

Impaired Cerebellar-Dependent Eyeblink Conditioning in First-Degree Relatives of Individuals With Schizophrenia

Amanda R. Bolbecker¹⁻³, Jerilyn S. Kent¹, Isaac T. Petersen¹, Mallory J. Klaunig³, Jennifer K. Forsyth³, Josselyn M. Howell³, Daniel R. Westfall³, Brian F. O'Donnell¹⁻³, and William P. Hetrick^{*,1-3}

¹Department of Psychological and Brain Sciences, Indiana University, Bloomington, IN; ²Department of Psychiatry, Indiana University School of Medicine, Indianapolis, IN; ³Larue D. Carter Memorial Hospital, Indianapolis, IN

*To whom correspondence should be addressed; Department of Psychological and Brain Sciences, Indiana University, 1101 E. 10th Street, Bloomington, IN 47405, US; tel: 812-855-2620, fax: 812-856-4544, e-mail: whetrick@indiana.edu

Consistent with reports of cerebellar structural, functional, and neurochemical anomalies in schizophrenia, robust cerebellar-dependent delay eyeblink conditioning (dEBC) deficits have been observed in the disorder. Impaired dEBC is also present in schizotypal personality disorder, an intermediate phenotype of schizophrenia. The present work sought to determine whether dEBC deficits exist in nonpsychotic first-degree relatives of individuals with schizophrenia. A single-cue tone dEBC paradigm consisting of 10 blocks with 10 trials each (9 paired and 1 unpaired trials) was used to examine the functional integrity of cerebellar circuitry in schizophrenia participants, individuals with a first-degree relative diagnosed with schizophrenia, and healthy controls with no first-degree relatives diagnosed with schizophrenia. The conditioned stimulus (a 400 ms tone) coterminated with the unconditioned stimulus (a 50 ms air puff to the left eye) on paired trials. One relative and 2 healthy controls were removed from further analysis due to declining conditioned response rates, leaving 18 schizophrenia participants, 17 first-degree relatives, and 16 healthy controls. Electromyographic data were subsequently analyzed using growth curve models in hierarchical linear regression. Acquisition of dEBC conditioned responses was significantly impaired in schizophrenia and first-degree relative groups compared with controls. This finding that cerebellar-mediated associative learning deficits are present in first-degree relatives of individuals with schizophrenia provides evidence that dEBC abnormalities in schizophrenia may not be due to medication or course of illness effects. Instead, the present results are consistent with models of schizophrenia positing cerebellar-cortical circuit abnormalities and suggest that cerebellar abnormalities represent a risk marker for the disorder.

Key words: schizophrenia/eyeblink conditioning/cerebellum/relatives/associative learning/reflex conditioning/conditioned response/cognition/psychosis

Introduction

Motor abnormalities have long been observed in schizophrenia, even from its earliest conceptualization.¹ The cerebellum has historically been identified as an integral structure for coordinated movement and motor learning, and accumulating evidence points to an important role in nonmotor psychological processes as well—including cognition.²⁻⁵ Motor dysmetria is commonly observed subsequent to cerebellar lesions. However, consistent with evidence of cerebellar contributions to nonmotor processes, cerebellar lesions can also result in cognitive and behavioral symptoms, including impaired visuospatial memory, blunted affect or disinhibited, contextually inappropriate behavior, impaired executive function, and inattention.^{6,7} These symptoms are remarkably similar to those observed in schizophrenia, contributing to theoretical evidence that the cerebellum may play a role in the disorder.

Empirical evidence of cerebellar dysfunction in schizophrenia has been revealed through postmortem and neuroimaging studies, which report reduced cerebellar volume in chronic,⁸⁻¹¹ neuroleptic-naïve,¹² adolescent,¹³ first-episode,¹⁴⁻¹⁶ and childhood-onset¹⁷ schizophrenia (but for exceptions see Cahn and colleagues¹⁸ and Levitt and colleagues¹⁹). Postmortem studies report reduced size and density of Purkinje cells in schizophrenia.²⁰⁻²² Functional neuroimaging studies have found abnormal cerebellar blood flow at rest^{8,23,24} and during cognitive tasks²⁵⁻²⁷ in schizophrenia patients. Finally, cerebellar abnormalities

are associated with clinical symptoms, cognitive deficits, and outcome measures in schizophrenia.^{10,28–30} For example, deficits in working memory and mental flexibility correlate with cerebellar volume,³¹ and fronto-cerebellar metabolic abnormalities are associated with anhedonia and ambivalence.³² Moreover, increased connectivity between frontal-parietal and cerebellar regions predicts better cognitive performance in controls and schizophrenia, and schizophrenia patients with improved connectivity have fewer disorganization symptoms.³³

Our recent studies also indicate performance deficits in schizophrenia on a number of tasks linked to cerebellar function,^{34–36} most notably delay eyeblink conditioning (dEBC).^{37–40} Importantly, we have found significant dEBC associative learning deficits in schizotypal personality disorder.⁴⁰ This finding is consistent with reports of cerebellar white and gray matter abnormalities in individuals at high genetic risk for schizophrenia.⁴¹ First-degree relatives of individuals with schizophrenia have also been found to have structural and functional cerebellar abnormalities. For example, the developmental trajectory of the cerebellum is altered in relatives of individuals diagnosed with schizophrenia.⁴² Functional connectivity of cerebellum to areas including hippocampus, inferior frontal gyrus, and insula are also significantly reduced in schizophrenia and sibling relatives compared with controls.⁴³ Taken together, the foregoing evidence suggests that cerebellar abnormalities may serve as risk markers for schizophrenia.

dEBC provides a well-validated method to investigate the function of the cerebellum and related structures. In dEBC, a conditioned stimulus (ie, a tone) becomes associated with an unconditioned stimulus (ie, an air puff) after repeated paired presentations. Subjects demonstrate learning when an eyeblink (the conditioned response) occurs prior to the onset of the unconditioned stimulus. The neural circuits that underlie dEBC—where onset of the conditioned stimulus precedes that of the unconditioned stimulus, but they coterminate—are well characterized, and extensive evidence indicates that the cerebellum is essential to both the development and the manifestation of the eyeblink conditioned response.^{44,45} While additional cortical and subcortical brain areas *modulate* dEBC acquisition and response latency (ie, hippocampus, medial septum, and frontal cortex [see Christian and Thompson⁴⁶ for review]), convincing evidence suggests that the cerebellum is the essential site of neuroplasticity underlying expression of the eyeblink conditioned response.⁴⁷ Therefore, dEBC presents a useful method to assess the functional integrity of the cerebellum and related brain circuits.

This study set out to determine the extent to which cerebellar abnormalities may represent risk markers for schizophrenia by studying dEBC in nonpsychotic first-degree relatives of individuals with the disorder. The primary hypothesis was that schizophrenia and first-degree

relative groups would show impaired dEBC compared with healthy comparison subjects.

Methods

Participants

Participants were 18 individuals (5 female) diagnosed with either schizophrenia ($n = 14$) or schizoaffective disorder ($n = 4$; schizophrenia group), 18 individuals (11 female) with a first-degree relative diagnosed with schizophrenia or schizoaffective disorder (first-degree relative group), and 18 control participants (10 female) with no personal or family history of schizophrenia spectrum diagnoses (control group). With the exception of 1 participant with schizophrenia who had a relative that was also included in the study, no other participants in the study were related to each other. The patient sample was recruited through outpatient and inpatient units at community and state hospitals. Healthy controls were recruited using fliers posted in the community and from newspaper advertisements. First-degree relatives were recruited using contact information obtained from a larger sample of schizophrenia patients who were willing to provide such information. The Family Interview for Genetic Studies⁴⁸ was used to ascertain whether there were probable schizophrenia spectrum diagnoses in relatives of potential control participants. If a probable diagnosis was identified for a first-degree relative within the potential control participant's family, the participant was excluded in cases where the diagnosis of the probable family member with the disorder could not be ruled out through an in-person diagnostic interview, eg, the probable family member with schizophrenia or schizoaffective disorder did not wish to participate in the study. The potential control participant's group assignment was changed to first-degree relative if the relative was determined to meet diagnostic criteria for schizophrenia or schizoaffective disorder.

Of the 18 participants per group recruited for each study, data for 2 controls and 1 relative were excluded from the analysis because their data did not fit a positive linear growth model. (See data analysis section below for complete explanation).

Table 1 shows demographic, clinical, and medication information for the remaining 51 participants. The mean age of schizophrenia participants, controls, and relatives did not differ ($F(2,48) = 0.022$, $P = .98$), and sex was not significantly different across groups ($\chi^2(2) = 4.17$, $P = .13$). Education level was available for all except for 3 participants (1 in each group) and was found to differ across groups, $F(2,45) = 11.31$, $P < .001$. Bonferroni corrected comparisons showed that controls had more education than both the relative and schizophrenia groups.

Diagnostic status for the schizophrenia group was determined using the Structured Clinical Interview for Diagnostic and Statistical Manual of Mental

Table 1. Demographic, Clinical, and Medication Information

	Schizophrenia	Relatives	Controls
Age (y); <i>M</i> (<i>SD</i>)	36.0 (12)	35.9 (13)	36.8 (13)
Education Level ^a ; <i>M</i> (<i>SD</i>)	3.2 (1)	3.1 (1)	4.5 (1)
Sex (M:F)	13:5	7:10	7:9
PANSS total score; <i>M</i> (<i>SD</i>)	61 (12)	—	—
Positive; <i>M</i> (<i>SD</i>)	17 (5)	—	—
Negative; <i>M</i> (<i>SD</i>)	15 (4)	—	—
General; <i>M</i> (<i>SD</i>)	30 (7)	—	—
Past alcohol dependence ^b	7	3	0
Past drug dependence ^c	8	3	0
Psychotropic medication			
No medication	2	16	18
Atypical antipsychotic	13	0	0
Typical antipsychotic	3	0	0
Anticonvulsant	2	0	0
Antidepressant	7	2	0
Anticholinergic	6	0	0

Note: PANSS, Positive and Negative Syndrome Scale.

^aEducation level included self-report data on completion of grade school (1), junior high school (2), high school (3), some college (4), bachelor’s degree (5), master’s degree (6), and doctoral degree (7).

^bFive schizophrenia patients and 1 relative met criteria for both past alcohol and other drug dependence.

^cOther drug dependence included cannabis (*n* = 5 schizophrenia, *n* = 3 relatives) and cocaine (*n* = 4 schizophrenia).

Disorders-IV Axis I Disorders (SCID-I)⁴⁹ sections for mood disorders, psychotic disorders, and substance abuse disorders, as well as chart review. Kappa interrater reliability in our lab has been 0.95 for schizophrenia vs mood disordered, or other diagnoses in patients who have been prescreened for showing psychosis. Participants in the relative group were evaluated using the SCID-II for Axis II disorders⁵⁰ and the SCID-I. Control participants were interviewed using the nonpatient version of SCID-I⁵¹ sections for mood, psychotic, and substance abuse and the SCID II to exclude psychiatric disorders. Participants with schizophrenia underwent symptom assessment using the Positive and Negative Syndrome Scale (PANSS).⁵² All assessments occurred within 30 days of dEBC testing, and all but 4 occurred within 7 days of testing (*M* = 5.6 days; *SD* = 6.6). The average PANSS total score indicated that the patient group as a whole fell in the mildly-to-moderately ill range (*M* = 61.3, *SD* = 11.8; see table 2).⁵³

Exclusion criteria for all participants included a history of neurological or cardiovascular disease, clinically documented hearing loss, head injury resulting in loss of consciousness, electroconvulsive therapy, diagnosis of alcohol or other substance dependence within 3 months, and intelligence quotient (IQ) below 70. Control participants were excluded if they had a history of psychotic or mood disorder. Recruiting for this study occurred within the context of a larger effort in which approximately 30% of participants interviewed as potential controls were excluded using these criteria. Of the remaining 70%, approximately 8% were enrolled as first-degree relatives based on information obtained during interviews. Family

Table 2. Parameter Estimates for the Hierarchical Linear Modeling Growth Curve Model for Percentage of Conditioned Responses

	Value (SE)	<i>df</i>	<i>t</i> Value	<i>P</i> Value
<i>R</i> ² = 0.75				
Intercept	48.9 (5.6)	456	8.71	.000
SZ–HC	–12.8 (7.7)	48	–1.66	.103
Rel–HC	–13.1 (7.8)	48	–1.67	.101
Slope	4.4 (0.6)	456	7.50	.000
SZ–HC	–2.0 (0.8)	456	–2.54	.012*
Rel–HC	–2.4 (0.8)	456	–2.95	.003*

Note: SZ = schizophrenia, Rel = relatives, HC = healthy controls.
*Indicates significance at *P* < .0125.

members were not excluded for a diagnosis of schizotypal personality disorder (*n* = 1), schizoid personality disorder (*n* = 1), depression (*n* = 2), dysthymia (*n* = 1), or past alcohol or other substance dependence (*n* = 6), because these disorders may reflect expression of risk factors also associated with schizophrenia. Ten schizophrenia patients met criteria for past alcohol or other substance dependence. The Indiana University Human Subjects Institutional Review Board approved all study procedures, and written informed consent was obtained from all participants.

Eyeblink Conditioning Procedure

Participants completed a single-cue tone dEBC task. The conditioned stimulus was a 400ms, 1000 Hz (80 dB sound

pressure level) tone, which, on paired trials, coterminated with a 50 ms air puff to the eye, the unconditioned stimulus. Subjects were presented with 8 unconditioned stimulus-alone trials (intertrial interval = 15 s), followed by 10 blocks of conditioning trials (mean intertrial interval = 15 s; range = 10–20 s). Each trial block consisted of 9 paired trials, in which the tone and air puff were presented together, and 1 tone-alone trial. To maintain the participants' attention throughout the experiment, neutral photographs selected from the International Affective Picture System⁵⁴ were presented (2-s duration) between each trial, and participants rated the pleasantness of the images by pressing a response pad button. In addition, participants were observed via a closed circuit monitor to ensure that their eyes remained open. The experiment was briefly suspended if signs of fatigue were observed so that the examiner could interact with the participant.

Procedure

Two bipolar electromyographic electrodes (4 mm Ag/Ag–Cl) were placed within 1 cm below the left eyelid, centered under the pupil, and placed 1 cm apart. These recorded eyeblinks from the orbicularis palpebrarum muscle of the eye. A ground electrode was placed on the forehead. The inside corner of the left eye was presented with an unconditioned stimulus air puff (50 ms, 10 lb psi at source) delivered via copper tubing (fused to the rim of lens-less glasses) connected to a regulator delivering air via plastic tubing (120 in.). The conditioned stimulus tone was delivered via ear inserts (E-A-RLINK—Aearo Company Auditory Systems). Electromyographic recordings were made continuously (2.5 KHz A/D rate; high-pass filter = 1 Hz; low-pass filter = 500 Hz; gain = 1000) throughout the experiment and stored offline.

Data Processing

Continuous data files for each subject were divided into 1086 ms epochs starting 500 ms prior to conditioned stimulus onset. After a 28 Hz (6 dB/octave) high-pass filter was applied, the data were rectified and smoothed using a 41-point Gaussian weighted moving average. Data were entered into DataMunch, a Matlab computer program purposely written for EBC data analysis^{37,39,40,55–58} for further analysis of the 90 paired trials. Alpha responses, which are reflexive, nonassociative orienting electromyographic responses to the tone conditioned stimulus, were assessed between 25 and 100 ms after conditioned stimulus onset. On a subject-by-subject basis, responses were recorded as blinks if the amplitude exceeded five standard deviations above the baseline (baseline window for each trial = 125 ms prior to conditioned stimulus onset). The analysis incorporated a “bad trial” window that was used to exclude trials where electromyogram activity was increased immediately before (–75 ms) and shortly

after conditioned stimulus onset (+25 ms). If a participant exhibits electromyographic blink activity during this interval, it is unlikely that a conditioned response can be emitted immediately thereafter. That is, spontaneous blinks occurring during this “bad trial” window may interfere with the subsequent execution of a conditioned response. Blinks recorded in this window are considered spontaneous blinks because they occur too early in reference to conditioned stimulus onset to be considered either tone related or conditioning related. Accordingly, the number of “bad trials” can be used as a rough index of spontaneous blink rate. The average number of “bad trials” rejected from analysis did not differ between groups, $F(2,48) = 1.27, P = .29$.

Conditioned responses were recorded if the blink occurred between 100 and 350 ms after conditioned stimulus onset, which corresponded to a period beginning 250 ms before the onset of the unconditioned stimulus. The onset latency was calculated as the point in time when the conditioned response exceeded 0.5 SD from the baseline.

Hierarchical Linear Modeling

Electromyographic data were analyzed using growth curve models in hierarchical linear modeling (HLM). HLM is superior to repeated measures ANOVA because it takes into account the dependence of nested (multilevel) data. In HLM, the best-fitting line through each individual's trajectory is identified while taking into account the trajectories of other members of the sample. In this way, using multiple iterations, HLM increases the accuracy of the fit and decreases measurement error for both the population and for each individual. While heterogeneity of variance and violations of sphericity are treated as nuisance factors to be eliminated in traditional statistical approaches, including ANOVA, HLM has fewer assumptions than ANOVA and can accommodate unequal variances. This point is important because heterogeneity of variance can be potentially meaningful, and ignoring it can obscure significant interaction effects.⁵⁹

Statistical Analysis

For each individual, the percentage of conditioned responses was calculated for each of the 10 blocks of the experiment, and the best-fitting line was generated for these 10 data points, producing 54 lines—one line for each individual initially included in the study. Three individuals (2 controls and 1 relative) were then excluded from further analysis because they displayed negative learning curves, operationally defined as a decrease in conditioned responding of more than 20% from the first to the last block of the experiment using the fitted values of the model. (A figure including best-fitting line for

all participants, as well as complete parameter estimates and significance tests with these 3 outliers included in the model, can be found in the [online supplementary material](#).) Therefore, 51 participants were included in the final analysis.

To model learning curves of conditioned responses, we used the `lme` function of the `nlme` package⁶⁰ in R 3.0 (R Development Core Team, 2009) for growth curve modeling in HLM. In this study, HLM accounted for the dependency of data within individuals due to the repeated measures. Models used maximum likelihood estimation, except when testing whether effects should be fixed or random, in which case restricted maximum likelihood was used as suggested by Singer and Willett.⁶¹ We examined linear and nonlinear forms of change with nested model comparisons using the likelihood ratio test. Model fit was examined with pseudo- R^2 ,⁶¹ which was calculated by the squared correlation between the model's fitted and observed values, representing the proportion of variance in the outcome explained by the predictors.

After examining various forms of change, we settled on linear trajectories with random intercepts and slopes, which fit the data well. Linear growth curves were fit to each participant's learning curve across blocks and estimated whether groups differed in two parameters: intercepts and slopes. The intercepts reflected model-fitted performance during the first block and tested whether groups differed in their initial values at block 1. Slopes reflected change across blocks and tested whether groups differed in their learning rates.

The intercepts and slopes of the schizophrenia and relatives groups were compared with the control group, resulting in 4 separate statistical tests of between-group differences. Results with $P < .05$ are reported, but only those that survived Bonferroni-corrected alpha levels of $P < .0125$ ($P < .05/4$ comparisons) are deemed significant. The same analysis procedures were then applied to each of the following dEBC measures as secondary analyses: conditioned response onset latency, conditioned response amplitude, unconditioned response amplitude, and unconditioned response peak latency.

Results

Baseline Unconditioned Response Amplitude

In order to rule out blink performance differences between groups as a source of differences in percentage of conditioned responses, responses to 8 solitary unconditioned stimuli presented prior to the conditioning phase of the procedure were analyzed. Neither the average peak unconditioned response amplitudes ($F(2,48) = 0.60$, $P = .56$) nor latencies ($F(2,48) = 0.93$, $P = .40$) were significantly different across diagnostic categories, suggesting that differences in conditioned response rates between groups are unlikely to be due to deficits in sensorimotor processing or motor

responding in the schizophrenia and first-degree relative groups.

Conditioned Responses

Percentage of Conditioned Responses. The model fit the data well, with a pseudo- $R^2 = 0.75$. Individual fitted lines and the average line fit for each group can be seen in [figure 1](#), along with the 10 block averages for each group. Collapsing across groups, performance improved on average as the experiment progressed, $t(456) = 7.50$, $P < 0.001$, $SE = 0.58$. Although there were no significant differences between groups at block 1, the rate of learning varied between groups. Specifically, both the schizophrenia ($t(456) = -2.54$, $P = .012$, $SE = 0.80$) and relatives ($t(456) = -2.95$, $P = .003$, $SE = 0.81$) groups had significantly smaller increases compared with controls, indicating impaired acquisition of dEBC associative learning. Complete information regarding parameter estimates and model fit can be found in [table 2](#).

Conditioned Response Onset Latency and Amplitude. Neither the schizophrenia group nor the relatives group differed from controls on conditioned response onset latency or amplitude with respect to the intercept or slope. Moreover, collapsing across groups, neither onset latency nor amplitude showed a significant effect of slope, indicating that neither measure systematically changed on average as the experiment progressed.

Unconditioned Responses (Paired Trials)

Unconditioned Response Latency and Amplitude. When the slopes for unconditioned response latency and amplitude for both the schizophrenia and relatives groups were compared to controls, no significant differences emerged. Likewise, no significant differences between groups were found with respect to the intercept for unconditioned response latency. The schizophrenia group had a higher intercept compared with controls on unconditioned response amplitude, indicating larger amplitude responses from the start of the experiment, but this difference did not survive Bonferroni correction ($P = .018 > .0125$). Collapsing across groups, the unconditioned response latencies became longer on average as the experiment progressed ($t(456) = -2.26$, $P = .024$, $SE = 0.75$). Similarly, unconditioned response amplitude decreased significantly over time ($t(456) = -2.90$, $P = .004$, $SE = 1.37$).

Discussion

The present results illustrate a striking dEBC associative learning deficit in first-degree relatives of schizophrenia patients that is remarkably similar in magnitude to individuals who have been diagnosed with the disorder. This finding adds to a growing literature documenting cognitive and learning deficits in first-degree

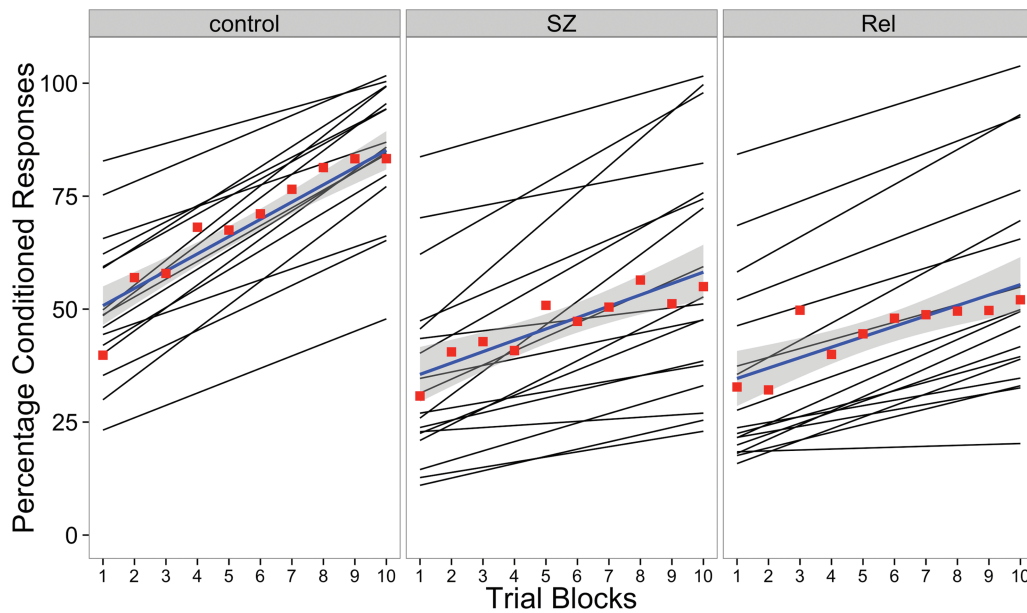


Fig. 1. Fitted lines for the percentage of conditioned responses across the 10 blocks of the experiment for each individual (black lines) with group averages for each block (red squares) and group averaged line fits.

relatives of individuals with schizophrenia, but extends previous findings by demonstrating an impairment on a fundamental form of cerebellar-dependent associative learning—dEBC.

The observed dEBC abnormalities in first-degree relatives and in schizophrenia may arise from aberrant cerebellar processing. Specifically, on the basis of an extensive animal and human literature, these deficits implicate anomalies both in the cerebellar cortex and deep interpositus nucleus and, perhaps, in the function of the Purkinje cells that project from the cortex to the deep nuclei of the cerebellum in both individuals with schizophrenia and their first-degree relatives. Animal studies have shown that the critical association between the conditioned and unconditioned stimuli occurs in the anterior interpositus nucleus,^{62–64} one of the deep cerebellar nuclei. The cortex of the cerebellum is thought to control the expression of conditioned responses and modulate both their timing and amplitude.^{45,65,66} The anterior lobe, through Purkinje cells projecting to the interpositus, appears to play a critical role in response timing, delaying the onset of conditioned responses until just prior to the unconditioned stimulus onset.^{67,68} Findings from these animal studies are supported by studies in humans suggesting that abnormalities in cerebellar structure are associated with dEBC performance.^{69,70}

Our findings are consistent with reports of cerebellar neurotransmitter, neurochemical, and genetic abnormalities in schizophrenia patients^{46,53,71–73} and with previous reports of structural cerebellar anomalies in family members of individuals with schizophrenia.⁴² Taken with the foregoing evidence of critical cerebellar involvement in EBC, the finding that a similar pattern of dEBC deficit occurs in first-degree relatives of individuals with

schizophrenia suggests that these abnormalities may be a core feature of schizophrenia.

An advantage of the cross-sectional design of this study is that any lingering questions about the origins of dEBC abnormalities observed in schizophrenia can now be more directly addressed. Because dEBC deficits are observed in first-degree relatives they are unlikely to be associated with the onset of schizophrenia, but instead may represent risk factors. Investigations such as the present one that examined nonpsychotic first-degree relatives are powerful tools in the identification of risk markers of schizophrenia for several reasons. First, familial studies indicate that first-degree relatives have elevated risk for schizophrenia spectrum disorders.⁷⁴ Second, first-degree relatives share genetic (and some environmental) risk factors with affected probands without exhibiting psychotic disorder and are estimated to have at least 10-fold greater risk for developing the disorder.^{75–77} Finally, compared with individuals with schizophrenia, first-degree relatives have far less exposure to psychotropic medications, especially antipsychotic drugs. Therefore, the observation of dEBC deficits in schizophrenia *and* in first-degree relatives strengthens the conclusion that dEBC deficits observed in schizophrenia in this study (and others)^{37–40,78} are unlikely to be due psychotropic medications. This point is important because a recent review⁷⁹ urged caution in ascribing dEBC abnormalities to illness mechanisms and argued that antipsychotic medication could account for conditioning deficits. The present findings of dEBC deficits in relatives also suggest that impaired dEBC in schizophrenia is unlikely to be due to course of illness variables.

Taken together with findings from this study, given that (1) schizophrenia is highly heritable,⁸⁰ (2) first-degree relatives are at substantially higher risk for the onset of schizophrenia,⁷⁴ (3) cerebellar abnormalities have been found using a wide variety of methodologies and paradigms and at multiple levels of analysis in schizophrenia (see Andreasen and Pierson⁸¹ for review), and (4) first-degree relatives show alterations in the cerebellum,^{42,43,82,83} these findings converge on the conclusion that dEBC abnormalities may represent a risk factor for schizophrenia.

Conclusions, Limitations, and Future Directions

While it seems likely that the observed deficits in dEBC are related to a failure in first-degree relative and schizophrenia to acquire a cerebellar-mediated conditioned response, a distributed network of brain regions participate in this form of learning. Therefore, neuroimaging methods can more definitively identify the extent to which these deficits are uniquely related to cerebellar alterations, and such studies are currently underway in our laboratory.

General intelligence has also been linked to cerebellar structure and function. For example, there is evidence linking cerebellar volume to IQ^{84–86} and our own data have shown an association between IQ and dEBC that was evident in controls but not in individuals with schizophrenia.⁸⁷ Reduced IQ is commonly found in schizophrenia,⁸⁸ and nonpsychotic relatives from multiplex families also reportedly have lower IQ compared with healthy control samples.⁸⁹ Therefore, the present findings suggest that a careful study of the relationship between cognition, symptom profiles, genetic loading for schizophrenia, and cerebellar measures is warranted.

The findings reported here are important because they suggest cerebellar abnormalities may represent potential risk markers of schizophrenia although it has not previously been commonly considered as such. In the context of the present findings and given the known role of the cerebellum in motor coordination and motor learning, it is worth noting that early motor abnormalities have convincingly been associated with worsening of prodromal symptoms in high-risk groups^{90,91} and the subsequent development of schizophrenia spectrum disorders.^{92–94} Early evidence suggests that consideration of novel treatment strategies focused on improving cerebellar function could represent a novel therapeutic approach. For example, we and others have shown the efficacy of pharmacological⁵⁵ and repetitive transcranial magnetic stimulation⁹⁵ targeting of cerebellar-dependent deficits. Our laboratory has studies underway that are aimed at characterizing the sensitivity of an ensemble of candidate biomarkers to cerebellar abnormalities in schizophrenia. This biomarker validation will provide an essential framework for translational research and intervention development.

Supplementary Material

Supplementary material is available at <http://schizophrenia.bulletin.oxfordjournals.org>.

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