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Targeting G-quadruplex DNA as cognitive function therapy for ATR-X syndrome

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Abstract

Alpha-thalassemia X-linked intellectual disability (ATR-X) syndrome is caused by mutations in ATRX, which encodes a chromatin-remodeling protein. Genome-wide analyses in mouse and human cells indicate that ATRX tends to bind G-rich sequences with high potential to form G-quadruplexes. Here, we report that Atrx mutation induces aberrant upregulation of Xlr3b expression in mouse brain, an outcome associated with neuronal pathogenesis displayed by ATR-X model mice. We show that ATRX normally binds to G-quadruplexes in CpG islands of the imprinted Xlr3b gene, regulating its expression by recruiting DNA methyltransferases. Xlr3b binds dendritic mRNAs, and its overexpression inhibits dendritic transport of CaMKIIα mRNA, promoting synaptic dysfunction. Notably, treatment with 5-ALA, which is converted into G-quadruplex-binding metabolites, reduces RNA polymerase II recruitment and represses Xlr3b transcription in ATR-X model mice. 5-ALA treatment also rescues decreased synaptic plasticity and cognitive deficits seen in ATR-X model mice. Our findings suggest a potential therapeutic strategy to target G-quadruplexes and decrease cognitive impairment associated with ATR-X syndrome.
Introduction

ATR-X syndrome (OMIM Entry #301040) a severe intellectual disability, is caused by ATRX mutations \(^{1-3}\). \textit{ATRX} encodes the switch/sucrose nonfermentable (SWI/SNF)-like chromatin remodeling protein ATRX, which contains two signature motifs. One is a plant homeodomain (PHD) designated the ATRX-DNMT3-DNMT3L (ADD) domain, which binds histone H3 tails, specifically at H3K4me0K9me2/3 \(^{4-6}\). The other includes seven helicase subdomains that confer ATPase activity \(^7,8\).

Genome-wide analysis combining chromatin immunoprecipitation with next-generation sequencing (ChIP-seq) in both primary human erythroid cells and mouse embryonic stem cells (ESCs) shows ATRX enrichment at G-rich variable number tandem repeats (VNTRs), some of which form non-B DNA structures, including G-quadruplexes \(^9\). ATRX functions as a part of a histone chaperone complex that deposits the histone variant H3.3 onto pericentromeric heterochromatin and telomeres collaborating with death domain-associated protein (DAXX) in HeLa cells \(^{10}\) and murine ESCs \(^{11,12}\). ATRX/H3.3 co-localization also occurs on the DNA-methylated allele of many imprinted genes and is associated with differentially methylated regions (DMRs) in mouse ESCs \(^{13}\). Some imprinted genes show upregulated expression in forebrain of \textit{Atrx} conditional knockout (cKO) mice \(^{14,15}\), suggesting that ATRX silences the active allele. Moreover, expression of the autism-related gene \textit{Nlgn4} significantly decreases in forebrain of \textit{Atrx} cKO mice \(^{16}\). Imprinting loss in neurons leads to various mental retardation syndromes, including Prader-Willi and Angelman syndromes \(^{17}\). However, the relationship between abnormal expression and cognitive dysfunction in ATR-X syndrome remains unclear.
Results

ATRX regulates Xlr3b expression in mouse brain

Atrx\textsuperscript{ΔE2} mice, which are engineered to lack Atrx exon 2, show cognitive defects, among other phenotypes \textsuperscript{18}, and express a mutant protein that corresponds to a variant with an Arg37Stop (R37X) mutation in exon 2 seen in human ATR-X syndrome \textsuperscript{19-21}. Moreover, Atrx\textsuperscript{ΔE2} mice show 80% reduction in ATRX protein levels \textsuperscript{18,19}, similar to outcomes seen in 27 individuals with ATR-X syndrome \textsuperscript{8}. We employed DNA microarrays to assess transcriptional profiles at post-natal day (P) 90 in hippocampus of wild-type (WT) and Atrx\textsuperscript{ΔE2} mice. To identify differentially-expressed genes, we used an algorithm combining false discovery rate (FDR) and fold-change in expression, and identified 31 genes (8 upregulated and 23 downregulated) in WT versus Atrx\textsuperscript{ΔE2} samples. Among them was Atrx itself, which was downregulated in Atrx\textsuperscript{ΔE2} mice (Supplementary Table 1 for list of genes with an FDR < 0.05 and a log2 fold-change of > 0.5 or < −0.5). Among genes markedly upregulated in Atrx\textsuperscript{ΔE2} mice were a member of the lymphocyte regulated (Xlr) gene family, Xlr3a and the imprinted gene Xlr3b \textsuperscript{22-24} (Fig. 1a, FDR < 0.05 and log2 fold-change of > 0.5). Xlr3a and Xlr3b genes show 94% protein similarity, and DNA microarrays are limited in their ability to distinguish related factors. Thus, we carried out reverse transcription-PCR (RT-PCR) analysis with a common forward primer and subtype-specific reverse primers (Supplementary Fig. 1a). Primer efficiency was confirmed by amplifying respective cDNAs (Supplementary Fig. 1b), and distinct Xlr3a and Xlr3b amplicons were detected on gels (Supplementary Fig. 1c, lanes 1 and 2). Then, using these primers, we detected Xlr3b, but not Xlr3a mRNA, in prefrontal cortex, hippocampus, hypothalamus and cerebellum of WT and Atrx\textsuperscript{ΔE2} mice (Supplementary Fig. 1c). Interestingly, quantitative RT-PCR (RT-qPCR) indicated Xlr3b transcript upregulation in some brain areas, including hippocampus, of Atrx\textsuperscript{ΔE2} mice, but not in peripheral tissues (Fig. 1b).

Next, in order to investigate changes in Xlr3 protein expression, we generated an Xlr3 antibody. To confirm its specificity, we performed Xlr3 knockout (KO)-validation using Clustered Regularly Interspaced Short Palindromic Repeats (CRISPR)/Cas9 in cultured hippocampal neurons. To do so, we transduced cultured hippocampal neurons with lentivirus harboring U6gRNA-Cas9-2A-GFP plus guide Xlr3 sgRNA (Xlr3-Cas9-GFP) or control lentivirus expressing GFP. Microscopy analysis revealed loss of Xlr3 immunoreactivity in MAP2-positive neurons infected with Xlr3-Cas9-GFP but not control GFP virus (Supplementary Fig. 1d). We also confirmed Xlr3b KO efficacy by RT-qPCR in cultured hippocampal neurons (Supplementary Fig. 1e).
Western blot analysis with the Xlr3 antibody also showed significantly decreased protein expression in KO relative to WT cultured hippocampal neurons (Supplementary Fig. 1f), confirming antibody specificity. Next, we used the Xlr3 antibody to examine Xlr3 protein expression in Atrx\textsuperscript{AE2} mouse prefrontal cortex, hippocampus, hypothalamus and cerebellum tissues by western blot, and observed significant increases in Xlr3 protein levels in brain of Atrx\textsuperscript{AE2} relative to WT mice (Fig. 1c).

CpG island (CGI) methylation by DNA methyltransferases (DMNTs) generally represses transcription\textsuperscript{25}, and cKO mice lacking \textit{Dnmt1} or both \textit{Dnmt1} and \textit{Dnmt3a} in forebrain neurons show increased \textit{Xlr3b} expression\textsuperscript{26, 27}. Although ATRX lacks a canonical DNA methyltransferase motif, human \textit{ATRX} mutations are associated with altered DNA methylation patterns at various genomic loci\textsuperscript{28}. Moreover, \textit{Xlr3} gene CGIs display similar sequences, and a sequence potentially forming a G-quadruplex structure is specific to \textit{Xlr3b} (Supplementary Fig. 2a). Thus, we asked whether ATRX recruits DNMTs to methylate \textit{Xlr3b} CGIs. To do so, we employed bisulfite sequencing to compare DNA methylation levels in \textit{Xlr3b} CGIs of whole brain samples from WT or Atrx\textsuperscript{AE2} mice. Relative to WT mice, Atrx\textsuperscript{AE2} mice showed significantly decreased DNA methylation at \textit{Xlr3b} CGIs, while we observed no difference in DNA methylation levels in CGIs of other Xlr3 subfamily genes between genotypes (Fig. 1d, Supplementary Fig. 2b). Circular dichroism (CD) spectra of a synthetic oligodeoxynucleotide containing sequences potentially forming a G-quadruplex in \textit{Xlr3b} CGIs (Xlr3b-ODN) showed a spectrum characteristic of parallel G-quadruplexes in 10 and 100 mM KCl, with maximum absorbance at 265 nm and minimum at 240 nm (Supplementary Fig. 3a). This conformational change in physiological conditions (100 mM KCl) was reflected by decreased DNA mobility in gel shift assays performed using native polyacrylamide gel electrophoresis (PAGE) gels (Supplementary Fig. 3b). We then undertook dimethyl sulphate (DMS) footprinting of Xlr3b-ODN to assess formation of an intramolecular parallel G-quadruplex consisting of a core of three stacked G-quartets and three loops (Fig. 1e, Supplementary Fig. 3c). This analysis confirmed that \textit{Xlr3b} CGI sequences can form parallel G-quadruplex structures.

To assess how ATRX regulates \textit{Xlr3b}, we performed luciferase reporter assays. To do so, we first cloned three different \textit{Xlr3b} genomic sequences into a pGL3 luciferase reporter vector. They included: a 2.0 kb fragment (designated pGL3-2K) upstream of the \textit{Xlr3b} ATG initiation codon; a pGL3-2K deletion mutant lacking G-quadruplex-forming sequences (pGL3-2K\textasciitors{Δ}G4); and a 1 kb \textit{Xlr3b} upstream sequence (pGL3-1K\textasciitors{Δ}CGI) that lacks CGIs. Mouse neuroblastoma Neuro-2a cells transfected
with pGL3-2K showed significantly reduced luciferase activity when co-transfected
with ATRX; conversely, luciferase activity relative to controls increased in cells
co-transfected with ATRX shRNA (Fig. 1f). However, cells transfected with either
pGL3-2KΔG4 or pGL3-1KΔCGI showed activity comparable to cells co-transfected
with ATRX or ATRX shRNA, suggesting that ATRX regulates Xlr3b through the
G-quadruplex structure (Fig. 1f). We confirmed ATRX shRNA knockdown efficacy by
immunoblotting for endogenous ATRX protein in Neuro-2a cells (Supplementary Fig
4a). Moreover, Neuro-2a cells co-transfected with pGL3-2K plus DNMTs (DNMT1 or
DNMT3A) showed significantly decreased reporter activity relative to respective
controls (Fig. 1g). However, cells co-transfected with pGL3-2K plus DAXX or H3.3
showed comparable reporter activity. Importantly, cells co-transfected with
pGL3-2KΔG4 plus DNMTs showed no change in reporter activity relative to
pGL3-2KΔG4 alone. Finally, we methylated pGL3-2K, pGL3-2KΔG4 and
pGL3-1KΔCGI constructs using CpG DNA methyltransferase M.SssI and assessed the
luciferase activity. We observed decreased luciferase activity in methylated pGL3-2K
and pGL3-2KΔG4, not in methylated pGL3-1KΔCGI, supporting the idea that CGI
methylation inhibits Xlr3b expression (Fig. 1g).

To determine whether ATRX and interacting factors target Xlr3b CGIs in mouse
hippocampus, we performed ChIP-qPCR with an ATRX antibody. We confirmed that
hippocampal samples from AtrxAE2 mice contained detectable levels of ATRX protein:
those lysates showed an 80% reduction in ATRX protein levels relative to lysates from
WT mice, and the ATRX antibody recognized both WT and mutant ATRX protein as
described in 18, 19 (Supplementary Fig. 4b). Co-immunoprecipitation in hippocampal
lysates revealed that ATRX interacts with DNMT1, DNMT3A, DAXX and H3.3 from
both WT and AtrxAE2 mice, and relatively low amounts of DNMT1, DNMT3A, H3.3
and DAXX were detected in immunoprecipitated samples of AtrxAE2 mice
(Supplementary Fig. 4c). We detected substantial ATRX enrichment at Xlr3b CGIs
containing G-quadruplexes based on ChIP analysis with primers targeting Xlr3b region
R1 in chromatin isolated from WT mouse hippocampus, while ATRX interaction with
chromatin at this site was greatly decreased in AtrxAE2 hippocampus (Fig. 1h). Levels of
DNMT1, DNMT3A, DAXX and H3.3 also significantly decreased in AtrxAE2 relative to
WT chromatin in the R1 region. Also, quantitative profiling of ATRX, DNMTs, DAXX
and H3.3 across an unrelated R2 region revealed small peaks, demonstrating enrichment
of these proteins at Xlr3b CGIs (Fig. 1h). These observations suggest overall that
ATRX binds to parallel G-quadruplexes in Xlr3b CGIs along with DNMTs, DAXX and
H3.3, where it regulates Xlr3b gene expression.
Neuronal RNA granules contain Xlr3b

We next undertook a proteomic screen to search for Xlr3 interaction partners by performing liquid chromatography-tandem mass spectrometry (LC-MS/MS) of proteins pulled down from lysates of P60 WT mouse brain with an Xlr3 antibody. LC-MS/MS analysis revealed an endogenous Xlr3 complex containing multiple components, including actin/myosin-related proteins, dynein motor complex (DYNLL, DYNLRB, DYNLT and DYNC1L12) proteins, ribonucleoproteins (hnRNP A/B and hnRNP D), 40S ribosomal proteins (RPS10 and RPS25), and others (Fig. 2a, Supplementary Table 2). We then confirmed association of purified GST-Xlr3b protein with hnRNP A/B, hnRNP D, DYNLL and DYNC1L12 by in vitro pull-down assays in mouse hippocampal lysates (Fig. 2b). Purified Xlr3b protein did not bind F-actin, and its presence did not alter actin polymerization (Supplementary Fig. 5a, 5b). To confirm interactions in vivo, we performed immunoprecipitation of mouse hippocampal extracts with an Xlr3 antibody followed by immunoblotting with hnRNP A/B, hnRNP D, DYNLL, or DYNC1L12 antibodies and observed co-precipitation of all with Xlr3 (Fig. 2c).

To determine Xlr3 cellular localization, we undertook immunofluorescence of mature MAP2-positive cultured neurons with an Xlr3 antibody. Xlr3 was localized to nuclei and perinuclear areas and seen in granules in MAP2-positive dendrites and in synaptic synaptophysin-positive puncta (Fig. 2d). Xlr3-positive structures of this type were also observed in MAP2-positive dendrites in mouse hippocampal CA1 analyzed at P60 (Fig. 2e). In dendrites, Xlr3-positive puncta co-localized with hnRNP A/B and cytotoxic granule-associated RNA binding protein 1 (TIA1), both found in RNA granules 29, and with SYTO14-visualized RNA. Xlr3-positive granules did not co-localize with DCP1a, a marker of RNA-processing bodies 29 (Fig. 2f).

Xlr3b exhibits a predicted nuclear localization signal (NLS) at amino acid (AA) residues 2-11; residues 74-200 also constitute a conserved Cor1/Xlr/Xmr region (Cor1 domain). To assess function of those sequences, we transfected Neuro-2a cells with FLAG-tagged Xlr3b or similarly-tagged deletion mutants shown in Figure 2g and analyzed their location by confocal microscopy (Fig. 2g). Based on FLAG expression, full length (FL)-Xlr3b significantly co-localized with TIA1 in nuclear regions compared to that of non-transfected cells. FL-Xlr3b also accumulated with hnRNP A/B in nuclei, but not with DCP1a (Fig. 2g, Supplementary Fig. 6a, 6b). Cells transfected with constructs lacking AA 2-11 (ΔNLS) or 74-123 (Δ74-123) showed partial loss of Xlr3b nuclear localization, but localization of the interacting proteins TIA1 and hnRNP A/B
was unchanged. These localization patterns were similar to those seen in FL-Xlr3b-transfected cells (Fig. 2g, Supplementary Fig. 6a, 6b). Deletion of AA 124-200 (Δ124-200) caused loss of nuclear Xlr3b expression and localization of tagged protein in cytoplasmic ubiquitin-positive inclusion-like structures, in which DCP1a, TIA1 and hnRNP A/B were also colocalized. In addition, the Δ158-170 mutant was distributed similarly to the Δ124-200 mutant (Fig. 2g, Supplementary Fig. 6a, 6b and 6c). Importantly, nuclear localization of TIA1 and hnRNP A/B in cells transfected with Δ124-200 or Δ158-170 constructs was very similar to patterns observed in non-transfected cells (Fig. 2g, Supplementary Fig. 6a).

To confirm interactions of Xlr3b AA 158-170 with RNA binding proteins (RBPs), we used Xlr3b inhibitory peptide (XIP), a 29-AA peptide containing Xlr3b AA 158-170 plus the cell-permeable antennapedia homeodomain (ANTP) peptide (Supplementary Fig. 7a). When FL-Xlr3b-transfected cells were treated with XIP (1μM for 4 h), TIA1 did not accumulate in the nucleus (Supplementary Fig. 7b). We confirmed that XIP, but not the ANTP control peptide, bound to RBPs, TIA1 and hnRNP A/B using pull-down assays of mouse hippocampal lysates with an ANTP antibody (Supplementary Fig. 7c). Immunoprecipitation confirmed that FLAG-tagged FL, Δ74-123, Δ124-200 and Δ158-170 mutants were immunoprecipitated with a FLAG antibody. Immunoblotting with TIA1 and hnRNP A/B antibodies detected immunoreactive TIA1 and hnRNP A/B bands in samples transfected with FL and Δ74-123, but not in Δ124-200 and Δ158-170. Remarkably, interactions with TIA1 and hnRNP A/B were not seen in FL-Xlr3b-transfected cells treated with 1μM XIP for 4 h before immunoprecipitation (Fig. 2h), suggesting that Xlr3b residues 158-170 are critical for RBP binding. Ubiquitination of Xlr3b Δ124-200 and Δ158-170 may prevent binding to RBPs, regardless of co-localization 30. Thus, to assess potential Δ124-200 and Δ158-170 ubiquitination, we performed immunoprecipitation of lysates of cells transfected with FLAG-tagged FL, Δ74-123, Δ124-200 and Δ158-170 mutants with a FLAG antibody followed by immunoblotting with a ubiquitin antibody. We confirmed ubiquitination in immunoprecipitants of constructs Δ124-200 and Δ158-170, but ubiquitination was little seen in FL or Δ74-123 proteins (Supplementary Fig. 6d).

Phylogenetic analysis identified the FAM9 gene family as human Xlr orthologues, and FAM9 proteins are predicted to have a Cor1 domain 32. BLAST searches indicated that Xlr3b residues 158-170 are highly conserved in Cor1 domains of FAM9 family proteins, including FAM9A and FAM9B (Supplementary Fig. 6e). We observed nuclear localization of FLAG-tagged FAM9A overexpressed in Neuro-2a cells, while a similar construct lacking residues analogous to Xlr3b 158-170 was found in
cytoplasmic inclusion body-like structures in a pattern similar to Xlr3b Δ158-170 transfected cells (Supplementary Fig. 6e).

**Aberrant Xlr3b expression in neurons decreases synaptic plasticity by inhibiting transport of dendritic CaMKIIα mRNA**

In neurons, RNA granules are rapidly transported to dendrites, a process regulated by motor proteins. Some RBPs including hnRNPA/B recognize a specific *cis*-acting element termed the hnRNP A2 response element (A2RE) in mRNA 3' untranslated regions (3' UTRs); these include Ca\(^{2+}\)/calmodulin-dependent protein kinase IIα (CaMKIIα), activity-regulated cytoskeleton-associated protein (Arc) and brain-derived neurotrophic factor (BDNF). Because Xlr3b interacts with the dynein motor complex and RBPs, we hypothesized that Xlr3b regulates dendritic mRNA transport via a dynein-mediated transport mechanism.

To test this hypothesis, we undertook RNA immunoprecipitation in mouse hippocampal lysates to determine whether an endogenous Xlr3 complex associates with dendritic mRNAs. After immunoprecipitation with an Xlr3 antibody, we isolated total RNA from precipitates and analyzed it by qPCR with primers specific to CaMKIIα, Arc and BDNF mRNA 3' UTRs. Immunoprecipitates contained all three mRNAs, and XIP treatment (1μM for 2h) of cell lysates significantly blocked these interactions (Fig. 3a).

Atrx\(^{AE2}\) mice show cognitive impairment and CaMKIIα dysfunction. Thus we focused on dendritic CaMKIIα mRNA. To determine whether Xlr3b protein is co-transported into dendrites with CaMKIIα mRNA, we used dual-color time-lapse imaging with mCherry-Xlr3b and GFP-MS2-labeled CaMKIIα mRNA (GFP-CaMKIIα 3' UTR)\(^{34}\). mCherry-Xlr3b co-localized with the GFP-CaMKIIα 3' UTR, and both were co-transported to dendrites of cultured neurons (Fig. 3b). By contrast, co-localization of mCherry-Xlr3b mutants (Δ124-200 and Δ158-170) with GFP-CaMKIIα 3' UTR immunofluorescence was rarely seen (Supplementary Fig. 8a). To analyze dynamics of dendritic CaMKIIα mRNA transport, we investigated movement of the GFP-CaMKIIα 3' UTR in proximal dendrites (20-100μm away from the cell body). Granules moved in anterograde (white arrowhead) and retrograde (yellow arrowhead) directions (Fig. 3c, top panel, left). Fig. 3c (top panel, right) shows a representative kymograph corresponding to 15-min of live imaging generated from a proximal dendrite that GFP-CaMKIIα 3' UTR (also see Supplementary Video 1). Imaging of GFP-CaMKIIα 3' UTR movement revealed that approximately half of the observed CaMKIIα mRNA granules were immobile, and the few mobile granules showed bidirectional movement in cultured WT neurons. The number of bidirectionally-mobile
CaMKIIα mRNA granules significantly decreased in cultured neurons of transgenic mice overexpressing Xlr3b driven by the neuron-specific Thy1.2 promoter (Thy1-Xlr3b TG mice; Supplementary Fig. 9a-9d), relative to effects seen in WT neurons. Moreover, in cultured neurons of AtrxΔE2 mice, the number of bidirectionally-mobile CaMKIIα mRNA granules significantly decreased. Xlr3b-shRNA transfection or XIP treatment antagonized decreases in dendritic CaMKIIα mRNA transport seen in AtrxΔE2 neurons, suggesting that Xlr3b acting through RBPs inhibits CaMKIIα mRNA transport (Fig. 3c, bottom). We confirmed efficacy of Xlr3b knockdown by immunoblotting of endogenous Xlr3 protein in cultured neurons (Supplementary Fig 9e). In addition, dynamics of CaMKIIα mRNA transport in distal dendrites (100-200μm from the cell body) was similar to that seen in proximal dendrites: bidirectionally-mobile CaMKIIα mRNA granules significantly decreased in Thy1-Xlr3b TG and AtrxΔE2 neurons relative to effects seen in WT neurons. Furthermore, Xlr3b-shRNA transfection or XIP treatment rescued decreases in dendritic CaMKIIα mRNA transport seen in AtrxΔE2 neurons (Supplementary Fig. 8b, Supplementary Video 2).

RNA granules localize beneath postsynaptic sites in dendrites, and CaMKIIα mRNA is located in dendrites and active postsynaptic regions. Indeed, the number of double-positive protein puncta containing the GFP-CaMKIIα 3′ UTR and the postsynaptic marker PSD95 significantly decreased in cultured neurons of Thy1-Xlr3b TG and AtrxΔE2 mice relative to the number seen in WT mice, and cultured neurons from AtrxΔE2 mice transfected with Xlr3b shRNA or treated with XIP showed a significant increase in the number of double-positive puncta (Fig. 3d). Next, we performed western blot analysis for CaMKIIα protein in synaptosomal membrane fractions, containing PSD95 isolated from cultured hippocampal neurons (Supplementary Fig. 9f). CaMKIIα protein was decreased in the synaptosomal membrane fractions from cultured neurons of Thy1-Xlr3b TG and AtrxΔE2 mice relative to that seen in WT mice. XIP treatment in AtrxΔE2 neurons significantly increased in the synaptosomal CaMKIIα protein levels without changes in both CaMKIIα and PSD95 levels in total cell lysate fractions between groups (Supplementary Fig. 9f).

Dendritic CaMKIIα mRNA translation is regulated by synaptic activity, and postsynaptic CaMKIIα protein regulates hippocampal long-term potentiation (LTP), which is critical for learning and memory. To assess a potential effect on LTP, we undertook electrophysiological analysis of brain slices from hippocampal CA1. Consistent with our previous study, we observed markedly reduced high frequency stimulation (HFS)-induced LTP in AtrxΔE2 mice, which was significantly rescued by a 2 h bath application of XIP to AtrxΔE2 mouse samples, but not following application of an
ANTP control peptide, suggesting that impaired LTP seen in AtrxΔE2 mice is in part Xlr3b-dependent (Fig. 3e). Notably, Thy1-Xlr3b TG mice also showed impaired hippocampal LTP (Supplementary Fig. 9g). We observed no significant difference in basic electrophysiological properties, among them input-output relations and paired-pulse ratio, between groups (Supplementary Fig. 9h). LTP induction, which strengthens synapses, results in CaMKII phosphorylation throughout the dendritic area, suggesting widespread CaMKII activation including in spines. In WT mice, levels of CaMKIIα phosphorylation significantly increased following hippocampal LTP induction. Consistent with LTP impairment, CaMKIIα phosphorylation did not increase following LTP induction in hippocampus of AtrxΔE2 and Thy1-Xlr3b TG mice. Notably, bath application of XIP, but not ANTP, significantly restored LTP-induced CaMKIIα phosphorylation in the AtrxΔE2 hippocampus (Supplementary Fig. 9i).

To assess whether Xlr3b overexpression affects cognitive function, we employed a novel object recognition task. In training trials, we observed no differences in the discrimination index using the same object between WT and Thy1-Xlr3b TG mice (data not shown). After a 24 h retention interval between trial and test sessions, Thy1-Xlr3b TG mice showed a significantly lower discrimination index for a novel object than did WT mice (Supplementary Fig. 9j). To assess contextual memory, we undertook a fear-conditioned passive avoidance task and observed no significant differences in latency to enter a dark room in the absence of foot shock between WT and Thy1-Xlr3b TG mice (data not shown). However, latency to enter the dark compartment was markedly decreased 1-week after foot shock in Thy1-Xlr3b relative to WT mice (Supplementary Fig. 9k). We then used a Y-maze to test spatial working memory and observed that Thy1-Xlr3b mice showed impairment based on the percentage of alternation behaviors relative to WT mice without a change in the total number of arm entries (Supplementary Fig. 9l).

5-ALA treatment reduces RNA polymerase II recruitment and represses Xlr3b transcription by modifying G-quadruplex structure.

G-quadruplex-binding small molecules such as porphyrin function as transcriptional repressors. Thus, we asked whether Xlr3b repression via porphyrin binding its G-quadruplex structure would antagonize synaptic dysfunction and cognitive deficits seen in AtrxΔE2 mice. To do so, we first assessed intracellular generation of porphyrins following administration of 5-aminolevulinic acid (5-ALA), which can be metabolized to porphyrins, protoporphyrin IX (PpIX) and hemin (Fig. 4a). We confirmed PpIX and hemin binding to Xlr3b-ODN G-quadruplexes using UV melting
analysis. The $\Delta T_m$ for PpIX and hemin was 2.0 and 2.3°C, respectively, indicating that
PpIX and hemin both moderately bind $Xlr3b$ G-quadruplexes structure. The $\Delta T_m$ for
TMPyP4, a well-known G-quadruplex binding ligand, was higher than 4°C
(Supplementary Fig. 10a). To determine whether 5-ALA treatment alters $Xlr3b$ gene
expression, we transfected Neuro-2a cells with the pGL3-2K reporter, treated cells with
varying concentrations of 5-ALA or TMPyP4, and assessed effects of $Xlr3b$ expression
based on luciferase assays. As expected, treatment with 5-ALA or TMPyP4 reduced
luciferase activity dose-dependently but had no effect on cells transfected with
pGL3-2KΔG4 (Fig. 4b, Supplementary Fig. 10b). Decreased luciferase activity in
5-ALA treated cells was efficiently rescued by co-treatment with the 5-ALA
dehydratase inhibitor succinylacetone (Fig. 4c). Next, we asked whether treatment with
G-quadruplex ligands would inhibit aberrant $Xlr3b$ gene expression seen in Atrx$^{AE2}$
mouse brain. To do so, we administered 5-ALA (p.o. daily from P30 to P90) or
TMPyP4 (i.p. twice weekly from P30 to P90). Either treatment significantly blocked
increases in hippocampal $Xlr3b$ mRNA and $Xlr3$ protein seen in vehicle-treated Atrx$^{AE2}$
mice (Fig. 4d and 4e, Supplementary Fig. 10c and 10d).
To investigate mechanisms underlying these effects, we undertook bisulfite
sequencing to compare DNA methylation levels. $Xlr3b$ CGIs in hippocampus and
prefrontal cortex of Atrx$^{AE2}$ mice showed significantly decreased DNA methylation
relative to WT mice, an effect unchanged by 5-ALA treatment (Fig. 4f, Supplementary
Fig. 10e). Next, we performed ChIP-qPCR of ATRX and RNA polymerase II (Pol II)
within the $Xlr3b$ gene, as G-quadruplex structures hinder Pol II passage $^{43, 44}$. The
prediction of quadruplex structures within the $Xlr3b$ gene was supported by the
G-quadruplex analysis tool QGRS Mapper $^{45}$ (Fig. 4g, top). In WT mouse hippocampus,
ATRX and Pol II accumulated at $Xlr3b$ CGIs containing G-quadruplexes, based on
analysis using primers targeting amplicon 1, but little signal was seen at amplicons 2, 3
and 4 (Fig. 4g, bottom). However, in Atrx$^{AE2}$ mouse, we detected an elevated Pol II
signals across all amplicons in the $Xlr3b$ gene body as well as CGIs, indicating
enhanced $Xlr3b$ transcription. Furthermore, elevated Pol II signals were significantly
attenuated following 5-ALA treatment (Fig. 4g, bottom). These results suggest that the
porphyrin metabolites PpIX and hemin, which are derived from 5-ALA, bind $Xlr3b$
G-quadruplex structures, partially reducing Pol II recruitment.

Treatment with G-quadruplex ligands counteracts cognitive deficits seen in
Atrx$^{AE2}$ mice.
Next, we asked whether 5-ALA treatment rescued decreases in dendritic CaMKIIα mRNA transport seen in Atrx^{AE2} neurons. Imaging of GFP-CaMKIIα 3’ UTR movement revealed that the decreased number of bidirectionally-mobile CaMKIIα mRNA granules was significantly rescued in 5-ALA treated Atrx^{AE2} neurons (7 days, 1μM) at proximal and distal dendrites (Fig. 5a, Supplementary Fig. 11a). The decreased number of double-positive protein puncta containing GFP-CaMKIIα 3’ UTR and PSD95 in Atrx^{AE2} neurons was also significantly rescued (Fig. 5b). Moreover, reduced HFS-induced LTP in hippocampus of vehicle-treated Atrx^{AE2} mice was significantly restored in 5-ALA-treated Atrx^{AE2} mice (p.o. daily from P30 to P90) without a change in basic electrophysiological properties (Fig. 5c, Supplementary Fig. 11b). Finally, 5-ALA treatment significantly blocked decreased CaMKIIα protein level at synaptosomal membrane fractions in cultured hippocampal neurons from Atrx^{AE2} mice, and restored LTP-induced CaMKIIα phosphorylation in hippocampus of Atrx^{AE2} mice (Supplementary Fig. 11c).

We next administered 5-ALA (p.o. daily from P30 to P90) or TMPyP4 (i.p. twice weekly from P30 to P90) to Atrx^{AE2} mice and subsequently assessed memory-related behaviors. We observed no differences in a novel object recognition task in the discrimination index using the same object in all groups in training trials (data not shown). After a 24h retention interval between trial and test sessions, Atrx^{AE2} mice showed a significantly lower discrimination index for a novel object than did WT mice. The discrimination index for the novel object in Atrx^{AE2} mice treated with 5-ALA or TMPyP4 was significantly higher than that seen in vehicle-treated groups (Fig. 5d, Supplementary Fig. 11d). In a fear-conditioned passive avoidance task, we observed no significant differences in latency to enter a dark room in the absence of foot shock in all groups (data not shown). However, latency to enter a dark compartment was markedly decreased 1-week after foot shock in Atrx^{AE2} relative to WT mice. 5-ALA or TMPyP4 administration also significantly rescued reduced latency time (Fig. 5e, Supplementary Fig. 11e). In a Y-maze test, Atrx^{AE2} mice showed impairment based on the percentage of alternation behaviors relative to WT mice without a change in the total number of arm entries. 5-ALA or TMPyP4 administration significantly increased the percentage of spontaneous alternation behaviors in Atrx^{AE2} mice (Fig. 5f, Supplementary Fig. 11f). Many patients with ATR-X syndrome exhibit abnormal social behaviors, especially shyness and social withdrawal 46. Interestingly, in interactions with WT mice, Atrx^{AE2} mice showed enhanced passivity, greater escape duration and decreased active social behaviors, such as following and sniffing. These changes were dramatically ameliorated in Atrx^{AE2} mice (Supplementary Fig. 11g).
Acute administration of a single 5-ALA dose (10mg/kg, p.o.) did not ameliorate impaired learning and memory-related behaviors in Atrx^{AE2} mice (data not shown).

To confirm that TMPyP4 was present in brain, we measured TMPyP4 fluorescence levels, as described in 47. Chronic intraperitoneal injection of TMPyP4 (i.p. twice weekly from P30 to P90) in mice increased fluorescence levels in some tissues, including brain, suggesting that treatment allows intracerebral transferability and enables TMPyP4 accumulation in brain tissue (Supplementary Fig. 11h). However, we stopped chronic TMPyP4 administration on day 60 due to toxicity, as evidenced by reduced body weight (Supplementary Fig. 11i). Unlike the case with TMPyP4, 5-ALA pharmacokinetics and biodistribution, including in brain, has been reported in humans 48 and rodents 49. We confirmed that 5-ALA crosses the blood brain barrier after oral administration. In mouse brain, 5-ALA (3mg/kg, p.o.) levels peaked at 30min and then decreased to basal levels by 24h after administration (Supplementary Fig. 11j).

To investigate expression of other genes underlying these effects, we performed DNA microarray analysis of RNA extracted from hippocampus of P90 WT, Atrx^{AE2}, or 5-ALA treated Atrx^{AE2} mice, and focused on 31 genes differentially expressed (8 upregulated and 23 downregulated) in WT versus Atrx^{AE2} samples using a threshold of FDR <0.05 and log2 fold-change of > 0.5 or < -0.5 (Supplementary Table 1). Rescued or not rescued genes were assessed using the difference between the average value in log2 global normalization. 5-ALA treatment of Atrx^{AE2} mice significantly blocked differential expression of 71.0% of genes including Xlr3b (Fig. 5g, Supplementary Table 3). Following analysis of mice treated with TMPyP4, we selected 138 genes differentially expressed (38 upregulated and 100 downregulated) in WT versus Atrx^{AE2} hippocampal samples using a threshold of log2 fold-change of > 1.0 or < -1.0. TMPyP4 treatment of Atrx^{AE2} mice blocked differential expression of 92.8% of genes including Xlr3b, leaving only 10 of 138 genes remained differentially expressed (Supplementary Table 4).
Discussion

Here, we make several critical observations relevant to activities of ATRX (Supplementary Fig. 12). (1) ATRX binds to parallel G-quadruplexes in Xlr3b CGIs together with DNMTs, DAXX and H3.3, where it regulates Xlr3b gene expression. (2) Cognitive deficits seen in AtrxΔE2 mice are accompanied by Xlr3b upregulation, which inhibits CaMKIIα mRNA transport in neuronal dendrites. (3) Abnormal neuronal phenotypes exhibited by AtrxΔE2 mice are rescued by treatment with 5-ALA, a G-quadruplex ligand that inhibits Xlr3b expression.

Among genes altered in AtrxΔE2 mouse brain, the imprinted gene Xlr3b is significantly overexpressed. ATRX is enriched with DNMTs at G-quadruplexes in Xlr3b CGIs. We observed that reduced ATRX levels at a given site were accompanied by reduced DNMT levels and substantial DNA demethylation, suggesting that ATRX regulates DNA methylation by DNMT recruitment to G-quadruplexes. Notably, forebrain-specific cKO of either DNMT1 or both DNMT1 and DNMT3A reportedly promotes increased Xlr3b expression in excitatory neurons 26, 27. Moreover, recent genome-wide analysis revealed preferential ATRX localization at the DNA-methylated allele of many imprinted DMRs in mouse ESCs and that ATRX-bound CGI sequences were methylated to a greater extent than that seen in other parts of the genome 13. ATRX deficiency also correlates with reduced H3.3 incorporation and with Pol II stalling at G-rich intragenic sites, indicating that ATRX influences Pol II-mediated transcription 16. Accordingly, we detected Pol II accumulation at Xlr3b CGIs harboring G-quadruplexes in WT hippocampus. We also found that cells in the AtrxΔE2 hippocampus showed enriched Pol II signals across the Xlr3b gene relative to cells from WT brain, and elevated Pol II signals in AtrxΔE2 hippocampus were significantly attenuated following 5-ALA treatment without changing DNA methylation levels. These results suggest that the porphyrin metabolites PpIX and hemin, which are derived from 5-ALA, bind G-quadruplex structures, partially inhibiting Pol II recruitment, a mechanism supported by biophysical analysis of G-quadruplexes in the HIF1A, KRAS, CMYB and CMYC genes 41. The G-quadruplex structure formed behind advancing Pol II may be recognized by other factors, including positive transcription elongation factor b 50. In this context, further studies are required to define dynamic transcriptional structures interacting with the region bearing G-quadruplex motifs and how they regulate ATRX-mediated DNA methylation and Pol II recruitment.

We also report a novel role for Xlr3b as a component of RNA granules containing RBPs, which inhibit CaMKIIα mRNA transport into neuronal dendrites. Xlr3b interacted with the dendritic mRNAs CaMKIIα, Arc and BDNF via a 13 AA Cor1
domain, which binds RBPs. This sequence is required for CaMKIIα mRNA transport, suggesting that RBP binding is also necessary for transport. Interestingly, that Xlr3b sequence is highly conserved in human FAM9 proteins, which also exhibit a Cor1 domain. Thus, binding ability may be common to proteins containing Cor1/Xlr/Xmr motifs. In addition, our fluorescence imaging and electrophysiological analysis supports the idea that Xlr3b negatively regulates synaptic plasticity by inhibiting dendritic CaMKIIα mRNA transport in Atrx^{ΔE2} mice. Mice engineered to lack the CaMKIIα mRNA 3′ UTR showed reduced levels of CaMKIIα transcripts at postsynaptic densities, reduced LTP, and impaired spatial memory. Dendritic transport and local translation of BDNF and Arc mRNA in synapses also enhances neuronal activity, strongly suggesting that Xlr3b-mediated dendritic mRNA transport is related to learning and memory functions. 39X^{pO} mice, a murine model of the female developmental disorder Turner syndrome, show aberrantly high Xlr3b expression in brain, an outcome associated with difficulties in performing a reversal learning test, relative to 40XX and 39X^{pO} mice, suggesting that Xlr3b governs mouse behaviors. However, mechanisms underlying these activities remain unknown. We also demonstrated that Thy1-Xlr3b TG mice show impaired LTP and memory deficits, indicating that aberrant neuronal Xlr3b expression partially affects dendritic mRNA transport and learning behaviors in Atrx^{ΔE2} mice.

We also show that administration of the G-quadruplex ligands 5-ALA or TMPyP4 significantly rescues neuronal phenotypes seen in Atrx^{ΔE2} mice by inhibiting Xlr3b transcription. However, recovery of cognitive deficits in Atrx^{ΔE2} mice following 5-ALA administration may be due in part to the binding of ligand to G-quadruplexes of other ATRX targets, although this activity remains unknown. G-quadruplex motifs broadly influence transcription and translation. In addition, ATRX can either repress or activate gene expression through G-quadruplexes. Although potential off-target effects remain to be investigated, our findings suggest a novel therapeutic strategy aimed at blocking Xlr3b expression through small molecule binding to G-quadruplex DNA in patients with ATR-X syndrome. Among candidate ligands, 5-ALA has been applied clinically with minimal risk and approved for use following intracranial tumor resection in Europe, Canada, and Japan, where it has been used as a photosensitizer in photodynamic diagnostics applied in neurosurgery. 5-ALA is also safe when administered chronically up to 200 mg per day in humans. Thus, its approval time would be short and the cost of clinical trials would be reduced, as pre-existing absorption, distribution, metabolism, excretion (ADME) and toxicity data is available. Finally, the risk of failure is reduced as data relevant to 5-ALA safety and
pharmacology is available.

In summary, our study indicates that G-quadruplex ligands regulate gene expression through a transcriptional mechanism involving Xlr3b and could be used to treat symptoms associated with ATR-X syndrome.

**Accession codes availability:** Microarray raw data are available at Gene Expression Omnibus (GEO); accession numbers, GSE103031 and GSE103032.

**Data Availability Statement:** Summary of all statistical data was shown in Supplementary Table 5. The other data that support the findings of this study are available from the corresponding authors on reasonable request.

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**Author contributions:** N.S., Y.Y., K.Y., M.O., and Y.L. performed experiments. K.K., H.T., N.O., T.E., H.S. and T.W. provided critical advice. N.S. and K.F. wrote the manuscript and designed the study. All authors discussed the results and commented on the manuscript.

**Competing interests:** The authors declare no competing financial interests.


Figure 1. ATRX regulates Xlr3b gene expression. a, Heat map summarizing expression data, and a list of genes exhibiting differential expression between WT and Atrx^ΔE2 mouse hippocampus at post-natal day (P) 90 (an FDR < 0.05 and a log2 fold-change of > 0.5). See also Supplementary Table 1 (a list of genes with an FDR < 0.05 and a log2 fold-change of > 0.5 or < −0.5). n = 7 mice each. b, Quantitative real-time RT-PCR showing Xlr3b mRNA expression in mouse lysates. **P < 0.01 by two-way ANOVA with Bonferroni’s post hoc test; n = 8-12 mice at P90. PC, (n=8 each); HP, (n=10 each); HT, (WT, n=10; Atrx^ΔE2 n=8); CE, (WT, n=9; Atrx^ΔE2, n=10); lung, liver, heart, and kidney, (n=12 each). PC, medial prefrontal cortex; HP, hippocampus; HT, hypothalamus; CE, cerebellum. c, (top) Representative immunoblot of P90 mouse brain lysates probed with indicated antibodies. (bottom) Densitometric analysis of Xlr3 normalized to β-tubulin (arbitrary units, A.U.). **P < 0.01, *P < 0.05 by two-way ANOVA with Bonferroni’s post hoc test; n = 4 mice each. d, Percentage of 5-methylcytosine (5-mC) as determined by bisulfite sequencing analysis in whole brain from male P90 WT or Atrx^ΔE2 mice. **P < 0.01 by two-sided unpaired t test. n = 12 clones each. (n = 3 mice each, 4 clones were sequenced per mouse). See also Supplementary Fig. 2. e, Xlr3b-ODN showing intramolecular parallel G-quadruplexes, which formed three separate G-quartets stacked 5′ to 3′ with three loops. f and g, Luciferase activity of Neuro-2a cells co-transfected with plasmids identified in left margin plus other indicated plasmids. Luciferase activity is presented relative to its activity in mock cells. **P < 0.01 by one-way ANOVA with Bonferroni’s post hoc test; n = 6 biologically independent samples. h, (top) Schematic diagram of Xlr3b: G-quadruplex-forming sequence (yellow), CGI (gray) and R1 and R2 regions. (bottom) Analysis of R1 and R2 regions in chromatin from hippocampus of Atrx^ΔE2 and WT mice by chromatin immunoprecipitation with indicated antibodies. Results are expressed as percent input. **P < 0.01, *P < 0.05 by two-sided unpaired t test. n = 6 mice at P90. Full-size scans of western blots shown in Supplementary Fig. 13.

Figure 2. Xlr3b is found in neuronal RNA granules. a, Pull-down assay and LC-MS/MS analysis of Xlr3-interacting proteins from P60 mouse brain lysates using an Xlr3 antibody. (left) Silver-stained SDS-PAGE gel of protein A sepharose column eluates with rabbit IgG (lane 1) or an Xlr3 antibody (lane 2) in mouse brain lysates. (right) Distribution of identified proteins in a shotgun proteomics experiment. b, Immunoblot analysis of GST affinity-column eluates using P60 mouse hippocampal
lysates. Aliquots of extracts (1% input) and eluates were subjected to SDS-PAGE and immunoblotted with indicated antibodies. The experiments were repeated three times with similar results. **c**, Immunoprecipitation (IP) assay with an Xlr3 antibody from P60 mouse hippocampal lysates. Precipitated proteins and inputs were immunoblotted with indicated antibodies. Aliquots of extracts (1% input) and eluates were subjected to SDS-PAGE. The experiments were repeated three times with similar results. **d**, Confocal images showing co-localization of Xlr3 (green) with synaptophysin (red) and MAP2 (blue) in primary mouse cultured neurons at day 21 in vitro. Arrowheads indicate synaptophysin-positive puncta. Scale bars: (top) 10μm, (bottom) 5μm. **e**, Confocal images show co-localization of Xlr3 (green) with MAP2 (magenta) in mouse hippocampal CA1. Images at bottom are high-magnification. Scale bars, 10μm. **f**, Confocal images showing co-localization of Xlr3 (magenta) with hnRNP A/B, TIA1, SYTO14 and DCP1a (green) in primary mouse cultured neurons at day 21 in vitro. Images at left are enlarged from corresponding boxed areas. Scale bars, 10μm. **g**, (top) Schematic representation of Xlr3b construct. Xlr3b contains a predicted typical nuclear localization signal (NLS) from residues 2-11 and a Cor1/Xlr/Xmr conserved region (Cor1) domain at residues 74-200. (bottom), Confocal images show co-localization of FLAG-tagged Xlr3b constructs (green) with TIA1 (red) in Neuro-2a cells. Nuclear DNA is labeled with DAPI (blue). Scale bars, 10μm. **h**, Neuro-2a cells were transfected with FLAG-tagged Xlr3b constructs, cell lysates were immunoprecipitated (IP) with anti-FLAG antibody, and western blots (WB) were probed with indicated antibodies. To validate an inhibitory effect of XIP, cells were treated 4 h with XIP peptides (1μM). In **d-h**, the experiments were repeated three times with similar results. Full-size scans of western blots shown in Supplementary Fig. 13.

**Figure 3. Aberrant Xlr3b expression perturbs synaptic plasticity by inhibiting dendritic CaMKIIα mRNA transport.** **a**, Xlr3 interacts with CaMKIIα, Arc and BDNF mRNAs in an RNA immunoprecipitation assay. Total mRNA was precipitated from P90 mouse hippocampal lysates with an Xlr3 antibody. cDNA was analyzed with primers amplifying CaMKIIα, Arc and BDNF mRNAs. **P < 0.01** by one-way ANOVA with Bonferroni's post hoc test; n = 5 mice each. **b**, Time-lapse images of mCherry-Xlr3b (red) and GFP-CaMKIIα 3’ UTR (green) co-transported in a dendrite of cultured neuron at day 21 in vitro. Granules moved in both anterograde (white arrowhead) and retrograde (yellow arrowhead) directions. Scale bars, 3μm. The experiments were repeated three times with similar results. **c**, (top, right) Representative
kymograph of movement of GFP-CaMKIIα 3’ UTR in a proximal dendrite. Scale bars, 5μm (x-axis) and 5min (y-axis). (top, left) Granules moved in anterograde (white arrowhead) and retrograde (yellow arrowhead) directions. Scale bars, 2μm. See also Supplementary Video 1. (bottom) Relative frequency of movement of GFP-CaMKIIα 3’ UTR granules. **P < 0.01 by two-way ANOVA with Bonferroni’s post hoc test; n = 5 neurons each, a proximal dendrite (20-100μm away from the cell body) per neuron to measure relative frequencies. Cells were treated with XIP (1μM) 4 h before imaging.

Figure 4. 5-ALA represses Xlr3b transcription with RNA polymerase II recruitment by modifying G-quadruplex structure. a, 5-ALA metabolism to indicated porphyrins. Succinylacetone inhibits 5-ALA dehydratase (ALAD). PBG, porphobilinogen; PpIX, protoporphyrin IX; FECH, ferrochelatase. b and c, Luciferase activity in Neuro-2a cells transfected with pGL3-2K or pGL3-2KΔG4 and treated with 5-ALA (0.1, 0.3, 1, 3 or 10μM) (b), or co-administered 10μM 5-ALA and succinylacetone at 1, 10, 100 or 1000μM (c) for 48h. Luciferase activity is shown relative to that in vehicle-treated cells. **P < 0.01 by one-way ANOVA with Bonferroni’s post hoc test; n = 3 biologically independent samples. d, Quantitative real-time RT-PCR showing Xlr3b mRNA expression in P90 mouse hippocampal lysates. **P < 0.01 vs. vehicle-treated WT mice, ##P < 0.01 vs. vehicle-treated AtrxΔE2 mice by one-way ANOVA with Bonferroni’s post hoc test; n = 6 mice each. e, (top) Representative immunoblot of P90 mouse brain lysates probed with indicated antibodies. (bottom) Densitometric analysis of Xlr3 normalized to β-tubulin (arbitrary units, A.U.). **P < 0.01 vs. vehicle-treated WT mice, ##P < 0.01 vs. vehicle-treated AtrxΔE2 mice by one-way ANOVA with Bonferroni’s post hoc test; n = 4 mice each. f, Percentage of 5-methylcytosine (5-mC) as determined by bisulfite sequencing analysis
in medial prefrontal cortex (PC) and hippocampus (HP) from male P90 WT or AtrxΔE2 mice. **P < 0.01 by two-way ANOVA with Bonferroni’s post hoc test; n = 9 clones. (n = 3 mice. 3 clones were sequenced per mouse.) See also Supplementary Fig. 10e, g, (top) The G-quadruplex prediction tool GQRS mapper was used to identify potential G-quadruplex forming sequences in the entire Xlr3b genomic DNA sequence, which is shown with exons highlighted in black and introns highlighted in gray. GQRS mapper provides a G-score (plotted in blue) which indicates the likelihood of G-quadruplex formation. Image is adapted from QGRS mapper (http://bioinformatics.ramapo.edu/QGRS/index.php). (bottom) Analysis of indicated amplicons in chromatin from hippocampus of P90 mice by chromatin immunoprecipitation with indicated antibodies. Results are expressed as percent input. **P < 0.01, *P < 0.05 vs. vehicle-treated WT mice, ###P < 0.01, #P < 0.05 vs. vehicle-treated AtrxΔE2 mice by one-way ANOVA with Bonferroni’s post hoc test; n = 6 mice each. Full-size scans of western blots shown in Supplementary Fig. 13.

**Figure 5. Treatment with 5-ALA counteracts cognitive deficits seen in AtrxΔE2 mice.**

a, (top) Representative kymograph of movement of GFP-CaMKIIα 3’ UTR in a proximal dendrite. Scale bars, 5μm (x-axis) and 5min (y-axis). (bottom) Relative frequency of movement of GFP-CaMKIIα 3’ UTR granules. Cells were treated with 5-ALA (1μM) for 7 days before imaging. *P < 0.05 vs. vehicle-treated WT neurons, ###P < 0.01 vs. vehicle-treated AtrxΔE2 neurons by two-way ANOVA with Bonferroni’s post hoc test; n = 5 neurons each, a proximal dendrite (20-100μm away from the cell body) per neuron to measure relative frequencies. Imm., immobile; Bidirect., bidirectional movement; Antero., anterograde movement; and Retro., retrograde movement. b, The number of GFP-CaMKIIα 3’ UTR (green) and PSD95 (magenta) double-positive puncta in a proximal dendrite. **P < 0.01 vs. vehicle-treated WT neurons, ###P < 0.01 vs. vehicle-treated AtrxΔE2 neurons by one-way ANOVA with Bonferroni’s post hoc test; n = 20 neurons each, a proximal dendrite per neuron. Whiskers represent minima and maxima. c, (left) Representative field excitatory post-synaptic potentials (fEPSPs) were recorded from the hippocampal CA1 region of mice. (middle) Changes in fEPSP slope recorded following high frequency stimulation (HFS) were attenuated in AtrxΔE2 mice, and 5-ALA treatment significantly rescued impaired LTP in hippocampal CA1. (right) fEPSP slope changes following HFS at 1 or 60 min, as shown at left. **P < 0.01, *P < 0.05 vs. vehicle-treated WT mice, ###P < 0.01, #P < 0.05 vs. vehicle-treated AtrxΔE2 mice by two-way ANOVA with post hoc.
Bonferroni’s multiple comparison test; WT + vehicle, Atrx^{ΔE2} + vehicle, and Atrx^{ΔE2} + 5-ALA, n = 8 mice; WT + 5-ALA, n=6 mice. d-f, 5-ALA treatment decreased cognitive deficits in Atrx^{ΔE2} mice based on memory-related tests. Novel-object recognition (NOR) test (d), latency time in retention trials in a passive avoidance (PA) test (e), and alternations in a Y-maze test (f). **P < 0.01 vs. vehicle-treated WT mice, ###P < 0.01, #P < 0.05 vs. vehicle-treated Atrx^{ΔE2} mice by one-way ANOVA with Bonferroni’s post hoc test; Respective sample sizes are indicated. g, Heat map summarizing expression data (left), and a list of rescued or not-rescued genes (right) between WT, Atrx^{ΔE2}, and 5-ALA treated Atrx^{ΔE2} mouse hippocampus at post-natal day (P) 90. The list was focused on 31 genes differentially expressed (8 up-regulated and 23 down-regulated) in WT versus Atrx^{ΔE2} samples in Supplementary Table 1 (an FDR < 0.05 and a log2 fold-change of > 0.5 or < 0.5). See also Supplementary Table 3, which includes 5-ALA treated Atrx^{ΔE2} group. Rescued or not rescued genes were assessed using the difference between the average value in log2 global normalization. WT, n=7 mice; Atrx^{ΔE2}, n=7 mice; Atrx^{ΔE2} + 5-ALA, n=4 mice. Full-size scans of western blots shown in Supplementary Fig. 13.
On-line Methods

Animals

Male mice (C57BL/6J) were used for all experiments. Mice were housed under climate-controlled conditions with a 12-h light/dark cycle and provided standard food and water ad libitum. Animal studies were conducted in accordance with the Tohoku University institutional guidelines. Ethical approval has been obtained from the Tohoku University Institutional Animal Care and Use Committee. Generation of homozygous AtrxΔE2 mice is described in 19. To generate Thy1-Xlr3b TG mice, Xlr3b cDNA was subcloned into the XhoI site of a mouse Thy1.2 expression cassette (Supplementary Fig. 9a). The Thy1 promoter construct was a gift of Joshua Sanes (Addgene plasmid # 20736). The Thy1.2-Xlr3b expression cassette was excised with NotI/PvuI and injected into fertilized eggs of B6 mice at Oriental Bio Service (Kyoto, Japan). Hemizygous Thy1-Xlr3b TG mice were generated by mating founder mice with B6 mice, and offspring were screened by PCR using genomic tail DNA. Primers used for amplification of Xlr3b sequences were ThXl(FW) (5′-GGTATTCATCATGCTCCG-3′) and ThXl(RV) (5′-GTTTCTGCCTCTCCTCACAG-3′), and the detected PCR product was 479 bp in founder animals (Supplementary Fig. 9b).

Cell culture

Neuro-2a mouse neuroblastoma cells (ATCC, CCL-131, authenticated by the provider using short tandem repeat profiling) were grown in Dulbecco's minimal essential medium (DMEM) supplemented with 10% heat-inactivated fetal bovine serum (FBS) and penicillin/streptomycin (100units/100μg/ml) in a 5% CO2 incubator at 37°C. Cells were routinely tested for mycoplasma contamination. Transfection was performed using Lipofectamine 2000 (Invitrogen, Carlsbad, CA, USA) transfection reagent according to the manufacturer's protocol. Primary cultures of neurons were established using previously described methods with slight modifications 56. Briefly, hippocampal tissue was dissected from embryonic day 18 mice and dissociated by trypsin treatment and trituration through a Pasteur pipette. Neurons were plated on coverslips coated with poly-L-lysine in Minimum Essential Medium (Invitrogen) supplemented with 10% FBS, 0.6% glucose (Wako, Osaksa, Japan), and 1mM pyruvate (Sigma-Aldrich, St. Louis, MO, USA). After cell attachment, coverslips were transferred to dishes containing a glial cell monolayer and maintained in Neurobasal medium (Invitrogen) containing 2% B27 supplement (Invitrogen) and 1% GlutaMax (Invitrogen). 5μM Cytosine
β-D-arabinofuranoside (Sigma-Aldrich) was added to cultures at DIV3 after plating to inhibit glial proliferation. Primary neurons were transfected with expression vectors and shRNAs using electroporation (NEPA21; NEPAGENE Co., Ltd. Chiba, Japan) at DIV0.

**Plasmid constructs and lentiviral transduction**

Plasmids expressing cDNAs encoding Xlr3a, Xlr3b, FAM9A, DNMT1, DNMT3A, H3.3 and DAXX were obtained from Kazusa DNA Research Institute (Promega, Madison, WI, USA) and cDNAs were inserted into pCMV vector. GST-Xlr3b was generated by cloning Xlr3b cDNA into pGEX-4T-1 (Pharmacia Biotech, Piscataway, NJ, USA). ATRX cDNA plasmid (pEGFP-C2-ATRX-HA) was kindly provided by Dr. David Picketts (University of Ottawa, Canada). ATRX shRNA (pSUPER-shATRX1) plasmid was kindly provided by Dr. Nathalie Berube (University of Western Ontario, Canada). GFP-MS2-nls and MS2 binding site (MS2bs)-CaMKIIα 3′UTR constructs were kindly provided by Dr. Kenneth S. Kosik (University of California Santa Barbara, Santa Barbara, USA). Xlr3b shRNA was purchased from Sigma-Aldrich (MISSION TRC-Mm2.0 TRCN0000255204). The non-targeting hairpin control SHC002 (Sigma-Aldrich), which containing a sequence that does not target any known human or mouse gene, was used as a negative control (shControl). FLAG-Xlr3b (FL, ΔNLS, Δ74-123, Δ124-200, and Δ158-170) and FLAG-FAM9A (FL and Δ292-304) were generated using the KOD-Plus Mutagenesis kit (Toyobo, Osaka, Japan) according to the manufacturer's protocol. For construction of Xlr3b promoter plasmids, genomic DNA isolated from mouse brain was obtained using a DNeasy Tissue Kit (Qiagen, Valencia, CA, USA). PCR was carried out to amplify a fragment containing the Xlr3b promoter region from nucleotides −2085 to +1 (where +1 is the A of the translation initiation codon). The fragment was then subcloned into the pGL3-basic luciferase reporter (Promega) vector (pGL3-2K) and sequenced. pGL3-2KΔAG4 and pGL3-1KΔCGI were generated using the KOD-Plus Mutagenesis kit (Toyobo). The CRISPR-Cas9 and GFP fusion protein expression lentivirus, U6gRNA-Cas9-2A-GFP plus guide Xlr3 sgRNA (Xlr3-Cas9-GFP), was purchased from Sigma-Aldrich. The Xlr3 guide sequence was TCATCTTTCAGTGCCATGG. Lentiviral transduction was performed according to the manufacturer's protocol (Sigma-Aldrich).

**Antibodies and peptides**

The following primary antibodies were used; anti-Xlr3 (1:500, generated commercially (MBL, Nagoya, Japan) against TDTAGRHSMDPNLLSDC peptide, anti-ATRX (1:100, Santa Cruz Biotechnology, H300 #sc-15408, Santa. Cruz, CA, USA),
anti-DNMT1 (1:1000, clone 60B1220.1, Abcam #ab13537, Cambridge, UK),
anti-DNMT3A (1:1000, clone 64B1446, Abcam #ab13888), anti-β-tubulin (1:5000,
clone AC-15, Sigma-Aldrich, #A5441), anti-hnRNP A/B (1:200: Santa Cruz
Biotechnology, G-10 #sc-376411), anti-hnRNP A/B (1:200: Santa Cruz Biotechnology,
M-15 #sc-82628), anti-DYNC1LI2 (1:1000, clone EPR11230(2), Abcam, #ab178702),
anti-DYNLL1 (1:200: Santa Cruz Biotechnology, #sc-136287), anti-histone H3.3
(1:1000, Abcam, #ab62642), anti-DAXX (1:200, Santa Cruz Biotechnology, M-112
#sc-7152), anti-hnRNP D (1:200, clone D6O4F, Cell Signaling Technology, Beverly,
MA, USA, #12382), anti-TIA1 (1:200, Santa Cruz Biotechnology, C-20, #sc-1751),
anti-FLAG (1:1000, clone M2, Sigma-Aldrich, #F1804), anti-synaptophysin (1:1000,
clone SVP-38, Sigma-Aldrich, #S5768), anti-MAP2 (1:10000, Abcam, #ab92434),
anti-RNA polymerase II (1:1000, clone CTD4H8, Millipore, MA, USA, #05-623),
anti-PSD95 (1:500, clone 6G6-1C9, Abcam, #ab2723), anti-phospho-CaMKII
(Thr286/Thr287, 1:5000, 18) and anti-CaMKIIα/β (1:5000, 18). For immunoblotting, the
following secondary antibodies were used: anti-mouse IgG antibody horseradish
peroxidase conjugated (1:5000, Southernbiotech, Birmingham, AL, USA, #1031-05),
anti-rabbit IgG antibody horseradish peroxidase conjugated (1:5000, Southernbiotech,
#4050-05) and anti-goat IgG antibody horseradish peroxidase conjugated (1:5000,
Rockland Immunochemicals Inc., Limerick, PA, USA, #605-4302). For
immunocytochemistry and immunohistochemistry, the following secondary antibodies
were used: Alexa488-conjugated donkey anti-rabbit (1:500, Invitrogen, #A-21206)
Alexa594-conjugated donkey anti-rabbit (1:500, Invitrogen, #A-21207), Alexa
488-conjugated donkey anti-mouse (1:500, Invitrogen, #A-21202), Alexa
594-conjugated donkey anti-mouse (1:500, Invitrogen, #A-21203), Alexa
488-conjugated donkey anti-goat (1:500, Invitrogen, #A-11055), Alexa 594-conjugated
donkey anti-goat (1:500, Invitrogen, #A-11058) and DyLight 405-AffiniPure donkey
anti-chicken IgY (1:500; Jackson ImmunoResearch, West Grove, PA, USA,
#703-475-155). XIP and ANTP control peptides were obtained from the
peptide-synthetic service at MBL.

**Luciferase assay**

Neuro-2a cells were co-transfected with pGL3 plasmids (pGL3-2K,
pGL3-2KΔG4 and pGL3-1KΔCGI) plus a pRL-TK plasmid, which contains Renilla
luciferase under control of the herpes simplex virus thymidine kinase promoter, with or
without other plasmids (ATRX, ATRX shRNA, control shRNA, DNMT1, DNMT3A,
H3.3 or DAXX). For *in vitro* DNA methylation assays, pGL3 plasmids were incubated
with 3U CpG methyltransferase (M. SssI) (New England Biolabs, Hitchin, UK) for 4h at 37°C in the presence of 1mM S-adenosylmethionine, following the manufacturer’s instructions. Firefly and Renilla luciferase activities were measured using the Dual-Luciferase Reporter Assay System (Promega) with a luminometer (Gene Light 55 Luminometer, Microtech, Chiba, Japan). The ratio of Firefly to Renilla luciferase luminescence was calculated.

Identification of Xlr3 interaction proteins by mass spectrometry
Mouse brain tissues at P60 were lysed in buffer containing 50 mM Tris-HCl (pH 7.5), 0.15 M NaCl, 0.1% Triton X-100, 4 mM EDTA, 4 mM EGTA, 1 mM Na3VO4, 50 mM NaF, 1 mM DTT, and protease inhibitors (trypsin inhibitor, pepstatin A, and leupeptin), followed by centrifugation at 15,000 × g for 10 min. Supernatants were collected and incubated with Protein A Sepharose column (Protein A HP SpinTrap, GE Healthcare Life Sciences, Piscataway, NJ, USA) with TBS buffer (50 mM Tris-HCl, 0.15 M NaCl, pH 7.5) containing an Xlr3 antibody (10 μg) at 4°C for 4 h with constant rotation. Bound proteins were then washed with TBS and eluted with 2.5% acetic acid. To confirm specific binding, samples were run on SDS-PAGE and the gel was stained using a Silver Stain Kit (Wako). All samples were then analyzed by LC-MS/MS at Oncomics Co., Ltd, (Nagoya, Japan), as a custom service. Proteins identified in control samples pulled down with rabbit IgG (10 μg) were subtracted from identified proteins.

Preparation of proteins binding to GST-Xlr3b
Preparation of proteins was performed using a GST Protein Interaction Pull-Down Kit (Pierce; Thermo Fisher Scientific, Rockford, IL, USA) according to the manufacturer’s instructions. GST and GST-Xlr3b were prepared in E. coli BL21 and immobilized to glutathione affinity resin. Mouse hippocampal tissues were lysed in buffer containing 50 mM Tris-HCl (pH 7.5), 0.15 M NaCl, 4 mM EDTA, 4 mM EGTA, 1 mM Na3VO4, 50 mM NaF, 1 mM DTT, and protease inhibitors (trypsin inhibitor, pepstatin A, and leupeptin), followed by centrifugation and incubation with immobilized glutathione affinity resin containing GST fusion proteins at 4°C for 4 h with constant rotation. Bound proteins were then washed, eluted with glutathione elution buffer, and run on SDS-PAGE.

Immunoprecipitation and immunoblotting
Immunoprecipitation and immunoblotting analysis was performed as described. Briefly, tissues or cells were homogenized in buffer containing 50 mM Tris-HCl, pH
7.5, 0.5% Triton X-100, 0.15 M NaCl, 4 mM EDTA, 4 mM EGTA, 1 mM Na\textsubscript{3}VO\textsubscript{4}, 50 mM NaF, 1 mM DTT, and protease inhibitors (trypsin inhibitor, pepstatin A, and leupeptin). For immunoprecipitation, lysates were incubated 2 h at 4°C with indicated antibodies with protein A-Sepharose CL-4B (GE Healthcare Life Sciences) in homogenization buffer. Subsequently, immunoprecipitates were washed three times with homogenization buffer, equivalent amounts of protein were electrophoresed on SDS-PAGE and proteins then transferred to an Immobilon polyvinylidene difluoride membrane. After blocking with TTBS solution (50 mM Tris–HCl, pH 7.5, 150 mM NaCl, and 0.1% Tween 20) containing 5% fat-free milk powder for 1 h at room temperature, membranes were incubated overnight at 4°C with indicated primary antibodies. After washing, membranes were incubated with the appropriate horseradish peroxidase-conjugated secondary antibody diluted in TTBS. Blots were developed using an ECL immunoblotting detection system (Amersham Biosciences, NJ, USA). Immunoreactive bands were visualized using a luminescent image analyzer LAS-4000 (Fuji Film, Tokyo, Japan) and quantified using Image Gauge version 3.41 (Fuji Film).

Subcellular fractionation.

Primary hippocampal neurons at DIV21 were homogenized in ice-cold lysis buffer containing the following (in mM): sucrose 320, HEPES 4, pH 7.4, EGTA 1, and protease inhibitors. The homogenate was centrifuged at 1,000 x g for 10 min at 4°C. The supernatant (S1, postnuclear supernatant) was centrifuged again at 12,000 x g for 15 min to obtain crude synaptosomal fraction (P2). For separating synaptosomal cytosol (LS1) and synaptosomal membrane (LP1), the pellet (P2) was hypo-osmotically lysed (5% lysis buffer; 95% distilled water containing 5% CHAPS and protease inhibitors) for 30 min and centrifuged at 100,000 x g for 60 min. The pellet (LP1) was resuspended in lysis buffer containing 1% Triton X-100 and sonicated.

Immunocytochemistry and immunohistochemistry

Immunocytochemistry and immunohistochemistry were performed as described Briefly, fixed cells or brain slices with 4% paraformaldehyde in phosphate-buffered saline (PBS) were treated with PBS containing 0.1% Triton X-100 for 10 min. Samples were incubated overnight at 4°C with indicated primary antibodies and then washed in PBS and incubated with appropriate secondary antibodies. Nuclei were stained with 4, 6-diamidino-2-phenylindole (DAPI) (Sigma-Aldrich). To visualize RNA molecules, living neurons were incubated with 50 nM SYTO14 dye (Life Technologies, Carlsbad, CA, USA, #S7576). Fluorescence intensities and images were analyzed by confocal
laser scanning microscopy (LSM700, Carl Zeiss, Thornwood, NY).

**Live imaging**

Live images of granule movement were captured in dendrites for 15 min (proximal dendrites) or 10 min (distal dendrites) at 15-s intervals in primary neurons at DIV21. Granule movement was analyzed in proximal dendrites (20-100μm) and in distal dendrites (100-200μm) away from the cell body. Relative frequency of movement of GFP-CaMKII 3’ UTR granules was analyzed for a subset of moving granules. Granules that changed direction during the tracking period were defined as bi-directional. Granules were classified as immobile when their movement was < 5 μm during the 15 min (proximal dendrites) or 10 min (distal dendrites) observation period. Kymographs were created using NIH ImageJ software (http://rsb.info.nih.gov/ij/).

**RT-PCR and quantitative RT-PCR (RT-qPCR) analysis**

Total RNA was purified from P90 mouse brain using an RNeasy Mini Kit (Qiagen) according to the manufacturer’s protocol. RNA was reverse-transcribed into single-stranded cDNA using an oligo(dT) primer (Promega) and Moloney murine leukemia virus-reverse transcriptase (Invitrogen), and then subjected to RT-PCR with gene-specific primers. RT-qPCR analysis was performed as described in 48-well plates (Mini Opticon real-time PCR system, Bio-Rad) using iQ SYBR Green Supermix 2× (Bio-Rad). Gene expression was assessed using the differences in normalized Ct (ΔΔCt) method after normalization to GAPDH. Fold-change was calculated by $2^{-ΔΔCt}$.

The following primers were used for RT-PCR and RT-qPCR:

In Fig. 1b, Fig. 4d, supplementary Fig. 1e, supplementary Fig. 9c, and supplementary Fig.10c.

- RT-qPCR-Xlr3b(FW) (5’-CAGAAAAAGGAAGGCCACTG-3’)
- RT-qPCR-Xlr3b(RV) (5’-GTTTTTCTTCTCCTGGCCTGT -3’)
- RT-qPCR-GAPDH(FW) (5’-TGTGTCCGTCGGATCTGA-3’)
- RT-qPCR-GAPDH(RV) (5’-CACCACCTTTCTTGATGTCATCATAC-3’)

In supplementary Fig. 1a,

- RT-PCR-Xlr3a(FW) (5’-AGCCGAGACCCGACCAAGTG-3’)
- RT-PCR-Xlr3a(RV) (5’-TGTTAGCTGGGTCTCTG-3’)
- RT-PCR-Xlr3b(FW) (5’-AGCCAAGGCCCCGACCAAGTG-3’)
- RT-PCR-Xlr3b(RV) (5’-TAGCTGTGCTGACTGACCTC-3’)

**Bisulphite sequencing**
Genomic DNA isolated from whole brain, medial prefrontal cortex and hippocampus of WT and Atrx^{ΔE2} mice at P90 were subjected to bisulphite-mediated C to U conversion using BisulFast (Toyobo, Osaka, Japan) and then used as template for PCR. The PCR products were ligated into the pCR2.1 vector by using a TOPO-TA cloning kit (Invitrogen), according to the manufacturer's instruction. The following primers specific for bisulfite-treated DNA were used for PCR:

Xlr-bisul(FW) (5′-GATTAAGTGGGATGAATTTTTGAGT-3′)
Xlr-bisul(RV) (5′-CCCAAACTAAAAATTTTCTCATTC-3′)

Chromatin immunoprecipitation (ChIP) assay
Chromatin solutions derived from P90 mouse hippocampus were prepared using the SimpleChIP Plus Enzymatic Chromatin IP Kit (Cell Signaling Technology) and then immunoprecipitated overnight at 4°C using 2 μg indicated antibodies. Assays included normal rabbit IgG as an antibody specificity control. The following antibodies and primers were used:

R1(FW) (5′-CCAAAGTGGAACCTCTGTATG-3′)
R1(RV) (5′-CCGGCGGAGTCCCACTGTGAACGT-3′)
R2(FW) (5′-CAGAAGTACTATACTAGAAAC-3′)
R2(RV) (5′-GGTTGGTCATACCTATGTAGGAAC-3′)

in Fig. 1

amplicon 1(FW) (5′-CCAAAGTGGGATGAACCTCTGTATG-3′)
amplicon 1(RV) (5′-CCGGCGGAGTCCCACTGTGAACGT-3′)
amplicon 2(FW) (5′-GCCACAGAGGGCAAGGCAAGGGAG-3′)
amplicon 2(RV) (5′-AAGTCAGGGAGGATGACCAGTCTC-3′)
amplicon 3(FW) (5′-AGTATCAGGCTTAGATTTAAAGGA-3′)
amplicon 3(RV) (5′-AAGTCAGGGAGGATGACCAGTCTC-3′)
amplicon 4(FW) (5′-CTGTGCAGGACTCCATGATACCC-3′)
amplicon 4(RV) (5′-CGTAAGCCCTTCAGCATACAAATCT-3′)

RNA immunoprecipitation (RIP) assay
Mouse hippocampal lysates were immunoprecipitated with an Xlr3 antibody (15 μg per sample) followed by RNA isolation using the RiboCluster Profiler RIP-Assay kit (MBL) according to the manufacturer's protocols. Immunoprecipitated RNA was reverse-transcribed into single-stranded cDNA using an oligo(dT) primer (Promega) and
Moloney murine leukemia virus-reverse transcriptase (Invitrogen), and subjected to RT-qPCR with the following gene-specific primers:

- **BDNF(FW)** (5′-TGGCCTAACAGTTTGCAG-3′)
- **BDNF(RV)** (5′-GGATTTGAGTGTGTTCTCC-3′)
- **CaMKIIα(FW)** (5′-GACACCAAAGTGCACATGG-3′)
- **CaMKIIα(RV)** (5′-GCGAAGCAAGGACACGG-3′)
- **Arc(FW)** (5′-AGCAGCAGACCTGACATCC-3′)
- **Arc(RV)** (5′-GGCTTGTCTTCACCTTCAG-3′)
- **GAPDH(FW)** (5′-TGTGTCCGTCGTGGATCTGA-3′)
- **GAPDH(RV)** (5′-CACCACCTTCTTGATGTCATC-3′)

**Gene microarray analysis**

For the oligo-DNA microarray analysis, we used the “3D-Gene” Mouse Oligo chip 24k and Human Oligo chip 25k arrays (Toray Industries Inc, Tokyo, Japan). Total RNA was purified from mouse hippocampus using an RNeasy Mini Kit (Qiagen) according to the manufacturer’s instructions. Total RNA was Cy5-labeled using the Amino Allyl MessageAMP II aRNA Amplification Kit (Applied Biosystems, CA, USA). Labeled aRNA pools were then hybridized 16 h in buffer using the supplier’s protocols (www.3d-gene.com). Hybridization signals were scanned using a ScanArray Express Scanner (PerkinElmer) and processed using the GenePixPro version 5.0 software (Molecular Devices).

**Electrophysiology**

Preparation of hippocampal slices was performed as described. Briefly, brains were rapidly removed from ether-anesthetized mice and chilled in ice-cold oxygenated artificial cerebrospinal fluid (ACSF: 126 mM NaCl, 5 mM KCl, 26 mM NaHCO₃, 2.4 mM CaCl₂, 1.3 mM MgSO₄, 1.26 mM KH₂PO₄, and 10 mM D-glucose). Transverse hippocampal slices (400-μm thickness) were cut using a vibratome (Microslicer DTK-1000, Dosaka EM, Kyoto, Japan) and transferred to a recording chamber where they were allowed to recover at least 1 h at 34°C before recording. ACSF maintained at 34°C was used during the experiment. A concentric bipolar stimulating electrode was placed in the stratum radiatum of CA1 to stimulate the Schaffer collateral pathway. High-frequency stimulation of 100 Hz with 1-s duration was applied twice with a 20-s interval. Traces were digitized with an A/D converter (PowerLab 200; AD Instruments, Castle Hill, Australia) and a computer (Windows, Measurement and Analysis System for LTP: FAL-3000). Paired-pulse responses were measured with inter-stimulus
intervals of 20-500 ms and expressed as the ratio of second stimulated amplitude to the first stimulated amplitude. The hippocampal input/output (I/O) ratio was determined for each group by measuring fEPSP amplitude in response to a series of stimuli with intensities ranging from 0.1 to 1.1 mA. After recording, slices were transferred to a plastic plate on ice to dissect the CA1 region under a microscope. CA1 regions were frozen in liquid nitrogen and stored at -80°C until biochemical analysis.

**Drugs**

5-ALA (COSMO BIO co., ltd. Tokyo, Japan), and sodium ferrous citrate were kindly provided by SBI Pharmaceuticals Co., Ltd., (Tokyo, Japan). TMPyP4 (5, 10, 15, 20-Tetrakis (1-methyl-4-pyridinio)porphyrin tetra) was purchased from Sigma-Aldrich. For administration to Atrx<sup>AE2</sup> mice, mice were randomized into 7 groups as follows: 5-ALA (3 and 10 mg/kg, p.o. with sodium ferrous citrate (20:1 mol. ratio) dissolved in distilled water) or vehicle (sodium ferrous citrate dissolved in distilled water) was administered daily from P30 to P90. TMPyP4 (10 and 30 mg/kg, i.p. in saline) or vehicle (saline) was administered twice weekly from P30 to P90. For acute administration, 5-ALA (10 mg/kg, p.o.) was administered 3 h before experiments. Measurement of biodistribution in TMPyP4 and 5-ALA was performed as described in 47, 49.

**Behavioral analysis**

Adult male mice at P90 were used for behavioral analysis. Animals were subjected to behavioral tests including Y-maze, step-through passive avoidance, novel object recognition and social interaction tasks. All behavioral experiments were conducted with the experimenter blind to mouse genotypes and treatments. In the Y-maze task, spontaneous alternation behavior in a Y-maze was assessed as a task of spatial reference memory. The apparatus consisted of three identical arms (50×16×32 cm<sup>3</sup>) of black plexiglas. Mice were placed at the end of one arm and allowed to move freely through the maze during an 8-min session. The sequence of arm entries was manually recorded. An alternation was defined as entries into all three arms on consecutive choices. The maximum number of alternations was defined as the total number of arms entered minus two, and the percentage of alternations was calculated as actual alternations/maximum alternations×100. The total number of arms entered during the session was also determined. In step-through passive avoidance task, training and retention trials of passive avoidance tasks were conducted in a box consisting of dark (25×25×25 cm<sup>3</sup>) and light (14×10×25 cm<sup>3</sup>) compartments. The floor was constructed
with stainless steel rods, and rods in the dark compartment were connected to an
electronic stimulator (Nihon Kohden, Tokyo, Japan). Mice were habituated to the
apparatus the day before passive avoidance acquisition. During training, a mouse was
placed in the light compartment, and when it entered the dark compartment, the door
was closed and an electric shock (0.4 mA for 2 s) was delivered from the floor. The
mouse was removed from the apparatus 30 s later. After a 7-day interval, each mouse
was placed in the light compartment and step-through latency was recorded over 300 s
to assess retention. In the novel object recognition task, mice were individually
habituated to an open-field box (35×25×35 cm$^3$) for 2 consecutive days. During
acquisition phase, two objects of the same material were placed symmetrically in the
center of the chamber for 10 min. 24 h later, one object was replaced by a novel object,
and exploratory behavior was analyzed again for 5 min. After each session, objects were
thoroughly cleaned with 70% ethanol to prevent odor recognition. Exploration of an
object was defined as rearing on the object or sniffing it at a distance of less than 1 cm,
touching it with the nose, or both. Successful recognition was reflected by preferential
exploration of the novel object. Discrimination of spatial novelty was assessed by
comparing the difference between exploratory contacts of novel and familiar objects
and the total number of contacts with both, making it possible to adjust for differences
in total exploration contacts. A social interaction test was established using methods
described in $^{57}$ with slight modifications. WT and Atrx$^{AE2}$ mice were housed in new
partitioned cages such that the Atrx$^{AE2}$ mouse occupied one compartment and a
weight-matched WT mouse (non-littermate) occupied the other. Forty-eight hours later,
the partition was removed and social interaction between mice was videotaped for
10 min. Scored behaviors were divided into four groups. Active social behavior,
initiated by the experimental mouse toward the partner, includes: (1) sniffing of any part
of partner's body including the anogenital area; (2) following including direct aggressive
attacks accompanied by bites toward the partner's back. Passive social behavior, which
occurs as a reaction to active behavior of the partner toward the experimental mouse,
includes (3) receptive responses when the experimental mouse tolerates sniffing by the
partner but shows no defensive or submissive behavior, (4) escape in response to a
partner's following or aggressive act. Data were analyzed as percentage of time spent in
various social behaviors during a 10-min interaction test. The videotapes for all
behavioral analysis were scored by a trained observer blind to genotype and treatment.

**Actin assays**

An *in vitro* F-actin sedimentation assay was performed using the Actin Binding
Protein Spin Down Assay (nonmuscle) kit (BK013; Cytoskeleton, Inc., Denver, CO, USA). Briefly, recombinant Xlr3b protein (1μM) purified from *E. coli* or α-actinin were mixed with polymerized nonmuscle actin (10μM) and incubated 30 min at 22°C. Mixtures were centrifuged to sediment F-actin, and supernatant and pellet fractions were analyzed by SDS–PAGE, followed by Silver staining. Actin polymerization was measured using a pyrene-actin polymerization kit (BK003, Cytoskeleton, Inc.) following the manufacturer's instructions.

CD spectra

All oligonucleotides were purchased from Sigma-Aldrich without further purification. Oligonucleotides for CD spectra were prepared in Li⁺ solution or K⁺ solution (20 mM lithium cacodylate, pH 7.0, 10 or 100 mM KCl). Annealing was performed by heating to 95°C for 5 min and cooling down slowly to room temperature. CD spectra were measured in 0.5-nm steps from 340 to 220 nm using a JASCO J-805LST Spectrometer and 1-cm quartz cuvettes.

UV melting experiment

UV–Vis melting temperature analyses were performed on a V-650 spectrophotometer (JASCO, Japan) with a thermocontrolled PAC-743R cell changer (JASCO, Japan) and a refrigerated and heating circulator F25-ED (Julabo). Oligonucleotides (ODN) (5 μM) in K⁺ solution (20 mM lithium cacodylate, pH 7.0, 5 mM KCl) and 1 eq compounds were added. Annealing was performed by heating to 95°C for 5 min and slowly cooling to room temperature. Melting temperature (*T*_m) of each sample was measured at 295 nm from 15 to 95°C at a rate of 1.0 °C /min. Δ*T*_m was calculated using *T*_m values of samples with and without compounds.

Gel electrophoresis

DNA samples (ODN) for gel electrophoresis were prepared similarly to CD spectra samples. Annealing was performed by heating to 95°C for 5 min and cooling slowly to room temperature. Each sample was analysed by native gel (8%) in 1×TBE buffer containing 10 or 100 mM KCl at room temperature. Single-stranded DNA was stained with SYBR® Green I Nucleic Acid Stain (50513, Lonza, Japan). Gels were then imaged by FLA-3000 (FUJIFILM).

DMS footprinting

Samples (ODN) for DMS footprinting were prepared similarly to samples of CD
spectra. DNA samples (10 μL) were then mixed with 1 μL of dimethyl sulfate solution (DMS:ethanol; 4:1, vol/vol). Reactions were quenched with 9 μL stop buffer (3 M -mercaptoethanol:water:NaOAc; 1:6:7, vol/vol). After ethanol precipitation and piperidine cleavage, reactions were separated on 12% denaturing polyacrylamide gels using a Hitachi DNA sequencer. Electrophoresis was conducted under 1.5 kV, ca. 25 mA, and 40°C.

**Statistical analysis**

To determine the sample size in our experiments, we followed the standard sample sizes used in similar experiments in each of the relevant fields in the literature. All values were expressed as means ± s.e.m. Comparisons between two experimental groups were made using the two-sided unpaired t test. Statistical significance for differences among groups was tested by one-way or two-way ANOVA with post-hoc Bonferroni's multiple comparison test. P < 0.05 was considered significant. All the statistical analyses were performed using GraphPad Prism 7 (GraphPad Software, Inc., San Diego, CA, USA). All statistical data were presented in the **Supplementary Table 5**.

Figure legends (for Supplementary materials only)

Supplementary Fig. 1. Identification of Xlr3b in mouse brain. a, Sequence of Xlr3a/Xlr3b mRNA and position of PCR primers. b, Xlr3a and Xlr3b primer specificity was confirmed by amplifying respective cDNAs. Serial dilutions of cDNA were prepared and Q-PCR performed. A negative correlation was found between increasing concentrations of cDNA and Ct. The R squared value (coefficient of determination, R²) was used to determine amplification efficiency. The experiments were repeated twice with similar results. c, Xlr3a and Xlr3b mRNA levels in mouse brain tissue as determined by RT-PCR. Xlr3a (lane 1) and Xlr3b (lane 2) cDNAs served as positive controls. W, wild-type mice; A, AtrxΔE2 mice; PC, prefrontal cortex; HP, hippocampus; HT, hypothalamus; CE, cerebellum. d, Cultured hippocampal neurons were transduced with U6gRNA-Cas9-2A-GFP plus guide Xlr3 sgRNA (Xlr3-Cas9-GFP) lentivirus (top) or control lentivirus expressing GFP (bottom) and immunostained 21 days later. Confocal images revealed loss of Xlr3 (red) immunoreactivity in MAP2-positive (blue) neurons infected with Xlr3-Cas9-GFP but not control GFP virus. In c and d, the experiments were repeated three times with similar results. e, Quantitative real-time RT-PCR showing Xlr3b mRNA expression in cells shown in d. **P < 0.01 by two-sided unpaired t-test. n = 4 biologically independent samples. f, Immunoblot (left) and corresponding quantitative (right) analysis of Xlr3 protein in cells shown in d. Densitometric analysis of Xlr3 normalized to β-tubulin (arbitrary units, A.U.). **P < 0.01 by two-sided unpaired t-test. n = 5 biologically independent samples.

Supplementary Fig. 2. Bisulfite sequencing evaluation of Xlr3 CGI methylation in AtrxΔE2 mouse brain lysates. a, (top) Schematic showing clusters of Xlr genes on the C57BL/6J X chromosome. See also Raefski and O'Neill, (2005) 23. (bottom) Location of Xlr3 CpG sites analyzed. Sequences potentially forming G-quadruplex is shaded in gray. b, Methylation status of Xlr3 CpG sites. Open circles, unmethylated CpGs; closed circles, methylated CpGs. Male P90 mice were used. n = 3 mice each. 4 independent clones of each sample were sequenced. Roman numerals correspond to those of the CpG sites shown in a and b.

Supplementary Fig. 3. G-quadruplex formation by sequences of Xlr3b CGI (Xlr3b-ODN). a, CD spectra of Xlr3b-ODN in Li⁺ or K⁺ solutions. b, Xlr3b-ODN primarily formed intramolecular, parallel G-quadruplexes based on native gel
electrophoresis. c, DMS footprinting of G-quadruplexes formed on Xlr3b-ODN in the Li⁺ or K⁺ solutions. In a-c, the experiments were repeated twice with similar results.

Supplementary Fig. 4. ATRX interacts with DAXX, DNMT1, DNMT3A and H3.3 in mouse hippocampus. a, Effect of ATRX shRNA in Neuro-2a cells. Immunoblot analysis (left) and densitometric quantification (right) of protein expression. Densitometric analysis of ATRX normalized to β-tubulin (arbitrary units, A.U.). **P < 0.01 by two-sided unpaired t-test. n = 3 biologically independent samples. b, (left) Representative immunoblot of P90 mouse hippocampal lysates probed with indicated antibodies, (right) Quantitative densitometry analyses. **P < 0.01 by two-sided unpaired t-test. n = 6 mice each. c, ATRX was immunoprecipitated (IP’d) from P90 WT and AtrxAE2 hippocampal extracts, and western blot analysis performed for indicated proteins. Control reactions were performed with IgG. Extracts were assessed as 1% input. The experiments were repeated three times with similar results.

Supplementary Fig. 5. Purified Xlr3b protein does not bind F-actin or alter actin polymerization. a, F-actin binding assay. Supernatant (S) and pellet (P) fractions were collected and samples separated on a SDS-gel subsequently stained using a Silver Stain Kit. Reactions were set up as indicated at top. Most α-actinin was found in pellet in the presence of F-actin filaments (lanes 9 and 10), while Xlr3b remained in the supernatant in presence of F-actin filaments (lanes 5 and 6). b, Actin polymerization as measured by enhanced fluorescence of pyrene-conjugated actin. Xlr3b addition to purified actin did not alter polymerization. In a and b, the experiments were repeated twice with similar results.

Supplementary Fig. 6. Xlr3b co-localizes with hnRNP A/B. a-c, Confocal images showing co-localization of FLAG-tagged Xlr3b constructs with hnRNP A/B (a), Dcp1a (b) and ubiquitin (c) in Neuro-2a cells. Nuclear DNA is labeled with DAPI (blue). Scale bars, 10μm. d, Neuro-2a cells were transfected with FLAG-tagged Xlr3b constructs, cell lysates were immunoprecipitated (IP) with anti-FLAG antibody, and western blot (WB) was probed with ubiquitin antibody. e, (top) Sequence comparison of Xlr3b residues 158-170 and comparable sequences from the Xlr human orthlogs FAM9A and FAM9B. Identical AAs are in blue and similar in light green. (bottom), Confocal images show localization of FLAG-tagged FAM9A constructs (green) in Neuro-2a cells. Nuclear DNA is labeled with DAPI (blue). Scale bars, 10μm. In a-e, the experiments were repeated three times with similar results.
Supplementary Fig. 7. Interaction of Xlr3b AA 158-170 and RBPs. **a**, Diagram of permeabilization control peptide (antennapedia homeodomain (ANTP)) and Xlr3b inhibitory peptide (XIP). The latter is a 29-AA peptide that contains Xlr3b AA 158-170 plus ANTP. **b**, FL-Xlr3b-transfected cells were treated with XIP (1μM for 4h) and immunostained for TIA1. Scale bars, 20μm. **c**, Pull-down assays with an ANTP antibody assessing TIA1 and hnRNP A/B in P90 mouse brain lysates. Eluted proteins and inputs were immunoblotted with indicated antibodies. Extract samples served as 1% input. IP, immunoprecipitation. In **b** and **c**, the experiments were repeated three times with similar results.

Supplementary Fig. 8. Dynamics of CaMKIIα mRNA transport in distal dendrites. **a**, Confocal images of mCherry-Xlr3b Δ124-200 or Δ158-170 (red) and GFP-CaMKIIα 3′ UTR (green) in cultured neurons at day 21 in vitro. Scale bars, 10μm. The experiments were repeated three times with similar results. **b**, (top, left) A confocal image of GFP-CaMKIIα 3′ UTR in primary mouse cultured neurons at day 21 in vitro. Scale bar, 50μm. Images at right is enlarged from corresponding boxed area. Scale bar, 10μm. See also Supplementary Video 2. (bottom) Relative frequency of movement of GFP-CaMKIIα 3′ UTR granules. **P < 0.01 by two-way ANOVA with Bonferroni's post hoc test; n = 5 neurons each, a distal dendrite (100-200μm away from the cell body) per neuron to measure relative frequencies. Cells were treated with XIP (1μM) 4 h before imaging. Imm., immobile; Bidirect., bidirectional movement; Antero., anterograde movement; and Retro., retrograde movement.

Supplementary Fig. 9. Generation of Thy1-Xlr3b transgenic mice. **a**, Genomic organization of Thy1 gene (top) and the transgenic construct (bottom). The Xlr3b cDNA was subcloned into a XhoI site of the Thy1.2 expression cassette. Blue boxes, untranslated exons. **b**, Representative PCR genotyping using tail DNA of transgenic founders (#13 and #57). M, size marker. **c**, Quantitative real-time RT-PCR showing Xlr3b mRNA expression in P90 mouse hippocampal lysates. **P < 0.01 by one-way ANOVA with Bonferroni's post hoc test; n = 6 mice each. TG, Xlr3b transgenic mouse. **d**, (top) Representative immunoblot of mouse hippocampal lysates probed with Xlr3 and β-tubulin antibodies, (bottom) Densitometric analysis of Xlr3 normalized to β-tubulin (arbitrary units, A.U.). **P < 0.01 by two-sided unpaired t-test; n = 5 mice each. **e**, Effect of Xlr3b shRNA in cultured neurons. Immunoblot analysis (top) and densitometric quantification (bottom) of protein expression. Densitometric analysis of
Xlr3 normalized to β-tubulin (arbitrary units, A.U.). *P < 0.05 by two-sided unpaired t-test; n = 3 biologically independent samples. f, (left) Method for isolation of synaptosomal membrane fractions from cultured neurons. The procedure for the subcellular fractionation is described in Methods. P1, nucleus/cell debris; S1, postnuclear supernatant; S2, cytosol fraction; P2, crude synaptosomal fraction; LS1, synaptosomal cytosol fraction; LP1, synaptosomal membrane fraction. Immunoblot showing CaMKIIα and postsynaptic marker, PSD95 between S2, LS1, and LP1 in cultured neurons from WT mice. (middle and right) Immunoblot and corresponding quantitative analysis of CaMKIIα and PSD95 proteins at LP1 and whole cell lysates in cultured neurons. Densitometric analyses of CaMKIIα normalized to PSD95 (arbitrary units, A.U.). **P < 0.01 by one-way ANOVA with Bonferroni's post hoc test. (In WT vs. Xlr3b-TG, **P < 0.01 by two-sided unpaired t-test.) LP1, n = 5 biologically independent samples; whole cell lysates, n = 4 biologically independent samples. g, (top) Representative field excitatory post-synaptic potentials (fEPSPs) were recorded from the hippocampal CA1 region of mice. (left), Changes in fEPSP slope following high frequency stimulation (HFS) were attenuated in Xlr3b-TG mice in hippocampal CA1. (right), Changes in fEPSP slope following HFS at 1 or 60 min. **P < 0.01 by two-way ANOVA with Bonferroni's post hoc test; n = 8 mice each. h, Paired pulse facilitation (left) and input-output relationship (right) were recorded. n = 5 mice each. There were no significant changes between the groups. i, LTP-induced CaMKIIα phosphorylation in the hippocampus. (left) Representative images of immunoblots using antibodies against phosphorylated CaMKIIα (pCaMKIIα) and total CaMKIIα. (right) Densitometric analysis of pCaMKIIα normalized to total CaMKIIα (arbitrary units, A.U.). **P < 0.01, *P < 0.05, vs. WT mice before high-frequency stimulation (HFS), ##P < 0.01, #P < 0.05, before HFS vs. after HFS in each group by two-way ANOVA with Bonferroni's post hoc test. Respective sample sizes are indicated. j-l, Xlr3b-TG mice show memory deficits. Novel-object recognition (NOR) test (j), Latency time in retention trials in a passive avoidance (PA) test (k), Alternations in a Y-maze test (l) are shown. **P < 0.01, *P < 0.05 by two-sided unpaired t-test. Respective sample sizes are indicated.

Supplementary Fig. 10. TMPyP4 treatment inhibits Xlr3b expression. a, The binding effect of Protoporphyrin IX (PpIX), hemin and TMPyP4 on G-quadruplexes formed by Xlr3b-ODN based on a UV melting experiment. b, Luciferase activity of Neuro-2a cells transfected with pGL3-2K or pGL3-2KΔG4 and treated with TMPyP4 (1, 3, 10, 30 and 100μM) for 48 h. Luciferase activity is shown relative to activity in
vehicle-treated cells. **P < 0.01 by one-way ANOVA with Bonferroni’s post hoc test; n = 3 biological replicates. 

**c, Quantitative real-time RT-PCR showing Xlr3b mRNA expression in mouse hippocampal lysates. **P < 0.01 vs. vehicle-treated WT mice, ##P < 0.01 vs. vehicle-treated Atrx<sup>ΔE2</sup> mice by one-way ANOVA with Bonferroni’s post hoc test; n = 4 mice each.

**d, (top) Representative immunoblot of mouse hippocampal lysates probed with Xlr3 and β-tubulin antibodies, (bottom) Densitometric analysis of Xlr3 normalized to β-tubulin (arbitrary units, A.U.). **P < 0.01 vs. vehicle-treated WT mice, ##P < 0.01 vs. vehicle-treated Atrx<sup>ΔE2</sup> mice by one-way ANOVA with Bonferroni’s post hoc test; n = 4 mice each.

**e, Methylation status of Xlr3b CpG sites. Open circles, unmethylated CpGs; closed circles, methylated CpGs. Male P90 mice were used. n = 3 mice each. three independent clones of each sample were sequenced.

**Supplementary Fig. 11. TMPyP4 treatment rescues cognitive deficits seen in Atrx<sup>ΔE2</sup> mice.**

**a, Relative frequency of movement of GFP-CaMKIIα 3’ UTR granules in distal dendrites. Cells were treated with 5-ALA (1μM) for 7 days before imaging, **P < 0.01, *P < 0.05 vs. vehicle-treated WT neurons, ##P < 0.01 vs. vehicle-treated Atrx<sup>ΔE2</sup> neurons by two-way ANOVA with Bonferroni’s post hoc test; n = 5 neurons each, a distal dendrite (100-200μm away from the cell body) per neuron to measure relative frequencies. Imm., immobile; Bidirect., bidirectional movement; Antero., anterograde movement; and Retro., retrograde movement.

**b, Paired pulse facilitation (left) and input-output relationship (right) were recorded. n = 5 mice each. There were no significant changes between the groups.**

**c, (left) Immunoblot and corresponding quantitative analyses of CaMKIIα and PSD95 proteins at LP1 (synaptosomal membrane fractions) and whole cell lysates in cultured neurons. Densitometric analyses of CaMKIIα normalized to PSD95 (arbitrary units, A.U.). *P < 0.05 vs. vehicle-treated WT neurons, #P < 0.05 vs. vehicle-treated Atrx<sup>ΔE2</sup> neurons by one-way ANOVA with Bonferroni’s post hoc test; n = 5 biologically independent samples. (right) LTP-induced CaMKIIα phosphorylation in the hippocampus. Representative images of immunoblots using antibodies against phosphorylated CaMKIIα (pCaMKIIα) and CaMKIIα. Densitometric analysis of pCaMKIIα normalized to total CaMKIIα (arbitrary units, A.U.). *P < 0.05, vs. WT mice before high-frequency stimulation (HFS), ##P < 0.01, before HFS vs. after HFS in each group by two-way ANOVA with Bonferroni’s post hoc test; n = 4 mice each.**

**d-f, TMPyP4 treatment rescued cognitive deficits in Atrx<sup>ΔE2</sup> mice based on memory-related behavioral tests. Novel-object recognition (NOR) test (d), latency time in retention trials in a passive avoidance (PA) test (e), alternations in a Y-maze test (f) are shown. **P < 0.01 vs. vehicle-treated WT mice, #P < 0.05, ##P < 0.01 vs. vehicle-treated WT mice, ##P < 0.01 vs. vehicle-treated Atrx<sup>ΔE2</sup> mice by one-way ANOVA with Bonferroni’s post hoc test; n = 4 mice each.
0.01 vs. vehicle-treated Atrx<sup>AE2</sup> mice by one-way ANOVA with Bonferroni's post hoc test; Respective sample sizes are indicated. **g, Atrx<sup>AE2</sup> mice show withdrawal in social interactions with WT mice. Atrx<sup>AE2</sup> mice showed enhanced passivity, higher escape duration and decreased social activity, such as following and sniffing behaviors, in social interactions with WT mice. These behaviors are dramatically improved by 5-ALA treatment. **<i>P</i> < 0.01, *<i>P</i> < 0.05 vs. vehicle-treated WT mice, ##<i>P</i> < 0.01, #<i>P</i> < 0.05 vs. vehicle-treated Atrx<sup>AE2</sup> mice by one-way ANOVA with Bonferroni's post hoc test; Respective sample sizes are indicated.

**h, Measurements of TMPyP4 fluorescence levels.** Chronic intraperitoneal injection of TMPyP4 in P90 mice increased fluorescence levels in some tissues, including brain. Respective sample sizes are indicated. **i, Measurements of body weight following chronic TMPyP4 administration on day 60 (i.p. twice weekly from P30 to P90).** Respective sample sizes are indicated. **j, Measurements of 5-ALA levels in P90 mouse brain after oral administration (3mg/kg, p.o.).** Respective sample sizes are indicated.

**Supplementary Fig. 12. Schematic showing outcomes following Atrx mutation.** ATRX binds to parallel G-quadruplexes in Xlr3b CGIs along with DNMTs, DAXX and H3.3, regulating Xlr3b gene expression through DNA methylation in cooperation with DMNTs. CaMKIIα mRNA locates in neuronal dendrites, and its transport is dynamically regulated by the motor protein, dynein. CaMKIIα translation enhances synaptic efficacy postsynaptically, which is critical for learning and memory. In control neurons, ATRX binds G-quadruplex-forming DNA in Xlr3b CGIs, inhibiting Xlr3b expression through DNA methylation. In Atrx mutant neurons, aberrant expression of Xlr3b protein occurs through DNA de-methylation at the site. Xlr3b protein has RNA binding capacity and cooperates with RNA binding proteins (RBPs), and inhibits dendritic transport of CaMKIIα mRNA, resulting synaptic dysfunction. Treatment with G-quadruplex (G4) ligand 5-ALA represses Xlr3b transcription, antagonizing both synaptic dysfunction and cognitive deficits in Atrx mutant mice.

**Supplementary Fig. 13. Full-size scans of western blots shown in figures.**

**Supplementary Table 1. List of genes exhibiting differential expression in hippocampus of P90 WT and Atrx<sup>AE2</sup> mice.**

**Supplementary Table 2. Xlr3 interaction partners identified in proteomic screen using LC-MS/MS analysis.**
Supplementary Table 3. List of genes exhibiting differential expression in P90 WT, AtrxΔE2 and 5-ALA treated AtrxΔE2 mouse hippocampus.

Supplementary Table 4. List of genes exhibiting differential expression in P90 WT, AtrxΔE2 and TMPyP4 treated AtrxΔE2 mouse hippocampus.

Supplementary Table 5. Summary of all statistical data.

Supplementary Video 1. Time-lapse imaging of GFP-MS2-labeled CaMKIIα mRNA (GFP-CaMKIIα 3′ UTR) in a proximal dendrite of a cultured WT neuron.

Supplementary Video 2. Time-lapse imaging of GFP-MS2-labeled CaMKIIα mRNA (GFP-CaMKIIα 3′ UTR) in a distal dendrite of a cultured WT neuron.
**Fig. 1**
Fig. 2
Fig. 3
Fig. 4
Fig. 5