Is there a role for DAZL in human female fertility?

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ABSTRACT: The RNA binding protein deleted in azoospermia-like (Dazl) is a key determinant of germ cell maturation and entry into meiosis in rodents and other animal species. Although the complex phenotype of Dazl deficiency in both sexes, with defects at multiple stages of germ cell development and during meiosis, demonstrates its obligate significance in fertility in animal models, its involvement in human fertility is less clear. As an RNA binding protein, identification of the in vivo mRNA targets of DAZL is necessary to understand its influence. Thus far, only a small number of Dazl targets have been identified, which typically have pivotal roles in germ cell development and meiotic progression. However, it is likely that there are a number of additional germ cell and meiosis-relevant transcripts whose translation is affected in the absence of Dazl. Efforts to identify these RNA targets have mainly been focused on spermatogenesis, and restricted to mice. In women, prophase I occurs in fetal life and it is during this period that the ovarian follicle pool is established, thus factors that have a role in determining the quality and quantity of the ovarian reserve may have significant impact on reproductive outcomes later in adult life. Here, we suggest that DAZL may be one such factor, and there is a need for greater understanding of the role of DAZL in human oogenesis and its contribution to lifelong female fertility.

Key words: DAZL / meiosis / female fertility / oocyte quality / DAZL RNA targets

Introduction

Female reproductive potential is governed by two key factors: oocyte quality and quantity, with the loss of quality underpinning the exponential increase in age-related aneuploidy, and the loss of quantity underlying the age at menopause (Broekmans et al., 2007; Schmidt et al., 2012; Nelson et al., 2013). These factors are increasingly important for women where societal and demographic changes have influenced the age at which they wish to have their families. Indeed, the maternal age at first birth across the Western world has steadily risen, with the number of first-time mothers over the age of 35 years being an all-time high (Schmidt et al., 2012). While an age-related decline in fertility is much less marked in men, defects in spermatogenesis are present in one-third to half of couples who experience infertility (Matzuk and Lamb, 2008; Miyamoto et al., 2015). Thus, reduced gamete quality and quantity is a key issue in both sexes, and although in some cases, techniques such as ICSI and array comparative genomic hybridization to identify euploid embryos can provide ways of circumventing the problem, they do not address the underlying issues.

The establishment of the ovarian reserve occurs during fetal life and begins with the migration of primordial germ cells into the gonadal ridge during early gestation (reviewed in Findlay et al., 2015), although post-natal oogenesis is also debated (Grieve et al., 2015). Here the germ cells undergo mitotic divisions with incomplete cytokinesis, producing a surplus of interconnected oogonia within germ cell ‘nests’ (Gondos et al., 1971; Pepling and Spradling, 1998). Ceasing proliferation, the oogonia enlarge and develop into primary oocytes, which commence meiosis I and progress through leptotene, zygotene and pachytene before arresting at diplotene of prophase I. The ‘nest’ structure undergoes breakdown, in which most oocytes are lost through apoptosis, while the remainder are enclosed by a layer of pregranulosa cells, thus forming primordial follicles from 17 weeks gestation. These oocytes remain quiescent in a state of meiotic arrest until, should they escape the atresia that befalls >99% of all follicles, they are reactivated at the time of ovulation: this will not be months or years, but decades later. Greater understanding of how oocytes maintain long-term meiotic arrest with chromosomal fidelity is critical to our knowledge of lifelong reproductive potential in women, and may provide an opportunity for both diagnosis and therapy.

DAZL (deleted in azoospermia-like)

DAZL and homologue members DAZ and BOLL (previously known as BOULE) belong to the DAZ family of genes encoding RNA binding proteins, and have essential roles in gametogenesis. It is likely that BOLL is the ancestral gene, first identified in Drosophila (where it is known as bol), and DAZL is considered to have arisen from BOLL through a gene duplication event (Castrillon et al., 1993). While BOLL appears to be strongly...
evolutionarily conserved, DAZL is only found in vertebrates and DAZ is limited to just humans and old world monkeys (Reijo et al., 1995). DAZL is a single-copy autosomal gene and encodes a protein that is specifically expressed in germ cells at all stages of oogenesis. In addition, DAZL expression has been reported in human and mouse granulosa cells, human theca interna cells and in granulosa-luteal cells of the human corpus luteum; however, these remain controversial (Reijo et al., 1997; Dorfman et al., 1999; Nishi et al., 1999; Pan et al., 2002).

In mouse fetal gonads, Dazl transcripts are first detected from embryonic day (e) 11.5 in post-migratory germ cells (Fig. 1), and this activation of Dazl appears to be a consequence of the genome-wide loss of DNA methylation that occurs in germ cells at this stage (Hackett et al., 2012). Dazl expression dramatically increases at e13.5 in females, coinciding with the onset of meiosis (Seligman and Page, 1998), and by e17.5 transcript expression in the ovary steadily decreases as the number of oocytes is depleted. In the human fetal ovary, a marked increase in DAZL mRNA and protein is observed between 9 and 14 weeks gestation (Ruggiu and Cooke, 2000; Anderson et al., 2007; He et al., 2013), that immediately precedes and is coincident with the onset of meiosis. At this time, DAZL undergoes a dynamic redistribution from the germ cell nucleus (where it may have a role in RNA processing and storage) to the germ cell cytoplasm where it is expected to function (Reynolds and Cooke, 2005; Anderson et al., 2007). Expression of DAZL is also found in oocytes of newly formed primordial follicles (He et al., 2013) and in mouse, Dazl expression has been demonstrated to persist through later stages of oocyte maturation, including through to zygote formation, where Dazl-dependent translation is necessary for spindle assembly, the metaphase I–II transition and early embryo development (Chen et al., 2011).

The phenotype of Dazl deficiency has been investigated in detail in the mouse. The deletion of Dazl results in a loss of germ cells in both male and female gonads, with evidence of increased apoptosis, reduced expression of germ cell markers and aberrant chromatin structure (Ruggiu et al., 1997; Schrans-Stassen et al., 2001; Lin and Page, 2005). The loss of Dazl leads to a reduction in the number of post-migratory primordial germ cells, defects in sexual differentiation of the germ cells, a failure to erase and re-establish genomic imprints, and defects in meiotic progression (Haston et al., 2009; Gill et al., 2011). The defect in meiosis has been shown by immunohistological analysis to be a failure to progress beyond leptotene of meiotic prophase I. Although axial elements are formed [visualized using an antibody against synaptonemal complex protein 3 (Sycp3)], complete synaptonemal complexes are not, giving a precise point beyond which meiosis cannot progress without Dazl (Saunders et al., 2003). Transplantation of ROSA/LacZ-labelled spermatogonia into the testes of Dazl−/− mice confirms that the somatic tissues are capable of supporting gamete development, indicating that it is only germ cell function that is compromised in these animals (Rilianawati et al., 2003).

Many RNA binding proteins are recognized as having roles in translational control during gametogenesis. During oogenesis, growing oocytes accumulate significant quantities of mRNAs, but transcription ceases during the final stages of maturation and subsequent protein production is dependent upon a well-orchestrated programme of dynamic modulation of mRNA poly(A) tail length, recruitment to polysomes and translational activation (Seydoux and Braun, 2006; Radford et al., 2008). Indeed, DAZL is associated with actively translating polysomes (Tsui et al., 2000b; Maegawa et al., 2002) and sucrose gradient analysis of translation intermediates from a 3′ untranslated region (UTR) tethering reporter...
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assay in Xenopus laevis oocytes revealed that mDazl acts to specifically stimulate translation through regulation of the initiation stage (Collier et al., 2005). DAZL contains two functional domains: a highly conserved RNA recognition motif (RRM) and a single DAZ domain unique to DAZ family proteins. Investigation into the binding specificity of Dazl using RNA homopolymers demonstrated that the RRM of Xenopus Dazl preferentially binds poly(U) and poly(G) RNA in vitro (Houston et al., 1998), and this was also the case for human DAZL (Tsui et al., 2000b). An improvement to the description of this binding element was achieved using SELEX, a powerful RNA ligand enrichment-based strategy, in addition to a three-hybrid screen, to show that mDazl binds oligo(U) stretches interspersed by G or C nucleotides (Venables et al., 2001). When the G or C nucleotides were replaced by A, only a weak interaction was observed between Dazl and the binding site, and replacement with a U abolished this interaction completely. Resolution of the RRM crystal structure of mDazl with and without 3′ UTR target sequences has revealed the involvement of an extended, kinked pair of β-strands in the recognition of a GUU triplet in the 3′ UTR of Dazl targets (Jenkins et al., 2011). The nucleotide at position 4 of this refined Dazl consensus sequence varies between complexes (U or C), and mutation of bases within the GUU triplet reduces the affinity of binding. Searching for DAZL targets based on this binding site alone is likely to return many false positives given that such short sequences may occur relatively frequently in many mRNAs. Therefore, the presence of multiple DAZL-binding elements, which can enhance translational stimulation (Collier et al., 2005; Fox et al., 2005), partner proteins such as DAZAP1 or PUM2 (Tsui et al., 2000a; Moore et al., 2003), which may modulate DAZL stimulation, and appropriately timed germ cell expression should all be taken into consideration when attempting to uncover DAZL targets.

**DAZL in fertility: action through its mRNA targets**

Efforts to identify in vivo mRNA targets of mammalian DAZL have mainly been focused on mouse. The results of these inquiries further attest to Dazl’s role in fertility as many identified targets are key players in germ cell development and gametogenesis, particularly during meiosis. DAZL could potentially have distinct mRNA targets at different stages of germ cell development. The first screen for Dazl targets, carried out as part of an investigation addressing the absence of DAZL proteins and spermatogenic defects, identified Tpx-1, a testicular cell adhesion protein (Jiao et al., 2002). The significance of Tpx-1 breakpoints in male fertility, together with its temporal expression pattern, suggests this gene is essential for the progression of spermatogenesis, and Dazl has a key role in regulating it. Tpx-1 is not expressed in the female, but a study in mouse oocytes has identified Tpx-2 as a Dazl target (Chen et al., 2011). TPX2 activates a regulator of centrosome and spindle pole assembly, and itself participates in spindle pole organization (Gruss et al., 2001; Bayliss et al., 2003). Morpholino down-regulation of Tpx-2 in oocytes causes spindle assembly disruption and chromosome condensation failure; phenotypic changes similar to those observed post-Dazl knockdown in oocytes (Brunet et al., 2008), suggesting Tpx2 protein is responsible for these defects and may be a key target for Dazl in post-natal mouse oocytes. These studies used in vivo matured germinal vesicle, metaphase I and metaphase II mouse oocytes and also confirm Tex19.1, which has a role in both male and female fertility in mice (Ollinger et al., 2008), as a bona fide Dazl target.

Tex19.1 had previously been reported as a potential Dazl target during spermatogenesis along with Sycp3 and Vasa (formally known as DDX4, or Mvh in mouse) (Reynolds et al., 2005). These genes were identified as potential targets for translational regulation by Dazl using a combination of immunoprecipitation and microarray in male mouse germ cells from wildtype and Dazl−/− animals, making these studies the first attempt to isolate endogenous Dazl–RNA complexes rather than exploiting interactions with recombinant proteins or mRNA libraries (Reynolds et al., 2005, 2007). Sequence analysis showed that Sycp3 and Vasa transcripts contain Dazl binding elements, and RNA binding and luciferase assays were used to demonstrate that Dazl stimulates transcript translation via the 3′ UTR; in the case of Sycp3, this response was diminished upon mutation of the Dazl binding sites. VASA is an ATP-dependent RNA helicase of the DEAD-box family (Raz, 2000), and is widely utilized as a germ cell marker, while SYCP3 is a major component of the lateral elements of the synaptonemal complex in meiotic germ cells (Schrann et al., 2011). Defects in translation of Vasa, Sycp3 and Tex19.1 could all contribute to the defects in meiotic progression reported in Dazl−/− spermatocytes (Saunders et al., 2003), as Mvh−/−, Sycp3−/− and Tex19.1−/− spermatocytes all have defects in progression from leptotene through to pachytene stages of meiosis I (Tanaka et al., 2000; Ollinger et al., 2008). However, some aspects of the male Dazl−/− phenotype, such as the defects in spermatogonial differentiation (Schrann-Stassen et al., 2001), the reduction in post-migratory germ cell number (Haston et al., 2009), extensive germ cell apoptosis during male fetal development (Lin and Page, 2005) and the failure of fetal oocytes to progress through meiotic prophase (Ruggiu et al., 1997; Saunders et al., 2003), are not reported in any of the Mvh−/−, Sycp3−/− or Tex19.1−/− mice, suggesting that additional as yet unidentified Dazl targets are likely contributing to the complex Dazl−/− phenotype.

DAZL may also impact fertility through its mRNA targets with known functions in the regulation of pluripotency, differentiation and apoptosis. Through an association with Tex1, Dazl has an essential role in mediating the hydroxylation of methylated cytosine residues in mouse embryonic stem cells that are actively reprogramming to a pluripotent ground state (Welling et al., 2015). In primordial germ cell-like cells derived from mice embryonic stem cells, DAZL was demonstrated to block the translation of core pluripotency factors Sox2 and Sal4, which are essential for stem cell self-renewal, as well as of Suz12, which is required for the differentiation of pluripotent cells (Chen et al., 2014). In addition, Dazl inhibited the translation of proapoptotic caspases, which is consistent with the Dazl−/− phenotype, in which embryos have a drastic reduction in germ cell number due to increased apoptosis (Lin and Page, 2005; Chen et al., 2014). Although it is mostly recognized to enhance translation, this is one of few instances where a repressive function has been reported for Dazl. It is possible that the observed translational repression by Dazl in these investigations is an indirect effect, as some of Dazl’s partner proteins, for example, Pumilio-2 (Moore et al., 2003; Fox et al., 2005), are known to bind RNA and function as translational repressors. Nevertheless, it appears that DAZL has a more complex role in regulating its downstream target transcripts than was previously believed, although these data also highlight the need for an understanding of which effects are through direct binding of the mRNA target and which are indirect.
Is DAZL relevant to human female fertility?

In humans, direct demonstration of a causative role is not possible, but observation of the association of DAZL with reproductive outcomes provides some evidence for the involvement of this gene in human fertility. However, given the protracted duration of oogenesis and the inaccessibility of the female gamete, the majority of findings pertaining to DAZL function have been restricted to the testis in both experimental animals and humans. In males, it is estimated that 10–15% of cases of azoospermia and oligozoospermia result from deletions within the Y chromosome, specifically the region which contains the DAZ gene cluster (Reijo et al., 1995), but DAZL is considered to be a prime candidate implicated in roughly 60% of idiopathic male infertility, given its high similarity to DAZ, its germ-cell specific expression and the suspected involvement of autosomal recessive mutations in these men (Lilford et al., 1994; Visser and Repping, 2010). Altered methylation of the DAZL promoter has been associated with male infertility (Navarro-Costa et al., 2010; Li et al., 2013). Furthermore, polymorphisms in both the promoter and coding sequence of DAZL have been correlated with total sperm count, sperm motility, age at menopause and primary ovarian insufficiency (POI), also known as premature ovarian failure, in fertile men and women, respectively (Teng et al., 2002; Tung et al., 2006a), although other studies have been unable to find such associations (Bartoloni et al., 2004; Zerbetto et al., 2008). Interestingly, many of the allelic variants studied exhibited opposing effects in men and women, which may be indicative of the different genetic requirements of male and female germ cells (Tung et al., 2006b). This sex-dependent effect was also observed in knockout mice, where the Dazl-/- phenotype could be partly rescued by a human DAZL transgene in male mice, but not in females (Vogel et al., 2002). Ultimately, these observations suggest that hypomorphic polymorphisms and single-nucleotide polymorphisms that quantitatively reduce DAZL expression or function in human germ cells could potentially have consequences for fertility.

Thus far, the study by Tung et al. (2006a, b) has been the only report of DAZL’s relevance to human female fertility and while Tung et al. suggest that DAZL might affect oocyte quantity in women (through influencing age at POI or menopause), there has been no examination as to whether the quality of oocytes is affected too. Given that DAZL was shown to influence total sperm count and sperm motility, quantity and quality parameters respectively, it seems possible that the same should occur in women; a key significant difference being the developmental period at which the effect happens. The generation of sperm via spermatogenesis, and thus meiosis, is a continuous process beginning at the onset of puberty and continuing throughout male adult life. However, in females, the process of oogenesis is initiated in the fetus, with the formation of stored oocytes that are used periodically over a defined reproductive lifespan. It is not clear if the genetic associations between DAZL polymorphisms and age at menopause and POI reflect a role for DAZL in fetal oocyte development or maintenance of oocytes during their prolonged post-natal arrest, but these associations are consistent with the fetal oocyte death reported in Dazl null mice (Ruggiu et al., 1997; Sanders et al., 2003), and provide a link between DAZL function during fetal/early post-natal life and reproductive success later in adult life.

The increase in the incidence of infertility and miscarriage from the mid-fourth decade of female life onwards (Schmidt et al., 2012) is in great part attributed to segregation errors in meiosis I in the female (Hassold and Hunt, 2001). This is likely because oocytes that are ovulated in late reproductive life differ from their earlier counterparts in the duration of prophase I arrest. The requirement for DAZL during meiotic progression in fetal mouse oocytes suggests that less severe hypomorphic mutations and/or polymorphisms in DAZL in humans could perturb meiotic progression sufficiently to impair oocyte quality. Accurate chromosome segregation in adult oocytes depends on successful establishment of meiotic crossovers between paired and synapsed homologous chromosomes during fetal development, and the subsequent maintenance of those crossovers post-natally (Herbert et al., 2015; MacLennan et al., 2015). Defects in meiotic progression during fetal oocyte development, such as those reported in female mice carrying mutations in the Dazl-target Syccp3, can result in aneuploidy arising during the meiotic chromosome segregations and reduced oocyte quality in adulthood, although this is not age-related (Yuan et al., 2002). However, analysis of chromosomies in oocytes of naturally aged mice demonstrated that premature bivalent separation into univalents is the primary defect responsible for age-related aneuploidy (Sakakibara et al., 2015). This was also found in human oocytes, where there was an increase in sister kinetochore separation, rotated bivalents and merotelic attachments, suggesting in fact there are multiple age-related changes in chromosome architecture with advanced maternal age (Zielinska et al., 2015). Therefore, the potential role of DAZL and DAZL-targets, in meiotic progression in human fetal oocytes may be able to impact on human female fertility in adulthood through influencing oocyte quality. Experiments in mice suggest that Dazl also has functions in fully grown oocytes in adult life where it is essential for the resumption of meiosis I in germinal vesicle oocytes and the transition to meiosis II, as in its absence, meiotic microtubules become disorganized causing meiotic spindle failure; these oocytes cannot be fertilized (Chen et al., 2011). Consequently, factors acting during oogenesis at the time of meiosis initiation, arrest and completion can potentially have fundamental roles in determining oocyte quality as well as quantity, thereby influencing both components of reproductive lifespan: a women’s ability to conceive later in life as well as her age at menopause. We suggest that DAZL may be one such factor, and for this reason, highlight the need for specifically female-oriented studies to investigate this notion further.

Experimental strategies to investigate the role of DAZL in women

The identification of DAZL mRNA targets illustrates the growing spectrum of processes in gametogenesis that this gene is proposed to regulate. However, the only work in humans to date, where overexpression of DAZL elevated endogenous VASA and SYCP3 levels (Kee et al., 2009), has been in embryonic stem cells rather than germ cells. The study of DAZL in women is complicated by scarce access to female gametes. However, it is possible to obtain fetal ovarian tissue after elective termination of pregnancy, while mature adult oocytes may be accessed in the course of IVF or ICSI cycles, and advances in follicle culture (Xiao et al., 2015) may allow access across a range of developmental stages. Further understanding of the role of DAZL in oogenesis will require knowledge of its target mRNAs at different stages throughout development. The most ideal way to achieve this would be to adopt an RNA immunoprecipitation approach similar to that of Reynolds and Cooke.
(2005) in order to isolate RNAs bound to endogenous DAZL. These RNAs can then be identified using microarray or high-throughput RNA sequencing; advances in these technologies now make this possible with limited amounts of starting material (Kanter and Kalisky, 2015). However, limitations include the sensitivity of the assay, as well as the strength of the DAZL–RNA interaction. Cell lysis can also lead to associations between molecules that are not representative of interactions that occur in vivo, and for this reason, newly identified targets should be investigated in vitro using RNA binding assays and polysome profiling to confirm DAZL’s ability to regulate target RNA translation. Lastly, gene manipulation of DAZL using RNA interference in cultured human fetal ovaries or mature oocytes provides a powerful strategy to explore the functional significance of novel RNA targets during meiosis, fertilization and early embryo formation. This approach has been used to study the roles of growth differentiation factor 9 and proliferating cell nuclear antigen in newborn hamster ovaries (Wang and Roy, 2011), and of DMRT-Like Family A2 (DMRTA2) in the human (Poulin et al., 2014). Alternatively, silencing of DAZL or RNA target gene expression in mature oocytes can be achieved through the injection of specific antisense morpholino oligonucleotides (Chen et al., 2011).

Concluding remarks

The RNA binding protein DAZL has an essential role in translational control during gametogenesis, and its absence or altered function is associated with infertility in several organisms, including humans. Although many targets of DAZL have been investigated in vitro and in vivo in a number of different species, many, or indeed probably most of its mRNA targets have yet to be identified, especially in women. As a key factor in the initiation of meiosis, DAZL target mRNAs involved with this are of particular significance, given the critical issues in meiosis specific to female gamete formation. Therefore, identification of mRNAs that depend on DAZL for regulation of their translation will increase our understanding of the fundamental processes that underpin oocyte quality and quantity, which in turn determine a woman’s reproductive potential and reproductive lifespan.

Authors’ roles

R.R. wrote the manuscript. I.R.A. and R.A.A. edited the manuscript.

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Conflict of interest

None declared.

References

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references

anderson ra, fultz n, cowan g, counts s, saunders pt. conserved and divergent patterns of expression of dazl, vas a and oct4 in the germ cells of the human fetal ovary and testis. bmc dev Biol 2007; 7: 136.
bartoloni l, cazzadore c, ferlin a, garolla a, foresta c. lack of the t54a polymorphism of the dazl gene in infertile italian patients. mol hum reproduc tion 2004; 10: 613–615.
bayliss r, sardon t, vernos i, conti e. structural basis of aurora-a activation by tpx2 at the mitotic spindle. mol cell 2003; 12: 851–862.
brunet s, dumont j, lee kw, kinoshita k, hikal p, gruss oj, maro b, verlhac mh. meiotic regulation of tpx2 protein levels governs cell cycle progression in mouse oocytes. plos one 2008; 3: e3338.
castillon dh, gonzcy p, alexander s, rawson r, eberhart cg, viswanathan s, dinardo s, wasserman sa. toward a molecular genetic analysis of spermatogenesis in drosophila melanogaster: characterization of male-sterile mutants generated by single p element mutagenesis. genetics 1993; 135: 489– 505.
chen j, melcon c, suh n, oh js, horner k, xie f, sette c, blieloch r, conti m. genome-wide analysis of translation reveals a critical role for deleted in azoospermia-like (dazl) at the oocyte-to-zygote transition. genes dev 2011; 25: 755–766.
chen hh, welling m, bloch db, munoz j, mientjes e, chen x, tramp c, wu j, yabuuchi a, chou yf et al. dazl limits pluripotency, differentiation, and apoptosis in developing primordial germ cells. stem cell rep 2014; 3: 892–904.
collier b, gorgoni b, loveridge c, cooke hj, gray nk. the dazl family proteins are pabp-binding proteins that regulate translation in germ cells. embo j 2005; 24: 2656–2666.
dorfman dm, genest dr, reijo pera ra. human dazl1 encodes a candidate fertility factor in women that localizes to the prenatal and postnatal germ cells. hum reproduc tion 1999; 14: 2531–2536.
findlay jk, hutt kj, hickey m, anderson ra. how is the number of primordial follicles in the ovarian reserve established? biol reproduc tion 2015; 93: 1–7.
fox m, urano j, reijo pera ra. identification and characterization of rna sequences to which human pumilio-2 (pum2) and deleted in azoospermia-like (dazl) bind. genomics 2005; 85: 92–105.
ill me, hu yc, lin y, page dc. licensing of gametogenesis, dependent on rna binding protein dazl, as a gateway to sexual differentiation of fetal germ cells. proc natl acad sci usa 2011; 108: 7443–7448.
grieve km, mclaughlin m, dunlop ce, telfer ee, anderson ra. the controversial existence and functional potential of oogonial stem cells. matutinas 2015; 82: 278–281.
gruss oj, carazo-salas re, schatz ca, guarguaglini g, kast j, wilim m, le bot n, vernos i, karsenti e, mattaj iw. ran induces spindle assembly by reversing the inhibitory effect of importin alpha on tpx2 activity. cell 2001; 104: 83–93.
hackett ja, reddington jp, nestor ce, dunican ds, branco mr, reichmann j, reik w, surani ma, adams ir, meehan rr. promoter dna methylation couples genome-defence mechanisms to epigenetic reprogramming in the mouse germline. development 2012; 139: 3623–3632.
haston km, tung yj, reijo pera ra. dazl functions in maintenance of pluripotency and genetic and epigenetic programs of differentiation in mouse primordial germ cells in vivo and in vitro. plos one 2009; 4: e5654.


Reynolds N, Collier B, Bingham V, Gray NK, Cooke HJ. Translation of the synaptonemal complex component Scy3 is enhanced in vivo by the germ cell specific regulator Dazl. RNA 2007;13:974–981.


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