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Authors
Hua, Lisa L
Mikawa, Takashi

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Mitotic antipairing of homologous and sex chromosomes via spatial restriction of two haploid sets

Lisa L. Hua* and Takashi Mikawaab,1

*Cardiovascular Research Institute, University of California, San Francisco, CA 94158; and bDepartment of Anatomy, University of California, San Francisco, CA 94158

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Pairing homologous chromosomes is required for recombination. However, in nonmeiotic stages it can lead to detrimental consequences, such as allelic misregulation and genome instability, and is rare in human somatic cells. How mitotic recombination is prevented and how genetic stability is maintained across daughter cells is a fundamental, unanswered question. Here, we report that both human and mouse cells impede homologous chromosome pairing by keeping two haploid chromosome sets apart throughout mitosis. Four-dimensional analysis of chromosomes during cell division revealed that a haploid chromosome set resides on either side of a meridional plane, crossing two centrosomes. Simultaneous tracking of chromosome oscillation and the spindle axis, using fluorescent CENP-A and centrin1, respectively, demonstrates collective genome behavior/segregation of two haploid sets throughout mitosis. Using 3D chromosome imaging of a translocation mouse with a supernumerary chromosome, we found that this maternally derived chromosome is positioned by parental origin. These data, taken together, support the identity of a model/paradigm and provides implications for maintaining mitotic fidelity.

Mitotic DNA recombination can be dangerous because of the high number of repeat sequences in the human genome that results in structural rearrangements and genome instability (1–3). Notably, recombination events can lead to loss of heterozygosity at specific gene loci and have been implicated in cancer initiation and progression (4, 5). To avoid this danger, recombination between nonallelic positions must be suppressed. One possible way of managing recombination may be limiting the potential pairing sites for recombination. Although homologous pairing in meiosis has been described in detail (6, 7), we know much less about the absence of pairing in mitosis.

It has been controversial whether homologous chromosomes are positioned antiparallel (8, 9) or distributed at random (10, 11) in human prometaphase cells. In this study, we revisit this unsolved but tantalizing question with many downstream implications. Methodologies that have been previously employed to analyze chromosome organization during mitosis include differential nucleotide detection by autoradiography (12), insertion of fluorescent repressive-operator arrays (13–15), photo-bleaching/photoactivation of fluorescently labeled histones (16–18), centromere marker-based tracing (19–21), and contrast-generating optical microscopy (22). Although these approaches provide a general observation of global chromosome organization, they lack detection for individual homologous chromosome pairs. To date, chromosome painting (23–25), the application of DNA probes specific to nonrepetitive sequences, has been a common and versatile technique to achieve simultaneous identification and visualization of homologous pairs. Employing this method, we systematically analyzed the spatial organization of all autosomal and sex chromosome pairs throughout mitosis.

Here, we report that homologous chromosomes are spatially segregated albeit without a defined position. We show that the spatial segregation is achieved by sequestering two haploid sets along specific cellular axes. Our data reconcile the current two contradictory models for the spatial organization of homologous pairs and propose a mechanism of antipairing between homologs during mitosis.

Results

Homologous Chromosomes Are Spatially Segregated Throughout Mitosis. For our initial survey of chromosome organization at prometaphase, the same stage analyzed in previous publications (8–11), chromosome paint probes were applied to human primary cells growing in a 2D cell-culture environment (Fig. 1 and SI Appendix, Fig. S1). We painted and mapped homologous pairs of various sizes (chromosomes 3, 4, 9, 11, and 18) in a simple epithelial cell type: human umbilical vein endothelial cells (HUVECs) (Fig. 1 A and SI Appendix, Fig. S1 B–G). Two-dimensional culture of HUVECs displays clear apical/basal polarity (26). The spatial relationship between a homologous pair was determined by interhomolog angular measurement. At each homologous pair is positioned antiparallel to each other, an optimum/maximal angle of 180° will be obtained (8, 9). Our 3D reconstruction analysis of chromosome paint data revealed that

Significance

Mitotic recombination must be prevented to maintain genetic stability across daughter cells, but the underlying mechanism remains elusive. We report that mammalian cells impede homologous chromosome pairing during mitosis by keeping the two haploid chromosome sets apart, positioning them to either side of a meridional plane defined by the centrosomes. Chromosome oscillation analysis revealed collective genome behavior of noninteracting chromosome sets. Male translocation mice with a maternal-derived supernumerary chromosome display the tracer chromosome exclusively to the haploid set containing the X chromosome. This haploid set-based antipairing motif is shared by multiple cell types, is doubled in tetraploid cells, and is lost in carcinoma cells. The data provide a model of nuclear polarity through the antipairing of homologous chromosomes during mitosis.

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1To whom correspondence should be addressed. Email: takashi.mikawa@ucsf.edu.

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Autosome pairs exhibited angular measurements of <90° (SI Appendix, Fig. S1A), indicating the homologous chromosomes deviated from a theoretically defined antiparallel configuration. A rose plot, displaying the frequency of angular distribution for the different homologous chromosomes, shows high angular variations (n = 37, $\chi^2$ test $\chi^2 = 0.10$; SI Appendix, Fig. S1A). Thus, the data are not consistent with the antiparallel/fixed position model for individual homologous pairs.

In no case, however, did we find two homologs occupying the same position within a prometaphase rosette. For example, for chromosome 4 each homolog was consistently separated from one another with an average distance of 3.6 ± 1.2 μm SD and angular orientation of 73.5° ± 44.3° SD, 3.3 ± 1.8 μm SD and 82.4° ± 50.9° SD, and 2.9 ± 1.4 μm SD and 88.4° ± 50.1° SD, respectively. The blue asterisk indicates a measurement representative of homologous chromosome pairing. (F) A connected scatterplot portraying the relative frequency distribution of minimal distance between chromosome 4 and its homologous partner (green line) or a nonhomologous chromosome (X or Y, gray dotted lines) at metaphase (square)/anaphase (triangle) ($P = 0.030$, $P < 0.001$ for metaphase/anaphase). (G–J) Select frames of a high-resolution 4D time-lapse movie of a cell labeled with CENP-A GFP and centrin1-GFP (arrowheads) to visualize the centromeres and centrosomes, respectively, undergoing mitosis from a top view (G–J) or a side view (G–J′). Six centromeres are labeled (colored spots) at an initial time point at (G and G′) prometaphase and tracked (colored lines) until metaphase (H and H′). Other centromere tracks are designated in gray. Note that at the beginning of prometaphase the centromeres are above/below the chromosome rosette. (L and L′) As in H and H′ but from metaphase to telophase. (K–K′) Tilted view of a HUVEC at anaphase stained with DAPI (blue) and γ-tubulin (white) visualized by immunofluorescence with the center of mass of the two sister anaphase chromosome masses labeled (white pixels) from confocal optical sections. Note that the centrosome, or mitotic spindle, axis is coincident with a nuclear meridional plane in mitotic cells. (K′) top view. (K′) side view. (L) Schematic delineating the $x$, $y$, and $z$ axes in a cell at anaphase. The $x$ axis was determined by the line crossing the center of mass of the two sister anaphase chromosome masses, the $z$ axis was determined as the perpendicular line to the coverslip, and the $y$ axis was defined as perpendicular to both $x$ and $z$ axes. (Scale bars: 1 μm.)
C, E, and F) (null hypothesis: homologs positioned at random relative to each other and similar to a nonhomologous partner, Student’s t test, P = 0.030, P < 0.001, for metaphase/anaphase; Fig. 1F). These data demonstrate that homologous chromosomes are not organized in an antiparallel orientation to each other. However, they did not come into contact and did appear to segregate from one another, thus being antipaired in fashion, throughout mitosis.

The above findings—that homologous chromosomes are predominantly distant from each other and not often close at different stages of mitosis—prompted us to question whether this reflects a level of organization in the mitotic chromosomes and/or overall cellular organization. With the emergence of new tools and the ability to define axes in 3D with cellular markers, it is now possible to accurately map the positions of individual chromosomes along a subcellular axis/plane. To develop and validate a 3D coordinate system to reliably map the homologs, the extent of chromosome movement, such as translational/rotational movements, was determined and the x, y, and z axes were defined during mitosis.

We tracked and analyzed the position and movement of individual chromosomes throughout mitosis in real time, using a human epithelial cell line (RPE1) (19) that stably expresses CENPA-GFP and centri1-GFP to identify the centromeres/chromosomes and centrosomes/nuclear division axis, respectively. The 3D, live-cell analysis revealed that from prometaphase to metaphase chromosomes displayed unstable movements along the centrosome axis (n = 5; Fig. 1 G–H and Movie S1), most likely due to concurrent formation of the mitotic spindle (27). In contrast, from metaphase to anaphase chromosomes exhibited stable movements along the nuclear division axis (n = 14; Fig. 1 I and J and Movie S2). Individual chromosomes predominantly moved parallel to, or along, the centrosome axis with little perpendicular fluctuation. Perpendicular fluctuation and rotation of chromosomes along the centrosome axis became apparent following late anaphase (n = 7; Fig. 1 J and Movie S3). As the nuclear envelope reforms during telophase, chromosomes in the sister nuclei act independently and asynchronously (Movie S3). The small displacement changes from metaphase to early anaphase suggested that individual chromosomes maintained their relative position to one another within the mitotic chromosome mass. These results also corroborate previous studies showing limited movements when chromosomes are attached to the mitotic spindle (19, 27, 28). Therefore, the absence of major displacement changes at metaphase to midanaphase allowed us to quantitatively map individual chromosomes in a 3D coordinate system (Fig. 1 K–L). The z axis was fixed as the optical path of the microscope, a perpendicular line to the coverslip, and along the apical–basal axis. The x axis is perpendicular to the z axis and coincident with the centrosome axis (Fig. 1 K, K’, and K’’); the origin (0, 0, 0) was then defined by a line crossing the center of the mitotic chromosomal mass, and the y axis was defined as perpendicular to both x and z axes (Fig. 1L). At prometaphase, a lack of a definable y axis along the plane of the chromosome rosette prevented establishment of a coordinate system.

To test whether there was a conserved position/address for each pair of homologous chromosomes that could be responsible for the antipairing organization of homologs we systematically mapped individual chromosomes in a 3D axial coordinate system (SI Appendix, Fig. S2 A–E). We quantified chromosome position using both 3D, superimposed, homolog contour map data and a normalized 2D axial coordinate system (SI Appendix, Fig. S2 F–I). The summation of all homologous chromosomes in physical space was plotted on a 3D contour map, upon which grayscale saturation represents the density of localization when all of the homologous chromosomes are overlaid (SI Appendix, Fig. S2 F and H). Isolines were then generated that connect contours that share the same gray value/intensity, revealing hotspots of maximal overlap. If the homologous chromosomes were in fixed positions there would be a clustered area of high density/occurrence that reflects the colocalization frequency for each of the homologs. In addition, the mapped position of each homologous pair in a coordinate system would also be identical for every cell. Neither analysis identified any evidence of fixed localizations for the homologous chromosomes based on the coordinate system in the present study (SI Appendix, Fig. S2 F–I and Movie S4). However, there was a modest correlation of chromosome size to position based on median values for individual chromosome populations (n = 578 cells, R² = 0.6119, P < 0.05; SI Appendix, Fig. S1J), corroborating previous work (29). These results suggest an alternative mechanism that is not regulated by a defined address that drives the antipairing organization of homologs.

**Homologous and XY Chromosomes Segregate to Opposite Nuclear Hemispheres During Mitosis.** Although our superimposed 3D matrix analysis demonstrated that a pair of homologs does not display fixed positions, surprisingly a distinct pattern did emerge. The 3D overlay data revealed that each autosome in a pair was localized to separate hemispheres, on either side of a “meridional plane” (n = 15/20 metaphase cells and n = 28/38 anaphase cells, Fig. 2 A and B for chromosome 4; see SI Appendix, Fig. S3 A–AR for all other autosomes). The meridional plane, an imaginary z plane horizontal to the metaphase/anaphase equatorial plate, was coincident with the centrosome, or x, axis, and was found to traverse along the apical/basal, or z, axis (n = 28 cells; Fig. 1 K–L). Midpoint calculation for pairs of homologous chromosomes confirms the meridional plane to consistently partition individual homologs to separate hemispheres (Fig. 2C). To test whether such an arrangement occurs by random chance, a statistical binomial test (30, 31) was performed on the positional coordinates of the center of mass for each chromosome (P = 0.021, P = 0.003 at metaphase/anaphase) (null hypothesis: homologous chromosomes are randomly arranged in the two nuclear hemispheres). The data support the presence of an axis-dependent antipaired configuration of one homolog per nuclear hemisphere at metaphase that persists throughout anaphase in dividing cells.

To evaluate whether sex chromosomes followed the same pattern observed for autosomes, female-derived HUVECs were examined with X chromosome-specific probes at metaphase and anaphase (SI Appendix, Fig. S3 AS–AX). The pair of X chromosomes showed the identical one homologous partner per hemisphere motif, rather than a random or overlapping distribution (n = 18/24 metaphase cells, P = 0.011 and n = 32/42 anaphase cells, P = 0.001; SI Appendix, Fig. S3 AS–AX). To explore whether the idea of a sequence homology based-repulsion between a pair of homologs is required for this organization, we visualized the sex chromosomes in male cells as each partner has minimal levels of sequence homology (32). The X and Y chromosome preserved the same pattern of one homolog per hemisphere in male cells (n = 19/28 metaphase cells, P = 0.044 and n = 20/28 anaphase cells, P = 0.001; Fig. 2 D–F and Movie S5). This result suggests that sequence homology is not required for establishing the one-homolog-per-hemisphere motif. Simultaneous visualization of sex and autosomal chromosomes demonstrates that both XY and homologous pairs follow the same topological hemisphere segregation (Fig. 2 G and H). Consequently, this discrete chromosomal organization produces a compartmentalization of two complete haploid chromosome sets along the nuclear division axis. In addition, another primary human cell type displayed a similar haploid set segregation motif (human fibroblasts, HFFs; SI Appendix, Fig. S4 A–F), suggesting the pattern may be conserved broadly for multiple cell types and is cell-type-independent.
Consistent with these observations, when chromosomes were assigned to groups based on their initial start position along the centrosome axis at metaphase and tracked to anaphase using the CENP-A/centrin1-GFP cell line, we found minimal mixing of individual chromosomes between the two groups (Fig. 2I and J and Movie S6). Detailed quantification of chromosomes at the interface between the two labeled groups demonstrates that the majority of chromosomes remain largely in their original positions relative to the chromosome mass (Fig. 2K). The data conclude that, in the absence of major chromosome fluctuations perpendicular to the centrosome axis during mitosis, the axis-dependent segregation between a pair of homologous chromosomes is conserved from metaphase to anaphase. These results thus confirm that the sets of chromosomes are showing a collective behavior during mitosis and support that collective genome segregation occurs as a haploid set.

A possible mechanism underlying the haploid set-based segregation is that each haploid compartment contains chromosomes of the same parental origin. Since the male and female pronuclei do not immediately fuse and are separated at early stages after fertilization (21, 33–39), it is plausible this parental origin may be spatially inherited as the identity for each haploid set. In support of this parental origin hypothesis, we tested two different experimental conditions: (i) compartmentalization of a haploid chromosome set is spatially preserved in naturally occurring tetraploid cells (4n), and (ii) genetic tracing of chromosomes reveals haploid set derived by parental origin.

Haploid Set Compartmentalization Is Conserved in Naturally Occurring Tetraploid Cells (4n). To first test whether a haploid set compartmentalization is spatially preserved and tightly regulated, we took advantage of naturally occurring tetraploid cells...
(4n) in human primary culture that undergo a whole-genome duplication event (40, 41). If individual haploid chromosome sets are sequestered from each other, we might expect this motif would be preserved following an additional duplication without nuclear separation in normal cells. We examined the sex chromosome distribution of male tetraploid HUVECs during mitosis to test whether tetraploid cells maintained compartmentalization for each of the four haploid sets (karyotype: XXYY). From prometaphase to anaphase, all of the four individual sex chromosomes were segregated from each other (n = 4/4 prometaphase cells, n = 16/20 metaphase cells, and n = 8/10 anaphase cells; Fig. 3 A–C). Quantitation of the distance and angular orientation of the XX and YY sex chromosome pairs revealed no pairing/overlap between the identical chromosome pairs within a tetraploid cell (n = 19/20 metaphase cells; Fig. 3D). Surprisingly, 3D overlay data showed each sex chromosome segregated to separate quadrasspheres along an axis parallel to the meridional plane (n = 15/20 cells; Fig. 3E). When the distance between both XY pairs in a single tetraploid cell was compared by subtraction, with zero representing identical distances, approximately similar distances for XY pairs were found with an average distance of 1.2 ± 1.1 μm SD (n = 20 cells; Fig. 3F). Taken together, these data suggest the compartmentalization of four individual haploid sets (1n) in tetraploid cells (4n) along the nuclear division plane (Fig. 3G). The data are not only consistent with previous findings of haploid set separation in triploid cells (9, 42) but also with the parental origin model, in which each haploid set is spatially discrete and compartmentalized.

**In Vitro and in Vivo Segregation of Mouse Chromosomes.** To directly test whether each haploid set is derived by parental origin, we labeled the maternal genome using a translocation mouse, Ts65Dn (43–47), which produces an extra supernumerary marker chromosome, to genetically distinguish the maternal genome set from the paternal genome set (Fig. 4G). However, first we had to determine whether homologous chromosomes in the mouse share the same haploid set-based antipairing organization as in humans. Our chromosome paint assessment of primary mouse embryo-derived fibroblast cells (MEFs) confirmed that, similar to humans, the homologs are not paired and segregated to separate hemispheres (n = 28/36 anaphase cells, P = 0.001; Fig. 4A–D). The sex chromosomes also displayed the one-homolog-per-hemisphere motif (n = 28/42 cells, P = 0.022; Fig. 4E and F). Taken together, these results demonstrate the haploid set-based pattern is conserved in the mouse.

The translocated chromosome 17<sup>16</sup> was distinguished from intact chromosome 16 by its size in anaphase cells (Fig. 4H). Each homologous chromosome 16 localized separately to either nuclear hemisphere, one of which contained the translocation chromosome 17<sup>16</sup> (n = 28/42 cells, P = 0.022; SI Appendix, Fig. S5 A and B). To identify which X-, or Y-containing haploid set was most likely associated with the translocation chromosome, we first determined the approximate boundary between the haploid sets by finding the midpoint of the pair of homologous chromosome 16. We then mapped the position of the translocated chromosome (Fig. 4I). In the majority of cases we examined, the hemisphere containing the translocated chromosome 17<sup>16</sup> also

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**Fig. 3.** Haploid set compartmentalization is conserved in normally occurring tetraploid (4n) cells. (A) Top view of male tetraploid (XXYY) HUVEC at prometaphase stained with TO-PRO3 (gray) and painted for the X (red) and Y (cyan) chromosome. (B and C) As in A but at metaphase and anaphase, respectively. (D) Distance and angular orientation of the XX and YY sex chromosome pairs (n = 19/20 cells at metaphase) with an average distance of 3.9 ± 2.3 μm SD and 3.1 ± 1.26 μm SD and angular orientation of 95.9° ± 47.9° SD and 68.3° ± 58.5° SD, respectively. (E) A 3D overlay of sex chromosome distribution of multiple male tetraploid (XXXX) cells at metaphase (n = 20). Each sex chromosome, X or Y, was assigned to be yellow/green/cyan/white based on its sequential order along the y axis and mapped to generate 3D overlay data. (F) Distance differences for a pair of XY chromosomes between two nuclear hemispheres with mean and SD error bars (n = 20). (G) Model for haploid set compartmentalization in a tetraploid cell. (Scale bars: 1 μm.)
contained the maternally derived X chromosome (n = 28/42 cells, P = 0.022; Fig. 4H–L and Movie S7). If the 17\textsuperscript{16} chromosome behaves similarly to other wild-type autosomes, these results are consistent with the model wherein one haploid set of an individual cell contains the maternally derived chromosomes while the other haploid set contains the paternally derived chromosomes.

To address whether the antipairing organization of homologs seen in cultured cells reflects the chromosome organization in their physiological environment, we examined the distribution of homologous chromosomes in neuroepithelial cells of E9.5 mouse embryos. Analyses of anaphase cells revealed consistent separation of homologs (n = 22/28 cells; Fig. 4M–P), in accord with cultured cells (Figs. 1–4). These data indicate that the establishment and maintenance of separation for homologous chromosomes are also conserved in the embryo.

**Discussion**

Genomes are dynamic entities that can be changed by processes such as recombination (1–3). As genome complexity increases, recombination events during mitosis minimize and are limited to meiosis (48). Therefore, homologous recombination must be tightly regulated to avoid genomic instability at mitosis but allow for precisely programmed genetic diversity during meiotic differentiation (49).

Our study has demonstrated a haploid chromosome set-based antipairing organization of homologous chromosomes during mitosis in human cells. A previous study has shown that translocations can occur independently of cell cycle stage through experimental induction (50), therefore suggesting the presence of a mechanism during mitosis to limit DNA exchange. Notably, direct evidence for the antipairing principle is supported by high-throughput screens that have elucidated candidate genes that antagonize somatic homologous pairing in the fly (51, 52). Spatial segregation of two haploid chromosome sets to discrete nuclear hemispheres may function to keep individual homologous partners apart, preventing, or at least minimizing, allelic misregulation and/or genetic recombination in complex genomes during mitosis. The purpose of this antipairing was indirectly tested when we observed a cancer cell line (53), which exhibited abnormal homologous pairing and consequently lost the haploid set-based antipairing pattern (Fig. 5).

How the haploid set-based groupings emerge still remains an interesting question yet to be answered. Although our data support the paradigm (13, 14, 47) that haploid sets are derived by parental origin, there are limitations to our analysis. The
reciprocal genetic cross with the Ts65Dn male mice to follow the paternally derived supernumerary chromosome proved to be challenging, as they are sterile (5, 54). However, a selective breeding of rare fertile males for unknown reasons were identified (55). The paternally derived translocation chromosome localized to the boundary between two nuclear hemispheres, making it difficult to determine its association to either X- or Y-containing hemispheres (SI Appendix, Fig. S5 C–I). Our study does not rule out the possibility that the supernumerary chromosome does not behave like a wild-type autosome, as it lacks a homologous partner. These rare fertile Ts65Dn male mice may be genetically unstable and exhibit chromosome characteristics more in common with diseased states such as cancer rather than native/endogenous cells. It is intriguing that both the paternally derived supernumerary chromosome (SI Appendix, Fig. S5 H and I), and paired homologous chromosomes in cancer cells (Fig. 5 J and J) are nonrandomly positioned along the boundary between the nuclear hemispheres.

Genome-wide 3D proximity maps and DNA FISH analyses at interphase (G1/G2) have shown a low frequency of interaction between homologs (56–58). These previous studies hint that the haploid set-based antipairing segregation may also be present at interphase, although homologs have been found to come in contact after induction of DNA damage in nonproliferative cells (59). We therefore examined whether the two haploid chromosome sets remain spatially distinct at interphase. However, interphase nuclei often rotate in a variety of cell types (60–62), which may complicate interpretation of chromosome mapping data. To avoid such complexity we examined monocytes, which allowed us to assign decondensed chromosome location along the longitudinal axis of a horseshoe-shaped, interphase nucleus (SI Appendix, Fig. S4 G and H). Three-dimensional overlays showed that both homologous autosomes and XY chromosomes did not segregate to separate nuclear halves (SI Appendix, Fig. S4 G and H). Although our chromosome paint approach is not effective in complete euchromatin detection, the above data suggest
that interphase monocyte nuclei lack a clear hemisphere-based haploid separation.

Another technical limitation/challenge in testing our haploid set-based segregation model in mesenchymal cells is due to the lack of apparent polarity that prevents application of our coordinate system to accurately map chromosome positions between cells. In the future, it would be necessary to reevaluate chromosome mapping in mesenchymal cells including mouse ES cells (25) and immortalized nonepithelial cell lines.

Many possible mechanisms can be implicated for this haploid set chromosome sequestration. These include intrinsic chromosome identity, such as preexisting parental identity of haploid sets, and/or cytoskeletal components to sequester one haploid chromosome set from the other while keeping both sets together without nuclear fragmentation. A recent paper has shown involvement of the cytoskeleton in parental genome segregation at fertilization (26). Although this paper attributes the parental genome segregation to the two mitotic spindles which disappear after the eight-cell stage, it would be interesting to interrogate other cytoskeletal factors that may persist to regulate haploid set segregation. What underlying mechanism(s) must exist for one haploid set to encompass all autosomes and one sex chromosome, however, still remains to be explored. Our study reinforces/renews a foundation to further explore a conserved mechanism in mitotic cells that sequesters haploid chromosome sets along a subcellular axis for antipairing.

Materials and Methods

Cell Preparation and Culture. Primary mouse embryonic fibroblasts were established from harvested E13.5 embryos (63) as described. HUVEC cells (PCS-100-013; ATCC) were cultured in MCDB-131 complete media (Life Technologies) supplemented with LVES (Life Technologies), 1% penicillin-streptomycin, and 1% L-glutamine. Human foreskin fibroblasts (CCLZR211; UCSF Cell Culture Core Facility) were cultured in DMEM H-21 media supplemented with 10% FBS, 1% penicillin-streptomycin, and 1% L-glutamine. The human epithelial cell line (CENP-A/centrin1-GFP RPE1) (24) was a gift from Alexey Khodjakov, New York State Department of Health, Albany, NY. CENP-A/centrin1-GFP RPE1 cells were cultured in DMEM/F12 (1:1) media supplemented with 10% FBS and 1% penicillin-streptomycin. All cells were cultured in a 37 °C incubator in a 10% CO2 atmosphere. For high-resolution imaging, cells were grown on custom diagnostic slides with 8-mm fields (Azerscientific) to 75–80% confluence. Human CD14+ monocytes (PB011; All Cells) were immediately spun down using a CytoSpin (Shandon 4; Thermo Scientific) on Superfrost slides (Fisher Scientific) for use.

Mouse Lines and Genotyping. Female Ts65Dn translocation mice (4–8) and wild-type mice (C57BL/6) were both purchased from The Jackson Laboratory. Ts65Dn translocation females were a gift from Roger Johns Hopkins University, Baltimore. Female/male Ts65Dn mice or wild-type mice were crossed to male/female wild-type mice to generate F1 embryos for collection at E9.5 and E13.5. Genotyping for the chromosome translocation and determination for embryonic sex was employed using PCR on purified genomic DNA obtained from yolk sac samples using previously reported primers (64, 65). All animal protocols were approved by the University of California, San Francisco Institutional Animal Care and Use Committee and in agreement with all institutional and federal guidelines.

Fluorescence in Situ Hybridization. HUVECs, HFFs, and MEFs were grown on slides and human monocytes were grown down on slides for chromosome painting/DNA-FISH as previously described (63). Mouse embryos were fixed at E9.5 with freshly made 4% paraformaldehyde diluted in PBS, pH 7.4. Preparation for embryonic cyrosections was performed as described with modifications to the antigen retrieval step using 10 mM NaCitrate, pH 6 (66). Slides were then processed with the combination of the ethanol series referenced in the above method (63). Slides were denatured/odenatured with Whole Chromosome Paint probes (Applied Spectral Imaging) at 80 °C with 70% formamide/2× SSC for 5 min (for HUVECs/HFFs/human monocytes), 7.5 min (MEFs), or 12 min (E9.5 embryonic mouse cyrosections).

Immunofluorescence. HUVECs were grown on slides, fixed with ice-cold –20 °C 100% methanol for 5 min at –20 °C, rinsed three times with cold PBS, and incubated with permeabilization solution (PFS) (0.7% fish skin gelatin, 0.025% saponin dissolved in PBS) for 1 h at room temperature (RT). Slides were then incubated with primary antibodies to rabbit γ-tubulin (1:1,000, TS26-2000UL; Sigma) and DAPI (1:1,000) diluted in PFS for 1.5 h at RT, washed four times for 5 min each with PFS at RT, and incubated with secondary antibodies to goat anti-rabbit Alexa Fluor 488 (Invitrogen) for 1 h at RT. Final washing was completed at RT with PFS four times for 5 min each and PBS three times for 5 min each then mounted with Prolong Antifade (Invitrogen).

Image Capture and Acquisition. Fixed mitotic cells were imaged with a confocal microscope (TCS SP2; Leica) using a 63× oil immersion objective with a digital 2× zoom. The image data were captured sequentially in a multitrack, four-channel mode. The Z-stacks were acquired using a frame size of 1,024 × 1,024 with a two-frame average and processed with Leica Application Suite (Advanced Fluorescence Lite 2.3.5 build 3379 software).

For high-resolution live imaging, cells were grown on 35-mm glass-bottom dishes (Ibidi). Cells were then placed in an incubation chamber (Okolab enclosure and Bold line incubator) on the motorized microscope stage (Piezo Z-drive) at 37 °C under a 10% CO2 atmosphere and imaged with an inverted fluorescence microscope (Nikon Ti) attached to a spinning disc confocal (CSU-22) using a Plan Apo VC 100×/1.4 oil immersion objective. The image data were captured in a single-channel mode. The Z-stacks were acquired using a frame size of 512 × 512 every 10 s for an hour with an electron-multiplying CCD camera (Evolve Delta; Photometrics) and processed with Nikon Elements (4.5 build). CENP-A/centrin1-GFP was excited with a 488-nm laser (150-mW Coherent OBIS). Brightness and color were adjusted using Photoshop (Adobe Systems).

Mitotic Cell Selection for Fixed Cell Analysis. Prometaphase rosettes/cells which displayed an unbroken radial, symmetrical configuration of chromosomes (and, consequently, a centrosome) were identified as a consistent zone (an absence of TO-PRO3 signal) were analyzed. Rosettes that were asymmetrical, or displayed a distorted shape, were not included in the analysis due to regional variations in DNA condensation within the cell (67). Inclusion of these variations could thus lead to skewed measurements in favor of homologous chromosome randomization. Metaphase cells were identified by linearly aligned chromosomes characteristic of this cell cycle stage (32, 68). Early/mid anaphase cells were identified by incomplete nuclear division (contiguous TO-PRO3 signal between two sister chromosome masses), and by an absence of a constriction characteristic for cytokinesis during late anaphase (67). Typically 0.02% of cells were in early/mid anaphase under our culture conditions. For example, for one experimental group of HUVECs, of 56,547 cells examined 11 mitotic cells were in early/mid anaphase under our culture conditions. For example, for one experimental group of HUVECs, of 56,547 cells examined 11 mitotic cells were in early/mid anaphase under our culture conditions. For example, for one experimental group of HUVECs, of 56,547 cells examined 11 mitotic cells were in early/mid anaphase under our culture conditions. For example, for one experimental group of HUVECs, of 56,547 cells examined 11 mitotic cells were in early/mid anaphase under our culture conditions. For example, for one experimental group of HUVECs, of 56,547 cells examined 11 mitotic cells were in early/mid anaphase under our culture conditions. For example, for one experimental group of HUVECs, of 56,547 cells examined 11 mitotic cells were in early/mid anaphase under our culture conditions. For example, for one experimental group of HUVECs, of 56,547 cells examined 11 mitotic cells were in early/mid anaphase under.
performed by finding the angle between two vectors. A vector was defined as the longest axis of an ellipsoid contained within each homologous chromosome. Therefore, a homologous pair was identified as being paired if the distance was ≤0.36 μm, the spatial resolution limit, and angular orientation = 0°.

Three-Dimensional Centromere Tracking of Time-Lapse Analysis. For live-cell analysis, sample drift correction (translational) was performed computationally for each cell by using the fixed z axis defined by the laser line and the centromere axis as reference points. Three-dimensional centromere movements were analyzed using Imaris software (spot object tracking algorithm) to track individual centromere trajectories over time. Spots were defined by CENP-A GFP signal as 0.5 μm in diameter. A minority of spots (~10%) that overlapped and/or had low signal due to fluorophore bleaching over time were not included in the centromere trajectory analysis.

Three-Dimensional Reconstruction and Overlay for Fixed Cell Analysis. Mitotic cells were individually cropped in each optical section and reconstructed into 3D and analyzed using Imaris software (Bitplane). For fixed cell analysis fluorescent images were thresholded by highlighting the brightest 50% of the image for all channels. Three-dimensional surfaces were then rendered and center of mass for each homologous chromosome was determined.

For 3D overlays, the position of center of mass of homologous chromosomes was mapped to a 3D axial coordinate system for metaphase/anaphase cells. The y axis was defined as a line crossing the furthest edges of the metaphase plate, the z axis was determined as the perpendicular line to the coverslip, and the x axis was defined as perpendicular to both y and z axes. For anaphase cells, the 3D axial coordinate system (Fig. 1L and SI Appendix, Fig. S2A–E) was defined as follows. The x axis was determined by the line crossing the center of mass of the two sister anaphase chromosomes. The y axis was determined as the line perpendicular to the coverslip, and the y axis was defined as perpendicular to both x and z axes.

As no directionality of individual chromosome pairs along the y axis can be predetermined, all X chromosomes or one of homologous chromosomes in closest proximity to y = 0 were given a positive value along the y axis for calculation. A homologous chromosome of a pair was assigned to be either a color/white based on its proximity to the y axis, when y = 0 (a color was assigned to most the proximal, and white the most distal) and mapped to generate the 3D overlay data.

Midpoint Analysis Using a 2D Coordinate System. A 2D coordinate system was established along the y axis, or a line crossing the furthest edges of the metaphase/anaphase cell based on TO-PRO staining. The minimum maximum position along the y axis was based on TO-PRO staining was assigned ±50%, and the center of each mitotic chromosome mass was defined as 0%. The midpoint was calculated as the middle value between the relative positions of each homologous pair along the y axis. Relative positions were determined by using the position of the center of mass of individual homologous chromosomes, divided by the length of TO-PRO signal along the y axis to normalize for cell size.

Three-Dimensional Superimposed Homolog Contour Map Data. Contour maps were generated in ImageJ. The grayscale saturation represents the density of localization when the homologous chromosomes are overlaid in 3D. Isolines were generated with lines that connect contours that share the same gray value (i.e., one contour outlines an area with the same intensity). The 3D superimposed contour map was thresholded by highlighting the brightest 20% of the image for contrast. Individual homologous chromosomes of a pair were labeled and sorted according to proximity to the nuclear meridional plane. The scale bar reflects inverted pixel measurements. White/black regions indicate a level of maximum/minimum localizations, respectively. If the homologous chromosomes were in fixed positions, there would be a clustered area of high density/occurrence that reflects the colocalization frequency for each of the homologs. In addition, the mapped position of each homologous pair would also be identical for every cell.

Two-Dimensional Positional Analysis of Homologous Chromosomes. The 2D coordinate system was established along the y axis as described previously. The relative positions of individual homologous chromosomes were determined using the closest point to the center of the chromosome mass at metaphase, or the mitotic spindle at anaphase, to reflect the centromere position normalized to the TO-PRO staining for the particular mitotic cell. As stated previously, determining the centromere position for individual homologs minimizes variations that could be attributed to an asymmetry in chromosome/rosette condensation. An example for the calculation of the normalization for centromere position at 3.92 μm for a mitotic chromosome mass that spans 8.96 μm along the y axis is (3.92 μm/8.96 μm) - 0.5*100 = -16%.

Statistical Analysis. A rose plot was generated in MATLAB to present the distribution/frequency of angles observed between multiple homologous chromosome pairs of various sizes (SI Appendix, Fig. S1A). The observed values were categorized to 10° intervals (18 total) within a range of 180°. The vector length reflects the number of mitotic cells that falls within each group. For example the expected values for a random distribution for 66 cells would be calculated as 66 cells/18 total 10° intervals = 3.67. A χ² analysis (36) was then performed for the observed values against the expected values of 3.67 for each 10° interval and calculated for significance (P value).

A connected scatterplot graph for frequency distribution shows the relationship between a chromosome and its homologous partner in comparison to a nonhomologous partner (Fig. 1F). Our null hypothesis is that chromosomes have a random distance to their homologous partners, thus reflected in similar distribution to a heterologous chromosome. The values were calculated based on the relative frequency (number of mitotic cells measuring minimal distance at a particular interval/total mitotic cells) for a chromosome compared to its homologous partner or nonhomologous partner (X or Y chromosome). A Student’s t test (36) was performed for the minimum distance values between chromosome 4 and its homolog compared with chromosome 4 to a heterologous chromosome (X or Y) and calculated for significance (P value). Note that 0-0.4 μm were grouped into a single distance interval as it is defined by the limit in confocal microscopy resolution.

The normal probability distribution (35, 36) was employed for statistics wherein the null hypothesis is that sex and homologous chromosome pairs are randomly distributed in two hemispheres of a mitotic cell at metaphase/anaphase. For example, if distributions of chromosomes are random between two hemispheres of equal size in a cell at metaphase/anaphase, the probability of the first chromosome of a given pair’s being positioned in one hemisphere will be 0.5. The probability for the second chromosome’s being positioned in the other hemisphere is 0.5. Therefore, the probability, or P, that an individual of a pair of sex or homologous chromosomes is positioned to two separate hemispheres, or x, which consists of multiple cells, or n, can be calculated by the binomial formula: b(x; n, P) = (n!/(x!(n-x)!)) * P*x * (1– P)* n– x. If the P value is ≤0.05, this would allow for rejection of our null hypothesis. Center of mass calculations were used for hemisphere localization; however, chromosomes that were positioned with the majority of their staining along the boundary of the hemispheres, due to the limits of resolution, were scored to support the null hypothesis, which goes against our model, in the calculation of the one-tailed P value which gives rise to an understimation of statistical significance.

We used linear regression analysis (36) to determine whether there is a relationship between chromosome size and position within the mitotic cell. The hypothesis is chromosome size determines position and can be visualized/represented with a trendline with a slope of 0. Subsequently, we then used a R analysis to test whether the linear regression/trendline between chromosome size (the independent variable) can predict position (dependent variable) within the mitotic cell. The R analysis reveals how close the actual data variation fits the modeled regression/trendline of the mean. For example, R² values of 0 provide no explanation of variation of the mean (i.e., chromosome size does not determine position in mitotic cell).

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