Multiple recent sex chromosome fusions in *Drosophila virilis* associated with elevated satellite DNA abundance

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ABSTRACT

Repetitive satellite DNA is highly variable both within and between species, and is often located near centromeres. However, the abundance or array length of satellite DNA may be constrained or have maximum limits. *Drosophila virilis* contains among the highest relative satellite abundances, with almost half its genome composed of three related 7 bp satellites. We discovered a strain of *D. virilis* that has 15% more pericentromeric satellite DNA compared to other strains, and also underwent two independent centromere-to-centromere sex chromosome fusion events. These fusions are presumably caused by DNA breakage near the pericentromeric satellites followed by repair using similar repetitive regions of nonhomologous chromosomes. We hypothesized that excess satellite DNA might increase the risk of DNA breaks and genome instability when stressed, which would be consistent with the apparent high rate of fusions we found in this strain. To directly quantify DNA breakage levels between strains with different satellite DNA abundances, we performed the comet assay after feeding flies gemcitabine and administering low-dose gamma radiation. We found a positive correlation between the rate of DNA breakage and satellite DNA abundance. This was further supported by a significant decrease in DNA breakage in an otherwise genetically identical substrain that lost the chromosome fusion and several megabases of satellite DNA. We find that the centromere-to-centromere fusions resulted in up to a 21% nondisjunction rate between the X and Y chromosomes in males, adding a fitness cost. Finally, we propose a model consistent with our
data that implicates genome instability as a critical evolutionary constraint to satellite abundance.
INTRODUCTION

Satellite DNA consists of long arrays of tandemly repeated sequences, and is often located near centromeres in heterochromatin (reviewed in Thakur et al. 2021). Satellite DNA varies greatly in sequence and abundance within and between species (Subirana et al. 2015; Wei et al. 2018; Cechova et al. 2019), and this can be partially explained by high rates of copy number mutation (Flynn et al. 2017). Although previously assumed to be inert “junk,” recent work has shown that satellite DNA is involved in essential processes in the cell, thus variation in it may be biologically important (Jagannathan et al. 2018; Mills et al. 2019). Although satellite DNA differences between some species have been linked to reproductive incompatibilities (Ferree and Barbash 2009; Jagannathan and Yamashita 2021), the biological implications of intraspecies abundance variation has not been explored. Satellite DNA can vary in abundance by several megabases among individuals of the same species, including in flies and humans (Miga et al. 2014; Wei et al. 2014; Flynn et al. 2020). Satellite DNA appears to be constrained by maximum limits, with no species studied so far having more than about half of their genome made up of satellite DNA (Gall and Atherton 1974a; Fry and Salser 1977; Petitpierre et al. 1995). The genomic abundance of transposable elements (TEs), the other highly pervasive type of repetitive DNA, seems to be less constrained than satellite DNA with many genomes over 50% and some containing up to 85% TE content (Anderson et al. 2019). The nature of satellite DNA with long tandem arrays of the same sequence, may impose instability that prevents it from expanding beyond a threshold, compared to more diverse sequences interspersed in the genome.

Drosophila virilis is an excellent model for studying satellite DNA variation. D. virilis has the highest relative abundance of simple satellite DNA (defined as satellites with unit length <=20 bp) compared to any other studied species. Three 7 bp satellites, AAACTAC, AA ACTAT, and AAATTAC take up over 40% of the genome in D. virilis, and they form arrays tens of megabases long in the pericentromeric region (Gall et al. 1971; Gall and Atherton 1974; Flynn et al. 2020). The extremely high relative abundance of satellite DNA in D. virilis makes it an ideal system in which to ask whether there are constraints or maximum limits on satellite abundance. One
strain in particular, vir00 (15010-1051.00), contains 15% more pericentromeric satellite DNA compared to other *D. virilis* strains (Flynn *et al.* 2020 Figure 4B). In past modeling efforts, satellite DNA arrays have been proposed to be weakly deleterious until they reach a maximum length beyond which they are not tolerated by selection (Charlesworth *et al.* 1986). Slow DNA replication or development time have been suggested as mechanisms to enforce strong negative selection against long satellite arrays, however empirical evidence for this has been limited (but see Bilinski *et al.* 2018). Here, we propose genome instability as a constraint on excessively long or abundant satellite DNA arrays.

Genome instability is characterized by DNA damage that often results in large-scale mutations or rearrangements, and is a fundamental driver of karyotype evolution, chromosomal disorders, and cancer rearrangements (Black and Giunta 2018; Mayrose and Lysak 2021). The first step is spontaneous DNA damage such as double-stranded DNA breaks, one of the most dire events to occur in a cell (Featherstone and Jackson 1999). Even if DNA breaks are repaired, they often result in large-scale genome rearrangements. Most DNA breaks leading to rearrangements occur near the centromere and within or near satellite DNA, including in human genomes (Black and Giunta 2018; Balzano *et al.* 2020). This may be due to intrinsic instability of satellite DNA arrays caused by replication stress of polymerases progressing through highly repetitive sequences, or the formation of unstable DNA topology (Barra and Fachinetti 2018). Robertsonian translocations, one of the most common rearrangements in medical genetics and evolution, occur when there are breaks near the centromere of acrocentric chromosomes and when they are repaired they are fused to each other (Mayrose and Lysak 2021). Robertsonian translocations are associated with multiple miscarriages in humans and aneuploidic disorders like Patau and Down syndromes (Braekeleer and Dao 1990), with increased rates of aneuploidy driven by increased nondisjunction of Robertsonian or fused chromosomes (Schulz *et al.* 2006). We hypothesize that variation in pericentromeric satellite DNA abundance influences the risk of genome instability events. Specifically, excess satellite DNA might increase the risk of genome instability and genome rearrangements.
Sex chromosome evolution has been studied for decades, mainly making use of sex chromosomes that have arisen in different time periods (Charlesworth and Charlesworth 2000). In Drosophila, when an autosome fuses to either an X or Y chromosome, a so-called neo-Y chromosome is formed. Because either the fused or unfused version of the chromosome will only be present in males and male Drosophila do not undergo recombination, mutations immediately begin to accumulate through Hill-Robertson interference and other linked-selection processes (Charlesworth and Charlesworth 2000). In the genus Drosophila, autosomes have fused to sex chromosomes multiple independent times (Nozawa et al. 2021): D. pseudoobscura (10 million years), D. miranda (1 million years; Bachtrog 2013), D. albomicans (0.24 million years; Wei and Bachtrog), and D. americana (29 thousand years; Vieira et al. 2006). Neo-sex chromosomes formed by de novo sex chromosome fusions are rare and actually under-represented compared to autosomal fusions in Drosophila (Anderson et al. 2020), and have never been discovered at their infancy before detectable divergence has occurred. How new sex chromosome fusions become stable, and how they compete with the ancestral karyotype within a species is unknown.

Here, we describe the discovery of two independent and extremely recent sex chromosome-autosome fusions in one D. virilis strain, vir00. We hypothesize that this strain has been more prone to DNA instability events, possibly caused by its excessive satellite DNA abundance. After applying DNA replication and physical stressors, we measured DNA damage levels directly and demonstrate that the DNA damage response is associated with satellite abundance. We use two genetically identical strains that differ only by a chromosome fusion and satellite DNA abundance, and demonstrate that the strain with more satellite DNA has significantly increased DNA damage when stressed. Finally, we propose a model that genome instability may impose a constraint on satellite abundance, a model that is entirely consistent with our data.

RESULTS

Two novel sex chromosome Robertsonian translocations in D. virilis strain vir00
In summer 2019, we performed DNA fluorescence in-situ hybridization (FISH) on larval neuroblast nuclei of the vir00 strain that we had obtained several months earlier from the National Drosophila Species Stock Center. We discovered it contained a Y-autosome fusion (Figure 1A). The Y chromosome is recognizable in *D. virilis* because it has a distinct DAPI staining intensity pattern, and contains a distinct arrangement of satellite DNA (Flynn *et al.* 2020). However, all other chromosomes are difficult to distinguish in metaphase spreads, so we could not immediately determine the fusion partner. We observed that the fused chromosome contained the same centromere-proximal satellite as the Y chromosome, AAACAT. To determine if this chromosome fusion was present in other strains from similar geographical locations, we imaged larvae from strains vir08, vir86, and vir48, which were collected from different localities in California and Mexico (vir00 was collected from California). No larvae screened in these other strains contained any chromosome fusions. We designate the vir00 substrain with the Y fusion as vir00-Yfus.

In Fall 2019, we obtained a second stock of vir00 from the National Drosophila Species Stock Center. We set up single pair crosses and performed larval neuroblast squashes to karyotype multiple male progeny of each cross. Surprisingly, we found that the Y-autosome fusion was not present in any of the larvae karyotyped. However, 3/10 crosses karyotyped contained a different fusion. This new fusion contained the brightly-dapi-staining AAATTAC satellite as the centromere-proximal satellite, indicating that it is completely distinct from the Y fusion and involved different chromosomes. We hypothesized that this new fusion involved the X chromosome for several reasons; 1) based on centromere satellite identity, it had a 67% probability (Flynn *et al.* 2020); 2) it was found only in single copy in male larvae, but sometimes two copies in female larvae; 3) it was associated with observed X-Y nondisjunction events, such as the presence of an XXXY female (Figure S1). We performed single-pair crosses and screened the resulting progeny until we isolated a substrain fixed for the X fusion, and called this substrain vir00-Xfus. We maintained one of the cross descendants from the 2019 stock that did not have any evidence of the X fusion, which we designate vir00-Nofus. We inferred that both X and Y fusions likely represent canonical Robertsonian translocations, in which two acrocentric
chromosomes that underwent DNA breakage were fused together at the centromere during repair.

**Figure 1.** Discovery and genetic validation of two independent fusion events in vir00. A) DNA-FISH in metaphase chromosome squashes demonstrating the Y fusion. B) DAPI staining of metaphase chromosome squashes demonstrating the X-4 fusion. C) Genetic validation of the Y-3 fusion and D) the X-4 fusion. Red chromosomes: *D. novamexicana*; orange: wildtype *D. virilis* autosomes; green: autosome containing a GFP marker; light blue: X chromosome; dark blue: Y chromosome.

Genetic validation revealed Y-3 and X-4 fusions

We designed two separate two-generation crossing experiments to validate the fusions genetically and identify the autosome each sex chromosome is fused to. Both experiments exploited autosomal markers which we could determine if they were segregating non-independently of sex. The first experiment to validate the Y-autosome fusion used crosses between vir00-Yfus and *D. novamexicana* and scoring of microsatellite loci on each candidate autosome. We scored 82 F2 male progeny for the Chr3 marker, and 79/82 contained both alleles, whereas the null hypothesis was 50% should contain both alleles (Figure 1C, Figure S2).
The three progeny that did not contain both alleles were determined to have nondisjunction events and did not contain the Y chromosome (Figure S3). We concluded that the Y chromosome is fused to Chr3, and that the fusion was fixed in this subline. The other markers on Chr2, Chr4, and Chr5 segregated independently of sex and acted as negative controls (Table S1). We also did a negative control with the same crossing scheme except with vir08 instead of vir00 (Chr3 chi-square p = 0.39, N=22, Table S1).

The second experiment to validate the X-autosome fusion used crosses to *D. virilis* transgenic lines containing GFP markers on one of each of the candidate autosomes. For the crosses to Chr4-GFP (vir95), we phenotyped 304 F2 female progeny, and found that progeny containing a GFP signal were significantly depleted compared to the Mendelian expectation of 50% (Figure 1D). This indicated that the X chromosome was fused to Chr4. This crossing scheme with two other candidate autosomes did not show association of GFP signal with sex (Chi-square p-value > 0.1, Table S2). We also performed a negative control with the *D. virilis* genome strain instead of vir00-Xfus crossed to the Chr4-GFP line and found the GFP signal was independent of sex (Chi-square p-value = 0.92, Table S2).

We validated that vir00-Yfus and vir00-Xfus are the same genetic line and not the result of contamination from other lines either in our lab or the stock center. We made use of medium-coverage whole genome sequencing from Flynn et al. (2020) to design primers to amplify singleton insertion/deletion variants present only in vir00 (the version that was sequenced was, in hindsight, vir00-Yfus) and not in any other wildtype *D. virilis* strain present in the stock center (Table S3). We designed primers to amplify four loci on chromosomes 2, 3, 5, and 6 which contain a homozygous 12-13 bp deletion in vir00 compared to the reference and the other strains (Table S4). We found that both versions of vir00 contained the deletion at each of these loci, confirming that these substrains are indeed the same strain and not a contamination (Figure S4). Although we did the above indel experiment first, the whole genome resequencing SNP analysis (below) was also concordant with these substrains being the same genetic line.
Satellite DNA decreased in vir00-Nofus compared to vir00-Yfus

We used Illumina to resequence the three vir00 substrains: vir00-Yfus, vir00-Xfus, and vir00-Nofus. We then used k-Seek to quantify the satellite abundance in each substrain (Wei et al. 2014). Assuming the most ancestral strain was vir00-Yfus, we found that satellite abundances decreased in the two derived lines vir00-Xfus and vir00-Nofus (Table S5). In vir00-Xfus, there was an 8% decrease only in the centromere proximal satellite of ChrX and Chr4 (AAATTAC).

vir00-Nofus had an overall 12% loss of satellite DNA compared to vir00-Yfus (Figure 3b). vir00-Nofus had a similar 8% decrease in AAATTAC, in addition to a 10% loss in the pericentromeric satellite (AAACTAC) and a 13% loss in the centromere-proximal satellite of ChrY and Chr3 (AAACTAT). These data are consistent with our interpretation that vir00-Nofus is the result of breaking apart of the fusion chromosomes, and we conclude that significant satellite DNA was lost from all three pericentromeric and centromeric satellites in vir00-Nofus.

DNA damage levels in response to stress is related to satellite abundance in D. virilis

The vir00 strain, with the highest abundance of satellite DNA, contained two recent independent Robertsonian translocations. There are multiple steps required to detect a Robertsonian translocation: DNA breakage near the centromere, repair and fusion with another acrocentric chromosome, retention of a functional centromere, and intergenerational retention. To isolate the first step and the fundamental process in genome instability, we sought to measure DNA breakage directly. Since vir00-Yfus also contained the highest abundance of satellite DNA compared to the other strains, we wanted to test if variation in DNA breakage level after stress was associated with satellite DNA abundance. We therefore measured DNA damage levels in 7 different D. virilis strains, including vir00-Yfus and vir00-Nofus, with varying abundances of satellite DNA in response to replication stress and low-level radiation.
Figure 2. Comet assay images of nuclei isolated from *D. virilis* adult testes of the genome strain vir87 (A/C) and from vir00-Yfus (B/D) with and without stress (gemcitabine + radiation). Blue arrows point to the comet head and yellow brackets indicate the comet tails. The comet head and tail distribution for GDvir are of similar size and intensity. In vir00-Yfus, the comet tail is largely diffuse and expanded in the stress conditions compared to the control. These strains represent the lowest and highest satellite DNA abundances which have the least and greatest increase in DNA damage in response to stress, respectively.

Since spontaneous DNA breakage events are rare and we did not want to confound our data with breaks that occur as part of meiotic crossing over, we used stressors to elevate the rate of DNA damage. This would allow us to potentially detect a difference in the phenotype of interest between strains and also amplify possible types of stress imposed by excess satellite DNA content. We fed 0-1 day old adult flies the nucleoside analog gemcitabine for 8 days, which stalls replication forks and acts as a sensitizer for radiation via the Rad51 pathway (Kobashigawa et al. 2015). We then irradiated these sensitized flies with gamma rays at low level radiation (10 Gy). For each of seven strains tested, we included a control which was fed with the same liquid food with no gemcitabine and did not receive radiation treatment. We used the comet assay or single-cell gel electrophoresis to measure DNA damage in the male
germline in each line (Figure 2). We quantified comets using OpenComet software and used the olive moment measurement as the statistic representing DNA damage (Gyori et al. 2014) (Table S6). To detect differences in DNA damage in response to stress between strains, we took the mean difference in olive moments between the control and stress treatments (see Methods).

The strain with the lowest satellite abundance, the genome strain, contained the lowest DNA damage response and this was significantly lower than all other strains tested (Figure 2A,C, Figure 3A). The strain with the highest satellite abundance, vir00-Y fus, had among the highest DNA damage responses (Figure 2B,D), but was not significantly different from two other strains vir8 and vir48 (Figure 3A). Like most other phenotypes, there are likely multiple genetic factors contributing to the variation we found. Ideally, to demonstrate that satellite DNA is a causal factor, we would manipulate satellite DNA abundance and test the DNA damage phenotypes, but multi-megabase long arrays of satellite DNA cannot be manipulated with traditional genome editing. However, vir00-Y fus and vir00-No fus differ only by a chromosome fusion and 12% satellite abundance. Thus, if there is a difference in DNA damage response between these substrains, it may be caused by differing satellite abundance. vir00-No fus, which contained 12% less satellite DNA than vir00-Y fus, had a significantly reduced DNA damage response, which is concordant with our expectations that satellite DNA plays a causal role (Figure 3B).
Figure 3. Satellite DNA is associated with DNA breakage in response to stress. A) Olive moment, a statistic of the comet assay, measuring DNA damage for 7 different strains paired with a control and stress treatment. N indicates the number of nuclei analyzed for each treatment. Flies treated with gemcitabine and radiation are suffixed with “rad” and control flies are suffixed with “con”. The lower panel shows the unpaired mean difference between “rad” and “con” for each fly strain (black dot) and 95% confidence intervals (black line) produced from dabestr (5000 bootstrap method). Groups a, b, and c indicate samples with overlapping confidence intervals. B) The satellite DNA abundance of each D. virilis strain (x axis) and its unpaired mean difference between treatment and control (y axis). vir00-Nofus experienced a 12% loss in satellite DNA compared to vir00-Yfus, which is associated with a significantly decreased DNA damage in response to stress. C) Model that is consistent with our observations and data.

Extremely minimal degradation of the neo-Y chromosome

The Y-fused version of Chr3 is expected to accumulate mutations independently of the autosomal version of Chr3 over time because of the halt in recombination in male flies. This
would be represented in the mapping of short-read sequencing data by elevated heterozygosity on Chr3. Specifically, if the mutations arose after the fusion of Chr3 to the Y, they would not be present in any other strains of virilis, assuming no recurrent mutation. Thus, we used GATK genotyping and found heterozygous singletons unique to vir00-Yfus compared to other non-vir00 strains on each autosome (Table S3). We found that the number of heterozygous singletons was modestly but significantly enriched on Chr3 in vir00-Yfus: 2.45 SNPs/Mb more than other autosomes (permutation test, p < 0.001). We suggest this elevated density of singleton heterozygous sites may be due to the fusion with the Y chromosome and lack of recombination over several generations. Assuming the enrichment was caused only by the Y-fusion and that the neo-Y (Chr3) evolved clonally in a single lineage, we roughly estimate the Y-3 fusion occurred 1000-2000 generations ago.

Because the X-4 fusion was segregating with the no-fusion karyotype, recombination in heterozygotes would prevent degeneration of the neo-Y version of Chr4. Furthermore, we believe the X-4 fusion occurred between 2018-2019 since it was not present in the stock we obtained in 2018. We did not observe an enrichment of heterozygous singletons on Chr4 in vir00-Xfus. However, there was still a slight enrichment on Chr3 in vir00-Xfus (1.45 SNPs/Mb, permutation test p = 0.031), supporting our assumption that vir00-Xfus is the result of the breaking apart of the Y fusion and forming a new fusion.

**Nondisjunction between the X and Y chromosomes is highly elevated in chromosome fusion lines**

Nondisjunction occurs when homologous chromosomes fail to separate at meiosis, and results in aneuploidy in the progeny. Autosomal and X chromosome aneuploidy is lethal in flies; but Y chromosome aneuploidy is viable: females that have a Y chromosome (XXY) are fertile, males with no Y chromosome (XO) are sterile, and males with two Y chromosomes (XYY) are fertile. Elevated rates of X-Y nondisjunction represent a fitness cost because zygotes with infertile or lethal karyotypes will form at increased frequency. We found some evidence of nondisjunction

Nondisjunction
in the vir00-Yfus genetic validation (Figure S3), and also common Y chromosome aneuploidy in
the stock of vir00-Xfus (Figure S1). Since both fusions involve the sex chromosomes, we tested
the rate of primary X-Y nondisjunction in males in all three versions of vir00, along with the D.
*virilis* genome strain as a control. We first fully isolated the vir00-Nofus strain to ensure no
fusion chromosomes were segregating, which even at low frequency could increase the rate of
nondisjunction in the line.

We crossed individual males of each strain to genome strain females and genotyped for the
presence or absence of the Y chromosome in progeny. Female progeny containing a Y
chromosome indicate XY sperm from the father, and male progeny lacking a Y chromosome
indicate nullisomic sperm from the father. Since nondisjunction was extremely high in vir00-
Xfus, 2/7 fathers tested were of XYY karyotype, which we could infer if more than half of his
female progeny contained a Y chromosome (Maggert 2014). We eliminated these fathers’
progeny from the primary nondisjunction rate calculation. No fathers tested from other
sublines were determined to be XYY. Males without a Y chromosome would not produce
progeny. We found that vir00-Yfus had a slightly elevated nondisjunction rate of 4.5%,
compared to the genome strain control of 1.2%, but it was not statistically significant with the
sample sizes we used (Table 1). vir00-Xfus had an extremely high primary nondisjunction rate of
21% (e.g. Figure S5), which was significantly higher than that of all other substrains (p < 0.004,
pairwise proportion test, Table 1). Surprisingly, vir00-Nofus had a significantly elevated
nondisjunction rate compared to the genome strain control, at 5.7%. This was not statistically
different from vir00-Yfus.
Table 1: Nondisjunction of the X and Y chromosomes in males is elevated in chromosome fusion lines.

| Line          | Nondisjunction rate | Fathers with nondisjunction/total fathers | Aneuploid progeny/total progeny | Significance group |
|---------------|---------------------|------------------------------------------|--------------------------------|--------------------|
| GDvir         | 1.2%                | 1 / 10                                   | 3 / 248                        | a                  |
| vir00-Yfus    | 4.5%                | 4 / 7                                    | 9 / 198                        | ab                 |
| vir00-Nofus   | 5.7%                | 4 / 6                                    | 11 / 192                       | b                  |
| vir00-Xfus*   | 21%                 | 5 / 5                                    | 27 / 127                       | c                  |

*only including primary nondisjunction

Discussion

*D. virilis* has the highest relative satellite DNA abundance of any studied species. Is there a maximum limit of satellite DNA a genome can tolerate before there are negative consequences? We found that vir00, the strain with 15% more pericentromeric satellite DNA than other strains recently underwent two independent chromosome fusion events. *vir00-Yfus* has the highest satellite DNA abundance and among the highest DNA damage level in response to stress. *vir00-Nofus*, with 12% less satellite DNA, which was presumably lost in the breaking apart of the chromosome fusions, has significantly lower DNA damage level in response to stress. We propose a model that places a constraint on satellite DNA abundance and can explain our findings with the *vir00*-line (Figure 3C). If satellite DNA expands, the risk of DNA breakage and genome instability increases and chromosome fusions may occur. These chromosome fusions may increase the nondisjunction rate and make the affected line less fit. The chromosome fusions may later break apart; and as this occurs pericentromeric satellite DNA may be lost (Figure 3C). This in turn results in a decreased risk of genome instability. The
question of whether satellite DNA abundance influences genome instability has had considerable interest recently (Arora et al. 2021), however a convenient system in which to test this question has been lacking.

There are several possible mechanisms that might cause excess satellite DNA to increase the risk of DNA breakage. Satellite DNA may form complex structures, loops, or non-B DNA, and increasing the length of the arrays may make these regions more unstable in cis (Barra and Fachinetti 2018). It may be challenging for polymerases to replicate many megabases of tandem satellite sequence, and longer arrays may have a higher risk of stalling polymerases and thus DNA breaks in cis (Barra and Fachinetti 2018). Finally, increased satellite DNA may titrate away binding proteins that maintain genome stability, in trans (Francisco and Lemos 2014; Brown et al. 2020a; Giunta et al. 2021). Whether there is a maximum threshold of satellite array length or abundance, as suggested in Charlesworth et al. (1986), or whether risk of genome instability varies continuously with satellite abundance will require further study.

Chromosome rearrangements such as Robertsonian translocations are rare occurrences, but are commonly the cause of karyotype evolution between species (Mayrose and Lysak 2021). In humans, rearrangements resulting from breaks near the centromere are associated with miscarriages and in the case of somatic rearrangements, cancer (Barra and Fachinetti 2018). Previous studies have found repetitive element involvement in chromosome rearrangements in multiple species (Paço et al. 2015; Reis et al. 2018). However, genetic variation of repetitive DNA within species has never been associated with increased risk of genome instability. Here our data suggests, at least in D. virilis strain vir00, that elevated satellite DNA may play a causal role in increasing the risk of DNA breakage, which can lead to deleterious rearrangements.

We identified elevated nondisjunction rates as a cost of chromosome fusions. Surprisingly, the rate of nondisjunction was 5-fold higher in vir00-Xfus compared to vir00-Yfus. With a nondisjunction rate of 21%, a significant proportion of abnormal karyotypes will be produced, such as XO (sterile), XYY (viable and fertile), and XXY or XXYY (viable and fertile), all of which we
found in cytological samples. Further mating between these abnormal karyotypes will produce significant proportions of sterile or inviable karyotypes like XYYY or XXX, which will further decrease the fitness of this line. Furthermore, karyotypes with extra Y chromosomes such as XYY and XXY have been found to have decreased lifespan (Brown et al. 2020b). Although we did not assay the nondisjunction rate of the fused autosome, it is possible the nondisjunction rate of Chr4 is also elevated in vir00-X fus, which would further decrease the fitness of the line because Chr4 aneuploidy is expected to be lethal (Lindsley et al. 1972). The extreme nondisjunction rate indicates X-Y pairing is severely disrupted due to the X-4 fusion, but only slightly if at all due to the Y-3 fusion. When the Y fusion presumably broke apart and the X fusion formed, it is possible other rearrangements on the X and/or Y occurred that disrupted pairing. Furthermore, the nondisjunction rate did not decrease to a level similar to the genome strain in vir00-Nofus. We believe this indicates a remaining rearrangement in vir00-Nofus affecting the pairing and disjunction of the X-Y. We only found a modest difference in estimated rDNA copy number between the substrains (Table S7), which has been found to mediate pairing of the X and Y (McKee and Karpen 1990). Detailed analysis of structural rearrangements in the heterochromatin will be required to determine the mechanism of the elevated nondisjunction rates.

Our study has several limitations. First, the vir00-Nofus flies we used for resequencing and for DNA damage assays had the X-4 fusion segregating at low frequency (<15%), which was unknown to us until we were able to correct it for the nondisjunction assays. However, we believe our results hold firmly because the main comparison was with vir00-Yfus, the substrate with the most satellite DNA. For testing our hypothesis, the fusion status in the other substrains matters less than the satellite DNA abundance, which was markedly lower in the vir00-Nofus substrate we used. Furthermore, the increase in damage level between the control and stressed flies may not be directly applicable to the risk of DNA breaks and genome instability in natural conditions. Although we find a difference in the DNA damage in response to stress between vir00 substrains with different abundances of satellite DNA, their fusion status is also different. We cannot eliminate the possibility that the presence of the Y fusion itself increased the rate of
DNA damage instead of the abundance of satellite DNA. Finally, we cannot eliminate the possibility that satellite DNA increased in vir00-Yfus after the fusion occurred and not prior to as we suggest in our model.

We believe the system we discovered will be useful for a variety of future studies. The vir00 fusion substrains will be useful for studying centromere identity. In both the Y-3 and X-4 fusions, two spherical regions of satellite DNA are present at the centromere of these fusions, representing one from each acrocentric chromosome (Figure 1). We note that the X-4 fusion in vir00 is homologous to an independent X-4 fusion in *D. americana* 29 thousand years ago, a species only 4.5 million years diverged. In the X-4 fusion of *D. americana*, there is only one discrete region of centromeric satellite (Flynn et al. 2020), unlike what we found here. In female meiosis where chromosomes can compete to get into the oocyte rather than the polar body, “stronger” centromeres with more satellite may have an advantage. A “supercentromere” resulting from a centromere-centromere fusion is one way to do this, and in *D. americana* the X-4 fusion has biased transmission into the egg (Stewart et al. 2019). Finally, since both centromere-centromere fusions had matching centromere-proximal satellites (AAACTAT for Y-3 and AAATTAC for X-4), we suggest that the centromeric satellite identity is important for DNA repair for Robertsonian translocations, or for stability and retention of the centromere.

**METHODS**

Scripts required to reproduce the computational results are available here:

https://github.com/jmf422/D-virilis-fusion-chromosomes

**Neuroblast squashes and satellite DNA FISH**

We dissected brains from wandering 3rd-instar larvae and performed the fixation steps as in (Larracuente and Ferree 2015). Specifically, we placed brains in sodium citrate solution for 6 minutes before fixation. After fixation and drying of slides, we applied Vectashield dapi
mounting medium. We performed DNA-FISH on vir00-Yfus, which allowed us to confidently identify the Y chromosome based on its unique satellite DNA composition. We used the same fixation and staining protocol as Flynn et al. (2020). We imaged metaphase cells using a 100x oil objective on an Olympus fluorescent microscope and Metamorph capture system at the Cornell Imaging Facility.

Y-autosome fusion validation

We designed an experiment that would both validate the Y chromosome fusion and to distinguish which autosome is fused. We first designed primers flanking microsatellite loci on all four autosomes that met the following criteria: 1) had 100% conserved non-repetitive and unique priming sites between *D. virilis* and *D. novamexicana*; 2) amplicon length differed between the species by at least 15 bp as to be distinguished on an agarose gel; 3) locus contained a mono or tri nucleotide repeat; 4) locus length ~200 bp. We next set up a two-generation crossing scheme (Figure 1A). We crossed *D. novamexicana* virgin females with vir00-Yfus males and selected the male progeny, which we backcrossed to *D. novamexicana* virgin females. We then genotyped the male F2 progeny from this cross at the 4 sets of primers corresponding to the four non-dot autosomes (Chr2, 3, 4, 5) (Table S4). We performed single-fly DNA extraction in strip tubes with Tris-EDTA buffer and 0.2 mg/mL proteinase K. We did 12 uL standard PCR reactions (3 min at 95, 30 cycles of 30 sec 95, 30 sec 55, 50 sec 72, final extension 5 min). Each primer on each PCR plate had a homozygous (*D. novamexicana*) and heterozygous (*D. novamexicana-D. virilis* F1 hybrid) control. We then ran the PCR product on 2.5% agarose gels. If there was indeed an autosome fused to the Y chromosome, we would expect to see 100% of the male progeny being heterozygous for the *virilis* and *novamexicana* alleles (except for rare cases of non-disjunction). For the autosomes that are not fused, we would expect to see 50% of the progeny being homozygous for the *novamexicana* allele, and half heterozygous, due to Mendel’s law of random segregation. We successfully validated the existence of the Y fusion, and found that it is fused to chromosome 3 (Muller D) (Table S1, Figure 1). There were 3 male progeny that were homozygous for the Chr3 *novamexicana* allele. We verified that these
were cases of nondisjunction (opposed to the Y fusion not being fixed in this subline) by finding that the Y chromosome was absent in controlled Y chromosome PCR assays (Figure S3).

**Isolation of vir00-Xfus and vir00-Nofus**

The X fusion was found to be segregating with a no fusion substrain in the 2019 stock of vir00. We wanted to isolate these into two separate substrains where the karyotype is fixed. From the progeny of the three original crosses in which we found the X fusion, we made 10 single pair crosses and did neuroblast squashes of 6-8 larval progeny per cross, including both sexes. By chance, we should be able to find a cross in which the mother had two copies of the fusion and the father had a single copy – in which the derived line would be fixed for the fusion. If all progeny imaged contained the fusion (and females contained two copies of the fusion), then it is likely that this was the case. We created this line, and call it vir00-Xfus. We maintained a line isolated from the 2019 stock that had no evidence of the X fusion and called it vir00-Nofus. We later found that vir00-Nofus still had the X fusion segregating at low frequency. We isolated a fixed Nofus version in the same way as above for nondisjunction assays, because a low frequency fusion could increase the nondisjunction rate greatly.

**X-autosome fusion validation**

We obtained transgenic strains with GFP (or Blue) insertions which are expressed in the eye and larval brain from the National Drosophila Species Stock Center (vir95, vir121, vir117). Stern et al. (2017) found the insertion sites of these lines. We chose lines which contained the GFP marker on candidate autosomes Chr2, Chr4, and Chr5. Chr3 was not a candidate because it is fused to the Y in vir00-Yfus and contains a different centromeric satellite. Before setting up crosses, we screened 10-20 larvae of each line with a fluorescent microscope to ensure the transgene had not drifted to low frequency. Larvae containing the transgene demonstrated the GFP signal in their brain. We chose to phenotype at the larval stage since we would be crossing GFP strains to wildtype red-eyed flies and the visibility of GFP in the adult eye would be low. We
then designed a crossing scheme which would allow us to both validate that the X chromosome was fused and distinguish which autosome it was fused to (Figure 1B). We crossed GFP-line males to vir00-Xfus virgin females. We then selected the male F1 progeny and crossed them to virilis genome strain virgin females. We then phenotyped F2 larvae, classifying each as either GFP positive or negative. When the phenotyped flies emerged, we sexed and counted them. If the candidate autosome is fused to the X, we would expect sex to segregate with the GFP marker: all female progeny will be GFP negative, and all male progeny will be GFP positive (except for phenotyping errors or rare nondisjunction events). For all other lines, sex should not segregate with GFP status. We performed negative control crosses in which the parental cross was replaced by genome strain virgin females, to ensure the crossing scheme produced the expected results (Table S2). We found that the X chromosome is fused to Chr4 (Muller B).

Validation of the three versions of vir00 with private fixed indels

We used GATK recommended practices to do genotyping of our low-coverage whole genome sequencing data from Flynn et al. (2020). We used vcftools to subset singletons present only in vir00, which was, in hindsight, vir00-Yfus. We then used GATK’s SelectVariants to select only non-reference homozygous indels 12 bp or more with a depth of at least 10 in vir00 and at least 2 in the other strains. We then manually inspected each potential candidate in IGV to ensure: no reads in other strains supported the indel, all reads in vir00 supported the indel, and there were no nearby indels in other strains. We then designed primers for the four loci (on Chr 2, 3, 5, 6) that met these criteria and also had enough SNP-free sequence flanking the indel in order to design primers that would amplify a locus 100-200 bp equally in all strains. We performed PCR and gel electrophoresis (2.5% gel, 98 V, 90 min).

DNA damage assays

We chose two stressors that would moderately increase the rate of DNA breaks and allow us to potentially detect differences between strains. Gemcitabine is a nucleoside analog that induces
replication stress by stalling polymerases, and also sensitizes cells to radiation via the RAD51 pathway (Kobashigawa et al. 2015). We selected dosage and a fly-feeding regime based on (Kislukhin et al. 2012). Ionizing radiation has long been used to increase the rates of DNA breaks in flies for mutagenesis. We chose a dose \( \frac{1}{4} - \frac{1}{2} \) of what has been typically used in mutagenesis (Carlson and Southin 1962).

We collected male flies 0-1 days old and fed them gemcitabine (0.718 mM) mixed with liquid food in vials with 8-12 adult flies as in (Kislukhin et al. 2012). Liquid food consisted of 12.5g sucrose, 17.5 g dry yeast, 5 mL corn syrup, and 95 mL PBS (autoclaved for 30 min immediately after adding the yeast). Flies were fed the drug for 7-9 days before radiation. Controls were fed with the same liquid food for 7-9 days, except no gemcitabine was added. Flies were moved to fresh vials every 3-4 days. For radiation treatment, we transferred flies into 50 mL conical tubes with 5 mL agar because these tubes were compatible with the radiation source. Control flies were also transferred to new tubes. We used a J.L. Shepherd & Associates Mark I Irradiator with 1,100 Ci of Cs-137, and flies were irradiated at approximately 400 rad/min for a total of 10 Gy. In one case, for the GDvir stress treatment, the radiation was not stopped on time so 4 extra Gy were applied. We believe this did not affect our results, especially because the GDvir strain had the lowest DNA damage increase with gemcitabine and radiation stress. We did comet assays to measure DNA damage (Angelis et al. 1999) over three different dates (Table S6), but ensured experimental conditions were practically identical each time. For some samples we had to combine results from two different dates to have enough nuclei for statistical analysis (Table S6). We dissected testes from approximately 8 flies from each treatment within one hour of radiation treatment to minimize the opportunity for breakage repair (Shetty et al. 2017). We then homogenized the testes tissue using a dounce, filtered the homogenate through a 40 micron filter to remove debris, and centrifuged and resuspended the cells to approximately \( 10^5 \) cells/mL.

We next performed the alkaline comet assay as directed by the Enzo comet kit (ADI-900-166), which provides higher sensitivity than the neutral comet assay (Angelis et al. 1999). We imaged
slides on a metamorph imaging system at 10x magnification using a fluorescent green filter to
detect the CyGreen dye included in the comet kit. We quantified damage levels using the
software OpenComet as a plugin in ImageJ (Gyori et al. 2014). We filtered called nuclei that
were not comet shapes or contained background interference. We used the measure of “olive
moment,” which is the product of the percent of DNA in the tail and distance between
intensity-weighted centroids of head and tail (Gyori et al. 2014) as the statistic to compare
between strains and treatments.

**Resequencing sublines to determine differences in their satellite abundance.**

Pools of 6 male flies were DNA extracted with Qiagen DNeasy blood and tissue kit. PCR-free
libraries were then prepared with Illumina TruSeq PCR-free library prep. Libraries were
sequenced on a NextSeq 500 single end 150 bp. We removed adapters and poly-G signal with
fastp and then ran k-Seek to count satellite abundances (Wei et al. 2014). We used average
read depth to normalize the kmer counts. We also mapped the reads to the *D. virilis* rDNA
consensus sequence ([http://blogs.rochester.edu/EickbushLab/?page_id=602](http://blogs.rochester.edu/EickbushLab/?page_id=602)) to estimate the
rDNA copy number in the three vir00 substrains as well as a vir08 as a control (Table S3, S7).

**Using sequencing data to estimate the age of the Y-3 fusion**

Scripts for this section are available here: [https://github.com/jmf422/D-virilis-fusion-chromosomes/tree/main/simulate_degradation](https://github.com/jmf422/D-virilis-fusion-chromosomes/tree/main/simulate_degradation). We used the sequencing data from Flynn et al.
(2020) in addition to data produced here for vir00-Yfus, vir00-Xfus, and vir00-Nofus and 10
other *D. virilis* strains. We mapped the data to the RS2 genome assembly using bowtie2. We
then genotyped with GATK following standard procedures (McKenna et al. 2010). We extracted
heterozygous singleton sites for vir00, and counted how many occurred on each autosome. We
calculated the enrichment on Chr3 in vir00-Yfus based on the difference from the average SNP
density on the other autosomes (excluding the dot chromosome Chr6). To determine whether
this enrichment of SNPs was significant based on the size of the chromosome and the number
of mutations, we randomly permuted the total number of heterozygous singleton SNPs on all autosomes and calculated the proportion falling on Chr3, and repeated this 1000 times.

We then performed simple simulations to determine approximately how many generations of mutation accumulation without recombination or selection would result in the enrichment we observed. Since heterozygous singletons are challenging for the genotyper to call with moderate coverage sequencing data, we incorporated this into our simulation. First, we made the genome assembly diploid then used mutation-simulator (Kühl et al. 2020) to simulate random mutations (transition/transversion ratio 2.0) at a rate of $2 \times 10^{-9}$ per bp per generation for 500, 1000, 2000, and 5000 generations on one copy of Chr3 only. We then simulated Illumina reads with ART (Huang et al. 2012) at the same depth as we have for vir00-Yfus in our real data (23 x haploid or 11.5 x diploid). We next used standard GATK genotyping and selected out heterozygous singletons on Chr3. We repeated the simulation 10 times for each number of generations to get a range of values. The empirical enrichment fell in between what we found in the simulations for 1000 and 2000 generations.

**Nondisjunction assays**

We crossed a single male from vir00-Yfus, vir00-Xfus, vir00-Nofus (fixed), and GDvir (control) to one or two GDvir (genome strain vir87) females. We collected the virgin progeny from each cross, extracted DNA with a squish-proteinase K prep, and genotyped with PCR and gel electrophoresis for the presence or absence of the Y chromosome in up to 16 female and 16 male progeny. We amplified a locus unique to the Y chromosome (primers designed by Yasir Ahmed-Braimah for a different project, Table S4). For a subset of individuals, we also performed multiplex controls with an autosomal locus. Otherwise, we performed DNA extractions in large batches with the same proteinase K mixture to minimize the chance of DNA extraction failure. A very small quantity of DNA is required for a standard PCR with robust primers. To control for the completeness of the PCR mastermix, we included male samples in the same batch as female samples. If a male lacked a Y chromosome, we inferred the father’s
sperm was missing the Y chromosome (nullisomic), and if a female contained a Y chromosome, we inferred the father’s sperm contained both X and Y. We used R prop.test to evaluate whether there were any differences between nondisjunction proportions for the different strains. After finding this highly significant, we used pairwise.prop.test in R with Holm-Bonferroni multiple test correction to determine which pairs of substrain nondisjunction rates were significantly different from each other.

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