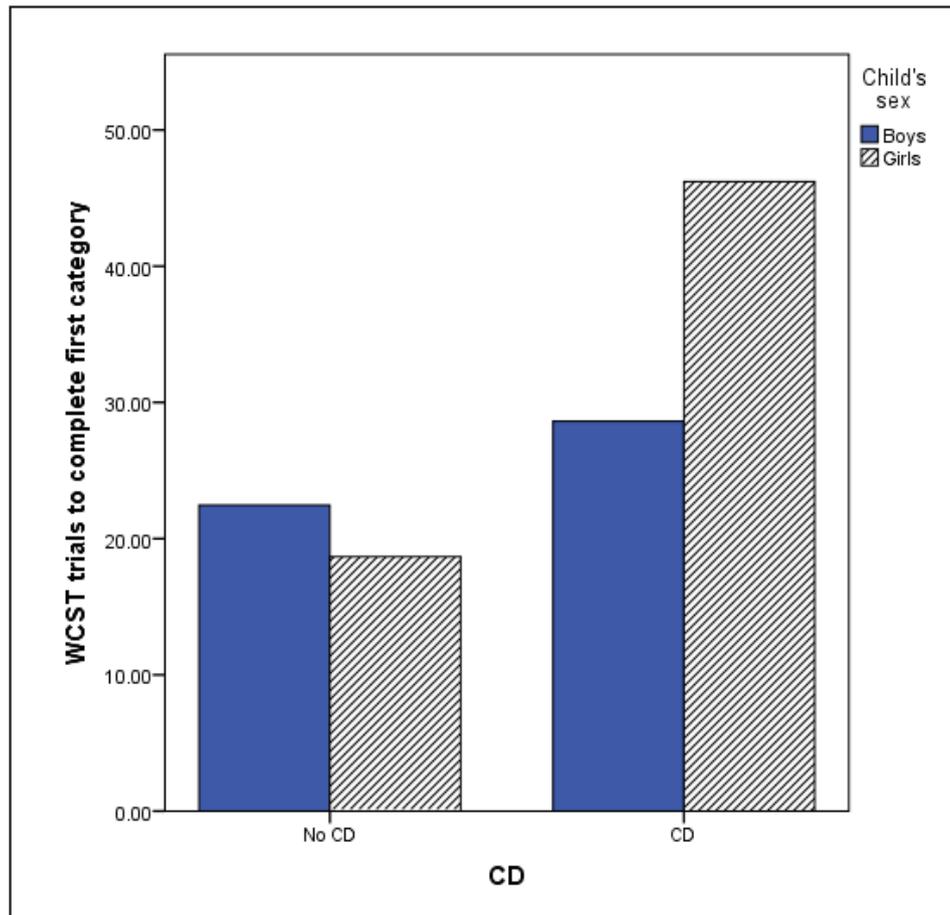
The Full Scale IQ for the group was within the average range (Tables 2a, 2b, and 2c). No significant IQ differences were found between boys and girls (FSIQ:  $F(1,325) = .081$ ,  $p = .776$ ). Children with ODD were found to have significantly higher scores on Full Scale IQ ( $F(1,325) = 4.77$ ,  $p = .030$ ), working memory ( $F(1,249) = 6.78$ ,  $p = .010$ ; with ODD:  $M = 95.64$ ,  $SD = 11.05$ ; without ODD:  $M = 91.58$ ,  $SD = 13.05$ ), and processing speed ( $F(1,249) = 7.37$ ,  $p = .007$ ; with ODD:  $M = 99.59$ ,  $SD = 13.47$ ; without ODD:  $M = 94.98$ ,  $SD = 13.16$ ) indices of WISC-IV compared to children without ODD. Significant differences were found between children with CD and without CD on verbal index when sex was used as a covariate ( $F(3,326) = 4.014$ ,  $p = .046$ ), children with CD having obtained lower scores compared to children without CD (with CD:  $M = 92.65$ ,  $SD = 14.02$ ; without CD:  $M = 96.83$ ,  $SD = 13.76$ ).

### Comorbid Disorders and Executive Function

**CPT.** Table 3 presents the CPT data. The MANCOVA results showed a significant main effect of anxiety disorders ( $F(6,337) = 2.693$ ,  $p = .014$ ) and age ( $F(6,337) = 19.654$ ,

$p < .001$ ). Children with anxiety disorders obtained higher T scores on commission errors, RT standard error, and RT inter-stimuli-interval change. No significant multivariate effect of CD ( $F(6,338) = 1.574$ ,  $p = .154$ ) or ODD ( $F(6,340) = .767$ ,  $p = .596$ ) was observed.

**WCST.** The MANCOVA using WCST variables (Table 4) showed no significant multivariate effect of anxiety disorders ( $F(6,348) = .645$ ,  $p = .694$ ) or ODD ( $F(6, 348) = 1.023$ ,  $p = .410$ ). However, significant multivariate CD ( $F(6,345) = 2.55$ ,  $p = .020$ ), age ( $F(6, 345) = 10.087$ ,  $p < .001$ ) and CD by sex interaction effects were observed ( $F(6, 345) = 2.173$ ,  $p = .045$ ). Children with CD made more perseverative and non-persistent errors, completed fewer categories and required more trials to complete the first category indicating poorer performance. Sex by CD interaction was found for non-persistent errors and number of trials to complete the first category. Boys with and without CD obtained similar scores on the non-persistent errors ( $M = 94.02$ ,  $SD = 16.17$ ;  $M = 94.71$ ,  $SD = 14.69$ ), while girls with CD made more errors ( $M = 84.55$ ,  $SD = 10.85$ ) compared with girls without CD ( $M = 96.86$ ,  $SD = 14.35$ ) or boys in either

**Figure 2. WCST trials to complete the first category**

Lower scores correspond to better performance; WCST = Wisconsin Card Sorting Test; CD = Conduct Disorder

group ( $p = .016$ ; Figure 1). Girls with CD also required more trials to complete the first category compared to boys and girls without CD, or boys with CD ( $p = 0.14$ ; Figure 2). This indicated that girls with CD had difficulty with initial concept formation.

**ToL.** On the planning task, a significant main effect of sex was observed ( $F(1,350) = 11.127, p = .001$ ). Boys, obtained higher scores ( $M = 111.28, SD = 13.57$ ) than girls ( $M = 105.04, SD = 16.93$ ). No significant effect of anxiety disorders ( $F(1, 353) = 1.856, p = .174$ ), CD ( $F(1,353) = .026, p = .873$ ) or ODD ( $F(1,353) = .080, p = .778$ ) was observed.

**Working Memory.** When sex and age were included in the MANCOVA model no significant effect of anxiety disorders was seen ( $F(2, 349) = 1.458, p = .234$ ). However if confounding variables were not used, a significant effect of anxiety disorders was observed ( $F(2,351) = 3.387, p = .035$ ). Since FW scores are standardised the FW was examined without controlling for age or sex. Univariate analysis of variance showed that children with anxiety disorders obtained lower scores on the FW task ( $F(1,353) = 6.62, p =$

$.010$ ). However, when only sex was used in MANCOVA the significant multivariate effect was no longer present ( $F(2,350) = 1.46, p = .552$ ). No other significant effects of comorbid disorders were observed (CD:  $F(2,352) = .497, p = .609$ ; ODD:  $F(2,352) = .433, p = .649$ ).

When multiple comorbidities were examined, such as anxiety disorders with CD or anxiety disorders with ODD, no significant differences were found between children with multiple comorbid disorders and those without multiple comorbid disorders.

## Discussion

The current study examined attention, inhibition and EF in children with ADHD, both with and without comorbid disorders in a large sample of children clinically diagnosed with ADHD. The initial hypotheses were partially supported as our findings show that the presence of comorbid CD or anxiety disorders affect children's EF performance. First, we found several differences in performance between children with and without anxiety disorders. Specifically, contrary to

previous reports (Bloemsa, et al., 2013) children with anxiety disorders had more difficulty with response inhibition, resulting in higher number of commission errors on CPT. While their overall reaction time for correct responses (RT-T) and change in reaction time across the test (RT-Block change) did not differ from children without anxiety disorders, children with anxiety disorders were more inconsistent in response speed (i.e. had a more erratic response style throughout the test as demonstrated by RT standard error) and as the speed between targets increased, their reaction time decreased (RT-ISI). Inconsistency in performance is now well recognized by clinicians working with the ADHD population. Our findings suggest that this inconsistency may be more pronounced in children who are also affected by anxiety disorders. Children with ADHD also have difficulties when faced with tasks that they deem challenging. In this study we observed that in laboratory setting, when the demands of the environment are increased, children with ADHD and anxiety disorders have difficulty adjusting to this change and as a result slow down their response. Contrary to other reports, we found no interaction between anxiety disorders and sex with respect to performance on the CPT task (Newcorn et al., 2001).

With respect to comorbid anxiety disorders and working memory our findings were in line with previous findings (Bedard & Tannock, 2008; Vance, Ferrin, Winther, Winther, & Gomez, 2013) as children with comorbid anxiety disorders had more working memory difficulties compared to those without anxiety disorders. This however was captured by only one of the working memory measures: the FW, and only when confounding variables (sex and age) were not included in the model. Difficulties with working memory may have a detrimental effect on children's ability to learn and their functioning in the classroom or in day-to-day activities. These findings further emphasize the significance of comorbid anxiety disorders. Anxiety disorders should be routinely evaluated in children diagnosed with ADHD. Furthermore, if children present with ADHD and anxieties in clinical setting, clinicians should be encouraged to assess the child's working memory abilities and if affected put in place intervention to decrease anxiety and accommodations allowing the child to compensate for anxieties as well as difficulties associated with working memory.

Presence of CD poses significant difficulties for children, their families and the society at large, due to associated risk factors. The findings of the current study bring to light potential deficits that may shape the phenotypic expression of the disorder. Contrary to other reports (Oosterlaan, Scheres & Sergeant, 2005), children in this study who were diagnosed with comorbid CD had more difficulties on the set-shifting task (WCST), making more perseverative and non-perseverative errors compared to their peers diagnosed with ADHD but without the comorbid CD. In addition, girls with CD had particular difficulties with initial concept formation and problem solving, as they required more trials

to complete the first category on the WCST. In addition to differences on the set-shifting task, children with comorbid CD had lower verbal IQ as compared to children without CD. This finding supports the recent report by Murray and Farrington (2010), who found low IQ to be a predictor of CD.

It is well established that the presence of CD is associated with antisocial outcomes and substance use later in life (Pardini & Fite, 2010), thus identifying the clinical profile of children with ADHD and CD is important for prevention, early intervention and effective treatment. Further investigation is required to confirm whether low verbal IQ, in combination with increased hyperactivity symptoms, and difficulties with cognitive flexibility are specific risk factors for developing CD in childhood as was seen in our study. It is possible that having more severe ADHD symptoms combined with difficulties in verbal communication may exacerbate behavioural problems, resulting in the kinds of disruptive behaviour seen in a diagnosis of CD. Prevention and treatment programmes should focus on intervention targeting children's language abilities, both expressive and receptive and cognitive flexibility, thus allowing children to develop necessary skills to respond in a socially adaptive manner to the demands of their environment.

In addition to sex differences seen between children with comorbid CD, we found sex differences in planning, as boys with ADHD performed better on the Tower of London task than did girls. Similar results were reported by O'Brien, Dowell, Mostofsky, Denckla and Mahone (2010) who compared girls and boys with ADHD using various EF tasks and found that girls demonstrated poorer performance on the planning test. The sex differences observed in this study may explain the differences in clinical presentation of ADHD between boys and girls, and could be explained by the neuroanatomical brain differences seen between boys and girls (Mahone & Wodka, 2008).

Finally, the overall clinical profile of the sample was similar to those reported in the literature. As expected, many children presented with at least one comorbid disorder (Gau et al., 2010). Multiple comorbidities were also frequent, which is similar to reports in the literature (Larson, Russ, Kahn, & Halfon, 2011), however having more than one comorbid disorder did not affect the EF task performance.

This study has several strengths, the first of which lays in addressing some of the methodological limitations in current literature. The large sample size allowed us to investigate subgroups of children with comorbidities, as well as examine sex differences. All the children in this study underwent a thorough diagnostic process, including clinical diagnosis by a child psychiatrist and vigorous parent interviews that included the level of impairment and interference with functioning in more than one setting as part of its diagnostic algorithm. The standardised EF assessment

procedure allowed to control for potential effects of medication and fatigue on EF task performance.

Alongside these strengths, the study has limitations as well. As we did not include a control group, no comparisons can be made with typically developing children. While the assessments for the comorbid disorders were made using a thorough diagnostic tool used with parents, the study would have benefited from the inclusion of a child report measure of comorbid disorders. Finally, no teacher ratings of ADHD symptoms were included in this analysis and the ADHD symptoms were based only on parent report.

## Conclusion

Our findings demonstrate that the presence of comorbidities, EF deficits, and sex differences may explain the heterogeneous nature of ADHD among elementary school-age children. These differences can also contribute to the inconclusive findings previously reported in the literature. Given the reported relationship between EF and social functioning (Miller & Hinshaw, 2010; Rinsky & Hinshaw, 2011) and EF and academic functioning (Miller & Hinshaw, 2010; Rogers, Hwang, Toplak, Weiss, & Tannock, 2011) in individuals with ADHD and comorbid disorders, identifying EF deficits should be an integral part of ADHD assessment. EF and cognitive deficits and comorbid disorders should also be central in intervention design for this population. In conclusion, to maximize treatment success, children diagnosed with ADHD should receive tailored treatment as opposed to one focusing solely on addressing ADHD symptoms. For example, children comorbid CD and with difficulties in the area of cognitive flexibility and language would benefit from skill training programme that will address these areas. Similarly, children whose working memory is affected can use strategies to compensate for this difficulty which will aid their functioning at school and in social settings. Further research is needed focusing on children with CD, particularly girls diagnosed with CD and ADHD and the developmental trajectories associated with development of CD in ADHD population.

## Acknowledgments / Conflicts of Interest

The authors would like to thankfully acknowledge the Douglas Institute ADHD research team for data collection and data entry work, specifically Sandra Robinson and Jacqueline Richard for conducting the EF assessments, and Nicole Pawliuk for her assistance. This study was supported by the Canadian Institutes of Health Research (CIHR) operating grant and CIHR doctoral scholarship awarded to the first author. Most of all, we would like to thank all children and their families who graciously agreed to participate in this study.

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