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Birth Weight and Cognition in Children with Epilepsy

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Summary

Objective—Birth weight is an important indicator of prenatal environment and subtle variations of birth weight within the normal range have been associated with differential risk for cognitive and behavioral problems. Therefore, we aimed to determine if there are differences in birth weight between full term children with uncomplicated new/recent-onset epilepsies and typically-developing healthy controls. We further examined the relationships between birth weight and childhood/adolescent cognition, behavior, and academic achievement.

Methods—108 children with new/recent-onset epilepsy and 70 healthy controls underwent neuropsychological assessment. All participants were born full-term (>37 weeks) without birth complications. Parents were interviewed regarding their child's gestation, birth and neurodevelopmental history.

Results—Birth weight of children with epilepsy was significantly lower than healthy controls ($p=0.023$). Whereas birth weight (covaried with age, sex, handedness, and mother's education) was significantly associated with cognition in controls in multiple domains (intelligence, language, aspects of academic achievement), this relationship was absent in children with epilepsy. Birth weight was not associated with clinical epilepsy variables (age of onset, epilepsy syndrome) and was not predictive of a variety of other academic or psychiatric comorbidities of epilepsy.

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We confirm that we have read the Journal's position on issues involved in ethical publication and affirm that this report is consistent with those guidelines.

Significance—Although the origin of lower birth weight in children with epilepsy is unknown, these findings raise the possibility that abnormal prenatal environment may impact childhood-onset epilepsy. Furthermore, the positive relationship between birth weight and cognition evident in healthy controls was disrupted in children with epilepsy. However, birth weight was not related to academic and psychiatric comorbidities of childhood epilepsy.

Keywords

Birth weight; cognition; academic achievement; new-onset epilepsy; localization-related epilepsy; idiopathic generalized epilepsy

Introduction

There is now substantial evidence indicating that neurobehavioral comorbidities of childhood-onset epilepsy can be evident at or prior to the time of epilepsy onset,¹⁻⁶ but the neurodevelopmental origins of these cognitive and behavioral complications are unknown. Human and non-human animal studies have shown that variance in cognitive and behavioral maturation can be traced to fetal development.^{7; 8} Birth weight has been established as a sensitive indicator of prenatal environment and even subtle differences within the normal range of birth weight have been associated with differential risks for cognitive problems and psychopathology in later life.^{9; 10} Specifically, in children with normal birth weight for gestational age, lower birth weight was significantly associated with poorer cognitive function in later life, with predictive significance beyond that predicted by socio-economic status.¹⁰ More recently, twin studies have shown modest but significant effects of birth weight on later cognitive ability in full term children.^{11; 12}

Extremely low birth weight (less than 1000g), premature birth, and concomitant pre- and perinatal complications have been associated with a variety of neurological disorders, including autism¹³ and cerebral palsy¹⁴ as well as poor neurodevelopmental outcomes broadly defined.^{15; 16} Abnormally low birth weight for gestational age and concomitant pre- and peri-natal complications has also been found in individuals with epilepsy.^{17; 18} Therefore, it would be expected that these children with complicated births and severe epilepsies would have poor cognitive outcomes. However, an important but unanswered question is whether children born full term with uncomplicated epilepsies will have perturbed prenatal environment as indexed by lower birth weight but within the normal range for gestational age. Given that a majority of children with epilepsy have uncomplicated births, close examination of this issue may have public health implications by raising the possibility of very early neurodevelopmental origin of childhood-onset epilepsies. Therefore, our study aimed to compare birth weight in children with uncomplicated epilepsies, normal gestation, delivery, and normal birth weight for gestational age to birth weight of healthy controls. Further, neurobehavioral comorbidities of epilepsy including abnormalities in academics, cognition, emotion, and social function, are frequently observed in children with epilepsy.^{6; 19-22} Thus, we investigated the relationship between variations in birth weight and risk for subsequent cognition, social, and psychiatric complications in the children with epilepsy.

Methods

Participants

The initial sample comprised 203 participants aged 8-18 including 124 children with new/recent-onset epilepsy (CWE) and 79 typically-developing healthy controls (HC). All CWE were followed for active epilepsy by a neurologist. Inclusion criteria were a diagnosis of epilepsy within the past 12 months, no other developmental disabilities or neurological disorders, normal neurological examinations and normal neuroimaging results. A board-certified pediatric neurologist (blinded to neuropsychological and interview data) confirmed that each patient had an idiopathic epilepsy diagnosis and provided independent confirmation of syndrome diagnosis. HC participants were age- and gender-matched first-degree cousins of epilepsy participants. HC participants presented with no history of seizures, early initial precipitating injuries (e.g., febrile convulsions), other developmental or neurological disease, or loss of consciousness greater than 5 minutes.

Given that the focus of the current study is on normal birth weight and full-term children, further we applied the following exclusion criteria determined through clinical interview for both healthy controls and completion of an abbreviated version of the Yale Neuropsychological Assessment Scales²³ and review of medical records included children with epilepsy: 1) preterm birth (<37 weeks); 2) low birth weight (at or below 10th percentile for normal gestation period, assessed separately for males and females following published birth weight norms²⁴); or significant pre-/peri-/postnatal difficulties (e.g., hypoxia/ischemia, cerebral hemorrhage, need for supplemental oxygen or intubation, use of incubator). Finally, one control child was excluded based on a full-scale IQ < 70. In Data for the exclusion criteria were obtained through clinical interview and completion of an abbreviated version of the Yale Neuropsychological Assessment Scales²³ and review of medical records. Exclusion rates were not significantly different between groups ($p=0.749$): in the epilepsy group, 16/124 (12.9%) of children in the initial sample were excluded, while 9/79 (11.4%) of children in the control group were excluded. Thus our final normal birth weight (NBW) sample consisted of 108 CWE and 70 typically-developing HC. Demographic and clinical epilepsy characteristics of the sample are given in Table 1.

Standard protocol approvals, registrations, and patient consents

Research approval was obtained from the Health Sciences Institutional Review Board at the University of Wisconsin Medical School. Written informed consent was obtained from legal guardians of participating children and adolescents, written informed consent was obtained from participants over age 18, and written informed assent was obtained from participants age 8-17. All procedures were consistent with the Declaration of Helsinki.²⁵

Procedures

Children underwent comprehensive neuropsychological testing. Parents underwent a clinical interview and completed questionnaires including the abbreviated Yale Neuropsychological Assessment Scales²³ to characterize gestation, delivery, neurodevelopment, and seizure. All pertinent medical records were obtained after parents signed a release of information. Parents were questioned through a structured interview about their child's

school progress, including any academic problems and any special educational services provided in order to address those problems. This interview was conducted blind to cognitive and behavioral results. Past and current psychiatric status was determined via a semi-structured interview, the *Kiddie-Schedule for Affective Disorders and Schizophrenia – Present and Lifetime Version*.²⁶ Rates of anxiety, depression, and attention deficit hyperactivity disorder (ADHD) were assessed within each group (HC, CWE). Finally, each participating parent completed the Child Behavior Checklist for children aged 6-18 (CBCL/6-18) from the Achenbach System of Empirically Based Assessment.²⁷

Neuropsychological assessment and analysis

All participants were administered a comprehensive test battery that included measures of intelligence, academic achievement, language, verbal memory, executive function, and speeded fine motor dexterity. Table 2 gives a complete list of tests by domain. Independent samples t-tests were used to compare CWE and HC groups on birth weight as well as demographic and cognitive variables. Partial correlations were used to examine the relations between birth weight and raw neuropsychological test scores, using age, gender, handedness, and mother's educational level (completion of 4- year college degree: yes/no) as covariates.

Results

Demographic and cognitive differences between groups

As shown in Table 1, at the time of testing CWE had lower full-scale IQ scores ($M = 102.92$, $SD = 12.77$) than did HC ($M = 107.67$, $SD = 11.59$), $p=0.013$, although it should be noted that the IQ of CWE was within the normal or average range. CWE were also more likely than HC to have received special services in school (41.12% versus 20.90%), $p=0.008$. Groups did not differ in age, gender, or mother's educational level. Clinical characteristics of the epilepsy group are given in Table 1. The average duration of epilepsy was 8.59 months ($SD = 3.78$), with an average age of onset of 11.45 years ($SD = 3.35$). CWE performed significantly worse than HC on 12/15 neuropsychological tests of general IQ, academic achievement, memory, language, psychomotor function, and executive function. The only tests in which performance of CWE was *not* significantly worse than performance of HC were in academic achievement (reading and spelling) and delayed verbal memory. Means and standard deviations of all test scores are presented by group in Table 2.

Birth weight

CWE had significantly lower birth weights ($M = 3440.46$ grams, $SD = 486.25$) than HC ($M = 3620.84$, $SD = 550.89$), $p=0.023$. The distributions of birth weight for both CWE and HC are shown in Figure 1. Kurtosis statistics were normal for each group (CWE: .561, $SE = .461$; HC: -.002, $SE = .566$). Likewise, the HC group skewness statistics was in the normal range (.232, $SE = .287$). However, the CWE group distribution was somewhat positively skewed (.483, $SE = .233$). Note that lower birth weight in CWE compared to HC reported here excluded individuals who were born pre-term, significantly underweight, or with severe pre-/perinatal complications. In both groups, birth weight was unassociated with current

weight or head circumference. Additionally, current weight at time of testing was not different between groups.

Birth weight and cognition

Partial correlations were computed to assess relations within each group (CWE, HC) between birth weight and raw cognition scores, controlling for age, gender, handedness, and mother's level of education. For 7/15 tests, birth weight in HC was significantly positively correlated with better performance (r 's ranging from 0.26 – 0.40). Notably, these cognitive correlates of birth weight were most evident in domains of general intelligence (full-scale IQ, verbal IQ, performance IQ) and language abilities (confrontation naming, expressive naming, and receptive language), with the exception of an arithmetic test of academic achievement. Other cognitive domains including executive function, memory, motor function, and remaining academic achievement (reading, spelling) were not associated with birth weight in HC, although trend level positive correlations (p 's between 0.05 and 1.00) were found for speeded fine motor dexterity (Grooved Pegboard—dominant hand) and problem-solving skills (D-KEFS correct sorts); see Table 2. Conversely, CWE showed no significant relations between birth weight and test scores from any functional domain (r 's ranging from -0.16 – 0.11). The differential relationships with birth weight and cognition between CWE and HC, are exemplified in Figure 2 and 3 with side-by-side group comparisons of these partial correlations for full-scale IQ and expressive naming, respectively.

Birth weight and common comorbidities of epilepsy

In CWE, birth weight was not associated with presence of academic difficulties including ADHD, receiving special services in school, having a specific educational plan in place, and presence of learning problems before diagnosis of epilepsy (all p 's > 0.60). Birth weight was also unassociated with presence of anxiety and depression, both at the time of testing and prior to epilepsy diagnosis (p 's > 0.70). Furthermore, in children with epilepsy birth weight was not associated with duration of epilepsy, age at diagnosis, epilepsy syndrome (IGE/LRE), or current number of anti-epilepsy medications (p 's > 0.16). AED's, all p 's > 0.16. To further examine the possible effects of AED's on birth weight and cognition, we then reran our correlations between birth weight and cognitive variables separately for those children with epilepsy on 0 AED's versus those on 1 or more AED's. We found that these correlations remained non-significant across groups with and without AED exposure, all p 's > 0.30. Finally, birth weight was not associated with CBCL scales of Total Competence, Total Problems, Internalizing, or Externalizing in either group (p 's > 0.60).

Discussion

Three primary findings were evident from this investigation. First, children with uncomplicated idiopathic localization-related and generalized epilepsies exhibited significantly lower birth weight compared to healthy controls. It is important to note that this comparison was made using only those participants who were of normal birth weight, who were not born preterm, and whose mothers did not have pregnancy complications. Second, increasing birth weight was related to significantly better cognitive abilities across multiple

functional domains later in childhood/adolescence in healthy controls, while this relationship was completely absent in children with epilepsy. Third, contrary to expectations, birth weight in children with epilepsy was predictive neither of common problematic comorbidities including academic difficulties and psychiatric complications, nor of clinical epilepsy features such as duration of epilepsy, age of disease onset, and broad epilepsy syndrome. Each of these points is reviewed in turn below.

Birth weight in children with epilepsy

This is among the first demonstrations that birth weight is significantly lower in children with epilepsy compared to healthy controls. Importantly, the distribution of birth weight was normal in both groups indicating that the effect was not driven by outliers (see Figure 1). Furthermore, excluded from both the epilepsy and control groups were children who were identified as extremely low birth weight, preterm deliveries, or documented pre-/peri-natal complications, rates of which were comparable in the epilepsy (12.9%) and control (11.4%) groups. In children born full term, the average birth weights for both groups fell in the normal range, but were significantly lower in the children with epilepsy regardless of whether they were followed for localization-related or idiopathic generalized syndromes.

Although the origin of lower birth weight in CWE is unknown, these findings raise the possibility that abnormal prenatal environment may impact childhood-onset epilepsy. The brain undergoes rapid growth during in utero and the dynamic of neurodevelopment significantly influences eventual birth weight. For example, Graca and colleagues examined cerebral volumes during the first post-natal week in 128 infants and found brain volumes are highly correlated with birth weight.²⁸ Further, fetal growth as documented by ultrasound during the first²⁹ and second³⁰ trimesters has been shown to influence post-natal birth weight. Given the important association between prenatal brain development and postnatal birth weight, our findings of modest birth weight reduction in CWE may provide the earliest indicator for the origin of abnormal neurodevelopment. Future studies should focus on factors that contribute to fetal growth and birth weight in childhood onset epilepsies, which may provide targets for early intervention.

Birth weight and cognition

Similar to findings from prior investigations of normally-developing children and adults,¹⁰⁻¹² our healthy control participants demonstrated a significant association between birth weight and neuropsychological status. Consistent with those results,¹⁰ these relationships were most evident across broad measures of intelligence (full-scale IQ, verbal IQ, performance IQ), as well as one area of academic achievement (arithmetic). We also found relations between birth weight and multiple language-dependent abilities (confrontation naming, expressive naming, receptive language). These analyses, and comparable analyses in the children with epilepsy, controlled for known important covariates including age, gender, handedness, and mother's educational level.^{10; 11; 31} In contrast, the relationship between birth weight and cognition was completely absent among the children with epilepsy, with no significant relationship between birth weight and measures of intelligence, language, academic achievement, memory, executive function, or cognitive/psychomotor speed. In summary, our study demonstrated that the presence of

childhood epilepsy, even uncomplicated idiopathic epilepsy, is associated with disruption of normal birth weight—cognition associations. These results are consistent with other findings from our group, a recent example being a significantly larger difference in IQ between parent and child IQ in CWE dyads relative to their HC counterparts.³² Taken as a whole, this pattern of findings suggests that some typical determinants of variance in cognitive maturation of normally-developing children may be weaker or absent altogether in CWE.

Birth weight and the comorbidities of childhood epilepsy

It is now widely appreciated that cognitive, academic, behavioral and psychiatric comorbidities of childhood epilepsy can be identified at or near the time of the first recognized seizure and diagnosis, often prior to initiation of drug treatment, and even prior to the recognition of seizures and the diagnosis of epilepsy.^{1; 2; 5; 6} The antecedent factors responsible for such findings largely remain to be characterized. There has been demonstration of family aggregation of some comorbidities,³³⁻³⁷ and it would be reasonable to hypothesize that factors related to pregnancy, birth and development might be pertinent as they have been shown to be in the general population.¹¹ Surprisingly, birth weight was not only unrelated to cognition as discussed above, but unrelated to other problematic comorbidities of childhood epilepsy including academic complications (e.g., learning problems, need for extra educational services, etc.) and psychiatric diagnoses of ADHD, anxiety, and mood disorders. These findings were not anticipated. While birth weight is reduced in children with epilepsy, it appears that other factors, yet to be identified, account for variance in later cognitive ability and other neurobehavioral comorbidities.

Limitations and future directions

There are limitations associated with this study. First, our sample was not population-based, although participants were recruited from medical centers covering broad portions of the state of Wisconsin. Second, our sample size precluded analysis of the relation between birth weight and cognition in specific syndromes of IGE and LRE (e.g., Juvenile Myoclonic, Absence, BECTS). We continue to recruit participants in an attempt to make such analyses possible. Third, we did not include some features of the socioeconomic status (SES) of our participating families (e.g., total family income). However, mother's education level—an important factor when assessing SES³⁸—was used as a covariate in all of partial correlation analyses. Within the healthy control group, birth weight was still significantly associated with cognition later in life even after controlling for mother's education level. Additionally, our sample of healthy controls is presumably similar in SES to participants with epilepsy, given that controls participants were first-degree cousins of participants with epilepsy. Despite these limitations, our study provides additional evidence that childhood-onset epilepsies are associated with early neurodevelopmental complications. These antecedent factors might have a negative impact on cognitive development by altering normal positive associations between birth weight and cognition. Note that the current study examined cognitive correlates of birth weight near the diagnosis of epilepsy. The cognitive impact of birth weight may accrue over time and thus inspection of longer term outcomes will be informative. Another pertinent question is whether lower birth weight in CWE is predictive of abnormal brain development. Therefore, future studies will examine relationships

between birth weight and prospective changes in gray/white matter volumes and cortical thickness.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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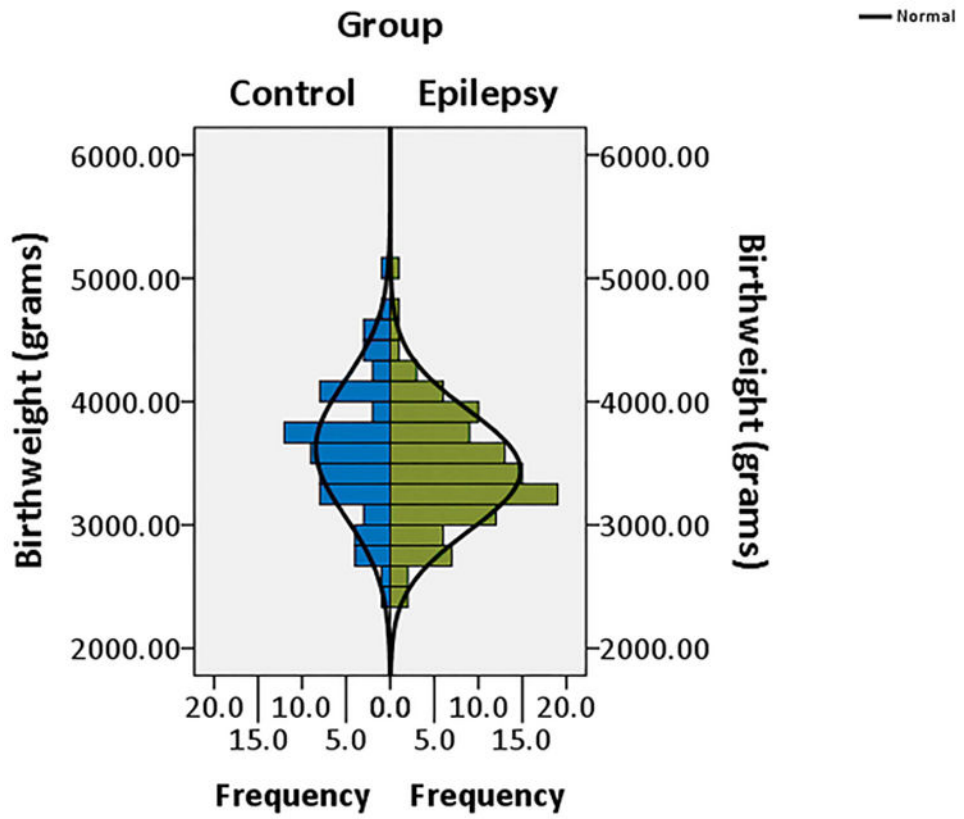


Figure 1. Distribution of birth weight by participant group (children with epilepsy; typically-developing healthy controls).

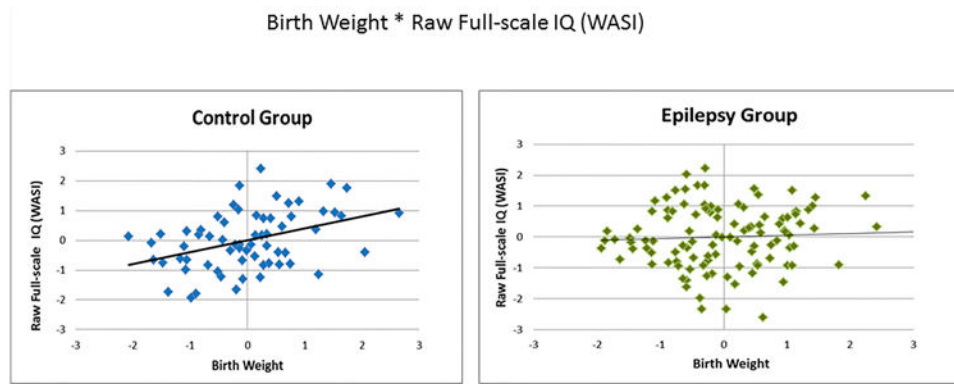


Figure 2. Birth weight by Full-scale IQ partial correlations. Both variables are presented as standardized residuals (covariates: age, gender, handedness, mother's education level.) WASI: Wechsler Abbreviated Scale of Intelligence

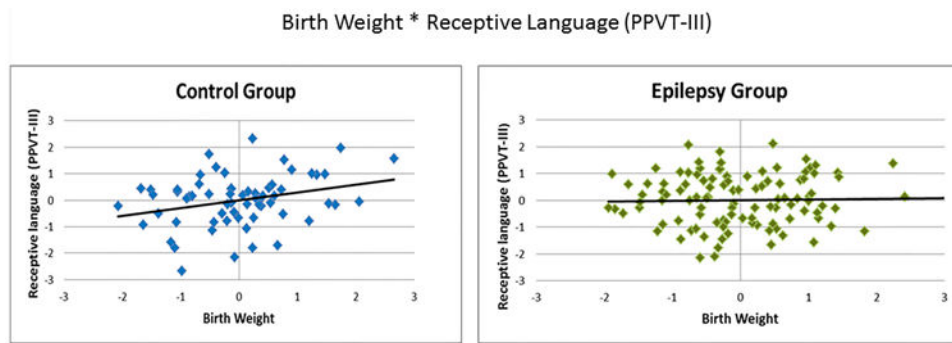


Figure 3. Birth weight by Receptive language partial correlations. Both variables are presented as standardized residuals (covariates: age, gender, handedness, mother's education level.) PPVT-III: Peabody Picture Vocabulary Test-III

Table 1

Participant characteristics by group (means and standard deviations).

Variable	Children with Epilepsy (n=108)	Healthy Controls (n=70)
Age (years)	12.33 (3.24)	12.63 (3.16)
Gender (#/% female)	49 (45.4%)	35 (50%)
Birthweight (grams) ^a	3440.46 (486.25)	3620.84 (550.89)
FSIQ ^a	102.92 (12.77)	107.67 (11.59)
Academic Problems(+/-) ^{ab}	44/63	14/53
Mother's Educational Level (4-year college graduate: yes/no) ^c	37/71	28/36
Age of seizure onset (years)	11.45 (3.35)	--
Epilepsy duration (months)	8.59 (3.78)	--
Epilepsy Syndrome (IGE/LRE) ^b	48/58	--
Number of antiepileptic drugs (0/1/2+)	21/81/6	--

^a *p* 0.05^b Information regarding academic problems was not available for 3 healthy controls and 1 child with epilepsy.^c Mother's educational level not available for 6 participants.^d 2 children with epilepsy were not classifiable as IGE or LRE but were included in all analyses.

FSIQ: Full-scale intelligence quotient

IGE: Idiopathic generalized epilepsy

LRE: Localization-related epilepsy

Table 2

Test scores and test score – birth weight partial correlations by functional domain for CWE and HC.

Domain	Ability	Test	CWE Mean (SD)	CWE Partial r BW * * Test Score ^a	HC Mean (SD)	HC Partial r BW * * Test Score ^a
Intelligence	Full scale	WASI	102.92 (12.77) ^b	0.05	107.67 (11.59)	0.40 ^c
	Verbal	WASI (verbal IQ)	103.17 (12.33) ^b	0.04	107.14 (12.91)	0.26 ^c
	Performance	WASI (performance IQ)	101.62 (13.97) ^b	0.05	106.53 (12.29)	0.39 ^c
Academic Achievement	Letter/word recognition	WRAT-3 (reading)	102.60 (12.60)	0.09	105.59 (10.79)	0.12
	Letter/word writing	WRAT-3 (spelling)	101.94 (14.19)	0.03	104.93 (12.91)	0.23
	Basic arithmetic	WRAT-3 (arithmetic)	98.14 (12.95) ^b	0.11	107.06 (11.04)	0.30 ^c
	Confrontation naming	Boston Naming Test	11.50 (2.17) ^{b,d}	0.02	12.33 (1.68)	0.27 ^c
Language	Expressive naming	Expressive Vocabulary Test	98.21 (13.84) ^b	-0.05	103.46 (13.54)	0.27 ^c
	Receptive language	Peabody Picture Vocabulary Test-III	106.86 (13.15) ^b	0.04	110.73 (10.13)	0.29 ^c
	Immediate verbal memory	CMS (word lists learning)	8.69 (3.05) ^{b,e}	0.11	9.80 (2.59)	0.01
Memory	Delayed verbal memory	CMS (word lists delayed)	9.06 (3.23) ^f	0.05	9.93 (2.85)	-0.03
	Problem-solving	D-KEFS (confirmed correct sorts)	9.06 (2.61) ^{b,e}	0.02	10.13 (1.97)	0.24
Executive Function	Response inhibition	D-KEFS (color-word interference test-Inhibition)	9.79 (2.78) ^{b,e}	-0.03	10.96 (2.47)	-0.17
	Speeded fine motor dexterity	Grooved Pegboard	85.10 (20.88) ^{b,d}	-0.16	72.54 (16.00)	0.23
Motor Function	Psychomotor speed	WISC-III (digit symbol-coding)	8.24 (2.65) ^{b,d}	-0.02	10.39 (2.96)	0.05

^a Covariates: age, gender, handedness, parent education

^b CWE<HC ($p < 0.05$)

^c $p < 0.05$

^d Raw score

^e Standardized score: M = 10, SD = 3

CWE: Children with epilepsy

HC: Healthy controls

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CWE: Children with epilepsy

WRAT-3: Wide Range Achievement Test - 3

WASI: Wechsler Abbreviated Scale of Intelligence

D-KEFS: Delis-Kaplan Executive Function System

CMS: Children's Memory Scale

WISC-III: Wechsler Intelligence Scale for Children - III