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Female human primordial germ cells display X-chromosome dosage compensation despite the absence of X-inactivation

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Abstract

X-chromosome dosage compensation in female placental mammals is achieved by X-chromosome inactivation (XCI). An exception are human pre-implantation embryos, where dosage compensation occurs by X-chromosome dampening (XCD). Here, we examined whether XCD extends to human prenatal germ cells given their similarities with naïve pluripotent cells. We found that female human primordial germ cells (hPGCs) display reduced X-linked gene

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Author contributions

T.C., K.P., and A.C. designed the experiments. T.C. and R.C. conducted immuno-RNA FISH experiments on tissues and *in vitro* PSC-derived cells. T.C., R.K., and E.P. generated female hiPSC lines from fibroblasts and conducted human blastocyst experiments. D.C. contributed to hPGCLC differentiation experiments. F.H. created interactive web-site to explore our scRNA-seq datasets. S.L. and K.S.-L. provided the human fetal tissues. T.C. performed scRNA-seq experiments, and the resulting data were analyzed by T.C. and I.D. T.C. analyzed bulk RNA-seq data. T.C., K.P. and A.C. interpreted all data and wrote the manuscript.

Ethics declarations

Competing interests

The authors declare no competing interests.

expression before entering meiosis. Moreover, in hPGCs, both X-chromosomes are active and express the long non-coding RNAs *XACT* and *XIST*, the master regulator of XCI, which are silenced upon entry into meiosis. These findings uncover *XACT* as hPGC-marker, describe XCD associated with *XIST*-expression in hPGCs, and suggest that XCD evolved in humans to regulate X-linked genes in pre-implantation embryos and PGCs. Additionally, we found a unique X-chromosome regulation in human primordial oocytes. Therefore, future studies of human germline development must consider the sexually dimorphic X-chromosome dosage compensation mechanisms in the prenatal germline.

Dosage compensation of genes on the X-chromosome is an essential epigenetic event that equalizes the X-linked gene imbalance between males and females^{1–8}. In mice, dosage compensation is mediated by X-chromosome inactivation (XCI)^{1–7}, which is established early during development, first in an imprinted form and after a brief phase of reactivation in naïve pluripotent epiblast cells, through the random form by silencing either the maternal or paternal X-chromosome^{1–7}. Molecularly, XCI is mediated by the long non-coding RNA (lncRNA) *XIST* (*X-inactive specific transcript*), which coats the X-chromosome *in cis* to establish and maintain a silencing compartment over the X-chromosome territory^{1–7}. Once established, XCI is stably maintained in female somatic cells, yet in female mouse primordial germ cells (mPGCs) the inactive X-chromosome (Xi) is reactivated, coincident with global epigenetic reprogramming^{9–11}. Following epigenetic reprogramming and as mPGCs differentiate into meiotic cells in females or pro-spermatogonia in males, germ cells display X-chromosome dosage excess or X-chromosome dosage decompensation, respectively¹², highlighting a sexually dimorphic regulation of gene expression on the X-chromosome in mouse germ cells.

Despite the conservation of *XIST* and XCI across placental mammals¹³, it is now appreciated that there is no evidence for the imprinted form of XCI in human pre-implantation embryos^{14–16}. Moreover, in human female pre-implantation blastocyst, *XIST* is uniquely expressed from both X-chromosomes^{14,15,17} and the expression of X-linked genes on both alleles is transcriptionally reduced, yet not silenced, a compensated state that is referred to as ‘X-chromosome dampening’ (XCD)¹⁴. Another striking difference in humans is the existence of *XACT* (X active coating transcript), a primate-specific lncRNA that is expressed from the active X-chromosome(s) in both human pre-implantation embryos and pluripotent stem cells that may oppose *XIST*'s function^{14,17,18}. The expression of *XACT* in human PGCs (hPGCs) is unknown.

The unique state of X-chromosome dosage compensation in human pre-implantation embryos has been puzzling and XCD is speculated to resolve to XCI within 1–2 weeks¹⁹. Therefore, it has been postulated that the transient accumulation of *XIST* on both X-chromosomes with XCD represents the initiating stages of the normal process of XCI²⁰. This interpretation is consistent with bi-allelic *XIST* expression observed in differentiating female mouse embryonic stem cells (ESCs) during the XCI initiation process²¹, yet inconsistent with other findings that suggest a transient state with two active X-chromosomes (Xa's) without *XIST* expression in cells transitioning between XCD and XCI^{22,23}. Since mouse and human PGCs capture many of the epigenetic features of epiblast

cells from the pre-implantation embryo^{24–26}, yet hPGCs stably persist for around two months during development, we hypothesized that female hPGCs could serve as an alternate model to evaluate the possibility that XCD is an independent and stable regulatory mechanism for X-chromosome dosage compensation in humans.

Results

Female hPGCs express *XACT* from two active X-chromosomes

Since expression of the X-linked lncRNA *XACT* is associated with the unique X-chromosome state of human pre-implantation embryos^{14,17}, we performed RNA fluorescence in situ hybridization (FISH) for *XACT* on prenatal ovaries together with immunostaining for the germ cell-specific protein Deleted in azoospermia like (DAZL), which marks hPGCs²⁷. We observed *XACT* transcripts in nearly all female hPGCs (DAZL-positive cells) at weeks 7–8 post-fertilization (pf), whereas somatic cells (DAZL-negative) lacked *XACT* expression (Fig. 1a,b). We identified two *XACT* clouds in approximately 60–70% of hPGCs, each likely associated with the X-chromosome the RNA is transcribed from (Fig. 1c). Starting from 10 weeks pf, hPGCs heterogeneously differentiate into meiotic germ cells in females²⁸, repressing pluripotency genes such as *NANOG*²⁹. We therefore additionally detected NANOG to distinguish NANOG+/DAZL+ hPGCs from NANOG-/DAZL+ meiotic germ cells. We discovered that *XACT* is still expressed in the majority of NANOG+/DAZL+ female hPGCs at week 14 pf (Fig. 1d–e). In contrast, the majority of NANOG-/DAZL+ meiotic germ cells are *XACT*-negative (Fig. 1d–e). Thus, *XACT* is expressed from both X-chromosomes in hPGCs, and is not expressed by ovarian somatic cells.

To evaluate whether additional genes are biallelically expressed in hPGCs, we performed RNA FISH for the X-linked genes *HUWE1* and *ATRAX*, detecting their nascent transcription foci in week 8 pf ovaries. We found that the majority of hPGCs marked by *XACT* also biallelically expresses *HUWE1* and *ATRAX* (Fig. 1f–h). In contrast, the somatic cells express these genes from a single allele due to XCI (Fig. 1f–h). These results are consistent with prior allelic analysis of X-linked gene expression from 53 sorted female germ cells³⁰, which revealed biallelic expression of X-linked genes (Extended Data Fig. 1a). Consequently, we conclude that female hPGCs carry two active X-chromosomes.

To confirm the active state of the X-chromosome in female hPGCs, we assessed a chromatin mark specific for the Xi. In mouse, newly specified female PGCs have an enrichment of H3K27me3 on the Xi which is progressively lost as the cells undergo Xi-reactivation^{9,11}. Given that 23% of female hPGCs have one *XACT* cloud (Fig. 1c), we evaluated whether female hPGCs show any evidence of nuclear H3K27me3 accumulation, indicative of an Xi^{31,32}. At week 4 pf, we identified early stage hPGCs entering the genital ridge epithelium before induction of DAZL expression through OCT4 expression (Extended Data Fig. 1b). We found that around 10% of OCT4+/DAZL- female hPGCs have an Xi-like nuclear accumulation of H3K27me3, which is reduced to less than 5% at weeks 7 and 12 pf, once the hPGCs have settled in the ovary (Extended Data Fig. 1b–e). The lack of a H3K27me3 Xi-like enrichment suggests that the monoallelic expression of *XACT* observed in a portion of female hPGCs is not linked to residual XCI.

Since a naïve-like pluripotent expression program is common to both male and female hPGCs³³, we explored whether *XACT* also marks male hPGCs. Using published bulk RNA-seq data of sorted male and female hPGCs^{24,25}, we detected *XACT* expression in hPGCs but not somatic cells (Fig. 1i). We confirmed this result using RNA FISH on a fetal testis at 13 weeks pf, detecting *XACT* in 80% of NANOG+ hPGCs (Fig. 1j–k). Thus, as is the case for human pre-implantation embryos, both female and male hPGCs express *XACT*.

XACT* is expressed in male and female hPGCLCs *in vitro

The expression of *XACT* in male and female hPGCs *in vivo* suggested that *XACT* may be expressed from the moment of hPGC specification. Given that hPGCs are thought to be specified between week 2–3 pf³⁴, it is not possible to study hPGC specification *in vivo*. Therefore, we modelled hPGC specification using the differentiation of hPGC-like cells (hPGCLCs)³⁵ from male (UCLA2 ESCs³⁶) and female (MZT04-J iPSCs³⁷) pluripotent stem cells, by analyzing *XACT* distribution in ITGA6+/EPCAM+ hPGCLCs and somatic cells at day 4 of differentiation (Fig. 2a). We discovered that most male hPGCLCs had one *XACT* cloud, whereas most female hPGCLCs had two (Fig. 2b–g). In contrast, >90% of the somatic cells were *XACT*-negative (Fig. 2b–g). The detection of biallelic *XACT* in female hPGCLCs is consistent with maintenance of the eroded Xi in differentiating hESCs²³. Collectively, these data uncover the lncRNA *XACT* as a new marker of male and female hPGCs *in vivo* and *in vitro*.

***XACT* expression is epiblast-specific in human embryo attachment cultures**

Given that *XACT* is expressed in NANOG+ hPGCs/hPGCLCs, we next investigated *XACT* distribution during the peri-implantation window of human development using human embryo attachment culture^{38,39}. Specifically, we explored the pattern of *XACT* expression in NANOG+ epiblast cells versus NANOG-negative trophoderm (TE) and primitive endoderm (PE) cells. We additionally detected the lncRNA *XIST* to uncover changes in its distribution. First, we performed RNA FISH for *XACT* and *XIST* in combination with immunostaining for NANOG on D6 pf female and male human blastocysts. At D6, both *XIST* and *XACT* clouds could be identified on all X-chromosomes (one in males and two in females) in 75% of female and 55% of male NANOG+ epiblast cells (Fig. 3a,b,d,e). Additionally, 30% of male epiblast cells also express only *XACT* (Fig. 3e). Conversely, NANOG-negative TE/PE cells have more diverse *XIST* and *XACT* expression states with *XACT* being more often repressed (Fig. 3c,f). These observations indicate that *XACT* expression is more strongly associated with the NANOG+ epiblast. Using human embryo attachment culture^{38,39} grown to D12 pf, the legal limit for human embryo culture in California, we discovered that the majority of female NANOG+ epiblast cells continued to display *XIST* and *XACT* clouds on both X-chromosomes (Fig. 3g,h), whereas male epiblast cells maintained expression of *XACT* but not of *XIST* (Fig. 3j,k). In the majority of female NANOG-negative cells, *XACT* was repressed and *XIST* was expressed from one X-chromosome (Fig. 3i), suggesting that these cells have transitioned to the initiation of XCI. In contrast in male embryos, both lncRNAs were silenced in 70% of NANOG-negative cells (Fig. 3l). Together, these data reveal that *XACT* is expressed by the majority of NANOG+ epiblast cells in the pre-implantation and early peri-implantation stages of male and female human embryo development. In contrast, *XIST* is rapidly repressed in male NANOG+

epiblast between D6 and D12, highlighting differences in *XIST* regulation in male and female human peri-implantation development.

Female hPGCs dampen expression from the active X chromosomes

Since female hPGCs express *XACT* from both X-chromosomes, our next goal was to determine whether XCD is occurring in hPGCs. We performed single cell (sc) RNA-seq on single cell suspensions of five prenatal ovaries and five prenatal testes from week 6 to 16 pf (Fig. 4a–c). This unbiased approach yielded a total of ~50,000 prenatal gonadal cells, with ~281 male and 1938 female germ cells for the analysis of the X-chromosome/autosome gene expression ratio (X/A ratio) (summarized in Supplemental Table 1).

Calculating the X/A ratio across individual cells per developmental age revealed that female germ cells consistently had a higher X/A ratio than male germ cells (Extended Data Fig. 2a,b). This difference between female and male germ cells is arising from significantly lower X-linked gene expression in males (Extended Data Fig. 2c). In contrast, we found no difference in the X/A ratio between male and female gonadal somatic cells (Extended Data Fig. 2a,b). We confirmed these results with published scRNA-seq data set of cKIT/size-selected germ cells⁴⁰, where germline cells are called fetal germ cells (FGCs) (Extended Data Fig. 2d,e). Thus, the X/A ratio is higher in female germ cells relative to the male, whereas male and female gonadal somatic cells are equivalent.

Since germ cell differentiation into meiotic cells is heterogeneous, we created an unsupervised developmental trajectory⁴¹, which ordered female human germ cells across 11 clusters (Fig. 4d,e). Clusters 0–5 represent hPGCs expressing the transcription factors *NANOG* and *OCT4*, together with the hPGC markers *NANOS3*, *PRDM1* and *SOX17* (Fig. 4d). Starting in cluster 6, we observed downregulation of naïve-like pluripotency genes and upregulation of the meiotic licensing gene *STRA8*⁴² and of genes encoding RNA binding proteins such as *DAZL* and *DDX4* (*VASA*) (clusters 6–7). This was followed by expression of the meiotic prophase I genes *SPO11* and *SYCP1* in clusters 8–9, and ultimately, in cluster 10, up-regulation of primordial oocyte (PO) genes including the *Zona Pellucida Protein 3* (*ZP3*)⁴³ (Fig. 4d). Thus, consistent with previous reports^{24–26,28,30,40}, our data capture the heterogeneous differentiation of female hPGCs into meiotic germ cells beginning around week 9–10 pf, which results in a complex mixture of germ cells including hPGCs, meiotic germ cells and primordial oocytes in a given prenatal ovary (Fig. 4f).

Analysis of the X/A ratio in female germ cells along the developmental trajectory (Fig. 5a) showed that as hPGCs begin differentiating into meiotic germ cells (cluster 6 onwards), the X/A ratio increases, reaching maximal levels in cluster 9 before precipitously dropping in primordial oocytes (cluster 10). Moreover, the X/A ratio in hPGCs (clusters 0–6) is lower than in gonadal somatic cells (Fig. 5a, Extended Data Fig. 3a–c). Overall X/A ratio in hPGCs (clusters 0–5) is significantly lower compared to meiotic germ cells (clusters 6–9) but higher than primordial oocytes (cluster 10) (Fig 5b). These changes were largely due to changes in X-linked gene expression (Extended Data Fig. 3d,e). We confirmed these observations in the FGC scRNA-seq data⁴⁰. Similar to our analysis, female FGCs displayed a slightly lower X/A ratio than gonadal somatic cells in the hPGC state, and a subtle but significant increase in the X/A ratio after expression of *STRA8* (Extended Data Fig. 3f,g).

Based on these results, we conclude that the dosage of X-linked genes is dampened in female hPGCs and that this dosage compensation is erased as cells enter prophase I of meiosis I.

To evaluate X/A ratios in male prenatal germ cells, we ordered the cells along a developmental trajectory, which divided the male germ cells from week 6–16 pf gonads into five clusters (0–4), with the hPGC program corresponding to clusters 0–3 and cluster 4 capturing differentiating germ cells (prospermatogonia) (Extended Data Fig. 4a,b). Unlike female germ cells that exhibit X-chromosome dosage excess with exit from the hPGC state, the X/A ratio does not change when male hPGCs differentiate into prospermatogonia (Extended Data Fig. 4c–h), which was confirmed with published data sets⁴⁰ (Extended Data Fig. 4i,j). Therefore, an increase in X/A ratios as hPGCs differentiate, is a female-specific phenomenon associated with entrance into prophase I of meiosis I.

XCD in female hPGCs is associated with *XIST* expression

To evaluate whether XCD in female hPGCs is associated with expression of *XIST*, as in female human pre-implantation embryos, we examined *XIST* in individual female germ cells along the developmental trajectory. We discovered that *XIST* expression is significantly higher in hPGCs (clusters 0–5) compared to meiotic germ cells (clusters 6–9) and primordial oocytes (cluster 10) (Fig. 5c,d). This result was also validated in the female FGC data set⁴⁰ (Extended Data Fig. 5a). In agreement with the low expression of *XIST* in primordial oocytes, the levels of the RNA are also low in adult oocytes⁴⁴ (Extended Data Fig. 5b). Thus, the increase in the X/A ratio from hPGCs to meiotic germ cells is accompanied by a reduction in *XIST* (Fig. 5b,d and Extended Data Fig. 5c). Consistent with this result, when analyzing hPGCs and meiotic germ cells together (clusters 0–9) based on *XIST* expression, *XIST*⁺ cells displayed a significantly lower X/A ratio than *XIST*-negative cells. Primordial oocytes (cluster 10) have an even lower X/A ratio, yet express *XIST* at a very low level (Fig. 6b,d, Extended Data Fig. 5d).

Next, we evaluated the localization of *XIST* in female hPGCs by RNA FISH from week 7 to week 14 pf, using *XACT* as a marker of hPGCs. We found that *XIST* is detectable in the vast majority of *XACT*-expressing hPGCs, with diverse patterns of the *XIST* signal (Fig. 6a–c). These include (i) an eroded cloud pattern, where *XIST* is restricted over one X-chromosome in a pattern characteristic of Xi-localization, yet less enriched compared to the Xi in somatic cells, combined with a nascent transcription spot of *XIST* on the second X-chromosome; (ii) a dispersed configuration where the *XIST* signal is detected throughout a large portion of the nucleus albeit in the vicinity of both *XACT* clouds, suggesting that *XIST* is expressed from both X-chromosomes; (iii) one dot in the vicinity of one of the two *XACT* signals, indicating expression from one X-chromosome; and (iv) two dots representing the nascent transcription sites of *XIST* on both X-chromosomes (Fig. 6a). Quantification of *XIST* expression patterns in female hPGCs with two *XACT* clouds, which enabled the localization of both X-chromosomes, revealed that the majority of hPGCs (58%) at week 7 pf had a dispersed *XIST* signal (Fig. 6c). Around 6% of cells with biallelic *XACT* expression displayed an eroded *XIST* cloud pattern and 19% and 15%, respectively, exhibited mono- and bi-allelic nascent *XIST* transcription foci (Fig. 6c). At later stages of

embryo development (week 8 and week 14 pf), *XACT*-positive female hPGCs displayed similar patterns of *XISTR* RNA with an increasing fraction of *XIST*-negative cells (Fig. 6b,c).

Since we observed an Xi-like distribution of *XIST* in 6% of week 7 and 8 pf hPGCs with two *XACT* clouds, we tested whether female hPGCs with one *XACT* cloud represent cells that have undergone XCI. In this case, *XACT* should be expressed from the Xa and *XIST* from the Xi. However, we found that *XIST* is typically expressed from the same X-chromosome as *XACT* in cells with monoallelic *XIST/XACT* expression (Fig. 6d), providing additional evidence for XCD instead of XCI in female hPGCs.

To confirm that the loss of *XIST* expression was associated with differentiation of NANOG⁺/DAZL⁺ hPGCs into NANOG⁻/DAZL⁺ meiotic germ cells, we evaluated a fetal ovary at 14 weeks pf by RNA FISH. We found that *XIST* was detectable in the majority of NANOG⁺/DAZL⁺ hPGCs, with the dispersed pattern being most prominent. In contrast, the majority of NANOG⁻/DAZL⁺ germ cells were negative for *XIST* and the subset of cells with *XIST* expression displayed the dot-like distribution (Fig. 6e,f). The quantification suggests that cells transition from the “dispersed *XIST*” pattern to the “2 *XIST* dot” and “1 *XIST* dot” patterns before *XIST* is turned off during germ cell differentiation. Furthermore, evaluating *XIST* localization around the X-chromosomes in female hPGCs relative to female human pre-implantation embryos, revealed a higher degree of *XIST* dispersal in hPGCs (compare Fig. 3/6), suggesting that the association of *XIST* with chromatin differs between the two cell types.

In addition to detecting *XIST* expression in female hPGCs, *XIST* transcripts could also be detected in male hPGCs based on scRNA-seq data, albeit at much lower level than in female cells (Extended Data Fig. 5 e,f). Furthermore, a small but significant reduction in the X/A ratio in male hPGCs correlated with significant increase in *XIST* levels (Extended Data Fig. 4j and 5f). Thus, a subset of male hPGCs can express *XIST* and display slight dampening of X-chromosome dosage. Moreover, female germ cells that lacked *XIST* displayed an increase, albeit not significant, in the *XACT* cloud size compared to cells with *XIST* expression (Extended Data Fig. 5g). Taken together, these data suggest that *XIST* may mediate dampening of X-linked gene expression in hPGCs.

Discussion

In this study, by analyzing human pre-implantation embryos, human embryo attachment culture, hPGCLC differentiation *in vitro*, and hPGCs *in vivo*, we revealed that the lncRNA *XACT* is expressed in pluripotent epiblast cells and hPGCs/hPGCLCs (Fig. 6g, Extended Data Fig. 6). Mechanistically, this result may be explained by the presence of an enhancer that threads *XACT* into the pluripotency network common to these cell types⁴⁵. From this analysis, we describe *XACT* as a unique marker of hPGCs, and speculate that it could be used to trace hPGCs from the time of lineage specification by RNA FISH. Additionally, our RNA FISH analysis of the X-linked genes *XACT*, *ATRX* and *HUWE1*, together with absence of H3K27me3 accumulation in the nucleus of most female hPGCs demonstrates that female hPGCs harbor two Xa's from at least week 4 pf onwards.

Although our imaging approaches demonstrated the presence of two active X-chromosomes in female hPGCs, the scRNA-seq data revealed that the X/A ratio is reduced in female hPGCs compared to female meiotic germ cells (Fig. 6g). These results indicate that X-linked dosage compensation in female hPGCs is regulated by the XCD mechanism, similar to female naïve human pluripotent stem cells²² and female human pre-implantation embryos¹⁴. While in pre-implantation embryos XCD is a transient state, in case of hPGCs, we show that XCD is not a transitional state into XCI, rather a stable state lasting at least 6 weeks until the point of meiotic initiation (Fig. 6g). Similar to female mouse PGCs¹², we also show that X/A ratio excess occurs as female hPGCs initiate meiosis. Following prophase I of meiosis I, the X/A ratio rapidly declines coincident with primordial oocyte formation indicating a third unique state of X-chromosome dosage compensation, which we have called X^o (Fig. 6g). Intriguingly, the loss of XCD in the female germline upon meiotic entry is linked to the silencing of *XIST* (Fig. 6g), suggesting that *XIST* is the mediator of XCD. In contrast, the further decline in the X/A ratio in primordial oocytes occurs in the absence of *XIST* expression. It remains unclear whether this regulation is achieved by XCD or other mechanisms.

Given that male hPGCs have a reduced X/A ratio compared to female and male somatic cells and female hPGCs, we refer to the active X-chromosome in male prenatal germ cells as Xa* (Fig. 6g). The lower X/A ratio in male hPGCs relative to female hPGCs could be due to inefficient dampening of X-linked gene expression from both X-chromosomes by *XIST* in female hPGCs, such that the levels in female hPGCs are higher than males. In support of this hypothesis, *XIST* is very dispersed in female hPGCs, which may lead to less efficient XCD compared to the more cloud-like distribution of *XIST* reported for the pre-implantation embryo^{14,15}. On the other hand, a higher X/A ratio in female and male somatic cells could be explained by upregulation of single Xa in somatic cells^{12,46,47}, which may not be occurring in female or male hPGCs. Combined with XCD occurring on the X-chromosomes in female hPGCs, this alternate explanation would explain lower X-linked gene expression in female hPGCs compared to somatic cells.

Since germline specification in humans takes days compared to hours in mice, and hPGC development is a much lengthier process than in mice, it is conceivable that the maintenance of X-chromosome dosage compensation in germ cells between the two species are different. It is likely that a primate specific X-chromosome regulation mechanism might be necessary to compensate dosage of X-linked genes in the human embryo during PGC specification and the first trimester of pregnancy. However, how *XIST* and *XACT* contribute to X-chromosome gene regulation in the developing human germline will need to be studied mechanistically. Achieving this goal will require new *in vitro* cell models of hPGC development that are able to reliably promote the differentiation of hPGCLCs into meiotic germ cells and primordial oocytes combined with functional approaches⁴⁸.

X-chromosome dosage regulation might be extremely important for Turner (XO) and Klinefelter (XXY) syndrome patients, who suffer from infertility associated with loss of germline cells^{49,50}. Although germline development in fetuses diagnosed with Turner syndrome is morphologically normal, oocyte loss occurs within the first few months after birth⁵¹. Potentially, meiosis is not occurring correctly in Turner syndrome patients due to

diminished levels of critical X-linked genes in differentiating XO hPGCs compared to XX hPGCs. Thus, upregulation of X-linked gene expression with entrance into meiosis may be necessary for the formation of mature oocytes.

In summary, with the demonstration of XCD, our work draws parallels between the X-chromosome state of human epiblast and hPGCs. Our work sheds light on mechanisms that regulate X-linked gene expression in hPGCs prior to meiosis, and reveal a unique X-chromosome state in oocytes, which potentially could be important for oocyte formation and zygote development downstream.

Methods

Human fetal tissues

Prenatal gonads from 4 to 16 weeks post-fertilization were obtained from either the University of Washington Birth Defects Research Laboratory (BDRL) or the University of Tübingen. At BDRL, prenatal gonads were obtained with regulatory oversight from the University of Washington IRB approved Human Subjects protocol, combined with a Certificate of Confidentiality from the Federal Government. BDRL collected the fetal testes and ovaries and shipped them overnight in HBSS with ice pack for immediate processing at UCLA. Prenatal samples from University of Tübingen were delivered to UCLA 24–48 hours after the procedure. The research project was also approved by the research ethics committee of the University of Tübingen (IRB# 584/2018BO2 and 634/2017BO1). All human fetal tissue used in the current study were obtained following informed consent. The donated human fetal tissue sent to UCLA did not carry any personal identifiers. No payments were made to donors and the donors knowingly and willingly consented to provide research materials without restrictions for research and for use without identifiers. Developmental age was documented by BDRL and University of Tübingen as days post fertilization using a combination of prenatal intakes and carnegie staging. A total of 16 fetal samples was used for this study.

Human pre-implantation embryos

Use of human embryos in this research project followed California State law and upon review by the Institutional Review Board (IRB) and the human embryonic stem cell research oversight committee (ESCRO) at UCLA. The ESCRO committee at UCLA approves human pluripotent stem cell and human embryo work at UCLA based on 2016 ISSCR guidelines. Together, these committees approve the process of informed consent, and experiments using human embryos for research purposes on an annual basis. Patients were not paid for participation, and all donors were informed that the embryos would be destroyed as part of the research study. All research with human embryos in this study complied with the principles laid out in the International Society for Stem Cell Research. Frozen human blastocysts at day 6–7 pf were used in this study and thawed using Vit Kit-Thaw (Irvine Scientific) according to manufacturer protocol. After thawing, embryos were cultured overnight in 5% O₂, 6% CO₂ at 37°C, and the zona pellucida was removed with Tyrode's acidified solution (Irvine Scientific). A total of 28 human blastocysts were used here. The

sex of the blastocysts was determined by cloud counts of lncRNA *XIST* and *XACT* expression from a single (male) or both X chromosomes (female).

Tissue processing for scRNA-seq

Fetal tissues were processed 24–48 hours after termination. Upon arrival, tissues were gently washed with PBS and placed in dissociation buffer containing of Collagenase IV 10mg/ml (Life Technologies #17104–019), Dispase II 250 ug/ml (Life Technologies #17105041), DNaseI 1:1000 (Sigma 4716728001), 10% FBS (Life Technologies 10099141) in 1x PBS. Tissues were dissociated for 15 minutes at 37°C. After every 5 minutes, the tissues were pipetted against the bottom of Eppendorf tube. Subsequently, cells were centrifuged for 5 minutes at 500g, resuspended in 1x PBS with 0.04% BSA, and strained through 40-micron strainer and counted using automated cell counter (Thermo Fisher, Countess II). The cell concentration was adjusted to 800–1200 cells per microliter and immediately used for scRNA-seq.

scRNA-seq library preparation and sequencing

scRNA-seq libraries were generated using the 10X Genomics Chromium instrument and Chromium single cell 3' reagent kit V2. Each individual library was designed to target 6000 cells. Libraries were generated according to manufacturer's instructions and library fragment size distribution was determined by BioAnalyzer instrument. Libraries were pooled together and sequenced on Illumina Novaseq 6000 platform, at average depth of 400–420 million reads per sample.

scRNA-seq data analysis

scRNA-seq reads were aligned to human hg38 genome assembly using 10X Genomics Cell ranger v2.2. Expression matrixes generated by Cell Ranger were imported to Scanpy⁵⁴ for downstream analysis. First, all libraries were merged, and cells were subjected to the same filtering. All genes expressed in less than five cells were discarded and cells with less than 250 detected genes were filtered out. The UMI counts were then normalized for each cell by the total expression, multiplied by 10000 and log-transformed. Using Scanpy's default method, highly variable genes were identified, and data were scaled to regress out variation from UMI counts and mitochondrial genes. Cells were clustered by Louvain algorithm⁵⁵ and the UMAP package was used to visualize cells in 2D plot⁵⁴. Germ cell clusters were identified by expression of germ cell-specific markers such *NANOS3*, *DAZL*, *DDX4*, *SYCP1*. Gonadal somatic cells were annotated by previously published literature⁴⁰. The female and male germline trajectories were created by partition-based graph abstraction (PAGA)⁴¹. The Li at al dataset⁴⁰ was analyzed through the same pipeline as described above. Gene expression matrixes of female and male germ cells were exported from Scanpy and X linked gene expression ratios with autosomal gene expression ratio per cell were calculated with custom R script.

Tissue processing and cryo-sectioning

Upon arrival, fetal tissues were gently washed with 1x PBS and fixed with freshly made 4% Paraformaldehyde in 1x PBS for 3–4 hours on a rotator at room temperature. Tissues were

washed with 1xPBS 3 times for 5 minutes and moved through increasing concentrations of sucrose, 10% sucrose 1 hour. 20% 1 hour and 30% overnight at 4°C. Afterwards, tissues were embedded in OCT (Tissue-Tek O.C.T) and 7micron sections were cut. Sections and tissue blocks were kept at -80°C.

Human ESC culture

Human ESC/hiPSC lines used in this study include UCLA2 (46, XY)³⁶ and MZTO4 iPSC (46, XX)³⁷. hESCs/hiPSCs were cultured on mitomycin C-inactivated mouse embryonic fibroblasts (MEFs) in hESC media, which is composed of 20% knockout serum replacement (KSR) (GIBCO, 10828-028), 100mM L-Glutamine (GIBCO, 25030-081), 1x MEM Non-Essential Amino Acids (NEAA) (GIBCO, 11140-050), 55mM 2-Mercaptoethanol (GIBCO, 21985-023), 10ng/mL recombinant human FGF basic (R&D systems, 233-FB), 1x Penicillin-Streptomycin (GIBCO, 15140-122), and 50ng/mL primocin (InvivoGen, ant-pm-2) in DMEM/F12 media (GIBCO, 11330-032). The hESCs and iPSCs were split every 7-8 days using Collagenase type IV (GIBCO, 17104-019). The hESC line used in this study is registered with the National Institute of Health Human Embryonic Stem Cell Registry and available for research use with NIH funds. Mycoplasma test (Lonza, LT07-418) was performed every month to all cell lines used in this study. All experiments were approved by the UCLA ESCRO Committee.

hPGCLC differentiation

hPGCLCs were induced from primed hESCs/hiPSCs as described before⁵³ with Stem Cell Factor (SCF) omitted from the differentiation media and starting with human pluripotent stem cells grown on MEFs. Briefly, hESCs/hiPSCs were dissociated into single cells with 0.05% Trypsin-EDTA (GIBCO, 25300-054) and plated on Human Plasma Fibronectin (Invitrogen, 33016-015)-coated 12-well- plates at the density of 200,000 cells/well in 2mL/well of iMeLC media - 15% KSR (GIBCO, 10828-028), 1x NEAA (GIBCO, 11140-050), 0.1mM 2-Mercaptoethanol (GIBCO, 21985-023), 1x Penicillin-Streptomycin-Glutamine (GIBCO, 10378-016), 1mM sodium pyruvate (GIBCO, 11360-070), 50ng/mL Activin A (Peprotech, AF-120-14E), 3mM CHIR99021 (Stemgent04-0004), 10mM of ROCKi (Y27632, Stemgent, 04-0012-10), and 50ng/mL primocin in Glasgow's MEM (GMEM) (GIBCO, 11710-035). After 24 hrs, iMeLCs were dissociated into single cells with 0.05% Trypsin-EDTA and plated into ultra-low cell attachment U-bottom 96-well plates (Corning, 7007) at the density of 3,000 cells/well in 200ml/well of hPGCLC media, which is composed of 15% KSR (GIBCO, 10828-028), 1x NEAA (GIBCO, 11140-050), 0.1mM 2-Mercaptoethanol (GIBCO, 21985-023), 1x Penicillin-Streptomycin- Glutamine (GIBCO, 10378-016), 1mM sodium pyruvate (GIBCO, 11360-070), 10ng/mL human LIF (Millipore, LIF1005), 200ng/mL human BMP4 (R&D systems, 314-BP), 50ng/mL human EGF (R&D systems, 236-EG), 10mM of ROCKi (Y27632, Stemgent, 04-0012-10), and 50ng/mL primocin in Glasgow's MEM (GMEM) (GIBCO, 11710-035).

hPGCLC sorting and preparation for RNA FISH

hPGCLC aggregates were dissociated 0.05% Trypsin-EDTA (GIBCO, 25300-054) for 10 min at 37°C. The dissociated cells were stained with conjugated antibodies, washed with FACS buffer (1% BSA in PBS) and resuspended in FACS buffer with 7-AAD (BD

PharMingen, 559925) as viability dye. The conjugated antibodies used in this study include ITGA6 conjugated with BV421 (BioLegend, 313624, 1:60), EPCAM conjugated with 488 (BioLegend, 324210, 1:60). The single cell suspension was sorted for further experiments using BD FACSAria fluorescence-activated cell sorting machine. FACS data was analyzed with FlowJo v10 software. Double positive cells for ITGA6 and EPCAM (hPGCLCs) and negative cells (non-hPGCLCs) were collected in hPGCLC media and plated on human plasma fibronectin coated coverslips overnight. Following morning, RNA FISH was performed using the coverslips.

Immunofluorescence

Slides of paraffin-embedded sections were deparaffinized by successive treatment with xylene and 100%, 95%, 70% and 50% ethanol. Antigen retrieval was performed by incubation with 10mM Tris pH 9.0, 1mM EDTA, 0.05% Tween at 95C for 40 min. The slides were cooled and washed with 1xPBS (phosphate buffered saline) and 1xTBS (PBS + 0.2% Tween). Cryo sections and Blastocysts attached to ibidi chambers were fixed with 4% PFA for 10 minutes and washed with 1xPBS. Afterwards paraffin-embedded, cryo sections and blastocysts were treated similarly. The samples were permeabilized with 0.5% Triton X-100 in 1xPBS, then washed with 1xTBS and blocked with 1% BSA in 1xTBS. Primary antibody incubation was conducted 1% BSA for 1h at RT. Samples were again washed with 3xTBS-tween and incubated with fluorescent secondary antibodies at 1:200 for 45 min, then washed and counterstained with DAPI for 5 min and mounted using Vectashield. The primary antibodies used for immunofluorescence in this study are listed in supplemental table 1 under Antibody List tab. The secondary antibodies used in this study were all from Life technologies used at 1:400 dilution. Images were taken using LSM 880 Confocal Instrument (Zeiss) or Zeiss Axio Imager M1. For image processing and analysis Fiji (ImageJ) was used. For signal quantification, images were converted into 8-bit images and afterwards performed using profile plot tool. Intensity values were exported as a csv file and afterwards R Studio and ggplot2 package was used for plotting.

RNA FISH

hPGCLCs and non-hPGCLCs after sorting were attached to fibronectin coated 18mm circular glass coverslips (Fisher Scientific, 12-545-100) overnight. Next morning, the coverslips were washed with DPBS, fixed with 4% formaldehyde for 10 min, permeabilized with cold (4°C) 0.5% Triton X-100 in DPBS for 10 min, and serially dehydrated with cold (4°C) 70–100% ethanol. Coverslips were air dried and hybridized with labeled DNA probes in a humidified chamber at 37°C overnight, washed for three 5-min intervals with 50% formamide in 2x SSC, 2x SSC, then 1x SSC at 42°C, counterstained with DAPI and mounted with Vectashield (Vector labs: H-1000). Double- stranded DNA probes were generated from full length cDNA constructs or BACs as described previously⁵⁶. The BACs used include *XIST* (RP11-13M9), *XACT* (RP11-35D3), *ATRX* (RP11-1145J4), and *HUWE1* (RP11-975N19). Every new batch of probes was tested on normal human dermal fibroblasts before use in experiments.

Immuno-RNA FISH

Immuno-RNA FISH on cryo sections was performed as described previously described⁵⁶. Briefly, immuno staining was performed first on cryo sections and blastocysts as described above. Samples were post fixed with 4% PFA after IF staining and RNA FISH was performed after post fixation as described in RNA FISH section above.

Immuno-RNA FISH on human blastocysts was performed as described previously²³, with the following modifications: Embryos at day 5 and day 6 post fertilization were thawed for these experiments and cultured for 24 hours before staining. First zona pellucida was removed with Tyrode's acid and blastocysts were washed with 6 mg/mL BSA (Sigma) in RNase-DNase-free PBS. They were then individually transferred to ibidi chambers (Ibidi 80827), coated with polylysine (Sigma P4832–50ML). fluid aspirated until dry and blastocysts were fixed with ice-cold 4% paraformaldehyde (PFA) for 10 min at room temperature. Immunostaining was performed as described above. In all buffers and antibody solutions RNaseOUT 1:200 (Thermo 10777019) was added to preserve RNA. Before performing RNA FISH samples were post fixed with 4% PFA for 10 min at RT. Afterwards RNA FISH was performed using DNA probes as described above.

Attached blastocyst culture

Human embryo attachment culture was performed following the procedures of (Deglincerti et al., 2016³⁸; Shahbazi et al., 2016³⁹). Briefly, cryopreserved human blastocysts were received vitrified from the IVF clinic following consent and thawed using Vit Kit-Thaw (Irvine Scientific) according to manufacturer protocol. The embryos were cultured in drops of Continuous Single Culture Complete medium pre-supplemented with HSA (Irvine Scientific) under mineral oil (Irvine Scientific) overnight at 37°C, 6% CO₂ and 5% O₂. The zona pellucida was removed using Tyrode's solution acidified (Irvine Scientific) and plated onto an m-Slide 8-well chamber slide (ibidi) in IVC-1 medium (Cell Guidance Systems) and incubated for 2 days at 37°C and 5% CO₂ to allow for attachment. Media was half replaced on the second and third day with IVC-1. From the fourth day and onward, media was fully replaced with IVC-2 medium (Cell Guidance Systems) until the appropriate developmental day was reached up to a maximum of Day 12, which includes the blastocyst stage plus days in culture. For these experiments, 9 blastocysts were cultured up to day 12 post fertilization, out of which 5 were used for immuno-RNA FISH experiments.

Bulk RNA-seq data analysis

Published Raw population RNA-seq data sets^{25,52,53} of male and female hPGCs and somatic cells were downloaded and realigned to hg19 genome as described previously¹⁷ for lncRNA *XACT* expression analysis. Expression tracks were generated using pyGenomeTracks package⁵⁷.

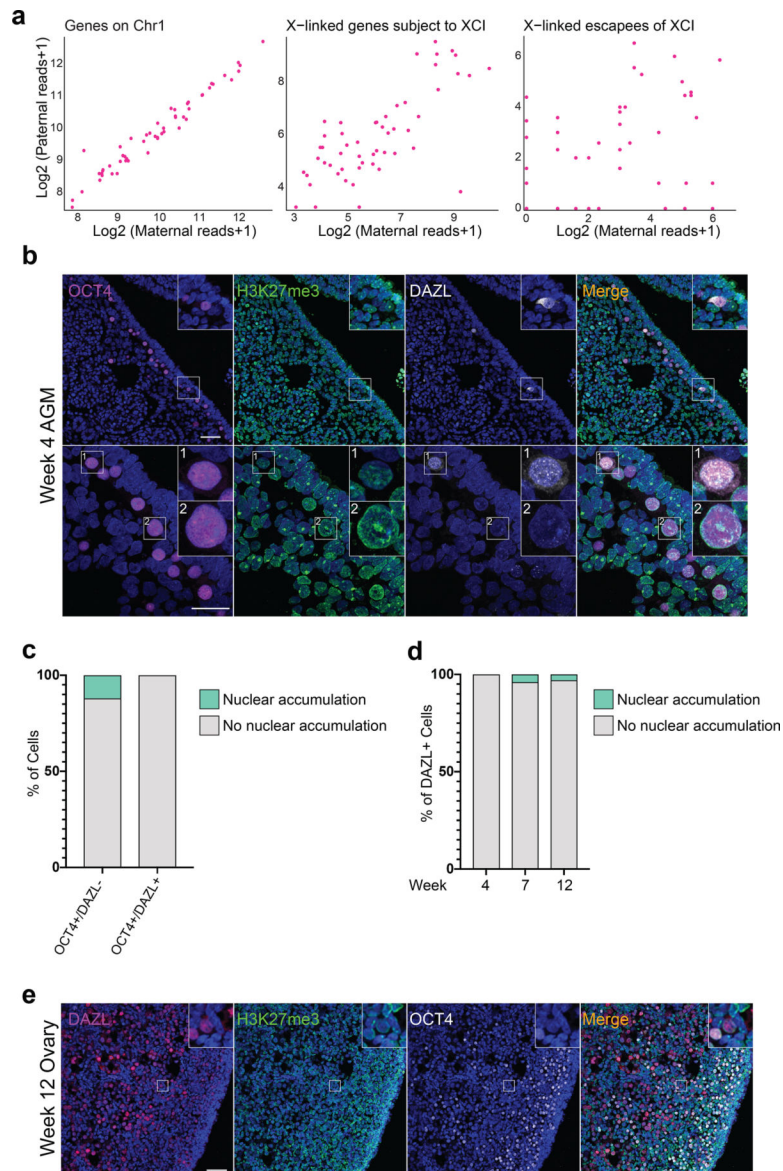
Code availability

Custom scripts used for aligning population RNA-seq, scRNA-seq, data processing and plotting are available upon request.

Statistics and Reproducibility

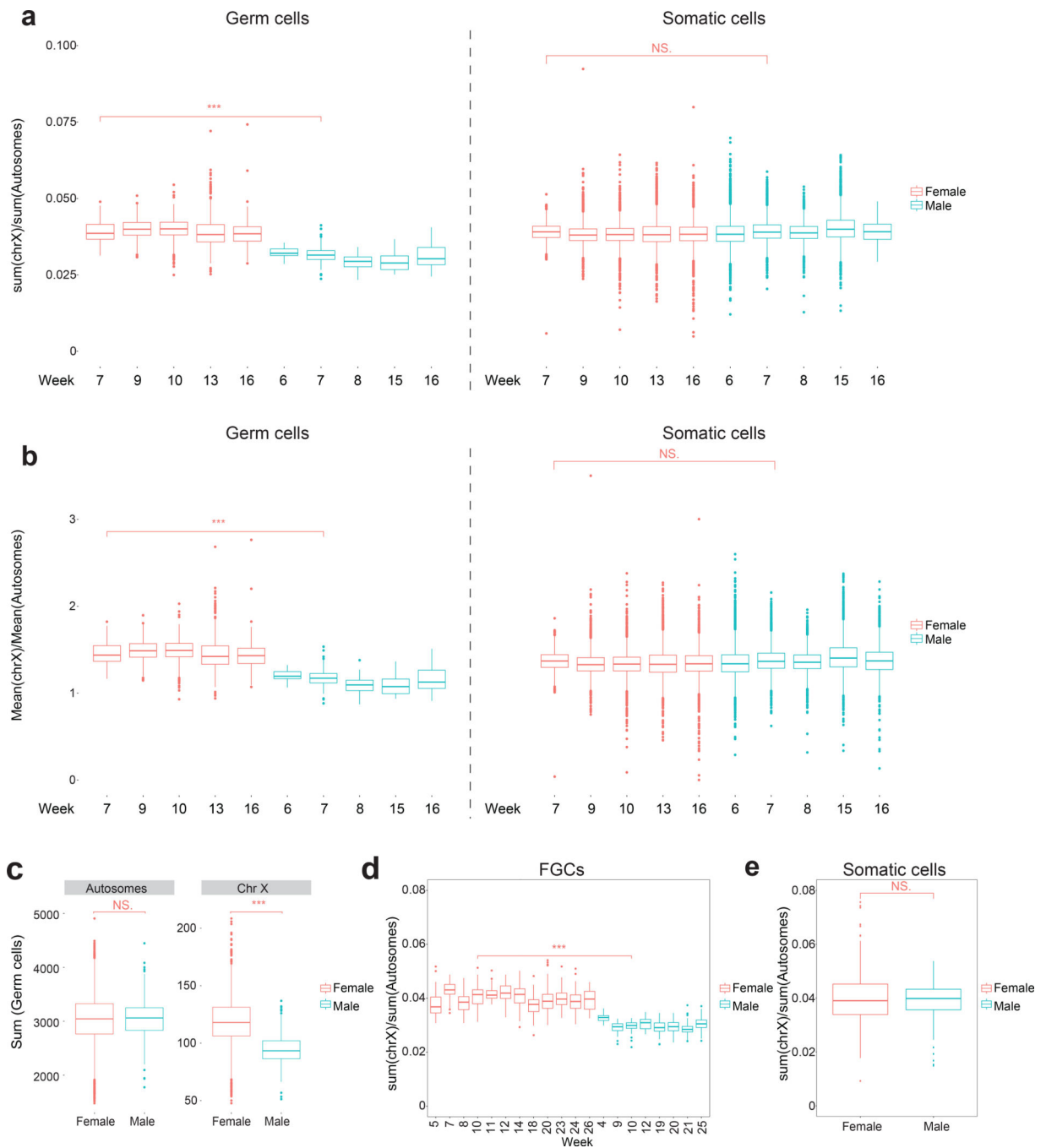
In the quantitative data, significance was assessed using Wilcoxon tests; * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$, *NS*- non-significant. Statistical analyses are described in detail for each panel. Immuno-RNA FISH experiments were performed in 2 independent experiments with similar results, unless specified otherwise in the legends. scRNA-seq datasets were pooled from 10 independent experiments. No statistical methods were used to predetermine the sample size, rather sample size was limited by availability of prenatal tissues. Signal intensity measurement details are described in Immunofluorescence section. For plotting and statistical analysis of scRNA-seq datasets and immune-RNA FISH quantifications ggsignif and ggplot2 R packages were used.

Extended Data



Extended Data Fig. 1. Female hPGCs from week 4 pf ovaries have lost the H3K27me3 nuclear accumulation

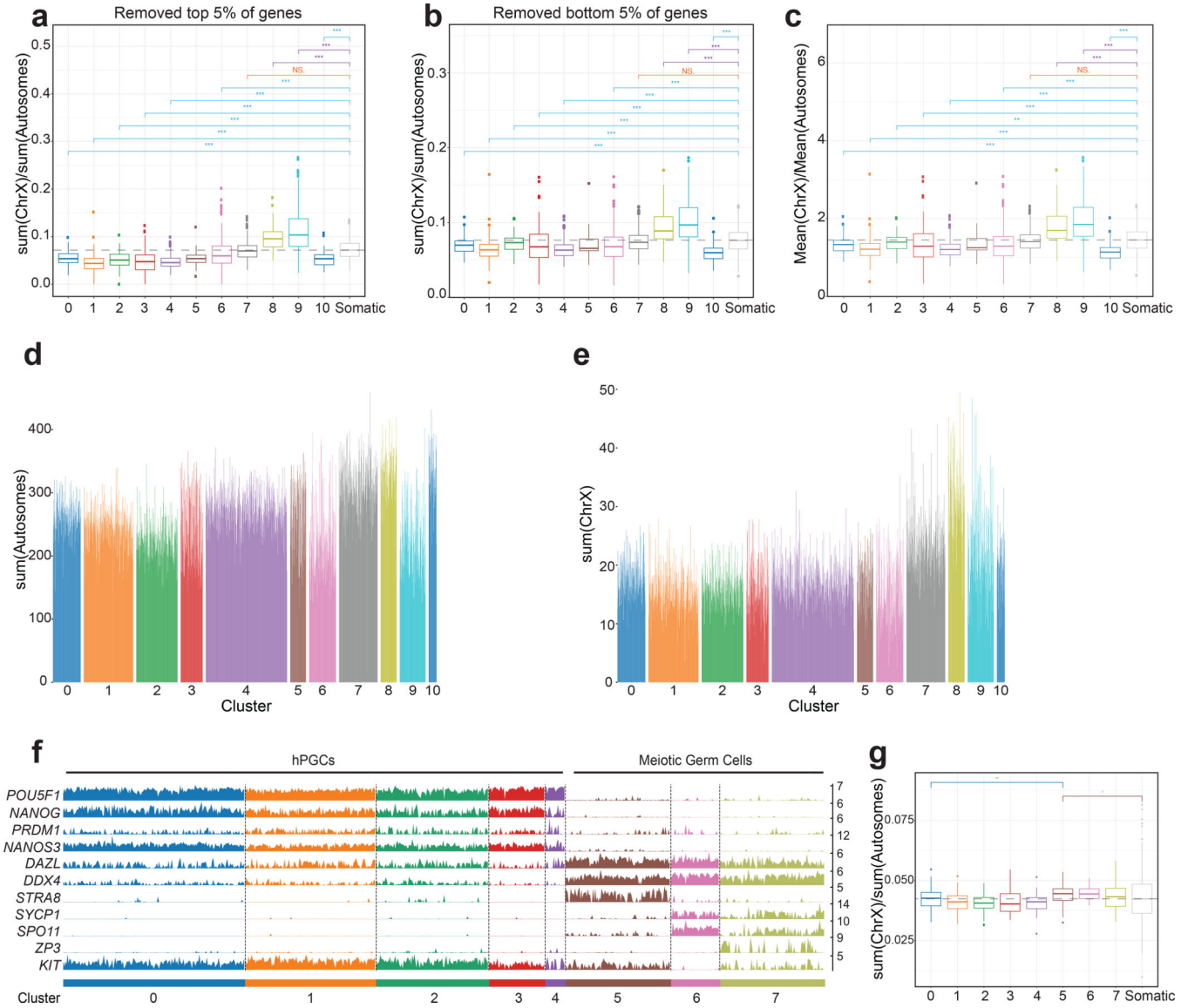
a. Distribution of single nucleotide polymorphisms (SNPs) from maternally inherited (*x*-axis) and paternally inherited (*y*-axis) alleles in gene expression data of female hPGCs. Each dot represents sum of all detected SNPs per cell for genes on chromosome 1 (Chr1), X-linked genes subject to XCI and escapees of XCI, respectively, based on published scRNA-seq data³⁰. **b.** Representative immunofluorescence staining of OCT4 (magenta), H3K27me3 (green), DAZL (grey) and DAPI (blue) on female hPGCs at week 4 pf prior to gonad formation, when hPGCs are migrating through the aorta-gonad-mesonephros (AGM) (1 sample was analyzed). Insets show a rare OCT4+/DAZL+ cell with no nuclear accumulation of H3K27me3 (inset 1) and an OCT4+/DAZL- cell with H3K27me3 accumulation (inset 2) along the genital ridge of the AGM. Scale bar upper panel 50 microns, lower panel 30 microns. **c.** Percentage of OCT4+/DAZL+ and OCT4+/DAZL- cells with an Xi-like nuclear accumulation of H3K27me3 from the experiment shown in (c); (n=58 cells from 1 AGM). **d.** Quantification of the proportion of DAZL+ female hPGCs at weeks 4, 7 and 12 pf with an Xi-like nuclear accumulation of H3K27me3 (n=50–100 cells per sample in 2 replicates). **e.** Representative immunofluorescence staining of a fetal ovary at week 12 pf with DAZL (magenta), H3K27me3 (green), OCT4 (grey) and DAPI (blue). Inset shows a DAZL+/OCT4 negative female germ cell that is negative for H3K27me3 (1 pair of ovaries were analyzed), scale bar 50 microns. Statistical source data are provided in Source data extended data fig. 1.



Extended Data Fig. 2. The X/A ratio is higher in female germ cells than in male germ cells

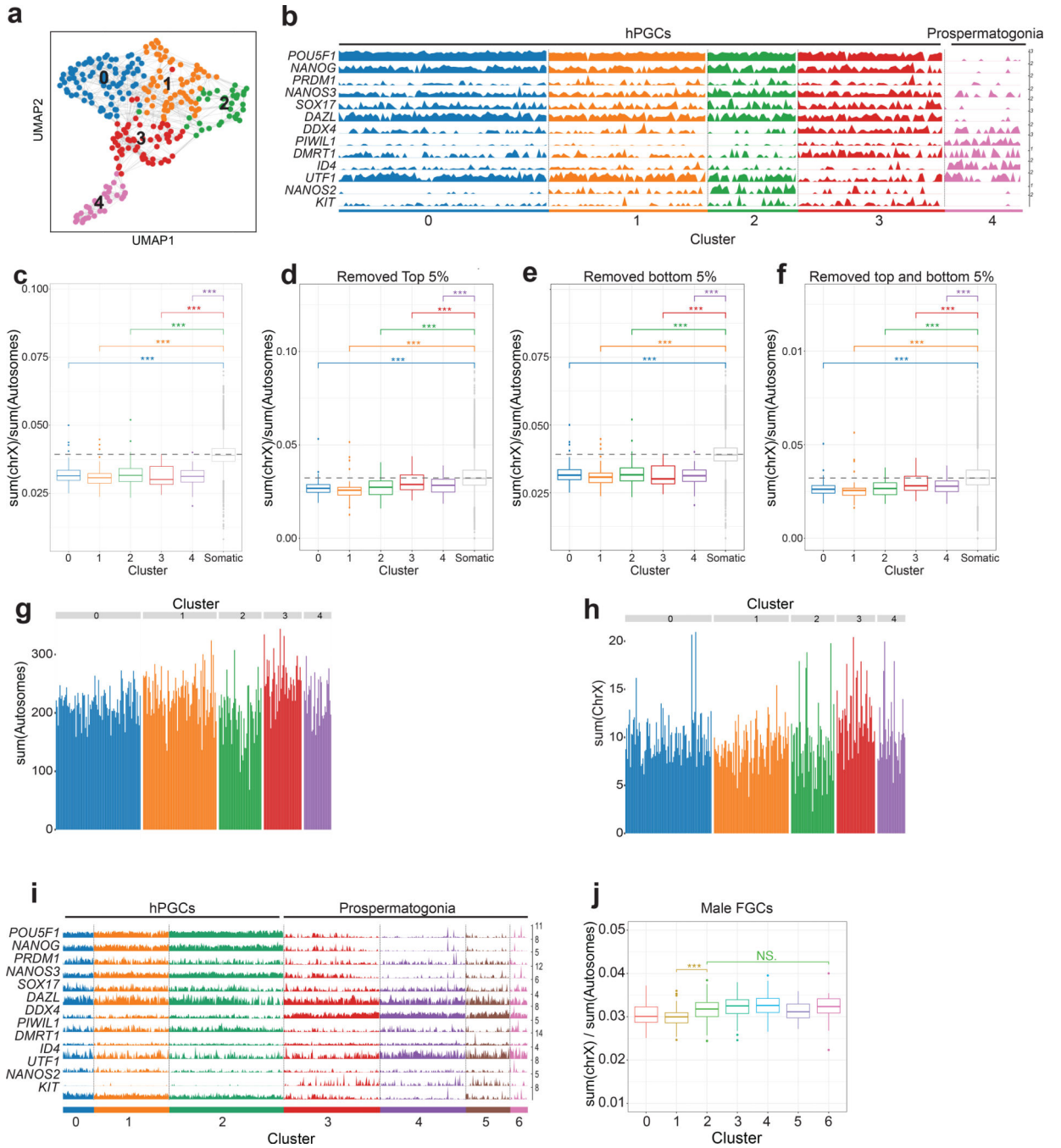
a. Boxplots presenting the X/A ratio, calculated from the sum of X-linked gene expression and the sum of autosomal gene expression, of individual female (red) or male (cyan) germ cells (left panel) and surrounding somatic cells (right panel) obtained from gonads harvested from indicated developmental timepoints (week). **b.** As in (a), except that the X/A ratio was determined from the mean expression levels of X-linked and autosomal genes per cell. **c.** Boxplot showing the distribution of the sum of all autosomal (left) and X-linked (right) gene expression, respectively, in individual female and male germ cells across

for developmental time points shown in (a). **d.** As in (a), except that the X/A ratios in female and male FGCs across developmental time from a published study are shown⁴⁰. **e.** X/A ratio per single cell in female and male gonadal somatic cells from all developmental ages accompanying the data shown in (d). Wilcoxon statistical testing between age matched samples, NS- Not Significant, * p<0.05, ** p<0.01, *** p<0.001. a-c: n = 49528 cells analyzed across 10 independent experiments and d-e: 1016 cells analyzed from published dataset⁴⁰ in total.



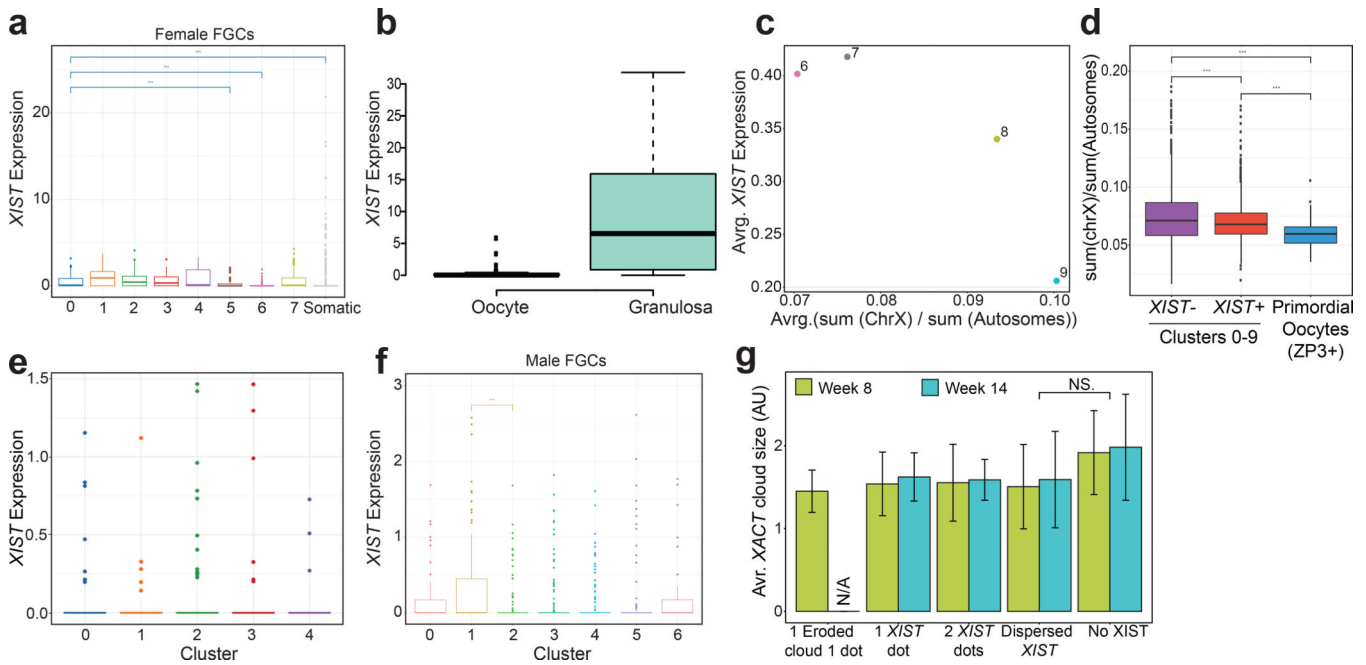
Extended Data Fig. 3. Female hPGCs dampen X linked gene expression before entering meiosis
a. Boxplots of the X/A ratios of female germ cells along the developmental trajectory and in female gonadal somatic cells, as described in Figure 5, except that the top 5% highest expressed genes were excluded from the analysis. **b.** As in (a), except that the bottom 5% of expressed genes were excluded from the analysis. **c.** Boxplots of the X/A ratios in female germ cells and female gonadal cells as described in Figure 5, except that the X/A ratios were

calculated from the mean value of X-linked and autosomal gene expression per cell. **d.** Sum of all autosomal gene expression normalized counts in female germ cells organized by clusters along the developmental trajectory. **e.** Sum of all X-linked gene expression in female germ cells organized by clusters along the developmental trajectory. X-linked gene expression increases in clusters 7–9 coincident with entrance into meiosis and repression of the naïve-like pluripotency program. **f.** Germline trajectory analysis of previously published scRNA-seq data from female FCGs⁴⁰. The hPGCs state with pluripotency program expression is captured with clusters 0–4, and meiotic entry in cluster 5–7. **g.** Boxplots of the X/A ratios for female germ cells and female gonadal somatic cells for the data set shown in (f). From cluster 5 onwards, X/A ratios in differentiating female germ cells are higher than gonadal somatic cells. Wilcoxon statistical testing for (a), (b), (c), (g). NS- Not Significant, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.



Extended Data Fig. 4. Male hPGCs do not change X/A ratio upon sex specific differentiation
a. Male germ cells from the scRNA-seq data shown in Figures 4a–c were projected along the developmental trajectory, and five clusters (0–4) were identified (n= 282 cells pulled from 5 samples). The pluripotency program is repressed in cluster 4, coincident with increased expression of prospermatogonia genes and exit from the cell cycle. **b.** Expression of marker genes along the developmental trajectory of male germ cells defined in (a). **c.** Box plots showing that X/A ratios in male germ cells along the developmental trajectory and in surrounding male somatic cells. **d.** As in (c), except that the top 5% highest expressed genes

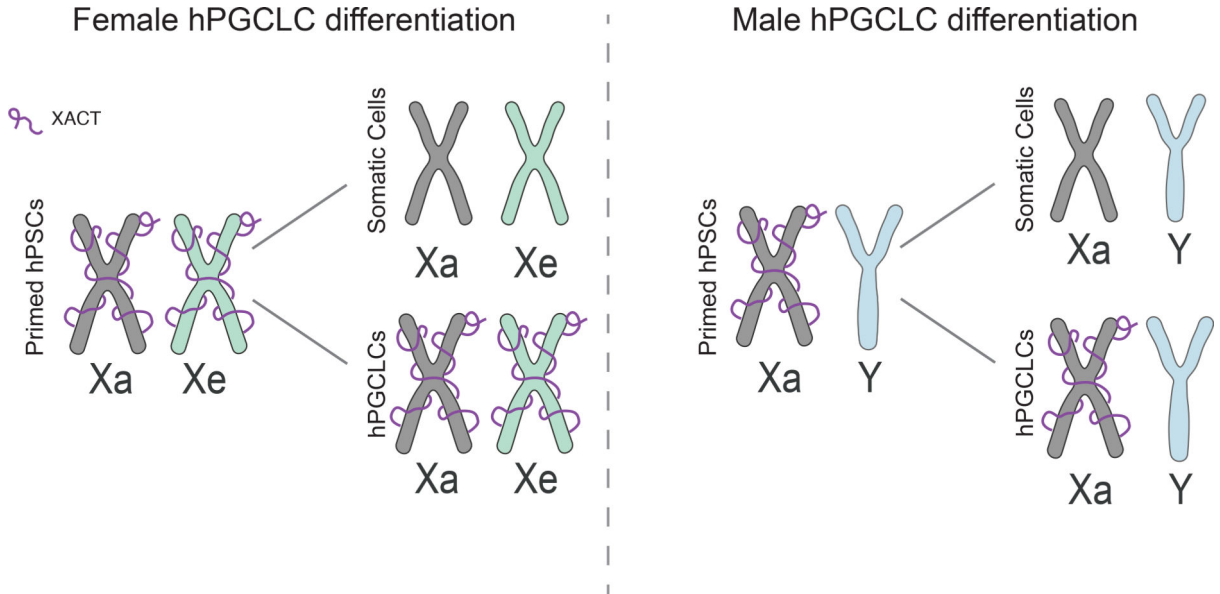
were excluded from the analysis. **e.** As in (c), except that bottom 5% of genes were excluded. **f.** As in (c), except that the top and bottom 5% of expressed genes were excluded. In total $n = 24740$ cells analyzed across 5 independent experiments in c-f. **g.** Sum of all autosomal gene expression per cell in male germ cells along the developmental trajectory, showing no dramatic differences across the clusters. **h.** As in (g), except for X-linked gene expression, showing no dramatic differences across the clusters. **i.** Germline trajectory analysis for male FGCs⁴⁰, identified 7 clusters (0–6). Marker gene expression is given for these clusters. **j.** Box plots of the X/A ratios in male FGCs along the developmental trajectory defined in (i), showing an increase in cluster 2 relative to cluster 1. In total, $n = 779$ cells analyzed from published dataset⁴⁰ in **j.** Wilcoxon statistical testing used for (c-f) and (j). NS- Not Significant, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.



Extended Data Fig. 5. *XIST* expression correlates with the X/A ratio

a. Boxplot depicting the expression of *XIST* in female germ cells organized by clusters along the developmental trajectory defined for the published FCG data set⁴⁰ in Extended Data Fig. 3f,g. *XIST* expression is significantly reduced from cluster 5 onwards. **b.** *XIST* expression in mature oocytes and granulosa cells from scRNA-seq data of adult ovary⁴⁴. **c.** Scatter plot of average *XIST* expression (y-axis) and average X/A ratio (x-axis) for female germ cells clusters 6–9 (from Figure 4d), capturing the entrance into meiosis. **d.** X/A ratios in female hPGCs and meiotic germ cells clustered based on expression of lncRNA *XIST*, *ZP3+* primordial oocytes are clustered separately. **e.** Boxplots depicting *XIST* expression in male germ cells ordered along the developmental trajectory defined in Extended Data Fig. 4a–h, indicating that *XIST* transcripts are rarely detected in male germ cells. **f.** As in (e), except for male FGCs⁴⁰ from Extended Data Fig 4i,j. **g.** Average *XACT* cloud size in week 8 and 14 pf hPGCs with different patterns of *XIST* expression, error bars show standard deviation of the cloud sizes (76 cells analyzed in total from 2 independent experiments). Wilcoxon statistical testing used for (a), (d), (f), (g). NS- Not Significant, * $p < 0.05$, **

$p < 0.01$, *** $p < 0.001$. Number of cells analyzed across 5 independent experiments: **a.** $n = 1016$ cells⁴⁰, **b.** $n = 148$ cells⁴⁴, **d.** $n = 1938$ cells, **f.** $n = 779$ cells⁴⁰.



Extended Data Fig. 6. *XACT* marks male and female hPGCLCs *in vitro*

Summary of the hPSC differentiation figure shown in Figure 2. Due to XCI erosion, *XACT* is expressed from the Xa and the eroded X-chromosome in primed, female human pluripotent stem cells. The Xe state is transmitted into differentiated cells and upon hPGCLC differentiation. Moreover, *XACT* is maintained in hPGCLCs, whereas somatic cells silence *XACT*. Consequently, female hPSC-derived PGCLCs carry two *XACT* clouds and male hPSC-derived PGCLCs one.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Data availability

The accession number for the scRNA-seq data of prenatal tissues reported in this paper are GSE143380 (female cell data) and GSE143356 (male cell data). scRNA-seq data sets are also available for interactive exploration at <http://germline.mcdb.ucla.edu>.

Previously published RNA-seq data of male and female hPGCs and somatic cells^{25,52,53} and single-cell RNA-seq data from female germ cells³⁰ and from female FCGs⁴⁰ that were re-analyzed here are available under accession codes: GSE63392²⁵, GSE60138⁵², GSE93126⁵³, GSE79280³⁰ and GSE86146⁴⁰ respectively. All other data supporting the findings of this study are available from the corresponding author on reasonable request.

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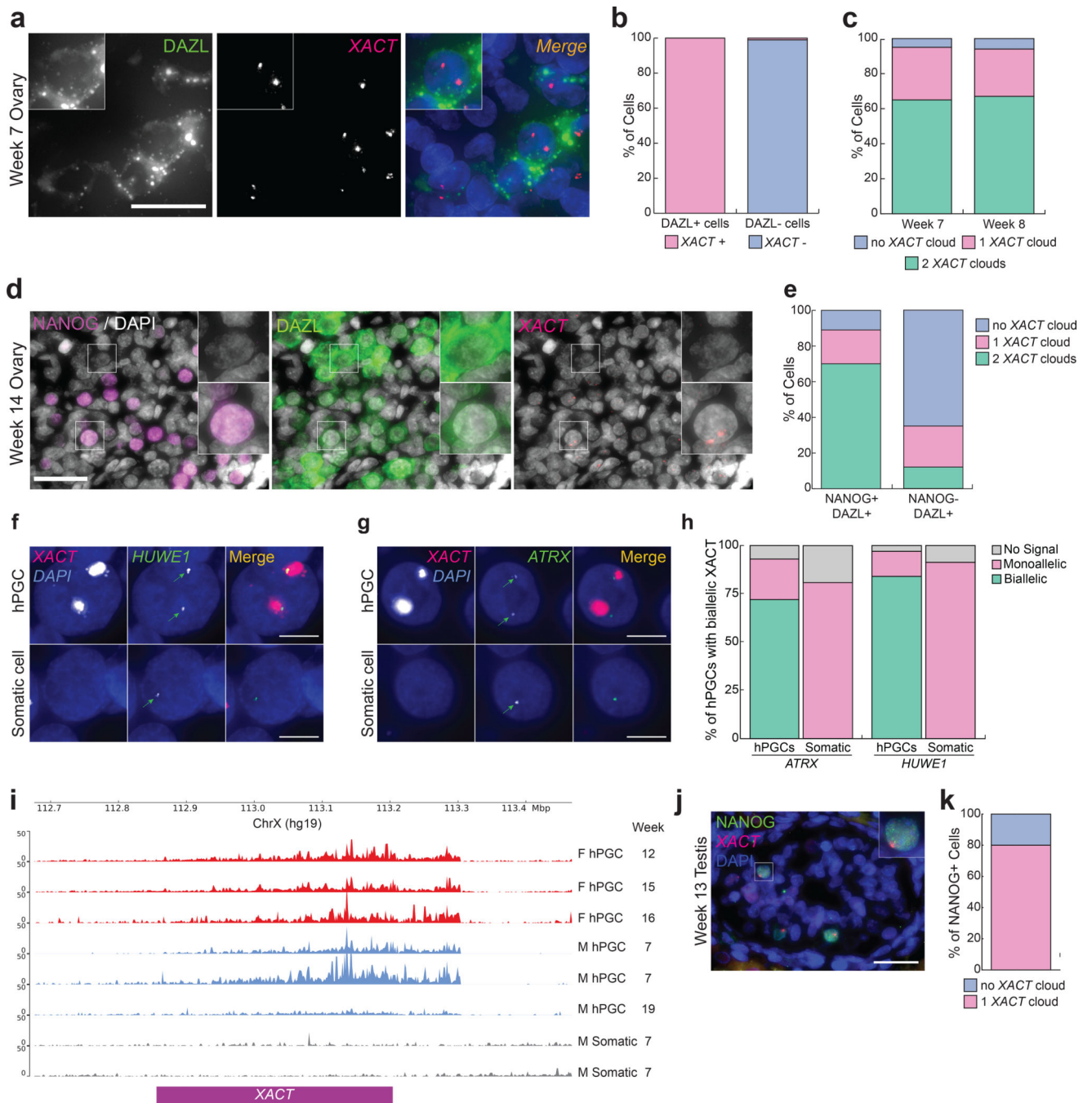


Fig. 1. Male and female hPGCs express lncRNA *XACT* and female hPGCs carry two active X chromosomes *in vivo*

a. Immuno-RNA FISH for *DAZL* (green) and *XACT* (red) in a week 7 pf embryonic ovary (1 pair of ovaries were analyzed) with DAPI staining (blue) to detect nuclei, scale bar 30 microns. **b.** Quantification of cells with *XACT* clouds based on the experiment in (a) (n=100 cells). **c.** Quantification of the number of *XACT* clouds in *DAZL*+ hPGCs at week 7 and 8 pf (n=100 cells/timepoint). **d.** Immuno-RNA FISH for *XACT* (red), *NANOG* (magenta), *DAZL* (green) and DAPI (grey) in a week 14 pf fetal ovary (1 pair of ovaries were

analyzed). Top inset shows a NANOG-/DAZL+ hPGC negative for *XACT*, lower inset a NANOG+/DAZL+ hPGC with two *XACT* clouds; scale bar 30 microns. **e.** Quantification of the proportion of cells with a different *XACT* cloud patterns in hPGCs (NANOG+/DAZL+) and differentiating hPGCs (NANOG-/DAZL+) from (d). 92 and 95 cells, respectively, were assessed. **f-g.** Representative RNA FISH images for detection of nascent transcripts of the X-linked genes *HUWE1* and *ATRX*, respectively, both normally subject to XCI, in a week 8 pf ovary. hPGCs are marked by *XACT* expression. The experiments have been performed twice with similar results. **h.** Signal quantification for (g); n=60 and n=70 cells for *ATRX* and *HUWE1*, respectively. **i.** Published bulk RNA-seq read mapped to the *XACT* genomic locus in female hPGCs (red, isolated using cKIT^{bright} or using INTa6/EpCAM), male hPGCs (blue, enriched for TNAP/KIT expression) and gonadal somatic cells (grey)^{25,52,53}. **j.** Immuno-RNA FISH in fetal male testis at week 13 pf (1 pair of testis were analyzed) with *XACT* (red), NANOG (green) and DAPI (blue), scale bar 30 microns. **k.** Quantification of the proportion of cells with one *XACT* cloud in NANOG+ male hPGCs from (j) (n=75 cells). Statistical source data are provided in Source data fig. 1.

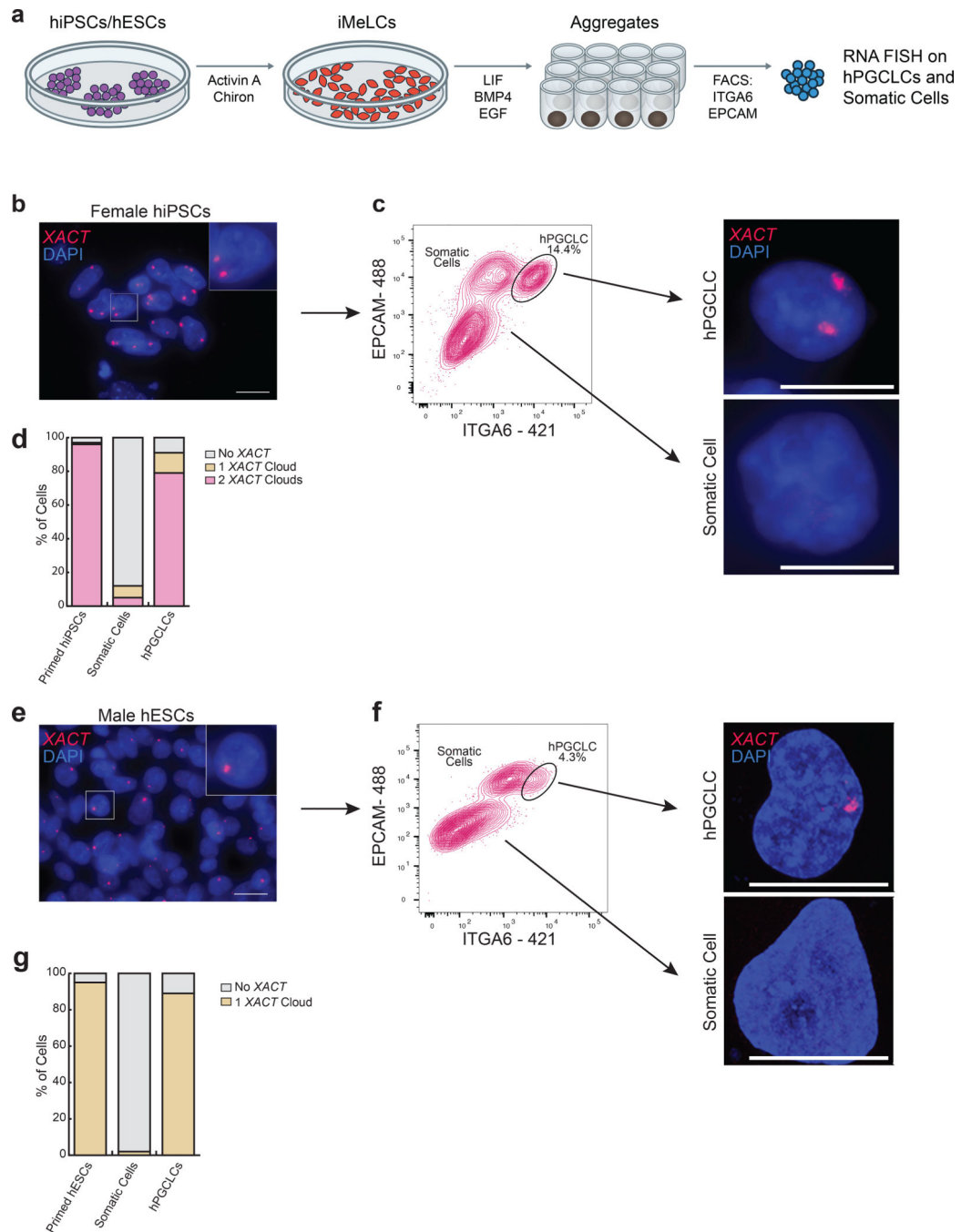


Fig. 2. The lncRNA *XACT* is restricted to male and female hPGCLCs and is not expressed in somatic cells *in vitro*

a. Differentiation of hPGCLCs from hiPSCs or hESCs through a incipient mesoderm like cells (iMeLC) intermediate. PGCLCs and somatic cells within the aggregates are separated at day 4 (D4) by FACS using antibodies that recognize EPCAM and ITGA6. **b.** RNA FISH for *XACT* in primed female hiPSCs (MZT04-J), that harbor a Xa and an eroded X-chromosome (Xe) with *XACT* (red) and DAPI (blue) stainings. *XACT* clouds are detected from the Xa and Xe; scale bar 20 microns. The experiments have been performed twice with

similar results. **c.** Female hiPSCs are differentiated to hPGCLCs and isolated from the aggregates by FACS at D4. The hPGCLC population is indicated. Right panel: *XACT*RNA FISH in hPGCLCs and somatic cells. The experiments have been performed twice with similar results; scale bar 10microns. **d.** Quantification of the proportion of cells with different numbers of *XACT* clouds in starting hiPSCs (n=100 cells), hPGCLCs (n=82 cells) and somatic cells (n=100 cells) from (c). **e.** *XACT*RNA FISH (red) in male hESCs (UCLA2); scale bar 20 microns similar to (b). **f.** As in (c), except for UCLA2 hESCs; the experiments have been performed twice with similar results, scale bar 10microns. **g.** Quantification of the proportion of cells with a different number of *XACT* clouds in UCLA2 hESCs (n=100 cells) and derived hPGCLCs (n=92 cells) and somatic cells (n=100 cells pooled from the two experiments). Statistical source data are provided in Source data fig. 2.

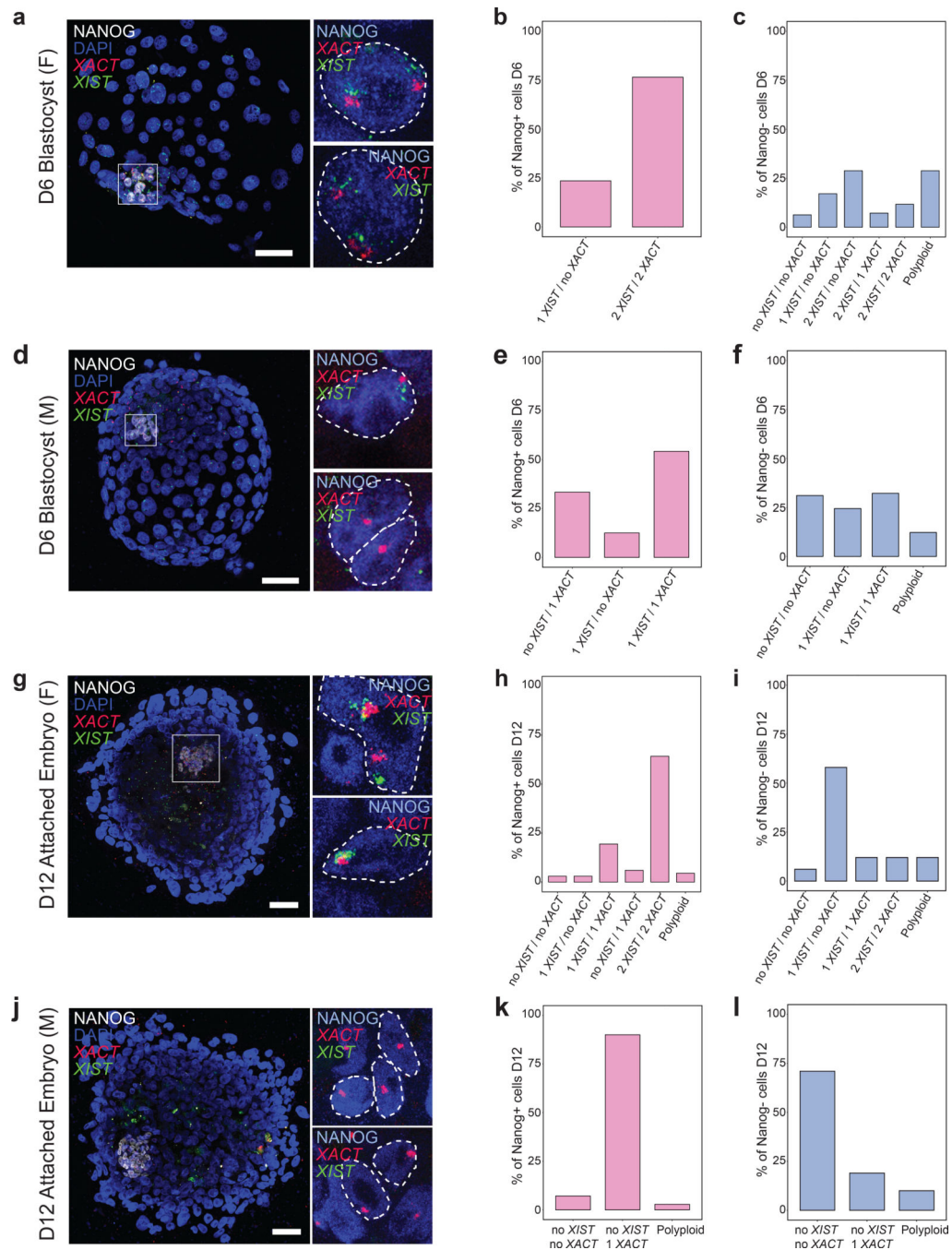


Fig. 3. XACT is predominantly expressed in NANOG+ pre- and post-implantation epiblast cells a, d, g, j. Immunofluorescence for NANOG (white), XIST (green) and XACT (red) in: **a.** day 6 (D6) intact female (F) pre-implantation blastocysts (2 blastocysts were analyzed); **d.** D6 male (M) pre-implantation blastocysts (3 blastocysts were analyzed); **g.** female embryo cultured to D12 using human embryo attachment culture (3 embryos were analyzed); **j.** male human embryos cultured to D12 using human embryo attachment culture (2 embryos were analyzed); scale bar 30 microns. Insets for a, d, g, j are showing NANOG+ (blue) nuclei with XIST and XACT clouds. **b.** Quantification of the proportion of cells with different

numbers of *XIST* and *XACT* clouds in *NANOG*⁺ epiblast cells from female blastocysts in (a); 17 cells analyzed in 2 blastocysts. **c.** As in (b), except for quantification of the RNA pattern in *NANOG*-negative trophoctoderm and primitive endoderm cells, with 111 cells quantified from the two blastocysts. **e.** As in (b), except for that *NANOG*⁺ epiblast cells from male blastocysts in (d) were assessed; with 24 cells from 3 blastocysts counted. **f.** Quantification as in (c), except for *NANOG*-negative trophoctoderm and primitive endoderm cells from male blastocysts in (d); 180 cells from 3 blastocysts were counted. **h.** Quantification as in (b) for female D12 embryos from (g); 67 cells from 3 embryos were counted. **i.** Quantification as in (c) for female D12 embryos in (g); with 180 cells from 3 embryos counted. **k.** Quantification as in (b) for male D12 embryos from (j); with 70 cells from 2 embryos assessed. **l.** Quantification as in (c) for male D12 embryos in (j); with 120 cells from 2 embryos quantified. Statistical source data are provided in Source data fig. 3.

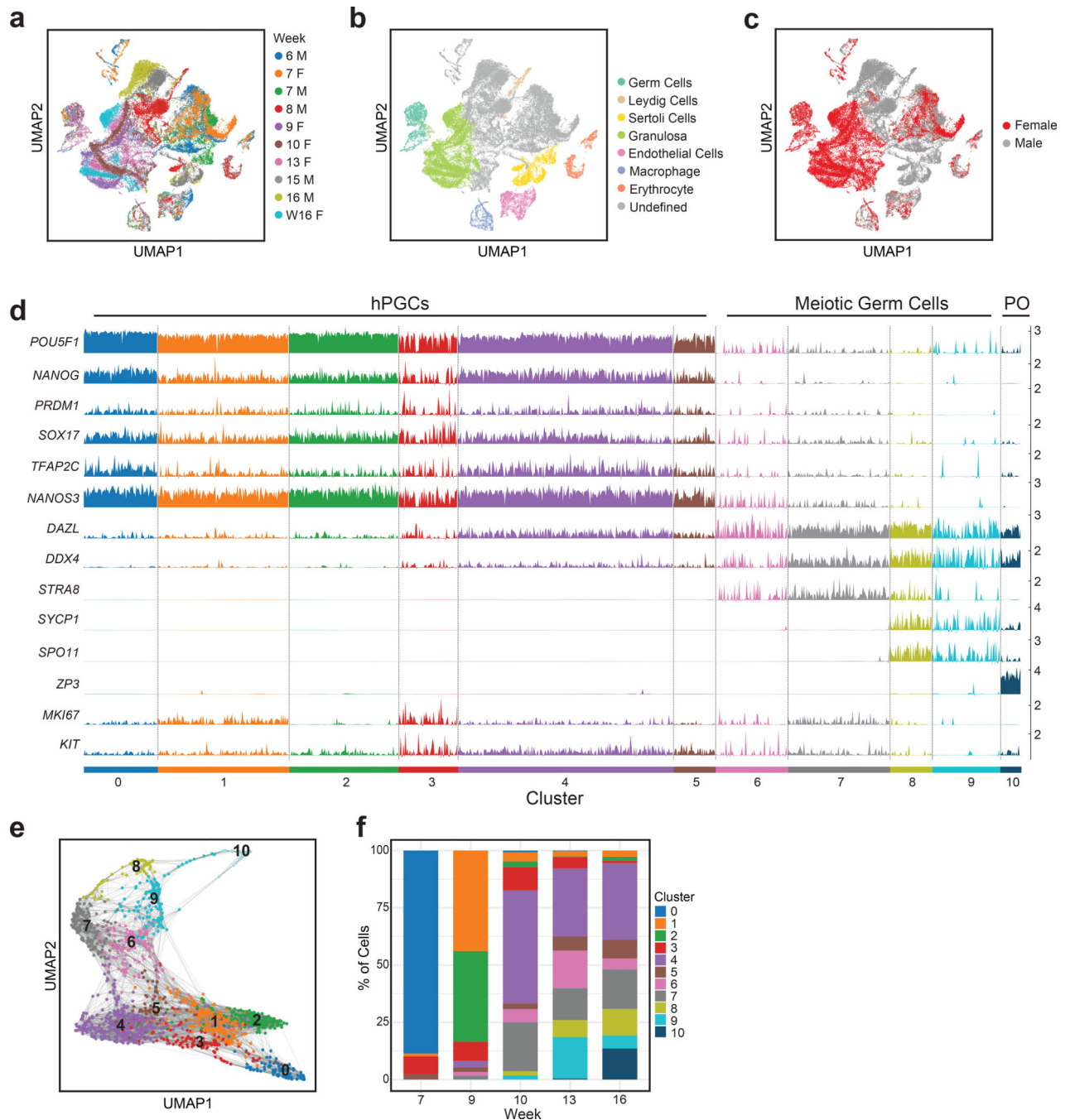


Fig. 4. Heterogeneous differentiation of female hPGCs into meiotic germ cells begins around week 9–10 pf

a. Distribution of single cell data derived from scRNA-seq from five prenatal male and five prenatal female gonads from 6–16 weeks pf on UMAP ($n=49674$ cells). **b.** Annotation of gonadal cell types in the map from (a), based on the expression of cell type-specific markers. **c.** Distribution of male and female cells on the map from (a). **d.** Ordering of female germ cells along the developmental trajectory from cluster 0 to cluster 10, with classification into hPGCs (clusters 0–5), meiotic germ cells (clusters 6–9) and primordial oocytes (PO, cluster

10), based on diagnostic germ cell marker gene expression. Each cluster contains many individual cells (columns), for which expression of indicated marker is given (rows). n=1938 cells from n=5 samples. **e.** Female germ cells displayed on UMAP plot, labelled by their cluster assignment from (d). **f.** For each of the five female gonads described in (a), the proportion of cells in the clusters defined in (d) is given. These data show that the repression of the pluripotency expression program and meiotic licensing (expression of *STRA8*) begins between 9–10 weeks pf, and that all germ cells at week 7 are in the hPGC state. Statistical source data are provided in Source data fig. 4.

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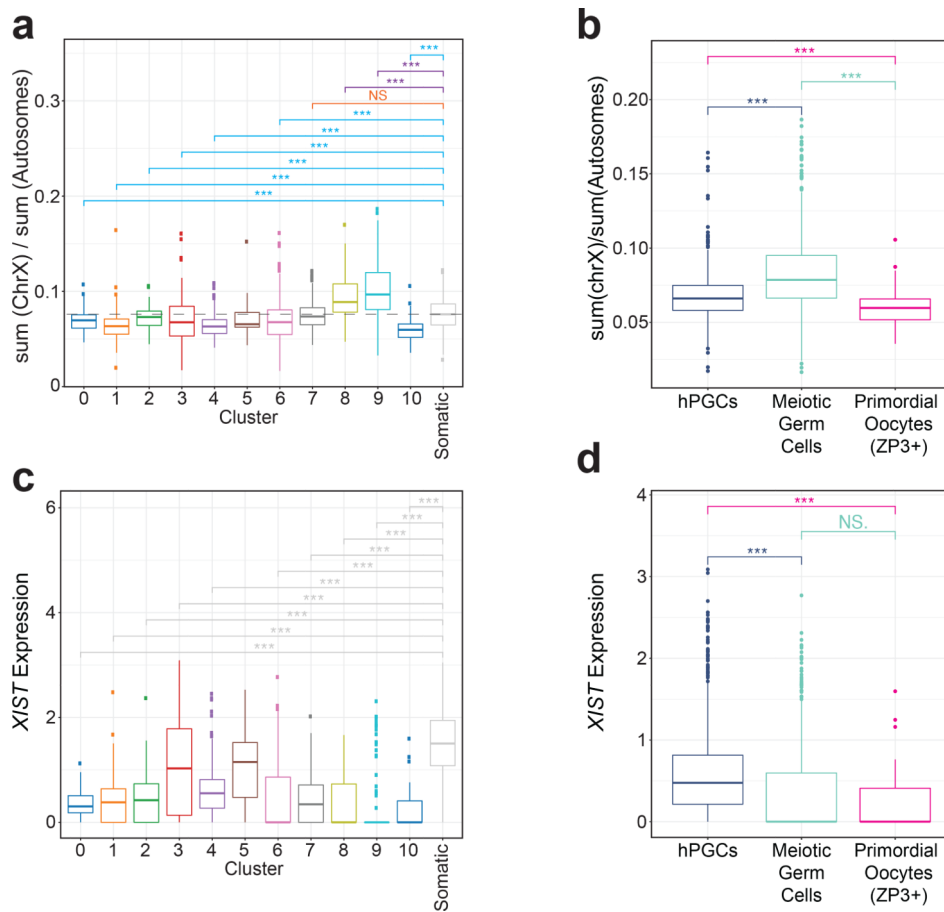


Fig. 5. *XIST* expression is associated with XCD in female hPGCs

a. Box plots of the ratios of X-linked gene to autosomal gene expression in single female germ cells ($n=1938$ cells), for each cluster along the developmental trajectory described in Figure 4d. In addition, the X/A ratio of gonadal somatic cells are given. The dotted line indicates the median level of the X/A ratio in female somatic gonadal cells. Significance testing given with blue lines shows clusters in which the X/A ratio is significantly lower compared to that in the somatic cell cluster; the orange line shows that the difference was not significant; and the purple line shows comparisons where the X/A ratio was significantly higher than in the somatic cell cluster. **b.** Box plots of the X/A ratios of individual female germ cells in the hPGC-state, meiotic germ cell-state and *ZP3+* primordial oocytes, merged based on developmental classification (hPGC-state, clusters 0–5; meiotic germ cells, clusters 6–9; and *ZP3+* primordial oocytes, cluster 10). **c.** Box plots of normalized counts of *XIST* transcripts in individual female germ cells for each cluster along the developmental trajectory described in Figure 4d and female gonadal somatic cells. These results show female hPGCs express significantly lower levels of *XIST* and an abrupt loss of *XIST* coincident loss of the pluripotency program in cluster 6. **d.** Box plots of normalized *XIST* transcript counts in individual female germ cells, as defined in (b). Significance was assessed with the Wilcoxon Test; NS-Not Significant, *** $p < 0.001$. $n=1938$ cells analyzed across 5 independent experiments.

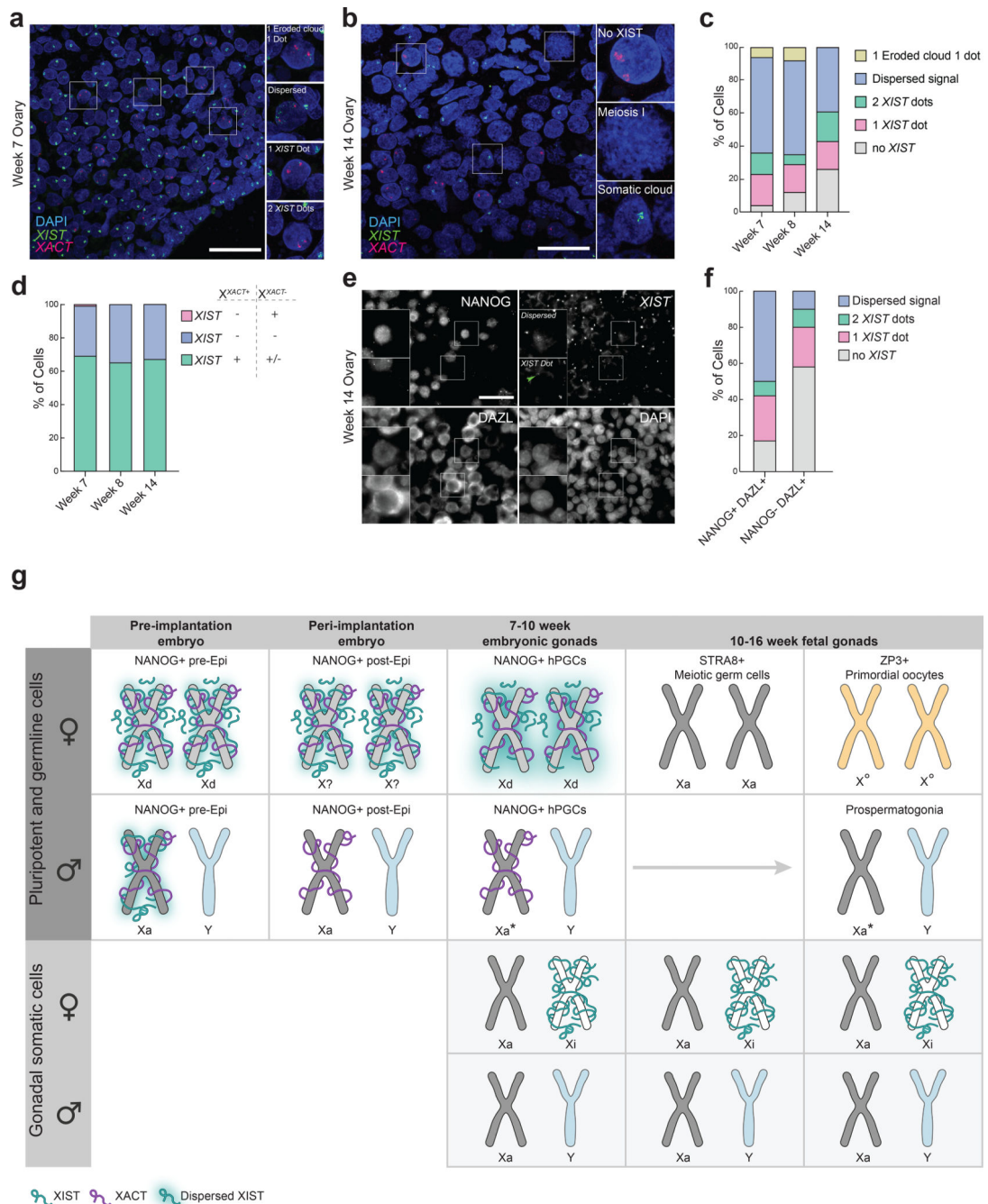


Fig 6. XIST is repressed as hPGCs enter meiosis

a. RNA FISH for *XIST* (green) and *XACT* (red) with DAPI (blue) identifying nuclei (week 7 n=1 pair of ovaries); scale bar 30 microns. **b.** As in (a), except for week 14 pf 1 pair of ovaries. **c.** Proportion of cells with indicated *XIST* expression patterns in female hPGCs with 2 *XACT* clouds (for each time point: n=100 cells from 1 pair of ovaries). **d.** Similar to (c) except in female hPGCs with a single *XACT* cloud. Cells with dispersed *XIST* signal where it was not clear to identify if *XIST* was expressed from one or both alleles are included in green category (total n=42 cells from 3 ovaries). **e.** Immuno-RNA FISH image for NANOG,

DAZL and *XISTRNA* in week 14 pf ovary (1 pair of ovaries); scale bar 30 microns. **f.** Similar to (c) in female hPGCs (NANOG+/DAZL+) and meiotic germ cells (NANOG-/DAZL+) from (e) (n=104 and 98 cells, respectively from 1 sample). **g.** Female NANOG+ pre- and post-implantation epiblast (Epi) cells predominantly express *XACT* and *XIST* from both X-chromosomes. While genes on both X-chromosomes in pre-Epi cells are dampened due to XCD (Xd)¹⁴ it is unclear whether XCD also occurs in female post-Epi cells. However, expression of both *XACT* and *XIST* from both X-chromosomes indicates that XCI has not occurred yet. NANOG+ female hPGCs similar to pre-Epi exhibit XCD with expression of *XIST* and *XACT* from both alleles. Notably, *XIST* is more dispersed in the nucleus of hPGCs (depicted by glow around the X-chromosome) than pre/post-Epi cells. Upon advancement to meiosis they silence both *XIST* and *XACT* and upregulate X-linked gene expression, transitioning to the Xa-state. Next, in primordial oocytes, the X/A ratio is reduced lower than in female hPGCs without expression of *XIST*. This *XIST*-independent repression of the X/A ratio we call ‘oocyte-specific X-chromosome regulation’ (X°). Male hPGCs harbor an Xa but exhibit a lower X/A ratio compared to male somatic cells, and this state is retained upon differentiation into prospermatogonia. We refer to this state as Xa*. Gonadal somatic cells display XCI at each stage analyzed.