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Wac

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Wac: a new Augmin subunit required for chromosome alignment but not for acentrosomal microtubule assembly in female meiosis

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The bipolar spindle forms without centrosomes naturally in female meiosis and by experimental manipulation in mitosis. Augmin is a recently discovered protein complex required for centrosome-independent microtubule generation within the spindle in Drosophila melanogaster cultured cells. Five subunits of Augmin have been identified so far, but neither their organization within the complex nor their role in developing organisms is known. In this study, we report a new Augmin subunit, wee Augmin component (Wac). Wac directly interacts with another Augmin subunit, Dgt2, via its coiled-coil domain. Wac depletion in cultured cells, especially without functional centrosomes, causes severe defects in spindle assembly. We found that a wac deletion mutant is viable but female sterile and shows only a mild impact on somatic mitosis. Unexpectedly, mutant female meiosis showed robust microtubule assembly of the acentrosomal spindle but frequent chromosome misalignment. For the first time, this study establishes the role of an Augmin subunit in developing organisms and provides an insight into the architecture of the complex.

Introduction

To build a functional bipolar spindle, it is critical to control the site and timing of microtubule polymerization within the cell. The centrosome is the primary site of microtubule nucleation in cells, but it is not the only site where microtubule polymerization is initiated (Luders and Sterns, 2007). A bipolar spindle can be assembled without centrosomes naturally in female meiosis or by experimental manipulation in mitosis (McKim and Hawley, 1995; Khodjakov et al., 2000; Megraw et al., 2001; Basto et al., 2006). New microtubule generation, either by nucleation or rescue, has been observed along spindle microtubules (Mahoney et al., 2006). Tubulin dimers are incorporated at the plus end of microtubules at kinetochores (Mitchison et al., 1986; Maiato et al., 2004). However, it is not fully understood how microtubule polymerization at these various sites is controlled spatially and temporally.

A recent genome-wide RNAi screen using a Drosophila melanogaster cell line (S2 cells) has revealed a full set of genes that contribute to spindle assembly in mitosis (Goshima et al., 2007). Among them, the products of five genes (Dgt2–Dgt6) were very recently shown to form a novel complex called Augmin (Goshima et al., 2008). At least one subunit (Dgt6) is conserved in humans (Goshima et al., 2008; Zhu et al., 2008). RNAi of any subunit of Augmin in S2 cells results in a marked reduction in new microtubule generation and the γ-tubulin signal within the spindle. Further analysis demonstrated that Augmin is responsible for most of the centrosome-independent spindle microtubule assembly in S2 cells.

In a parallel study, we have adopted a new strategy to systematically uncover novel microtubule regulators (Hughes et al., 2008). Microtubule-associated proteins (MAPs) were purified from D. melanogaster embryos, and the microtubule interactome was determined by mass spectrometry. RNAi study of all 83 previously uncharacterized MAPs in S2 cells revealed 13 MAPs essential for proper spindle morphology, including three genes that failed to be detected by the aforementioned genome-wide RNAi approach.

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Abbreviations used in this paper: dsRNA, double-stranded RNA; MAP, microtubule-associated protein; MBP, maltose-binding protein; Wac, wee Augmin component.

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In this paper, we report the study of a previously uncharacterized MAP named wee Augmin component (Wac), which was identified uniquely through our functional proteomics approach. We show that Wac is a new Augmin subunit, which directly interacts with another Augmin subunit, Dgt2, via its coiled coil. We have generated a deletion mutant of Wac, which is viable but female sterile. Unexpectedly, we found that Wac is not required for acrosomal spindle assembly but for chromosome alignment in female meiosis. Therefore, oocytes have an efficient microtubule assembly pathway independent from centrosomes and Augmin but require Augmin specifically for correct chromosome alignment and segregation.

Results and discussion

Wac is a new component of the Augmin complex

In this report, we have studied the products of two genes, CG13879 (that we call wac) and CG16969 (Dgt2), which we identified in our functional proteomics study of the microtubule interactome (Hughes et al., 2008). In our original study, Wac and Dgt2 showed an identical RNAi phenotype and were highlighted as a potential interacting MAP pair through a bioinformatics analysis of large-scale yeast two-hybrid data (Hughes et al., 2008). Dgt2 has also recently been identified in an independent study (Goshima et al., 2008) as part of a new protein complex, Augmin. The estimated size of the complex is larger than the total mass of five known subunits. Therefore, we hypothesized that Wac may be a previously unidentified subunit of Augmin.

To test this hypothesis, we first raised specific antibodies against these two proteins (Fig. S1). The Dgt2 protein has only been studied by epitope tagging (Goshima et al., 2008), and Wac has not previously been characterized. Using these antibodies, we showed that both Wac and Dgt2 bind to microtubules in vitro and uniformly associate with spindle microtubules in S2 cells (Fig. S1).

We then found that Wac and Dgt2, a known Augmin subunit, reciprocally coimmunoprecipitated from an S2 cell extract (Fig. 1 A). We also noticed that the amount of Wac in S2 cells decreased when Dgt2 was depleted by RNAi and vice versa. Furthermore, we showed that Wac levels were also reduced when other known Augmin subunits (Dgt2–6) were depleted from S2 cells (Fig. 1 B). It is already known that depletion of any Augmin subunit generally results in destabilization of other subunits in the complex (Goshima et al., 2008). Therefore, our results strongly suggest that Wac is also an integral component of the Augmin complex.

Wac is required for microtubule generation within the spindle in S2 cells

To establish whether Wac has a similar role to known Augmin subunits, we cytologically examined the RNAi phenotype of Wac in detail. As reported for other Augmin subunits (Goshima et al., 2008), immunostaining revealed a significant increase in cells that have a bipolar spindle with reduced microtubule density and in cells with a monopolar spindle (Fig. 1 C). Live imaging using GFP–α-tubulin showed that control and Wac-depleted cells both initially formed a comparable bipolar spindle (Fig. 1 D and Videos 1 and 2). Then, in control cells, the density of spindle microtubules quickly increased. In Wac-depleted cells, this increase did not take place, and instead, the spindle elongated (Fig. 1, D and E). Furthermore, anaphase onset was significantly delayed in Wac-depleted cells (147 ± 61 min vs. 41 ± 17 min in control, P < 0.01, n ≥ 19; Fig. 1, D and E, arrowheads). Monopolar spindles accumulated as a result of a failure both in centrosome separation and in conversion into a bipolar spindle, as seen in depletion of other Augmin subunits (Goshima et al., 2008), not because of the collapse of a bipolar spindle.

Codepletion of Wac with centrosomin, a protein required for centrosomal γ-tubulin recruitment, severely disrupted spindle assembly (Fig. 1 F). This indicates that Wac, like other Augmin subunits, has a major role in centrosome-independent microtubule assembly. It has previously been proposed that Augmin is required for the localization of γ-tubulin to the spindle (Goshima et al., 2008). Our quantification showed a reduction of the γ-tubulin signal on the spindle after Wac depletion but only to a degree comparable with that of the α-tubulin signal (Fig. 1 G), even at earlier time points (not depicted). Therefore, it is not possible to conclude which is the primary defect, a decreased γ-tubulin localization or decreased microtubule density. Nevertheless, Wac depletion showed similar defects to the depletion of other Augmin subunits.

In summary, Wac physically interacts with a known Augmin subunit, and its stability depends on all other known components of the Augmin complex. In addition, Wac shares functional similarity and microtubule-binding property with other Augmin subunits. Based on these findings, we concluded that Wac is an integral component of the Augmin complex.

Wac directly interacts with a known Augmin subunit, Dgt2, via its coiled-coil domain

We have identified a sixth subunit of the Augmin complex, but neither this nor previous work (Goshima et al., 2008) have so far offered insights into the molecular organization within the complex. To address this question, we tested whether the interaction between Wac and Dgt2 was direct. Bacterially produced maltose-binding protein (MBP)–Dgt2 specifically interacts with Wac translated in vitro using reticulocyte lysate (Fig. 2 A). We further confirmed the direct interaction between Wac and Dgt2 using a wheat germ in vitro translation system (Fig. S2). By using a series of Wac truncations, we mapped the region of Wac responsible for its interaction with Dgt2 to 61 residues containing a predicted coiled-coil region (Fig. 2 A). Systematic replacement of conserved residues to alanines defined four conserved amino acids (LERF) required for this interaction (Fig. 2, B and C).

Wac expression is associated with cell proliferation during development

So far, all studies of Augmin subunits have been performed using cultured cell lines (Goshima et al., 2007, 2008; Hughes et al., 2008; Somma et al., 2008; Zhu et al., 2008). Thorough if not exhaustive searches for mutants defective in mitosis have not yet identified a mutation in any Augmin subunit. Therefore, we decided to study Augmin subunits Wac and Dgt2 in developing organisms.
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(Fig. 3 C). These expression patterns are consistent with a role for Wac and Dgt2 in cell division.

**Wac is dispensable for viability**

To further understand the role of the Augmin complex during development, we generated a clean wac deletion mutant (Δ12); hereafter called wacΔ) by imprecise excision of a P element (Fig. 4 A). We found that homoyzogotes or hemizygotes (over a deficiency) of this deletion mutation are viable. Visual inspection of adult
bodies showed few morphological defects typically seen in mitotic mutants, such as rough eyes, missing bristles, or abnormal cuticle defects, indicating few cell division failures in imaginal tissues. The absence of the Wac protein in wacΔ was confirmed by immunoblots (Fig. 4 B). The amount of Dgt2 was also greatly reduced in wacΔ (Fig. 4 B). Therefore, we concluded that Wac is not essential for viability in D. melanogaster.

wac deletion has a limited impact on chromosome segregation in somatic mitosis and male meiosis

To elucidate the role of Wac in somatic mitosis, we first examined mitosis in the larval central nervous system (CNS) of the wacΔ mutant by chromosome staining. Compared with the control CNS, chromosome condensation appeared to be generally higher in wacΔ, but overcondensation was rarely observed. The frequency of mitosis was significantly higher and the frequency of anaphases among mitotic cells was significantly lower in the mutant (Fig. 4 C), although we found only a low level of aneuploid cells (see Materials and methods). These results indicated that mitotic progression was delayed but not blocked before anaphase onset and that chromosomes are properly segregated in general.

To understand the cause of this mitotic delay, CNSs were immunostained. Unlike RNAi in S2 cells, few monopolar spindles with mitotic chromosomes were observed in the wacΔ mutant, and spindle length did not seem to differ from that in wild type. However, bipolar spindles in the wacΔ mutant appear to have a lower microtubule density (Fig. 4 D). In summary, spindle defects in somatic cells of the wac deletion mutant were milder than those seen after wac knockdown in S2 cells.

Next, we chose male meiosis as a system to test spindle function and the accuracy of chromosome segregation, as it depends on centrosomes but does not have a robust spindle checkpoint (Rebollo and Gonzalez, 2000). Wild-type and wacΔ homozygotes both showed a similarly narrow distribution of nuclear size (2.3 ± 0.1 μm; Fig. 4 E) in spermatids at the postmeiotic onion stage. A genetic test detected 7% of missegregation of the sex chromosomes in wacΔ (0.5% in a control). These results indicate that most chromosomes are properly segregated in male meiosis in the wacΔ mutant. We concluded that loss of Wac has a significant but limited impact on somatic mitosis and male meiosis based on the apparent lack of cell division failure in imaginal tissues and the largely correct segregation of the chromosomes in larval CNSs and testes.

Wac is dispensable for acentrosomal microtubule assembly in female meiosis

We found that the wacΔ mutant was male fertile but female sterile. Mutant adult females developed fully matured ovaries and laid eggs that failed to hatch. In female meiosis, a bipolar spindle forms without centrosomes in a γ-tubulin–dependent manner (Tavosanis et al., 1997). In S2 cells, simultaneously depleting Augmin and an essential centrosome component severely compromised spindle assembly to a degree comparable to that of γ-tubulin depletion (Goshima et al., 2008).

To investigate roles for Wac in female meiosis, we examined the metaphase I spindle in matured oocytes from the wacΔ mutant by immunostaining. Unexpectedly, wacΔ mutant oocytes assembled a robust bipolar spindle (Fig. 5 A). Confirming this observation, quantification of tubulin signal intensity suggested that there was only a small decrease in spindle microtubule density in the wacΔ mutant (1.3 ± 0.8 in wacΔ vs. 1.9 ± 0.9 in wild type, P = 0.05). This is in clear contrast to S2 cells, in which simultaneously depleting Wac and impairing centrosome function severely compromises spindle microtubule assembly.

To probe the spindle architecture, the localization of the pole protein D-TACC (D. melanogaster transforming acidic coiled coil) was examined. Immunostaining showed that D-TACC properly accumulated at both of the acentrosomal poles (Fig. 5 A), confirming that the bipolarity of the spindle is established in the absence of Wac. In addition, some spindles were longer (25% increase in the mean length) and/or had minor morphological
defects (Fig. 5 A), which may be a direct influence of the wac deletion and/or a secondary consequence of chromosome misalignment (see following paragraph).

**Wac is essential for correct kinetochore positioning in female meiosis**

We found that chromosome positioning was disrupted within the bipolar spindle in the majority of wacΔ oocytes (57%; $n = 40$). Meiotic chromosomes were often spread and/or stretched along the spindle in wacΔ rather than being clustered together at the equator region as seen in wild type (Fig. 5 A).

To define the nature of this chromosome misalignment, FISH was used to locate the centromeres of the X chromosome. In wild-type, one dot was found at each opposite edge of the chromosome mass at the equator, representing two pairs of sister centromeres attached to opposite poles (Fig. 5 B).

![Image of molecular biology figures and text](image-url)
In the wacΔ mutant, the number of X centromere dots remained at two, demonstrating that oocytes have a diploid complement and that sister chromatid cohesion has been established and maintained. One X centromere dot was located within each half of the chromosome mass (in 35 out of 40 spindles), indicating that homologous kinetochores were mostly bioriented. However, the mean distance between the X centromere dots doubled in wacΔ (Fig. 5 B), indicating an altered balance of the forces acting on kinetochores.

To see the consequence of this defect, we examined meiotic progression in activated oocytes. We found chromosome missegregation in both meiotic divisions, disorganized meiosis II spindles, and fragmented meiotic products in the oocytes (Fig. S3). We then immunostained older eggs laid by wacΔ mothers. The majority of eggs contained only a few clusters of chromosomes without proliferation of free centrosomes even after being aged for an hour, which is consistent with severe meiotic defects. A few exceptional embryos in which nuclei had divided several times contained chromosomes with misshapen spindles and multiple free centrosomes (Fig. S3). We concluded that Wac is not required for acentrosomal spindle microtubule assembly but for chromosome alignment and segregation in female meiosis.

Dgt2 is concentrated at the polar regions of the acentrosomal meiotic spindle
To understand how Augmin contributes to this acentrosomal spindle function, we attempted to examine the localization of the Augmin subunits Wac and Dgt2 in the female meiotic spindle. Our Wac antibody did not give significant signals above background. Interestingly, the Dgt2 signal was detected at the polar regions of some acentrosomal meiotic spindles (Fig. 5 C). This is in contrast to the uniform staining observed over mitotic spindles in syncytial embryos and S2 cells (Fig. 3 C and Fig. S1).

The essential role of Wac, a new Augmin subunit, in female meiosis
In this study, we have identified Wac as a new subunit of Augmin that directly interacts with Dgt2 via conserved residues in its coiled-coil domain. Augmin was previously shown to be
responsible for centrosome-independent microtubule generation within the spindle in S2 cells (Goshima et al., 2008). In this study, we have defined, for the first time, the role of an Augmin subunit in developing organisms. Unexpectedly, Wac is largely dispensable for somatic mitosis and the bulk microtubule assembly of acientrosomal spindles in female meiosis I but is required for chromosome alignment and segregation in female meiosis. Therefore, unlike mitosis in S2 cells, female meiosis must have an efficient way to generate spindle microtubules in a centrosome- and Augmin-independent manner. We found an increased level of monooriented homologous kinetochores in metaphase I, suggesting some defects in kinetochore-microtubule attachment. However, most homologous kinetochores were bioriented but further apart than in wild type. Assuming that Wac does not affect chromosome elasticity, this phenotype in metaphase I can be explained either by the increase of the pulling force on kinetochores or the decrease of the polar ejection force on chromosomes. We currently favor the latter possibility, as the increase of a force in the absence of Wac is less likely. The polar ejection force is thought to be important for chromosome alignment and generated by microtubule motors or polymerization (Rieder & Salmon, 1994). The accumulation of Augmin at acientrosomal poles may generate polar microtubules important for the polar ejection force. This may take place gradually after spindle formation, which can explain chromosome congression observed during metaphase I (Gilliland et al., 2009).

Materials and methods

Molecular and protein techniques

Standard DNA manipulation and protein techniques were used (Harlow & Lane, 1988; Sambrook et al., 1989). Polyclonal antibodies against Wac and Dgt2 were raised in rabbits or rats using full-length proteins fused to GST or MBP produced in Escherichia coli and were used after affinity purification. In vitro translation was performed using the T7 TNT Quick Coupled system (Promega) in the presence of [35S]-methionine. The Proteins wheat germ extract system (Tobaco) was also used for in vitro translation. For the expression of Wac truncations, relevant regions were amplified, and the T7 promoter was added by PCR. Alkaline substitution constructs were generated using overlapping PCR. Pull-down assays were performed as described previously (Dzhindzhev et al., 2005). S2 cell extracts were prepared by sonication in lysis buffer using 106 cells, and immunoprecipitation was performed as described previously (Lancaster et al., 2007). Microtubule cosedimentation experiments were performed as described previously (Kellogg et al., 1989) after S2 cell extract prepared in BB880 buffer was incubated with 100 μg microtubules prepolymerized in BRB80 buffer containing 20 μM taxol. The developmental Western blot was performed as described previously (Cullen et al., 1999).

Cell culture and RNAi

Culture of S2 cells and RNAi were performed according to published methods (Dzhindzhev et al., 2005; Hughes et al., 2008). Double-stranded RNA (dsRNA) corresponding to F. coli β-lactamase was used as a control.

Fly techniques

Standard fly techniques (Ashburner et al., 2005) were used. w1188 was used as wild type. wac mutants were generated by remodeling of a P element inserted 0.15 kb upstream of the wac gene. The original insertion line (PV05778) was viable and fertile. Imprecise excision events were selected and analyzed over Df(3R)BSC125 by PCR. Three lines (Δ12, Δ44, and Δ53), which delete most of the wac gene without deleting the neighboring genes, are all viable and fertile. wacΔ12 was used in this study. For the genetic test of mis segregation of the sex chromosomes, each wacΔ male carrying a B-marked Y chromosome was crossed with w1188 females. The frequency of mis segregation (i.e., sperm with no or both sexes chromosomes) was estimated from the frequency of non-B males or B females in the progeny (5.3 and 1.6% in wacΔ vs. 0.3 and 0.5% in control sibling heterozygotes).

Cytological techniques

Immunostaining was performed and examined as previously described for S2 cells (Dzhindzhev et al., 2005; Hughes et al., 2008), larval CNSs (Cullen et al., 1999), nonactivated oocytes (Cullen and Ohkura, 2001), activated oocytes (Cullen et al., 2005), and embryos (Cullen et al., 1999) except that an Axio Imager attached to an Exciter (LSM5; Carl Zeiss, Inc.) was used for confocal analysis. Orcein staining of larval CNSs and phase-contrast microscopy of spermatozoids were described previously (Cullen et al., 1999). The frequency of aneuploidy in larval CNSs was estimated by orcein staining of squashed CNSs after incubation with colchicine (3 μg/ml in 0.7% NaCl) and hypotonic shock (in 0.5% sodium citrate). Squashed cells that clearly had six large chromosomes were counted as diploids, whereas cells with greater or less than six were counted as hyperploids or hypoploids, respectively. wacΔ and wild type showed 0.7 and <0.4% of hyperploids and 8.2 and 7.7% of hypoploids (n > 250), although most of the apparent hypoploids were likely to be caused by the squashing of brains (i.e., artifacts). Live S2 cells in the culture media on a concanavalin A– coated coverslip were examined at room temperature by a microscope (Axiovert; Carl Zeiss, Inc.) attached to a spinning-disc confocal head (Yokogawa) using Volocity (PerkinElmer). Immunostaining of centromere identified showed equal segregation of centromeres in most spindles with telophase appearance in Wac-depleted S2 cells. We also see both genuine anaphase and pseudoanaphase cells with chromosomes scattered within a spindle. Intensity of GFP-tubulin signals was estimated by averaging the signal intensity within three equal-sized squares of 0.4 μm2 and subtracting the background signal. The intensities were normalized using the signal intensity at the poles in prophase. Signals of γ- and α-tubulin were measured in cells immunostained with GTU-B8 and YOL1/34 (Sigma- Aldrich). For both signals, the mean intensity in the spindle (mean of three squares of 0.16 μm2) was normalized using the mean intensity at the pole after being background corrected. Significance was analyzed using the Wilcoxon test. For FISH in oocytes, stage 14 oocytes prepared as described previously (Cullen and Ohkura, 2001) were passified with 4% formaldehyde, and hybridization was performed as described in Dernburg et al. (1996) using two Alexa Fluor 488–conjugated 40-mer oligonucleotides corresponding to the 359-bp repeats found at the X chromosome centromere as probes. Digital images were imported to Photoshop (Adobe) and contrast/brightness was changed uniformly across the field.

Online supplemental material

Fig. S1 shows Wac and Dgt2 associate with spindle microtubules in mitosis. Fig. S2 shows that Wac and Dgt2 directly interact. Fig. S3 shows abnormal syncytial embryonic division and meiotic chromosome missegregation in the wacΔ mutant. Video 1 shows an S2 cell expressing GFP-tubulin after control RNAi (Fig. 1D). Video 2 shows an S2 cell expressing GFP-tubulin after wacΔ RNAi (Fig. 1D). Images for both videos were taken every 30 s. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200811102/DC1.

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