Effects of regulatory BCI RNA deletion on synaptic plasticity, learning, and memory

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Brain cytoplasmic (BC) RNAs are a nonprotein-coding subtype of small cytoplasmic RNAs. BCI RNA (BC200 in human and nonhuman primates) localizes to synapto-dendritic domains in neurons and regulates protein synthesis by targeting the initiation phase of translation (Tiedge et al. 1991, 1993; Wang et al. 2002, 2005; Eom et al. 2011, 2014). BCI RNA may contribute to maintaining translation homeostasis at synapses because it (1) represses translation stimulated by group 1 metabotropic glutamate receptor (mGluR) activation, and (2) group 1 mGluR-stimulated translation of synaptic proteins is exaggerated in the absence of BCI RNA (Wang et al. 2002; Lin et al. 2008; Eom et al. 2014). In contrast to the substantial knowledge of the molecular actions of small cytoplasmic RNAs, relatively little is known about the role of BC RNAs in neural function and behavior. Here, using BCI knockout (KO) mice, we investigated whether absence of BCI RNA alters hippocampus-dependent cognitive behavior. Two lines of BCI KO mice were used; each derived from independent mutant ES cell lines. Each line initially had a mixed C57BL/6J and 129/SvJ background (Skryabin et al. 2003). The first “BC1 KO-13” line still has the mixed C57BL/6J and 129×1/SvJ background. The second, “BC1 KO-B” line, mostly has a C57BL/6J background, having been backcrossed over 15 generations. The corresponding WT-13 and WT-B wild-type (WT) mice were of the respective identical background. The mice were bred in house and were 8–12 wk old at the time of the physiological and behavioral assays. All procedures were carried out in accordance with the Public Health Service Policy on Humane Care and Use of Laboratory Animals and were approved by the SUNY, Downstate Medical Center and the New York University Institutional Animal Care and Use Committees.

To test whether the absence of BCI RNA alters mGluR-stimulated hippocampus synaptic function, we investigated mGluR-stimulated long-term depression DHPG-induced LTD of CA3-to-CA1 Schaffer collateral synapses (Huber et al. 2001). Ex vivo electrophysiology was performed as described (Alarcon et al. 2004; Nichols et al. 2008). Right dorsal transverse hippocampal slices (400 μm) were cut in ice-cold artificial cerebrospinal fluid (ACSF, in mM: NaCl 119, KCl 4, MgSO4 1.5, CaCl2 2.5, NaHCO3 26.2, NaH2PO4 1.0, and glucose 11, saturated with 95% O2, 5% CO2). Slices recovered at 36°C for 45 and 60 min at room temperature in ACSF and were then set in an interface recording chamber perfused with ACSF at 36°C. Field excitatory postsynaptic potentials (fEPSPs) from the CA1 “stratum radiatum” were evoked (pulse duration: 50 μs, pulse sampling: 0.017 Hz, pulse intensity: 40% of the maximum fEPSP slope) and detected using a pair of stimulation bipolar (FHC & Co) and recording borosilicate glass pipette (5–10 mΩ filled with ACSF; Sutter Instruments) electrodes. The group 1 mGluR agonist 3,5-dihydroxyphenylglycine (DHPG, 50 μM; Sigma-Aldrich) was used to induce DHPG-induced LTD (Huber et al. 2001). fEPSP responses were acquired using PClamp (Molecular Devices) and analyzed using Origin (Microcal Software). ANOVA with repeated measures, followed by the Holm–Šidák test for multiple comparisons were performed using Prism (GraphPad Software).

Expression of DHPG-induced LTD was quantified as the reduction in fEPSP slope 40 min after DHPG treatment relative to the 15-min baseline prior to application of DHPG (Fig. 1). Because DHPG-induced LTD was indistinguishable for WT-13 mice and WT-B mice, they were combined into a single WT group. This was statistically justified because there were 5 mice per group and one-way ANOVA power analysis with three groups, α = 0.05 and β = 0.95, and the observed effect size f = 0.75, calls for group sizes of four. BC1 KO-B animals showed enhanced DHPG-induced LTD expression (65.18 ± 14.06%) compared with WT mice (86.45 ± 6.42%) and it was even larger in BC1 KO-13 mice (40.64 ± 10.20%). These observations were confirmed by a significant effect of genotype (F2,23 = 29.96; P < 0.0001), and according to post hoc tests, the ranking of DHPG-induced LTD enhancement was BC1 KO-13 mice > BC1 KO-B mice > WT mice. These data indicate a role of BC1 RNA regulation of group 1 mGluR-dependent synaptic function, which is greater in BC1 KO-13 mice than BC1 KO-B mice.

Next we evaluated cognitive behavior in these two BCI-KO lines by testing them in the hippocampus synaptic plasticity-dependent active place avoidance task, learning and memory were impaired in the BCI-KO line with the more severely altered DHPG-induced LTD. These findings indicate a role for BC1 RNA control of mGluR-dependent synaptic function in hippocampus and associated cognitive ability.

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Article is online at http://www.learnmem.org/cgi/doi/10.1101/lm.045617.117.
are indicated by black asterisks, and relative to BC1 KO-B mice by gray asterisks (***).

DHPG-induced LTD is exaggerated in slices from BC1 KO-13 mice, in comparison with those from WT and BC1 KO-B mice. (A) The graph shows the time course of CA3–CA1 synaptic responses during baseline recordings, induction of LTD by DHPG (50 µM for 20 min, solid gray line), and the washout period (40 min). DHPG-induced LTD is exaggerated in slices from BC1 KO-13 mice, in comparison with those from WT and BC1 KO-B mice. (B) Traces illustrate representative examples of evoked responses. 

Locomotion was similar, measured by the total distance moved across the three training trials, all genotypes decreased the number of times they entered the shock zone, indicating place learning (Fig. 2B). The WT and BC1 KO-B mice also showed the typical decrease in the number of entries across the two conflict sessions in all the genotypes, but long-term memory was not expressed in BC1 KO-13 mice as their path to the first entrance did not appear to increase, indicating impaired cognitive flexibility. These impressions were confirmed by two-way genotype × trial ANOVAs, with repeated measures on the factor trial. The effects of genotype (entrances: $F_{(2,20)} = 9.73, P < 0.005$; path to first enter: $F_{(2,20)} = 3.55, P < 0.05$) and trial (entrances: $F_{(6,120)} = 158.7, P < 0.0001$; path to first enter: $F_{(6,120)} = 6.50, P < 0.0001$) were significant for both the measure of learning (Fig. 2B) and the measure of long-term memory retention (Fig. 2C). The interaction was significant for the number of entrances ($F_{(12,120)} = 2.63, P < 0.005$) but not for the path to the first entrance ($F_{(12,120)} = 1.26, P > 0.25$). Post hoc tests confirmed no differences between WT and BC1 KO-B mice, and relative to these genotypes, the BC1 KO-13 mice were impaired on the initial learning and conflict trials (Fig. 2B). The BC1 KO-13 mice expressed impaired long-term memory relative to WT and BC1 KO-B mice on the 24-h retention and the two conflict trials once long-term memory was strongly expressed by the other genotypes (Fig. 2C). BC1 KO-13 mice express cognitive impairments in active place avoidance not observed in BC1 KO-B mice. While there is no evidence that DHPG-induced LTD abnormality causes the cognitive deficits, a larger abnormality in DHPG-induced LTD is observed in BC1 KO-13 mice, and we do not know what additional consequences the loss of BC1 RNA throughout the brain may have had (e.g., compensatory changes). The present findings demonstrate functional abnormality after loss of BC1 RNA by demonstrating, for the first time, synaptic plasticity changes and parallel impairment of behavior known to involve the altered synapses.

Prior behavioral work contrasts with the present findings. In that work, BC1-mutant mice expressed impaired spontaneous exploratory behavior and higher anxiety-like behavior but no differences in learning or memory assessed by a test battery that included the Barnes, Water, Radial 8-Arm, Multiple-T, and alley mazes (Leweijohann et al. 2004). The task variant we used is no more stressful than exploration of a familiar space (Lesburgueres
set-shifting task was also recently reported in these BC1 KO strains (Iacoangeli et al. 2017).

Absence of BC1 RNA resulted in exaggerated DHPG-induced LTD. This follows because dendritically transported BC1 RNA interacts with the translation initiation machinery by binding to eu-karyotic initiation factors eIF4A (an RNA helicase) and eIF4B (a modulator of eIF4A function), thereby blocking the recruitment and formation of 48S initiation complexes (Wang et al. 2002; Lin et al. 2008; Eom et al. 2014). While we did not experimentally block translation in the present studies, anisomycin was used in prior investigations of BC1-mutant mice that confirmed BC1 RNA represses translation downstream from MEK–ERK signaling in a sequential-independent manner to the fragile X mental retardation protein (FMRP) (Zhong et al. 2010). Consequently, BC1 RNA absence results in hippocampal CA3 hyperexcitability following group 1 mGluR stimulation and audiogenic seizures, both of which are blocked by pretreatment with anisomycin (Zhong et al. 2009), and are further exaggerated when FMRP is also absent (Zhong et al. 2010). While it is beyond the present scope to determine whether other forms of synaptic function and plasticity are also altered, diverse abnormalities are predicted given the broad expression of BC1 RNA and its role in repressing translation.

The present findings add to prior reports of increased synthesis of synaptic proteins (Cheng et al. 1996), increased prolonged CA3 epileptiform discharges, increased susceptibility to audiogenic seizures, and excessive cortical $\gamma$ oscillations in the absence of BC1 RNA, all of which are triggered by group 1 mGluR stimulation (Zhong et al. 2009, 2010; Iacoangeli and Tiedge 2013). Given that BC1 RNA and FMRP both repress translation, albeit by different mechanisms, it is noteworthy that all of these abnormalities have also been observed in Fmr1 KO mice that lack FMRP to model the FXS genetic defect (Huber et al. 2001; Chuang et al. 2005; Zhong et al. 2010; Bhakar et al. 2012; Gonçalves et al. 2013; Radwan et al. 2016). Causal relationships remain to be confirmed among these abnormalities that span the molecular, cellular, network, and behavioral levels of biological organization.

The two BC1 KO strains differed in the severity of the abnormalities that we measured. Prior work with the BC1 KO-B mice showed a mild active place avoidance deficit (Zhong et al. 2010), using a harder to learn protocol of three 10-min sessions per day for 3 d (Tsokas et al. 2016), however, the deficit was not as severe as we observed in the BC1 KO-13 strain. It is well documented that both physiology and behavior can be impacted by the background strain of a mouse (Yoshiki and Motoiwa 2006; Spencer et al. 2011; Darrell 2014; Ishimura et al. 2014; Lei et al. 2014). Phenotypic differences including fear-learning deficits due to Fmr1 deletion are weaker in C57Bl/6 mice compared with the 129×1/SvJ strain (Paradee et al. 1999; Moy et al. 2009; Spencer et al. 2011). The strain differences we observed could arise because BC1 RNA translational control is extraordinary in the C57Bl/6 strain compared with other strains (Darrell 2014; Ishimura et al. 2014), and thus less severe phenotypic consequences of BC1 RNA absence could manifest in the BC1 KO-B line. Known complex molecular interactions may also play a role. Specifically, BC1 RNA binds to FMRP to increase the protein’s mRNA target specificity (Zalfa et al. 2003). In addition, FMRP expression is higher in the absence of BC1 RNA, measured with both background strains (Zhong et al. 2009). Thus the differences we detected in the phenotypic expression of the two BC1 KO strains highlight the potential importance of genetic background. In the present case, there is both the possibility of background-strain dependent partial compensation for loss of BC1-mediated translation repression by FMRP as well as exaggerated FMRP loss of function in the absence of BC1 RNA (Zalfa et al. 2003; Zhong et al. 2009). In addition, the differences may also result from the nonlinear relationships between deficits in molecular signaling, physiological, and
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behavioral consequences, a fact that presents a major challenge to understanding how molecular mechanisms of disease and dysfunction cause clinical symptoms (Mitchell et al. 2013). The present findings add to the evidence that translation regulation in neurons, including by nonprotein coding RNA, is a mechanism of complex neural circuit and brain function, the disruption of which is sufficient to result in synaptic plasticity and cognitive dysfunction (Darnell et al. 2011).

Acknowledgments
We thank Henri Tiedge for providing mice. This work was supported in part by National Institutes of Health (NIH) grants NS081625 and NS091830 (J.M.A.), MH099128 (A.A.F.).

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Received March 17, 2017; accepted in revised form August 17, 2017.