A quantitative and semiautomated method for determining misaligned and lagging chromosomes during mitosis

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ABSTRACT Accurate chromosome alignment at metaphase facilitates the equal segregation of sister chromatids to each of the nascent daughter cells. Lack of proper metaphase alignment is an indicator of defective chromosome congression and aberrant kinetochore–microtubule attachments which in turn promotes chromosome missegregation and aneuploidy, hallmarks of cancer. Tools to sensitively, accurately, and quantitatively measure chromosome alignment at metaphase will facilitate understanding of the contribution of chromosome segregation errors to the development of aneuploidy. In this work, we have developed and validated a method based on analytical geometry to measure several indicators of chromosome misalignment. We generated semiautomated and flexible ImageJ2/Fiji pipelines to quantify kinetochore misalignment at metaphase plates as well as lagging chromosomes at anaphase. These tools will ultimately allow sensitive and systematic quantitation of these chromosome segregation defects in cells undergoing mitosis.

INTRODUCTION Faithful segregation of chromosomes relies on their accurate alignment at the metaphase plate during mitosis (Kuniyasu et al., 2018). Errors in this process result in chromosome missegregation and aneuploidy, which is defined as an abnormal number of chromosomes for a given cell type or organism. Aneuploidy drives the development of cancer, at least in part through promoting chromosomal instability (CIN), a state in which the cell displays a high rate of gain and loss of whole chromosomes (Weaver et al., 2007; Gordon et al., 2012; McGranahan et al., 2012; Storchova, 2018; Soto et al., 2019). CIN is a hallmark of cancer, at least in part through promoting chromosomal instability (Kuniyasu et al., 2018). To prevent the emergence of aneuploidy and CIN, eukaryotic cells align duplicated sister chromatids at metaphase in a bioriented (also known as amphitelic) manner. The process by which chromosomes are transported to the spindle equator, known as chromosome congression, is thought to promote mitotic fidelity by ensuring that chromosomes enter anaphase in a spatially coordinated manner, thereby preventing random chromosome segregation (Maiato et al., 2017). By aligning at the spindle equator, chromosomes are forced to come to a common separation and the subsequent poleward movement from the same relative position. Chromosome alignment also increases the likelihood of kinetochore capture by microtubules from both spindle poles, thus favoring a bioriented geometry for chromosome attachments (Cheeseman, 2014; Joglekar and Kukreja, 2017; Musacchio and Desai, 2017; Hinshaw and Harrison, 2018). Finally, chromosome congression promotes correction of erroneous and unstable attachments; proximity to kinase activity at the spindle poles can result in phosphorylation at the kinetochore that weakens the interaction with spindle microtubules (Barisic et al., 2014; Chmtal et al., 2015; Ye et al., 2015). Therefore, chromosome alignment is critical to the cause of tumor heterogeneity, is associated with poor patient prognosis, and has been linked to drug resistance (McGranahan et al., 2012). To prevent the emergence of aneuploidy and CIN, eukaryotic cells align duplicated sister chromatids at metaphase in a bioriented (also known as amphitelic) manner. The process by which chromosomes are transported to the spindle equator, known as chromosome congression, is thought to promote mitotic fidelity by ensuring that chromosomes enter anaphase in a spatially coordinated manner, thereby preventing random chromosome segregation (Maiato et al., 2017). By aligning at the spindle equator, chromosomes are forced to come to a common separation and the subsequent poleward movement from the same relative position. Chromosome alignment also increases the likelihood of kinetochore capture by microtubules from both spindle poles, thus favoring a bioriented geometry for chromosome attachments (Cheeseman, 2014; Joglekar and Kukreja, 2017; Musacchio and Desai, 2017; Hinshaw and Harrison, 2018). Finally, chromosome congression promotes correction of erroneous and unstable attachments; proximity to kinase activity at the spindle poles can result in phosphorylation at the kinetochore that weakens the interaction with spindle microtubules (Barisic et al., 2014; Chmtal et al., 2015; Ye et al., 2015). Therefore, chromosome alignment is critical to the...
fidelity of mitosis, and approaches to systematically and rapidly identify the degree of chromosome misalignment are required.

Chromosome alignment and congression can be followed using live-cell imaging or in fixed samples by indirect immunofluorescence. Alignment is typically visually scored in both cases through counting displacement of chromosomes from the equator of the mitotic spindle. The result is often provided as a binary readout of either aligned or misaligned cells. Alternatively, the alignment phenotype is subdivided into multiple poorly defined categories. This approach is prone to bias and may lack reproducibility as it depends on the subjective decision of what constitutes misalignment.

Currently, methods to reliably and reproducibly quantify chromosome misalignment and missegregation are limited. One approach described by Lampson and Kapoor (2005) measures and normalizes the distance of individual kinetochores from the nearest spindle pole along the pole–pole axis; while this gives a relative distance for individual kinetochores, this method currently lacks automation (Lampson and Kapoor, 2005). More recently, Fonseca and Stumpff (2016) measured average intensity of each pixel column in the fluorescence channel corresponding to kinetochores along the axis of the metaphase plate and along the entire length of the mitotic spindle (Fonseca and Stumpff, 2016). Here, the output is a distribution profile comprising the intensities of kinetochores along the mitotic spindle. From the Gaussian function fit to this distribution, the full width at half maximum (FWHM) is calculated. Each analyzed cell provides a single FWHM value that can be used to plot a sigmoidal curve comprising the fraction of the cellular population analyzed by FWHM. In this manner, statistical comparisons between sigmoidal curves reflect the comparisons between different conditions in an experiment, giving a global picture of chromosome alignment in the conditions examined.

Here, we developed a method based on analytical geometry to directly quantify misaligned kinetochores at metaphase and lagging chromosomes in anaphase. The method relies on determining the position of the two spindle poles, followed by the establishment of an alignment region based on segmentation of the area between the poles. Kinetochores lying outside a user-defined alignment zone are considered misaligned at metaphase and those lying inside as lagging in anaphase cells. The same approach was adapted to directly measure the distance of individual kinetochores away from a theoretical metaphase plate. We automated the evaluation of chromosome alignment and lagging chromosomes using a series of user-friendly ImageJ/Fiji macros (freely available at https://github.com/Elowesab/elowelab) which can be used to rapidly generate quantitative and robust readouts for these mitotic defects.

RESULTS

Cell geometry visualization

To determine chromosome alignment defects, cells were arrested in metaphase using approaches previously described in the literature (Kapoor et al., 2006; Fonseca and Stumpff, 2016). Briefly, cells were synchronized in mitosis using drugs that interfere with microtubule dynamics or formation of a bipolar spindle (such as nocodazole, or the Eg5 inhibitor S-Trityl-L-Cysteine, STLC). This was followed by release into media containing the proteasome inhibitor MG132, which prevents anaphase onset due to maintenance of high CDK1 activity and persistent sister-chromatid cohesion. Chromosomes were allowed to align at the metaphase plate before being fixed and stained for kinetochores (using CREST antibodies, or antibodies against CENP-C or another kinetochore protein) and spindle poles (γ-tubulin or a centrosome marker) as a minimum. To ensure that only the population of cells where the spindle and spindle poles are found in the same focal plane were sampled, cells were additionally stained with α-tubulin to facilitate visualization of the mitotic spindle (Figure 1A). Cells with compromised chromosome congression were not expected to properly align chromosomes at metaphase, which was manifest as a disorganized chromosome mass (visualized by DAPI or Hoechst staining) and misaligned kinetochores (marked here by CREST, Figure 1A).

Modeling cell geometry

To facilitate modeling of the geometry, cells are spatially translated into a Cartesian plane (Figure 1B). This allows mathematical definition of an alignment zone encompassing the theoretical metaphase plate with kinetochores localized outside this region considered unaligned (Figure 1C). To achieve this, we developed an approach that is broadly divided into three steps: identification of spindle pole position, determination of a user-defined alignment zone, and finally, enumeration of kinetochores outside the defined alignment zone.

After the cell of interest is selected, the method relies on two main inputs from the user required at the beginning of the operation. The first is to identify the position of the two spindle poles in the image (Figure 1D). The significance of the poles lies in the assumption that a theoretical metaphase plate (represented by the metaphase line, green in Figure 1D) is perpendicular to and intersects at the midpoint of the line connecting the two spindle poles (the spindle line, yellow in Figure 1D). To this end, \((x_{P1}, y_{P1})\) and \((x_{P2}, y_{P2})\) represent the coordinates of the first and second spindle poles, respectively. Given the two points, the equation of the spindle line that passes through both poles and thus the midpoint in between the poles can be deduced (Figure 1, E and F). This midpoint represents the centroid of the alignment zone.

The second user-defined input is a pair of values that are used to determine a variable we call the Range. For identifying chromosome alignments defects, the Range essentially corresponds to the alignment region, which is eventually excluded in order to quantify misaligned kinetochores lying outside this zone. Based on the premise that metaphase is a polygon centered on the spindle midpoint (Figure 2A), the user inputs 2 Range parameters, which we call the “total segments” and “aligned segments,” to define the width of the Range relative to the distance between the poles. Here, the value for total segments represents the number of equal segments into which the area between the spindle poles is divided, whereas aligned segments are the number of central segments encompassing the desired Range. In this manner, the line between the poles is divided into equal segments, where the segment(s) in the middle represent the Range (Figure 2B). The values for both Range parameters are arbitrary and user-defined, with some limitation. To ensure that the alignment zone is central, if the spindle length is divided into an odd number of segments, the alignment zone should also be divided into an odd number of central segments. In contrast, if the total number of segments is an even number, the value for aligned segments should also be even. For instance, if the spindle line is divided into five equal segments, the narrowest Range can be represented by the single central segment. A broader metaphase may be desired in which case the three central segments would be considered as the alignment zone, and the value provided for aligned segments would be 3. Likewise, if the spindle line is divided into six equal segments, where the four central sections are considered by the user to be the alignment region, then the values for total segments are 6, and 4 for aligned segments (Figure 2B). The choice for the number of segments in each category would thus allow for precise definition.
FIGURE 1: Defining mitotic cells in a Cartesian plane. (A) Representative images of mitotic cells with aligned and misaligned chromosomes. HeLa S3 cells were fixed and immunostained with CREST serum to mark the kinetochores, γ-tubulin for the spindle poles, and α-tubulin for the mitotic spindle. DNA was stained with Hoechst. The degree of misalignment displays a phenotype that is relatively difficult to quantify. Scale bar = 5 µm. (B) A mitotic cell represented in a Cartesian plane as a function of x- and y-axes. (C) A cartoon representation of the key inputs that are provided by the user: positions of the spindle poles and parameters for identifying the alignment zone (Range, red rectangle) around the metaphase line (dashed green line). (D) Inferring the metaphase line from the position of the spindle poles. The metaphase line is assumed to be perpendicular to the spindle line that passes through the two spindle poles. (E) The equation of the spindle line is deduced from the coordinates of the spindle poles in the Cartesian plane. Spindle pole centroids are indicated by white circles. (F) The midpoint coordinate (green circle) is deduced from the equation of the spindle line.

Setting the Range parameters, the points that delimit the alignment region along the spindle line (i.e., the width of the alignment zone), defined as \((x_1; y_1)\) and \((x_2; y_2)\) in the Cartesian plane, are calculated as a function of the Range based on analytical geometry (Figure 2C; see Materials and Methods).

Defining the alignment zone polygon in the Cartesian plane

Once the points that delimit the alignment zone width along spindle line are determined, the outer limits of this region are defined by establishing the points that delimit the rectangular alignment zone centered at the midpoint (Figure 3A). To do this, lines parallel to the metaphase line and that pass through \((x_1; y_1)\) and \((x_2; y_2)\), defined here as alignment lines, are calculated (Figure 3, B and C). The width (W) along the spindle line of the rectangle is set by the position of the alignment lines, but its length (L) is not a function of the spindle, and here we set them to incorporate the entire cell as metaphase plate dimensions can vary (Figure 3D). By applying the endpoints described above to the equations of the alignment lines, the four points of the rectangle defining the alignment zone are obtained (see Materials and Methods). Because the slope of the spindle line in the Cartesian plane can be positive or negative, the endpoint values on the x- and y-axes of the plane used to determine the limits of the rectangle are set accordingly in the macro described below (Figure 3, D and E). In sum, our geometrical model of the cell requires as minimal inputs by the user the coordinates of the spindle poles if they are being manually defined and arbitrary Range parameters to determine the alignment region.

The method described above classifies kinetochores as either aligned or misaligned. However, the distance between kinetochores and the metaphase line can give an alternative readout for chromosome alignment (Figure 3F). To measure kinetochore distances, the metaphase line is defined as indicated above based on the position of the spindle poles. The kinetochores are then either manually or automatically detected and the shortest distance between individual kinetochores and the metaphase line is computed (see Materials and Methods).

Semiautomated ImageJ macro for quantification of misaligned chromosomes

To implement and automate the above methods for detection of kinetochore misalignment, we created Fiji (ImageJ2) macros to count...
kinetochore misalignment as well as measure distances of individual kinetochores away from the metaphase line. The first step of the macro for kinetochore misalignment is a prompt for the user to open images (single plane or a projected stack) that will be used to identify the spindle poles, kinetochores, and the chromatin in that particular order, which is predetermined to facilitate automation of the method. Next, the user is prompted to select the region/cell of interest (Figure 4Ai). Signals outside of this region are subsequently excluded to ensure that only kinetochores from the cell of interest contribute to the final kinetochore count. Once the region of interest (ROI) is selected, the user is prompted to set the parameters necessary to determine the Range, the number of total segments and aligned segments. The y-axis limit, which defines the length of the alignment zone, is set to a default 15,000 pixels and can be manually changed if desired, depending on the parameters used for image acquisition (Figure 4Aii). Signals outside of this area are subsequently excluded to ensure that only kinetochores from the cell of interest contribute to the final kinetochore count. Once the region of interest (ROI) is selected, the user is prompted to set the parameters necessary to determine the Range, the number of total segments and aligned segments. The y-axis limit, which defines the length of the alignment zone, is set to a default 15,000 pixels and can be manually changed if desired, depending on the parameters used for image acquisition (Figure 4Aii). Signals outside of this area are subsequently excluded to ensure that only kinetochores from the cell of interest contribute to the final kinetochore count. Once the region of interest (ROI) is selected, the user is prompted to set the parameters necessary to determine the Range, the number of total segments and aligned segments. The y-axis limit, which defines the length of the alignment zone, is set to a default 15,000 pixels and can be manually changed if desired, depending on the parameters used for image acquisition (Figure 4Aii).

The next step is to identify the position of the spindle poles. The user is prompted to identify the poles either automatically or manually in the appropriate channel (Figure 4Bi). Automatic detection is accomplished with the built-in ImageJ function Analyze Particles, as indicated in the Materials and Methods (Figure 4Bii), whereas the multi-point function is used for manual selection. The position of the poles as well as Range parameters are then used to determine the alignment zone (Figure 4Ci); the signals inside this zone are deleted with the Clear function in the kinetochore image, and only the objects (misaligned kinetochores) outside the alignment region remain (Figure 4Cii). These kinetochores are then counted automatically with ImageJ/Fiji built-in functions. The user is prompted to evaluate and can manually threshold the remaining signal in the kinetochore channel to facilitate accurate quantification of the number of objects remaining (Figure 4Ciii and iv).

Measurement of kinetochore distances from the metaphase line does not require definition of an alignment zone or Range parameters. Instead, after the poles are identified (either automatically or manually) and the metaphase line computed as above (Figure 4D and E), the user is prompted to choose between manual or automatic kinetochore detection (Figure 4F). Once kinetochores are
Identified, the distance of each kinetochore to the metaphase line is computed (see Materials and Methods).

Evaluation of chromosome alignment in Aurora B and CENP-E inhibited cells

To generate chromosome misalignments and reliably test our pipeline, we took advantage of the phenotypes associated with the inhibition of two key proteins that regulate chromosome alignment during mitosis: the kinase Aurora B and the kinesin CENP-E. Aurora B (the catalytic subunit of the chromosomal passenger complex) localizes mainly to the centromere during early mitosis and can phosphorylate multiple targets at kinetochore proximal centromeres and kinetochores to ultimately regulate the microtubule–kinetochore interaction status (Welburn et al., 2010; Hindriksen et al., 2017; Lampson and Grishchuk, 2017). Moreover, Aurora B activity can destabilize kinetochore–microtubule attachments by inhibiting protein phosphatases responsible for stabilizing microtubule binding (Liu et al., 2010; Foley et al., 2011; Nijenhuis et al., 2014; Meppelink et al., 2015). CENP-E is a processive plus-end motor protein that moves its chromosome cargo along spindle microtubules, thereby promoting congression to the metaphase plate (Schaar et al., 1997; Espeut and Abrieu, 2015; Yu et al., 2019). CENP-E is also thought to promote the formation of end-on stable kinetochore–microtubule attachments to maintain chromosome alignment at the metaphase plate (Shrestha and Draviam, 2013). Inhibition or depletion of either Aurora B or CENP-E in human cells causes alignment defects and lagging chromosomes in anaphase (Yen et al., 1991; Harborth et al., 2001; Ditchfield et al., 2003; Hauf et al., 2003; Tanudji et al., 2004; Cimini et al., 2006; Bennett et al., 2015).

To evaluate the sensitivity of the analysis pipeline, we used it to determine chromosome alignment and metaphase kinetochore distances from the metaphase plate in HeLa S3 cells in which Aurora B or CENP-E were inhibited with ZM447439 or GSK923295, respectively, using the lowest working concentrations commonly found in the literature (See Figure 5A for experimental timeline).
FIGURE 4: Key steps in the execution of chromosome alignment and kinetochore-metaphase line distance macros. (A) (i) The initial step is selection of the ROI in the DAPI/Hoechst channel. Signal outside this selection is deleted from the CREST and γ-Tubulin images. (ii) This is followed by selection of the Range parameters. (B) (i) Selection of the spindle poles is then initiated in the appropriate channel (here, in the image stained for γ-tubulin). (ii) The spindle poles (red triangles) are then detected automatically in Fiji by the built-in function Analyze Particles. The centroid from each area is used as pole coordinates for the mathematical operations. (C) (i) The alignment region is delimited by the dotted lines in the image corresponding to kinetochores. (ii) In the chromosome alignment macro, the signal inside the selection is removed, with only the signal from misaligned kinetochores remaining. (iii) The Threshold function is then used to identify the remaining kinetochores. (iv) These are then counted in the macro using Analyze Particles functions. (D) To measure kinetochore distance from the metaphase plate, ROI selection occurs in the image corresponding to the spindle poles. As in A, signals outside the ROI are deleted in the images corresponding to the poles and kinetochores. (E) Spindle poles are detected as in B. (F) All kinetochores (after thresholding) are detected and their distance from the metaphase line is computed.

Measurement of the distances of individual kinetochores from the metaphase line in cells confirmed an increase in kinetochore distances from the metaphase line of cells treated with either 0.5 μM ZM447439 or 10 nM GSK923295 as expected (Figure 5, B and D). Similar conclusions were reached when the mean kinetochore distance from the metaphase line was calculated for each cell examined (Figure 5C).

To test robustness and versatility, we determined kinetochore misalignment using 3 variations of the Range, each with progressively increasing width (Figure 5E): (i) 1 aligned segment out of 5, (ii) 2 aligned segments out of 6, and (iii) 4 aligned segments out of 6. As expected, low levels of misaligned chromosomes were observed in cells treated with DMSO, while cells treated with either inhibitor presented a marked and statistically significant increase in chromosome misalignment (Figure 5E) for almost all Range values (see below). In addition, the absolute number of misaligned kinetochores counted from the same dataset decreased with increasing alignment zone (Range) size for all conditions tested (compare the values for the same condition among Figure 5E, i–iii). This is as expected, given that a wider alignment zone/Range would result in more kinetochores being considered as aligned, further validating our approach. Moreover, the data also confirm that the choice of values for the total segments and alignment segments inputted determines the extent of misalignment and should be optimized carefully by the user. For example, using a Range of 1 out of 5 and 2 out of 6 revealed differences in chromosome misalignment with higher statistical significance compared with a Range of 4 out of 6 in cells treated with either ZM447439 or GSK923295. Once optimized, the Range parameters should be maintained throughout the entire analysis of any given experiment.
To provide further support for the robustness of the method in assessing diverse degrees of chromosome alignment, we measured the kinetochore distances from metaphase plates and quantified chromosome misalignment in cells treated with increasing concentrations of ZM447439. Using the macro and selecting the Range ratio of 4 aligned segments out of 6, the average number of misaligned kinetochores increased incrementally with increasing doses of ZM447439 until a plateau was reached at 10 µM (Figure 5F). In agreement, measurement of kinetochore distances also demonstrated larger displacement from the metaphase line with increasing doses of the Aurora B inhibitor, at the level of individual kinetochores as well as individual cells (Figure 5, G and H). Collectively, these observations strongly correlate with the effect of these inhibitors on chromosome alignment documented in the literature and validate the methodology presented above.

**Evaluation of lagging chromosomes**

Chromosome lag in anaphase is a well-known consequence of improper kinetochore–microtubule attachments and can result in erroneous segregation as well as damage to lagging chromosomes trapped in the cleavage furrow or micronuclei (Holland and Cleveland, 2012). Quantification of lagging chromosomes usually relies on manual counting (Godek and Compton, 2018), which can be prone to human error and fatigue. We reasoned that translation of the cell into a Cartesian plane and its segmentation can be simply and rapidly adapted (and automated in Fiji/ImageJ2) to measure lagging chromosomes in anaphase. In this context, instead of using the poles to delimit the spindle line, the two anaphase chromosome masses are used (Figure 6A). After selecting the ROI, the user is again prompted to define the Range as above. Next, the center of each chromosome mass in the DAPI/Hoechst image is manually or automatically detected (see Materials and Methods). Once these anchor points have been defined, the same geometric procedure described above is used to delimit the user-defined Range in the center, which now represents the ROI for evaluating the number of lagging chromosomes and is thus retained, whereas the area outside is excluded. To validate this method and the associated macro, we treated HeLa S3 cells entering mitosis with ZM447439 to generate chromosome attachment errors (Figure 6B). Quantification of anaphase-lagging chromosomes in control and ZM447439-treated cells using the macro and the user-defined segmentation of the area between the divided chromosome masses (in this case, a Range of 3 out of a total of 5) confirmed these observations and demonstrated a statistically significant difference in the number of lagging chromosomes between control and ZM447439-treated cells (Figure 6, C and D).

**DISCUSSION**

Here, we used a Cartesian plane model of the cell and analytical geometry to facilitate the quantification of misaligned kinetochores and anaphase-lagging chromosomes, common mitotic defects that are often reported in the literature. Moreover, we have automated the operations in a series of user-friendly, flexible macros (https://github.com/Elowesab/elowelab). We have validated this approach and the associated macros using data acquired from cells treated with Aurora B and CENP-E inhibitors, both of which have been consistently reported to induce perturbations in chromosome alignment at metaphase and segregation at anaphase (Yen et al., 1991; Ditchfield et al., 2003; Tanudji et al., 2004; Cimini et al., 2006). Our results are fully consistent with the literature and confirm the utility of the approach developed here for automated scoring of these defects.

In addition to providing an objective quantification of chromosome alignment errors, one of the key strengths of this approach is the versatility in Range parameters which allows for standardized definition of the alignment zone. Range parameters can be tailored and optimized by the user to help improve interexperiment variation, for example. It is important, however, to retain the same Range parameters within a single experiment to ensure comparable analysis of control and test conditions. Considering that the Range is a constant fraction of spindle length, the inherent variability of different cell sizes in a heterogeneous cell population is accounted for in the execution of the method. These characteristics, together with the semi-automated nature of this methodology, will undoubtedly reduce user errors and increase reproducibility.

We show here also that the method is flexible and can be adapted to measure lagging chromosome in anaphase (Figure 6), an important mitotic defect that is currently only manually curated. Furthermore, although limited to images of fixed cells in this manuscript, this approach can in principle be expanded to analysis of video microscopy of cell division; by determining the poles and kinetochore positioning in each frame from a live-cell movie, the dynamics of kinetochore positioning in relation to a theoretical metaphase plate can also be inferred. If the relative position of each kinetochore is calculated for each time point, it would be possible to evaluate chromosome congression in real time.

One limitation of the approach presented here is that it assumes a symmetric spindle with the alignment region centered between

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**FIGURE 5:** Validation of the chromosome alignment and kinetochore to metaphase line distance operations and macros. (A) Experimental timeline for validation of the chromosome alignment kinetochore to metaphase plate distance pipelines. (B) Distance of individual kinetochores from the metaphase line in cells treated as indicated. Individual data points are shown from three replicate experiments, with 25–142 kinetochores per cell and a minimum of 20 cells measured per condition. Large circles represent the means ± SE. Statistical differences were measured by a repeated-measures one-way ANOVA. Exact P values are shown were determined by a paired t test between control and test conditions. (C) Average kinetochore distance per cell for the data in B. Individual data points are shown from three replicate experiments, with a minimum of 20 cells per condition. Large circles represent the means ± SE of individual experiments and statistical differences were measured by a repeated-measures one-way ANOVA. Exact P values are shown. (D) Representative images of the data are shown in B and C. Scale bar = 5 µm. (E) Measurement of kinetochore misalignment under the conditions indicated using Range parameters of (i) 1 out of 5, (ii) 2 out of 6, and (iii) 4 out of 6. Individual data points are shown from three independent experiments, with a minimum of 25 cells per condition. Large circles represent the means ± SE of individual experiments. Statistical differences were measured by a repeated-measures one-way ANOVA. Exact P values are shown. (F) The number of misaligned kinetochores, (G) kinetochore distance from metaphase line, and (H) average kinetochore distance from metaphase line in cells treated with increasing doses of the Aurora B inhibitor ZM447439. A range of 4 out of 6 was used for evaluation of the alignment errors. Individual data points from two independent experiments are shown, with a minimum of 30 cells per condition. For G, 25–95 kinetochores were examined per cell. Large circles represent the means ± SD of individual experiments.
the two half-spindles and thus misalignment from severely malformed or asymmetric spindles may not be accurately detected automatically, although this may still be possible manually. As with any other automated or semiautomated image analysis methods, reliable immunostaining (of kinetochores and spindle poles in this case) are key to generating reproducible results. However, this is countered by the flexibility offered in detection of the anchor points (spindle poles and the anaphase DNA centroids), which in the macro can be identified either manually or automatically with built-in Fiji functions. This feature may be useful in the absence of pole staining, or in cases where poor staining of centrosome or spindle pole markers in the image makes automatic detection problematic. In this case, pole coordinates could be inferred (albeit less precisely) from mitotic spindle architecture and positioning using α-tubulin staining, for example. Moreover, manual selection of kinetochores and thresholding at key points during pauses in the operation allow for further user optimization, as necessary. In conclusion, the method outlined in this study provides a reliable, semiautomated and versatile tool for quantification of chromosome segregation defects in mitosis and will serve as a useful tool for researchers in the cell cycle and aneuploidy fields.

**MATERIALS AND METHODS**

**Cell culture and immunofluorescence**

HeLa S3 cells were cultured in DMEM (HyClone) supplemented with 10% (vol/vol) of bovine growth serum and Pen/Strep (100 μg/ml, HyClone) at 37°C and 5% CO₂. For measuring chromosome alignment, cells seeded onto coverslips were arrested with STLC (Sigma, 5 μM) for 12 h. Subsequently, cells were released into MG132 (Calbiochem, 20 μM) for 2.5 h and then fixed with PTEMF (0.2% Triton X-100, 20 mM PIPES, pH 6.9, 1 mM MgCl₂, 10 mM EGTA, and 4% formaldehyde). Cells were also treated with either Aurora B

**FIGURE 6:** Adaptation and of the Range operations for the measurement of lagging chromosomes. (A) Steps used by the macro to count the number of lagging chromosomes. After the desired cell is selected and outside signal deleted, the center of each anaphase mass in the chromatin channel is determined using Fiji built-in functions. The lagging region is determined as a function of the Range, and signals outside this are excluded in the image corresponding to the kinetochores. Kinetochore thresholding and detection then facilitate counts of objects/kinetochores within the zone defined by the Range. (B) Experimental timeline for validation of the lagging chromosome macro. (C) Representative immunofluorescence images from the experiment outlined in B. Kinetochores from lagging chromosomes are marked with yellow arrowheads. Scale bar indicate 5 μm. (D) The number of lagging chromosomes per cell in cells treated as indicated. The experiment was assessed with a Range of 3 out 5, which is represented in A. Data points from three independent experiments are shown, with a minimum of 20 cells per condition. Large circles represent the means ± SE of individual experiments. Statistical significance is tested by the paired t test and exact p values indicated.
Macro and image analysis

Image analysis was performed using ImageJ2 (Fiji version 1.52i). The initial steps integrated into the macros provided with this article are as follows: First, on starting the macro, the user is prompted to open the single plane or projected image files of the kinetochore, chromosomes, and spindle poles in a particular order which is retained to facilitate automation. Next, the cell of interest is selected using the Oval selection tool, followed by the deletion of signals outside the cell with the Edit>Clear outside command. The user is then prompted to input the Range parameters: total number of segments and aligned segments as well as the y-axis inferior limit. For quantifying the number of misaligned chromosome (chromosome alignment.im), the selection of each pole is prompted by the macro, either manually or automatically. If the user choses to manually select the poles, the Multi-point tool (right click on the point tool to select multi-point) is used. The macro then initiates the Analyze>Measure command to output the centroid coordinate for each pole. Alternatively, if the user chooses the option of automatic detection of the poles, the macro initiates the Analyze>Analyze Particles command. Once the poles and Range are defined, the four points defining the corners of the alignment region are calculated (see below) and the resulting polygonal selection created in the image corresponding to the kinetochores. Signal from inside the polygon is subsequently deleted with the Clear function resulting in an image containing only the misaligned kinetochores outside the alignment zone. The kinetochore image is filtered using the set Auto threshold function, followed by Make Binary and Watershed. Further manual thresholding at the discretion of the user can be implemented to optimize kinetochore detection. The number of kinetochores/objects and their positions are detected with the Analyze Particles function. As with thresholding, the parameters for the Analyze Particles function can be additionally optimized by the user. The final output from the macro is a summary table with the number of misaligned kinetochores as well as their average size and a second table with a detailed description of each kinetochore identified including the area and position.

For measuring kinetochore distances away from the metaphase plate (kinetochore to metaphase distance.im), the selection of each pole is performed as described in the previous paragraph following the parameters inputted by the user for the Range. The image is then filtered through Process>Filters>Convolve and Image>Adjust>Threshold>MaxEntropy dark, followed by the detection of kinetochores with the Analyze Particles function. The shortest distances of individual kinetochores from the metaphase line are calculated (see below), and the final output are two windows: the first table contains the kinetochore identification number and its distance to the metaphase plate, and the second table lists the distance between the poles.

Finally, in the macro to identify lagging chromosomes in anaphase cells (lagging chromosomes.im), the desired cell is selected and the exterior is deleted as indicated above. Note that for the purpose of this macro, the background needs to be black which can be set under image>color>color picker. The coordinates at the centroids of the two anaphase chromosome masses can be selected manually using the Multi-point tool as indicated above in the image of the chromatin signal. Alternatively, the center of the anaphase chromosome masses is determined automatically by selecting the first mass, excluding the exterior image with Clear, and then using the Threshold function and Create selection to create a selection containing the first anaphase chromatin mass. The Measure function will then output the centroid coordinates corresponding to the center of the chromatin mass. The same procedure is repeated in the macro for the second chromatin mass. After obtaining both coordinates and providing Range parameters, the polygon can be calculated and in the kinetochore image with the built-in Polygon function. The function Clear outside is then used to exclude all signal apart from the kinetochores inside the “lagging region” defined by the polygon. The kinetochores are counted with the Analyze particles function as indicated above for measuring chromosome alignment. The output includes a table with the number of lagging chromosomes, total area, and average size, and a second table lists the area and position of the chromosome masses.

Mathematical deductions

To determine the coordinates of the spindle, the cell is interpreted in a Cartesian plane. Given the two poles where \((x_{P1}, y_{P1})\) and \((x_{P2}, y_{P2})\) represent the coordinates of the first and second spindle poles, respectively, the equation of the line that passes through the poles (i.e., spindle line) is represented by:

\[
y = a_{spindle} x + b_{spindle}
\]

Where the slope of the spindle line \(a_{spindle}\) is given by the equation:

\[
a_{spindle} = \frac{y_{P2} - y_{P1}}{x_{P2} - x_{P1}}
\]

Using this equation, \(a_{spindle}\) is calculated and is then used to determine the slope of the metaphase line \(a\) which is perpendicular:

\[
a = -\frac{1}{a_{spindle}} = \frac{x_{P2} - x_{P1}}{y_{P2} - y_{P1}}
\]

To define the alignment zone, the Range parameters are user-defined as follows:

\[
Range = \frac{\text{Aligned segments}}{\text{Total segments}}
\]

where aligned segments indicate the number of central segments that constitute the alignment zone and total segments are the total number of segments that the area between the spindle poles is divided into. The offset factor \(F\), see Figure 2C) defines the points along the spindle line that demark the boundaries of the alignment zone and is a function of the Range. Thus:
The remaining points of the alignment polygon are subsequently calculated as above.

Finally, to calculate the distance of individual kinetochores away from metaphase line, the shortest distance between kinetochores and the metaphase line needs to be computed. To this end, the metaphase line is defined as:

\[ y = ax + b_{\text{metaphase}} \]

The midpoint, which is the point at which the spindle and metaphase lines intersect, is then calculated and used to determine the metaphase line. Thus:

\[ (x_{\text{middle}}; y_{\text{middle}}) = \left( x_{P1} + \frac{(x_{P2} - x_{P1})}{2}; y_{P1} + \frac{(y_{P2} - y_{P1})}{2} \right) \]

and

\[ b_{\text{metaphase}} = y_{\text{middle}} - ax_{\text{middle}} \]

Therefore, the distance \( d \) of any given point \((x, y)\) reflecting a kinetochore position relative to the metaphase line (Figure 4G) can be calculated as follows:

\[ d = \frac{|ax - y + b_{\text{metaphase}}|}{\sqrt{a^2 + 1}} \]

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