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A high proportion of genes involved in position effect variegation also affect chromosome inheritance

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Abstract

Suppressors and enhancers of position effect variegation (PEV) have been linked to the establishment and maintenance of heterochromatin. The presence of centromeres and other inheritance elements in hetero-chromatic regions suggests that suppressors and enhancers of PEV, *Su(var)*s and *E(var)*s [collectively termed *Mod(var)*s], may be required for chromosome inheritance. In order to test this hypothesis, we screened 59 ethyl methanesulfonate-generated *Drosophila Mod (var)*s for dominant effects on the partially compromised inheritance of a minichromosome (*J21A*) missing a portion of the genetically defined centromere. Nearly half of these *Mod(var)*s significantly increased or decreased the transmission of *J21A*. Analyses of homozygous mutant larval neuroblasts suggest that these mutations affect cell cycle progression and native chromosome morphology. Five out of six complementation groups tested displayed mitotic abnormalities, including phenotypes such as telomere fusions, overcondensed chromosomes, and low mitotic index. We conclude that *Mod (var)*s as a group are highly enriched for genes that encode essential inheritance functions. We propose that a primary function of *Mod(var)*s is to promote chromosome inheritance, and that the gene silencing phenotype associated with PEV may be a secondary consequence of the heterochromatic structures required to carry out these functions.

Introduction

The ability to distribute chromosomes properly during cell division is essential for the viability of all organisms. At the heart of the segregation process is the centromere, which serves as the nucleation site for the kinetochore (for review see Sullivan et al. 2001), where microtubules attach and help move sister chromatids or homologs to opposite poles during mitosis and meiosis. The centromere is typically embedded in heterochromatin, the portion of the genome rich in repeated sequences and poor in genes (for review see Hennig 1999). Heterochromatin is involved in other critical inheritance functions, including meiotic pairing (Dernburg et al. 1996; Karpen et al. 1996), sister chromatid cohesion (Bickel and Orr-Weaver 1996; Dej and Orr-Weaver 2000; Bernard et al. 2001; Hall et al. 2003) and interactions with the NOD kinesin-like protein (Afshar et al. 1995; Murphy and Karpen 1995a). Dissecting the roles of heterochromatin in centromere function and other inheritance

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processes will greatly increase our understanding of the *cis* and *trans* components of chromosome inheritance.

Significant information about heterochromatin has come from studying the phenomenon of position effect variegation (PEV; for review see Wallrath 1998). Position effect variegation refers to the clonal inactivation of gene expression that occurs when a gene is removed from its normal chromatin context (e.g. euchromatin) and is placed in an opposing chromatin context (e.g. heterochromatin) via chromosomal rearrangement or transgene insertion. A classic example of PEV in *Drosophila* is the expression of the *white* eye color gene in the chromosomal inversion *In(1)w^{m4}* (Muller 1930), in which the euchromatic *white* gene is placed immediately adjacent to X chromosome centric heterochromatin. The proximity to heterochromatin causes mosaic *white* repression, resulting in a mottled eye color phenotype. Multiple genetic screens for modifiers of PEV [*Mod(var)s*] have identified a large number of mutations that dominantly enhance [*E(var)s*] or suppress [*Su(var)s*] heterochromatin-mediated gene silencing (Reuter and Wolff 1981; Locke et al. 1988; Dorn et al. 1993).

In *Schizosaccharomyces pombe*, mutations that alleviate centromere-mediated silencing also induce elevated rates of minichromosome loss and missegregation (Allshire et al. 1995; Grewal et al. 1998; Freeman-Cook et al. 1999). In *Drosophila melanogaster*, mutations in the *Su(var)2–5* locus, which encodes Heterochromatin Protein 1 (HP1), exhibit recessive mitotic defects, including telomere fusions (Fanti et al. 1998) and chromosome undercondensation (Kellum and Alberts 1995). Other *Mod(var)* mutations — specifically *Su(var)2–8*, *Su(var)2–10* and *Su(var)3–6* — cause chromosome loss, undercondensation and overcondensation, respectively (Axton et al. 1990; Wines and Henikoff 1992; Hari et al. 2001). Mammalian HP1 can be delocalized as a consequence of prolonged exposure to deacetylase inhibitors or by mutations in the histone H3 methyltransferase SUV39, and this is accompanied by defects in chromosome segregation (Rea et al. 2000; Lachner et al. 2001; Taddei et al. 2001). Chromosome segregation defects associated with mutations in the *S. pombe* HP1 homolog, Swi6, appear to be caused by defects in cohesin recruitment to centromeric regions (Bernard et al. 2001; Nonaka et al. 2002). In *Drosophila*, a recent study has demonstrated that some *Mod(var)* mutations affect meiotic recombination (Westphal and Reuter 2002). These results indicate that *Mod(var)* mutations can impact chromosome inheritance and behavior; however, no comprehensive study of the inheritance roles of *Mod(var)* genes in metazoans has been reported.

The fully functional 1.3 Mb *Drosophila* minichromosome *Dp(1;f)1187* (*Dp1187*) has been a useful substrate for studies of centromere structure and function and the molecular biology of heterochromatin (Le et al. 1995; Murphy and Karpen 1995b; Karpen et al. 1996; Sun et al. 1997, 2003). Previous studies using this minichromosome and its radiation-induced deletion derivatives localized the functional centromere to a 420 kb region within the heterochromatin of *Dp1187*. Derivatives of *Dp1187* with deletions of the centromere region show reduced transmission. The derivative *J21A* is missing approximately one-third of the genetically defined centromere, and is transmitted to progeny half as well as intact minichromosomes (Sun et al. 1997). Its intermediate level of transmission makes it an ideal candidate to look for dominant interactions. Indeed, known inheritance mutations dominantly enhance or suppress the transmission of *J21A* and other ‘sensitized’ minichromosomes (Murphy and Karpen 1995a; Cook et al. 1997). Seventy-eight *P* element-induced mutations that dominantly affect minichromosome inheritance were isolated by screening for increased or decreased transmission of *J21A* (Dobie et al. 2001). *P* element insertions were recovered in genes previously characterized to play roles in inheritance and many of the novel homozygous lethal mutations affected the function of the normal *Drosophila* chromosome complement, further substantiating a role for these mutations in inheritance.

Here, we present evidence that suggests that a high proportion of *Mod(var)*s are required for normal chromosome inheritance. We describe the results of a genetic screen in which 59 *Mod(var)* mutations, representing 49 complementation groups, were tested for dominant effects on the transmission of *J21A*. Nearly half of the *Mod(var)* mutations dominantly increased or decreased *J21A* transmission to progeny. Cytological analysis demonstrated that endogenous *Drosophila* chromosomes display visible mitotic chromosome defects in homozygous lethal mutants. We conclude that a large percentage of modifiers of PEV have essential, diverse roles in chromosome inheritance. Our findings suggest that a primary function of heterochromatin is to promote chromosome inheritance, and that some properties of heterochromatin, such as gene silencing, may be the consequence of the DNA and protein structures required to ensure proper inheritance of chromosomes from parent to progeny.

Materials and methods

Fly stocks and crosses

J21A is the result of a terminal deletion that removed approximately one-third of the genetically defined centromere and the rest of the flanking heterochromatin from an inversion derivative (*Dp238*) of *Dp(1;f)1187*. It is 580 kb in size. *J21A* and modifiers of terminal deficiency associated PEV [*Mod(TDA-PEV)*] are described in more detail elsewhere (Murphy and Karpen 1995b; Donaldson et al. 2002). Previously identified *Su(var)*s (Reuter and Wolff 1981; Sinclair et al. 1992; Eissenberg et al. 1992) were obtained from B. Wakimoto, G. Reuter, and J. Eissenberg. Balancer-GFP (green fluorescent protein) stocks came from the Bloomington Stock Center. All mutations were crossed into a standard *y^l; ry⁵⁰⁶* background prior to use.

J21A monosome transmission assay

J21A is a *Dp1187* minichromosome deletion derivative marked with two insertions of a *ry⁺*-marked *P* element construct (Murphy and Karpen 1995b). Transmission tests for dominant effects on minichromosome inheritance were performed as described previously (Murphy and Karpen 1995a; Cook et al. 1997). Single, heterozygous mutant females bearing one copy of *J21A* were crossed to several *C(1;Y)1, y^l; ry⁵⁰⁶ (X⁺Y)* males; transmission frequency was determined by counting *ry⁺* (*J21A* present) and *ry⁻* (*J21A* absent) female progeny. The attached X–Y (*X⁺Y*) is utilized to produce *X / X⁺Y* female progeny, such that the presence of an extra *Y* suppresses the partial silencing of the *ry⁺* marker. Only vials containing at least 30 progeny were included in this analysis. Standard Student's *t*-tests were used to compare mutant results with *y^l; ry⁵⁰⁶* controls; *P* values of ≤ 0.05 were considered statistically significant.

Cytological analysis

Females and males of the genotype mutation/GFP-marked balancer were crossed to produce homozygous mutant, non-GFP larvae for cytological analysis of neuroblast mitoses. The oldest developing homozygous mutant larvae produced from each line were subsequently selected for dissection. Larval brains were dissected in 0.7% NaCl, fixed with 45% acetic acid for 30–60 s, squashed onto slides, and frozen in liquid nitrogen. After coverslip removal, chromosomes were stained with 1 μ g/ml 4',6-diamidino-2-phenylindole and scored on a Zeiss Axiophot Epifluorescence Microscope. Images were captured with a cooled CCD camera (Princeton Instruments) and IP Lab Spectrum software (Scanalytics). In addition to the above method, a subset of neuroblasts from some lines were also prepared with a 5 min soak in 0.8% sodium citrate after the dissection. The hypotonic swell is known to cause artificial chromatid arm separation (Gatti et al. 1994) and this phenotype was ignored in these neuroblasts. Mitotic figures from mutant larvae were compared with *y^l; ry⁵⁰⁶* control larvae. Since early instar larvae have fewer mitotic figures than later stage larvae, random

sampling can lead to sample bias. Therefore, the entire brain was analyzed for mitotic and anaphase figures. The mitotic index here is defined as the total number of mitotic and anaphase figures divided by an approximation of the total number of cells (calculated by multiplying the average number of cells from five randomly selected fields by the total number of fields scored). Standard Student's t-tests were used to determine the significance of differences between mutants and controls. The fields were scored at 100× magnification with a 1.25× optivar.

Results

A high proportion of previously identified *Su(var)*s dominantly affect *J21A* transmission

The minichromosome deletion derivative *J21A* is transmitted to only 28% of progeny as a monosome, compared with 50% transmission for a full-length *Dp1187* minichromosome (Murphy and Karpen 1995b; Cook et al. 1997). The partial function of *J21A* makes it an excellent substrate for identifying genes that play a role in chromosome inheritance, since a reduced level of a protein required for inheritance may not affect the normal complement of chromosomes, but may affect *J21A* transmission. Consequently, potential inheritance loci can be identified by observing dominant effects of heterozygous mutations on *J21A* transmission. This approach also circumvents the recessive lethality associated with many loss-of-function mutations in inheritance genes. Known inheritance mutations dominantly affect *J21A* transmission, but do not dominantly alter the transmission of minichromosomes with the entire genetically defined centromere, or the normal complement of chromosomes, demonstrating the sensitivity of *J21A* (Murphy and Karpen 1995a; Cook et al. 1997). The *J21A* assay has also been used successfully to identify new chromosome inheritance genes (Dobie et al. 2001).

We tested nine previously identified *Su(var)* mutations representing five different loci (Reuter and Wolff 1981; Wustmann et al. 1989; Eissenberg et al. 1992) for dominant effects on *J21A* minichromosome transmission (see Materials and methods). We also tested the effect of an extra *Y* chromosome, known to be a strong suppressor of many types of PEV (Spofford 1976). Three of the nine *Su(var)* mutations and the extra *Y* chromosome reduced transmission significantly ($P \leq 0.05$, Table 1), while one mutation increased transmission. *Su(var)205¹* and the extra *Y* chromosome displayed the most dramatic reductions of *J21A* transmission, to 5% and 8%, respectively. *Su(var)2–10* reduced transmission to 15%, and in-depth analysis demonstrated that *Su(var)2–10* affects chromosome condensation and nuclear organization (Hari et al. 2001). Thus, three out of five loci tested had at least one allele that altered *J21A* transmission significantly, suggesting a general role for *Su(var)*s in chromosome inheritance.

Nearly half of the *Mod(var)*s induced in an isogenic background significantly affect *J21A* transmission

Data from screening the previously identified *Su(var)*s suggested that this class of mutations would provide a rich source of genes involved in chromosome transmission. However, genetic background in the form of additional mutations induced on the parental chromosomes or accumulated in the balanced stocks could be responsible for the transmission phenotype. A large group of ethyl methanesulfonate (EMS)-induced modifiers of TDA-PEV (Donaldson and Karpen 1997; Donaldson et al. 2002) was generated in the same isogenic *y; ry* genetic background used for previous *J21A* transmission screens (Cook et al. 1997). Most of the mutations affected multiple examples of PEV (Donaldson et al. 2002). These mutations were used to test the generality of *Mod(var)* effects on inheritance.

Fifty modifiers of TDA-PEV, representing maximally 45 complementation groups (Donaldson et al. 2002), were tested for dominant effects on *J21A* transmission (Table 2). The original *y^l; ry⁵⁰⁶* stock used for the EMS screen was used as a control, and showed 27% transmission, as expected from previous studies. Twenty-three of the 50 (46%) lines tested altered *J21A* transmission significantly. A majority of the modifiers analyzed were *Su(var)s*. In total, 20 out of 45 *Su(var)* lines significantly altered *J21A* transmission. Seventeen of the 20 lines reduced transmission to 7–21%, representing decreases of 22–74% from the control transmission value of 27%. Three lines increased transmission to 37–41%, showing an increase of at least 50% from the control. Five enhancers of TDA-PEV were also tested. Three of five enhancers affected *J21A* transmission significantly. One line (*E1060*) reduced transmission weakly to 20%, while two lines (*E1178* and *E1377*) increased transmission to 39–40%. In total, 21 of 45 complementation groups tested (47%) altered *J21A* transmission significantly.

Combining the results from the analyses of previously identified *Su(var)s* and the isogenic collection of modifiers of TDA-PEV, 27 of 59 *Mod(var)* mutations affected *J21A* transmission significantly (Table 3). Twenty-four of the 54 *Su(var)s* significantly altered *J21A* transmission; the vast majority of these mutations (20/23, or 87%) decreased transmission. Of the three *E(var)* lines that altered *J21A* transmission, two increased transmission while the other decreased it. One complementation group [*Su(var)2–5*] is represented by both previously identified and newly induced *Mod(var)* mutations, thus lowering the total number of loci that affect *J21A* transmission to 23 of 49. These results show that a large proportion of *Mod(var)s* (23/49 loci, 47%) affect the inheritance of the *J21A* ‘sensitized’ minichromosome.

Mitotic defects are observed in homozygous lethal *Mod(var)* mutants

In previous studies, mutations in inheritance genes did not have dominant effects on the transmission of minichromosomes containing the entire genetically defined centromere (Cook et al. 1997); therefore, the *Mod(var)s* that have dominant effects on *J21A* transmission are unlikely to have dominant effects on the transmission of the normal complement of chromosomes. Nevertheless, the *Mod(var)s* may have recessive effects on the inheritance of normal, full-length chromosomes, which could lead to aneuploidy and lethality. Therefore, we examined larval neuroblasts from nine homozygous lethal *Mod(var)* mutations (six complementation groups) that altered *J21A* transmission for cytological defects in chromosome behavior.

Homozygous mutants for many of the lines had either small brains and/or diminutive or missing imaginal discs, phenotypes previously demonstrated to correlate with mitotic defects (Gatti and Baker 1989). Larval brains from lines 234, 1097, 1227, 1545, and 1552 displayed significantly lower frequencies of cells in metaphase and anaphase, in comparison with *y^l; ry⁵⁰⁶* controls and line 1259, which is homozygous viable and had no effect on *J21A* transmission (Table 4). The low mitotic indices may be due to cell cycle arrest in interphase, a prolonged interphase, or decreased length of mitosis. The rare metaphase nuclei in these lines had no obvious chromosomal abnormalities.

Mitotic chromosomes from two other complementation groups displayed striking cytological defects (Fig. 1). Telomere fusions were seen in second and third instar larval neuroblasts from *Su(var)1207*, including interchromosomal fusions of two or more chromosomes to form large rings, intrachromosomal fusions to form small rings, and multiple chromosomes fused to form linear ‘trains’. Such fusions have been observed in homozygous mutant larvae for *Su(var)2–5* (Fanti et al. 1998), and for *UbcD* (also known as *effete*, Cenci et al. 1997). 1207 and 1009 fail to complement *Su(var)2–5⁰²* for viability and contain nonsense mutations in the *Su(var)2–5* open reading frame, indicating that they are

new 2–5 alleles (Donaldson et al. 2002). Analysis of third instar larval brains from homozygotes of the *E(var)* line *E1060* revealed that mitotic chromosomes were substantially shorter and more compact than control chromosomes (Fig. 1). This overcondensed chromosome phenotype correlates well with the ‘tighter’ chromatin structure of genes silenced by PEV (Wallrath and Elgin 1995), which is likely further condensed by *E(var)* mutations.

Line *1126*, which decreased *J21A* transmission to 12% and is homozygous lethal, displayed a normal mitotic index and no obvious cytological defects. This mutation may not affect the activity of the gene enough to cause abnormal behavior of full-length chromosomes, or the defect may be too subtle to observe in cytological analysis.

In summary, cytological examination of homozygous lethal lines that disrupted *J21A* transmission revealed that mutations in five out of six complementation groups caused visible chromosome defects represented by three different phenotypes: low mitotic index, telomere fusions, and overcondensed chromosomes. These results demonstrate that mutations identified through dominant effects on *J21A* transmission also have recessive effects on the inheritance of normal *Drosophila* chromosomes.

Discussion

Here, we have analyzed 59 *Mod(var)* mutations for dominant effects on chromosome inheritance, and discovered that 47% of the loci tested had at least one allele that significantly affected the transmission of the *J21A* ‘sensitized’ minichromosome. The demonstration that a particular locus or protein is responsible for the transmission phenotype must await cloning of the gene and phenotypic rescue by *P* element constructs. However, the general conclusion that *Mod(var)*s are highly enriched for genes involved in chromosome inheritance is sound. Strong support for this comes from the observation that only 2.6% of approximately 3000 randomly selected *P* element insertions affected *J21A* transmission (Dobie et al. 2001); thus, *Mod(var)*s are enriched, perhaps as much as 20-fold, for factors involved in inheritance. In addition to pulling out known inheritance genes, cytological characterization of the Dobie et al. mutants revealed that they had mitotic defects, which further substantiates *J21A* as a legitimate tool for identifying chromosome inheritance factors. Finally, rescue experiments with the first of these loci to be cloned, *Su(var)2–10/dPIAS*, demonstrated that the dominant effect on *J21A* transmission and condensation defects map to that locus (Hari et al. 2001).

The 59 mutations investigated here correspond maximally to 49 loci, based on lethal complementation analysis (Donaldson et al. 2002). This is likely to be an overestimate of the number of loci, since many mutations are not homozygous lethal and cannot be definitively tested for complementation. On the other hand, we may be underestimating the proportion of *Mod(var)*s with effects on inheritance, since some mutations in inheritance genes may not reduce or alter protein function enough to observe dominant effects on *J21A* transmission. For example, the *Su(var)3–9* protein (Schotta et al. 2002) is homologous to the histone H3 K9 methyltransferases that play key roles in heterochromatin structure and function (for review see Jenuwein and Allis 2001). Line *1699* contains a mutation in *Su(var)3–9* and is the strongest *Su(var)* isolated in our screen (Donaldson et al. 2002), yet only reduced *J21A* transmission to 22%. It is likely that the absence of significant effects on *J21A* transmission result from redundancy in 3–9 function; *1699* and all other 3–9 alleles are homozygous viable. Nonetheless, our results demonstrate that proteins affecting heterochromatin function in *Drosophila* play a more important role in inheritance than previously appreciated.

Cytological analyses revealed that neuroblasts from homozygous mutant larvae displayed visible mitotic defects in five out of six *Mod(var)* complementation groups tested — specifically telomere fusions, overcondensation of full-length chromosomes and low mitotic indices. These results suggest that these *Mod(var)*s are bona fide chromosome inheritance mutations, and that their effects are not specific to *J21A*.

Several of the *Mod(var)* lines (234, 1097, 1227, 1545, 1552) had low mitotic indices, suggesting that the genes may affect cell cycle progression directly or indirectly. Defective chromosome inheritance may be the result of altered cell cycle timing or inactivation of a cell cycle checkpoint (for review see Clarke and Gimenez-Abian 2000). If mitosis is abbreviated, there may not be enough time for the functionally impaired *J21A* to assemble the required segregation machinery and attach securely to the spindle, although normal centromeres would be able to. In a similar fashion, if the spindle attachment checkpoint is defective (for review see Burke 2000), mitosis may not be delayed enough to allow time for the *J21A* to compensate for its partial inheritance defects. Further analysis is necessary to determine exactly how these mutations affect cell cycle progression or timing, and the roles that heterochromatin proteins play in these functions.

The telomere fusions observed in homozygotes of the *Su(var)2–5* mutant alleles 1009 and 1207 could cause chromosome loss. Telomeres are thought to play a role in nuclear organization (Gasser 2001; Hari et al. 2001) and are important for chromosome inheritance in many organisms (Nimmo et al. 1998; Cooper 2000). Fusions of telomeres can result in dicentric chromosomes, which are broken during anaphase (McClintock 1938). No abnormal chromosomal fragments were seen in anaphases from 1009 and 1207 homozygotes, suggesting that these telomere fusions are weak enough to be resolved, as observed previously for other HP1 alleles (Fanti et al. 1998). However, the fact that the telomeres undergo end-to-end fusions indicates that the telomeres are not functioning properly, and that defects in telomere function may cause lethality through mechanisms other than chromosome breakage. For example, *Su(var)2–10/dPIAS* mutations affect telomere-telomere clustering and telomere-lamina interactions, which may be responsible for the undercondensation and gene expression phenotypes (Hari et al. 2001).

We also observed that some *Mod(var)*s affect general chromosome structure. The overcondensation of whole chromosomes observed in homozygous *E1060* mutants is consistent with the enhancement of PEV seen in *E1060* heterozygotes. Gene silencing of euchromatic genes placed in or near heterochromatin may be due to the genes adopting a ‘closed’ or compacted, heterochromatin-like structure, and *E(var)* loci may promote further such ‘heterochromatinization’. *Su(var)2–10/dPIAS*, which also affects *J21A* transmission (Hari et al. 2001 and Table 1), has the opposite effect: homozygous mutant neuroblasts display undercondensed chromosomes, which also supports a correlation between effects on gene expression and chromosome condensation. However, there was not a strict correlation between the direction of the effects on PEV and *J21A* transmission; both *E(var)*s and *Su(var)*s were observed to cause decreases and increases in *J21A* transmission. This is not surprising, since in most cases we do not know what protein product and processes are affected. In some cases, effects on gene expression could be regulated differently from effects on inheritance.

The results presented here demonstrate that modifiers of PEV are highly enriched for factors that affect the inheritance of a minichromosome. Cytological evidence along with previously published *Mod(var)* data suggest that inheritance of the normal complement of chromosomes is also compromised. This further bolsters the idea that these genes play a crucial role in a variety of inheritance functions in a metazoan, as previously demonstrated for *S. pombe* (Allshire et al. 1995; Bernard et al. 2001). It is quite likely that other *Mod(var)*s

with effects on chromosome inheritance will disrupt other chromosomal processes, such as meiotic pairing (Dernburg et al. 1996; Karpen et al. 1996) and NOD-mediated antipoleward forces (Murphy and Karpen 1995a). Indeed, a study has recently been published demonstrating effects of *Mod(var)*s on meiotic recombination (Westphal and Reuter 2002). While some *Mod(var)* loci will likely encode protein components of heterochromatin, others will undoubtedly have indirect effects on heterochromatin function and chromosome transmission. For example, some *Mod(var)* mutations may alter the expression, function or localization of chromosome inheritance proteins. Still others may be involved in general aspects of nuclear organization and function critical for normal inheritance, as recently demonstrated for *Su(var)2–10* (Hari et al. 2001). The unusually high percentage of *Mod(var)*s that affect inheritance intriguingly suggests that the primary function of heterochromatin is to regulate chromosome inheritance and nuclear organization by determining chromosome structure, and that PEV is a secondary consequence of the chromatin structures required to carry out these functions. Further characterization of these *Mod(var)*s will provide important insights into inheritance mechanisms that involve heterochromatin.

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Fig. 1.

A–D Metaphase figures from homozygous lethal mutants demonstrate effects on chromosome behavior and morphology. **A** Wild type. **B, B'** Chromosome fusion phenotypes associated with line *1009*. Note chromosome ‘trains’ (**B**) and ‘rings’ (**B'**). **C, C'** Chromosome fusion phenotypes associated with line *1207*. *Arrows* point to telomere fusions. **D, D'** Chromosome overcondensation phenotype associated with line *E1060*. *Bar* represents 2 μ m

Table 1

The effects of previously identified *Su(var)s* on *J21A* transmission. (N, number of vials with >30 progeny)

Gene	<u><i>J21A</i> transmission</u>		<i>P</i> value
	(%±SD)	N	
<i>Su(var)205^l</i>	5±3	13	<0.01
Extra <i>Y</i>	8±5	10	<0.01
<i>Su(var)2-10²</i>	15±10	20	<0.01
<i>Su(var)2-1^l</i>	14±13	16	<0.01
<i>Su(var)2-10^l</i>	20±14	14	0.12
<i>Su(var)2-57^l</i>	23±8	16	0.13
<i>Su(var)2-1²⁻¹³</i>	23±12	9	0.43
<i>Su(var)2-48^l</i>	25±14	15	0.62
<i>Su(var)2-1⁹</i>	29±13	17	0.47
<i>Su(var)2-5⁵</i>	34±8	14	0.01

Table 2

Results of screening modifiers of position effect variegation (PEV) in an isogenic background for *J21A* transmission effects. (E, Enhancer of variegation; N, number of vials with >30 progeny; S, Suppressor of variegation)

Line	<i>J21A</i> transmission		<i>P</i> value
	(%±SD)	N	
Decreased transmission (18)			
1038	7±4	14	<0.01
224	10±6	14	<0.01
1116	11±9	14	<0.01
1126	12±7	7	<0.01
1025	13±6	14	<0.01
1009 ^a	13±8	12	<0.01
1474	14±8	12	<0.01
1453	14±6	13	<0.01
1447	14±8	11	<0.01
1539	16±10	13	<0.01
1315	16±9	10	<0.01
234 ^c	16±10	14	<0.01
1260	17±6	9	<0.01
600	17±8	13	<0.01
1207 ^a	19±7	15	<0.01
E1060 ^b	20±5	10	<0.01
1200	19±7	11	0.01
242	21±7	15	0.02
Increased transmission (5)			
1094	37±8	11	<0.01
1173	39±9	9	<0.01
1144	41±12	19	<0.01
E1178 ^b	40±16	15	0.01
E1377	39±13	9	0.03
No effect (27)			
25S, 2E Control	25±10	322	
y ^l ;ry ⁵⁰⁶	27±7	19	

Complementation groups:

^a 1227 and 1545 (no effect), 1009 and 1207

^b E1060, E1178

^c 234, 1551 (no effect)

Table 3

Summary of effects of *Su(var)* and *E(var)* mutations on *J21A* transmission

	Lines tested		Complementation groups tested	
	Altered transmission	Total	Altered transmission	Total
Previously isolated	4	9	3	5
<i>Su(var)</i> s				60
New <i>Su(var)</i> s	20	45	19	41
New <i>E(var)</i> s	3	5	2	4
Total	27	59	24(23) ^a	50(49) ^a

^a Alleles of *Su(var)2-5* are present among the previously isolated and new *Su(var)*s. Therefore, the total of 24/50 is reduced to 23/49

Table 4

Homozygous lethal mutations affecting *J21A* transmission display reduced mitotic index. (N, number of brains scored. See Materials and methods for how mitotic index was calculated)

Line	Mitotic Index ($\times 10^{-4}$)	N	P value	Larval stage
<i>y^l;ry⁵⁰⁶</i>	2 $\pm$ 2	7	N/A	1st
<i>y^l;ry⁵⁰⁶</i>	30 $\pm$ 10	7	N/A	2nd
234	6 $\pm$ 4	5	0.03	2nd
1097	3 $\pm$ 2	5	0.01	2nd
1227	4 $\pm$ 4	5	0.03	2nd
1545	0	6	0.03	1st
1552	5 $\pm$ 3	5	0.01	2nd
1259	40 $\pm$ 40	5	0.23	2nd
1126	3 $\pm$ 3	4	0.67	1st