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Retinoic Acid Signalling and the Control of Meiotic Entry in the Human Fetal Gonad

Andrew J. Childs1,*, Gillian Cowan1,*, Hazel L. Kinnell1, Richard A. Anderson2, Philippa T. K. Saunders1

1 Medical Research Council Human Reproductive Sciences Unit, The Queen’s Medical Research Institute, University of Edinburgh, Edinburgh, United Kingdom, 2 Division of Reproductive and Developmental Sciences, Centre for Reproductive Biology, The Queen’s Medical Research Institute, University of Edinburgh, Edinburgh, United Kingdom

Abstract

The development of mammalian fetal germ cells along oogonic or spermatogenic fate trajectories is dictated by signals from the surrounding gonadal environment. Germ cells in the fetal testis enter mitotic arrest, whilst those in the fetal ovary undergo sex-specific entry into meiosis, the initiation of which is thought to be mediated by selective exposure of fetal ovarian germ cells to mesonephros-derived retinoic acid (RA). Aspects of this model are hard to reconcile with the spatiotemporal pattern of germ cell differentiation in the human fetal ovary, however. We have therefore examined the expression of components of the RA synthesis, metabolism and signalling pathways, and their downstream effectors and inhibitors in germ cells around the time of the initiation of meiosis in the human fetal gonad. Expression of the three RA-synthesising enzymes, ALDH1A1, 2 and 3 in the fetal ovary and testis was equal to or greater than that in the mesonephros at 8–9 weeks gestation, indicating an intrinsic capacity within the gonad to synthesise RA. Using immunohistochemistry to detect RA receptors RARα, β and RXRα, we find germ cells to be the predominant target of RA signalling in the fetal human ovary, but also reveal widespread receptor nuclear localization indicative of signalling in the testis, suggesting that human fetal testicular germ cells are not efficiently shielded from RA by the action of the RA-metabolising enzyme CYP26B1. Consistent with this, expression of CYP26B1 was greater in the human fetal ovary than testis, although the sexually-dimorphic expression patterns of the germ cell-intrinsic regulators of meiotic initiation, STRA8 and NANOS2, appear conserved. Finally, we demonstrate that RA induces a two-fold increase in STRA8 expression in cultures of human fetal testis, but is not sufficient to cause widespread meiosis-associated gene expression. Together, these data indicate that while local production of RA within the fetal ovary may be important in regulating the onset of meiosis in the human fetal ovary, mechanisms other than CYP26B1-mediated metabolism of RA may exist to inhibit the entry of germ cells into meiosis in the human fetal testis.

Introduction

Primordial germ cells (PGCs) are the embryonic precursors of sperm and egg in the adult organism. Although initially bipotential, with the capacity to adopt spermatogenic or oogenic fates, the developmental trajectory of PGCs is dictated by the somatic sex of the gonad in which they find themselves following migration [1]. PGCs which find themselves in a female (ovarian) environment enter meiosis from embryonic day (e)13.5 in the mouse and 11 weeks gestation in the human, whilst germ cells in the developing testis progressively enter a state of cell cycle arrest, resuming proliferation and differentiation only after birth [2].

The mechanism(s) by which this dimorphism in meiotic entry is established has long been a matter of debate. Recent data have suggested that meiosis is initiated in the fetal mouse ovary by retinoic acid (RA) signalling, which acts on germ cells to promote the expression of Stimulated by Retinoic Acid (Stra8) [3,4], a protein required for pre-meiotic DNA replication [5]. Germ cells in the fetal mouse testes are shielded from the meiosis-inducing action of RA, first by somatic cell expression of the RA-metabolising enzyme Cyp26b1 [3,4], then subsequently by the action of the germ cell-expressed RNA-binding protein Nanos2, which acts to ‘lock in’ the male germ cell differentiation program and repress Stra8 expression in testicular germ cells following the downregulation of Cyp26b1 [6].

RA is a potent morphogen that exerts diverse effects during development and differentiation [7,9,9]. It is tightly regulated by a group of RA synthesising- and metabolising-enzymes. The retinaldehyde dehydrogenase enzymes (Aldh1a 1,2 and 3) are responsible for the oxidation of RA precursors to produce RA [10,11], while RA signalling is negatively regulated by three RA degrading enzymes, Cyp26a1, Cyp26b1 and Cyp26c1, which...
metabolize RA into hydroxylated polar derivatives [12]. Although undetectable in the fetal gonad itself, expression of the RA synthesis enzymes Aldh1a2 and Aldh1a3 has been demonstrated in mesonephroi of fetal mice between e11.5 and 13.5, and the mesonephros has been shown to synthesis high levels of RA at this stage [3]. A source-sink model of meiotic entry in the fetal mouse gonad has therefore been proposed [3,4], in which the delivery of mesonephros-derived RA into the anterior portion of the gonad, and its subsequent diffusion along the gonadal axis, results in the entry of germ cells in the fetal ovary into meiosis in a rostro-caudal (anterior-posterior) wave, with expression of PGC/pluripotency-associated markers such as Oct4 downregulated [13,14,15], and markers of meiosis such as Stra8, Synaptonemal Complex Protein 3 (Sycp3) and the Dosage suppressor of mck1 homologue 1 (Dmc1) [14,15] upregulated.

In contrast to the rodent however, germ cell differentiation in the human fetal ovary does not occur in a synchronized rostro-caudal wave. Germ cells at different developmental stages are instead arranged radially, with undifferentiated PGCLike OCT4-positive/VASA-negative germ cells present at the periphery of the organ and more differentiated OCT4-negative/VASA-positive germ cells deeper within the medulla [16,17]. Significantly, these subpopulations exist in parallel, overlapping in space and time such that undifferentiated OCT4-positive cells are still detectable at the periphery of the ovary several weeks after the first germ cells enter meiosis, and even beyond the onset of follicle formation [16]. This suggests that local control of germ cell differentiation may play a greater role in the human fetal ovary than occurs in the rodent at the equivalent developmental stage, a hypothesis supported by the recent demonstration of intrinsic RA synthesis and metabolism by the human fetal ovary [18].

The aim of this study was to determine whether similar mechanisms to those reported to control the initiation of meiosis in mouse fetal germ cells also operate to control this process in the human fetal gonad, by examining the expression and localization of key components of the retinoid synthesis, signalling and effector machinery in the human fetal tests and ovary across the period of meiotic initiation. We report differences in the expression and localization of RA metabolising enzymes in the developing human gonad to those reported in mice at similar developmental stages, notably greater expression of CYP26B1 in the ovary compared to the testis. We find germ cells to be the primary target of retinoid signalling in the human fetal ovary, but reveal RA receptor expression – and activation – to be widespread in the human fetal tests, indicating that RA metabolism does not fully shield human fetal testicular cells from RA signalling activity. Finally, we demonstrate that RA has the capacity to induce STRA8 expression in the human fetal testis, but does not increase expression of other genes associated with the initiation of meiosis. Together these data suggest that the control of meiotic initiation in the human fetal ovary may not be controlled exclusively by retinoid signalling and metabolism, and that there may be greater emphasis on the regulation of meiosis at a local, rather than whole-organ level in the human fetal ovary, than occurs in mouse.

**Results**

**Expression of the genes encoding retinaldehyde dehydrogenase enzymes during human fetal gonadal development**

The mesonephros of the developing mouse embryo is thought to be the site of synthesis of meiosis-inducing RA, and mesonephric (but not gonadal) expression of the genes encoding RA synthesising enzyme (Aldh1a2 and Aldh1a3) has been reported [3]. We examined the expression of RA synthesis enzymes ALDH1A1, ALDH1A2 and ALDH1A3 at three gestational stages (namely 8–9, 14–16 and 17–20 weeks gestation) which broadly reflect the timing of key events in the development of the human fetal gonad. At 8–9 weeks gestation, sex determination has occurred, yet male and female gonads contain only undifferentiated proliferating primordial germ cells. By 14–16 weeks, meiosis has initiated and syncitial germ cell nests are forming alongside ongoing germ cell proliferation in the fetal ovary. Finally, at 17–20 weeks, germ cell nests in the fetal ovary start to break down, primordial follicle formation commences and germ cells become arrested at the diplotene stage of meiotic prophase I. A progressive process of germ cell maturation also occurs in the fetal tests during this period, with a progressive loss of primordial germ cell-associated marker expression [16,19].

In the human fetal testis, ALDH1A1 expression increased significantly across the gestational range examined (p<0.05; Figure 1A). In contrast, there was a trend towards decreasing expression of ALDH1A1in the fetal ovary over the same period (Figure 1A) although this did not reach significance. We found no significant differences in expression of ALDH1A1 between gonads of different sexes at the same gestational age, or between fetal gonads and mesonephroi from 8–9 week fetuses. ALDH1A2 expression in the fetal testis was significantly higher at 8–9 weeks gestation than at 14–16 weeks (p<0.05) or 17–20 weeks gestation (p<0.01. Figure 1B). Furthermore, at 8–9 weeks gestation we detected a sexual dimorphism in ALDH1A2 expression, with transcript levels significantly higher in the testis than the ovary at this developmental stage (p<0.05). Notably, ALDH1A2 transcript levels were also higher in the 8–9 week human fetal testis than in mesonephroi from the same fetuses (p<0.01), indicating that in humans the gonad, rather than the mesonephros, may be the predominant site of RA synthesis. No significant differences in the expression of ALDH1A3 were detected either within or between sexes at any developmental stage (Figure 1C) although, as with ALDH1A1 and ALDH1A2, expression in both ovary and testis was not lower than in mesonephros. Together, these data support the hypothesis [18] that the human fetal gonad has an intrinsic capacity to produce retinoic acid.

**Expression and localisation of retinoid receptors in the human fetal gonads**

Retinoid signals are transduced by two families of receptors, RAR and RXR receptors, which can hetero- and homodimerise to regulate gene expression. Inhibition of RA receptor action has been shown to inhibit meiosis in RA-treated mouse testes [3,4] and in the human fetal ovary [18]. To identify the possible receptors involved in retinoid signalling in the human fetal gonad, we examined the expression of the RA receptors (RARα, β and γ) and the retinoid receptors (RXRα, β and γ) at the transcript and protein level in the developing human fetal ovary and testis.

We detected transcripts encoding all three RAR (Figure 2A–C) and RXR (Figure 2D–F) receptors in human fetal testes and ovaries. Interestingly however, we did not detect any significant changes in expression of any of the receptor isoforms either between gonads obtained from fetuses of the same sex at different gestational ages, or between those of different sexes at the same developmental stage (Figure 2A–F, n = 4–6 per group). Expression of the retinoid receptor machinery therefore appears not to be developmentally-regulated at the transcript level in human fetal gonads around the time of meiosis.

To establish the cellular targets of retinoid signalling in the human fetal tests and ovary, we performed immunohistochemistry using specific antibodies raised against RARα, RARβ and
RXRα on sections of second trimester human fetal ovaries and testes (Figure 3). RARα expression was widely distributed in the second trimester human fetal testis (Figure 3A). Expression was detected in germ cells, which displayed either nuclear or both nuclear and cytoplasmic staining. Sertoli cells were predominantly immunopositive, and displayed strong nuclear staining, although a subpopulation of these cells could be identified which did not express RARα. Peritubular myoid (PTM) cells were also mostly immunopositive, with the nucleus the predominant site of receptor localisation in this cell type. Interstitial cells were mostly immunopositive, although a subpopulation which showed no staining was also detectable. RARα expression in the human fetal ovary at a comparable developmental stage was present in germ cells in syncitial clusters (also known as germ cell nests; Figure 3B), and localised to both the cytoplasm and nuclei of these cells. Pregranolosa cells interspersed between germ cells also displayed strong nuclear staining for RARα. Expression of RARα in streams of mesenchymal cells between germ cell nests was variable, with some cells displaying nuclear staining and some being immunonegative.

Like RARα, the expression of RARβ in the second trimester testis was broadly distributed, with germ, Sertoli, PTM and interstitial cells all immunopositive (Figure 3C). In contrast to RARα however, the subcellular localisation of RARβ appeared to be predominantly nuclear in all cell types examined. In the fetal ovary, germ cells in syncitial clusters were again the predominant site of RARβ expression, with a mixture of large weakly-staining germ cells in which staining was cytoplasmic, and smaller germ cells with intensely immunopositive nuclei (Figure 3D). Pregranolosa cells displayed no staining, whilst RARβ staining was either weak or absent in the nuclei of mesenchymal somatic cells.

In the fetal testis, Sertoli and PTM cells displayed intense nuclear staining for RXRα (Figure 3E). Germ cells also expressed this receptor, but displayed weaker staining in both the cytoplasm and nucleus. A subpopulation of interstitial cells with immunonegative nuclei could also be discerned. In the fetal ovary, RXRα expression was similar to that of RARβ, with germ cells in nests displaying both nuclear and cytoplasmic staining (Figure 3F), whilst mesenchymal somatic cells and pregranolosa cells were immunonegative.

These data suggest that RA and its derivatives likely target a diverse range of cell types in the second trimester human fetal testis, whilst germ cells are the predominant target of retinoid action in the fetal ovary at a comparable developmental stage.

Expression of genes encoding STRA8, CYP26B1 and NANOS1-3 in the developing human fetal gonad

In the fetal mouse ovary, RA induces the expression of STRA8, which is required for pre-meiotic DNA replication and progression.
Germ cells in the fetal testis are shielded from the meiosis-inducing action of RA by Sertoli cell-expression of the RA metabolising enzyme Cyp26b1, and later by the expression of Nanos2 in germ cells which acts to ‘lock in’ the male fate. As conservation of the expression patterns of these genes in the human fetal gonad is likely to reflect functional conservation with their roles in mice, we compared the expression of STRA8, CYP26B1 and NANOS2 (and its paralogues NANOS1 and NANOS3) in the male and female human fetal gonad across the developmental window encompassing meiotic entry outlined above.

Expression of STRA8 was low/absent in the gonads of both sexes at 8–9 weeks gestation (Figure 4A), consistent with these tissues containing only pre-meiotic proliferating germ cells at this developmental stage. STRA8 expression increased significantly between 8–9 and 14–16 weeks in the human fetal ovary however (p<0.05), concomitant with the initiation of meiosis in this tissue. STRA8 expression remained very low in the testis at all gestations examined, and was significantly higher in the fetal ovary than testis at 14–16 weeks (p<0.0001) and 17–20 weeks (p<0.008) gestation. The developmental and sex-specific pattern of Stra8 expression therefore appears to be conserved between the fetal gonads of humans and mice at comparable developmental stages, as reported previously [18,20].

In contrast to STRA8, however, the expression of CYP26B1 in the human fetal gonad diverges significantly from that previously
reported in mouse. CYP26B1 expression was not significantly different between ovaries and testes at 8–9 weeks gestation, but was unexpectedly higher in 14–16 week ovaries than testis (p = 0.02, Figure 4B). To determine whether another member of the CYP26 family may be responsible for retinoid metabolism in the human fetal testis, we also examined the expression of CYP26A1 and CYP26C1, but were unable to detect transcripts for either gene at any gestational age examined in both fetal testis and ovary (data not shown). These data suggest that the ovary may have a greater capacity for RA metabolism than the fetal testis at this stage.

NANOS2 displayed both developmentally-regulated and sexually-dimorphic expression in the human fetal gonad (Figure 4C). NANOS2 expression was significantly higher in the fetal testis at 17–20 weeks gestation than at 8–9 weeks, consistent with the progressive commitment of germ cells to the male fate between these developmental stages. Whilst no difference in NANOS2 expression was detected between testes and ovaries at 8–9 weeks gestation, NANOS2 transcript levels were significantly higher in the fetal testis than ovary at 14–16 weeks (p = 0.01) and 17–20 weeks (p<0.01). The expression of NANOS2 in the human fetal testis therefore appears to mirror that reported for the homologous gene in mouse, indicating possible conservation of its functional role in reinforcing the male fate in testicular germ cells between these two species.

We also examined the expression of the related genes NANOS1 and NANOS3 in the developing fetal gonad. We found no differences in the expression of NANOS1 between testes and ovaries at any gestational age, nor between gonads of the same sex at different developmental stages (Figure 4D). Unexpectedly however, NANOS3 transcript levels were found to be significantly higher in ovaries than testes at 14–16 weeks (p<0.05) and at 17–20 weeks gestation (p = 0.02; Figure 4E), revealing a sexual dimorphism in expression levels in this gene that has not previously been reported in mice. Strikingly – and in stark contrast to the expression of Nanos3 in the fetal mouse ovary, which is downregulated shortly after the entry into meiosis from e14.5 onwards - the expression of NANOS3 increased significantly between 8–9 and 14–16 weeks gestation in the human fetal ovary (p<0.05), suggesting a possible role in meiosis for this protein.
Figure 4. Conserved and divergent patterns of expression of STRA8, CYP26B1 and NANOS1-3 in the human fetal gonad. qRT-PCR analysis of human fetal gonads reveals female-biased and developmentally-regulated expression of STRA8 (A). STRA8 expression increased significantly between 8–9 and 14–16 weeks in the human fetal ovary (a vs b, p < 0.05) consistent with the initiation of meiosis in the fetal ovary around 11 weeks gestation. Levels of transcripts encoding STRA8 were low and not significantly different between human fetal testes and ovaries at 8–9 weeks gestation, but were significantly higher in fetal ovaries than fetal testes at 14–16 weeks (c vs d, p < 0.0001) and 17–20 weeks (e vs f, p = 0.008). CYP26B1 (B) expression was not significantly different between samples of the same sex at different gestational ages, but was significantly higher at in the fetal ovary than the fetal testis at 14–16 weeks (a vs b, p = 0.02); suggesting the male-specific expression of CYP26B1 reported in mice at a comparable developmental stage is not conserved to humans. NANOS2 (C) expression was predominantly male-specific and developmentally-regulated, with expression increasing in the human fetal testis with increasing gestational age (a,b,c, p < 0.001). NANOS2 expression was also significantly higher in fetal testes than ovaries at 14–16 weeks (d vs e, p = 0.01) and 17–20 weeks (f vs g, p < 0.01), a result consistent with a role for this protein in repressing meiosis in the fetal male germline. 8–9, 14–16 and 17–20 denote the gestational age (in weeks) of specimens. No differences were detected in the expression of NANOS1 (D) between testis and ovaries at any gestational age, nor between gonads of the same sex at any developmental stage. NANOS3 expression (E) was significantly higher in the human fetal ovary at 14–16 weeks gestation than at 8–9 weeks gestation (a vs b, p < 0.05), in contrast to the downregulation of the homologous gene in the fetal mouse ovary at the comparable developmental stage. Expression of NANOS3 was also greater in the fetal ovary than in the fetal testis at 14–16 weeks (c vs d, p < 0.05) and 17–20 weeks (e vs f, p < 0.02). 8–9, 14–16 and 17–20 denote the gestational age (in weeks) of specimens, values denote mean ± s.e.m, 8–9, 14–16 and 17–20 denote the gestational age (in weeks) of specimens.

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Retinoic acid induces STRA8 expression in the second trimester human fetal testis, but does not affect the expression of other meiosis-associated genes

To determine whether the induction of STRA8 expression by RA is conserved between mouse and human, we investigated the effects of RA treatment on cells from the second trimester human fetal testis. To ensure germ cell exposure to RA, fetal testes (14–15 weeks gestational age, n = 6) were disaggregated to a single cell suspension (thus uncoupling germ cells from their associated somatic cells which are in the mouse thought to be the site of CYP26B1 expression) and cultured in serum free medium in the presence of either vehicle (DMSO) or 1 μM all-trans RA for 24 hours. Expression of STRA8, and of the meiosis markers SYCP3 and DMC1 was then assessed by qRT-PCR in control and RA-treated cultures. Treatment with RA for 24 hours resulted in a 2.2±0.3 fold increase in STRA8 expression relative to vehicle-treated controls (p<0.05, n = 5; Figure 5A); revealing conserved induction of STRA8 expression by RA in the human fetal testis. No significant differences were found between RA- and vehicle treated cultures in the expression of SYCP3 (94.4±5.7% of controls, n.s.; Figure 5B) or DMC1 (115.1±19.6% of controls, n.s.; Figure 5C) however, suggesting that whilst RA can selectively induce the expression of STRA8 in the human fetal testis, it may not be sufficient to induce additional meiosis-associated gene expression at this developmental stage, or in this experimental system.

Discussion

In the fetal mouse gonad, germ cell differentiation proceeds in a rostro-caudal wave, possibly reflecting the diffusion of mesonephros-derived retinoic acid along the long axis of the gonad [3,4,14,15]. Whilst meiosis is also thought to initiate at the cranial end of the fetal ovary in the human [21], within weeks a radial distribution of germ cells is detectable, with undifferentiated premeiotic PGC-like cells found at the periphery of the ovary and progressively more differentiated germ cells found towards the central cortex [16,17]. The existence of multiple subpopulations of germ cells at different developmental stages within the human fetal ovary indicates that differentiation is far less synchronous than in the mouse, and raises the question as to how this asynchrony is maintained. In this report we have examined the expression of key components of the retinoid synthesis and signalling apparatus in the human fetal gonad, and of key downstream effectors (STRA8) and antagonists (CYP26B1, NANOS2) of the meiosis-inducing action of RA. Whilst identifying conservation of some aspects of the regulation of meiotic entry between mice and humans, we have also identified significant species-specific differences in the expression of genes associated with the entry or inhibition of meiosis, which may contribute to or help explain the differing spatiotemporal organization of germ cell differentiation in the fetal ovary. Our findings also support the recent hypothesis that intrinsic RA synthesis within the ovary, rather than RA originating from the mesonephros, may be the primary driver of meiotic initiation in the human fetal ovary [18].

The genes encoding the retinoid synthesis enzymes Aldh1a2 and Aldh1a3 are expressed in the mesonephroi of the fetal mouse around the time of the initiation of meiosis, but their expression is undetectable within the gonad itself [3]. In this report however, we demonstrate the expression levels of all three ALDH1A genes in the gonads of both sexes to be at least equal to those detected in mesonephroi from fetuses at 8–9 weeks gestation, suggesting that unlike that of the mouse, the human fetal gonad has an intrinsic capacity to synthesis RA. We found the expression of ALDH1A2 to be sexually-dimorphic, being significantly greater in the fetal testis.
at 8–9 weeks gestation than in ovaries from age matched samples or compared to testis from fetuses at later gestational ages. We found no significant differences in the level of expression of Aldh1a1 between the ovary and testis however, in contrast to the male-specific expression of Aldh1a1 reported in the fetal mouse and chicken at a comparable developmental stage [22]. The male-biased expression of Aldh1a1 in the human fetal testis may replace the function of Aldh1a1 in these species. Our data also both compliment and contrast with that reported recently by Le Bouffant et al., who found the expression of Aldh1a1 to increase sharply in the ovary around 11 weeks post-ovulation (wpo), coincident with the onset of meiosis (yet was lower in all ovarian samples examined than the 9 wpo tests), whilst Aldh1a2 expression was relatively stable across the developmental range examined [18]. A direct comparison of our data with that of Le Bouffant et al. [18] is not straightforward, given the different systems of aging fetal specimens, the range of gestational ages examined and the number of independent biological replicates (fetal specimens) used for each time point examined. Despite these specific differences however, the data reported here and by Le Bouffant et al. [18] support the same conclusion; that unlike the mouse, the fetal human gonad has an intrinsic capacity to produce RA, and thus gonadal, rather than mesonephric RA synthesis may drive the initiation of meiosis in germ cells in the human fetal ovary. Furthermore, local production rather than diffusion of RA from the mesonephros may help explain the asynchronous entry of germ cells into meiosis in the human fetal gonad.

We have examined for the first time the expression of all of the genes encoding the RAR and RXR retinoid receptor proteins during human fetal gonadal development, and demonstrated protein expression of members of both receptor families (RARα, RARβ and RXRγ) in the germ cells of the fetal human ovary, and in a wide range of cell types in the human fetal testis. Our finding that germ cells in the human fetal ovary are transducing RA signals (as indicated by nuclear localization of the receptors) contrasts with that reported by Morita and Tilly in the fetal mouse ovary at a comparable developmental stage (e13.5), who (using pan-RAR and pan-RXR antibodies) reported only weak cytoplasmic expression of RAR proteins, and – in contrast to the data presented here - no expression of the RXR proteins [23]. In the same study, treatment of e13.5 fetal mouse ovaries with a relatively low concentration of RA (0.01 μM) resulted in the relocalisation of RAR proteins to germ cell nuclei, leading the authors to conclude that RA signalling must either be extremely low or absent in the fetal ovary in vivo at this time [23]. The absence of nuclear-localised RA receptors in the fetal mouse ovary around the time of the initiation of meiosis seemingly contradicts the model in which RA signalling in ovarian germ cells around e13.5 stimulates Stra8 transcription and subsequent meiotic entry, but suggests perhaps that only very low levels of RA (i.e insufficient to cause widespread receptor nuclear localization) are required to initiate meiosis in the fetal mouse ovary. The predominantly nuclear localization of RARs in human ovarian fetal germ cells may indicate the existence of higher local RA concentrations than are present in the mouse fetal ovary, perhaps arising from intrinsic ovarian, rather than mesonephric RA synthesis. Alternatively, it may reflect a broader role for RA signalling in germ cell development in the human beyond the regulation of meiotic entry, such as in the regulation of germ cell survival or proliferation [18].

The most significant aspect of the immunohistochemical analyses reported here however, is the identification of germ cells displaying nuclear staining for RARα, RARβ and RXRα in the human fetal testes. This strongly suggesting that they are both receiving and transducing retinoid signals and are therefore not effectively shielded by the action of the somatic cell-expressed metabolizing enzyme CYP26B1 as is believed to be the case in the fetal mouse testis; [3,4]. The widespread localization of the RA receptors in the human fetal testis also contrasts with reports that RARα and RARβ are undetectable in the developing rodent testis prior to e16 [24], providing further evidence of extensive divergence in testicular RA signalling between these species.

These data, coupled with greater expression of Cyp26b1 in the human fetal ovary than testis (this paper and [18]; again in contrast to the male-specific expression of Cyp26b1 in the fetal mouse testes), suggests a less important role for CYP26B1 in the regulation of meiosis in the developing gonad in the human than in the mouse. Although the inappropriate entry of testicular germ cells into meiosis in mice homozygous for targeted disruptions of Cyp26b1 [25,26], and in vitro cultures of fetal testes in which Cyp26b1 is inhibited with ketoconazole [3,4] provide compelling evidence of a role for Cyp26b1 in inhibiting the initiation of meiosis in testicular germ cells, aspects of this model have recently come under increasing scrutiny. The ketoconazole culture experiments have not been replicated by other groups [27], and ketoconazole is unable to ameliorate the inhibitory effects on ovarian germ cell meiotic entry of a testis co-cultured with an ovary [28], as would be expected if Cyp26b1 metabolism of RA was the key inhibitor of meiotic initiation. Together, these data argue that other mechanisms that inhibit the entry of germ cells into meiosis in the fetal testis are likely to exist. The recent identification of FGF9 as an inhibitor of RA-induced meiosis in the fetal mouse testis [29], and the demonstration that secretory pathways and their cargoes play a key role in determining germ cell sex determination [27,28] provide further evidence for the existence of additional mechanisms that inhibit meiotic entry in the fetal testis, or conversely promote it in the ovary.

Some aspects of meiosis initiation and germ cell sex determination do appear to be conserved between mouse and human, however. We find Stra8 expression to increase between 8–9 and 14–16 weeks gestation, consistent with previous reports of the expression of this gene increasing around the onset of meiosis in the human fetal ovary [18,20] and reflecting a comparable increase in Stra8 expression in the fetal mouse ovary from e13.5 onwards. We have also conducted the first comparative analysis of gene expression of the Nanos family in the developing human fetal ovary and testis. We find the expression of Nanos2 to be restricted to the human fetal testis, consistent with the male-specific expression and meiosis-inhibiting action of Nanos2 in the germ cells of the mouse testis [6,30]. In contrast, we detected the opposite pattern of expression for Nanos3, which we find to be expressed at higher levels in fetal ovaries than in testes. Nanos3 is expressed exclusively in pre-meiotic germ cells in the mouse and is downregulated shortly after the onset of meiosis [30], yet in the human fetal ovary we find the expression of Nanos3 to increase significantly between 8–9 and 14–16 weeks gestation, concomitant with the onset of meiosis. This raises the intriguing possibility that Nanos3 may be involved in the regulation or progression of meiosis in human fetal ovarian germ cells; a finding that warrants further investigation. Our finding that Nanos1 expression is maintained at a broadly constant level across the gestational range examined here is also unexpected, as Nanos1 protein is detectable only in the germ cells of the second trimester testis onwards [19]. The absence of Nanos1 transcripts in the first trimester human fetal gonad suggests therefore that Nanos1 mRNA may be subject to post-transcriptional regulation in human fetal germ cells.

We have demonstrated for the first time that RA can induce expression of Stra8 in the human fetal testis. Utilizing a serum-
free culture system used previously to investigate the effects of growth factor signalling in the human fetal ovary [31], we demonstrated increased STRA8 expression in response to RA in cultures of human fetal testes at 14–15 weeks gestation – disaggregated to ensure germ cells were uncoupled from the local RA-metabolising action of neighbouring CYP26B1-expressing somatic cells. We failed to detect any changes in the levels of transcripts encoding the meiosis-specific proteins SYCP3 and DMC1 however, indicating that whilst RA is sufficient to induce STRA8 expression, it cannot trigger widespread meiosis-associated gene expression in second trimester testicular germ cells. This may in part reflect our use of early second trimester tissues in this experiment, as expression of NANOS2, which acts to inhibit meiotic entry in testicular germ cells in a cell-intrinsic fashion [6], is at its peak at this time, or the relatively short (24 hours) period of culture used. Further experiments will need to be performed before conclusions as to whether RA can induce meiosis in the human fetal testis can be drawn.

In summary, we have characterized the expression of the essential components of RA signalling in the human fetal ovary and testis that may underpin the initiation of meiosis in the fetal ovary and its inhibition in the fetal testis. However, we identified key differences between humans and mice in the expression and distribution of components of the retinoic acid synthesis, signalling and effector machinery required for RA-regulation of sex-specific entry into meiosis. Whilst a key role for RA in the regulation of the initiation of meiosis in the human fetal ovary now appears indisputable [18], many aspects of this – particularly with respect to how this relates to the spatiotemporal organization of germ cell differentiation in the human fetal ovary and the apparent asynchronous entry of human fetal ovarian germ cells into meiosis – remain to be clarified. Detailed morphometric studies to establish the location of the first meiotic cells within the fetal ovary (aided by the development of antibodies to the human STRA8 and NANOS3 proteins) and the determination of the sites of retinoic acid synthesis and metabolism within the human fetal ovary and testis will be required to resolve these questions.

Materials and Methods

Ethics statement

Ethical approval for this study was obtained from Lothian Research Ethics Committee (study code LREC 08/S1101/1). All participants gave informed written consent in accordance with national guidelines [32].

Collection of human fetal tissues

Human fetal testes and ovaries were obtained following elective termination of pregnancy during the first (50–65 days gestation) and second (13–19 weeks gestation) trimesters, as dated from last menstrual period. No terminations were for reasons of fetal abnormality and all fetuses appeared morphologically normal. Termination was induced with mifepristone (200 mg, orally) in the same sample. Primer sequences can be found in Table S1. The expression level of each gene of interest was normalized to that of the 18S ribosomal RNA within the same sample. To determine the expression of STRA8, SYCP3 and DMC1 in cultured human fetal testis, first strand cDNA was prepared using the Superscript VILO mastermix kit (Invitrogen), and assessed by qRT-PCR using SYBR green technology on an ABI7900HT thermal cycler (Applied Biosystems) as described previously [35]. Primer sequences along with corresponding probe numbers are listed in Table S1. The expression level of each gene of interest was normalized to that of the housekeeping gene RPL32 within the same sample. Primer sequences can be found in Table S2.

Immunohistochemistry

Immunohistochemistry was performed on fixed sections of fetal ovary and testis tissue essentially as described previously [36]. Briefly, 5 μm thick sections of Bouin’s-fixed, paraffin embedded tissues were mounted on electrostatically charged glass slides (BDH Chemicals, Poole, UK), dewaxed and rehydrated using xylene and graded alcohols and antigen retrieval performed by pressure cooking in 0.01 M sodium citrate buffer (pH 6) for five minutes. Endogenous peroxidase activity was blocked using 3% (w/v) hydrogen peroxide (H2O2) in methanol for 30 minutes, and slides were blocked using the avidin/biotin blocking kit (Vector Laboratories, Inc., Peterborough, UK) and incubation in Tris Buffered Saline (TBS), supplemented with 5% BSA and 20% normal serum (NS). Antibodies (listed in Table S3) were diluted in 5% BSA/TBS and applied to the sections at 4°C overnight. Antibodies were detected using the appropriate biotinylated secondary antibodies (30 minutes, 1:500 dilution in BSA/TBS/NS) and incubation with avidin-biotin-HRP complex (Vector...
Laboratories. Bound antibodies were visualized using 3,3-diaminobenzidine tetrahydrochloride (DAB; DAKO, Cambridge, UK). Negative controls, incubated with blocking serum instead of primary antibody, were included in each experiment and displayed no staining (data not shown).

Statistical analyses

Data presented represent mean ± standard error of the mean (SEM) of at least four independent biological replicates. Data were analysed using either ANOVA or Student’s t test using Graphpad Prism Software. P values of less than 0.05 were considered statistically significant.

Supporting Information

Table S1 Oligonucleotide sequences and corresponding Roche Universal Probe Library numbers used in qRT-PCR assessment of gene expression in frozen human fetal tissues.

(DoC)

References


Table S2 Oligonucleotide primer sequences used in SYBR green qRT-PCR analysis of gene expression in cultured human fetal testes.

(DoC)

Table S3 Antibodies used for immunohistochemistry.

(DoC)

Acknowledgments

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Author Contributions

Conceived and designed the experiments: AJC GC RAATKPS. Performed the experiments: AJC GC HLK. Analyzed the data: AJC GC RAATKPS. Wrote the paper: AJC GC RAATKPS.