

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

3

4 Abstract

It is currently unclear why mutations in numerous chromatin-modifying enzymes cause neurodevelopmental disorders (NDDs). While some chromatin regulators are important for neuron-specific gene expression, loss of repressive chromatin regulators can also lead to the aberrant transcription of tissue-specific genes outside of their intended context. Because very few studies have characterized these aberrant, tissue-specific transcripts, the mechanisms behind their dysregulation and their functional consequences are largely unknown. Here, we explored this crisis in cellular identity in NDDs by investigating the ectopic expression of germline (e.g. sperm and egg) genes in mice lacking lysine demethylase 5c (KDM5C, also known as JARID1C or SMCX), an eraser of histone 3 lysine 4 di and tri-methylation (H3K4me2/3). We found male *Kdm5c* knockout (-KO) mice, which recapitulate key behavioral phenotypes of Claes-Jensen X-linked intellectual disability (MRXSCJ, OMIM: 300534), aberrantly express many liver, muscle, ovary, and testis genes within the amygdala and hippocampus. Many genes expressed in the *Kdm5c*-KO brain were typically unique to germ cells, indicating an erosion of the soma-germline boundary. Using germ cell-depleted RNA sequencing datasets, we curated a list of germline-enriched genes for unbiased characterization of KDM5C in germline gene repression. Germline genes are typically decommissioned in somatic lineages in the post-implantation epiblast, yet *Kdm5c*-KO epiblast-like cells (EpiLCs) aberrantly express key regulators of germline identity and meiosis, including *Dazl* and *Stra8*. We found that while KDM5C promotes the initial placement of DNA methylation at a subset of germline gene promoters in EpiLCs, germline genes can also become activated independent of direct KDM5C regulation. Intriguingly, many of the germline transcripts expressed in the *Kdm5c*-KO brain were late-stage spermatogenesis genes that are not directly bound by KDM5C. This suggests germline developmental programs can progress ectopically in the background of typical *Kdm5c*-KO development. Ultimately, this work provides novel insight into the demarcation of somatic and germline lineages in mammals while also characterizing the cellular identity crisis within a chromatin-linked neurodevelopmental disorder.

## 28 Introduction

29 A single genome holds the instructions to generate the myriad of cell types found within the adult organism.  
30 This is, in part, accomplished by chromatin regulators that can either promote or impede lineage-specific  
31 gene expression through DNA and histone modifications<sup>1,2</sup>. While many chromatin regulators were initially  
32 identified for their roles in shaping cellular and tissue identity<sup>3–5</sup>, recent advancements in next generation  
33 sequencing unexpectedly revealed many neurodevelopmental disorders (NDDs) are caused by mutations in  
34 chromatin regulators<sup>6</sup>. Several studies have suggested this connection between chromatin regulators and  
35 neurodevelopment is due to their regulation of brain-specific genes, such as orchestrating transcriptional  
36 programs for synaptic maturation<sup>7</sup> and transitioning between neuronal and glial fates during neural precursor  
37 differentiation<sup>8</sup>. However, loss of some chromatin regulators can also result in the ectopic transcription of  
38 tissue-specific genes outside of their target environment, such as the misexpression of liver-specific genes  
39 within adult neurons<sup>9</sup>. Very few studies have investigated this cellular identity crisis in chromatin-linked  
40 NDDs<sup>9,10</sup> and it is currently unknown if partial loss of brain identity contributes to neurodevelopmental  
41 impairments.

42 To elucidate the role of tissue identity in chromatin-linked neurodevelopmental disorders, it is essential  
43 to first characterize the types of genes dysregulated and the molecular mechanisms governing their de-  
44 repression. In this study, we characterized the aberrant expression of tissue-enriched genes with loss of  
45 lysine demethylase 5C (KDM5C). KDM5C, , also known as SMCX or JARID1C, is a chromatin regulator that  
46 can repress gene expression through erasure of histone 3 lysine 4 di- and trimethylation (H3K4me2/3)<sup>11</sup>,  
47 marks enriched at active gene promoters. Pathogenic mutations in *KDM5C* cause Intellectual Developmental  
48 Disorder, X-linked, Syndromic, Claes-Jensen Type (MRXSCJ, OMIM: 300534), whose predominant features  
49 include intellectual disability, seizures, aberrant aggression, and autistic behaviors<sup>12–14</sup>. *Kdm5c* knockout  
50 (-KO) mice recapitulate key MRXSCJ patient phenotypes, including hyperaggression, increased seizure  
51 propensity, and learning impairments<sup>10,15</sup>. Unexpectedly, RNA sequencing (RNA-seq) of the *Kdm5c*-KO  
52 hippocampus revealed ectopic expression of testis genes within the brain<sup>10</sup>. It is currently unknown what types  
53 of testis genes are dysregulated, when in *Kdm5c*-KO development this begins, and if other tissue-specific  
54 genes are also aberrantly expressed with KDM5C loss.

55 The testis contains both germ cells (cells that pass on their genetic material, e.g. sperm) and somatic  
56 cells (cells that perform all other bodily functions). Distinguishing between the germline and the soma is a  
57 key feature of multicellularity and that typically occurs during early embryogenesis. In mammals, chromatin  
58 regulators play a key role in decommissioning germline genes in somatic cells during the transition from naïve  
59 to primed pluripotency. Initially, repressive histone H2A lysine 119 monoubiquitination (H2AK119ub1)<sup>16</sup>,  
60 histone 3 lysine 9 trimethylation (H3K9me3)<sup>16,17</sup>, are placed at germline gene promoters in embryonic stem  
61 cells and are then decorated with DNA CpG methylation<sup>17–19</sup> in the post-implantation embryo. How KDM5C  
62 interacts with known germline gene silencers during this time point is currently unknown. Additionally, most

63 studies have focused on select genes important for early germ cell development rather than germline-enriched genes as a whole given the lack of a curated list of germline-enriched genes. Thus, it is unclear if the mechanism of repression differs for certain classes of germline genes, e.g. meiotic genes versus spermatid differentiation genes. Systematically characterizing KDM5C's role in germline gene repression during early embryogenesis will unveil key mechanisms underlying the demarcation between soma and germline identity and while also providing molecular footholds to test the impact of ectopic germline genes on neurodevelopment.

70 To illuminate KDM5C's role in tissue identity, here we characterized the aberrant transcription of tissue-enriched genes within the *Kdm5c*-KO brain and epiblast-like stem cells (EpiLCs), an *in vitro* model of the post-implantation embryo. We observed general dysregulation of tissue-enriched genes in both the *Kdm5c*-KO brain and EpiLCs, including misexpression of genes typically highly enriched within the testis, liver, muscle, and ovary. Both the *Kdm5c*-KO amygdala and hippocampus had significant enrichment of testis genes. We demonstrated testis and ovary-enriched genes are germline genes by generating a list of germline enriched-genes using germ cell-depleted RNA sequencing datasets. Using this curated list, we found *Kdm5c*-KO EpiLCs aberrantly expressed key drivers of germline identity and meiosis, including *Dazl* and *Stra8*, while the *Kdm5c*-KO brain primarily expressed germline genes important for late spermatogenesis. Additionally, KDM5C was highly enriched at germline gene promoters in EpiLCs but only bound to a subset of germline genes expressed in *Kdm5c*-KO cells, indicating germline-enriched mRNAs can be aberrantly transcribed through indirect mechanisms. Finally, we found KDM5C promotes the long-term silencing of germline genes in somatic cells by aiding the placement of DNA CpG methylation in EpiLCs. Thus, we propose KDM5C plays a fundamental role in the development of tissue identity during early embryogenesis, including the establishment of the soma-germline boundary.

## 85 Results

### 86 **Tissue-enriched genes, including testis genes, are aberrantly expressed in the 87 *Kdm5c*-KO brain**

88 Previous RNA sequencing (RNA-seq) in the adult hippocampus revealed ectopic expression of some testis genes within the *Kdm5c* knockout (-KO) brain<sup>10</sup>. Since tissue-enriched genes have not been systematically characterized in the *Kdm5c*-KO brain, if the testis is the only tissue type misexpressed and if loss of tissue identity a major consequence of KDM5C loss. Therefore, we globally assessed the expression of genes enriched in 17 mouse tissues<sup>20</sup> in our published mRNA-seq datasets of the adult amygdala and hippocampus in mice with constitutive knockout of *Kdm5c*<sup>21</sup>.

94 We found a large proportion of genes that are significantly upregulated within the *Kdm5c*-KO brain (DESeq2<sup>22</sup>, log2 fold change > 0.5, q < 0.1) are enriched in non-brain tissues in wild-type mice (Amygdala:

96 35%, Hippocampus: 24%) (Figure 1A-B). The majority of tissue-enriched differentially expressed genes  
97 (tissue-enriched DEGs) were testis genes (Figure 1A-C). Even though the testis has the largest total number  
98 of tissue-biased genes compared to any other tissue (2,496 genes), testis-biased DEGs were significantly  
99 enriched for both brain regions (Amygdala  $p = 1.83e-05$ , Odds Ratio = 5.13; Hippocampus  $p = 4.26e-11$ ,  
100 Odds Ratio = 4.45, Fisher's Exact Test). One example of a testis-enriched DEG is *FK506 binding protein 6*  
101 (*Fkbp6*), a known regulator of piRNA expression and meiosis in germ cells<sup>23,24</sup> (Figure 1C).

102 In addition to the high enrichment of testis genes, we also identified aberrant expression of other tissue-  
103 enriched genes within the *Kdm5c*-KO brain. Although the *Kdm5c*-KO mice we examined were male, we  
104 observed significant enrichment of ovary-biased DEGs in both the amygdala and hippocampus (Amygdala  
105  $p = 0.00574$ , Odds Ratio = 18.7; Hippocampus  $p = 0.048$ , Odds Ratio = 5.88, Fisher's Exact) (Figure 1D).  
106 Ovary-enriched DEGs included *Zygotic arrest 1* (*Zar1*), sequesters mRNAs in oocytes for meiotic maturation  
107 and early zygote development<sup>25</sup> (Figure 1D). Although not consistent across brain regions, we also found  
108 significant enrichment of DEGS biased towards two non-gonadal tissues - the liver (Amygdala  $p = 0.0398$ ,  
109 Odds Ratio = 6.58, Fisher's Exact Test) and the muscle (Hippocampus  $p = 0.0104$ , Odds Ratio = 6.95,  
110 Fisher's Exact Test). An example of a liver-biased DEG dysregulated in both the hippocampus and amygdala  
111 is *Apolipoprotein C-I* (*Apoc1*), a lipoprotein metabolism and transport gene<sup>26</sup> (Figure 1E). Testis, ovary,  
112 and liver-enriched DEGs showed little to no expression in the developing and adult wild-type brain, yet our  
113 mRNA-seq data indicate they are polyadenylated and spliced into mature transcripts (Figure 1C-E). Of  
114 note, we did not observe enrichment of brain-enriched genes (Amygdala  $p = 1$ ; Hippocampus  $p = 0.74$ ,  
115 Fisher's Exact), despite the fact these are brain samples and the brain has the second highest total number  
116 of tissue-enriched genes (708 genes). Together, these results suggest misexpression of tissue-enriched  
117 genes within the brain is a major effect of KDM5C loss.

## 118 **Germline genes are misexpressed in the *Kdm5c*-KO brain**

119 • Intriguingly, some of the ectopic testis transcripts identified within the *Kdm5c*-KO hippocampus have  
120 known functions unique to germ cells<sup>10</sup>, suggesting KDM5C may play a role in demarcating somatic  
121 versus germline identity.

122 The testis contains both germ cells (meiotic cells, e.g. spermatogonia) and somatic cells (non-meiotic,  
123 e.g. Leydig cells) that support hormone production and germline functions. To determine if *Kdm5c*-KO  
124 testis-enriched DEGs are somatic or germline genes, we first assessed their known functions through  
125 gene ontology analysis. We found *Kdm5c*-KO testis-enriched DEGs high enrichment of germline-relevant  
126 ontologies, including spermatid development (GO: 0007286,  $p.adjust = 6.2e-12$ ) and sperm axoneme  
127 assembly (GO: 0007288,  $p.adjust = 2.45e-14$ ) (Figure 2A).

128 To further validate if these testis DEGs are truly germline genes, we then evaluated their expression  
129 in somatic versus germ cells within the testis. We first compared their expression in the testis with germ

130 cell depletion<sup>27</sup>, which was accomplished by heterozygous *W* and *Wv* mutations in the enzymatic domain  
131 of *c-Kit* (*Kit*<sup>W/Wv</sup>) that prevent the maturation of germ cells<sup>28</sup>. Almost all *Kdm5c*-KO testis-enriched DEGs  
132 lost expression with germ cell depletion (Figure 2B). We then assessed testis-enriched DEG expression  
133 in a published single cell RNA-seq dataset that identified cell type-specific markers within the testis<sup>29</sup>. We  
134 found that while some testis-enriched DEGs were classified as specific markers for different germ cell types  
135 (e.g. spermatogonia, spermatocytes, round spermatids, and elongating spermatids) none marked somatic  
136 cells (Figure 2C). Together, these data demonstrate that the *Kdm5c*-KO brain aberrantly expresses germline  
137 genes, reflecting an erosion between somatic versus germline identity.

138 We then wanted to more deeply characterize germline gene misexpression with *Kdm5c* loss, but lacked  
139 a comprehensive list of mouse germline-enriched genes. To facilitate downstream analyses, we generated  
140 a list of germline-enriched genes using currently available RNA-seq datasets in *Kit*<sup>W/Wv</sup> mice. Wild-type  
141 and *Kit*<sup>W/Wv</sup> datasets included males and females at embryonic day 12, 14, and 16<sup>30</sup>, as well as adult  
142 male testes<sup>27</sup>. We defined genes as germline-enriched if their expression met the following criteria: 1)  
143 their expression is greater than 1 FPKM in wild-type gonads 2) their expression in any adult wild-type,  
144 non-gonadal tissue<sup>20</sup> does not exceed 20% of their maximum expression in the wild-type germline, and  
145 3) their expression in the germ cell-depleted germline, for any sex or time point, does not exceed 20% of  
146 their maximum expression in the wild-type germline. These criteria yielded 1,288 germline-enriched genes  
147 (Figure 2D), which was hereafter used as a resource to globally assess germline gene misexpression with  
148 *Kdm5c* loss (Supplementary table 1).

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

151 Germ cells are typically distinguished from somatic cells soon after the embryo implants into the uter-  
152 ine wall<sup>31,32</sup> when a subset of epiblast stem cells become the primordial germ cells (PGCs) while the  
153 remainder differentiate into the ectoderm, mesoderm, and endoderm to form the somatic tissues<sup>33</sup>. This  
154 developmental time point can be modeled *in vitro* through differentiation of embryonic stem cells (ESCs) into  
155 post-implantation epiblast-like stem cells (EpiLCs) (Figure 3A, top). While some germline-enriched genes  
156 are also expressed in embryonic stem cells (ESCs) and in the 2-cell stage<sup>34–36</sup>, they are silenced as they  
157 differentiate into EpiLCs<sup>17</sup>. Therefore, we tested if KDM5C was necessary for silencing germline genes in  
158 the post-implantation embryo by evaluating the impact of *Kdm5c* loss in EpiLCs.

159 We first identified *Kdm5c*-KO EpiLC DEGs in our previously published RNA-seq dataset<sup>37</sup> (DESeq2, log2  
160 fold change > 0.5, q < 0.1) and assessed tissue-enriched gene expression. Similar to the *Kdm5c*-KO brain,  
161 we observed general dysregulation of tissue-enriched genes, with the largest number of genes belonging  
162 to the brain and testis, although they were not significantly enriched (Figure 3B). Although we observed  
163 aberrant expression of many tissue-enriched genes, including 54 germline-enriched genes, we did not find

any significant difference in primed pluripotency genes, such as *Dnmt3b*, *Fgf5*, *Pou3f1*, and *Otx2* (Figure 3C). We also did not observe any gross changes in *Kdm5c*-KO cell morphology during ESC to EpiLC differentiation (Figure 3D), altogether indicating KDM5C loss does not impair EpiLC formation.

To determine if germline DEGs are constitutively dysregulated or if they can change over the course of development, we compared EpiLC germline DEGs to those expressed in the *Kdm5c*-KO brain. We found the majority of germline DEGs expressed in EpiLCs and the brain were unique, with only *Cyct* shared across all sequencing datasets (Figure 3E-F). We then evaluated the known functions of EpiLC and brain germline DEGs and found particularly high enrichment of meiosis-related gene ontologies in EpiLCs (Figure 3G), such as meiotic cell cycle process (GO: 1903046, p.adjust = 1.59e-08) and meiotic nuclear division (GO: 0140013, p.adjust = 9.76e-09). While there was modest enrichment of meiotic gene ontologies in both brain regions, the *Kdm5c*-KO hippocampus primarily expressed late-stage spermatogenesis genes, such those involved in the sperm axoneme structure.

Interestingly, DEGs unique to *Kdm5c*-KO EpiLCs included key drivers of germline identity, such as *Stimulated by retinoic acid 8 (Stra8)* and *Deleted in azoospermia like (Dazl)* (Figure 3H). These genes are typically expressed during embryonic germ cell development to commit PGCs to the germline fate, but are also expressed later in life to trigger meiotic gene expression programs<sup>38–40</sup>. Of note, some germline genes, including *Dazl*, are also expressed in the two-cell embryo<sup>35,41</sup>. However, we did not see misexpression of two-cell embryo-specific genes, like *Duxf3 (Dux)* ( $q = 0.337$ ) and *Zscan4d* ( $q = 0.381$ ), indicating *Kdm5c*-KO in EpiLCs do not revert back to a 2-cell state (Figure 3H).

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

Previous work suggests KDM5C represses germline genes during early development, as re-expression of KDM5C in knockout neuronal cultures fails to suppress two hippocampal germline DEGs<sup>10</sup>. There is some evidence KDM5C binds to select germline gene promoters in ESCs but is lost in neurons<sup>10</sup>. However, the lack of a comprehensive list of germline-enriched genes prohibited systematic characterization of KDM5C binding at germline gene promoters. Thus, it is currently unclear if KDM5C is enriched at germline gene promoters, what types of germline genes KDM5C regulates, and if its binding is maintained at any germline genes in neurons.

To further characterize KDM5C's role in germline gene silencing, we analyzed KDM5C chromatin immunoprecipitation followed by DNA sequencing (ChIP-seq) datasets in EpiLCs<sup>37</sup> and primary neuron cultures (PNCs) from the cortex and hippocampus<sup>15</sup>. EpiLCs had a higher total number of KDM5C peaks than PNCs (EpiLCs: 5,808, PNCs: 1,276, MACS2  $q < 0.1$  and fold enrichment  $> 1$ , removal of *Kdm5c*-KO false positive peaks) and KDM5C was primarily localized to gene promoters in both cell types (EpiLCs: 4,190, PNCs: 745 +/- 500bp from the TSS), although PNCs showed increased localization to non-promoter regions (Figure 4A).

The majority of promoters bound by KDM5C in PNCs were also bound in EpiLCs (513 shared promoters),

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

We then compared KDM5C binding at RNA-seq DEG promoters to determine if the germline mRNAs expressed in the *Kdm5c*-KO brain and EpiLCs are direct targets of KDM5C (Figure 4F). About one third of EpiLC-specific and brain-specific (hippocampus, amygdala, or both) germline DEGs were bound by KDM5C in EpiLCs (EpiLC only: 36%, Brain only: 33.3%). Again, we did not observe any KDM5C binding at germline gene promoters in PNCs, even for brain-specific DEGs (Figure 4H). Some notable differences in KDM5C binding for EpiLC-specific DEGs included *Dazl* and *Stra8*, two key drivers of germline identity discussed above. Although both mRNAs are expressed in *Kdm5c*-KO EpiLCs, KDM5C is only bound to *Dazl*'s promoter and not *Stra8*'s in EpiLCs (Figure 4F). In contrast to the unique DEGs, 3 out of the 4 genes dysregulated in both the brain and EpiLCs are bound by KDM5C around the TSS (Figure 4G). Altogether, this suggests KDM5C decommissions germline genes in EpiLCs but is no longer recruited to germline gene promoters in neurons. Furthermore, majority of germline mRNAs expressed in *Kdm5c*-KO cells are dysregulated independent of direct KDM5C recruitment to their promoters in EpiLCs.

## KDM5C promotes *de novo* DNA methylation at germline genes

KDM5C is generally thought to suppress transcription through erasure of histone 3 lysine 4 di- and trimethylation (H3K4me2/3)<sup>11</sup>.

recent studies in ESCs have suggested KDM5C's repression *Dazl* is independent of its catalytic activity<sup>41</sup>. Somatic repression of germline genes is typically established during the transition between naïve and primed pluripotency, which modeled *in vitro* as ESCs to EpiLC differentiation. In ESCs, chromatin regulators place repressive histone modifications at germline gene promoters, including histone 2A 119 monoubiquitination (H2AK119ub1) and histone 3 lysine 9 trimethylation (H3K9me3)<sup>16,17,42</sup>. Germline genes are then silenced long-term in EpiLCs by *de novo* placement of DNA CpG methylation (DNAme)<sup>17</sup>. Although KDM5C's catalytic activity is not required to suppress *Dazl* in ESCs, KDM5C may promote long-term germline gene silencing via H3K4me3 removal since H3K4me3 can impede DNAme placement<sup>43,44</sup> and DNAme is

234 lost at select germline gene promoters in the hippocampus<sup>10</sup>. Because KDM5C's role in germline gene  
235 repression has only been characterized in ESCs and in the mature brain, it is currently unclear to what extent  
236 KDM5C is involved during transition between ESCs and EpiLCs and if its catalytic activity is required for  
237 long-term silencing.

238 To elucidate KDM5C's role in germline gene silencing, we first characterized KDM5C's substrates, histone  
239 3 lysine 4 di- and trimethylation (H3K4me2/3) in our previously published ChIP-seq datasets in wild type and  
240 *Kdm5c*-KO amygdala<sup>21</sup> and EpiLCs<sup>37</sup>. In congruence with previous work in the *Kdm5c*-KO hippocampus<sup>10</sup>,  
241 we observed aberrant accumulation of H3K4me3 around the transcription start site (TSS) of germline genes  
242 in the *Kdm5c*-KO amygdala (Figure 5A). We additionally found a marked increase in H3K4me2 around the  
243 TSS of germline genes in *Kdm5c*-KO EpiLCs (Figure 5B).

244 We then evaluated KDM5C's embryonic role in germline gene silencing during ESC to EpiLC differentiation.  
245 We first characterized KDM5C's mRNA and protein expression at 0, 24, and 48 hours of differentiation  
246 (Figure 5C). While *Kdm5c* mRNA steadily decreased from 0 to 48 hours of differentiation (Figure 5D), KDM5C  
247 protein initially increased from 0 to 24 hours but then decreased to near knockout levels by 48 hours (Figure  
248 5E). We then performed whole genome bisulfite sequencing in wild-type and *Kdm5c*-KO ESCs and 96 hour  
249 extended EpiLCs (exEpiLCs) to determine KDM5C's role in the initial placement of DNA methylation at  
250 germline gene promoters. While wild-type cells accumulated high levels of DNA methylation at germline  
251 gene promoters over the course of ESCs to exEpiLC differentiation, DNA methylation was markedly reduced  
252 in *Kdm5c*-KO exEpiLCs (Figure 5F).

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

## 258 Discussion

259 In the above study, we demonstrate KDM5C's pivotal role in the development of tissue identity, resulting in  
260 substantial misexpression of tissue-enriched genes within the adult *Kdm5c*-KO hippocampus and amygdala.  
261 We defined tissue-enriched genes based on the criteria used in a previously published dataset of 17 C57/Bl6J  
262 mouse tissues<sup>20</sup>, which defined genes as tissue-enriched if they had more than 4-fold higher expression  
263 when compared to any other tissue. In addition to testis genes identified previously<sup>10</sup>, we found significant  
264 enrichment of muscle, liver, and even ovary-biased genes aberrantly expressed within the male *Kdm5c*-KO  
265 brain. It is currently unclear what the impact of these tissue-enriched genes are upon *Kdm5c*-KO brain

266 function and if they ultimately contribute to MRXSCJ-related phenotypes. While the majority of tissue-  
267 enriched DEGs were testis and ovary genes with no known brain functions, select liver and muscle-biased  
268 DEGs do have known roles in the wild-type brain. For example, *Apolipoprotein C-I* (*Apoc1*) is a lipid transport  
269 gene highly enriched within the liver, but is also lowly expressed in the wild-type brain. Intriguingly, *Apoc1*  
270 overexpression in the mouse brain can impair learning and memory<sup>45</sup> and is implicated in Alzheimer's  
271 disease in humans<sup>46</sup>. Our results suggest KDM5C fine-tunes the expression of tissue-enriched genes like  
272 *Apoc1* to match the level required for proper brain function. However, further studies are required to determine  
273 if these genes contribute to *Kdm5c*-KO neuronal impairments and MRXSCJ pathology.

274 A substantial portion of genes upregulated in *Kdm5c*-KO brain are testis genes that have no known  
275 function within the brain. Through the use of publicly available RNA-seq datasets, we demonstrated these  
276 testis-enriched DEGs are typically unique to germ cells. Misexpression of germline genes in the brain sug-  
277 gests *Kdm5c*-KOs fail to demarcate between somatic and germline lineages, a key feature of multicellularity  
278 and sexual reproduction. Previous studies have demonstrated germline genes are decommissioned in  
279 somatic cells by multiple chromatin-modifying enzymes, but have primarily focused on the repression of  
280 key marker genes. To globally characterize KDM5C's role in germline gene repression, we curated a list  
281 of mouse germline-enriched genes using publicly available germ cell-depleted RNA-seq datasets. This  
282 resource enabled us to identify 1) the extent of germline gene dysregulation in *Kdm5c*-KO cells, 2) the types  
283 of germline genes misexpressed at different developmental time points, and 3) which types of germline  
284 genes are directly or indirectly regulated by KDM5C.

285 Dysregulation of *Kdm5c*-KO tissue identity begins during early embryogenesis, as germline and other  
286 tissue-enriched genes are aberrantly expressed in epiblast-like cells (EpiLCs). *In vivo*, germline genes  
287 are typically decommissioned in epiblast stem cells and remain silenced as the epiblast differentiates  
288 into the body's somatic tissues<sup>33</sup>. However, a small subset of epiblast stem cells will receive signals to  
289 reactivate germline gene expression to become the primordial germ cells (PGCs) that will ultimately form  
290 the mature germline<sup>31,32</sup>. This process can be mimicked *in vitro* by differentiating EpiLCs into primordial  
291 germ cell-like cells (PGCLCs)<sup>47</sup>. Thus, misexpression of germline genes in *Kdm5c*-KO EpiLCs might  
292 suggest they are progressing beyond EpiLC differentiation and becoming PGCLCs. Yet, *Kdm5c*-KO EpiLCs  
293 had proper expression of EpiLC marker genes for primed pluripotency and we observed no difference in  
294 cellular morphology during ESC to EpiLC differentiation. Furthermore, we saw no significant change in *Otx2*  
295 expression, an epiblast stem cell marker that is known to repress epiblast and EpiLC differentiation into PGC  
296 and PGCLCs<sup>48</sup>. This, in conjunction with *Kdm5c*-KO mice being viable, suggests germline gene expression  
297 is occurring ectopically in conjunction with typical developmental programs, rather than a complete shift to  
298 germline identity.

299 • change first sentence: Other chromatin regulators involved in germline gene repression act in ESCs  
300 and EpiLCs. We observed this is true for KDM5C

301 We then globally characterized KDM5C binding at germline-enriched gene promoters through analysis

302 of KDM5C ChIP-seq in EpiLCs and primary hippocampus and cortex neuron cultures (PNCs). While we  
303 observed high enrichment of KDM5C peaks at germline genes in EpiLCs, KDM5C was not bound to germline  
304 gene promoters in PNCs. This suggests dysregulation of germline genes in the mature *Kdm5c*-KO brain is  
305 due to loss of repression during embryogenesis, which is consistent with previous work that found introducing  
306 human KDM5C into *Kdm5c*-KO PNCs does not repress germline transcripts<sup>10</sup>. Although KDM5C is enriched  
307 at germline gene promoters in EpiLCs, KDM5C was only bound to about a third of EpiLC germline-enriched  
308 DEGs. One notable DEG not bound by KDM5C is *Stra8*, a transcription factor activated by retinoic acid  
309 signaling in germ cells that promotes meiotic initiation<sup>49,50</sup>. Retinoic acid can only activate *Stra8* expression  
310 when DAZL is present, which is a direct target of KDM5C and is aberrantly transcribed and translated in  
311 *Kdm5c*-KO EpiLCs. This indicates germline genes can be aberrantly expressed in *Kdm5c*-KO cells indirectly  
312 of KDM5C regulation through activation by other ectopic germline programs. Consistent with this idea, many  
313 *Kdm5c*-KO EpiLC germline DEGs are important for early germ cell development and meiosis while those  
314 expressed in the mature *Kdm5c*-KO brain are involved in late sperm development. Altogether, this indicates  
315 ectopic germline programs are, to some extent, progressing through germ cell developmental stages over  
316 the course of *Kdm5c*-KO development.

- 317 • Talk about different motifs if we do see differences there and if it explains direct vs indirect dysregulation  
318 – Could also mention PRC1.6/KDM5C binds pcgf6 in ESCs  
319 • Maybe talk about other regulators/if anything is known about them long term? Or save for 2nd paper.  
320 – another germline repressor: <https://pubmed.ncbi.nlm.nih.gov/35345626/>  
321 • Might be good to look at for retinoic acid paper (WT germ expression): <https://journals.plos.org/plosone/article?id=10.1371/journal.pone.0005654>

323 It is important to note that while we focused on KDM5C's regulation of germline genes, some germline-  
324 enriched genes are also expressed in naïve ESCs and at the 2-cell stage due to their roles in pluripotency  
325 and self-renewal. For example, although primarily known for committing PGCs to the germline fate and  
326 regulating the translation of germline-specific RNAs, *Dazl* is also expressed in naïve ESCs *in vitro*<sup>34</sup>, the  
327 inner cell mass *in vivo*<sup>34</sup>, and at the 2-cell stage<sup>51</sup>, but is silenced when cells transition from naïve to primed  
328 pluripotency during ESC to EpiLC differentiation<sup>17</sup>. Very recently, two screens of *Dazl*-repressors in ESCs  
329 identified KDM5C as a direct regulator of *Dazl* expression<sup>41,52</sup>. Interestingly, one screen found *Kdm5c*-KO  
330 ESCs also aberrantly expressed 2-cell-specific genes, indicating KDM5C negatively regulates totipotency in  
331 ESCs<sup>41</sup>. We found that while KDM5C also represses *Dazl* expression in EpiLCs, *Kdm5c*-KO EpiLCs do not  
332 express 2-cell specific genes like *Dux* and *Zscan4c*. Out of the four 2-cell regulators characterized in ESCs,  
333 KDM5C was the only factor whose repression of *Dazl* was independent of *Dux* expression<sup>41</sup>. Together, this  
334 suggests the 2-cell-like state in *Kdm5c*-KO ESCs is an indirect consequence of KDM5C's primary role in  
335 germline gene repression, including germline genes that are also involved in pluripotency and self-renewal.

336 It is hypothesized distinct repressive chromatin marks are placed at germline gene promoters as the  
337 embryo transitions from naïve to primed pluripotency. Initially, germline genes are repressed by placement of  
338 histone 2A lysine 119 monoubiquitination (H2AK119ub1) and histone 3 lysine 9 trimethylation (H3K9me3)  
339 in ESCs and then gain *de novo* DNA CpG methylation (CpGme) in EpiLCs<sup>17–19,42,53,54</sup>. KDM5C may be  
340 instead required to remove an active mark, histone 3 lysine 4 trimethylation (H3K4me3), since H3K4me3  
341 can impede *de novo* CpGme placement<sup>43,44</sup>. This is supported by previous work in the *Kdm5C*-KO adult  
342 hippocampus, which found an increase in H3K4me3 around the transcription start site (TSS) of germline  
343 DEGs and loss of CpGme at at least two germline promoters<sup>10</sup>. However, KDM5C's role in embryonic  
344 germline gene silencing is still unclear, as KDM5C's catalytic activity was recently found to be dispensable  
345 for silencing DAZL in ESCs<sup>41</sup>. In this study, we observed a global increase in H3K4me3 around the TSS of  
346 germline-enriched gene in the *Kdm5c*-KO amygdala and an increase in H3K4me2 in *Kdm5c*-KO EpiLCs. We  
347 found KDM5C's expression is dynamically regulated during ESC to EpiLC differentiation and loss of KDM5C  
348 leads to impaired placement of CpGme in extended EpiLCs. Altogether, this suggests KDM5C is necessary  
349 during the transition from naïve to primed pluripotency to promote the initial placement of CpGme at CpG  
350 islands for the long-term silencing of germline genes.

- 351 • Talk about impact - Other germline gene chromatin regulators, NDDs, cancer. Maybe talk about myt1l  
352 • include cancer in there somewhere (Somatic misexpression of germline genes has been implicated in  
353 many cancers.)

354 In summary, we demonstrate loss of the X-linked intellectual disability gene KDM5C leads to widespread  
355 dysregulation of brain tissue identity, including misexpression of germline genes in the somatic brain. In  
356 EpiLCs, KDM5C directly represses key drivers of germline identity like *Dazl*, by promoting intial CpG methy-  
357 lation placement in the post-implantation embryo. However, germline genes can also become ectopically  
358 expresssed in *Kdm5c*-KO cells independent of direct KDM5C regulation, including the meiotic transcription  
359 factor *Stra8*. These ectopic germline developmental programs can, to some extent, mimic typical germ  
360 cell development, resulting in aberrant transcription early developmental and meiotic genes in *Kdm5c*-KO  
361 EpiLCs and late-stage spermatogenesis genes in the *Kdm5c*-KO brain. Altogether, these results define  
362 KDM5C's role in the demarcation between soma and germline identity and offers novel insight into how this  
363 dysregulation of tissue identity changes over the course of development. Additionally, this study provides  
364 the mechanistic foundation required to ultimately investigate the impact of aberrant germline identity upon  
365 neurodevelopment.

366 **Materials and Methods**

367 **Classifying tissue-enriched and germline-enriched genes**

368 Tissue-enriched differentially expresssd genes were determined by their classification in a previously  
369 published list of genes enriched in 17 male and female mouse tissues<sup>20</sup>. This study defined expression as  
370 greater than 1 Fragments Per Kilobase of transcript per Million mapped read (FPKM) and tissue enrichment  
371 as at least 4-fold higher expression than any other tissue.

372 We curated a list of germline-enriched genes using an RNA-seq dataset from wild-type and germline-  
373 depleted (Kit<sup>W/W<sup>v</sup></sup>) male and female mouse embryos from embryonic day 12, 14, and 16<sup>30</sup>, as well as adult  
374 male testes<sup>27</sup>. Germline-enriched genes met the following criteria: 1) their expression is greater than 1  
375 FPKM in wild-type germline 2) their expression in any wild-type somatic tissues<sup>20</sup> does not exceed 20%  
376 of maximum expression in wild-type germline, and 3) their expression in the germ cell-depleted (Kit<sup>W/W<sup>v</sup></sup>)  
377 germline, for any sex or time point, does not exceed 20% of maximum expression in wild-type germline.

378 **Cell culture**

379 We utilized our previously established cultures of male wild-type and *Kdm5c* knockout (-KO) embryonic  
380 stem cells<sup>37</sup>. Sex was confirmed by genotyping *Uba1* on the X and Y chromosomes with the following  
381 primers: 5'-TGGATGGTGTGCCAATG-3', 5'-CACCTGCACGTTGCCCTT-3'. Deletion of *Kdm5c* was  
382 confirmed through the primers 5'-ATGCCCATATTAAGAGTCCTG-3', 5'-TCTGCCTTGATGGACTGTT-3',  
383 and 5'-GGTTCTCAACACTCACATAGTG-3'.

384 Embryonic stem cells (ESCs) were initially cultured in primed ESC (pESC) media consisting of Knock-  
385 Out DMEM (Gibco#10829–018), fetal bovine serum (Gibco#A5209501), KnockOut serum replacement  
386 (Invitrogen#10828–028), Glutamax (Gibco#35050-061), Anti-Anti (Gibco#15240-062), MEM Non-essential  
387 amino acids (Gibco#11140-050), and beta-mercaptoethanol (Sigma#M7522). They were then transitioned  
388 into ground-state “naive” ESCs (nESCs) by culturing for four passages in N2B27 media containing DMEM/F12  
389 (Gibco#11330–032), Neurobasal media (Gibco#21103–049), Gluamax, Anti-Anti, N2 supplement (Invitro-  
390 gen#17502048), and B27 supplement without vitamin A (Invitrogen#12587-010), and beta-mercaptoethanol.  
391 Both pESC and nESC media were supplemented with the GSK3 inhibitor CHIR99021 (Sigma #SML1046-  
392 5MG), the MEK inhibitor PD0325901 (Sigma #PZ0162-5MG), and leukemia inhibitory factor (LIF, Milli-  
393 pore#ESG1107).

394 nESCs were differentiated into epiblast-like cells (EpiLCs, 48 hours) and extendend EpiLCs (exEpiLCs,  
395 96 hours) by culturing in N2B27 media containing DMEM/F12, Neurobasal media, Gluamax, Anti-Anti,  
396 N2 supplement, B27 supplement (Invitrogen#17504044), beta-mercaptoethanol, fibroblast growth factor 2  
397 (FGF2, R&D Biotechne 233-FB), and activin A (R&D Biotechne 338AC050CF).

398 **Immunocytochemistry (ICC)**

399      ICC of DAZL in EpiLCs was performed by washing cells thrice with phosphobuffered saline (PBS), fixing  
400      cells in 4% paraformaldehyde, washing thrice in PBS, blocking in PBS containing 0.3% Triton X-100, and 5%  
401      fetal bovine serum for 1 hour, washing thrice with PBS, and incubating in primary antibody (Rabbit anti DAZL,  
402      abcam ab34139, 1:200) in the blocking solution overnight at 4C. The next day cells were rinsed thrice with  
403      PBS, incubated in secondary antibody (Alexafluor 488 Invitrogen #710369, 1:1000) in blocking buffer, rinsed  
404      thrice in PBS, and then imaged. Images were taken blinded for genotype, chosen based on similar levels of  
405      DAPI signal, and then quantified via ImageJ.

406 **RNA sequencing**

407 **Published datasets**

408      All published datasets are available at the Gene Expression Omnibus (GEO) [https://www.ncbi.nlm.nih.g](https://www.ncbi.nlm.nih.gov/geo/)  
409      ov/geo. Previously published RNA sequencing datasets analyzed in this study included the male wild-type  
410      and *Kdm5c*-KO adult amygdala and hippocampus<sup>21</sup> (available at GEO: GSE127722) and male wild-type and  
411      *Kdm5c*-KO EpiLCs<sup>37</sup> (available at GSE: GSE96797).

412 **Alignment and analysis**

413      After ensuring read quality via FastQC (v0.11.8), reads were then mapped to the mm10 *Mus musculus*  
414      genome (Gencode) using STAR (v2.5.3a), during which we removed duplicates and kept only uniquely  
415      mapped reads. Count files were generated by FeatureCounts (Subread v1.5.0), and BAM files were  
416      converted to bigwigs using deeptools (v3.1.3) and visualized by the UCSC genome browser. RStudio (v3.6.0)  
417      was then used to analyze counts files by DESeq2 (v1.26.0)<sup>22</sup> to identify differentially expressed genes  
418      (DEGs) with a q-value (p-adjusted via FDR/Benjamini–Hochberg correction) less than 0.1 and a log2 fold  
419      change greater than 0.5. For all DESeq2 analyses, log2 fold changes were calculated with IfcShrink using  
420      the ashr package<sup>55</sup>. MA-plots were generated by ggpibr (v0.4.0), and Eulerr diagrams were generated by  
421      eulerr (v6.1.1). Boxplots and scatterplots were generated by ggpibr (v0.6.0) and ggplot2 (v3.3.2). Heatmaps  
422      of gene expression were generated using the base R functions scale and hclust and visualized using the R  
423      package ComplexHeatmap (v2.12.1). The Upset plot was generated via the package UpSetR (v1.4.0)<sup>56</sup>.  
424      Gene ontology (GO) analyses were performed by the R package enrichPlot (v1.16.2) using the biological  
425      processes setting.

426 **Chromatin immunoprecipitation followed by DNA sequencing (ChIP-seq)**

427      We analyzed our previously published KDM5C ChIP-seq experiments in primary neuron cultures from  
428      the cortex and hippocampus<sup>15</sup> (available at GEO: GSE61036) and EpiLCs<sup>37</sup> (available at GEO: GSE96797).

429 ChIP-seq of histone 3 lysine 4 dimethylation in male EpiLCs<sup>37</sup> is also available at GEO: GSE96797. ChIP-seq  
430 of histone 3 lysine 4 trimethylation in the male amygdala<sup>21</sup> are available at GEO: GSE127817.

431 ChIP-seq reads were aligned to mm10 using Bowtie1 (v1.1.2) allowing up to two mismatches. Only  
432 uniquely mapped reads were used for analysis. Peaks were called using MACS2 software (v2.1.0.20140616)  
433 using input BAM files for normalization, with filters for a q-value < 0.1 and a fold enrichment > 1. Common  
434 peak sets were obtained in R via DiffBind (v3.6.5), and count tables for the common peaks were generated  
435 with the Bedtools multicov command. We removed “black-listed” genomic regions that often give aberrant  
436 signals. Peak proximity to genome annotations was determined by ChIPSeeker (v1.32.1). Enriched motifs  
437 were identified using HOMER<sup>57</sup>. Gene ontology (GO) analyses were performed by the R package enrichPlot  
438 (v1.16.2) using the biological processes setting. Average binding across the genome was visualized using  
439 deeptools (v3.1.3). Bigwigs were visualized using the UCSC genome browser.

#### 440 **Whole genome bisulfite sequencing (WGBS)**

#### 441 **Data availability**

#### 442 **Acknowledgements**

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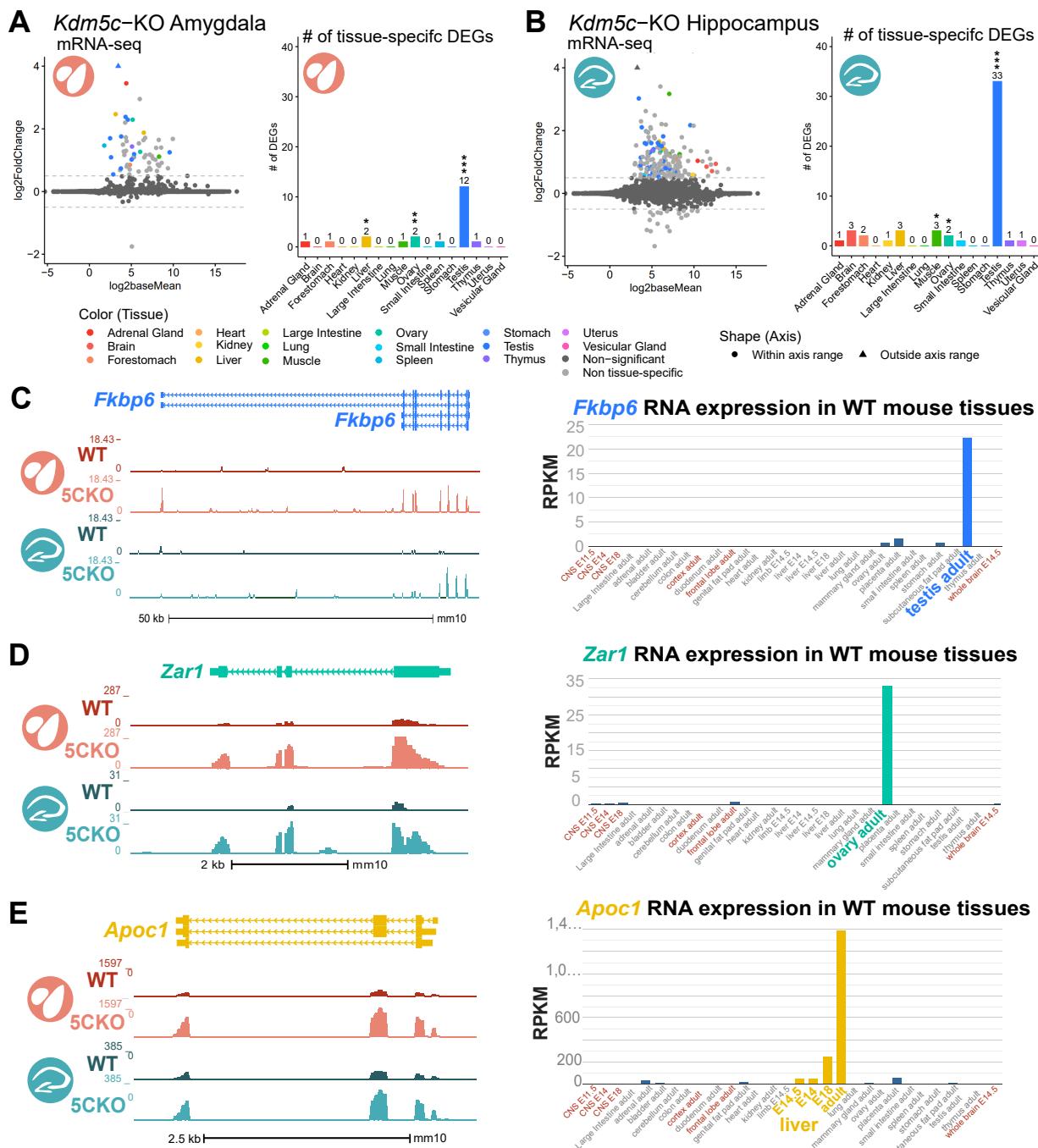
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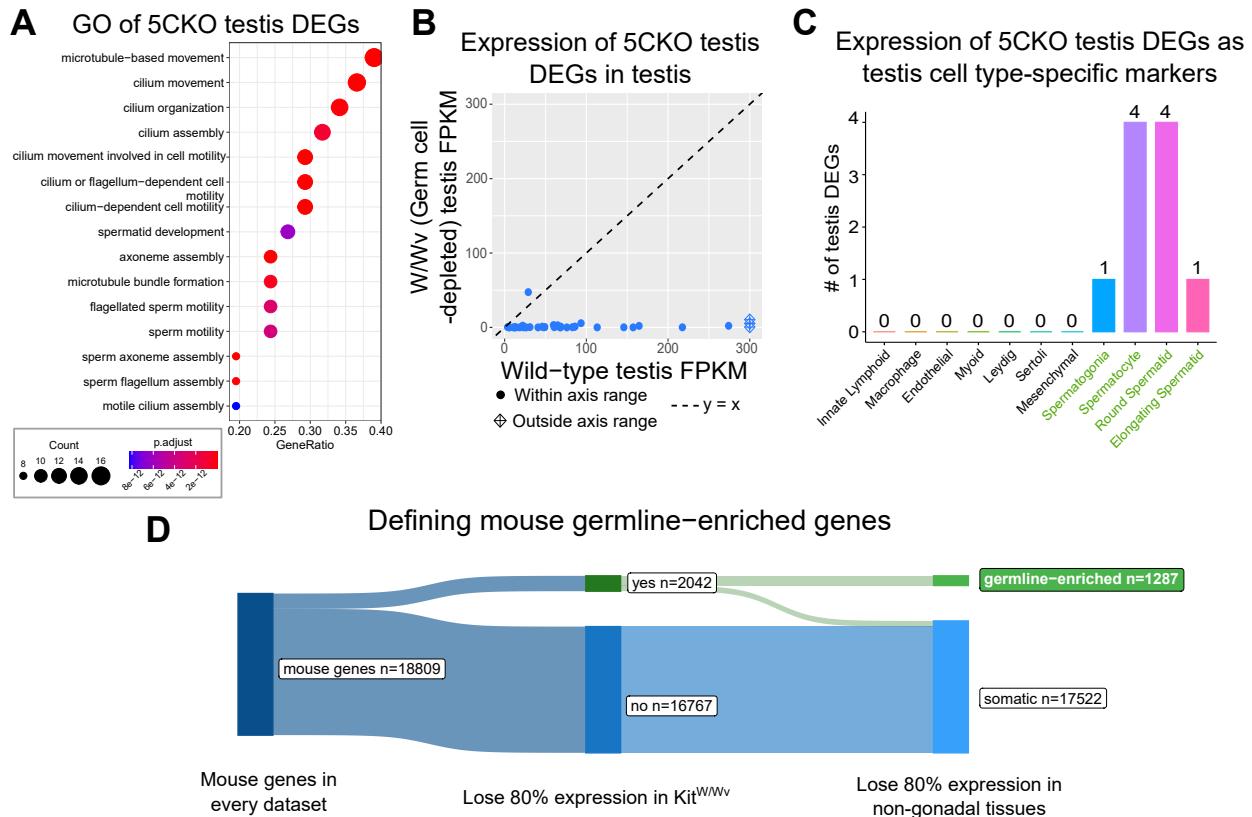
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562 **Figures and Tables**

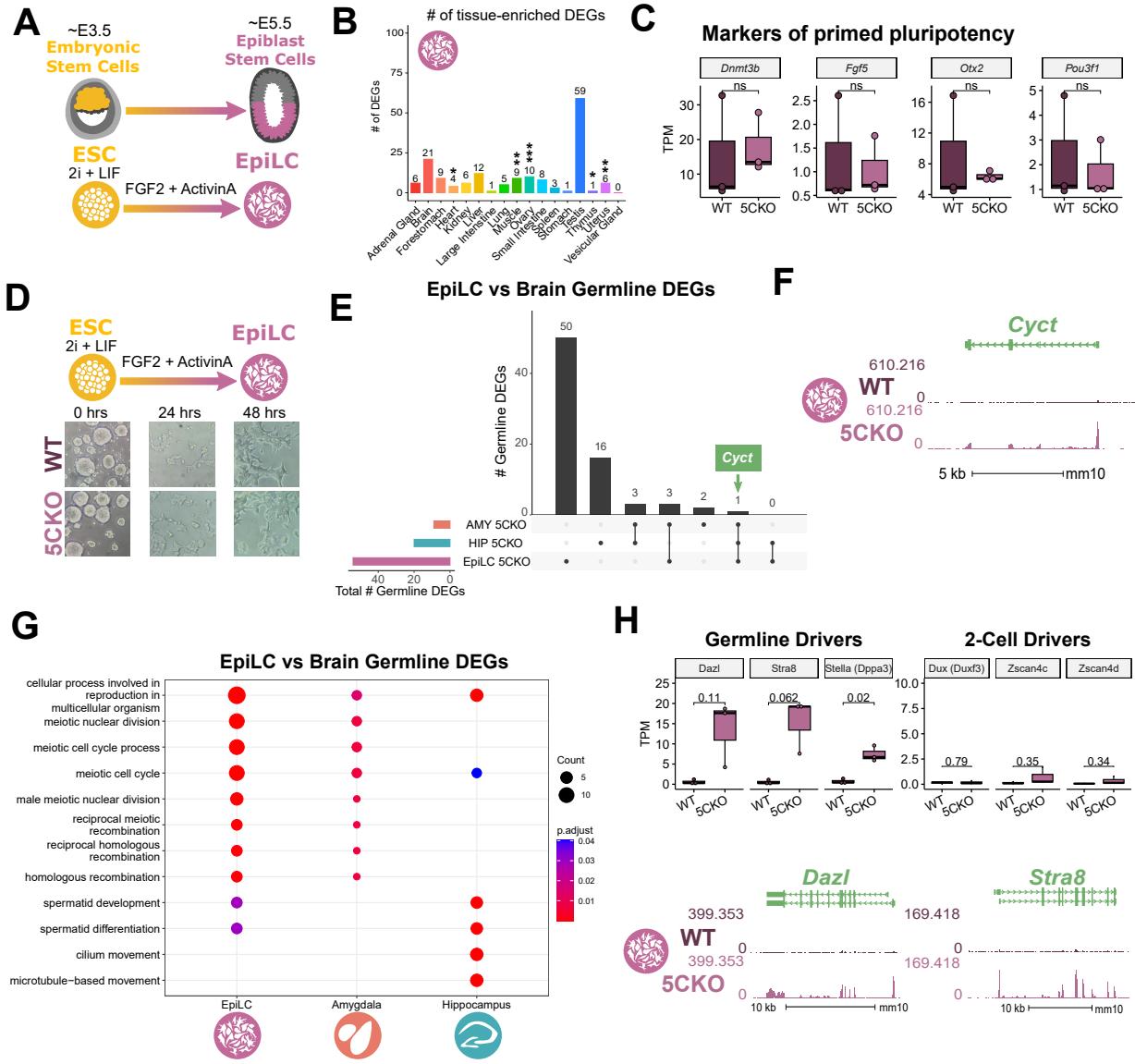
- 563     • Supplementary table 1: list of all germline genes.
- 564       – Columns to include:
- 565           \* KDM5C bound vs not
- 566           \* DEG in EpiLC, brain, both, neither (separate columns?)



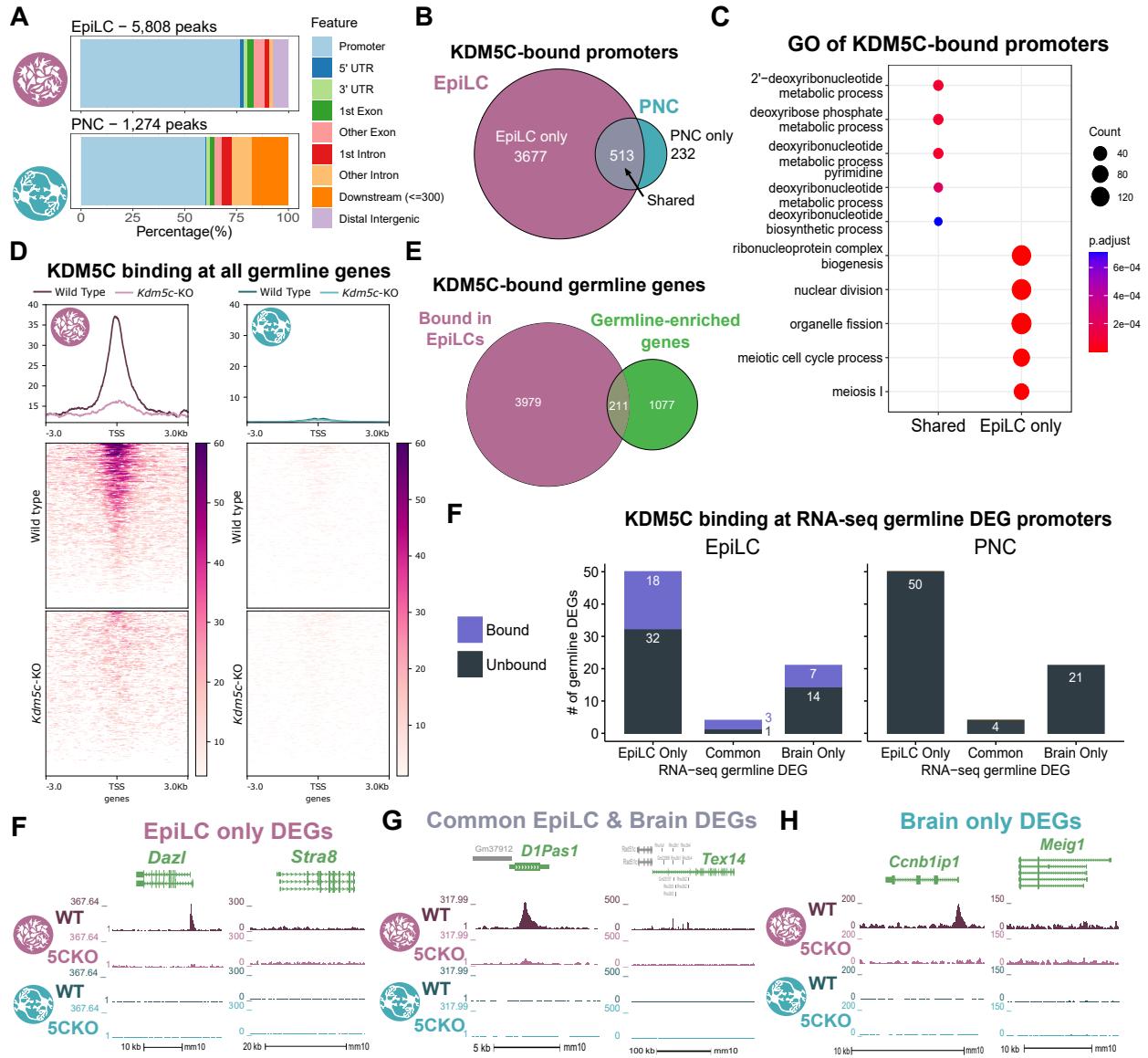
**Figure 1: Tissue-enriched genes are misexpressed in the *Kdm5c*-KO brain.** **A-B.** Expression of tissue-enriched genes (Li et al 2017) in the male *Kdm5c*-KO amygdala (A) and hippocampus (B). Left - MA plot of mRNA-sequencing. Right - Number of tissue-enriched differentially expressed genes (DEGs). \*  $p < 0.05$ , \*\*  $p < 0.01$ , \*\*\*  $p < 0.001$ , 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).

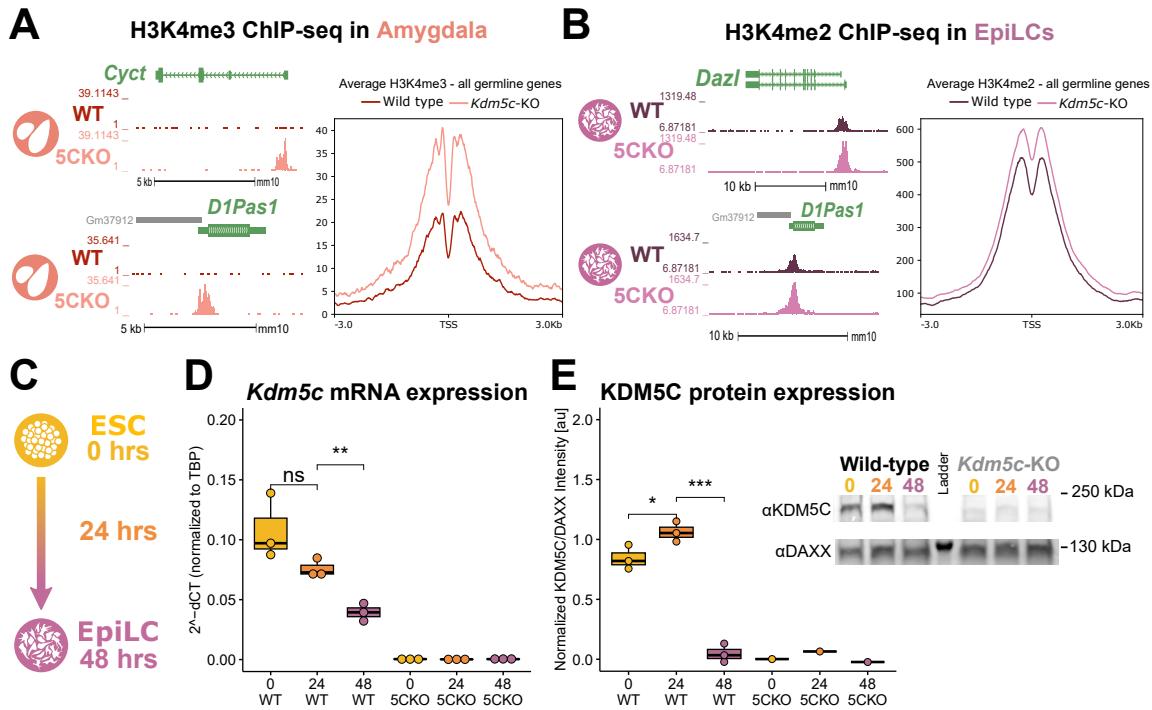


**Figure 3: Kdm5c-KO epiblast-like cells express key drivers of germline identity** **A.** Top - Diagram of *in vivo* differentiation of embryonic stem cells (ESCs) of the inner cell mass into epiblast stem cells. Bottom - *in vitro* differentiation of ESCs into epiblast-like cells (EpiLCs). **B.** Number 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



**F**

**G**

**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 transcripton 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 comparision 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

567 **Notes**

568 **Dazl**

569 We were particularly interested in the aberrant transcription of *Dazl*, since it is essential for germ cell  
570 development and promotes the translation of germline mRNAs<sup>58</sup>. A significant portion of germline transcripts  
571 misexpressed in *Kdm5c*-KO EpiLCs are known binding targets of DAZL, including *Stra8*<sup>59</sup> ( $p = 1.698e-07$ ,  
572 Fisher's Exact Test). This suggests expression of DAZL protein could promote the translation of other  
573 aberrant germline transcripts, influencing their ability to impact *Kdm5c*-KO cellular function. We thus tested  
574 DAZL protein expression in *Kdm5c*-KO EpiLCs through immunocytochemistry (Figure 3I). We observed  
575 about 25% of *Kdm5c*-KO EpiLCs expressed DAZL protein and it was localized to the cytoplasm ( $p = 0.0015$ ,  
576 Welch's t-test), consistent with the pattern of DAZL expression in spermatogonia<sup>59</sup>. Altogether these results  
577 suggest tissue-specific genes are misexpressed during *Kdm5c*-KO embryogenesis, including key drivers of  
578 germline identity that can be translated into protein.

579 **Discussion notes**

- 580 • We additionally found *Kdm5c*-KO EpiLCs ectopically express DAZL protein that is localized to the  
581 cytoplasm, similar to its morphology in spermatogonia<sup>59</sup>. **note: maybe just put in results.** Could  
582 move around depending upon if I get pheno working.
- 583 • end with something like: this indicates kdm5c not only modulates pluripotency and self-renewal in  
584 ESCs, but also has a role in long-term silencing of germline genes
  - 585 – then transition into the long term silencing mechanism paragraph
- 586 • KDM5C is dynamically regulated during the window of embryonic germline gene silencing, our EpiLC  
587 ChIP-seq is likely catching the tail end of KDM5C's main involvement.
- 588 • Our results indicate KDM5C is a pivotal repressor of germline programs in somatic cells.
- 589 • Studies on other chromatin regulators have focused on key marker genes to identify germline gene  
590 misexpression, such as *Dazl*.
- 591 • Using a germline-depleted mouse model, we curated a list of mouse germline-enriched genes to  
592 globally assess germline gene dysregulation.
- 593 • Many of the genes dysregulated in the mature *Kdm5c*-KO brain were important for late stages of  
594 spermatogenesis, such as those important for sperm flagellar structure. Contrastingly, *Kdm5c*-KO  
595 EpiLCs aberrantly expressed key drivers of germ cell identity, including *Dazl* and *Stra8*.

- 596 • While a significant portion of KDM5C-bound promoters are germline genes, many germline genes  
597 expressed during *Kdm5c*-KO embryogenesis are not directly bound by *kdm5c*.
- 598 • One notable example is *Stra8*, a transcription factor important for germ cell specific transcription and  
599 meiotic initiation
- 600 • The including the demarcation between soma and germline fates.
- 601 the *Kdm5c*-KO brain also Previous work identified testis-enriched genes in *Kdm5c* genes
- 602 –
- 603 – However unlike the gonadal-biased DEGs,
- 604 • The demarcation of the germline versus soma is a key feature of multicellularity and sexually dimorphic  
605 reproduction
- 606 • Anything known about tissue-biased gene expression in other H3K4me regulators?
- 607 • Our data suggests the germline developmental program is occurring ectopically as *Kdm5c*-KOs pro-  
608 gresses through somatic tissue development
- 609 • tissue-biased gene expression:
- 610 • Altogether, these data indicate that while some germline genes are misexpressed due to direct loss of  
611 KDM5C binding during emryogenesis, secondary downstream mechanisms can also promote their  
612 aberrant transcription.
- 613 • Papers to read/reference:
- 614 – Reconstitution of the Mouse Germ Cell Specification Pathway in Culture by Pluripotent Stem Cells:  
615 [https://www.cell.com/fulltext/S0092-8674\(11\)00771-9](https://www.cell.com/fulltext/S0092-8674(11)00771-9)
- 616 – 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/>

618 **Figure outline:**

- 619 **Figure 1: Misexpression of tissue-specific genes in the *Kdm5c*-KO brain** \* MA-plot and bar graphs of  
620 tissue-enriched genes \* Example testis-specific genes (NCBI and bigwigs) \* An example ovary tissue-specific  
621 gene \* An example muscle/liver tissue-specific gene (NCBI and bigwigs)
- 622 **Figure 2: The male *Kdm5c*-KO brain expresses male and female germline-enriched genes** \* Gene  
623 ontology of testis DEGs in the amygdala and hippocampus - ontologies are germline ontologies \* Expression  
624 of testis DEGs in germline-depleted testis (this is adult testis data) \* scRNASeq of testis - # of testis DEGs that  
625 are germline-specific markers \* Although far fewer, 5CKO brain also expresses ovary-enriched genes(NCBI

626 and bigwigs of Zar1) \* These ovary enriched genes are also germline specific (NCBI/Li tissues are in adult  
627 ovary, so it would be best to show they're oocyte-specific in adult ovary. But I don't think there's a published  
628 adult female W/Wv dataset. Could try looking at scRNASeq or just do TPM in embryonic W/Wv data since  
629 oocytes are developed at this point? Or both?) \* Defining what is/isn't a germline gene, and which are  
630 male/female biased using embryonic W/Wv data

631 **Figure 3: Kdm5c-KO epiblast-like cells express key drivers regulators of germline identity** \* A) ESC  
632 to EpiLC differentiation Left - Morphology is unchanged, \* B) 5CKO EpiLCs express EpiLC differentiation  
633 genes similar to WT lvs \* C) Male EpiLCs express germline genes (example Cyct again) \* Overlap between  
634 brain and EpiLC germline genes - show they're mostly unique \* GO of Brain and EpiLC germline genes  
635 (meiotic enriched) \* Bigwigs or TPM of master regulators \* Show that while some are also 2-cell genes  
636 (Dazl), 2-cell specific genes aren't dysregulated (Zscan4). Important point because published KDM5C dazl  
637 paper is saying KDM5C is a 2-cell regulator, but as far as I can tell only genes shared between germline and  
638 2-cell are dysregulated.

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

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

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

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

658 Introduction: \* Chromatin regulators are important for cellular identity \* H3K4me1-3 linked to active  
659 gene promoters and enhancers \* Surprisingly, mutations in Chromatin regulators lead to many NDDs  
660 (including many H3K4 regulators) \* Recent studies have shown some chromatin regulators are important  
661 for regulating neuron-specific gene expression/chromatin stat\_compare\_means \* However, loss of some

662 chromatin regulators can also lead to ectopic expression of tissue-enriched genes \* Very few studies have  
663 looked at these genes and it's unclear if these genes contribute to NDD impairments. \* Necessary to  
664 first characterize the mechanism behind their derepression to identify molecular footholds into testing their  
665 contribution to neuronal impairments and potential for therapeutic intervention

- 666 • Loss of KDM5C can result in the misexpression of genes typically only found in the testis
- 667 – Misexpression of tissue-enriched genes hasn't been systematically characterized - Unclear if  
668 these genes are exceptions or if other tissue-specific genes are dysregulated
- 669 – Interestingly, these genes (Cyct, D1pas1) typically function in the germline
- 670 – Germ cells (meiotic cells) are typically distinguished from somatic cells very early on in embryoge-  
671 nesis and is a key feature of multicellularity
- 672 – Chromatin regulators are very important for decommissioning germline genes and act successively  
673 the embryo implants into the uterine wall
- 674 \* Most studies have focused on ESCs, which have a similar transcriptome to germ cells / 2-cells
- 675 \* recently, KDM5C was shown to repress DAZL in ESCs, independent of its catalytic activity
- 676 \* However, DNA methylation is lost in the mature 5CKO brain, DNA methylation is placed later  
677 and it's Unclear if it's required for long-term repression (maybe too specific, just trying to go  
678 into the fact that the mechanism is partially understood but unclear)
- 679 – Systematic characterization of ectopic germline genes hasn't been done
- 680 \* unknown if other germline-enriched genes are dysregulated, including oocyte-specific genes
- 681 \* Crucially, it's unknown if misexpression of the germline program leads to functional conse-  
682 quences in 5CKO cells.

683 **Germline gene repression background:**

684 Interestingly, some of the ectopic testis transcripts identified in the *Kdm5c*-KO brain are typically ex-  
685 pressed in germ cells<sup>10</sup>. Unlike somatic cells, germ cells (e.g. sperm and eggs) undergo meiosis and pass  
686 on their genetic material to the next generation. The germline and the soma are typically distinguished during  
687 early embryogenesis, when germline genes are silenced in epiblast stem cells soon after implantation and  
688 only reactivated in a subset to form the germline. Chromatin regulators play a key role in decommissioning  
689 germline genes as the embryo transitions from naïve to primed pluripotency by placing repressive histone  
690 H2A lysine 119 monoubiquitination (H2AK119ub1)<sup>16</sup>, histone 3 lysine 9 trimethylation (H3K9me3)<sup>16,17</sup>, and  
691 DNA CpG methylation<sup>17–19</sup> at germline gene promoters. KDM5C may also be involved in this early decom-  
692 missioning of germline genes, as re-expression of KDM5C in neurons fails to suppress their dysregulation<sup>10</sup>.  
693 In support of this, KDM5C was very recently shown to repress *Deleted in azoospermia like (Dazl)*, a key  
694 regulator of germline development, in mouse embryonic stem cells (ESCs)<sup>41,52</sup>. In support of this, two  
695 independent screens in mouse embryonic stem cells (ESCs) recently identified KDM5C as a repressor of  
696 *Deleted in azoospermia like (Dazl)*, a key regulator of germline development. However, KDM5C's role in

697 embryonic germline gene repression is currently unclear, given that Dazl is also expressed in ESCs and in  
698 the 2-cell stage and germline gene misexpression has yet to be globally characterized during *Kdm5c*-KO  
699 embryogenesis.