

1 Erosion of somatic tissue identity with loss of the X-linked intellectual
2 disability factor KDM5C

3

4 **Abstract**

5 **Introduction**

6 A single genome holds the instructions to generate the myriad of cell types found within the adult organism. This is, in
7 part, accomplished by chromatin regulators that can either promote or impede lineage-specific gene expression through DNA
8 and histone modifications^{1,2}. Although many chromatin regulators were initially identified for their roles in shaping cellular and
9 tissue identity^{3,4}, recent advancements in next generation sequencing unexpectedly revealed many neurodevelopmental
10 disorders (NDDs) are caused by mutations in chromatin regulators⁵. Several studies have suggested this connection
11 between chromatin regulators and neurodevelopment is due to their regulation of brain-specific genes, such as orchestrating
12 transcriptional programs for synaptic maturation⁶ and transitioning between neuronal and glial fates during neural precursor
13 differentiation⁷. However, loss of some chromatin regulators can also result in the ectopic transcription of tissue-specific
14 genes outside of their target environment^{3,4,8}, such as the misexpression of liver-specific genes within adult neurons⁹. Very
15 few studies have investigated the misexpression of tissue-specific genes in chromatin-linked NDDs^{9,10} and it is currently
16 unknown if this partial loss of brain identity contributes to neurodevelopmental impairments.

17 To elucidate the role of tissue identity in chromatin-linked neurodevelopmental disorders, it is essential to first characterize
18 the types of genes dysregulated and the molecular mechanisms governing their de-repression. In this study, we characterized
19 the aberrant expression of tissue-enriched genes with loss of lysine demethylase 5C (KDM5C). KDM5C, , also known
20 as SMCX or JARID1C, is a chromatin regulator that can repress gene expression through erasure of histone 3 lysine 4
21 di- and trimethylation (H3K4me2/3)¹¹, marks enriched at active gene promoters. Pathogenic mutations in *KDM5C* cause
22 Intellectual Developmental Disorder, X-linked, Syndromic, Claes-Jensen Type (MRXSCJ, OMIM: 300534), whose predominant
23 features include intellectual disability, seizures, aberrant aggression, and autistic behaviors^{12–14}. *Kdm5c* knockout (-KO)
24 mice recapitulate key MRXSCJ patient phenotypes, including hyperaggression, increased seizure propensity, and learning
25 impairments^{10,15}. Unexpectedly, RNA sequencing (RNA-seq) of the *Kdm5c*-KO hippocampus revealed ectopic expression of
26 testis genes within the brain¹⁰. It is currently unknown if this dysregulation of brain tissue identity further impairs *Kdm5c*-KO
27 neurodevelopment and if ectopic gene expression within the *Kdm5c*-KO brain is unique to testis genes.

28 Intriguingly, some of the ectopic testis transcripts identified within the *Kdm5c*-KO hippocampus have known functions
29 unique to germ cells¹⁰, suggesting KDM5C may play a role in demarcating somatic versus germline identity. Distinguishing
30 between the germline (cells that pass on their genetic material, e.g. sperm and eggs) and the soma (cells that perform all other
31 bodily functions) is a key feature of multicellularity and occurs during early embryogenesis. In mammals, chromatin regulators
32 play a key role in decommissioning germline genes in somatic cells as the embryo implants into the uterine wall and transitions
33 from naive to primed pluripotency by placing repressive histone H2A lysine 119 monoubiquitination (H2AK119ub1)¹⁶, histone

34 3 lysine 9 trimethylation (H3K9me3)^{16,17}, and DNA CpG methylation¹⁷⁻¹⁹ at germline gene promoters. KDM5C may also be
35 involved in this early decommissioning of germline genes, as re-expression of KDM5C in knockout neurons fails to suppress
36 their dysregulation¹⁰. In support of this, KDM5C was very recently shown to repress *Deleted in azoospermia like (Dazl)*, a key
37 regulator of germline development, in mouse embryonic stem cells (ESCs)^{20,21}. However, KDM5C binding and *Kdm5c*-KO
38 germline gene misexpression has yet to be globally characterized during early embryogenesis. Given that *Dazl* and other
39 germline-enriched genes can also be expressed in ESCs and at the 2-cell stage, it is unclear if KDM5C has a direct role in the
40 long-term germline gene silencing that occurs in the post-implantation epiblast. Systematically characterizing KDM5C's role
41 in germline gene repression during early embryogenesis will unveil key mechanisms underlying the demarcation between
42 soma and germline identity and while also providing molecular footholds to test the impact of ectopic germline genes on
43 neurodevelopment.

44 To illuminate KDM5C's role in tissue identity, here we characterized the aberrant transcription of tissue-enriched genes
45 within the *Kdm5c*-KO brain and epiblast-like stem cells (EpiLCs), an *in vitro* model of the post-implantation embryo. We
46 observed general dysregulation of tissue-enriched genes in both the *Kdm5c*-KO brain and EpiLCs, including misexpression
47 of genes typically highly enriched within the testis, liver, muscle, and ovary. Both the *Kdm5c*-KO amygdala and hippocampus
48 had significant enrichment of testis genes that are typically uniquely expressed in germ cells. While the *Kdm5c*-KO brain
49 primarily expressed germline genes important for late spermatogenesis, *Kdm5c*-KO EpiLCs aberrantly expressed key drivers
50 of germline identity and meiosis, including *Dazl* and *Stra8*. KDM5C was highly enriched at germline gene promoters in
51 EpiLCs but only bound to a subset of germline genes expressed in *Kdm5c*-KO cells, indicating germline-enriched mRNAs
52 can be aberrantly transcribed through indirect mechanisms. Finally, we found KDM5C promotes the long-term silencing of
53 germline genes in somatic cells by aiding the placement of DNA methylation in EpiLCs through H3K4me2/3 removal. Thus,
54 we propose KDM5C plays a fundamental role in the development of tissue identity during early embryogenesis, including the
55 establishment of the soma-germline boundary.

56 Results

57 **Tissue-enriched genes, including testis genes, are aberrantly expressed in the *Kdm5c*-KO brain**

- 58 • note: should I compare amygdala and hippocampus? scandaglia only looked at hippocampus

59 Previous RNA sequencing (RNA-seq) in the adult hippocampus revealed ectopic expression of some testis genes within
60 the *Kdm5c* knockout (-KO) brain¹⁰. Since tissue-enriched genes have not been systematically characterized in the *Kdm5c*-KO
61 brain, it is currently unclear if this erosion of brain tissue identity is a major consequence of *Kdm5c* loss and if it is unique to
62 testis-enriched genes. Therefore, we first globally assessed the expression of genes enriched in 17 mouse tissues²² in our
63 published mRNA-seq datasets of the adult amygdala and hippocampus in mice with constitutive knockout of *Kdm5c*²³.

64 We found a large proportion of genes that are significantly upregulated within the *Kdm5c*-KO brain (DESeq2²⁴, log2 fold
65 change > 0.5, q < 0.1) are typically enriched in non-brain tissues (Amygdala: 35%, Hippocampus: 24%) (Figure 1A-B). The
66 majority of tissue-enriched differentially expressed genes (tissue-enriched DEGs) were testis genes (Figure 1A-C). Even
67 though the testis has the largest total number of tissue-biased genes compared to any other tissue (2,496 genes), testis-biased
68 DEGs were significantly enriched for both brain regions (Amygdala p = 1.83e-05, Odds Ratio = 5.13; Hippocampus p =
69 4.26e-11, Odds Ratio = 4.45, Fisher's Exact Test). One example of a testis-enriched DEG is *FK506 binding protein 6 (Fkbp6)*,
70 a known regulator of piRNA expression and meiosis in germ cells^{25,26} (Figure 1C).

71 In addition to the high enrichment of testis genes, we also identified aberrant expression of other tissue-enriched genes
72 within the *Kdm5c*-KO brain. Although the *Kdm5c*-KO mice were male, we observed significant enrichment of ovary-biased
73 DEGs in both the amygdala and hippocampus (Amygdala p = 0.00574, Odds Ratio = 18.7; Hippocampus p = 0.048, Odds
74 Ratio = 5.88, Fisher's Exact) (Figure 1D). Ovary-enriched DEGs included *Zygotic arrest 1 (Zar1)*, which was recently shown
75 to sequester mRNAs in oocytes for meiotic maturation and early zygote development²⁷ (Figure 1D). Although not consistent
76 across brain regions, we also found significant enrichment of DEGS biased towards two non-gonadal tissues - the liver
77 (Amygdala p = 0.0398, Odds Ratio = 6.58, Fisher's Exact Test) and the muscle (Hippocampus p = 0.0104, Odds Ratio = 6.95
78 Fisher's Exact Test). An example of a liver-biased DEG dysregulated in both the hippocampus and amygdala is *Apolipoprotein*
79 *C-I (Apoc1)*, which is involved in lipoprotein metabolism (Figure 1E). Testis, ovary, and liver-enriched DEGs showed little to no
80 expression in the developing and adult wild-type brain, yet our mRNA-seq data indicate they are polyadenylated and spliced
81 into mature transcripts (Figure 1C-E). Of note, we did not observe enrichment of brain-enriched genes (Amygdala p = 1;
82 Hippocampus p = 0.74, Fisher's Exact), despite the fact these are brain samples and the brain has the second highest total
83 number of tissue-enriched genes (708 genes). Together, these results suggest misexpression of tissue-enriched genes within
84 the brain is a major effect of KDM5C loss.

85 Germline genes are misexpressed in the *Kdm5c*-KO brain

86 The testis contains both germ cells (meiotic cells, e.g. spermatogonia) and somatic cells (non-meiotic, e.g. Leydig cells)
87 that support hormone production and germline functions. Intriguingly, many *Kdm5c*-KO testis and ovary enriched-DEGs have
88 germline-specific functions, suggesting *Kdm5c*-KO cells fail to distinguish between the soma and germline. To test if this
89 holds true for all *Kdm5c*-KO testis-biased DEGs, we first assed their known functions through gene ontology analysis. We
90 found *Kdm5c*-KO testis-enriched DEGs high enrichment of germline-relevant ontologies, including spermatid development
91 (GO: 0007286, p.adjust = 6.2e-12) and sperm axoneme assembly (GO: 0007288, p.adjust = 2.45e-14) (Figure 2A).

92 To further validate if these testis DEGs are truly germline genes, we then evaluated their expression in somatic versus germ
93 cells within the testis. We first compared their expression in the testis with germ cell depletion²⁸, which was accomplished by
94 heterozygous *W* and *Wv* mutations in the enzymatic domain of *c-Kit* (*Kit*^{W/Wv}) that prevent the maturation of germ cells²⁹.
95 Almost all *Kdm5c*-KO testis-enriched DEGs lost expression with germ cell depletion (Figure 2B). The only testis-enriched
96 DEG that did not show considerable downregulation with germline depletion was *FK506 binding protein 6 (Fkbp6)*, the
97 aforementioned testis gene that regulates piRNA expression and meiosis in germ cells^{25,26}. We then assessed testis-
98 enriched DEG expression in a published single cell RNA-seq dataset that identified cell type-specific markers within the
99 testis³⁰. We found that while some testis-enriched DEGs were classified as specific markers for different germ cell types
100 (e.g. spermatogonia, spermatocytes, round spermatids, and elongating spermatids) none marked somatic cells (Figure 2C).
101 Together, these data suggest the somatic *Kdm5c*-KO brain aberrantly expresses germline genes.

102 We then wanted to more deeply characterize germline gene misexpression with *Kdm5c* loss, but lacked a comprehensive
103 list of mouse germline-enriched genes. To facilitate downstream analyses, we generated a curated list of germline-enriched
104 genes using currently available RNA-seq datasets in *Kit*^{W/Wv} mice. Wild-type and *Kit*^{W/Wv} datasets included males and females
105 at embryonic day 12, 14, and 16³¹, as well as adult male testes²⁸. We defined genes as germline-enriched if their expression
106 met the following criteria: 1) their expression is greater than 1 FPKM in wild-type gonads 2) their expression in any adult
107 wild-type, non-gonadal tissue²² does not exceed 20% of their maximum expression in the wild-type germline, and 3) their
108 expression in the germ cell-depleted germline, for any sex or time point, does not exceed 20% of their maximum expression
109 in the wild-type germline. These criteria yielded 1,288 germline-enriched genes (Figure 2D), which was hereafter used as a

110 resource for assessing germline gene misexpression with *Kdm5c* loss (Supplementary table 1).

111 ***Kdm5c*-KO epiblast-like cells aberrantly express master regulators of germline identity**

112 Misexpression of germline genes in the adult brain suggests *Kdm5c*-KO cells fail to demarcate between germline and
113 somatic identity. Germ cells are typically distinguished from somatic cells soon after the embryo implants into the uterine
114 wall^{32,33} when a subset of epiblast stem cells become the primordial germ cells (PGCs) while the remainder differentiate into
115 the ectoderm, mesoderm, and endoderm to form the somatic tissues³⁴. This developmental time point can be modeled *in*
116 *vitro* through differentiation of embryonic stem cells (ESCs) into post-implantation epiblast-like stem cells (EpiLCs) (Figure 3A,
117 top). Previous studies have demonstrated that while some germline-enriched genes are also expressed in embryonic stem
118 cells (ESCs) and in the 2-cell stage³⁵⁻³⁷, they are silenced as they differentiate into EpiLCs¹⁷. Therefore, we assessed if
119 KDM5C was necessary for embryonic silencing of germline genes by evaluating the impact of *Kdm5c* loss in EpiLCs.

120 We first identified *Kdm5c*-KO EpiLC DEGs in of our previously published RNA-seq dataset³⁸ (DESeq2, log2 fold change
121 > 0.5, q < 0.1) and assessed tissue-enriched gene expression. Similar to the *Kdm5c*-KO brain, we observed general
122 dysregulation of tissue-enriched genes, with the largest number of genes belonging to the brain and testis, although they were
123 not significantly enriched (Figure 3B). Using the curated list of germline genes generated above, we found *Kdm5c*-KO EpiLCs
124 aberrantly expressed 54 germline-enriched genes, including the previously characterized hippocampal DEG¹⁰ *Cytochrome*
125 *C, testis-specific* (*Cyct*) (Figure 3C). Although we observed aberrant expression of many tissue-enriched genes, we did not
126 observe any significant difference in primed pluripotency genes (Figure 3D) or gross changes in *Kdm5c*-KO cell morpholgy
127 during differentiation (Figure 3E), indicating KDM5C loss does not impair EpiLC formation.

128 We then compared EpiLC germline DEGs to those expressed in the *Kdm5c*-KO brain to determine if all germline DEGs,
129 like *Cyct*, are constitutively dysregulated or if they can change over the course of development. We found this was primarily
130 not the case, as the majority of germline DEGs expressed in EpiLCs and the brain were unique, with only *Cyct* shared across
131 all sequencing datasets (Figure 3F). We then compared the known functions of EpiLC and brain germline DEGs and found
132 particularly high enrichment of meiosis-related gene ontologies in EpiLCs (Figure 3G), such as meiotic cell cycle process (GO:
133 1903046, p.adjust = 1.59e-08) and meiotic nuclear division (GO: 0140013, p.adjust = 9.76e-09). While there was modest
134 enrichment of meiotic gene ontologies in both brain regions, the *Kdm5c*-KO hippocampus primarily expressed late-stage
135 spermatogenesis genes, such those involved in the sperm axoneme structure.

136 Interestingly, DEGs unique to *Kdm5c*-KO EpiLCs included key drivers of germline identity, such as *Stimulated by retinoic*
137 *acid 8* (*Stra8*) and *Deleted in azoospermia like* (*Dazl*) (Figure 3H). These genes are typically expressed during embryonic
138 germ cell development to commit PGCs to the germline fate, but are also expressed later in life to trigger meiotic gene
139 expression programs³⁹⁻⁴¹. Of note, some germline genes, including *Dazl*, are also expressed in the two-cell embryo^{20,36}.
140 However, we did not see misexpression of two-cell embryo-specific genes, like *Duxf3* (*Dux*) (q = 0.337) and *Zscan4d* (q =
141 0.381), indicating *Kdm5c*-KO in EpiLCs do not revert back to a 2-cell state (Figure 3H).

142 We were particularly interested in the aberrant transcription of *Dazl*, since it is essential for germ cell development and
143 promotes the translation of germline mRNAs⁴². A significant portion of germline transcripts misexpressed in *Kdm5c*-KO
144 EpiLCs are known binding targets of DAZL, including *Stra8*⁴³ (p = 1.698e-07, Fisher's Exact Test). This suggests expression
145 of DAZL protein could promote the translation of other aberrant germline transcripts, influencing their ability to impact
146 *Kdm5c*-KO cellular function. We thus tested DAZL protein expression in *Kdm5c*-KO EpiLCs through immunocytochemistry
147 (Figure 3I). We observed about 25% of *Kdm5c*-KO EpiLCs expressed DAZL protein and it was localized to the cytoplasm (p =
148 0.0015, Welch's t-test), consistent with the pattern of DAZL expression in spermatogonia⁴³. Altogether these results suggest

149 tissue-specific genes are misexpressed during *Kdm5c*-KO embryogenesis, including key drivers of germline identity that can
150 be translated into protein.

151 KDM5C binds to a subset of germline gene promoters during early embryogenesis

152 Previous work suggests KDM5C represses germline genes during early development, as re-expression of KDM5C in
153 knockout neuronal cultures fails to suppress hippocampal germline DEGs and KDM5C is not bound to their promoters in
154 neurons¹⁰. There is some evidence KDM5C binds to select germline gene promoters in ESCs¹⁰, including *Dazl*^{20,21}. As
155 KDM5C's binding at germline gene promoters has not been systematically characterized, it is currently unclear if KDM5C is
156 enriched at germline gene promoters, what types of germline genes KDM5C regulates, and if its binding is maintained at any
157 germline genes in neurons.

158 To further characterize the mechanism behind *Kdm5c*-KO germline gene misexpression, we analyzed KDM5C chromatin
159 immunoprecipitation followed by DNA sequencing (ChIP-seq) datasets in EpiLCs³⁸ and primary neuron cultures (PNCs) from
160 the cortex and hippocampus¹⁵. EpiLCs had a higher total number of KDM5C peaks than PNCs (EpiLCs: 5,808, PNCs: 1,276,
161 MACS2 q < 0.1 and fold enrichment > 1, removal of *Kdm5c*-KO false positive peaks) and KDM5C was primarily localized to
162 gene promoters in both cell types (EpiLCs: 4,190, PNCs: 745 +/- 500bp from the TSS), although PNCs showed increased
163 localization to non-promoter regions (Figure 4A).

164 The majority of promoters bound by KDM5C in PNCs were also bound in EpiLCs (513 shared promoters), however
165 a large portion of gene promoters were only bound by KDM5C in EpiLCs (3677 EpiLC only promoters) (Figure 4B). We
166 then performed gene ontology analysis to compare the known functions of genes with KDM5C bound to their promoter in
167 EpiLCs and PNCs. While there were no ontologies significantly enriched for genes only bound by KDM5C in PNCs, gene
168 ontologies for genes shared between PNCs and EpiLCs were enriched for functions involving nucleic acid turnover, such
169 as deoxyribonucleotide metabolic process (GO:0009262, p.adjust = 8.28e-05) (Figure 4C). Germline-specific ontologies
170 were only enriched in promoters unique to EpiLCs, such as meiotic nuclear division (GO: 0007127 p.adjust = 6.77e-16)
171 and meiotic cell cycle process (GO:1903046, p.adjust = 5.05e-16) (Figure 3C). We then evaluated KDM5C binding around
172 the transcription start site (TSS) of all germline-enriched genes. While KDM5C was bound to a subset of germline gene
173 promoters in EpiLCs, it was not bound to any in PNCs (Figure 4D). Together, this suggests KDM5C is significantly enriched at
174 a subset of germline gene promoters in EpiLCs, including meiotic genes, but does not regulate germline genes in neurons.

175 We then compared KDM5C binding at RNA-seq DEG promoters to determine if the germline mRNAs expressed in the
176 *Kdm5c*-KO brain and EpiLCs are direct targets of KDM5C (Figure 4E). About one third of EpiLC-specific and brain-specific
177 (hippocampus, amygdala, or both) germline DEGs were bound by KDM5C in EpiLCs (EpiLC only: 36%, Brain only: 33.3%).
178 Some notable differences in KDM5C binding for EpiLC-specific DEGs included *Dazl* and *Stra8*, two key drivers of germline
179 identity discussed above. Although both mRNAs are expressed in *Kdm5c*-KO EpiLCs, KDM5C is only bound to *Dazl*'s
180 promoter and not *Stra8*'s in EpiLCs (Figure 4F). In contrast to the unique DEGs, 3 out of the 4 genes dysregulated in both
181 the brain and EpiLCs are bound by KDM5C around the TSS (Figure 4G). Again, we did not observe any KDM5C binding
182 at germline gene promoters in PNCs, even for brain-specific DEGs (Figure 4H). Altogether, this suggests the majority of
183 germline mRNAs expressed in *Kdm5c*-KO cells are dysregulated independent of direct KDM5C recruitment to their promoters
184 during embryogenesis.

185 **notes:** - do Direct vs indirect DEGs motif analysis

186 **KDM5C erases H3K4me3 to promote long-term repression of germline genes via DNA methylation**

187 Although KDM5C is generally thought to suppress transcription through erasure of histone 3 lysine 4 di- and trimethylation
188 (H3K4me2/3)¹¹, recent studies in ESCs have suggested KDM5C's repression *Dazl* is independent of its catalytic activity²⁰.
189 Somatic repression of germline genes is typically established during the transition between naive and primed pluripotency,
190 which modeled *in vitro* as ESCs to EpiLC differentiation. In ESCs, chromatin regulators place repressive histone modifications
191 at germline gene promoters, including histone 2A 119 monoubiquitination (H2AK119ub1) and histone 3 lysine 9 trimethylation
192 (H3K9me3)^{16,17,44}. Germline genes are then silenced long-term in EpiLCs by *de novo* placement of DNA CpG methylation
193 (DNAme)¹⁷. Although KDM5C's catalytic activity is not required to suppress *Dazl* in ESCs, KDM5C may promote long-term
194 germline gene silencing via H3K4me3 removal since H3K4me3 can impede DNAme placement^{45,46} and DNAme is lost at
195 select germline gene promoters in the hippocampus¹⁰. Because KDM5C's role in germline gene repression has only been
196 characterized in ESCs and in the mature brain, it is currently unclear to what extent KDM5C is involved during transition
197 between ESCs and EpiLCs and if its catalytic activity is required for long-term silencing.

198 To elucidate KDM5C's role in germline gene silencing, we first characterized KDM5C's substrates, histone 3 lysine 4
199 di- and trimethylation (H3K4me2/3) in our previously published ChIP-seq datasets in wild type and *Kdm5c*-KO amygdala²³
200 and EpiLCs³⁸. In congruence with previous work in the *Kdm5c*-KO hippocampus¹⁰, we observed aberrant accumulation of
201 H3K4me3 around the transcription start site (TSS) of germline genes in the *Kdm5c*-KO amygdala (Figure 5A). We additionally
202 found a marked increase in H3K4me2 around the TSS of germline genes in *Kdm5c*-KO EpiLCs (Figure 5B).

203 We then evaluated KDM5C's embryonic role in germline gene silencing during ESC to EpiLC differentiation. We first
204 characterized KDM5C's mRNA and protein expression at 0, 24, and 48 hours of differentiation (Figure 5C). While *Kdm5c*
205 mRNA steadily decreased from 0 to 48 hours of differentiation (Figure 5D), KDM5C protein initially increased from 0 to
206 24 hours but then decreased to near knockout levels by 48 hours (Figure 5E). We then performed whole genome bisulfite
207 sequencing in wild-type and *Kdm5c*-KO ESCs and 96 hour extended EpiLCs (exEpiLCs) to determine KDM5C's role in the
208 initial placement of DNA methylation at germline gene promoters. While wild-type cells accumulated high levels of DNA
209 methylation at germline gene promoters over the course of ESCs to exEpiLC differentiation, DNA methylation was markedly
210 reduced in *Kdm5c*-KO exEpiLCs (Figure 5F).

- 211 • Catalytic activity
212 • Altogether, this suggests KDM5C is dynamically regulated during ESC to EpiLC differentiation and promotes germline
213 gene silencing via removal of H3K4me2/3 during early embryogenesis.
214 • **note:** should I test if increase in H3K4me2 and H3K4me3 was highest in *Kdm5c*-KO cells at genes bound by KDM5C
215 at their promoter in EpiLCs? Don't think it's that impactful

216 **Discussion**

- 217 • The demarcation of the germ vs soma is a key feature of multicellularity
218 • Anything known about tissue-biased gene expression in other H3K4me regulators?
219 • Our data suggests the germline developmental program is occurring ectopically as *Kdm5c*-KOs progresses through
220 somatic tissue development

- 221 • tissue-biased gene expression:
- 222 – However unlike the gonadal-biased DEGs, many liver and muscle-biased DEGs have a known involvement in brain
223 function. For example, the liver-biased DEG is *Apolipoprotein C-I* (*Apoc1*), is important for lipoprotein metabolism
224 but has also been shown to influence learning and memory.
- 225 • Otx2 is expressed in EpiLCs and is known to repress PGC identity.
- 226 – It's properly expressed in *Kdm5c*-KO EpiLCs, further supporting they aren't just becoming PGCs <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6485399/>
- 228 • Altogether, these data indicate that while some germline genes are misexpressed due to direct loss of KDM5C binding
229 during emryogenesis, secondary downstream mechanisms can also promote their aberrant transcription.
- 230 – This shift from meiotic genes to later spermatogenesis genes in the hippocampus suggests the germline develop-
231 opmental program could occur ectopically as *Kdm5c*-KO cells progresses through somatic tissue development.
232 **note: this is strengthened by the ChIP-seq data since KDM5C is not directly bound to many brain/flagellar**
233 **DEGs. This point might be stronger in the ChIPseq figure**
- 234 • KDM5C is dynamically regulated during the window of embryonic germline gene silencing, our EpiLC ChIP-seq is likely
235 catching the tail end of KDM5C's main involvement.
- 236 • Papers to read/reference:
- 237 – Reconstitution of the Mouse Germ Cell Specification Pathway in Culture by Pluripotent Stem Cells: [https://www.cell.com/fulltext/S0092-8674\(11\)00771-9](https://www.cell.com/fulltext/S0092-8674(11)00771-9)
- 238 – two cell gene list used by Suzuki et al Max paper is based on 2 cell sequencing: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3395470/>

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334 **Figures and Tables**

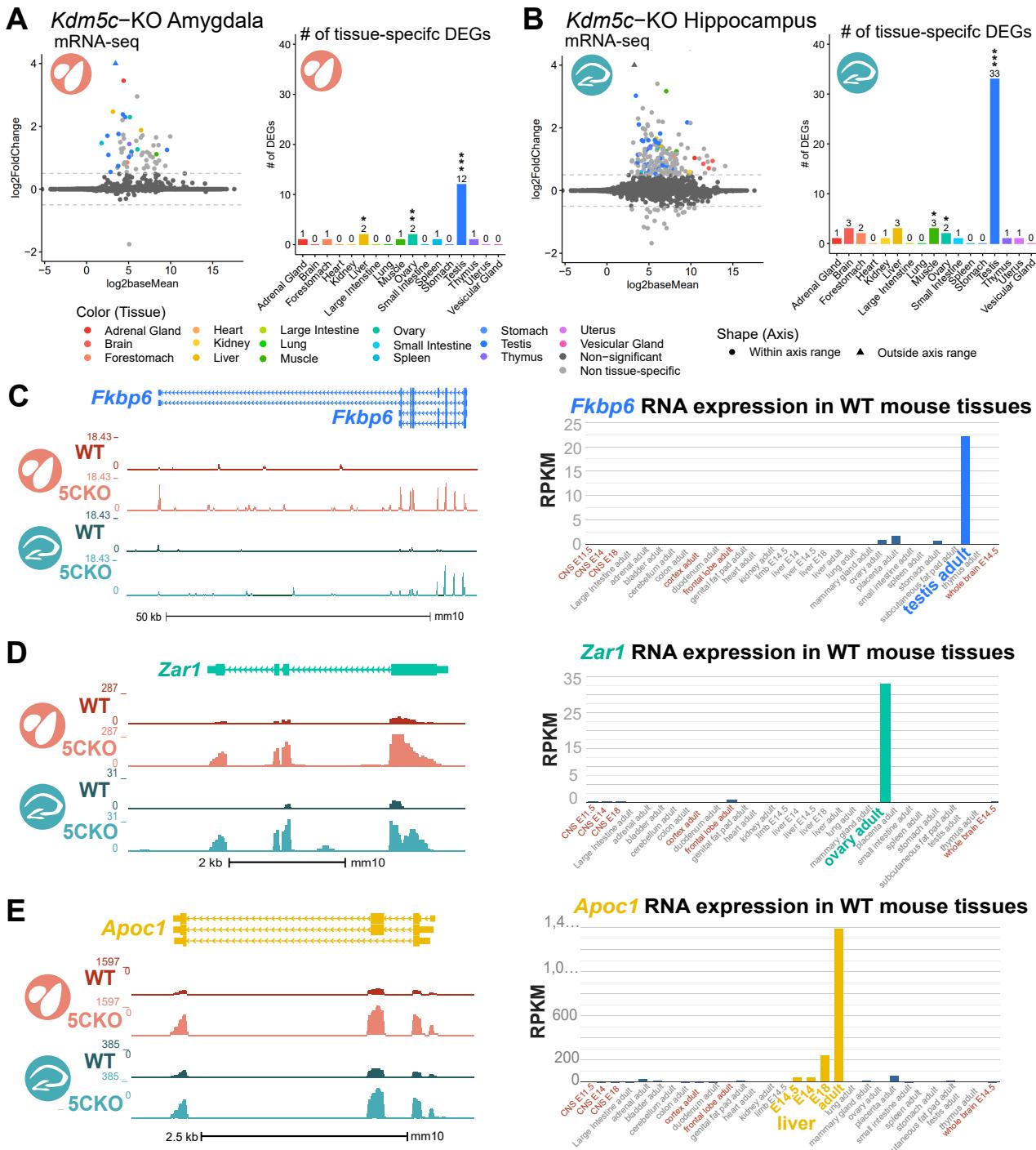


Figure 1: Tissue-enriched genes are misexpressed in the *Kdm5c*-KO brain. **A-B.** Expression of tissue-enriched genes (Li et al 2017) in the male *Kdm5c*-KO pY mouse amygdala (A) and hippocampus (B). Left - MA plot of mRNA-seq. Right - Number of tissue-enriched differentially expressed genes (DEGs). * $p<0.05$, ** $p<0.01$, *** $p<0.001$, Fisher's exact test. **C.** Left - Average bigwigs of an example aberrantly expressed testis-enriched DEG, *FK506 binding protein 6* (*Fkbp6*) in the wild-type (WT) and *Kdm5c*-KO (5CKO) amygdala (red) and hippocampus (teal). Right - Expression of *Cyct* in wild-type tissues from NCBI Gene, with testis highlighted in blue and brain tissues highlighted in red. **D.** Left - Average bigwigs of an example ovary-enriched DEG, *Zygotic arrest 1* (*Zar1*). Right - Expression of *Zar1* in wild-type tissues from NCBI Gene, with ovary highlighted in teal and brain tissues highlighted in red. **E.** Left - Average bigwigs of an example liver-enriched DEG, *Apolipoprotein C-I* (*Apoc1*). Right - Expression of *Apoc1* in wild-type tissues from NCBI Gene, with liver highlighted in orange and brain tissues highlighted in red.

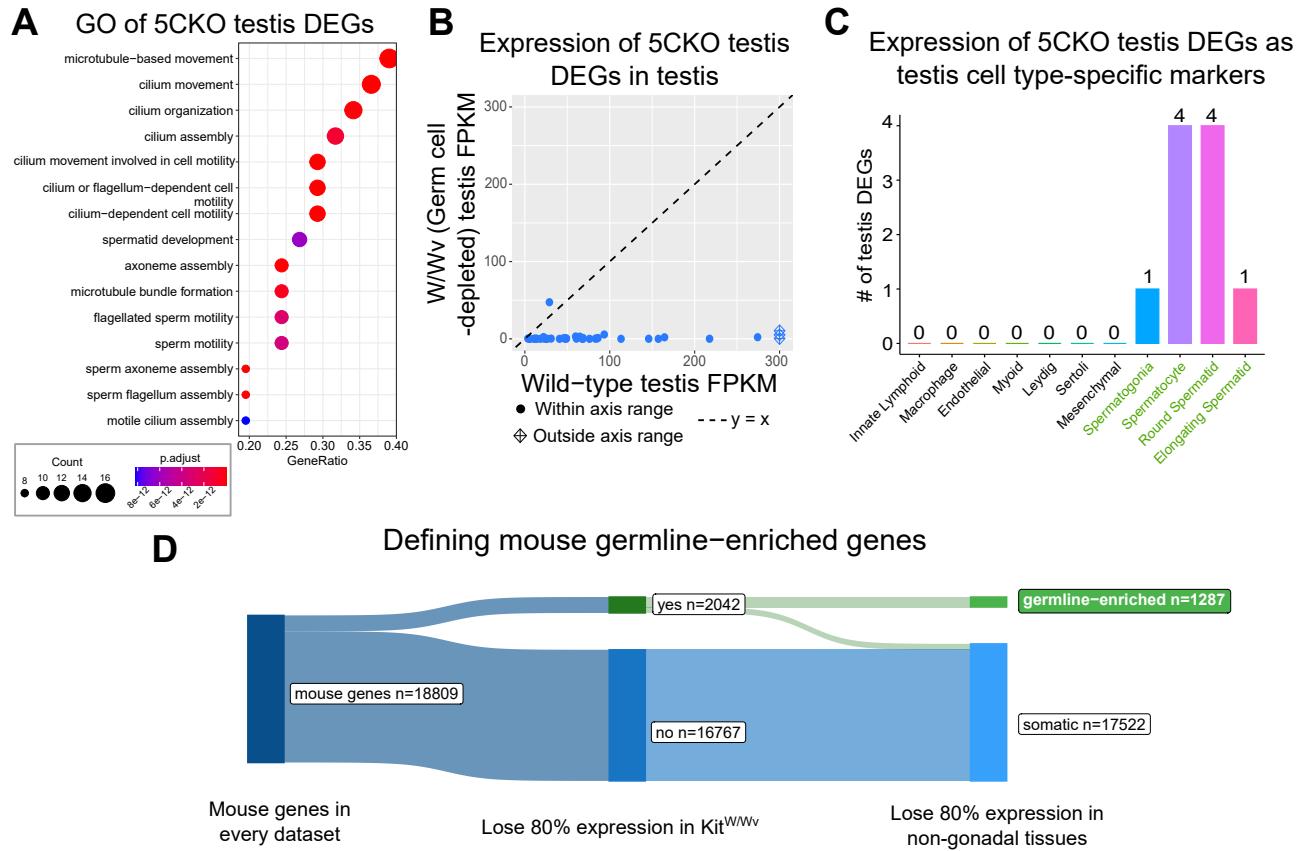


Figure 2: Aberrant transcription of germline genes in the *Kdm5c*-KO in the brain. **A.** enrichPlot gene ontology (GO) of *Kdm5c*-KO amygdala and hippocampus testis-enriched DEGs **B.** Expression of testis DEGs in wild-type (WT) testis versus germ cell-depleted (W/Wv) testis (Mueller et al 2013). Expression is in Fragments Per Kilobase of transcript per Million mapped reads (FPKM). **C.** Number of testis DEGs that were classified as cell-type specific markers in a single cell RNA-seq dataset of the testis (Green et al 2018). Germline cell types are highlighted in green, somatic cell types in black. **D.** Sankey diagram of mouse genes filtered for germline enrichment based on their expression in wild-type and germline-depleted mice and in adult mouse non-gonadal tissues (Li et al 2017).

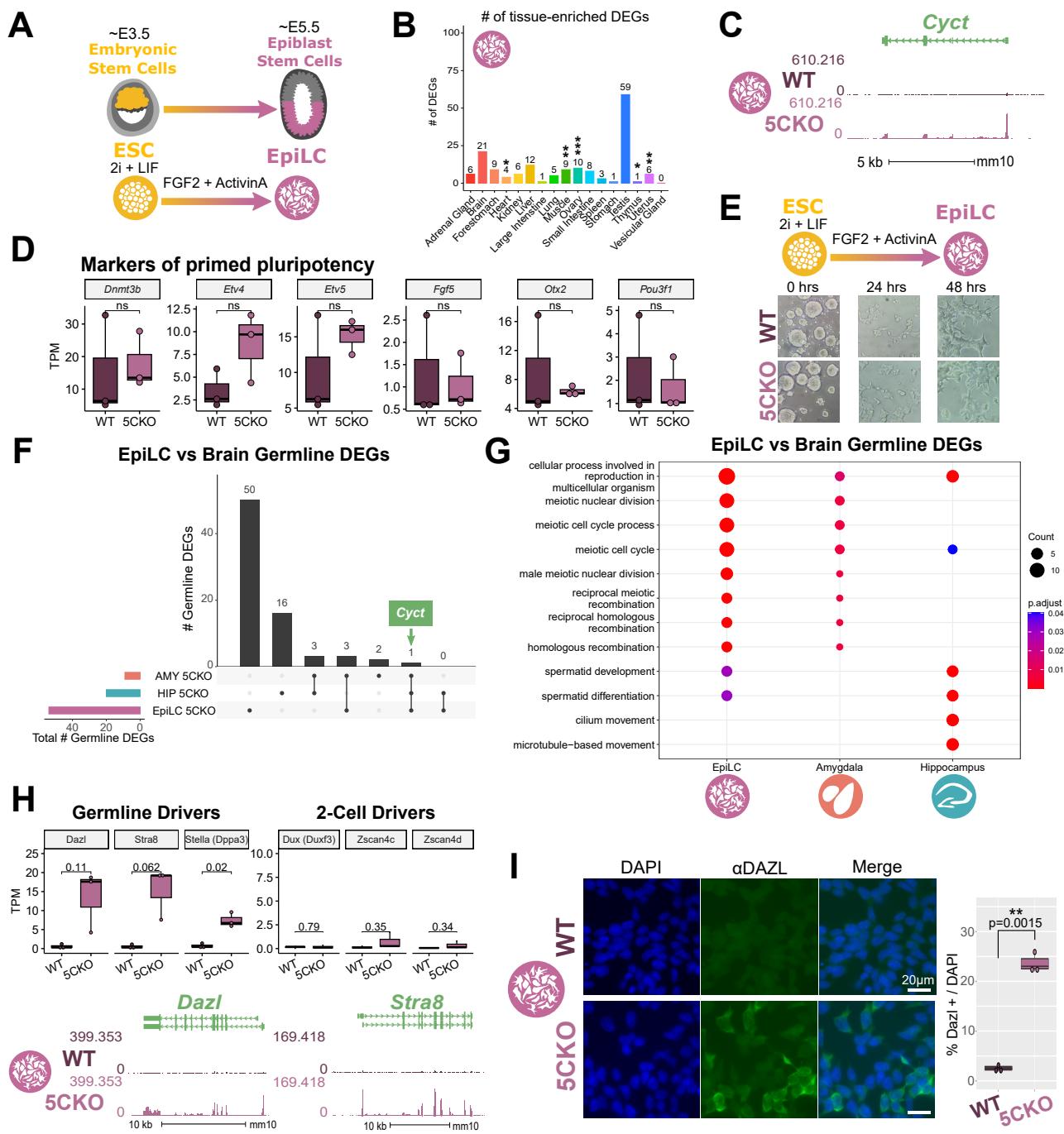


Figure 3: Kdm5c-KO epiblast-like cells express key drivers of germline identity

A. Top - Diagram of *in vivo* differentiation of embryonic stem cells (ESCs) of the inner cell mass into epiblast stem cells. Bottom - *in vitro* differentiation of ESCs into epiblast-like cells (EpiLCs).

B. Number of tissue-enriched differentially expressed genes (DEGs) in *Kdm5c*-KO EpiLCs. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, Fisher's exact test.

C. Average bigwigs of an example germline gene, *Cyct*, that is dysregulated *Kdm5c*-KO EpiLCs.

D. No significant difference in primed pluripotency marker expression in wild-type versus *Kdm5c*-KO EpiLCs. Welch's t-test, expression in transcripts per million (TPM).

E. Representative images of wild-type (WT) and *Kdm5c*-KO cells during ESC to EpiLC differentiation. Brightfield images taken at 20X.

F. Upset plot displaying the overlap of germline DEGs expressed in *Kdm5c*-KO EpiLCs, amygdala (AMY), and hippocampus (HIP) RNA-seq datasets.

G. enrichPlot comparing enriched biological process gene ontologies for *Kdm5c*-KO EpiLC, amygdala, and hippocampus germline DEGs.

H. Top left - Example germline identity DEGs unique to EpiLCs. Top right - Example 2-cell genes that are not dysregulated in *Kdm5c*-KO EpiLCs. p-values for Welch's t-test. Bottom - Average bigwigs of *Dazl* and *Stra8* expression in wild-type and *Kdm5c*-KO EpiLCs.

I. Immunocytochemistry of DAZL in male wild-type (WT) and *Kdm5c*-KO (5CKO) EpiLCs. Percentage of DAZL-positive cells normalized to number of DAPI-positive cells, p-value for Welch's t-test.

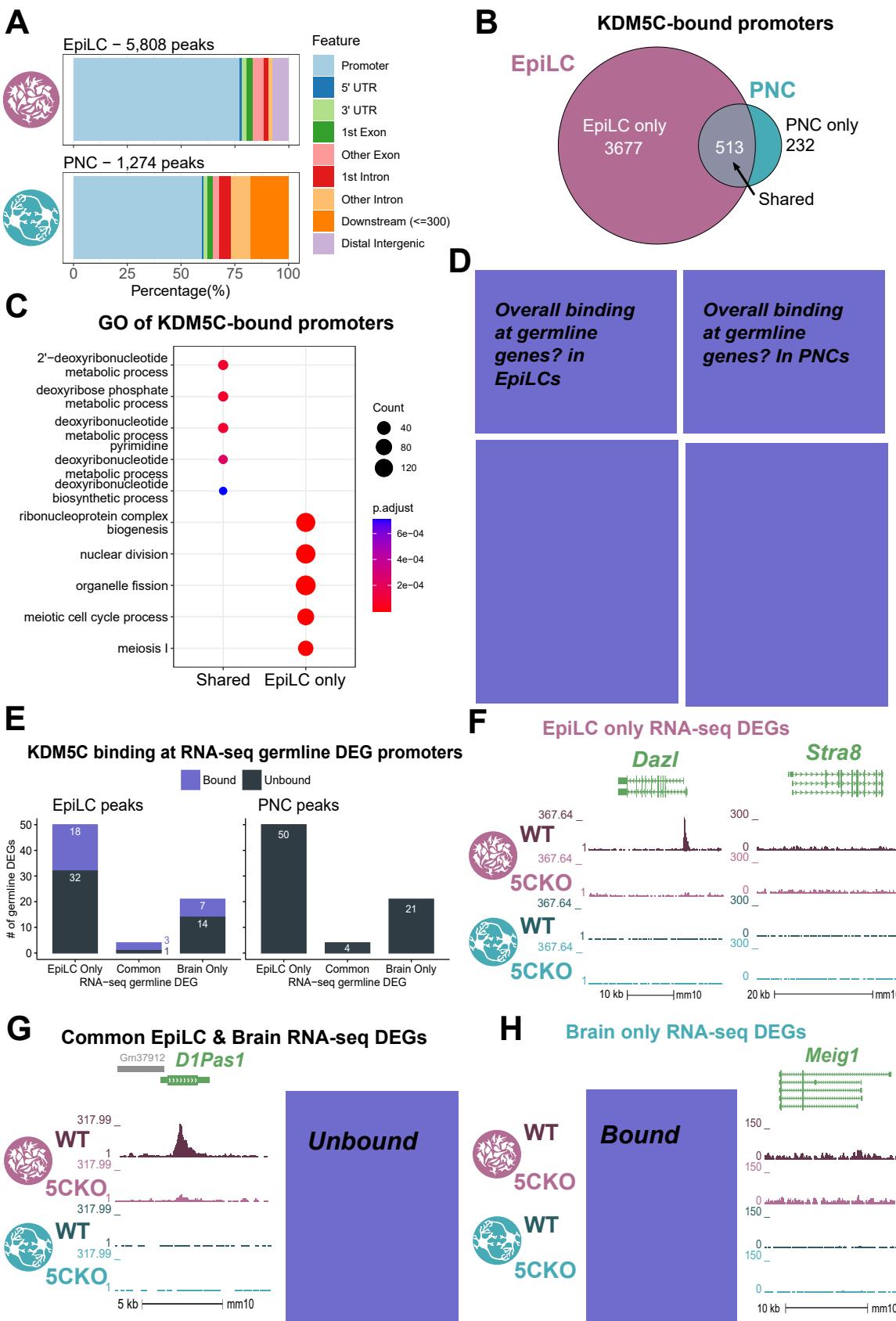


Figure 4: KDM5C binds to a subset of germline gene promoters during early embryogenesis **A.** ChIPseeker localization of KDM5C peaks at different genomic regions in EpiLCs (top) and hippocampal and cortex primary neuron cultures (PNCs, bottom). **B.** Overlap of genes with KDM5C bound to their promoters in EpiLCs (purple) and PNCs (blue). **C.** Gene ontology (GO) comparison of genes with KDM5C bound to their promoter in EpiLCs and PNCs. Genes were classified as either bound in EpiLCs only (EpiLC only), unique to PNCs (PNC only, no significant ontologies) or bound in both PNCs and EpiLCs (shared). **D.** Average KDM5C binding around the transcription start site (TSS) of all germline-enriched genes in EpiLCs (left) and PNCs (right). **E.** KDM5C binding at the promoters of RNA-seq germline DEGs. Genes were classified as either only dysregulated in EpiLCs (EpiLC only), genes dysregulated in the hippocampus or amygdala but not EpiLCs (brain only), or genes dysregulated in both EpiLCs and the brain (common). (Legend continued on next page.)

Figure 4: KDM5C binds to a subset of germline gene promoters during early embryogenesis. (Legend continued.) **F.** Example KDM5C ChIP-seq bigwigs of DEGs unique to EpiLCs. Although both are expressed in *Kdm5c*-KO EpiLCs, KDM5C is only bound to the *Dazl* promoter and not the *Stra8* promoter in EpiLCs. **G.** Example KDM5C ChIP-seq bigwigs of RNA-seq DEGs common between the brain and EpiLCs. KDM5C is bound to the *D1Pas1* promoter but not the *XXX* promoter in EpiLCs. **H.** Example KDM5C ChIP-seq bigwigs of RNA-seq DEGs unique to the brain. KDM5C is bound to the *XXX* promoter but not the *Meig1* promoter

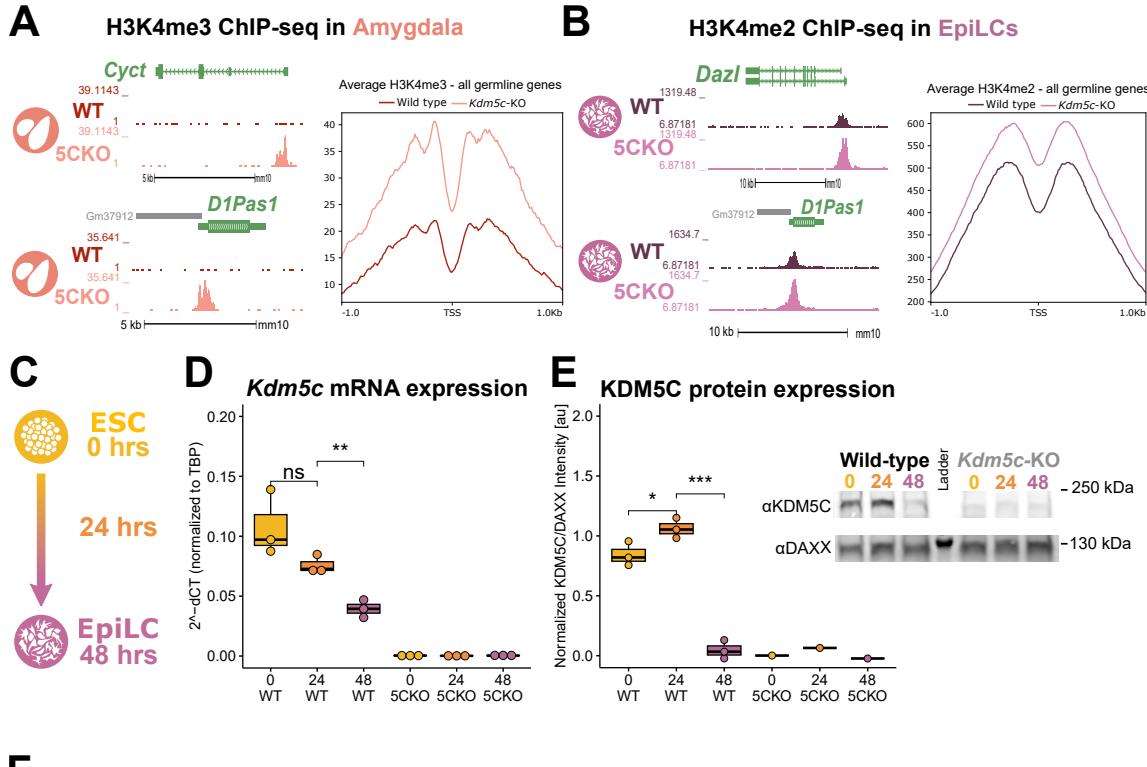


Figure 5: KDM5C's catalytic activity promotes long-term silencing of germline genes via DNA methylation. **A.** Left - Bigwigs of representative histone 3 lysine 4 trimethylation (H3K4me3) ChIP-seq peaks at two germline genes in the wild-type (WT) and *Kdm5c*-KO (5CKO) adult amygdala. Right - Average H3K4me3 at the transcription start site (TSS) of all germline-enriched genes in wild-type (dark red) and *Kdm5c*-KO (light red) amygdala. **B.** Left - Bigwigs of representative histone 3 lysine 4 dimethylation (H3K4me2) ChIP-seq peaks at representative germline genes in wild-type and *Kdm5c*-KO EpiLCs. Right - Average H3K4me2 at the TSS of all germline-enriched genes in wild-type (dark purple) and *Kdm5c*-KO (light purple) EpiLCs. **C.** Diagram of embryonic stem cell (ESC) to epiblast-like cell (EpiLC) differentiation protocol and collection time points for RNA and protein. **D.** Real time quantitative PCR (RT-qPCR) of *Kdm5c* RNA expression, calculated in comparison to TBP expression ($2^{-\Delta\Delta CT}$). * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, Welch's t-test. **E.** KDM5C protein expression normalized to DAXX. Quantified intensity using ImageJ (artificial units - au). Right - representative lanes of Western blot for KDM5C and DAXX. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, Welch's t-test **F.** XXX **G.** XXX

335 **Notes**

336 **Figure outline:**

337 **Figure 1: Misexpression of tissue-specific genes in the *Kdm5c*-KO brain** * MA-plot and bar graphs of tissue-enriched
338 genes * Example testis-specific genes (NCBI and bigwigs) * An example ovary tissue-specific gene * An example muscle/liver
339 tissue-specific gene (NCBI and bigwigs)

340 **Figure 2: The male *Kdm5c*-KO brain expresses male and female germline-enriched genes** * Gene ontology of
341 testis DEGs in the amygdala and hippocampus - ontologies are germline ontologies * Expression of testis DEGs in germline-
342 depleted testis (this is adult testis data) * scRNASeq of testis - # of testis DEGs that are germline-specific markers * Although
343 far fewer, 5CKO brain also expresses ovary-enriched genes(NCBI and bigwigs of Zar1) * These ovary enriched genes are
344 also germline specific (NCBI/Li tissues are in adult ovary, so it would be best to show they're oocyte-specific in adult ovary).
345 But I don't think there's a published adult female W/Wv dataset. Could try looking at scRNASeq or just do TPM in embryonic
346 W/Wv data since oocytes are developed at this point? Or both?) * Defining what is/isn't a germline gene, and which are
347 male/female biased using embryonic W/Wv data

348 **Figure 3: *Kdm5c*-KO epiblast-like cells express key drivers regulators of germline identity** * A) ESC to EpiLC
349 differentiation Left - Morphology is unchanged, * B) 5CKO EpiLCs express EpiLC differentiation genes similar to WT lvs *
350 C) Male EpiLCs express germline genes (example Cyct again) * Overlap between brain and EpiLC germline genes - show
351 they're mostly unique * GO of Brain and EpiLC germline genes (meiotic enriched) * Bigwigs or TPM of master regulators
352 * Show that while some are also 2-cell genes (Dazl), 2-cell specific genes aren't dysregulated (Zscan4). Important point
353 because published KDM5C dazl paper is saying KDM5C is a 2-cell regulator, but as far as I can tell only genes shared
354 between germline and 2-cell are dysregulated.

355 Staining of Dazl (+ Stra8 if I can get it to work)

356 **Figure 4: Loss of KDM5C's catalytic activity impairs DNAme placement and long-term silencing of germline
357 genes** * Increase in H3K4me3 in *Kdm5c*-KO amygdala at germline genes * Increase in H3K4me2 in EpiLCs at germline
358 genes * *Kdm5c* binding in EpiLCs vs PNCs to show that germline repression is happening in early embryo * Previous studies
359 only looked at ESCs, unknown if catalytic activity is required for long-term repression, especially since DNA methylation is
360 placed later). KDM5C RNA and protein ESC → EpiLC (increasing then decreasing) * RNA expression of germline genes with
361 catalytic dead rescue (Ilakkiya) * DNA methylation in WT and 5CKO EpiLCs (Ilakkiya)

362 **Figure 5: Ectopic, germline-like phenotypes in *Kdm5c*-KO ESCs/EpiLCs** * Sycp3 staining * DDX4 staining and
363 repression of retrotransposons * Cilia??

364 Gaps in knowledge addressed: * Are other tissue-enriched genes dysregulated, or only testis, germline genes? * Curating
365 a robust list of male and female germline genes * Should talk about 2-cell genes vs germline genes - way to systematically
366 categorize? * Mechanism behind long-term germline gene misexpression * Recent evidence suggests loss of KDM5C
367 in ESCs express some germline genes * Unclear if catalytic activity is required for long-term silencing * Unclear if their
368 dysregulation lasts throughout life or the same between brain or not * When in development does it begin? - Recent evidence
369 suggests some germline genes expressed in 5CKO ESCs but unclear if their dysregulation lasts throughout differentiation
370 and if the identity of germline genes are different compared to the brain * Are there functional consequences to germline
371 gene misexpression?

372 Introduction: * Chromatin regulators are important for cellular identity * H3K4me1-3 linked to active gene promoters and
373 enhancers * Surprisingly, mutations in Chromatin regulators lead to many NDDs (including many H3K4 regulators) * Recent

374 studies have shown some chromatin regulators are important for regulating neuron-specific gene expression/chromatin
375 stat_compare_means * However, loss of some chromatin regulators can also lead to ectopic expression of tissue-enriched
376 genes * Very few studies have looked at these genes and it's unclear if these genes contribute to NDD impairments. *
377 Necessary to first characterize the mechanism behind their derepression to identify molecular footholds into testing their
378 contribution to neuronal impairments and potential for therapeutic intervention

- 379 • Loss of KDM5C can result in the misexpression of genes typically only found in the testis
- 380 – Misexpression of tissue-enriched genes hasn't been systematically characterized - Unclear if these genes are
381 exceptions or if other tissue-specific genes are dysregulated
- 382 – Interestingly, these genes (Cyct, D1pas1) typically function in the germline
- 383 – Germ cells (meiotic cells) are typically distinguished from somatic cells very early on in embryogenesis and is a
384 key feature of multicellularity
- 385 – Chromatin regulators are very important for decommissioning germline genes and act successively the embryo
386 implants into the uterine wall
- 387 * Most studies have focused on ESCs, which have a similar transcriptome to germ cells / 2-cells
- 388 * recently, KDM5C was shown to repress DAZL in ESCs, independent of its catalytic activity
- 389 * However, DNA methylation is lost in the mature 5CKO brain, DNA methylation is placed later and it's unclear if
390 it's required for long-term repression (maybe too specific, just trying to go into the fact that the mechanism is
391 partially understood but unclear)
- 392 – Systematic characterization of ectopic germline genes hasn't been done
- 393 * unknown if other germline-enriched genes are dysregulated, including oocyte-specific genes
- 394 * Crucially, it's unknown if misexpression of the germline program leads to functional consequences in 5CKO
395 cells.

396 **Germline gene repression background:**

397 Interestingly, some of the ectopic testis transcripts identified in the *Kdm5c*-KO brain are typically expressed in germ cells¹⁰.
398 Unlike somatic cells, germ cells (e.g. sperm and eggs) undergo meiosis and pass on their genetic material to the next genera-
399 tion. The germline and the soma are typically distinguished during early embryogenesis, when germline genes are silenced
400 in epiblast stem cells soon after implantation and only reactivated in a subset to form the germline. Chromatin regulators
401 play a key role in decommissioning germline genes as the embryo transitions from naive to primed pluripotency by placing
402 repressive histone H2A lysine 119 monoubiquitination (H2AK119ub1)¹⁶, histone 3 lysine 9 trimethylation (H3K9me3)^{16,17},
403 and DNA CpG methylation¹⁷⁻¹⁹ at germline gene promoters. KDM5C may also be involved in this early decommissioning of
404 germline genes, as re-expression of KDM5C in neurons fails to suppress their dysregulation¹⁰. In support of this, KDM5C
405 was very recently shown to repress *Deleted in azoospermia like* (*Dazl*), a key regulator of germline development, in mouse
406 embryonic stem cells (ESCs)^{20,21}. In support of this, two independent screens in mouse embryonic stem cells (ESCs) recently
407 identified KDM5C as a repressor of *Deleted in azoospermia like* (*Dazl*), a key regulator of germline development. However,
408 KDM5C's role in embryonic germline gene repression is currently unclear, given that *Dazl* is also expressed in ESCs and in
409 the 2-cell stage and germline gene misexpression has yet to be globally characterized during *Kdm5c*-KO embryogenesis.