Developmental Electrophysiology Laboratory Yale Child Study Center

Background

- Psychiatric co-morbidities in autism spectrum disorder (ASD) are
- common, occurring in as many as 72% of children with ASD¹. • Children with ASD often have deficits in aspects of cognitive efficiency, specifically working memory, processing speed, and/or executive functioning, though findings are mixed².
- Psychiatric comorbidity has negative consequences on cognitive functioning, including executive functioning.
- Understanding whether secondary conditions or comorbidities have unique or shared impact on ASD compared to other diagnostic groups has important implications for assessment and treatment.

Objective: To study the effect of comorbidity on cognitive efficiency in individuals with ASD compared to individuals with other childhood psychiatric conditions.

Methods

Procedures:

- Use of archival clinic database from an academic medical center clinic specializing in developmental disability evaluations.
- Cognitive assessments were conducted by licensed psychologists, and final consensus diagnosis was determined by a multidisciplinary team consisting of psychology, psychiatry, and speech/language specialists following comprehensive evaluation.

Measures:

- Wechsler Intelligence Scale for Children, 3rd & 4th Edition (WISC-III, WISC-IV)
- Wechsler Adult Intelligence Scale, 3rd Edition (WAIS-III)
- Behavior Rating Inventory of Executive Function (BRIEF)

Inclusion/Exclusion Criteria:

- Primary diagnosis of ASD (ASD) or other psychiatric disorder (Non-ASD).
- Excluded: Children diagnosed with a primary or comorbid Intellectual Disability, cerebral palsy, or children with no clinical diagnosis.

Participant Demographics:							
	N		Sex (M, F)	Age (SD)			
ASD	307		225, 46	9.76 (3.75)			
Non-ASD	108	79, 14 10.90 (3.84))			
Non-ASD Group Diagnoses		n			n		
ADHD		17	Obsessive Compulsive Disorder		1		
Anxiety Disorders		17	Oppositional Defiant Disorder		1		
Conduct Disorder		2	Reactive Attachment Disorder		3		
Global Developmental Delay		5	Childhood Schizophrenia		3		
Major Depressive Disorder		6	Tourette's Syndrome		3		
Mood Disorders		5	Other		3		
Language Disorders		14					

• There was a significant difference in age between the ASD and Non-ASD group, *t*(399)=-2.66, *p*<.01.

• There was no significant difference in sex $[\chi^2(3,415)=3.77, p=.29]$.

• There was no significant difference in race $[\chi^2(6,415)=3.61, p=.73]$. ASD Group: Caucasian: 37%; Asian: 1%; African American: 1%; Indian: <1%; and Multi-racial: 2%. Non-ASD Group: Caucasian: 36%; Asian: 4%; African American: 2%; Indian: n/a; and Multi-racial: 3%.

The effect of comorbidity on cognitive efficiency in autism spectrum disorder

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Methods, cont.

Statistical Analyses:

Comparison of group differences were analyzed using factorial ANCOVAs with the independent variables of either diagnostic group (ASD and non-ASD) and/or comorbidity group (presence or absence of a comorbid or secondary diagnosis) controlling for age.

Results

Diagnostic Group Comparison

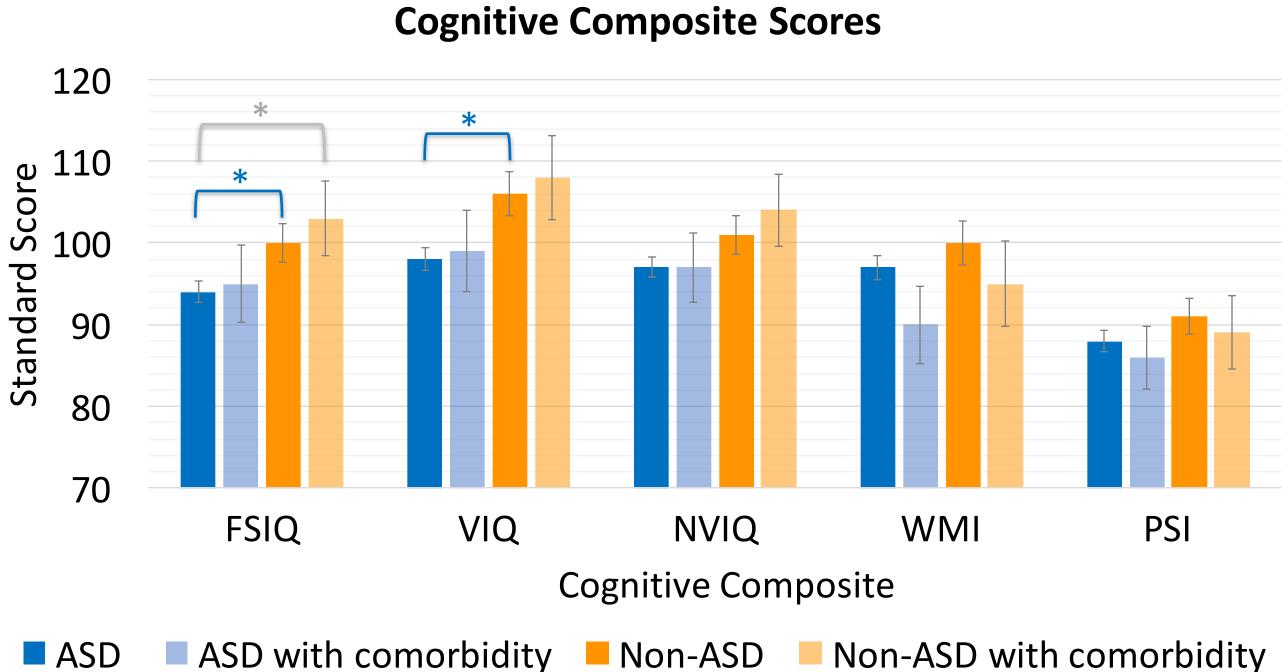
- The ASD group had significantly lower Full Scale IQ (FSIQ) than the Non-ASD group [*F*(1,381)=8.50, *p*<.01].
- Of the Verbal, Nonverbal, Working Memory and Processing Speed composite scores, there was only a significant difference in Verbal Standard Scores between the ASD and Non-ASD group [F(1,340)=9.59,*p*<.01].

Comorbid or Secondary Diagnosis Group Comparisons

• Comorbid conditions were diagnosed in 7% (*n*=22) of individuals with ASD and 20% (n=22) of individuals in the non-ASD group ($\chi^2=14.70$, p<.01).

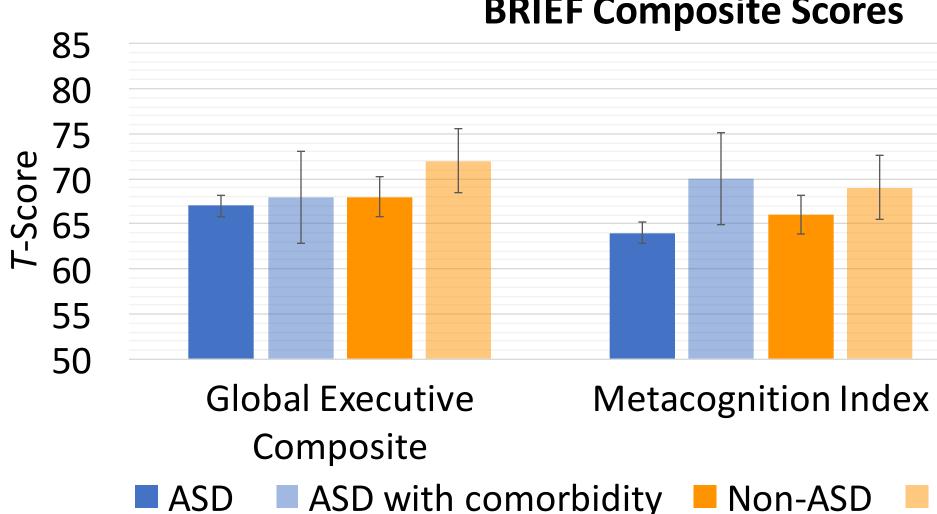
Comorbid or Secondary Diagnosis	ASD [<i>n</i> (%)]	Non-ASD [<i>n</i> (%)]
ADHD	5 (2%)	4 (4%)
Learning Disorder	9 (3%)	6 (6%)
Tourette's Syndrome	4 (1%)	1 (<1%)
Childhood Schizophrenia	1 (<1%)	0
Other developmental conditions	2 (<1%)	0
Mood or Anxiety Disorder	0	11 (10%)

- **FSIQ:** There was no significant interaction between diagnostic group and comorbidity status on FSIQ [F(1,379)=0.20, p=.66]. There was a main effect of diagnostic group [F(1, 379)=5.06, p=.03].
- Verbal: There was no main effect of comorbidity [F(1,338)=0.13, p=.72], but there was a main effect of diagnostic group [F(1,338)=5.04, p=.03] on Verbal Standard Scores. There was no significant interaction between comorbidity and diagnostic group [F(1,338)=0.01, p=.91].
- **Nonverbal:** There was no main effect of comorbidity status [*F*(1,338)=0.34, *p*=.56] or diagnostic group [*F*(1,338)=2.66, *p*=.10].
- Working Memory and Processing Speed: There was no main effect of comorbidity status or diagnostic group on WMI [(F(1,283)=2.30, p=.13), (F(1,283)=1.25, p=.26)] or PSI [(F(1,277)=0.48, p=.49), (F(1,277)=0.82, p=.26)] *p*=.37)].



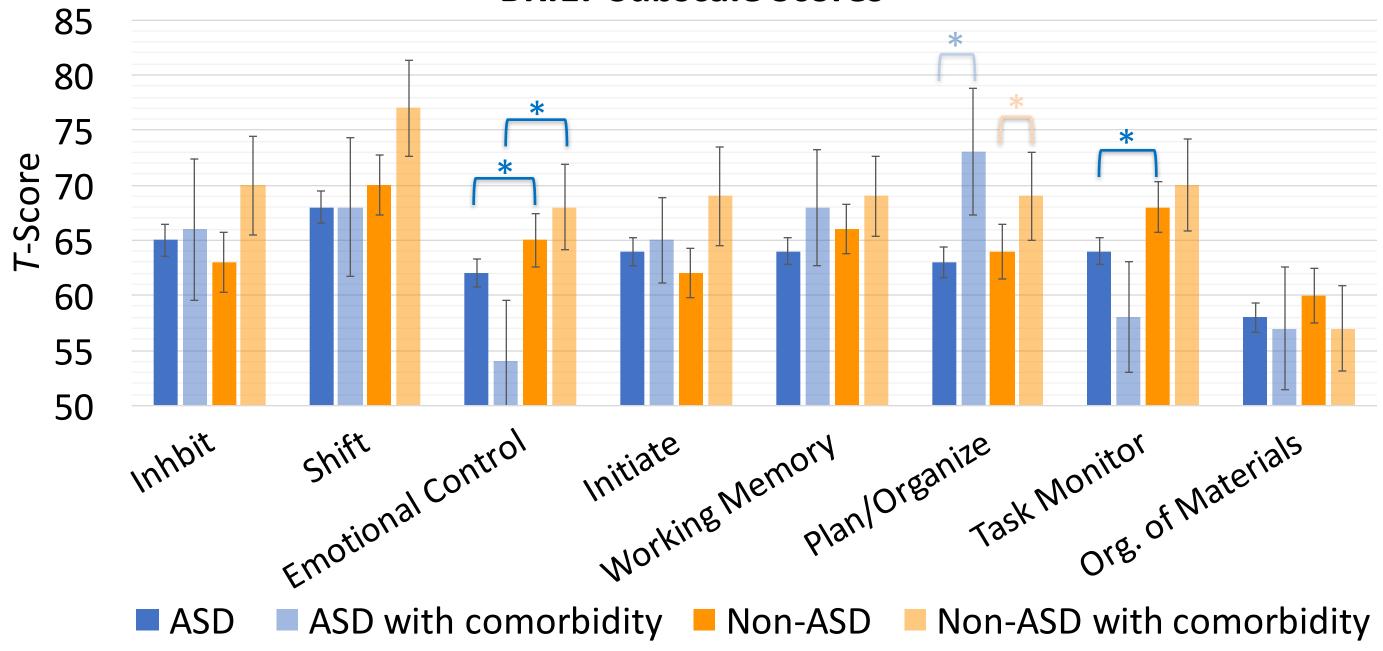
ASD

Executive Functioning (EF):



- 96)=5.27, *p*=.02] than the ASD group.
- $(M^{a}=54, SE=5.59)$ groups (p=.05).





1. Leyfer, O. T., Folstein, S. E., Bacalman, S., Davis, N. O., Dinh, E., Morgan, J., & Lainhart, J. E. (2006) Comorbid psychiatric disorders in children with autism: interview development and rates of disorders. Journal of Autism and Developmental Disorders, 36, 849–861. 2. Hill, E.L. (2004). Executive dysfunction in autism. *Trends Cogn Sci*, 8(1):26–32. doi:10.1016/j.tics.2003.11.003

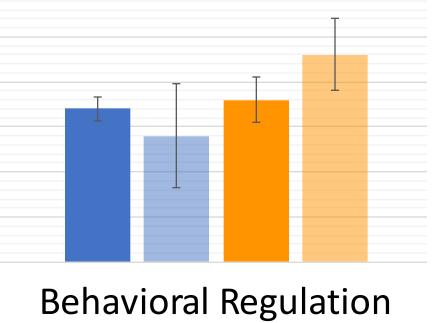
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Results, cont.

• There were no significant differences in overall executive functioning (GEC), Behavioral Regulation Index (BRI), or Metacognitive Index (MI). **BRIEF Composite Scores**



ASD with comorbidity Non-ASD with comorbidity • Children in the Non-ASD group had significantly greater deficits in Emotional Control [F(1,105)=4.68, p=.03] and Task Monitoring [F(1, 1)]

• Post hoc analyses revealed the same effect for Emotional Control between the Non-ASD with comorbidity (*M^a*=68, *SE*=3.9) and ASD with comorbidity

• Children with a comorbid condition in both diagnostic groups had

significantly greater deficits in Planning/Organizing [F(1,104)=4.19, p=.04]; there was no significant interaction [F(1,104)=0.39, p=.54].

Discussion

Comorbidity did not differentially impact core cognitive abilities or cognitive efficiency in children with ASD or another psychiatric condition. • Comorbidity increased impairment in one domain of EF in both ASD and Non-ASD groups, specifically planning and organizing work or activities. Analysis with a larger sample with comorbidity is needed.

• A limitation was the small sub-samples on the BRIEF, as well as within each comorbid condition that did not allow for more nuanced analysis of specific disorder(s) differential impact on cognitive processes.

References

