Germ-line intrachromosomal recombination restores fertility in transgenic MyK-103 male mice

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Males of the MyK-103 line of transgenic mice are fertile and sire litters of normal size, but they never transmit the transgene, whereas females transmit the transgene with normal frequency. The chromosome originally bearing the transgene can be transmitted through the male germ line, but only after the transgene is deleted or rearranged by intrachromosomal recombination. The transgene encodes a functional herpes simplex virus (HSV) thymidine kinase gene that causes sperm infertility when expressed in postmeiotic germ cells. Immunocytochemistry revealed clones of germ cells that fail to express HSV thymidine kinase. We postulate that these cells arose by transgene deletion in embryonic germ cells and postnatal spermatogonial stem cells and that they are responsible for the normal fertility of MyK-103 males. The frequency of recombination events at the integration locus suggests that it contains a hotspot for mitotic recombination.

[Key Words: recombinational hotspot; spermatogonial stem cell; transmission ratio distortion]

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The MyK-103 pedigree has a unique pattern of male germ-line transmission [Palmiter et al. 1984]. Males are fertile in natural matings but never transmit the integrant to their offspring. The block to germ-line transmission in MyK-103 males occurs prior to fertilization because males and females both have litters of normal size, but only females transmit the integrant to roughly half of their offspring. This pattern of germ-line transmission appears similar to the cases of transmission ratio distortion that have been studied at the mouse t complex and the Drosophila segregation distorter locus (Bennett 1975; Hartl and Hiraizumi 1976). In these cases, however, the germ-line transmission of one allele has a selective advantage over the other of up to 99%, but it is not absolute, as in MyK-103 males.

Initially, we proposed that transmission ratio distortion in MyK-103 males resulted from a selective disadvantage of sperm containing the transgene due to integration near a gene that was necessary for the formation of fertile spermatozoa [Palmiter et al. 1984]. The observation that there was a depletion of transgenic sperm in the cauda epididymus of some males supported this view. A complication of this interpretation is that all the descendants of a spermatogonial stem cell are connected by intercellular bridges during premeiotic divisions, meiosis, and postmeiotic maturation. These bridges are about 1 μm in diameter (Dym and Fawcett 1971) and would, in principle, allow sharing of macromolecules among all the cells in the clone. Thus, to account for transmission ratio distortion in MyK-103 males as due to insertional inactivation of a gene critical for spermatogenesis, we would also have to propose that development of the haploid transgenic spermatids could not be rescued by their meiotic partners bearing wild-type alleles.

We subsequently cloned the integration site of the transgene and found that it was flanked by a 5-kb duplication of DNA that was present only once at the preintegration site on chromosome 6 [Wilkie and Palmiter 1987]. This observation suggested the possibility that the transgene, which contains a functional copy of the metallothionein/herpes simplex virus (HSV) thymidine kinase fusion gene [pMK2], might frequently be lost by intrachromosomal recombination through the duplicated sequences. If the transgene, instead of resulting in insertional inactivation of an essential gene, resulted in expression of a gene product that caused sperm infertility, then the transgenic males would be sterile due to sharing of gene products among spermatids. However, recombination events that deleted the transgene could result in clones of wild-type spermatids that might develop normally and account for the fertility of the MyK-103 males.

This paper provides both genetic and immunocyto-
chemical data supporting the latter view. We used a restriction fragment length polymorphism (RFLP) within the 5-kb duplicated sequence that allowed us to distinguish male germ-line transmission of the chromosome bearing the transgene independently of the transgene itself. We show that deletion of the Myk-103 transgene by homologous recombination occurs at a high frequency in embryonic germ cells and mitotically active spermatogonial stem cells, resulting in the equal transmission of the recombinant MyK-103 chromosome and its meiotic partner. The evidence presented here, and independently by Braun et al. (1989, 1990), shows that macromolecules can be shared among syncytial spermatids and that postmeiotic expression of HSV thymidine kinase causes sperm infertility. The synthesis of these observations renders a satisfying explanation of the transduction distortion in MyK-103 males, one of the oldest lines of transgenic mice in existence (Palmiter et al. 1982).

Results

Homologous recombination in the male germ line deletes the transgene

The structure of the MyK-103 integrant is shown in Figure 1. Genetic instability of this integrant could result from deletion by intrachromosomal recombination between the flanking duplicated mouse sequence. We would not have detected male germ-line transmission of the hypothetical recombinant chromosome in the MyK-103 pedigree because it would be indistinguishable from the nontransgenic homolog in mice of the original C57BL/6 × SJL background. To test this hypothesis, we used a RFLP resulting from a 1.5-kb deletion in AKR MyK-103 mice. As diagramed in Figure 2, the 6A allele from their father and the 6M allele from their mother were chosen as test-cross males to be mated with AKR females. Considering all possibilities, the test-cross males (6A/6M) might transmit the 6A or 6M allele, as well as the 6C allele, which could be generated from the 6M allele by homologous recombination. Segregation of the 6A, 6M, and the recombinant 6C alleles was assayed in the test-cross progeny by Southern blot or dot hybridization of tail DNA. The Southern blot shown in Figure 3B compares the hybridization patterns of genomic DNA isolated from parental mice, test-cross males, and test-cross progeny. DNA was digested with BamHI and hybridized with the E-N probe (Fig. 3A), which maps completely within the deletion in the 6A allele. MyK-103 DNA (6M/6C) yielded two bands (6.5 and 5.5 kb) that hybridized to the E-N probe. The 5.5-kb band is twice the intensity of the 6.5-kb band because the former is found in both the 6M and 6C alleles, whereas the latter is derived only from the 6M allele. DNA from AKR mice (6A/6A) did not hybridize, as expected, and DNA from test-cross males (6M/6A) yielded

Figure 1. Structure of the Myk-103 integrant. The integrant is composed of 19 kb of DNA flanked on either side by a 5-kb duplication of mouse DNA that is normally found in single copy at this locus. The integrant contains four fragments of the plasmid pMK (w, x, y, and z). The sequences in pMK include pBR322 [thin line], the metallothionein I promoter (hatched box), the HSV thymidine kinase coding sequence (thick black line), and downstream sequence (shaded line; for further details, see Wilkie and Palmiter 1987). Note that the duplicated flanking sequences (box, arrow inclusive) are in the same relative orientation, as are the repeated DNA sequences of pBR322 present in the pMK fragments marked w, y, and z. The novel junctions formed by ligation of pMK fragments are marked J2 through J6. A 532-bp fragment of mouse repetitive DNA is positioned between J6 and J7 [Wilkie and Palmiter 1987]. The origin of the restriction fragments that were used to prepare hybridization probes is indicated: [BM] 1.5 kb BamHI, [BG] 1.6 kb BglII, [MT] 0.6 kb Stul-BglII, [TK] 2.0 kb BgIII-Smal, [EN] 0.9 kb EcoRI-NcoI, and [EP] 0.6 kb EcoRI-PvuII. The sequences of all probes except EP occur twice in the Myk-103 insert. [Diagram adapted from Wilkie and Palmiter 1987.]

Figure 2. MyK-103 × AKR test-cross. MyK-103 mice are heterozygous at the integration locus; one homolog (6M) contains the prototype Myk-103 insert and is flanked by the duplicated sequence, and the other homolog (6C) contains one copy of this sequence, as do C57BL/6 and SJL mice. AKR mice are heterozygous for a small deletion (6A) within the sequence that was duplicated in the 6M allele.
the 5.5-kb and 6.5-kb bands at equal intensity because only the 6M allele hybridized to the E-N probe. This blot also shows that when 11 test cross progeny were assayed, seven offspring inherited the 6A allele and four offspring inherited the 6C allele from their fathers. In total, 54 progeny sired by five test-cross males were assayed by dot hybridization, first with the E-N probe [Fig. 3C] and, after this probe was removed from the filter, with pBR322 [data not shown]. None of the progeny inherited the 6M allele, consistent with the observation that MyK-103 males never transmit the prototype MyK-103 insert. The number of test-cross progeny that inherited the 6A or 6C allele was nearly equal; 26 progeny inherited the 6A allele and the remaining 28 progeny inherited the recombinant 6C allele from their fathers. A Southern blot of sperm DNA from three of these test-cross males that was hybridized with the metallothionein I [MT-I] probe showed that roughly half of their sperm carried the transgene [Fig. 3D]. Thus, the vast majority of sperm produced by the test-cross males carried the 6M allele or its 6A meiotic partner, yet the recombinant 6C allele was transmitted at essentially equal frequency with the 6A allele. Therefore, sperm that carried the transgenic 6M allele or its 6A meiotic partner were infertile. The only fertile sperm that were produced by the test-cross males carried the recombinant 6C allele or its 6A meiotic partner and, relative to the transgenic sperm, were few in number.

**Mosaic expression of HSV thymidine kinase in MyK-103 testes**

The genetics indicated that the MyK-103 testis was a mosaic that contained both transgenic and recombinant nontransgenic germ cells. To visualize this mosaicism, we took advantage of the fact that HSV thymidine kinase mRNA was readily detected in testes [Fig. 4A] and the protein could be visualized by immunocytochemistry.

In MyK-103 males, HSV thymidine kinase expression in the testis appears to be restricted to round and elongating spermatids, with the heaviest staining in the latter cell type [Fig. 5B]. The immunocytochemical assay did not detect HSV thymidine kinase protein in the testes of nontransgenic littermates [Fig. 5A]. In the six MyK-103 hemizygous males that were assayed, some seminiferous tubules expressed HSV thymidine kinase in all spermatids. Considering that only half of the spermatids inherited the pMK transgene following meiosis in a hemizygous male, it is striking that even those spermatids that inherited the nontransgenic homolog of chromosome 6 contained HSV thymidine kinase protein. The equal distribution of HSV thymidine kinase between genetically distinct spermatids was presumably accomplished by movement of the mRNA, and perhaps protein as well, across the intercellular bridges in a syncytial cluster of spermatids [Braun et al. 1989, 1990]. Thus, all spermatids developing within a syncytium that expressed HSV thymidine kinase were phenotypically transgenic.

Male sterility has recently been correlated with testicular expression of HSV thymidine kinase in several different lines of transgenic mice [Al-Shawi et al. 1988; Iwakura et al. 1988; Heyman et al. 1989]. In the most definitive study, all lines that expressed relatively high levels of HSV thymidine kinase in the spermatids of hemizygous mice were sterile, whereas low-level expres-
Germ-line mosaicism via mitotic recombination

Figure 4. Northern analysis of MyK-103 testis RNA. Total RNA (10 μg) was isolated from either liver [L] or testis [T] from five adult males and hybridized with a probe specific to one of three probes. (A) HSV thymidine kinase [TK]. (B) The EP restriction fragment. (C) A 0.6-kb EcoRI fragment including coding sequence of the mouse protamine 1 gene [Prm-1; Peschon et al. 1987]. Male 96-1 was a nontransgenic offspring in the MyK-103 pedigree, males 90-5, 97-5, and 33-4 inherited the prototype MyK-103 insert, and male 100-1 inherited the rearranged transgene δMyK.

pression HSV thymidine kinase elicited intricate mosaic patterns in antibody-stained cross sections of Myk-103 testes. Some of the seminiferous tubules that were revealed by cross section did not express HSV thymidine kinase in any of their spermatids (data not shown). The absence of expression in certain tubules did not simply reflect developmental regulation of HSV thymidine kinase translation because elongating spermatids are present in all seminiferous tubules and elongating spermatids in each stage of development could express HSV thymidine kinase. Other seminiferous tubules appeared sectored by distinct patches of spermatids that did not express HSV thymidine kinase [Fig. 5B, tubule 1]. Juxtaposition of two patches of spermatids within a single tubule defines the boundary of at least two syncytial clusters of spermatids. Sectoring was also observed in the mosaic tubule shown in Figure 5D where the elongating spermatids expressed HSV thymidine kinase, but it was not expressed in the underlying round spermatids. A similar pattern was apparent in the tubules marked 2–5 in Figure 5B. Elongating and round spermatids develop in separate syncytia, prohibiting the diffusion of HSV thymidine kinase between these two cell types. The elongating spermatids that expressed HSV thymidine kinase were descendents of spermatogonia that committed to spermatogenesis in the seminiferous epithelium one cycle earlier than the underlying round spermatids within the same tubule.

On the basis of our immunocytochemical analysis, we can refine the interpretation of the genetic results presented above. We infer that the transgenic sperm and their meiotic partners which were infertile in MyK-103 males developed from syncytial germ cells that expressed HSV thymidine kinase. The spermatids that did not express HSV thymidine kinase were descendents of spermatogonial stem cells which, at some time during their development and regenerative divisions, deleted the pMK transgene via homologous recombination (Fig. 6). Those spermatogonial stem cells in mosaic MyK-103 testes that had deleted pMK produced fertile sperm.

The contrast between syncytia that did or did not express HSV thymidine kinase did not affect male germ-line transmission of the HSV thymidine kinase gene [Braun et al. 1990]. Immunocytochemical analysis of HSV thymidine kinase expression levels in the round and elongating spermatids of these transgenic mice was compared to expression in MyK-103 spermatids. The intensity of staining in MyK-103 testes was similar to that observed in testes from the sterile transgenic males [data not shown]. Some other unusual features seen in the sterile mice, such as multinucleated giant cells in the ducts of the cauda epididymis [Braun et al. 1990], were also found in some MyK-103 males. These similarities indicated that HSV thymidine kinase expression in MyK-103 spermatids contributes to, and may be the sole cause of, infertility of transgenic sperm and their meiotic partners. However, MyK-103 males are fertile because a significant fraction of their spermatid syncytia do not express HSV thymidine kinase.

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It is possible to estimate the deletion frequency of pMK per cell division in the MyK-103 male germ line given the age of the mouse when sacrificed and the percentage of nontransgenic spermatids, in addition to certain assumptions [see Hartl 1971]. The assumptions are (1) the deletion frequency was constant throughout em-
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Figure 5. Immunocytochemical analysis of HSV thymidine kinase expression. Testes from adult animals were fixed, sectioned, and stained with primary and secondary antibodies as described in Materials and methods. (A) Control testis; magnification, 90×. (B) Hemizygous MyK-103 testis (92-3) 11 weeks of age; magnification, 62×. (C) Hemizygous ΔMyK testis; magnification, 90×. (D) 92-3 testis, counterstained with hematoxylin; magnification, 360×. Expression of HSV thymidine kinase was restricted to round and elongating spermatids.

In addition to HSV thymidine kinase, MyK-103 testes also expressed another transcript that was not observed in nontransgenic siblings. To identify this transcript we screened restriction fragments of mouse genomic DNA that were isolated from A. clones that spanned either flank of the pMK insert (X2 and X21; see Wilkie and Palmer 1987). Northern blots of poly(A) RNA isolated from MyK-103 and nontransgenic testes were hybridized with the various restriction fragments. The hybridization probe EP identified an 800-nucleotide transcript specifically in MyK-103 testes (Fig. 4B). This probe was isolated from mouse DNA to the right of the pMK insert (Fig. 1) and is present in single copy per haploid genome; it does not cross-hybridize with the transgene pMK or to mouse repeat DNA. To assay the normal expression pattern of this transcript, the EP probe was hybridized to Northern blots of RNA that was isolated from a variety of germ cell sources, including primitive type A spermatogonia, A and B spermatogonia, spermatocytes, and spermatids (Thomas et al. 1989). We also screened Northern blots of poly[A] RNA from adult somatic tissues, but the 800-nucleotide transcript has never been identified in any of these samples; it remains a unique feature of the MyK-103 testis. Expression of this aberrant transcript in testis might be enhanced by proximity of the MT-1 promoter in MyK-103 mice and it may contribute to the MyK-103 phenotype.

Male germ-line transmission of rearranged MyK-103 transgenes
We have shown that sperm fertility is recovered when the entire transgene insert is deleted by recombination between the flanking duplicated mouse sequences. Fertile sperm might also be produced if the metallothionein/HSV thymidine kinase fusion genes were deleted...
by recombination between duplicated sequences of pBR322 within the insert. Although hemizygous males never transmit the prototype MyK-103 insert to offspring, we have observed male germ-line transmission of two distinct types of transgenic rearrangements. These rearrangements were distinguished from each other and the prototype MyK-103 insert by their restriction patterns on Southern blots. DNA from hemizygous mice containing the prototype MyK-103 insert produced five bands that hybridized to pBR322 when cut with P_{vul}II, whereas the type 1 rearrangement (ΔMyK) produced a single 6.6-kb restriction fragment (Fig. 7A) and the type 2 rearrangement (Δ + TK) produced two bands of 8.5 and 4.5 kb in length [Fig. 7B].

The structure of the rearrangements inherited by ΔMyK and Δ + TK mice [Fig. 8] was determined by additional Southern blot analysis. For example, the TK probe hybridized to the 8.5- and 5.9-kb bands from MyK-103 mice but only to the 8.5-kb band in Δ + TK [Fig. 7C] and it did not hybridize to ΔMyK (data not shown). It is clear that the 8.5-kb P_{vul}II restriction fragment in Δ + TK corresponds to the right junction fragment spanning J5, J6, and J7 in MyK-103, likewise, the 4.5-kb band, which only hybridizes to pBR322, corresponds to the left junction fragment that spans J2 (Wilkie and Palmiter 1987). To create Δ + TK, and thus delete the 5.9-kb band, homologous recombination probably occurred between the DNA sequences that were in the same relative orientation surrounding the P_{vul}II sites indicated in Figure 8.

The structure of the ΔMyK transgene was determined following additional Southern blot analysis of DNA from a hemizygous mouse compared side by side with DNA from a nontransgenic control and a prototype MyK-103 transgenic mouse. Somatic DNA from each animal was cut with B_{am}HI and E_{co}RI, in addition to P_{vul}II, and hybridized on separate filters with either radiolabeled pBR322 or the restriction fragments BG or BM [see Fig. 1]. pBR322 hybridized to a single restriction fragment generated by digestion with P_{vul}II (6.6 kb, see Fig. 7A), B_{am}HI (5.4 kb), or E_{co}RI (5.3 kb); the BG and BM probes hybridized to the expected fragments [data not shown]. ΔMyK apparently arose by homologous recombination between two copies of pBR322 that were also oriented in the same direction within the nucleotide sequences indicated in Figure 8.

In total, 7 of 44 fertile males in the MyK-103 pedigree transmitted rearranged transgenes. Each rearrangement apparently occurred independently because six of the seven males did not have the same mother [Table 1], and the two G₀ males that did have the same mother both inherited the prototype MyK-103 insert, as demonstrated by the hybridization pattern to somatic DNA [Fig. 7A and data not shown]. This inheritance pattern also suggests that the rearrangements must have occurred de novo in the germ line of G₀ males. In contrast, the rearrangement carried in the Δ + TK line probably occurred in a female during embryogenesis, because both male 35-1 and his brother 35-4 inherited Δ + TK from their mother [Fig. 7B; see Hartl 1971].

Given that ΔMyK and Δ + TK have retained some portion of pMK and the flanking duplicated sequences, it is apparent that the structure of the rearranged transgenes is very similar to the prototype MyK-103 insert. If transmission ratio distortion in MyK-103 males were caused by gene disruption, then ΔMyK, Δ + TK, and MyK-103 would most likely have similar phenotypes due to the overall similarity in structure between these three lines of mice. However, both the ΔMyK and Δ + TK inserts were transmitted through the germ line of hemizygous G₁ males without apparent bias [Table 1]. HSV thymidine kinase protein could not be detected by immunocytochemical staining of ΔMyK [Fig. 5C] or Δ + TK testes [data not shown]. Northern blot analysis corroborated that HSV thymidine kinase was not expressed in ΔMyK testes, and it is interesting that the 800-nucleotide transcript was also not expressed [Fig. 4A, B]. Unfortunately, the Δ + TK line was lost before Northern blots of these transcripts could be assayed. It is unclear why HSV thymidine kinase was not expressed in the testes of Δ + TK males because the MT-1 promoter and HSV thymidine
kinase coding sequences are apparently intact; only the 3' untranslated sequence is truncated in this copy of pMK, but well downstream (190 nucleotides) of the two poly(A) addition sites in the HSV thymidine kinase gene [see Wagner et al. 1981 and Fig. 7B]. Nevertheless, male germ-line transmission of these rearranged transgenes strengthens the conclusion that infertility of transgenic sperm and their mitotic partners is a dominant trait in MyK-103 males that is caused by expression of HSV thymidine kinase, possibly in combination with the 800-nucleotide transcript.

Discussion

Male mice in the MyK-103 pedigree are fertile but do not transmit the prototype transgenic insert to their offspring. This phenomenon occurs because the germ line of every MyK-103 male is a genetic mosaic consisting of both transgenic and nontransgenic cells. The transgenic germ cells that retained the prototype insert expressed HSV thymidine kinase during the haploid phase of spermatogenesis (Figs. 4 and 5), which apparently caused infertility of both transgenic sperm and their nontransgenic mitotic partners. MyK-103 males never sire offspring that inherit the prototype insert, and other lines of transgenic mice that expressed comparable levels of HSV thymidine kinase at the same stage of spermatogenesis are male sterile [Braun et al. 1990]. In contrast to the other transgenic mice, MyK-103 males are fertile because the transgene is deleted in a significant proportion of the spermatogonia stem cells which, following spermatogenesis, give rise to fertile sperm.

Genetic analysis demonstrated that deletion of the transgene occurred by homologous recombination, most often between the duplicated 5-kb sequence of mouse DNA that flanks either side of the insert (Fig. 3). Occasionally, internal rearrangements of the transgene, which were generated by homologous recombination between different fragments of pBR322 sequences in the prototype insert, were also transmitted through the male germ line [see Table 1]. In these cases, either both copies of the HSV thymidine kinase gene present in the prototype MyK-103 insert were deleted [as for the type 1 rearrangement in the ΔMyK line], or the single copy of the HSV thymidine kinase gene that remained in the type 2 rearrangement transmitted by the Δ+TK line [Fig. 8] did not express detectable amounts of protein. We inferred that the type 1 rearrangements occurred in the male germ line because each C0 male inherited the prototype MyK-103 insert from his mother [Fig. 7A]. The germ line in males 33-4, 34-1, 56-1, and 57-1 was apparently a complex mosaic because the rearranged transgene was transmitted at low frequency [Table 1]; most of the off-
Germ-line mosaicism via mitotic recombination

Table 1. Male germ-line transmission of transgene rearrangements

| Male   | Parent | Transgene | Germ-line transmission |
|--------|--------|-----------|------------------------|
| G₀     | 32-3 [ΔMyK] | type 1   | 26/70                  |
|        | 33-4    | type 1   | 2/17                   |
|        | 34-1    | type 1   | 1/11                   |
|        | 37-3    | type 1   | 1/2                    |
|        | 56-1    | type 1   | 1/11                   |
|        | 57-1    | type 1   | 2/10                   |
| G₁     | 91-3 [ΔMyK] | type 1   | 13/26                  |
|        | 35-4 [Δ + TK] | type 2   | 3/7                    |

Seven of 44 males in the MyK-103 pedigree have transmitted rearranged transgenes to their offspring. Six of these males are considered G₀ founder mice of new sublines because they inherited the prototype MyK-103 insert from their mothers but have transmitted type 1 rearrangements to offspring. Male mice in the ΔMyK transgenic line have transmitted a type 1 rearrangement for two generations. The type 2 rearrangement that was transmitted in the Δ + TK line was inherited from female 15-8 by male 35-1 and his sibling 35-4.

Spring from these males probably inherited the meiotic products of germ cells that had fully deleted the MyK-103 insert. However, the majority of germ cells in these mice presumably retained the prototype insert, as evident, for example, in the comparable levels of HSV thymidine kinase RNA in the testes of 33-4 and other MyK-103 males (Fig. 4A). On the basis of the transmission frequency that we observed for the various recombination products, we presume that the germ line in most MyK-103 males is a four-component mosaic, consisting mostly of germ cells that inherit the prototype insert, some that inherit the complete deletion between the flanking repeats, and fewer still that inherit either the type 1 or type 2 rearrangement. The germ line of male 32-3, who transmitted the type 1 rearrangement to half of his offspring (Table 1), is an obvious exception to this general observation, presumably because this rearrangement occurred relatively early in germ cell development.

Due to the limitations of the genetic and immunological assays, we primarily detected recombination events that occurred in embryonic germ cells, spermatogonia, and spermatogonial stem cells; homologous recombination events that might have occurred in germ cells committed to spermatogenesis were probably not detected. For example, if pMK were deleted during meiotic recombination, the resulting sperm would presumably still be infertile due to diffusion of HSV thymidine kinase from syncytial spermatids that developed from nonrecombinant spermatocytes (Fig. 5; see also Braun et al. 1989, 1990). By a similar argument, if deletion of pMK occurred in type A or type B spermatogonia, then only a fraction of the spermatids in a syncytium of this type would have expressed HSV thymidine kinase. As a result of diffusion within the syncytium, occasional patches of elongating spermatids might be recognized which stained very weakly with antibody against HSV thymidine kinase, but this was not observed. Furthermore, we would expect pMK to be transmitted through the male germ line at some low frequency because other hemizygous transgenic mice that expressed low levels of HSV thymidine kinase, as in the H-40 line reported by Braun et al. (1990), were fertile and transmitted the transgene to half of their offspring. However, not one of 636 offspring sired by 44 Myk-103 males has inherited the prototype MyK-103 insert. Immunocytochemistry revealed that almost all of the mosaic testicular tubules in the six hemizygous MyK-103 males that were analyzed had large patches of nontransgenic germ cells. The seminiferous tubules that were entirely devoid of HSV thymidine kinase provided evidence that deletion of pMK could occur during embryogenesis; that portion of the tubule containing only nontransgenic germ cells might have been colonized by nontransgenic spermatogonia. This is the more likely explanation in mice that were sacrificed at a young age, but in older mice it is also possible that several independent deletion events occurred throughout adult life, eventually converting all of the spermatogonial stem cells in a given length of tubule to nontransgenic stem cells.

In MyK-103 testes, the proportion of seminiferous tubules that were mosaic or entirely devoid of HSV thymidine kinase ranged from 5% to 80%, depending on the individual and his age. This high degree of mosaicism is probably required for fertility in MyK-103 males; an independent study showed that the threshold of fertility in male mice is at roughly 10% of the normal capacity for sperm production (Meistrich 1982). Given that fertile sperm only developed in those spermatid syncytia that did not express HSV thymidine kinase, we would expect MyK-103 males to be sterile if germ-line mosaicism was less pronounced. Germ-line mosaicism in MyK-103 males is significant in light of the fact that mosaicism was not detected in any of the sterile hemizygous males from the other transgenic lines that also expressed HSV thymidine kinase in spermatids (Braun et al. 1990). It is of further interest that 11 of the 16 founder males that were studied by Braun et al. (1990) were fertile but did not transmit the HSV thymidine kinase gene to their offspring, the remaining 5 males were each sterile. Immunocytochemical analysis of the fertile founder males revealed the mosaic expression of HSV thymidine kinase among spermatids, similar to the pattern seen in MyK-103 males. The fundamental difference was that these founder males were mosaic due to late integration of the transgene during embryogenesis (Wilkie et al. 1986), whereas mosaicism in hemizygous MyK-103 males was due to deletion of the transgene via homologous recombination.

Duplicated sequences and tandem repeats of genes and repetitive elements are inherently unstable in the genome. Several genetic diseases in humans are caused by deletions due to recombination between reiterated members of a multigene family, as observed in the globin locus (Higgs et al. 1989) and the red and green visual pigments (Nathans et al. 1986). Deletion breakpoints have also been identified between different Alu sequences in the LDL receptor gene (Lehrman et al. 1987), and frequently the entire steroid sulfatase gene is...
deleted by recombination between flanking repeats in patients with ichthyosis (Ballabio et al. 1990; Yen et al. 1990). Although many deletions undoubtedly occur during meiosis, analysis of deletions in the Duchenne muscular dystrophy gene demonstrates that mitotic recombination does occur, resulting in germ-line mosaicism and the transmission of new mutations to offspring [Bakker et al. 1987; Darras and Francke 1987]. Germ-line mosaicism was also found within certain repetitive DNA sequences of the VNTR family (Jeffreys et al. 1990).

A recombination rate for deletion of pMK from the germ line of MyK-103 males was estimated based on the extent of testicular mosaicism. To make this calculation, we assumed a constant recombination frequency per cell generation and that proliferation of transgenic stem cells was not under selection. Based on comparisons between other transgenic mouse lines, there is no evidence to suggest that the HSV thymidine kinase gene imparts any selective pressure on transgenic progenitor cells or stem cells (Wilkie et al. 1986; Heyman et al. 1989, Braun et al. 1990). Indeed, within the MyK-103 pedigree, the HSV thymidine kinase gene appears to be neutral during embryonic germ-line development prior to sexual dimorphism because females show no bias in germ-line transmission of pMK—148 transgenic offspring among 315 born to hemizygous MyK-103 females. Within the male germ line of MyK-103 mice, we estimated a frequency of mitotic intrachromosomal recombination of about 2 X 10^{-3} per cell division. Deletion of the transgene presumably occurred at a similar frequency in somatic cell lineages. Unfortunately, transcription of the HSV thymidine kinase gene is not prolific in somatic tissues of MyK-103 mice (Palmiter et al. 1984), which precludes an analysis of somatic mosaicism by immunocytochemistry. However, we documented one mosaic mouse among 148 transgenic offspring in the MyK-103 pedigree (Palmiter et al. 1984) that probably resulted from deletion of pMK early in embryogenesis (see Kelly et al. 1989). In contrast, somatic reversion at the murine d^ locus, which occurred by intrachromosomal recombination between duplicated 520-bp LTRs, was detected once among 1.1 million mice [Seperack et al. 1988].

Mitotic recombination rates between duplicated sequences similar in length and structure to MyK-103 have been estimated in mouse somatic cell culture to be 2 X 10^{-5} [Liskay et al. 1984] and in yeast to be between 2 X 10^{-5} [Jackson and Fink 1981; Aguilara and Klein 1989] and 2 X 10^{-4} [Holingsworth and Byers 1989]. Thus, the mitotic recombination frequency in the MyK-103 pedigree appears to be significantly greater than observed at other duplicated sequences of similar structure in eukaryotic cells, suggesting that MyK-103 integration locus contains a hotspot for homologous recombination. Most of the MyK-103 integrant and the flanking mouse DNA have been cloned, and it might be possible to reconstruct the pertinent elements of this region that would allow increased recombination frequency in tissue culture cells or transgenic mice. Furthermore, it should be possible to derive cell lines from MyK-103 mice and use the selectable HSV thymidine kinase gene to study recombination frequencies in somatic cells.

Materials and methods

DNA and RNA analysis

DNA manipulations and Southern blots were performed using standard procedures [Maniatis et al. 1982]. Dot hybridization of tail DNA was performed as previously described [Brinster et al. 1985]. RNA used for Northern blot analysis was extracted by homogenization in guanidinium isothiocyanate, followed by precipitation with lithium chloride as described [Cathala et al. 1983]. RNA samples were electrophoresed in agarose–formaldehyde gels, transferred to nitrocellulose paper, and hybridized with radiolabeled probes as described [Peschon et al. 1987]. All radiolabeled hybridization probes used in this study are indicated in Figure 1.

Histology

Histology was done on Carnoy’s fixed tissues, which were embedded in paraffin, then sectioned and stained with hematoxylin and eosin or PAS. Staining for HSV thymidine kinase was done on paraffin-embedded sections using a rabbit anti-HSV thymidine kinase primary antibody and a peroxidase-AEC staining system.

Calculation of the deletion frequency in the MyK-103 germ line

To calculate the deletion frequency of pMK per cell division in the MyK-103 germ line, we chose a representative cross section of the testis from a hemizygous male, 92-1, which was sacrificed at 11 weeks of age and stained for testicular HSV thymidine kinase expression by immunocytochemistry. The sample was photographed at 40 X magnification and a composite of the entire testis cross section was reconstructed [data not shown]. In this section, 320 seminiferous tubules were displayed, 20 of which were mosaic or entirely lacked HSV thymidine kinase expression. The mosaic tubules were randomly distributed in the cross section. On average, 70% of the spermatids in these 20 tubules did not express HSV thymidine kinase. According to our assumptions [see text], the proportion of nonexpressing spermatids is equal to the proportion of spermatogonial stem cells that have deleted the transgene. Therefore, the fraction of recombinant spermatogonial stem cells (As-R) in the total stem cell population is calculated as 0.7[20/320] = 4.4% As-R. Assuming that As divide, on average, once every 8 days [the time required for one cycle of the seminiferous epithelium in mice; Oakberg 1971], then the elongating spermatids that were scored at 11 weeks were descendents of As cells that had undergone approximately nine stem cell divisions. Deletion of the transgene presumably occurs not only in spermatogonial stem cells but also during exponential growth of the primordial germ cells. A newborn mouse has approximately 40,000 spermatogonial stem cells [20,000 per testis; de Rooij 1983], which would be generated in 16 synchronous divisions of exponential growth. Neonatal germ cell death has not been assayed in mouse but, in rats, the embryonic testis contains about twice the number of precursor cells as later develop into spermatogonial stem cells [Beaumont and Mandi 1963].

The frequency of pMK deletion can be stated as the probability of a daughter cell inheriting a recombinant chromosome sometime during embryonic germ cell proliferation and spermatogonial stem cell renewal. The probability is calculated...
from the series

$$P_k = \sum_{i=1}^{k} \mu(1 - \mu)^{k-1}$$

where $P_k$ is the probability that a cell is nontransgenic after $k$ divisions, and $\mu$ is the deletion rate per cell division. If $P_k$ is small, the deletion frequency is approximated as the percentage of nontransgenic spermatogonia divided by the number of generations in the stem cell lineage at the time the animal was sacrificed. The deletion frequency estimated from male 92-1, whose germ-line was 4.4% mosaic when sacrificed, was estimated to be $0.044/(16-1-9)$, or 2.5 x $10^{-3}$ per cell division. A similar calculation was made for male 33-4, which had roughly 25% nontransgenic germ cells at 87 weeks of age, giving a frequency of 2.5 x $10^{-3}$ per cell division.

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