

Table S2: Transcripts down-regulated greater than 2-fold in Dppa4-deficient ES cells with established or proposed functions in gametogenesis.

Gene Symbol	Annotation	Ref.
Syce1	„Synaptonemal central element 1“: structural component of meiotic synaptonemal complex	(4)
Mael	„Maelstrom“: component of <i>nuage</i> (germ-cell-specific organelle); essential for spermatogenesis	(20)
Sohlh2	„Spermatogenesis-and-oogenesis-helix-loop-helix-2“: germ-cell-specific transcription factor; essential for male and female gametogenesis	(3, 7, 23)
Dazl	„Deleted in azoospermia-like“: RNA-binding protein; essential for gametogenesis in males and females	(12, 19, 25)
Spesp1	„Sperm equatorial segment protein 1“: specific component of the sperm acrosome involved in sperm/egg binding	(24)
Ddx4 = Mvh	„Dead box polypeptide 4“ = „Mouse vasa homolog“: ATP-dependent RNA helicase; spermatogenesis is blocked in homozygous mutant mice, resulting in male infertility. Female mutant mice are fertile	(22)
Rhox5/Pem	„Reproductive hox 5“: imprinted gene encoding putative transcription factor; thought to be involved in regulation of gametogenesis; targeted deletion causes male subfertility associated with increased germ-cell apoptosis and decreased sperm count and motility	(6, 10, 14)
Hormad1	„HORMA domain 1“: strongly expressed in testis; contains putative chromatin-binding domain; homologous to Hop1, a yeast meiosis-specific protein, as well as to asy1, a meiotic synaptic protein in <i>Arabidopsis thaliana</i>	(2)
Stk31	„Serine threonine kinase 31“: poorly characterized; has been associated with female meiosis/germ cell development and spermatogenesis	(17)
Tex19	„Testis expressed gene 19“: expression restricted to germ cells and pluripotent cells; Tex19 knockout males exhibit impaired spermatogenesis	(11, 18)
Aard	„Alanine and arginine rich domain containing“: reported to show sexually dimorphic expression in fetal mouse gonads; specifically upregulated in Sertoli cells during	(21)

mouse testis differentiation

Aurkc	„Aurora kinase c“: single nucleotide deletion causes male infertility in humans owing to the production of abnormal sperm; knockout mice exhibit compromised fertility due to abnormal sperm	(5, 9)
Fthl17	„Ferritin heavy polypeptide-like 17“: X-linked, spermatogonially-expressed, germ-cell-specific gene; located near chromosomal breakpoints in 2 azoospermic patients	(13)
Cytct	„Cytochrome c testis“: testis-specific isoform of cytochrom c; sperm from Cytct-KO mice is less effective in fertilizing oocytes and contains large numbers of immotile spermatozoa; Cytct-KO mice exhibit early testicular atrophy	(16)
Hook1	Identical with „azh (abnormal spermatozoon head shape)“ gene: autosomal recessive mutation results in highly abnormal morphology of sperm heads and tails with resulting male infertility	(15)
Rhox6	„Reproductive hox 6“: member of the recently identified X-linked homeobox gene cluster, which is composed of genes selectively expressed in reproductive tissues; see also Rhox5/Pem	(8, 14)
Serpnb6c	„Serine protease inhibitor b6c“: member of the serpin family with sexually dimorphic expression in gonads	(1)

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