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Biochemical Society Transactions

The role of chromosome segregation and nuclear organisation in human subfertility

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20 **Abstract**

21 Spermatogenesis is central to successful sexual reproduction, producing large numbers of haploid
22 motile male gametes. Throughout this process, a series of equational and reductional chromosome
23 segregation precedes radical repackaging of the haploid genome. Faithful chromosome segregation
24 is thus crucial, as is an ordered spatio-temporal “dance” of packing a large amount of chromatin into
25 a very small space. Ergo, when the process goes wrong, this is associated with improper
26 chromosome number, nuclear position and/or chromatin damage in the sperm head. Generally,
27 screening for overall DNA damage is relatively commonplace in clinics, but aneuploidy assessment is
28 less so and nuclear organization studies form the basis of academic research. Several studies have
29 focussed on the role of chromosome segregation, nuclear organisation and analysis of sperm
30 morphometry in human subfertility observing significant alterations in some cases, especially of the
31 sex chromosomes. Importantly, sperm DNA damage has been associated with infertility and both
32 extrinsic (e.g. lifestyle) and intrinsic (e.g. reactive oxygen species levels) factors, and whilst some
33 DNA strand breaks are repaired, unexpected breaks can cause differential chromatin packaging and
34 further breakage. A “healthy” sperm nucleus (with the right number of chromosomes, nuclear
35 organization and minimal DNA damage) is thus an essential part of reproduction. The purpose of this
36 review is to summarise the state of the art in the fields of sperm aneuploidy assessment, nuclear
37 organization and DNA damage studies.

38 Introduction

39 Gametogenesis, the process of producing haploid gametes is central to successful sexual
40 reproduction, and in male mammals, spermatogenesis describes the transformation of germ cells
41 into spermatozoa. Taking place during three distinct phases, the mitotic proliferative phase, the
42 meiotic phase and the cytodifferentiation (spermiogenesis) phase, a series of events that includes
43 both equational and reductional chromosome segregation as well as radical repackaging of the
44 haploid genome occurs. Faithful chromosome segregation is thus crucial for the process to continue
45 normally, as is an ordered spatio-temporal “dance” of packing a large amount of chromatin into a
46 very small space. Given this, it is hardly surprising that, when the process goes wrong it is associated
47 with improper chromosome number, nuclear position or chromatin damage in the sperm head.

48

49 Given that infertility affects approximately one in six couples globally^[1], and that male factor
50 subfertility contributes to around 50% of these, there is an indisputable need for more research into
51 the male gamete to be undertaken to understand the role of chromosome segregation and
52 chromatin packaging in male infertility. To date however, studies have focused mostly on “spot
53 counting” i.e. interphase cytogenetics to establish the proportion of aneuploid cells in an ejaculate^[2],
54 studies to assess the overall levels of DNA damage in sperm heads^[3] and nuclear position of
55 chromosome territories^[4,5]. Whilst screening for overall DNA damage is relatively commonplace in
56 some IVF clinics, aneuploidy assessment is less so (although such techniques are nonetheless offered
57 by some companies). This review covers our current understanding of the importance of sperm
58 nuclear organisation and the mechanisms of chromosome segregation in human sperm, with a focus
59 on the differences between fertile and subfertile individuals. Given that the clinical definition of
60 infertility refers to one year of unwanted non-conception following unprotected intercourse in the
61 fertile phase of the menstrual cycle (WHO definition) and is sometimes used interchangeably with
62 sterility, here we use the term “subfertility” throughout to refer to any form of reduced fertility that
63 occurs over a prolonged period of time.

64 **Screening of sperm and a possible role for aneuploidy assessment**

65 When screening human semen for fertility evaluation, various different physical characteristics are
66 routinely assessed, including the volume, appearance, viscosity and pH of the ejaculate, as well as
67 the morphology of the sperm heads^[6]. Given that some studies have suggested that these routinely
68 assessed parameters are not entirely indicative of fertility^[7], it is clear that other screening methods
69 are necessary. Given that numerous studies have identified that there is a correlation between
70 sperm aneuploidy and male infertility^[8-12], irrespective of constitutional chromosome abnormalities
71 (i.e. men that have normal karyotypes, but compromised semen parameters), aneuploidy
72 assessment has been proposed as a potential alternative screening method that is currently not
73 routinely implemented clinically.

74

75 Infertile men who were previously unable to procreate are now able to, due to the development of
76 various methods of assisted reproductive technologies (ART) such as intracytoplasmic sperm
77 injection (ICSI). Potentially therefore, those men who fit the referral category for ICSI, in theory run
78 the risk of perpetuating aneuploidy to their offspring. Although the majority of autosomal
79 aneuploidies are maternal in origin, 7% are paternally derived (this equates to around 1 in 10,000
80 children with paternally derived Down syndrome for instance)^[13,14] and 50% of sex chromosome
81 aneuploidies also arise in the male gamete. That is, it has been shown that almost half of XYY, three
82 quarters of XO, 5% of XXX, and all XYY cases are a result of an aneuploid sperm^[15]. Aneuploid events
83 in sperm can be identified by fluorescence *in situ* hybridisation (FISH)^[8,16], which permits thousands
84 of sperm heads to be screened. The first reports which used FISH as a screening tool for human
85 fertility demonstrated that aneuploidy was far more common in men afflicted by severely comprised
86 semen parameters such as concentration (oligozoospermia), morphology (teratozoospermia) and
87 motility, (asthenozoospermia), together known as oligoasthenoteratozoospermia (OAT). The
88 presence, or not, of sperm aneuploidy in both fertile (normozoospermic) and infertile men has been
89 widely studied and the received wisdom is that all men produce a proportion of aneuploid sperm^[2].

90 As described above, the incidence of aneuploidy however has been positively correlated with
91 reduced semen parameters^[17], and these occurrences increase with the severity of the infertility.
92 Initial studies on sperm aneuploidy involved the analysis of karyotypes of those human sperm cells
93 that were capable of fertilising a hamster oocyte. Whilst this method permitted the detection of
94 both structural and numerical chromosome aberrations, this approach is challenging and time
95 consuming. Such studies revealed that structural chromosome abnormalities were more prevalent
96 than numerical incidences, and that non-disjunction events were most common in chromosomes 21,
97 22, X and Y compared to the rest of the chromosome complement. Given that meiotic
98 recombination assists homologous chromosomes to stay together and that these chromosomes
99 often only cross over once during recombination, it is not unexpected that these would be the most
100 affected pairs^[18].

101

102 The genetic quality of sperm cells used in ART must therefore be considered, and ultimately the
103 selection of a euploid sperm prior to ICSI is the ultimate goal^[19]. In the meantime, being aware of the
104 overall level of sperm aneuploidy (and hence the risks involved) is the primary option for patients.
105 Even though IVF clinics have the ability to screen sperm for aneuploidy (by outsourcing to a company
106 such as iGenomix), this is rarely performed. Given that aneuploid sperm are still capable of
107 fertilisation, and that aneuploidy has been estimated to be more likely in samples from subfertile
108 male^[8,15], it has been argued that such screening would be worthwhile. Of course another solution is
109 to screen the embryos of all ICSI patients by preimplantation genetic tests (PGT-A) and a recent
110 study^[20] has demonstrated that embryos from ICSI males have elevated levels of sex chromosome
111 aneuploidies. Aneuploid embryos can result from a non-disjunction event in the oocyte or sperm
112 cell, or via mitotic loss, mitotic gain or a non-disjunction event in the embryo itself. Thus, although
113 PGT-A is in itself controversial^[21] it is argued that severe male factor subfertility should be a referral
114 category for it.

115

116 **Sperm nuclear organisation**

117 Genome condensation is necessary prior to the transmission of the male genome to the offspring;
118 this involves the replacement of histones with a family of small, arginine-rich proteins, protamines to
119 ensure that the complexes occupy a minimal cell volume^[22]. This unique structure is important for
120 two reasons; protection from DNA damage and a fast and full unpacking of the male genome to the
121 oocyte^[23]. Faithful sperm chromatin packaging has been implicated as essential for the
122 establishment and continuation of a normal pregnancy^[5,10]. Some studies have suggested however
123 that the impact of abnormal sperm chromatin on embryo development is subject to not only how
124 severe the damage is, but also how efficient the oocyte is at repairing any abnormalities^[24].

125

126 The term nuclear organisation describes the spatiotemporal arrangement of the DNA and associated
127 proteins in the interphase nucleus. It is often assayed by establishing the specific positions occupied
128 by each chromosome territory (CT) and/or specific loci^[25,26]. In humans, investigations into the
129 organisation and spatial arrangement of CTs at interphase have provided valuable insights into
130 genome function, particularly when considering higher levels of control that transcend the impact of
131 the DNA sequence alone. Studies of nuclear organisation in somatic cells have also revealed a
132 correlation between the gene density of the chromosome and the radial positioning of CTs^[27]. In
133 many somatic cell types, it has long been established that gene-rich CTs are located towards the
134 interior of the nucleus, whereas gene-poor chromosomes are positioned in the peripheral regions^[28].
135 This arrangement has been shown to be cell-type and tissue-type specific, and is evolutionarily
136 conserved^[29]. The structure of CTs has been shown to be dynamic, and less physically constrained
137 than once thought^[30], thereby enabling genes to reposition from the periphery of the nucleus
138 towards the interior following a change in cell status caused by quiescence or senescence^[31]. It is
139 also evident that the organisation is imperative for cellular functions (such as transcription) to
140 proceed normally and it has been hypothesised that chromatin organisation may be associated with
141 epigenetic modifications^[32] (discussed later), genomic imprinting^[33] and X chromosome

142 inactivation^[34]. In human sperm, chromosomes are organised non-randomly^[5] and centromeres form
143 a chromo-centre (i.e. they cluster) in the nucleus interior, with telomeres positioned nearer the
144 periphery^[5]. This pattern is similar in many other mammalian species with the sex chromosomes also
145 clustering nearer the centre of the nucleus^[35]. Further to this, It has been well documented that
146 there is a functional significance for the ordered pattern of chromosomes in human sperm cells^[36],
147 and that, in turn, aberrant organisation is common in samples from subfertile men. Evidence
148 suggests that irregular chromosome organisation is correlated with delayed decondensation,
149 impacting the zygote's first mitotic division, and playing a role in sex chromosome aneuploidy
150 events. Such studies have been performed in both 2D and 3D^[4].

151

152 As described above, in recent years numerous studies have focussed on the role of chromosome
153 segregation, nuclear organisation and analysis of sperm morphometry in human subfertility^[5,27,37-39].
154 To date however, such studies are still few and far between in other mammalian species, particularly
155 those in agriculturally important species such as cattle (*Bos taurus taurus*), pigs (*Sus scrofa*
156 *domesticus*), goats (*Capra aegagrus hircus*) and sheep (*Ovis orientalis aries*)^[40-42]. A key goal in
157 animal production is to identification of subfertile animals (so that they can be removed from
158 breeding programmes in a timely manner). That is, animals with fertility problems have the potential
159 to produce reduced litter sizes throughout the breeding population, thereby impeding the
160 production of foodstuffs^[7]. Some of our current work aims to address this by comparing nuclear
161 topology and chromosome positioning in fertile and subfertile pig samples.

162

163 **DNA damage and the impact of epigenetic change**

164 Sperm DNA damage has been related to numerous different factors that can be both extrinsic (e.g.:
165 lifestyle factors)^[43,44] and intrinsic (e.g.: levels of reactive oxygen species (ROS)^[45,46]). Whilst some
166 DNA strand breaks are expected and subsequently repaired (for example those occur as part of
167 chromatin remodelling^[47]), unexpected breaks have the potential to cause chromatin to be packaged

168 differently, and may lead to further DNA breakage. Interestingly, it has been shown that men with
169 abnormal semen parameters present with high levels of an apoptotic protein, Fas^[48]. The presence
170 of this protein on spermatozoa is indicative of cells that had been set aside for programmed cell
171 death, but have evaded this due to the high numbers of cells present in the ejaculate. This
172 mechanism is known as 'abortive apoptosis' and can lead to oligozoospermia, azoospermia, or a high
173 number of abnormal sperm, which in turn may have an impact on successful fertilisation. It has been
174 well documented that men of reproductive age that are being, or that have been, treated with
175 chemotherapy present with impaired spermatogenesis, increased levels of sperm aneuploidy, and a
176 higher rate of DNA fragmentation^[49]. Whilst aneuploidy levels recover, levels of DNA damage may
177 not. Lifestyle factors such as smoking and obesity have also been correlated with reduced semen
178 quality and higher levels of DNA damage^[43,50]. Further to this, it has been shown that an increase in
179 the rate of DNA fragmentation is associated with lower natural, intrauterine insemination (IUI) and
180 IVF conception rates^[51,52].

181

182 There are several different ways in which the levels of sperm chromatin damage can be assessed, as
183 described in Figure 1. These include sperm nuclear matrix assays such as the chromatin dispersion
184 test^[53], the use of sperm chromatin structural probes such as the sperm chromatin structure assay
185 (SCSA)^[54], and DNA fragmentation assays such as the Terminal deoxynucleotidyl transferase dUTP
186 Nick-End Labelling (TUNEL)^[10] and comet assays^[53]. It has been shown that if an SCSA test detects
187 DNA fragmentation of over 30%, there is a far smaller chance for fertilisation to be a success via
188 natural pregnancy or IUI^[55]. This does not however seem to be the case for ICSI cases, and can
189 almost certainly be attributed to the fact that both the sperm and the subsequent embryo are
190 carefully selected prior to implantation. Nevertheless, pregnancy loss following IVF or ICSI has been
191 linked to abnormal levels of sperm DNA damage. It has been suggested that this is because
192 embryonic genome expression does not happen until the 4-8 cell stage, and therefore DNA damage

193 in sperm does not affect fertilisation *per se*, however there are only a limited number of studies that
194 have focussed on the effect of DNA fragmentation on IVF or ICSI outcomes^[51,52].
195 The role of epigenetics in human reproduction is an active and interesting field of research,
196 particularly due to the transgenerational effects attributed to epigenetic modifications in both male
197 and female gametes. Epigenetic patterns are metastable heritable changes in gene expression that
198 can change due to endogenous and environmental factors^[56,57]. For example, the epigenetics of
199 ageing sperm has been linked to increased frequency of neurocognitive disorders such as autism,
200 schizophrenia and other bipolar disorders, as well as metabolic dyshomeostasis and obesity in
201 offspring^[56]. At present, epigenetic modifications in sperm have been found to have an impact on
202 four key areas of reproduction: 1) spermatogenesis failure, 2) embryogenesis, 3) success rates and
203 overall outcomes associated with ART procedures and 4) long-term progeny effects^[57]. Whilst
204 several epigenetic modifications relating to DNA methylation and histone modifications have been
205 found to occur during various stages of spermatogenesis (mitosis, meiosis and spermiogenesis),
206 histone-protamine replacement has been found to be the main change in sperm cells^[57]. The
207 literature suggests that various features of male infertility, including oligozoospermia and OAT,
208 chromosomal aneuploidies, DNA fragmentation and chromatin packaging could all be linked to
209 epigenetic modifications occurring at various junctures of spermatogenesis. Paternal epigenetic
210 changes have also been associated with childhood cancers and imprinting diseases, and that such
211 changes are increased in offspring conceived via ART^[56], and further to this, it has been shown that
212 control of the paternal lifestyle (for example the use of dietary antioxidant supplements) before
213 conception may have a downstream impact^[57].

214

215 **Perspectives**

- 216 • **Importance of the field:** A “healthy” nucleus is an essential part of any cell or tissue. In
217 chromatin terms this can mean the correct number of chromosomes, the appropriate
218 organization of CTs and the absence of significant DNA damage. This is particularly apparent

219 in the sperm cell, in part because (being haploid and the end of a very specialized
220 developmental pathway) it does not have the opportunity to repair its DNA, eject offending
221 chromosomes, nor reorganize its chromatin. In this respect, analysis of sperm chromatin can
222 also be used to monitor the effects of toxic agents or environmental pollutants. Sperm are
223 our legacy to the next generation and thus, in this regard, with eggs, perhaps the most
224 important cells in our body.

225 • **Summary of current thinking:** Increased sperm aneuploidy, aberrant nuclear organization
226 and DNA damage have all been associated with male infertility and this is particularly
227 important for couples seeking fertility treatment such as ICSI. In this regard, the genetic
228 quality of the sperm is important as an indicator of the likely success of the procedure and
229 possibly could impact on the future health of the resultant child.

230 • **Comment on future directions:** One of the confounding factors in sperm head analysis is the
231 necessity to score a large number of cells. In sperm aneuploidy studies, this can mean up to
232 20,000 cells, which can be prohibitively laborious. Automated scoring is thus a priority and
233 adaptations of flow cytometry such as flow FISH are essential in this regard. Moreover, still
234 relatively little is known about the basic mechanisms that lead to chromatin damage in
235 sperm and this is a fascinating area that needs much deeper investigation, ultimately for
236 future patient benefit.

237

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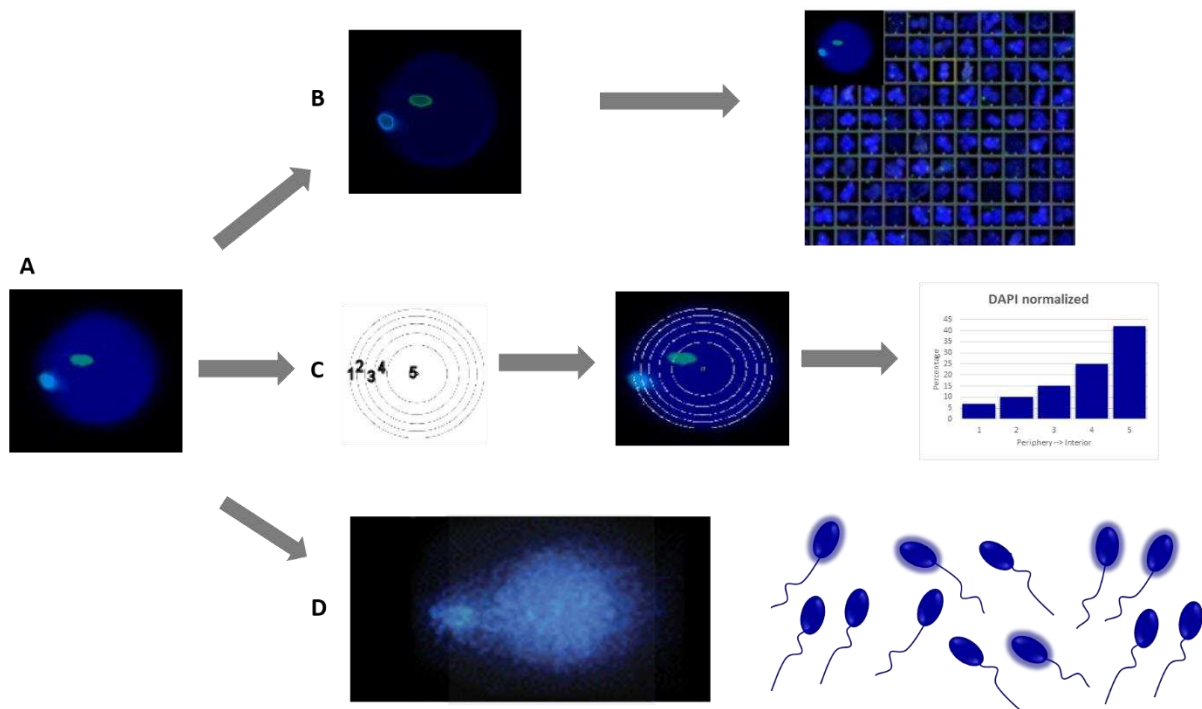


Figure 1: Assays to detect a “healthy” sperm nucleus. A: Sperm nucleus showing overall chromatin content (dark blue) plus two chromosomes highlighted in light blue and green. B: Chromosome signals can be detected and counted (either manually or using automated systems) then compared with numerous cells from the same ejaculate to establish the proportion that have extra or missing chromosomes (aneuploidy). Typically around 1000 cells are counted. C: The most common way to assess for chromosome territory (CT) position (and hence nuclear organization) is to overlay a template of five concentric shells and, taking into account overall DNA density, determine the proportion of signals that appear in each shell in around 50-100 cells, hence producing a histogram. D: Pre-treatment of the chromatin such as the COMET assay or sperm chromatin dispersion (SCD) test (both depicted) can give an indication of the proportion of cells with DNA damage. In general terms, assays for DNA fragmentation are commonplace in clinics, sperm aneuploidy testing is offered, but rarely, nuclear organization tests are still in the domain of research studies.