Adolescent alcohol exposure epigenetically regulates CREB signaling in the adult amygdala

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Binge alcohol drinking in adolescence leads to increased risk for alcohol use and other psychiatric disorders in adulthood. The transcription factor cAMP-response element binding (CREB) protein is involved in the neuronal response to adult ethanol exposure, but its role in the enduring effects of adolescent alcohol exposure in adulthood is unknown. We exposed male rats to adolescent intermittent ethanol (AIE) or saline (AIS) during post-natal days 28–41 and evaluated the epigenetic regulation of CREB dynamics in the adult amygdala. A subset of these adult rats was exposed to an acute ethanol challenge. AIE decreased CREB, phosphorylated CREB, CREB-binding protein (CBP) and p300 protein levels in adult amygdaloid brain structures. AIE exposure also causes deficits in Creb1, Cbp, and p300 mRNA expression in the amygdala of AIE adult rats which are normalized after acute ethanol exposure. Interestingly, occupancy of acetylated histone H3K9/14 proteins at specific locations in the Creb1, Cbp, and p300 gene promoter regions was decreased in the amygdala of AIE adult rats and was normalized by acute ethanol exposure. These results suggest that AIE exposure epigenetically reduces CREB and other related transcriptional activators in the amygdala in adulthood that may be associated with the behavioral effects of adolescent alcohol exposure.

Frequent binge drinking is a risk factor for the development of alcoholism, as ethanol consumption is more likely to result in later addiction if binge amounts are consumed early in life1,2. Binge drinking is associated with acute pathophysiological symptoms including learning and memory impairments and disturbances in sleep-wake patterns3–5. Long-term sequelae of adolescent binge drinking include increased alcohol use, decreased educational and economic attainments, and increased risk for anxiety and depression6–8. Animal models of binge ethanol exposure in adolescence recapitulate many of these phenotypes including increased alcohol consumption and anxiety in adulthood9–10. Repeated high-dose alcohol exposure in adolescence leads to neuroadaptations in specific brain circuits and signaling pathways that underlie behavioral changes seen in addiction11,12. The limbic system, and particularly the amygdala, plays a critical role in the neurobiology of alcoholism and the negative affective states, including anxiety, that are associated with alcohol abuse11,13,14. In humans, the amygdala undergoes a number of structural and connectivity changes during adolescent development15, and this process is disrupted by binge alcohol abuse in adolescence16. Furthermore, previous studies in rodent models of adolescent binge drinking have indicated that there are substantial changes in signaling pathways in the amygdala that may mediate the increased anxiety-like and alcohol drinking behaviors seen in adulthood in these models17,18.

The cAMP-response element binding protein (CREB) is an important transcription factor that is activated by G-protein-coupled receptor signaling cascades and intracellular calcium influx19. CREB signaling plays a critical role in neuronal activity and is dysregulated in pathological states such as addiction20,21. Neuronal activity causes an increase in CREB phosphorylation22,23, which leads to nuclear localization and recruitment of CREB binding protein (CBP), p300, and other transcription factors to the chromatin24–26. CBP and p300 possess intrinsic histone acetyltransferase (HAT) activity, catalyzing the addition of acetyl groups to N-terminal tails of histone27–29. This epigenetic modification is generally associated with increased gene transcription30.

Acute ethanol exposure in adult rats increases CREB phosphorylation in the amygdala, while withdrawal after chronic ethanol treatment decreases CREB and CBP expression31. Reduced levels of CREB and phosphorylated CREB...
CREB (pCREB) have been shown in the central nucleus of the amygdala (CeA) and medial nucleus of the amygdala (MeA) of alcohol-preferring (P) adult rats, which display heightened anxiety-like behavior and increased alcohol consumption as compared with alcohol non-preferring (NP) rats. The involvement of CREB, CBP, and p300 in alcohol exposure is not surprising given the intrinsic HAT activity of CBP and p300 and the widespread modulation of epigenetic mechanisms by alcohol. In addition, HATs dynamically interact with histone deacetylases (HDACs) to epigenetically regulate synaptic plasticity. We recently demonstrated that a specific HDAC isoform, HDAC2, is increased and histone H3K9 acetylation is decreased in the CeA and MeA after AIE in adulthood. Recently, we also observed that an acute ethanol challenge in adulthood attenuates AIE-induced anxiety-like behaviors. However, the changes in the CREB signaling in adulthood, including the HATs CBP and p300 and their epigenetic regulation, following adolescent alcohol exposure is currently unknown. Therefore, the present study explored CREB signaling and the epigenetic regulation of CREB-related molecules in the amygdala of rats after AIE in adulthood. We also examined the effects of an acute challenge of ethanol on the molecular manifestations of AIE in adulthood. Our results show for the first time that adolescent alcohol exposure causes lasting, epigenetically-encoded deficits in CREB signaling in the amygdala at adulthood.

Results
AIE decreases CREB and pCREB levels in the amygdala in adulthood. To explore the lasting influence of adolescent alcohol exposure on CREB signaling-related protein levels, we measured CREB and pCREB levels in the amygdala of AIE and AIS adult rats. We observed a significant reduction in CREB and pCREB protein levels in the CeA (CREB: t(10) = 12.26, p < 0.001; pCREB: t(10) = 12.52, p < 0.001) and MeA (CREB: t(10) = 13.17, p < 0.001; pCREB: t(10) = 10.28, p < 0.001), but not in the BLA, of AIE adult rats compared to AIS adult rats. Additionally, Creb1 mRNA levels are significantly decreased (t(10) = 4.15, p = 0.002) in the amygdala of AIE adult rats compared to AIS adult rats.

AIE increases histone acetyltransferases (HATs) CBP and p300 in the amygdala in adulthood. pCREB recruits CBP and the related molecule p300 which both act to transfer acetyl groups to histone proteins and drive transcriptional activity. Due to the effect of AIE on amygdala CREB and pCREB expression, as well as the involvement of histone acetylation mechanisms in the long-lasting effects of AIE, we measured protein levels of HATs (CBP and p300) in the amygdala following AIE. CBP is significantly decreased in the CeA (t(10) = 14.83, p < 0.001) and MeA (t(10) = 11.53, p < 0.001), but not the BLA, of AIE adult rats compared to AIS adult rats.
adult rats (Fig. 2a,b). Additionally, Cbp mRNA levels are decreased (t (13) = 2.41, p = 0.032) in the amygdala of AIE adult rats as compared with AIS adult rats (Fig. 2c).

We next measured p300 protein and mRNA levels in the amygdala of these rats. p300 protein levels are significantly decreased in the CeA (t (10) = 8.17, p < 0.001) and MeA (t (10) = 8.05, p < 0.001) of AIE adult rats when compared to AIS adult rats, and there was no significant difference in p300 protein levels in the BLA between groups (Fig. 3a,b). p300 mRNA levels are also decreased (t (13) = 2.19, p = 0.047) in the amygdala of AIE adult rats when compared to AIS adult rats (Fig. 3c).

**Acute alcohol challenge in adulthood normalizes AIE-induced changes in mRNA expression in the amygdala.** We exposed adult rats that had been previously treated with AIE or AIS in adolescence to an acute challenge of alcohol (2 g/kg). Previously, this acute alcohol challenge was shown to normalize the anxiety-like behaviors and transcriptional and epigenetic changes seen in the amygdala of AIE adult rats. We found that there was an overall group effect of AIE on mRNA levels of Creb1 (F(AIE),130 = 15.7, p < 0.001) among the groups. Also, significant effects of both acute ethanol exposure (F(acute EtOH),130 = 28.1, p < 0.001) and an interaction between AIE and acute ethanol (F(AIE x acute EtOH),130 = 43.5, p < 0.001) were observed. Post hoc comparison revealed that AIE significantly (p < 0.001) decreased Creb1 mRNA levels in the amygdala of AIE adult rats, which was normalized (p < 0.001) to control levels after acute ethanol exposure (Fig. 4a). We also found that Cbp mRNA levels were significantly modulated by AIE (F(AIE),130 = 46.7, p < 0.001) and acute ethanol exposure (F(acute EtOH),130 = 53.0, p < 0.001), with a significant interaction of AIE with acute ethanol exposure (F(AIE x acute EtOH),130 = 6.20, p = 0.019). Post hoc comparison indicated a significant reduction (p = 0.006) in Cbp mRNA levels in the amygdala of AIE adult rats as compared with AIS adult rats. Interestingly, Cbp mRNA levels were significantly increased in the amygdala by acute alcohol in both the AIS (p < 0.001) and AIE (p = 0.002) adult rats (Fig. 4a). Similarly, a two-way ANOVA revealed a significant effect of AIE as well as acute ethanol exposure on p300 mRNA levels (F(AIE),130 = 79.3, p < 0.001; F(acute EtOH),130 = 109.6, p < 0.001), and also a significant interaction between AIE and acute ethanol exposure (F(AIE x acute EtOH),130 = 22.5, p < 0.001) was observed. Furthermore, post hoc comparison indicated a significant decrease by AIE (p = 0.007) and an increase in p300 mRNA levels by acute alcohol exposure in both AIS (p < 0.001) and AIE (p < 0.001) adult rats (Fig. 4a). These results suggest that AIE produces long-lasting reductions in the mRNA levels of Creb1, Cbp, and p300 in the amygdala which are normalized by acute ethanol challenge in adulthood.
Figure 3. Effects of adolescent intermittent ethanol (AIE) exposure on p300 levels in the amygdala of adult rats. (a) Representative photomicrographs (Scale bar = 50 μm) of p300 immunolabeling in the amygdaloid brain structures of AIE and adolescent intermittent saline (AIS) exposed adult rats, and (b) Bar diagram showing quantification of p300 gold immunolabeling in the central (CeA), medial (MeA) and basolateral amygdala (BLA) of AIE and AIS adult rats. Values are presented as the mean ± SEM of the number of immuno-gold particles/100 μm². (c) Bar diagram showing fold changes in mRNA levels of p300 in the amygdala of AIS and AIE adult rats. Values are presented as the mean ± SEM of the fold changes relative to AIS control rats. *p < 0.05, **p < 0.01, Student's unpaired two-tailed t-test, n = 6/group (Gold immunolabeling), n = 7–8/group (mRNA studies).

AIE-induced deficits in histone acetylation of Creb1, Cbp, and p300 promoters in the amygdala at adulthood & reversal by acute ethanol challenge. Given the involvement of epigenetic mechanisms in AIE-induced neuroadaptation and our previous work showing decreased global histone acetylation in the CeA and MeA of AIE-exposed adults, we investigated the promoter regions of Creb1, Cbp, and p300 for the occupancy of acetylated H3K9/14 (H3K9/14ac) using the chromatin immunoprecipitation (ChIP) assay. We found that H3K9/14ac levels were significantly altered by both AIE exposure (F(AIE), 1,19 = 25.2, p < 0.001) and acute alcohol challenge in adulthood (F(acute EtOH), 1,19 = 45.6, p < 0.001), but not by the interaction between AIE and acute alcohol, in the Creb1 proximal promoter site (+593 base pairs [bp] upstream of transcription start site), with post hoc testing revealing a decrease in the amygdala of AIE animals at baseline (p < 0.001) and an increase by acute alcohol in both AIS (p < 0.001) and AIE (p < 0.001) adult rats (Fig. 4b). However, there were no significant differences between the groups at a separate Creb1 promoter (+375 bp) site containing a cAMP response element (CRE; Fig. 4b).

In addition, we found that H3K9/14ac levels in the amygdala were significantly altered by both AIE (F(AIE), 1,20 = 12.3, p = 0.002) and acute ethanol (F(acute EtOH), 1,20 = 31.0, p < 0.001), but not the interaction between AIE and acute ethanol, at the Cbp proximal promoter site (+3014 bp), with AIE adult rats showing a reduction (p = 0.001) at baseline compared to AIS adult rats, and both AIS (p = 0.014) and AIE adult rats (p < 0.001) exposed to acute alcohol displaying increased levels of H3K9/14ac compared to their saline-exposed counterparts (Fig. 4c). Additionally, H3K9/14ac levels at the Cbp proximal promoter site (+590 bp) were significantly altered by only acute ethanol exposure (F(acute EtOH), 1,20 = 5.95, p = 0.024). AIE rats exposed to acute ethanol displayed increased H3K9/14ac levels at the Cbp proximal promoter site (+590 bp) compared to AIE adult rats exposed to saline (p = 0.034; Fig. 4c).

At the p300 proximal promoter site (+1536 bp), we observed a main effect of altered H3K9/14ac occupancy in the amygdala by acute ethanol (F(acute EtOH), 1,20 = 12.2, p = 0.002), with AIS (p = 0.035) and AIE rats (p = 0.015) exposed to an acute ethanol challenge displaying increased H3K9/14ac levels at this site compared to saline-exposed AIS and AIE rats, respectively (Fig. 4d). The occupancy of H3K9/14ac was also significantly different between groups at the p300 proximal promoter (+198 bp) site with main effects of acute ethanol (F(acute EtOH), 1,20 = 49.5, p < 0.001) and the interaction between AIE and acute ethanol (F(AIE x acute EtOH), 1,20 = 8.55, p = 0.008), with post hoc testing showing a decrease in the amygdala of AIE animals at baseline (p = 0.006) and an increase by acute alcohol in both AIS (p = 0.009) and AIE (p < 0.001) adult rats (Fig. 4d). These epigenetic changes in the amygdala possibly explain some of the transcriptional alterations in Creb1, Cbp, and p300 mRNA produced by AIE in adulthood.
Discussion

In the present study, we found that AIE produced long-lasting deficits in CREB signaling in the amygdala via histone acetylation mechanisms. The CREB-related signaling factors CBP and p300, possessing intrinsic HAT activity, displayed a decrease in mRNA and protein levels in the CeA and MeA in the adult amygdala after AIE. The deficits in Creb1, Cbp, and p300 mRNA levels were normalized in the amygdala of AIE adult rats exposed to an acute challenge of ethanol or saline in adulthood. Occupancy of acetylated histone 3 lysine 9 and 14 (H3K9/14ac) protein was investigated using the chromatin immunoprecipitation (ChIP) assay and primers specific for two sites in the promoter regions of Creb1, Cbp, and p300 in the amygdala of AIE and AIE adult rats exposed to an acute challenge of ethanol or saline. Values are presented as the mean ± SEM of the fold changes relative to AIS + Saline control rats. *p < 0.05, **p < 0.01, ***p < 0.001, two-way ANOVA followed by Tukey's post hoc test, (n = 8–9/group for mRNA studies and n = 5–6/group for ChIP assay). CRE = cAMP response element; bp = base pairs.

Figure 4. Acute alcohol challenge in adulthood epigenetically normalizes Creb1, Cbp, and p300 mRNA expression in the amygdala of rats exposed to adolescent intermittent ethanol (AIE). (a) Bar diagram showing fold changes in mRNA levels of Creb1, Cbp, and p300 in the amygdala of adolescent intermittent saline (AIS) and AIE rats exposed to an acute challenge of ethanol or saline in adulthood. Occupancy of acetylated histone 3 lysine 9 and 14 (H3K9/14ac) protein was investigated using the chromatin immunoprecipitation (ChIP) assay and primers specific for two sites in the promoter regions of Creb1, Cbp, and p300 in the amygdala of AIE and AIE adult rats exposed to an acute challenge of ethanol or saline. Values are presented as the mean ± SEM of the fold changes relative to AIS + Saline control rats. *p < 0.05, **p < 0.01, ***p < 0.001, two-way ANOVA followed by Tukey's post hoc test, (n = 8–9/group for mRNA studies and n = 5–6/group for ChIP assay). CRE = cAMP response element; bp = base pairs.
the CeA decreased pCREB levels, provoked anxiety-like behaviors, and increased ethanol intake of adult NP rats\(^25\). Similarly, CREB haploinsufficient adult mice display increased alcohol drinking and anxiety-like behaviors compared to wild-type littermates\(^25\). Sprague-Dawley adult rats display decreased pCREB levels in the CeA and MeA and increased anxiety-like behavior during withdrawal after chronic ethanol exposure\(^24\). Sprague-Dawley adult rats consume less alcohol as compared to adult P rats, suggesting that these changes can occur regardless of genetic manipulation and further suggesting that both epigenetic changes and environmental influences such as age of first alcohol exposure contribute to the development of alcohol use disorders.

pCREB activates the downstream signaling molecules CBP and p300, which possess HAT activity\(^{29,30}\). Previously, we reported reduced CBP levels in the CeA and MeA and increased anxiety-like behavior in adult rats during ethanol withdrawal after chronic exposure\(^24\). Other studies have shown that p300 conditional knock-out mice exhibited reduced histone H3 acetylation in the perirhinal cortex\(^35\). The decrease in CBP and p300 in the present study corresponds with the decreased global H3K9 acetylation and increased HDAC2 expression reported in the CeA and MeA of AIE adult rats\(^8\). We have also reported reduced levels of acetylated H3K9/14ac at the promoters of brain-derived neurotrophic factor (Bdnf) and activity regulated cytoskeleton-associated protein (Arc), genes crucial for synaptic plasticity, and decreased dendritic spine density in the CeA and MeA\(^8\). Taken together, these results suggest that dynamic interactions between decreased HATs and increased HDACs lead to a condensed chromatin conformation and aberrant synaptic plasticity in the amygdala at adulthood following adolescent alcohol exposure that may be involved in adult psychopathology.

In a previous study, we showed that exposure to an acute challenge of ethanol (2 g/kg) in adulthood normalized AIE-induced anxiety-like behaviors\(^8\). Here, we show that acute ethanol challenge also rescues the mRNA expression deficits of Creb1, Cbp, and p300 in the amygdala. This is mirrored by a decrease in activating H3K9/14ac marks at specific promoter sites of these genes in the amygdala of AIE adult rats that returns to control levels after an acute ethanol challenge. Based on these data we postulate that the long-lasting AIE-induced neuroadaptations in the amygdala contribute to the phenotypes of increased anxiety and alcohol consumption\(^8\), and that acute exposure to ethanol normalizes these neuroadaptations by increasing H3K9/14ac. This opens the chromatin and allows for increased RNA polymerase binding and subsequent increased transcription of Creb1, Cbp and p300. In AIS-treated animals, we show increases in H3K9/14ac which is consistent with previous studies\(^{31,36}\) that demonstrate that acute ethanol exposure in ethanol naïve adult animals leads to open chromatin in the amygdala via increases in CBP and increased global H3K9ac. Our study expands on this previous data and shows that p300 and Cbp are increased after acute ethanol injection in AIS animals due to increases in H3K9/14ac levels in their promoters, suggesting epigenetic regulation of these signaling molecules. Further, the current study suggests that acute ethanol after AIE increases H3K9/14ac at the promoter regions of Cbp, Creb1 and p300 which rescues decreases in H3K9/14ac and decreased mRNA levels produced by AIE in adulthood. This is consistent with the rescue of anxiety-like behavior after an acute ethanol injection in AIE adult rats\(^8\). These findings are particularly interesting in the context of the theory that affective states such as anxiety drive increased alcohol consumption, suggesting the possibility that AIE rats may consume alcohol to restore Cbp, Creb1 and p300 levels in order to prevent the negative affective states\(^{13,14}\). As mentioned above, the decrease in histone acetylation may result from globally increased HDACs\(^6\) and decreased HATs as observed here in the amygdala in adulthood after AIE. Our results raise the possibility that epigenetic enzymes such as HDAC2 and CBP/p300 change the chromatin and affects neuroadaptations of chronic alcohol consumption, suggesting that both epigenetic changes and environmental influences such as alcohol consumption are involved in the development of alcohol use disorders.

In conclusion, AIE causes lasting changes to the CREB signaling pathway in the amygdala, including decreased CREB, pCREB, CBP and p300 protein levels. This is paralleled by decreased Creb1, Cbp, and p300 mRNA in the amygdala that returns to control like levels after an acute challenge of ethanol in adulthood, likely via alterations in H3K9/14ac levels at the respective promoter regions. Collectively, the decreased levels of signaling molecules in the critical CREB pathway seen in this study, along with the increased anxiety and alcohol preference previously reported in AIE adult animals\(^8\), add to the growing body of evidence that CREB/pCREB/CBP/p300 levels in the amygdala are negatively correlated with anxiety-like and alcohol-drinking behaviors. These novel results suggest that adolescent alcohol causes lasting modifications to the epigenetic dynamics controlling the expression of CREB and related signaling molecules in brain regions crucial for emotion and anxiety. These results may lead to the development of new interventions targeting the genome of the alcohol-drinking population who begin alcohol consumption in adolescence.

**Methods**

**Animals and adolescent intermittent ethanol (AIE) exposure.** Sprague-Dawley (SD) dams with pups or timed-pregnant female rats were purchased from Harlan Laboratories (Indianapolis, IN, USA) and housed under a 12:12-h light/dark cycle with ad libitum access to drinking water and food. All experimental protocols strictly adhered to the NIH guidelines for the Care and Use of Laboratory Animals and were approved by the University of Illinois at Chicago Animal Care and Use Committee. Male pups were weaned at post-natal day (PND) 21 and were group-housed (2–3 rats/cage) with ad libitum access to water and food maintained. Rats were randomly assigned to receive either adolescent intermittent ethanol (AIE) or adolescent intermittent saline (AIS) treatment. Adolescent male rats received one injection of ethanol (2 g/kg, 20% w/v) or volume-matched n-saline per day via intraperitoneal (i.p.) injection for two consecutive days, followed by 2 days without ethanol treatment for a total of 8 injections of ethanol during PND 28–41. Rats matured without further ethanol exposure until PND 92–102. This schedule of ethanol exposure has been used by our lab and other investigators previously\(^{32,33,39,40}\).
Baseline brain tissue collection. On PND 92, AIS and AIE animals were anesthetized (either with 50 mg/kg of i.p. pentobarbital or inhaled isoflurane) and decapitated, and brain tissues (amygdala) were dissected on ice and quickly frozen. Some rats were perfused with normal saline followed by 4% paraformaldehyde (PFA) solution prepared in 0.1 M phosphate buffer (PB; pH 7.4). Brains were isolated and post-fixed overnight in PFA and soaked in graded sucrose solution (10–20–30%) prepared in PB. All brains were frozen and kept at −80 °C until further use for either biochemical studies or immunohistochemistry.

RNA isolation and Real-Time Quantitative PCR. Total RNA was extracted from amygdalar tissue using the Qiagen miRNAeasy protocol and DNase Kit (Qiagen, Venlo, NED) by homogenizing frozen tissue on ice using phenol, guanidine isothiocyanate and chloroform. Samples were then treated with RNase-free DNase, and RNA was eluted in RNase-free water. RNA was reverse transcribed using the GeneAmp RNA PCR Core Kit (Life Technologies). Aliquots from each cDNA were amplified by Real-Time PCR using either a Mx3000p qPCR system (Agilent Technologies, Santa Clara, CA, USA) and SYBR Green master mix (Fermentas, Glen Burnie, MD, USA) or a CFX Connect qPCR system with IQ SYBR SuperMix (BioRad, Hercules, CA, USA). Gene of interest expression for Creb1, Cbp, and p300 mRNA was examined and glyceraldehyde-3-phosphate dehydrogenase (Gapdh) or hypoxanthine-guanine phosphoribosyltransferase 1 (Hprt1) mRNA was used as a reference gene. PCR conditions for all primers were 30 s at 95 °C; 30 s at 58 °C and 30 s at 72 °C for 40 cycles. Primer sequences are located in Table 1. mRNA data analysis was performed by subtracting the average Ct (crossing threshold) of the reference gene from the gene of interest Ct for each sample. Relative expression levels were then determined using the 2−ΔΔCt method to acquire individual fold change41. Data is presented as average fold change relative to AIS control animals.

Immunohistochemistry. Gold immunolabeling was performed as described previously by our laboratory24,42. Coronal brain sections (20 μm) were cut and washed in phosphate buffered saline (PBS), and incubated in RPMI 1640 medium containing l-glutamine (Invitrogen, Carlsbad, CA, USA) for 30 min. Sections were incubated with 10% normal goat serum (NGS) diluted in PBS containing 0.25% Triton X-100 (PBST) for 30 min at room temperature. After blocking the sections with 1% bovine serum albumin (BSA) in PBST, they were incubated with antibodies against CREB (1:500 dilution; Millipore, Billerica, MA, USA, catalog number 06-863), pCREB (1:500; Millipore 06-519), CBP (1:200; Santa Cruz Biotechnology, Dallas, TX, USA, catalog number sc-583) or p300 (1:200; Santa Cruz sc-585) diluted in 1% BSA prepared in PBST for 18 hrs at room temperature. Sections were then washed with PBS followed by 1% BSA in PBS and incubation with gold particle (1.4 nm)-conjugated anti-rabbit secondary antibody (1:200 dilution in 1% BSA in PBS; Nanoprobes, Inc., Yaphank, NY, USA) for 1 hr at room temperature. Gold particles were silver enhanced (Ted Pella Inc., Redding, CA, USA) for 12 to 20 minutes and washed. Quantification of the immunolabeled gold particles was performed by the computerized Image Analyzer Software (Loats Associates, Westminster, MD, USA). Multiple brain sections were taken for each animal in order to assure bregma-matching. A total of nine object fields from 3 amygdala sections from distinct slices were counted for each animal and each amygdaloid structure (CeA, MeA, BLA). Results are represented as the number of immuno-gold particles per 100 μm² area.

Acute ethanol challenge in adulthood. A subset of AIS and AIE animals were raised and exposed to adolescent alcohol or saline exposure as described above. Additionally, these animals were exposed to an acute challenge either ethanol (2 g/kg; i.p.) or volume-matched n-saline at PND 101-102 to generate 4 total groups (AIS + Saline, AIS + EtOH, AIE + Saline, & AIE + EtOH) as described previously43. One hr after acute ethanol or saline challenge, rats were given anesthesia then sacrificed for the dissection of the amygdala to be used for mRNA analysis (as described above) or chromatin immunoprecipitation (ChIP) assay.

Chromatin immunoprecipitation (ChIP) assay. The amygdala of animals subjected to acute challenge were also evaluated for acetylated histone 3 lysine 9 and 14 (H3K9/14ac) occupancy at the promoter regions of Creb1, Cbp, and p300 using ChIP, as described previously89. Amygdala tissue was dissected and fixed in 1% methanol-free formaldehyde at 37 °C for 5 min and then quenched with 1 M glycerol before homogenization in lysis buffer (1% (v/v) SDS, 10 mM EDTA, 50 mM Tris-HCl pH 8.0). Homogenate was then sonicated using the Covaris ME220 (Covaris, Woburn, MA, USA) to achieve sheared DNA fragments of 200–500 base pairs which was then clarified using centrifugation (17,000 × g for 10 min, 4 °C) to obtain a chromatin fraction. Aliquots of sonicated chromatin were removed as input for normalization in downstream analysis and the remaining sonicated chromatin was incubated with an antibody to H3K9/14ac (Millipore 06-599; 5 μg/sample) overnight at 4 °C. Magnetic Protein A Dynabeads (ThermoFisher Scientific, Waltham, MA, USA) was then added to chromatin samples and rotated for 1 hr 30 min at 4 °C. Chromatin was then washed five times using washing buffer (0.01% SDS, 1.1% Triton X-100, 1.2 mM EDTA, 16.7 mM Tris-HCl pH 8.0, 167 mM NaCl) prior to DNA purification using Chelex-100 resin (10% (w/v), BioRad) by boiling at 95 °C for 10 min. The input aliquot was precipitated using 100% ethanol, washed once with 75% ethanol then purified by using (10% (w/v) Chelex-100 resin at 95 °C for 10 min45. Purified DNA was analyzed by quantitative PCR using primers directed to specific sites in the Creb1, Cbp, and p300 promoter regions (Table 1). The data was analyzed using the ΔΔCt method, normalizing to input, and then data are expressed as fold change relative to AIS + Saline control rats.

Statistical analysis. Statistical analysis was conducted using the SigmaStat software suite (Systat Software Inc., San Jose, CA, USA). Differences between adult AIE and AIS groups at baseline were analyzed using Student's unpaired two-tailed t-test. For experiments involving analysis of acute ethanol challenge, two-way ANOVA (factors: AIE treatment, acute EtOH) followed by Tukey's post-hoc test for multiple comparisons was performed. p values of less than 0.05 were considered significant for all tests. Exact p values are reported except for cases in which p < 0.001.
Table 1. Primers sets used in this study for mRNA analysis and promoter analysis after chromatin immunoprecipitation.

| Primer name                        | Sequence (5’ to 3’)               |
|------------------------------------|-----------------------------------|
| Creb1 mRNA forward                 | AGAAGCAGCACGAAAGAGAG              |
| Creb1 mRNA reverse                 | CACCTGCACTGTGCTCTAAAA             |
| Cbp mRNA forward                   | TAATGGAGGTGGCAGTGGTGTAA           |
| Cbp mRNA reverse                   | GTCGCGGAGCCTGTTGTTTGTTG           |
| p300 mRNA forward                  | AAACACCAGACGAGAGAAGTGGGGA         |
| p300 mRNA reverse                  | TCCATGTTGGCGTAGATTTTTGGA          |
| Hprt1 mRNA forward                 | TCCTCAGACCGGTTTCCCCGG             |
| Hprt1 mRNA reverse                 | TCATACATCAATCTACAGAGGCTGG         |
| Gadph mRNA forward                 | ACAGATGTTGTAAGGTTGGTGGTA          |
| Gadph mRNA reverse                 | AGCCTCCCATTCCTCGACT               |

Sample sizes were not pre-calculated but resemble those commonly used in the field and used in previous studies. Sample sizes are listed in figure legends as n values and are indicative of the number of animals per group in each experiment. All mRNA and ChIP PCR experiments were calculated from technical triplicates or quadruplicates.

Data availability. All processed data are available in the manuscript. Raw data are available from the corresponding author on reasonable request.

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**Author Contributions**

S.C.P. conceived ideas for the studies. H.Z. and S.C.P. designed and planned all experiments. H.Z. and E.J.K. generated experimental animals. H.Z., E.J.K., J.P.B., D.M.K. and T.T. performed various experiments, generated and analyzed data. All authors participated in manuscript writing and approved the final manuscript.

**Additional Information**

**Competing Interests:** All authors report no competing interests except SCP reports that a US patent application entitled “Histone acetyltransferase activators and histone deacetylase inhibitors in the treatment of alcoholism” (serial number 60/848237 filed on September 29th, 2006) is currently pending.

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