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Heterogeneity in ADHD: Neurocognitive predictors of peer, family, and academic functioning

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ABSTRACT
Childhood attention-deficit/hyperactivity disorder (ADHD) is associated with impairments in peer, family, and academic functioning. Although impairment is required for diagnosis, children with ADHD vary significantly in the areas in which they demonstrate clinically significant impairment. However, relatively little is known about the mechanisms and processes underlying these individual differences. The current study examined neurocognitive predictors of heterogeneity in peer, family, and academic functioning in a well-defined sample of 44 children with ADHD aged 8–13 years (M = 10.31, SD = 1.42; 31 boys, 13 girls; 81% Caucasian). Reliable change analysis indicated that 98% of the sample demonstrated objectively-defined impairment on at least one assessed outcome measure; 65% were impaired in two or all three areas of functioning. ADHD children with quantifiable deficits in academic success and family functioning performed worse on tests of working memory (d = 0.68 to 1.09), whereas children with impaired parent-reported social functioning demonstrated slower processing speed (d = 0.53). Dimensional analyses identified additional predictors of peer, family, and academic functioning. Working memory abilities were associated with individual differences in all three functional domains, processing speed predicted social functioning, and inhibitory control predicted family functioning. These results add to a growing literature implicating neurocognitive abilities not only in explaining behavioral differences between ADHD and non-ADHD groups, but also in the substantial heterogeneity in ecologically-valid functional outcomes associated with the disorder.

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ADHD; Social; Family; Academic; Functioning; Heterogeneity

Attention-deficit/hyperactivity disorder (ADHD) is a complex, chronic, and heterogeneous disorder of brain, behavior, and cognition that affects approximately 5% of school-aged children (American Psychiatric Association, 2013; Polanczyk, Willcutt, Salum, Kieling, & Rohde, 2014) at an annual cost of illness of $42 billion in the United States (US) alone (Pelham, Foster, & Robb, 2007). Although long treated as error variance, heterogeneity in symptoms and impairments are being increasingly
recognized as important considerations for refining our understanding of ADHD pathogenesis and improving treatment outcomes (Koehler et al., 2013; Nigg, Willcutt, Doyle, & Sonuga-Barke, 2005). To this end, a growing body of research has identified factors associated with within-group heterogeneity in ADHD behavioral symptom presentation, including demographic characteristics such as gender (Gaub & Carlson, 1997) and age (Halperin, Trampush, Miller, Marks, & Newcorn, 2008), informant and setting factors (Valo & Tannock, 2010; Whalen et al., 1978), medical and behavioral treatment (van der Oord, Prins, Oosterlaan, & Emmelkamp, 2008), and neurocognitive abilities (Chhabildas, Pennington, & Willcutt, 2001; Halperin et al., 2008; Nigg, Blaskey, Huang-Pollock, & Rappley, 2002; Sarver, Rapport, Kofler, Raiker, & Friedman, 2015). In contrast, relatively less is known about the mechanisms and processes associated with heterogeneity in daily functioning among children with ADHD. This relative paucity of research is surprising given that functional impairments may be better predictors of long-term clinical outcomes than core ADHD behavioral symptoms (Pelham, Fabiano, & Massetti, 2005). The goal of the current study is to examine factors associated with heterogeneity in peer, family, and academic functioning in a well-defined sample of children with ADHD, with a particular focus on neurocognitive abilities that (a) are also characterized by significant within-group heterogeneity among children with ADHD (Fair, Bathula, Nikolos, & Nigg, 2012; Rajendran, O'Neill, Marks, & Halperin, 2015), and (b) have been shown previously to help explain the disorder’s behavioral symptoms and functional deficits relative to typically developing children at the group level (Rapport, Orban, Kofler, & Friedman, 2013).

Childhood ADHD is associated most frequently with impairments in three primary areas: peer, family, and academic functioning (Pelham et al., 2005). Interestingly, although impairment is required for diagnosis (American Psychiatric Association, 2013), children with ADHD vary significantly in the areas in which they demonstrate clinically significant impairment. For example, an estimated 50% to 80% of children with ADHD exhibit peer relational (social) problems (de Boo & Prins, 2007; Huang-Pollock, Mikami, Pfiffner, & McBurnett, 2009). Stated differently, these figures suggest that approximately 20% to 50% of these children are not viewed as experiencing clinically significant social problems. Similarly, rates of academic underachievement and learning difficulties are estimated to occur in 33% to 63% of children with ADHD across academic domains (Mayes & Calhoun, 2006). Impaired family functioning occurs in 62% to 87% of cases based on meta-analysis (Theule, Wiener, Tannock, & Jenkins, 2013), and includes a variety of difficulties involving parental perceptions of lower attachment, warmth, and connectedness in the parent–child relationship (Keown & Woodward, 2002), impaired parent–child communication (Cussen, Sciberras, Ukoumunne, & Efron, 2012; Keown & Woodward, 2002), and lower levels of parental confidence (Johnston & Mash, 2001) and parental involvement (Rogers, Wiener, Marton, & Tannock, 2009). Collectively, the significant variation in impairment rates among children with ADHD highlights the heterogeneity in functional consequences for these children and underscores the importance of understanding predictors of cross-domain impairment risk. However, with the exceptions reviewed below, relatively little is known about the mechanisms and processes underlying this heterogeneity (Nigg, 2005). This gap in turn constrains our ability to understand and ultimately
predict the extent to which individual children with ADHD are likely to develop impairments in each functional area.

Neurocognitive heterogeneity is a particularly appealing candidate to explain functional heterogeneity among children with ADHD for at least three reasons. First, the neurocognitive functions implicated in ADHD have been linked developmentally with a wide array of academic (Barry, Lyman, & Klinger, 2002; Thorell, 2007) and social/peer outcomes (Clark, Prior, & Kinsella, 2002; Dennis, Brotman, Huang, & Gouley, 2007; Holmes, Kim-Spoon, & Deater-Deckard, 2016). For example, developmental research suggests strong links between children’s working memory abilities and their social (Alloway et al., 2005) and academic functioning (Thorell, 2007). In particular, phonological working memory shows strong cross-sectional and longitudinal continuity with academic success in reading (Cain, Oakhill, & Bryant, 2004; Sarver et al., 2012; Sesma, Mahone, Levine, Eason, & Cutting, 2009), whereas visuospatial working memory may predict math productivity better than PHWM (Maybery & Do, 2003; Sarver et al., 2012). Similarly, inhibition has been linked with social functioning (Gewirtz, Stanton-Chapman, & Reeve, 2009; Nigg, 1999) as well as math (Thorell, 2007; Wåhlstedt, Thorell, & Bohlin, 2009), English, and science achievement (St. Clair-Thompson & Gathercole, 2006), and processing speed predicts academic performance in reading, math, and written expression in non-ADHD samples (Mayes & Calhoun, 2007).

Second, clinical research suggests that many but not all children with ADHD have deficits in any given aspect of neurocognitive functioning. For example, meta-analytic effect sizes indicate that up to 80% of children with ADHD may exhibit working memory deficits (Kasper, Alderson, & Hudec, 2012), while approximately 0% to 38% have inhibition deficits (Alderson, Rapport, & Kofler, 2007; Willcutt, Doyle, Nigg, Faraone, & Pennington, 2005), and 41% to 45% demonstrate slowed processing speed (Frazier, Demaree, & Youngstrom, 2004; Kofler et al., 2013) based on comparisons with typically developing groups (Zakzanis, 2001). Conservatively computed based on Cohen’s $d$ effect sizes as the percentage of non-overlap between the ADHD and non-ADHD population distributions (i.e., the percentage of children with ADHD scoring outside the typically developing range) as recommended (Zakzanis, 2001).

Third, experimental studies suggest that specific neurocognitive functions may account for ADHD-related deficits in behavioral symptoms and at least some aspects of functioning at the group level. Much of this research has focused on working memory, and suggests that experimentally increasing working memory demands evokes differential decreases in objectively-measured attentive behavior (Kofler, Rapport,
Bolden, Sarver, & Raiker, 2010) and increases in gross motor activity (hyperactivity; Kofler, Sarver, & Wells, 2015; Rapport et al., 2009) for ADHD relative to typically developing groups. In addition, cross-sectional mediation models suggest minimal differences between ADHD and typically developing groups with regard to impulsive responding (Raiker, Rapport, Kofler, & Sarver, 2012), inhibitory control (Alderson, Rapport, Hudec, Sarver, & Kofler, 2010), delay aversion (Patros et al., 2015), and response variability (Kofler et al., 2014) after accounting for working memory. Importantly, working memory deficits also appear important for explaining between-group differences in social functioning (Bunford et al., 2015; Kofler et al., 2011) and math performance (Antonini et al., 2016), suggesting an important role for this cognitive ability in ecologically-valid, functional outcomes for children with ADHD.

To our knowledge, however, no studies of childhood ADHD have simultaneously examined the role of multiple neurocognitive functions (e.g., inhibition, processing speed) in explaining impairments in social or family functioning.

To summarize, the impetus for examining the link between neurocognitive abilities and functional heterogeneity in ADHD comes from converging lines of research indicating that (a) neurocognitive abilities predict important functional outcomes in non-ADHD samples (Holmes et al., 2016; Thorell, 2007), (b) many but not all children with ADHD exhibit deficits in specific neurocognitive abilities and each area of functioning (Pelham et al., 2005; Rapport et al., 2013), and (c) neurocognitive deficits may explain behavioral and functional impairments in ADHD at the group level (Chacko, Kofler, & Jarrett, 2014).

To this end, emerging evidence suggests that neurocognitive abilities may help explain heterogeneity in social (Miller & Hinshaw, 2010), academic (Biederman et al., 2004; Preston, Heaton, McCann, Watson, & Selke, 2009), and global functioning (Cheung et al., 2015) among children and adolescents with ADHD. Specifically, individual differences in IQ among children with ADHD predict their reading, math (Alloway & Stein, 2014), and spelling success (Preston et al., 2009). Beyond this most general cognitive estimate (Dennis et al., 2009), individual differences in working memory components predict concurrent reading, math, and overall academic achievement (Alloway & Stein, 2014; Mayes & Calhoun, 2007; Rogers, Hwang, Toplak, Weiss, & Tannock, 2011) and longitudinally predict their reading abilities into young adulthood (Miller, Nevada-Montenegro, & Hinshaw, 2012). Similarly, children with ADHD with faster processing speed show higher attainment in reading (Jacobson et al., 2011), as well as math and written expression (Mayes & Calhoun, 2007). Further, subgroups of children with ADHD defined by the quantity of their neurocognitive deficits differ in academic attainment and grade retention (Biederman et al., 2004). In contrast, to our knowledge no ADHD study has examined the extent to which individual differences in behavioral inhibition predict academic heterogeneity, examined the relation between neurocognitive task performance and family functioning, or simultaneously examined the impact of multiple neurocognitive abilities on functional outcomes.

The current study is the first to examine neurocognitive predictors of heterogeneity in academic, peer, and family functioning among children with ADHD while also considering several known risk factors and correlates of academic and social difficulties in ADHD such as age, socioeconomic status (SES), ADHD subtype/presentation,
medication status, and gender. We selected global cognitive functioning (IQ) and four primary neurocognitive functions—phonological working memory, visuospatial working memory, behavioral inhibition, and processing speed—given the large bodies of research on these abilities in ADHD and developmental evidence linking each with one or more of the functional outcomes as described above. We predicted that a majority of children with ADHD would exhibit quantifiable, objectively-defined deficits in each area of functional impairment (peer, family, academic), and that children with deficits in each area would demonstrate identifiable neurocognitive profiles. We expected dimensional analyses to be consistent with these between-group findings (functional impairment vs. no impairment), such that working memory abilities would predict individual differences in social problems (Bunford et al., 2015; Kofler et al., 2011) and each of the neurocognitive constructs would predict individual differences in academic functioning given the developmental and clinical findings reviewed above. No predictions regarding family functioning were made due to the paucity of research.

**Method**

**Participants**

The sample comprised 44 children aged 8 to 13 years ($M = 10.31$, $SD = 1.42$; 31 boys, 13 girls) from the Southeastern US, who were consecutive referrals to a children’s learning clinic (CLC) through community resources for a psychoeducational assessment and participation in a behavioral ($n = 37$) or cognitive training ($n = 7$) treatment study. Pretreatment data were used in the current study. Working memory performance data were reported for a subset of the current sample in Kofler et al. (2015) to examine conceptually-unrelated hypotheses. Psychoeducational evaluations were provided to the parents of all participants. All parents and children gave informed consent/assent, and the university’s institutional review board approved the study prior to the onset of data collection.

**Group Assignment**

All children and their parents participated in a detailed, semi-structured clinical interview using the Kiddie Schedule for Affective Disorders and Schizophrenia for School-Aged Children (K-SADS; Kaufman et al., 1997). The K-SADS (2013 Update) assesses onset, course, duration, severity, and impairment of current and past episodes of psychopathology in children and adolescents based on Diagnostic and Statistical Manual of Mental Disorders - Fifth Edition (DSM-5; American Psychiatric Association, 2013) criteria. Its psychometric properties are well established, including inter-rater agreement of .93 to 1.00, test–retest reliability of .63 to 1.00, and concurrent (criterion) validity between the K-SADS and psychometrically established parent rating scales (Kaufman et al., 1997).

K-SADS interviews were supplemented with parent and teacher ratings scales from the Behavior Assessment System for Children – Second Edition (BASC-2; Reynolds & Kamphaus, 2004) and Child Symptom Inventory – Fourth Edition (CSI-IV; Gadow, Sprafkin, & Salisbury, 2004). Children with any ADHD subtype/presentation were
eligible given evidence of the instability of ADHD subtypes (Valo & Tannock, 2010) and previous research implicating neurocognitive processes in both inattentive (Kofler et al., 2010) and hyperactive (Rapport et al., 2009) symptom clusters.

A total of 44 children met the following criteria and were included in the ADHD group: (1) an independent diagnosis by the CLC’s directing clinical psychologist using DSM-5 criteria for ADHD based on K-SADS interviews; (2) parent ratings at least 1.5 SDs above the mean on the Attention Problems and/or Hyperactivity clinical syndrome scales of the BASC-2 parent form, or exceeding the criterion score for the parent version of the ADHD-Inattentive and/or ADHD-Hyperactive/Impulsive subscales of the CSI-IV; and (3) teacher ratings at least 1.5 SDs above the mean on the Attention Problems and/or Hyperactivity clinical syndrome scales of the BASC-2 teacher form, or exceeding the criterion score for the teacher version of the ADHD-Inattentive and/or ADHD-Hyperactive/Impulsive subscales of the CSI-IV. Four children with ADHD failed to meet the teacher cutoff criteria, likely due to behavior well controlled on medication. In these cases, previous psychoeducational evaluations were available that documented cross-setting behavioral symptoms and impairment. In accordance with DSM-5, all children had current impairments based on K-SADS parent interview.

Of the 44 children with ADHD (13 girls), 18 met “AND” criteria for Combined, 23 for Inattentive, and 3 for Hyperactive/Impulsive Presentation. The “AND” criteria required the child to meet symptom thresholds based on both parent and teacher report (Willcutt et al., 2012). For example, the Combined presentation was specified for children who met/exceeded symptom thresholds for both Inattentive and Hyperactive/Impulsive symptom clusters for both informants. To improve generalizability (Wilens et al., 2002), children with comorbidities were included. Comorbidities reflect clinical consensus best estimates based on parent and child K-SADS interviews, child psychoeducational testing, and multiple parent, child, and teacher norm-referenced questionnaires. In all cases, K-SADS interview indicated that the onset of ADHD symptoms preceded the onset of comorbid symptoms, and that the child’s inattention and/or hyperactive/impulsive symptoms could not be better accounted for by the comorbid condition. Comorbidities included oppositional defiant disorder (11%), depressive disorders (16%), and anxiety disorders (18%). None of the children screened positive for specific learning disorders in reading, math, or oral language based on DSM-5-recommended standard scores > 1.5 SDs below the normative sample mean (American Psychiatric Association, 2013, p. 69) on the Kaufman Test of Educational Achievement, Second or Third Edition (age norms; Kaufman & Kaufman, 2004, 2014); one child screened positive for deficits in written language. Child race/ethnicity included Caucasian non-Hispanic (81%), Hispanic English-speaking (7%), Asian (5%), African American (2%), and mixed racial/ethnic (5%) backgrounds.

Children were excluded from the study if they presented with (a) gross neurological, sensory, or motor impairment, (b) history of a seizure disorder, (c) psychosis, (d) autism spectrum disorder, (e) an FSIQ score less than 80, and/or (f) non-stimulant medications that could not be withheld for testing. Twenty-two of the 44 children with ADHD were prescribed psychostimulants; medication was withheld for a minimum of 24 hours prior to both research testing sessions given evidence of psychostimulant effects on processing speed and other non-executive aspects of neurocognitive task performance (cf. Rapport et al., 2013).
Procedures

All children participated in two consecutive Saturday testing sessions following the baseline psychoeducational assessment. Neurocognitive tasks were administered as part of a larger battery of laboratory tasks that required the child’s presence for approximately 3 hours per session. All tasks were counterbalanced across testing sessions to minimize order effects. Children were seated in a caster-wheel swivel chair approximately 0.66 m from the computer monitor for all tasks. Performance was monitored at all times by the examiner, who was stationed just out of the child’s view to provide a structured setting while minimizing performance improvements associated with examiner demand characteristics (Gomez & Sanson, 1994). All children received brief (2–3 minute) breaks after each task, and longer (10–15 minute) breaks after every two to three tasks to minimize fatigue.

Neurocognitive Performance

Phonological and Visuospatial Working Memory

The phonological and visuospatial working memory tasks developed by Rapport et al. (2008) were used for the current study. Previous studies of ADHD and typically developing children indicate large magnitude differences in these tasks (Kofler et al., 2014; Patros et al., 2015; Rapport et al., 2008), and performance on these tasks predicts ADHD-related impairments in objectively-measured activity level (Rapport et al., 2009), attentive behavior (Kofler et al., 2010), impulsivity (Patros et al., 2015; Raiker et al., 2012), inhibitory control (Alderson et al., 2010), and social dysfunction (Kofler et al., 2011). Evidence for the reliability and validity of these working memory tasks includes high internal consistency (α = .82 to .97), one- to three-week test–retest reliability of .76 to .90 (Sarver et al., 2015), and demonstration of the expected magnitude of relations (Swanson & Kim, 2007) with established measures of short-term memory (Raiker et al., 2012). Six trials were administered at each set size (3, 4, 5, or 6 stimuli) based on re-analysis of data demonstrating that all 6-trial versions correlate ≥ .90 with the corresponding 12-trial versions reported in Kofler et al. (2015). The 24 total trials (6 trials at each set size) were randomized and then grouped into two blocks of 12 trials each, such that the stimulus set size for a given trial was not predictable based on the preceding trial. Mixed presentation was selected given evidence that it results in higher central executive working memory demands due to memory set unpredictability relative to sequential presentation (Conway et al., 2005; Kofler et al., 2015). Five practice trials were administered before each working memory task; children were required to achieve 80% correct before advancing to the full task. Children received short breaks between each 12-trial block (approximately 1–2 minutes). Task duration was approximately 2.5 (visuospatial) to 3.5 (phonological) minutes per block for the phonological and visuospatial tasks described below.

Phonological Working Memory (PHWM) Task. The PHWM task is similar to the Letter-Number Sequencing subtest on the Wechsler Intelligence Scale for Children – Fifth Edition (WISC-V; Wechsler, 2014), and assesses PHWM based on Baddeley’s (2007) model. Children were presented a series of jumbled numbers and a letter at a
rate of 1 stimuli per second. The letter was never presented in the first or last position of the sequence to minimize potential primacy and recency effects, and was counter-balanced across trials to appear an equal number of times in the other serial positions (i.e., positions 2, 3, 4, or 5). Children were instructed to recall the numbers in order from smallest to largest, and to say the letter last (e.g., 4 H 6 2 is correctly recalled as 2 4 6 H). Two trained research assistants, shielded from the participant’s view, recorded oral responses independently (inter-rater reliability is 99.50%).

**Visuospatial Working Memory (VSWM) Task.** Children were shown nine squares arranged in three offset vertical columns on a computer monitor. The columns were offset from a standard 3 × 3 grid to minimize the likelihood of phonological coding of the stimuli (e.g., by equating the squares to numbers on a telephone pad). A series of 2.5 cm diameter dots (3, 4, 5, 6) were presented sequentially in one of the nine squares during each trial such that no two dots appeared in the same square on a given trial. All but one dot was black; the exception being a red dot that never appeared as the first or last stimulus in the sequence. Each dot was displayed for 800 ms, followed by a 200 ms interstimulus interval. Children were instructed to respond by pressing the corresponding squares on a modified computer keyboard, and to reorder the dot locations by indicating the serial position of the black dots in the order presented, followed by the serial position of the red dot last.

**Dependent Variables: Working Memory Task Performance.** Performance data were collected for each trial for each participant. The randomized trials were collated during post-processing to allow estimation of performance at each stimulus set size (3, 4, 5, 6). Partial-credit unit scoring (i.e., stimuli correct per trial) was used to index overall working memory performance at each set size, as recommended (Conway et al., 2005).

**Behavioral Inhibition and Processing Speed Stop-Signal Task.** The stop-signal task and administration instructions were identical to those described in Schachar, Mota, Logan, Tannock, and Klim (2000) and Alderson, Rapport, Sarver, and Kofler (2008). Psychometric evidence includes high internal consistency and three-week test–retest reliability (.72), as well as convergent validity with other inhibitory control measures (Soreni, Crosbie, Ickowicz, & Schachar, 2009). Go-stimuli were displayed for 1000 ms as uppercase letters X and O positioned in the center of a computer screen (500 ms interstimulus interval; total trial duration = 1500 ms). Xs and Os appeared with equal frequency throughout the experimental blocks. A 1000 Hz auditory tone (i.e., stop-stimulus) was presented randomly on 25% of trials. Stop-signal delay (SSD)—the latency between presentation of go- and stop-stimuli—was initially set at 250 ms and dynamically adjusted ± 50 ms contingent on participant performance. Successfully inhibited stop-trials were followed by a 50 ms increase in SSD, and unsuccessfully inhibited stop-trials were followed by a 50 ms decrease in SSD. The algorithm was designed to approximate successful inhibition on 50% of the stop-trials. In the current study, inhibition success was 53.9%, 54.3%, 51.0%, and 51.3% across the four experimental blocks. All participants completed two practice blocks and four consecutive experimental blocks of 32 trials per block (24 go-trials, 8 stop-trials per block).
Dependent Variables: Inhibition. SSD at each of the four blocks served as the primary indices of behavioral inhibition. SSD was selected based on conclusions from recent meta-analytic reviews that it is the most direct measure of behavioral inhibition in stop-signal tasks that utilize dynamic SSDs, given that SSDs change systematically according to inhibitory success or failure (Alderson et al., 2007; Ilieva, Hook, & Farah, 2015; Lijffijt, Kenemans, Verbaten, & van Engeland, 2005).

Dependent Variables: Processing Speed. Mean choice reaction time (MRT) to correct go-trials during each of the four stop-signal blocks served as the primary indices of processing speed. Anticipatory responses (reaction times < 150 ms) were excluded as recommended (Schmiedek, Oberauer, Wilhelm, Süß, & Wittmann, 2007).

Global Intellectual Functioning (IQ)

All children were administered the Wechsler Abbreviated Scales of Intelligence –Second Edition (WASI-II; Wechsler, 2011; n = 35), WISC-IV (Wechsler, 2003; n = 2), or WISC-V (Wechsler, 2014; n = 7) to obtain an overall estimate of intellectual functioning. Full Scale IQ (FSIQ) was not analyzed because FSIQ performance depends heavily on the neurocognitive constructs described above (Ackerman, Beier, & Boyle, 2005; Dennis et al., 2009). Following Rapport et al. (2008) and Kofler et al. (2013), we computed a residual FSIQ score by covarying the working memory, inhibition, and processing speed factor scores, described below, out of FSIQ ($R^2 = .30$, $p = .006$). This residual FSIQ score represents cognitive functions important for IQ test performance other than these neurocognitive constructs, and was examined as a potential predictor in the analyses described below. Importantly, our method of removing the influence of working memory from IQ assumes that working memory influences IQ rather than vice versa. This assumption is based on a large and compelling cognitive literature showing working memory to be an important predictor of global IQ (cf. Engle, Tuholski, Laughlin, & Conway, 1999; Giofrè, Mammarella, & Cornoldi, 2013; Tourva, Spanoudis, & Demetriou, 2016), as well as specific developmental evidence that age-related improvements in working memory lead directly to improvements in IQ (Tourva et al., 2016). Thus, we propose that it is reasonable to conclude that the shared variance between working memory and IQ is, in large part, attributable to working memory’s influence on IQ rather than vice versa.

We considered using the General Ability Index (GAI) rather than FSIQ$_{\text{residual}}$ given the conceptual interpretation of GAI as an IQ estimate that is free from the influence of working memory and processing speed (Wechsler, 2014). However, the construct validity of this interpretation appears to be limited. Specifically, the WISC-V Technical and Interpretive Manual (Wechsler, 2014, p. 74, table 5.1) indicates that GAI correlates .61 with the Working Memory Index (WMI), indicating significant influence of working memory on this “process-free” estimate (the WMI-FSIQ correlation of .72 is similarly high, despite the interpretive manual recommendation to conceptualize GAI as IQ without the influence of working memory). Similarly, the WASI-2 FSIQ, which is conceptually GAI because it is comprised of the same subtests used to calculate GAI on the WISC-IV, correlates .88 with the WISC-IV FSIQ (Wechsler, 2011, p. 131), again suggesting that the conceptual distinction between GAI and FSIQ is limited. We also note that according to
the Wechsler manuals, GAI was not created or verified via factor analysis like FSIQ but is rather a conceptually-derived estimate (Wechsler, 2014, p. 16). The statistical overlap between WMI and GAI suggests limited utility of this index for estimating process-free IQ abilities, and suggests that its raw inclusion in the model would result in removing significant variance attributable to working memory from working memory (cf. Dennis et al., 2007; Rapport et al., 2008).

**Neurocognitive Dimension Reduction**

**Control for the Task Impurity Problem.** To address the task impurity problem pervasive within neurocognitive measurement (Snyder, Miyake, & Hankin, 2015), we used a dimension reduction approach to isolate reliable variance associated with each neurocognitive construct and approximate the removal of all random and task-specific, non-construct error (Conway et al., 2005). Because no task is process pure, generalizability of results requires experimenters to use multiple measures and create factor scores to estimate common variance associated with each construct (for review and specific examples, see Shipstead, Redick, & Engle, 2010). Following Kofler et al. (2013), this involved creating a factor score for each neurocognitive construct using a principal components factor analysis on the 16 neurocognitive performance variables (4 blocks each for PHWM, VSWM, SSD, and MRT; 78.01% of variance accounted for; construct-specific factor loadings $r = .68$ to $.92$; Supplementary Table 1). The ratio of participants (44) to factors (4) was deemed acceptable (Hogarty, Hines, Kromrey, Ferron, & Mumford, 2005). By design, the intercorrelations among the derived PHWM, VSWM, Behavioral Inhibition, and Processing Speed variables were $r_{all} = .00$ ($p > .99$). Higher scores reflect better working memory and inhibition but slower processing speed.

**Peer, Family, and Academic Functioning**

Nationally standardized, psychometrically sound, and widely used instruments were used to obtain estimates of overall peer, family, and academic functioning. Parents and teachers were asked to consider the child’s behavior when off his or her medication.

**Peer (Social) Functioning**

The BASC-2 (Reynolds & Kamphaus, 2004) parent and teacher forms are 160- and 139-item scales, respectively, that assess internalizing and externalizing behavior problems in children aged 2 to 21 years. Raw scores are converted to age- and gender-specific $T$-scores based on the national standardization sample ($N = 1800$ per form). The parent and teacher Social Skills subscales each contain 9 items that index children’s peer/social functioning (six-week test–retest = $.84$ to $.86$, $\alpha = .87$ to $.92$). The parent and teacher social skills composite scores served as the primary indices of social functioning at home and school, respectively. Higher scores reflect better social functioning.

**Family Functioning**

The Parenting Relationship Questionnaire (PRQ; Kamphaus & Reynolds, 2006) is a 71-item parent report scale that assesses family functioning across seven domains (national standardization, $N = 4130$). $T$-scores are obtained for each factor according to age and gender; no total PRQ score is computed. Four subscales were selected that were thought
to be most relevant to the parent–child relationship: Parent-Child Attachment, Parent-Child Communication, Parent-Child Involvement, and Parenting Confidence (four- to five-week test–retest = .76 to .84, $\alpha = .82$ to .88). Higher scores reflect better perceived attachment, involvement, communication, and parenting efficacy.
**Academic Functioning**

The Academic Performance Rating Scale (APRS; DuPaul, Rapport, & Perriello, 1991) was completed by each child’s teacher to assess academic functioning (two-week test–retest = .93 to .95, α = .94 to .95). The APRS contains subscales that reflect academic productivity and success. The Academic Productivity scale is comprised of 12 items that assess academic efficiency (e.g., percentage of classwork completed correctly) and consistency, following group instructions, and completing work in a timely manner. The Academic Success subscale contains 7 items that assess the quality of reading and spoken work, how quickly children learn new material, and how well they retain new information; T-scores were obtained by comparing performance to the standardization sample (N = 487) according to age and gender. Higher scores reflect better academic functioning.

**Socioeconomic Status (SES)**

SES was estimated using the Hollingshead (1975) scoring based on the education and occupation of the caregiver(s).

**Data Analysis Overview**

The analytic plan was executed in two tiers. The first tier examined functional heterogeneity in ADHD by quantifying the extent to which our sample exhibited impairments in each functional area relative to published age and gender norms, and examining between-group differences in neurocognitive abilities across Impaired vs. Not Impaired subgroups for each functional outcome. Following Sarver et al. (2015), this involved applying the Jacobson and Truax (1991) model of reliable change to each child’s norm-referenced scores on each of the peer, family, and academic outcomes. This method was selected over static cut points (e.g., 1 SD below the mean) because it improves precision by explicitly accounting for measurement unreliability (Jacobson & Truax, 1991). Children were classified as Impaired or Not Impaired on each functional outcome based on whether their norm-referenced score was reliably below the normative sample (i.e., difference exceeded chance at p < .05). This classification was based on computation of the Reliable Change Index (RCI), which is the ratio of the difference between the child’s score and the test mean divided by standard error (computed using each measure’s reported test–retest reliability and the SD of the normative sample; Rule B; Jacobson & Truax, 1991) individually for each child for each outcome. Reported test–retest reliability across all tests/subscales was .76 to .95. The RCI is tested against the z distribution; impairment is defined as a score that is significantly worse than the test mean given the test’s SD and reported reliability. We then compared the neurocognitive performance of children defined as Impaired vs. Not Impaired on each functional outcome using bias-corrected, bootstrapped Cohen’s d effect sizes. Inspection of the RCI data indicated that the impairment cutoffs centered around 1 SD below the normative sample mean across measures; statistical significance was obtained at different cut points across measures dependent on each measure’s test–retest reliability (i.e., for tests with lower reliability, scores further from the mean were required to conclude with p < .05 certainty that the child’s score was more likely to come from the...
dysfunctional/impaired population than the functional population). To further probe individual differences in functioning and capitalize on the increased power of continuous vs. dichotomous variables, the second tier used a dimensional approach to examine neurocognitive predictors of norm-referenced T-scores for peer, family, and academic functioning among children with ADHD.

Bootstrapping

All analyses were completed utilizing a bias-corrected bootstrapping procedure to minimize Type II error, as recommended by Shrout and Bolger (2002). Bootstrapping is appropriate for total sample sizes as low as 20 (Efron & Tibshirani, 1993); the bias-corrected, bootstrapped 95% confidence intervals (CIs) were used to estimate effect magnitude and determine statistical significance for all comparisons. SPSS v22 (IBM Corp, 2013) was used for all analyses, and 10,000 samples were derived from the original sample (N = 44) by a process of resampling with replacement (Shrout & Bolger, 2002).

Results

Power Analysis

Given the relatively small sample size, we conducted a power analysis using GPower v3.1 (Faul, Erdfelder, Lang, & Buchner, 2007) to determine our sensitivity for detecting effects. The between-group analyses (Impaired vs. Not Impaired) are powered to detect large effects (d ≥ .80) based on our sample size of 44 for power = .80 and α = .05; we therefore report bias-corrected, bootstrapped Cohen’s d effect sizes and interpreted 95% CIs rather than p values, given their robustness to distributional characteristics (Fritz & MacKinnon, 2007). Minimal guidance was available for a priori selection of expected effect sizes due to the paucity of studies defining heterogeneity based on functional outcomes rather than cognitive or behavioral symptoms in ADHD. However, effects of this magnitude were considered reasonable based on meta-analyses indicating large magnitude relations between ADHD and neurocognitive abilities (e.g., working memory; Kasper et al., 2012), and between ADHD and each area of functional impairment as reviewed above. In addition, we supplemented the between-group analyses with linear regression to capitalize on the increased power associated with continuous relative to dichotomous variables. Power analysis for regression indicated that we were adequately powered to reliably detect effects of $\rho^2 = .27$ for power = .80, α = .05, and six predictors (PHWM, VSWM, SSD, MRT, IQ, and one covariate as described below) based on our sample size of 44.

Preliminary Analyses

Means and SDs for each outcome variable are shown in Table 1. All variables were screened for univariate/multivariate outliers and tested against $p < 0.001$. No significant outliers were found. One-sample T-tests revealed that the BASC-2 parent and teacher
Attention Problems scores (for both Combined and Inattentive presentations) and Hyperactivity scores (for the ADHD-Combined group) were significantly elevated relative to the scale’s T-score mean of 50 as expected (all $p < .0005$; Table 1). Age, SES, gender, ADHD subtype/presentation, and medication status were not significantly related to any of the peer, family, or academic outcomes (all 95% CIs substantially overlap 0.0; all $p \geq .20$), with the following exceptions: child age was related to Parent-Child Attachment ($r = .38, 95\% CI = .11$ to .60, $p = .02$), Parent-Child Communication ($r = .42, 95\% CI = .10$ to .67, $p = .009$), and Parent-Child Involvement ($r = .47, 95\% CI = .20$ to .72, $p = .002$); SES was related to all teacher-reported outcomes, including social functioning ($r = .37, 95\% CI = .09$ to .58, $p = .02$), Academic Success ($r = .40, 95\% CI = .11$ to .63, $p = .01$), and Academic Productivity ($r = .30, 95\% CI = .01$ to .54, $p = .07$); and medication status was related to Parent-Child Involvement ($r = -.31, 95\% CI = -.01$ to -.60, $p = .06$). These variables were therefore included as covariates in the models predicting outcomes with which they were correlated; all others reflect simple model results with no covariates.

**Tier 1: Functional Heterogeneity Subgroup Classification**

**Descriptive Statistics**

Following Sarver et al. (2015), each child was classified as Impaired or Not Impaired on each functional outcome using the Jacobson and Truax (1991) model of reliable change. As shown in Figure 1, the current sample displayed substantial heterogeneity in each functional outcome. Specifically, 98% ($n = 43$) of the sample displayed impairment in at least one measured domain, and 65% were impaired in two (41%) or all three domains (24%). The 2% characterized as Not Impaired reflect one participant who fell just below the criterion for academic impairment ($z = 1.80, p = .07$). Within functional domains, the proportion of ADHD children classified as Impaired was 70% for teacher-reported academic functioning, 62% for parent-reported family functioning, and 55% for teacher- or parent-reported social functioning.

Collectively, these descriptive analyses confirmed significant functional heterogeneity in the current sample that was similar to previous studies in terms of the proportion of children with ADHD classified as impaired in each domain (de Boo & Prins, 2007; Mayes & Calhoun, 2006). Of primary interest was the extent to which children with impairments in each functional domain demonstrated an identifiable neurocognitive profile. We therefore compared children defined as Impaired vs. Not Impaired on each functional outcome, and interpreted the bias-corrected, bootstrapped 95% CIs of the Cohen’s $d$ effect size for each between-group comparison as described above.

**Academic Functioning (Teacher Report)**

Comparisons of Impaired vs. Not Impaired children revealed that children defined by impairments in academic success exhibited large magnitude PHWM deficits ($d = 0.86, 95\% CI = 0.28$ to 1.44, $p = .007$). Similarly, children defined based on impairments in academic productivity demonstrated medium magnitude impairments in PHWM ($d = 0.54, 95\% CI = -0.02$ to 1.10, $p = .08$); however, the possibility of no effect for this comparison remained due to the narrow inclusion of 0.0 in the CI. We observed also a medium magnitude effect on global IQ for children with deficits in academic success ($d = 0.57,$
95% CI = −0.12 to 1.21, \( p = .09 \); however, the 95% CI included 0.0, suggesting the possibility of no effect. Effect sizes were small to minimal for all other comparisons (all \( d \leq 0.30 \), all 95% CIs centered around 0.0).

**Social Functioning (Parent and Teacher Report)**

Children defined as socially impaired based on parent report demonstrated slower processing speed (\( d = 0.53, 95\% \text{ CI} = 0.07 \) to 1.25, \( p = .04 \)). In addition, children with parent-reported social impairment demonstrated medium magnitude impairments in PHWM (\( d = 0.53, 95\% \text{ CI} = -0.17 \) to 1.22, \( p = .12 \)); however, the possibility of no effect for this comparison remained due to the narrow inclusion of 0.0 in the CI. Children with teacher-defined social impairments showed small to medium deficits in inhibitory control (\( d = 0.44, 95\% \text{ CI} = -0.28 \) to 1.09, \( p = .20 \)) and global IQ (\( d = 0.44, 95\% \text{ CI} = -0.18 \) to 1.06, \( p = .19 \)) with 95% CIs that leave open the possibility of no effect. Effect sizes were small to minimal for all other comparisons (all \( d \leq 0.21 \), all 95% CIs centered around 0.0).

**Family Functioning (Parent Report)**

Children defined based on impairments in parent–child attachment demonstrated large magnitude deficits in VSWM (\( d = 1.09, 95\% \text{ CI} = 0.47 \) to 1.66, \( p = .001 \)). A medium
magnitude effect size was also noted for PHWM ($d = 0.45$, 95% CI $= 0.52$ to 1.49, $p = .34$); however, the 95% CI substantially overlapped 0.0, suggesting a high likelihood of no effect. Children whose parents reported significantly impaired parenting confidence demonstrated large magnitude VSWM deficits ($d = 1.05$, 95% CI $= 0.01$ to 2.07, $p = .048$) and medium magnitude deficits in PHWM ($d = 0.68$, 95% CI $= 0.04$ to 1.33, $p = .04$). Effect sizes were small to minimal for all other comparisons (all $d \leq 0.44$, all 95% CIs centered around 0.0). No neurocognitive deficits were detected for children with impaired parent–child communication (all $d \leq 0.32$, all 95% CIs centered around 0.0).

**Tier 2: Dimensional Analyses**

**Academic Functioning (Teacher Report)**
The bias-corrected, bootstrapped regression model was significant for academic success ($R^2 = .49$, $p < .005$). Better-developed VSWM (partial $R^2 = .12$, $B = 3.01$, 95% CI $= 0.76$ to 5.52, $p = .03$), PHWM (partial $R^2 = .24$, $B = 3.98$, 95% CI $= 1.89$ to 6.35, $p = .001$), and IQ (partial $R^2 = .26$, $B = 0.32$, 95% CI $= 0.11$ to 0.50, $p = .002$) each predicted higher academic success. Processing speed showed similar relations with academic success (partial $R^2 = .11$, $B = 2.84$, 95% CI $= −0.19$ to 5.68, $p = .07$); however, the possibility of no effect for this predictor remained due to the narrow inclusion of 0.0 in the CI. Better-developed PHWM predicted higher academic productivity (partial $R^2 = .16$, $B = 3.17$, 95% CI $= 0.89$ to 5.51, $p = .01$), but the omnibus test for academic productivity was non-significant ($R^2 = .21$, $p = .14$). Inhibition did not predict either academic outcome (both $p > .15$, both 95% CIs substantially overlap 0.0, both $R^2 < .05$).

**Social Functioning (Parent and Teacher Report)**
The neurocognitive variables significantly predicted teacher-reported social functioning ($R^2 = .43$, $p = .005$), such that better-developed VSWM (partial $R^2 = .12$, $B = 4.02$, 95% CI $= 1.05$ to 7.00, $p = .03$) and processing speed (partial $R^2 = .16$, $B = 4.53$, 95% CI $= 1.18$ to 8.34, $p = .03$) each predicted better social functioning. Higher SES predicted better teacher-reported social functioning (partial $R^2 = .26$, $B = 0.56$, 95% CI $= 0.23$ to 0.91, $p = .002$); inhibition, PHWM, and IQ did not predict teacher-reported social functioning (all 95% CIs centered around 0.0, all $R^2 < .04$, all $p > .29$). In contrast, only PHWM predicts parent-reported social functioning (partial $R^2 = .11$, $B = 3.07$, 95% CI $= 0.17$ to 6.42, $p = .06$), but the omnibus test for parent-reported social functioning was non-significant ($R^2 = .12$, $p = .44$).

**Family Functioning (Parent Report)**
The omnibus tests were significant for parenting confidence ($R^2 = .39$, $p = .007$), parent-child attachment ($R^2 = .34$, $p = .04$), and parental involvement ($R^2 = .46$, $p = .008$). Better-developed inhibitory control predicted greater parent-reported attachment (partial $R^2 = .13$, $B = 3.02$, 95% CI $= 0.74$ to 5.62, $p = .03$) and parenting confidence (partial $R^2 = .26$, $B = 4.24$, 95% CI $= 1.89$ to 6.18, $p < .0005$). In addition, better-developed PHWM predicted greater parenting confidence (partial $R^2 = .20$, $B = 3.07$, 95% CI $= 0.37$ to 5.38, $p = .02$). Older age predicted more difficulties with parent-child attachment (partial $R^2 = .15$, $B = 1.89$, 95% CI $= 0.34$ to 3.22, $p = .03$) and parental...
involvement (partial $R^2 = .23$, $B = 2.50$, 95% CI = 0.45 to 4.21, $p = .01$); psychostimulant medication predicted lower reported parent-child involvement (partial $R^2 = .21$, $B = 6.70$, 95% CI = 0.81 to 12.72, $p = .05$). The omnibus test for parent-child communication was non-significant ($R^2 = .25$, $p = .18$). Processing speed and VSWM failed to predict any family outcomes (all $p > .12$, all 95% CIs substantially overlap 0.0, all $R^2 < .07$).

**Discussion**

The current study was the first to examine neurocognitive predictors of heterogeneity in each of the three primary areas of functional impairment associated with ADHD (Pelham et al., 2005). Overall, results add to our understanding of individual differences in neurocognitive abilities among children with ADHD, and reveal that this variation appears to play an important role in peer, family, and academic functioning. Specifically, working memory abilities were associated with ADHD-related heterogeneity in all three functional domains, processing speed predicted teacher-reported social functioning, and inhibitory control predicted caregiver perceptions of family functioning. These findings were generally consistent with the developmental literature (Holmes et al., 2016) and previous comparisons of ADHD and typically developing groups (Chhabildas et al., 2001; Rapport et al., 2013; Rucklidge & Tannock, 2002), and extend previous findings by demonstrating that specific neurocognitive abilities are important for understanding heterogeneity in functional impairments among children with ADHD.

Among the neurocognitive predictors, working memory abilities accounted for significant individual differences across several functional indicators, and were the only assessed neurocognitive abilities to predict outcomes across both informants and all three areas of functioning. This pattern implicates working memory dysfunction as a liability for broad-based functional impairment, and is consistent with previous studies linking individual differences in working memory components with academic attainment among children with ADHD (Alloway & Stein, 2014), as well as studies identifying cross-sectional (Mayes & Calhoun, 2007; Rogers et al., 2011) and longitudinal associations (Miller et al., 2012; Sarver et al., 2012) between working memory storage/rehearsal subcomponents and individual differences in specific academic domains. The current study extends these findings, and suggests that working memory may also be important for understanding heterogeneity in family and social functioning among children with ADHD. That is, children with ADHD who are better able to mentally store and process information are perceived by teachers and parents as more socially adept and more effectively parented. Conversely, underdeveloped working memory likely makes it extraordinarily difficult to engage in the give-and-take, listen-and-wait behaviors required for adept social interactions (Kofler et al., 2011). This explanation is consistent also with the observation that a majority of DSM-5 hyperactivity/impulsivity items refer to intrusive verbal behavior and the inability to maintain thoughts and forestall action (e.g., interrupting conversations, blurring out).

The current results were highly consistent with previous studies demonstrating strong continuity between working memory and social problems in ADHD (Bunford et al., 2015; Kofler et al., 2011), and provide new data suggesting that children’s working
memory abilities may influence perceptions regarding relationship quality and their ability to effectively parent their child with ADHD. Combined with the finding that better-developed inhibitory control predicts improved family functioning, these results are generally consistent with developmental models suggesting that child cognitive/intellectual assets may facilitate positive interactions with caring adults (Lerner, Phelps, Forman, & Biowers, 2009), which in turn may shape early executive function development (Cuevas et al., 2014) and buffer against adverse outcomes for at-risk children (Eccles & Gootman, 2002). The current study extends these findings by identifying specific neurocognitive abilities that influence parent–child interactions for children with ADHD. Alternatively, the distinct neurocognitive profiles associated with family vs. social (peer) impairments may suggest that the abilities and behaviors required for successful parent–child interactions differ somewhat from those required for successful peer interactions. That is, parents may have expectations for their children that require better developed working memory and inhibitory control (e.g., following multistep directions, inhibiting unwanted behaviors), whereas successful interactions with same-aged peers may rely to a greater extent on the rapid processing of social information (Phillips, Tunstall, & Channon, 2007). This hypothesis is consistent with the current finding that somewhat more children were classified as impaired in family functioning (62%) relative to social functioning (55%), as well as meta-analytic findings that working memory deficits may be more prevalent and/or of larger magnitude than processing speed deficits (Kasper et al., 2012; Kofler et al., 2013).

Inhibitory control was uniquely associated with family functioning. Its failure to predict academic functioning was surprising given our use of a psychometrically-supported inhibition task (Alderson et al., 2007; Snyder et al., 2015), as well as previous developmental studies suggesting a small but significant role for inhibitory control in academic functioning (St. Clair-Thompson & Gathercole, 2006; Thorell, 2007; Wåhlstedt et al., 2009). In contrast, the current findings were consistent with previous ADHD studies that failed to find links between inhibition and ADHD symptoms (Alderson et al., 2010; Rucklidge & Tannock, 2002), as well as meta-analytic conclusions that inhibitory control may be intact in ADHD (Alderson et al., 2007; Lijffijt et al., 2005). Interestingly, the zeitgeist regarding inhibition appears to be shifting in both the clinical and cognitive literatures. Whereas inhibitory control was once considered a core executive function (Miyake et al., 2000) with promise for offering a unifying explanation of ADHD (Barkley, 1997), it may now be considered a “dead end” in ADHD (Rommelse et al., 2007) and is no longer considered a core executive function in at least one influential model of human cognition (Miyake & Friedman, 2012). Notably, however, inhibitory control showed strong continuity with parental confidence and parent–child attachment in the current study, suggesting that it remains an important factor in understanding ADHD-related impairments even if inhibition deficits are not present at the group level (Alderson et al., 2007; Lijffijt et al., 2005).

**Limitations**

The unique contribution of the current study was its systematic examination of neurocognitive predictors of functional heterogeneity in a well-defined sample of children with ADHD. Several caveats merit consideration despite methodological refinement including
our approach to isolating reliable variance associated with each neurocognitive function and examination of multiple impairment domains. Generalization of findings from highly controlled laboratory experiments are always limited to some extent, and no conclusions regarding neurocognitive deficits can be drawn due to the lack of a typically developing comparison group. However, ADHD-related impairments in neurocognitive abilities are well documented (Kasper et al., 2012), and impairments in each functional outcome (Pelham et al., 2005) were quantified objectively using norm-referenced, psychometrically-sound tests. In addition, significant predictors of each functional outcome were detected, suggesting adequate power and supporting our a priori effect estimation. However, the significant unexplained variance in each outcome indicates a clear need for future research that includes larger samples as well as typically developing and clinical comparison groups to determine the extent to which the mechanisms associated with peer, family, and academic functioning differ across clinical and nonclinical populations.

In addition, several of the children with ADHD met criteria for comorbid behavioral and mood disorders; thus, the extent to which the findings generalize to children with “pure” ADHD is unknown. The inclusion of these common comorbidities, however, is expected to improve generalizability given that the sample is more representative of the larger population of children with ADHD (for which the majority have at least one comorbid diagnosis; Wilens et al., 2002). Fifty percent of our ADHD sample was prescribed stimulant medication, which was broadly consistent with epidemiological estimates (39% to 69%; Froelich et al., 2007; Visser et al., 2014). Although medication status was generally unrelated to our study variables, it may have dampened effect size estimates when juxtaposing neurocognitive performance off medication with parent and teacher perceptions that may be influenced by medication. The mean IQ of our sample was higher than the national average by approximately 1/3 to 2/3 SD; thus, the extent to which the findings generalize to children with average or lower intellectual abilities remains unknown. Finally, future research may benefit from examining the influence of informant source on impairment estimates (Valo & Tannock, 2010), as well as impairment indicators beyond those represented herein (e.g., health impairment, quality of life, sociometric standing) to further specify the mechanisms and processes underlying these impairments and identify mechanistic subtypes (Fair et al., 2012).

**Clinical and Research Implications**

Collectively, results of the current study suggest that neurocognitive processes are particularly important for understanding heterogeneity in daily functioning among children with ADHD. In particular, children with impairments in academic and family functioning showed large magnitude working memory deficits, whereas children with social impairments demonstrated slowed processing speed. If replicated, these findings suggest differential assessment and intervention targets depending on each child’s functional impairment profile. That is, in addition to direct remediation of each identified functional area, improved efficacy may be realized by adding interventions that target the specific mechanisms associated with the child’s identified functional impairment(s) (Chacko et al., 2014). For example, children with academic impairments may be likely to benefit from interventions that facilitate academic success and productivity (e.g., class-wide peer tutoring; DuPaul &
Weyandt, 2006) while concurrently targeting their underdeveloped working memory abilities. Similarly, we hypothesize that processing speed training may augment interventions that facilitate prosocial engagement (Mikami, Lerner, Griggs, McGrath, & Calhoun, 2010), and family-based interventions may see incremental benefits when combined with working memory and/or inhibitory control training. Unfortunately, extant medications and “working memory” training programs generally fail to improve working memory (Melby-Lervåg & Hulme, 2016; Rapport et al., 2013; Rubia et al., 2014; Shipstead, Hicks, & Engle, 2012), suggesting that this combined approach will have to wait until next-generation neurocognitive training programs and/or medications have been developed and shown to effectively improve the specific neurocognitive abilities they claim to target (Chacko et al., 2014). Nonetheless, the current results add to a growing literature implicating neurocognitive abilities not only in explaining behavioral differences between ADHD and non-ADHD groups, but also in the substantial heterogeneity in functional outcomes associated with the disorder.

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