Meiotic and radiation studies in four oligochiasmatic men

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SUMMARY The meiotic findings in four oligochiasmatic males are described. Radiation studies on the somatic cells of three of them failed to provide evidence for a reduced facility to repair DNA which might also have accounted for the observed failure of chiasma formation at meiosis. The data support the idea that the 'low chiasma count' condition in sterile men is of mixed aetiology.

Mutants which drastically modify the normal pattern of behaviour at meiosis have been described for a number of species, both plant and animal (see Baker et al., 1976; and Koulischer et al., 1970; Dutrillaux and Guéguen, 1971; Skakkebaek et al., 1973; Chaganti and German, 1974; Koulischer and Schouysman, 1974; Ferguson-Smith, 1976; Templado et al., 1976). At pachytene in the majority of these cases, the bivalents appear to be completely synapsed (desynaptic type), but in two patients, one described by Hultén et al. (1974a), the other by Ferguson-Smith (1976), the formation of the synaptonemal complex was defective and the sex vesicle not identified (synaptic type).

That some of these effects may have had a genetic cause is suggested by the fact that, in three cases, the men were from consanguineous marriages (Hultén et al., 1970; Dutrillaux and Guéguen, 1971; Ferguson-Smith, 1976). In a fourth case, a maternal uncle and a maternal male cousin of the patient were both sterile (Chaganti and German, 1974), and in a fifth case (Ferguson-Smith, 1976), two married brothers and one married sister of the patient were also sterile.

Among the most important generalisations to have emerged from the study of meiotic mutants in other species is that, if one, and in the case of the metabolic processes involved in meiotic development and recombination to appear to operate also in somatic cells to ensure chromosome stability, both spontaneously and after irradiation. The relationship is illustrated by the many reciprocal observations that mutants affecting chiasma frequency of recombination also often exhibit somatic chromosome instability, while mutants recovered because they are sensitive to irradiation or radiomimetic drugs often prove to have meiotic effects, particularly on recombination (Baker et al., 1976).

The idea that the same biochemical pathways could be involved in genetic recombination, repair replication, and the formation of chromosomal exchanges (Evans, 1966; Kihlm and Harley, 1968) has led to studies being undertaken to investigate the response to irradiation of peripheral blood lymphocytes or cultured fibroblasts from 'low chiasma count' men (Adams, 1970; Pearson et al., 1970; Clarkson, 1972; Ferguson-Smith, 1978). Aberration induction was assessed after x-rays, and levels of unscheduled DNA synthesis, indicative of repair replication were estimated in autoradiographic preparations of somatic cells exposed to 3H thymidine after UV or x-rays. In one published case (Pearson et al., 1970), the patient was a man with a mean chiasma count at meiosis of 9 per cell, who showed germ cell maturation arrest after metaphase...
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I and resulting azoospermia. The studies showed an approximately 50% reduction in levels of unscheduled DNA synthesis compared with controls after both UV and x-rays, and an aberration yield reduced about 20% below control levels. The authors suggested that these effects, and the observed failure of chiasma formation at meiosis, could all be accounted for by a reduced facility for repairing DNA in the cells of this particular subject. No other evidence for reduced repair capacity in the somatic cells of an asynaptic or desynaptic man, however, appears to exist. Three other cases, mentioned by Pearson et al. (1970), have been tested for UV and x-ray induced unscheduled DNA synthesis and all were found to be normal compared with controls (Adams, 1970; Clarkson, 1972). The case described by Ferguson-Smith (1976) was also normal in respect of response to UV irradiation when tested for unscheduled DNA synthesis (excision repair) (Ferguson-Smith, 1976) and postreplication repair (Lehmann et al., 1977). A fifth case, referred by J. German, in which the meiotic defect showed arrest of prophase at pachytene, also showed normal levels of postreplication repair after UV and normal survival after treatment with γ-rays, UV, and mitomycin C (Lehmann et al., 1977).

The opportunity to study four more cases of this kind has now arisen in our own laboratory and it seemed important, therefore, to test the response of their somatic cells to irradiation. The four patients, all of whom were phenotypically normal, were ascertained in a systematic study of patients undergoing testicular biopsy for investigation into the causes of their infertility. From one subject to another, they displayed in their spermatocytes at metaphase I a range of chiasma reductions from slightly subnormal to virtually achiasmatic, and in their testicular histology, degrees of spermatogenic impairment ranging from mild to severe. In no case was there reason to believe that exogenous factors were responsible for the meiotic irregularities, and in no case was an obvious abnormality observed in the somatic karyotype.

Case reports and meiotic investigations

CASE 1. ST 317

This case was a 32-year-old painter who presented at the Subfertility Clinic in May 1971 after two years of childless marriage. He had two sisters, one of whom was married with a child and the other unmarried, and one unmarried brother. Two seminal samples, taken one month apart, gave a mean sperm count of 8.5 m/ml, mean motility of 28%, and mean normal morphology of 65%. Histologically, the spermatogenic activity appeared normal and meiotic studies carried out on air-dried preparations showed all stages of spermatogenesis present on the slides. A distribution count on 200 dividing cells was within normal limits compared with findings for control patients (Table 1). No obvious pairing abnormalities were seen in the meiotic prophase stages, but at metaphase I a mean chiasma count of 40.6 per cell was recorded for 26 cells. This was subnormal compared with the normal mean frequency of 50-1 per cell recorded for human spermatocytes (Paris Conference, 1971). In 56% of cells, the X and Y chromosomes were present as univalents, a factor associated with maturation failure in germ cells (Miklos, 1974; Chandley et al., 1976).

CASE 2. ST 322

This case was a 26-year-old civil servant, childless for nearly 3 years before presenting at the Subfertility Clinic in August 1971. His wife later became pregnant and a baby was born in 1973. The patient then remarried in 1974 and returned to the Subfertility Clinic in 1976. Another baby was born later that year to his second wife. He had two unmarried brothers and one unmarried sister. On physical examination, both testes were found to be small, but no other phenotypic abnormality was recorded. The seminal analysis, based on two samples, showed a mean sperm count of 14.5 m/ml, mean motility of 51%, and mean normal morphology of 68%. Histologically, the biopsy was classified as normal and the air-dried meiotic preparations showed all stages of spermatogenesis on the slides. The distribution count gave evidence, however, for a proportion of cells failing to reach metaphase II compared with controls (Table 1). No obvious synaptic abnormalities were seen at meiotic prophase, but at metaphase I the mean chiasma count was low at 39.3 per cell for the 26 cells analysed. Dissociated X and Y chromosomes were seen in 76% of 46 cells analysed. Four of these cells also showed a single pair of autosomal univalents.

<table>
<thead>
<tr>
<th>Case</th>
<th>Spermatogonial metaphase (%)</th>
<th>Metaphase I (%)</th>
<th>Metaphase II (%)</th>
<th>Total cells analysed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Case 1</td>
<td>8.5</td>
<td>55.5</td>
<td>36.0</td>
<td>200</td>
</tr>
<tr>
<td>Case 2</td>
<td>5.5</td>
<td>68.5</td>
<td>26.0</td>
<td>200</td>
</tr>
<tr>
<td>Case 3</td>
<td>6.0</td>
<td>90.0</td>
<td>4.0</td>
<td>100</td>
</tr>
<tr>
<td>Case 4</td>
<td>31.0</td>
<td>67.0</td>
<td>2.0</td>
<td>100</td>
</tr>
<tr>
<td>Controls</td>
<td>10.9</td>
<td>52.8</td>
<td>36.4</td>
<td>14 020</td>
</tr>
</tbody>
</table>
CASE 3. ST 526
This case was a 32-year-old machine operator, married for 5½ years before presenting at the Subfertility Clinic in February 1977. He had two brothers and four sisters, all of whom were married with children. On physical examination, his left testis was found to be small and the right testis of normal size. A seminal analysis, based on a single specimen, showed a sperm count of 3 m/ml, motility of 30%, and normal morphology of 14%. The histological report noted that the tubules contained the normal content of spermatogonia and spermatocytes, but complete spermatogenesis was seen in only a small proportion of tubules. Many obviously degenerating primary spermatocytes were seen and few spermatids or spermatozoa were formed. The distribution count showed a severe reduction in numbers of cells reaching metaphase II. Prophase pairing appeared normal in all cells examined, but in metaphase I there was much obvious degeneration and many univalent chromosomes. A mean chiasma count of 28·1 was recorded for 15 cells which were still sufficiently healthy to analyse. All showed dissociated X and Y chromosomes, several large single chiasma bivalents, and from one to five pairs of autosomal univalents. However, the mean chiasma count recorded for this patient was spuriously high owing to the fact that cells with very low chiasma counts and many univalents, which made up the majority, were too degenerate to analyse. A relatively healthy C-banded metaphase I is shown in Fig. 1. This cell had a chiasma count of about 25 and 4 pairs of univalents including the dissociated XY pair. C-banded metaphase II division is shown in Fig. 1.

Fig. 1 (a) Metaphase I and (b) metaphase II from case 3. Autosomal univalents are marked U. C-banded preparations.
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Fig. 2  Histological section of seminiferous tubules from case 4.

Fig. 3  Testicular cells from the air-dried preparations of case 4.  
   a, spermatogonial metaphase;  
   b, pachytene;  
   c, diplotene;  
   d, metaphase I.
CASE 4. ST 537
This case was a 25-year-old soldier who presented at the Subfertility Clinic in March 1977 after 3½ years of childless marriage. He had one brother married with children. Physical examination showed that both testes were small and a seminal analysis based on a single specimen revealed azoospermia. Histologically, the testicular biopsy showed maturation arrest at the primary spermatocyte stage in the majority of tubules and the presence of spermatozoa in only a minority (Fig. 2). In the air-dried meiotic preparations there was an unusually high proportion of spermatogonial metaphases, but very few cells passed through meiosis to reach metaphase II (Table 1). The spermatogonial metaphases appeared normal (Fig. 3a), as did the majority of pachytene spermatocytes. However, in one or two cells at pachytene, the bivalents displayed synaptic irregularities, the chromosomes showing paired and unpaired segments (Fig. 3b). The appearance of these aberrant human pachytene cells was reminiscent of some of the pachytene spermatocytes we have observed in a male hinny, the sterile hybrid between the horse and donkey. In this animal, a range of pairing irregularities, attributable to parental karyotypic differences, are observed at meiotic prophase (Chandley et al., 1974). At diplo- tene in case 4, the bivalents appeared curly and generally desynapsed (Fig. 3c), and at metaphase I the majority of chromosomes were present as univalents (Fig. 3d). The cells appeared virtually achiasmatic and, in the majority of cases, fuzzy and degenerate.

Radiation studies
To test the response of somatic cells to irradiation, blood lymphocytes from case 2 and from two healthy control males with normal chiasma counts were irradiated with x-ray doses of 0, 75, 150, and 300 rad. (Blood from case 1 could not be obtained.) Irradiations were carried out in plastic syringes, before culture (GO), using a 250 kV Siemens machine operated at 6 mA filtered by a Thraeaus filter. The cells were cultured for 48 hours by a modification of the method of Hungerford (1965). Slides were stained with aceto-orcein, coded, and scored blindly by one observer (MK). A total of 100 cells per sample were analysed for dicentrics, rings, and deletions.

Levels of unscheduled DNA synthesis in peripheral blood lymphocytes from the same three patients were assessed autoradiographically after UV exposures of 0, 15, and 30 seconds from a Hanovia low pressure mercury vapour lamp. (Unfortunately, total doses delivered cannot be given, as at the time of conducting the experiments the record was not kept of the dose rate for the UV lamp.) Exposure of lymphocytes to UV was carried out in phosphate buffered saline (PBS), the cell suspension forming a 3 to 4 mm deep layer in the bottom of a glass petri dish; in order to obtain a uniform exposure, this was continuously stirred with an electromagnetic stirrer during irradiation. Immediately after irradiation, cells were transferred to 10 ml plastic tubes and 3 ml aliquots of medium were added. For the autoradiography, 10 µCi/ml H thymidine (specific activity 17 to 25 Ci/mm) was added to the medium and the cells incubated for 2 to 3 hours at 37°C. The medium was then spun off and the cells washed twice in PBS, fixed in acetic alcohol (3:1), and air-dried onto slides. The slides were stained with carbol fuchsin before filming with Ilford L4 emulsion. The filmed slides were then left to expose in the dark at 4°C for two weeks and developed in Kodak D19B developer for five minutes. Grain counts scores were counted independently by two observers, each of whom scored 50 lymphocytes from each patient. Further blood samples could not be obtained from case 3 or 4, but skin fibroblasts were cultured from these two patients along with fibroblasts from a fertile control male (chiasma count not known) and a control female, and a 7-year-old girl with severe manifestations of xeroderma pigmentosum. In the latter case, reduced levels of unscheduled DNA synthesis are expected since such subjects exhibit a greatly reduced capacity for repairing UV-induced DNA lesions (Cleaver, 1968, 1969). Levels of unscheduled DNA synthesis after UV doses of 30, 90, 300, 900, and 3000 ergs/cm² were assessed in autoradiographic preparations, which were prepared in a manner similar to that described above for lymphocytes. All slides were coded and randomized before scoring and grain counts over 100 cells were scored at each dose level for each subject by two independent observers. The numbers of heavily labelled S-phase cells in the cultures were also recorded, and the percentage of cells showing unscheduled DNA synthesis was noted in addition to the grain counts per cell. Attempts were made to grow up cells from the fibroblast cultures of cases 3 and 4 in order to test for postreplication repair after UV irradiation and γ-ray survival, but the plating efficiency was so poor that these studies had to be abandoned temporarily (C. Arlett, personal communication). Further attempts are, however, contemplated.

Results
The levels of aberration induction in blood lymphocytes after x-rays in case 2 and the two controls were
given in Table 2. No significant differences were found from controls. Levels of UV stimulated unscheduled synthesis in these three patients are given in Table 3. There was no significant difference between the case and the controls.

The results of the unscheduled synthesis experiments on fibroblasts from cases 3 and 4 and controls are shown in Fig. 4. Again, no significant differences were detected between cases and controls. Only in the case of the girl with xeroderma pigmentosum were low levels of incorporation of $^3$H thymidine found and this was expected for this particular condition.

Table 2 Percentages of aberrant metaphases seen in X-irradiated peripheral blood lymphocytes from case 2 and 2 control males with normal chiasma frequencies. (100 lymphocytes analysed at each dose for each subject)

<table>
<thead>
<tr>
<th>X-ray dose (rads)</th>
<th>0</th>
<th>75</th>
<th>150</th>
<th>300</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control 1</td>
<td>0</td>
<td>6</td>
<td>26</td>
<td>69</td>
</tr>
<tr>
<td>Control 2</td>
<td>1</td>
<td>10</td>
<td>---</td>
<td>48</td>
</tr>
<tr>
<td>Case 2 (ST 322)</td>
<td>1</td>
<td>8</td>
<td>26</td>
<td>61</td>
</tr>
</tbody>
</table>

*Culture failed.

Table 3 Mean grain count per lymphocyte after UV irradiation and $^3$H thymidine incorporation for case 2 and controls. (100 lymphocytes analysed at each dose for each subject)

<table>
<thead>
<tr>
<th>UV exposure time (sec)</th>
<th>0</th>
<th>15</th>
<th>30</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control 1</td>
<td>1.1</td>
<td>37.6</td>
<td>38.7</td>
</tr>
<tr>
<td>Control 2</td>
<td>2.8</td>
<td>30.8</td>
<td>38.3</td>
</tr>
<tr>
<td>Case 2 (ST 322)</td>
<td>1.7</td>
<td>38.7</td>
<td>32.2</td>
</tr>
</tbody>
</table>

Discussion

Our negative findings concerning somatic cell sensitivity to irradiation in these particular oligochiasmatic men did not enable us to gain further insight into the possible cause of the meiotic irregularities. It would seem, however, that unlike in the case studied by Pearson et al. (1970), the factor or factors responsible for impaired chiasma formation at meiosis are not involved in maintaining chromosomal stability at mitosis. This could also be said of at least five other cases of this kind which have been studied (Adams, 1970; Clarkson, 1972; Ferguson-Smith, 1976; Lehmann et al., 1977). It may, of course, be argued that only those subjects showing severe reductions in chiasma frequency at meiosis would be likely also to show mitotic effects. Certainly, in three cases tested for unscheduled synthesis by Clarkson (1972) and in two of our own cases (cases 2 and 3), chiasma frequencies were reduced by only a half or less. However, there still remains our case 4 and the case reported by Ferguson-Smith (1976), both of whom showed a virtually achiasmatic picture at meiosis, but no effects, either spontaneous or radiation-induced, in their somatic chromosomes.

The answer in these cases may lie, however, not in a failure to repair DNA, but in a defect of some other process operating specifically at meiosis along the synopsis/recombination pathway, which has no function in somatic cells. Possible candidates for this role might be zygote DNA synthesis, or the r-protein/lipoprotein complex, both of which, under normal conditions, ensure orderly synopsis of homologues at meiosis (Stern and Hotta, 1977). Neither appears to function in somatic cells (Hotta and Stern, 1971; Stern and Hotta, 1971, 1977), but
inhibition of either can result in asynapsis at meiosis (Roth and Ito, 1967; Hotta and Shephard, 1973). A third possibility is the formation or functioning of the enzymes necessary for breakage and repair of DNA to affect meiotic recombination (Howell and Stern, 1971). Failure of nicking or ligation of DNA strands at pachytene could result in the formation of achiasmatic bivalents at meiosis, but since these enzymes appear to function specifically during zygotene and pachytene (Howell and Stern, 1971), their malfunctioning would be expected to produce little or no effect on the somatic elements.

Finally, one interesting corollary to these investigations is provided by a study of meiosis in a man with severe manifestations of the skin condition xeroderma pigmentosum (Hultén et al., 1974b). In the particular case investigated, repair replication after UV light, as measured by levels of unscheduled DNA synthesis in cultured fibroblasts, was greatly reduced compared with normal controls. Meiosis, however, was normal and there was no evidence of defective pairing or impaired chiasma formation in the spermatocytes. This, therefore, provides another situation in which the metabolic pathways operating to repair UV-damaged DNA in somatic cells and those operating at meiosis to ensure normal chiasma formation were independent of each other.

Meiotic studies in other men showing repair defects or chromosomal instabilities in their somatic cells might help to give us clearer insight into the possible operation of common mitotic and meiotic metabolic pathways in man. Further radiation response studies in oligochiasmatic men would also undoubtedly be informative. Such men are rare in the population, however, and opportunities for this kind of study are therefore extremely limited. It does seem likely, however, as previous authors have suggested (Hultén et al., 1974a; Templado et al., 1976), that the 'low chiasma count' condition in sterile men is of mixed aetiology.

References


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