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Aurora A inhibition by MNL8054 promotes centriole elongation during *Drosophila* male meiosis

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Aurora A kinase plays an important role in several aspects of cell division, including centrosome maturation and separation, a crucial step for the correct organization of the bipolar spindle. Although it has long been shown that this kinase accumulates at the centrosome throughout mitosis its precise contribution to centriole biogenesis and structure has until now not been reported. It is not surprising that so little is known, due to the small size of somatic centrioles, where only dramatic structural changes may be identified by careful electron microscopy analysis. Conversely, centrioles of *Drosophila* primary spermatocytes increase tenfold in length during the first prophase, thus making any change easily detectable. Therefore, we examined the consequence of the pharmacological inhibition of Aurora A by MLN8054 on centriole biogenesis during early *Drosophila* gametogenesis. Here, we show that depletion of this kinase results in longer centrioles, mainly during transition from prophase to prometaphase of the first meiosis. We also found abnormal ciliogenesis characterized by irregularly growing axonemal doublets. Our results represent the first documentation of a potential requirement of Aurora A in centriole integrity and elongation.

Introduction

The proper separation of sister chromosomes during cell division and their faithful segregation at the opposite poles of the daughter cells require the correct organization of the meiotic and mitotic spindles. The assembly of these microtubule-based structures requires both centrosomal and acentrosomal routes. Although in some systems the centrosome function is redundant when the centrosomes are present they influence the shape and the orientation of the microtubule network. Moreover, besides its role in the organization of the microtubule network, the centrosome also represents an important coordination center of the cell because it contains cell-cycle regulatory, checkpoint and signaling proteins.^{1,2} Control of the centrosome number is essential during the cell life, since numerical alterations represent hallmarks of cell transformation and may contribute to genetic instability.³⁻⁵

Two orthogonally arranged centrioles, small cylindrical organelles with a beautiful and highly conserved 9 symmetry, are found at the heart of each centrosome.⁶ Because the centrioles mark the sites where the centrosomal material is recruited, the centrosome dynamics is closely correlated to the centriole duplication cycle.⁷ Therefore, the centriole duplication has to be accurately regulated so that centrioles and thus the centrosomes duplicate once and only once in even cell cycle, in concert with DNA replication, to

avoid the formation of multiple centrosomes and thus multipolar spindles.⁸⁻¹⁰

Centrioles have a double life and also act as templates for cilia and flagella. During interphase or in quiescent cells the centriole pair migrates to the periphery and the mother one docks to the cell membrane and converts in a basal body that nucleate the ciliary axoneme. Since the primary cilium is implicated in sensing environmental cues and signal transduction pathway, it represents an essential organelle required for animal development and adult homeostasis. Although centrioles and basal bodies represent different functional aspects of the same structure, the basal bodies cannot assemble primary cilia and organize the centrosomal material at the same time. The presence of a primary cilium seems, indeed, to prevent cell division,¹¹⁻¹⁴ although some exceptions have been described.^{15,16} Thus, the reenter in mitosis of vertebrate cells needs an additional control, namely the disassembly of the ciliary axoneme. This process releases the basal body from the plasma membrane that migrates inwards and allows the centriole to organize the functional centrosomes that manage the assembly of the bipolar spindle. Some cell cycle regulatory proteins, and among them the serin-threonine kinase Aurora A, have been implicated in the process of ciliary resorption in proliferating cells.^{17,18}

Aurora A was firstly described in *Drosophila* where the loss of function of this kinase leads to failure of centrosome separation

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and the formation of spindles with abnormally organized poles, including characteristic monopolar spindles.¹⁹ This characteristic phenotype has led to the widely accepted role of Aurora A in centrosome separation, even though the analysis of different model systems reported a range of apparent contradictory defects in the absence of Aurora A.^{20,21} Aurora A activity is also required to control centrosome maturation, mitotic entry, bipolar spindle assembly, chromosome congression, midzone formation at anaphase and cytokinesis.^{18,22} These pleiotropic functions of Aurora A depend on the interaction with different proteins that may modulate its activity.^{23,24}

CALK, a distant Aurora A orthologue, has been reported to control the disassembly of the flagellar axoneme in *Chlamydomonas* after ionic stress,²⁵ suggesting that phosphorylation-based signaling may play a key role in the mechanism of axoneme microtubule depolymerisation. In mammalian cells, the resorption of the primary cilium depends by the interaction of Aurora A with HEF1,²⁶ Pitchfork,²⁷ and calmodulin,²⁸ to activate the histone deacetylase-6 (HDAC-6) and determine the depolymerisation of the axonemal microtubules. Aurora A also negatively regulates primary cilia assembly during mitosis by interacting with the keratin intermediate filament protein trichoplein that is localized in the subapical region of the centriole.²⁹

Based on the importance of Aurora A in ciliary disassembly we ask whether the inhibition of this kinase activity could also lead to structural defects of the centriole that nucleate the ciliary axonemes. Although, Aurora A has been found at the centrosome in a variety of cells,^{30,31} including *Drosophila* embryos,³² no relationships with the centriole biogenesis and organization have been reported.

Since centrioles in somatic cells are very short, eventual defects in their organization may escape to conventional immunofluorescence observations and may be only detectable under careful EM analysis. To circumvent this issue we focused on the structure and dynamics of centrioles/CLRs complexes during *Drosophila* male gametogenesis. The centrioles of *Drosophila* mature primary spermatocytes are 10 times longer than somatic ones thus representing a good model to investigate structural modifications and length variations. In this study, we sought to examine the effects of Aurora A depletion on centriole/CLRs complexes during *Drosophila* spermatogenesis upon incubation in MLN8054, a small inhibitory molecule for this kinase.³³ MLN8054 is a particularly useful biochemical tool in this context, as it has been demonstrated that the Aurora A inhibition occurs rapidly and is more than 150-fold selective for Aurora A over the family member Aurora B in cultured cells.³³

Our data suggest that the Aurora A kinase may be involved in the control of centriole length during *Drosophila* male meiosis.

Results

Previous studies of *aurora* mutations in *Drosophila* have been restricted to the effects on the early mitoses in syncytial embryos obtained by homozygous females and in the neuroblasts divisions in third-instar larval brains. These studies revealed defects in

centrosome separation leading to the formation of typical monopolar spindles.¹⁹ In the attempt to clarify the role of the aurora kinase during the meiotic divisions we analyzed male gametogenesis in the heterozygous *aur²⁰⁹/aur²⁸⁷* pupae. However, we were unable to find a distinct phenotype except a slightly asynchrony of the germ cell divisions within the same cysts that did not affect the normal progression through meiosis. The ultrastructural analysis also failed to reveal specific abnormalities. Since the aurora mutants are weak hypomorphic alleles it is possible that a reduced amount of the protein could be sufficient to allow the proper meiotic progression. Thus, we decided to overcome this limitation by studying the pharmacological inhibition of the aurora kinase during male gametogenesis.

Germ cell line development in *Drosophila* males starts at the tip of the testis with the asymmetric division of the germ line stem cells that produce the primary spermatogonia. Spermatogonia undergo 4 round of mitosis to originate cysts of 16 primary spermatocytes that after 2 successive meiotic divisions form 64 round spermatids. Spermatids then differentiate in elongated sperm cells. Both cell division types rely on the proper organization of a spindle apparatus that supports the correct segregation of the sister chromosomes. Spermatogonia centrioles duplicate once during each cell cycle in concert with DNA replication, like centrioles of somatic tissues. Thus the early spermatocytes inherit at the end of fourth spermatogonial mitosis one centrosome with 2 orthogonally arranged centrioles that duplicate as the germ cells switch from the cell division program to the extended prophase. Each young primary spermatocyte has, therefore, at the beginning of prophase, 2 pairs of short centrioles that moved to the cell periphery to nucleate an axoneme that pushed against the plasma membrane to form a cilium-like region (CLR).

The product of the *uncoordinated* (*unc*) gene, previously identified as a protein involved in the process of centriole/basal body conversion,³⁴ represents a good marker for the centriole/CLR complex.¹⁶ Therefore, we examined the spermatogenesis in flies expressing an Unc-GFP fusion protein to look at the consequences on centriole and CLR organization upon treatment with MLN8054.

According to previous reports Unc-GFP labeling was not detected in asymmetric dividing germ line stem cells, during the spermatogonial mitoses and in early spermatocytes, but first appeared as 2 pairs of small close spots in apolar spermatocytes (Fig. 1A). As prophase progressed the centriole/CLR complexes elongated (Fig. 1B) and reached their full length in mature spermatocytes (Fig. 1C). The Unc labeling was localized in 3 distinct domains: the mid-apical region of the centriole, the CLR and an intermediate dot-like region (Fig. 1B). This distribution remained unchanged during transition from prometaphase (Fig. 1D) to metaphase (Fig. 1E) and persisted through the whole meiotic process.¹⁶

Therefore, the spermatocytes contain 4 ciliary processes, suggesting that all centrioles within the male germ cells have the same potential. Thus, the insect spermatocytes have 4 virtually mature centrioles that nucleate the ciliary axoneme. This is a remarkable condition since it is generally assumed that only the mother centriole is able to nucleate a ciliary axoneme. Vertebrate

cells contain, indeed, one mature centriole, the mother centriole that is able to nucleate a cilium, and an immature centriole, the daughter one that will take full maturation 1.5 cell cycle later.^{35,36}

The distribution of Unc-GFP on centrioles and CLRs of treated spermatocytes did not show significant differences (Fig. 1B–E). However, higher magnification revealed that the centrioles scored from late prophase to meiotic divisions

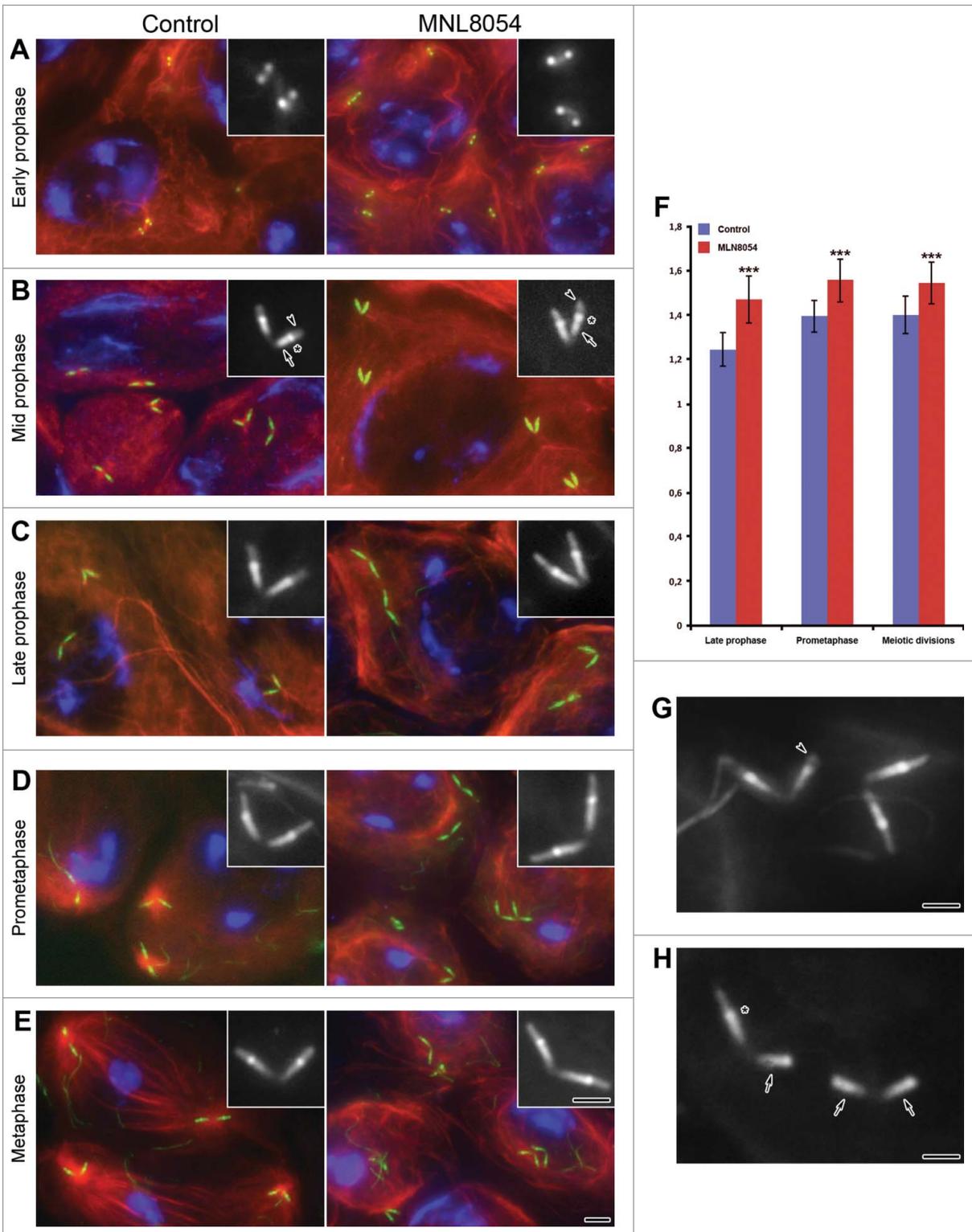


Figure 1. For figure legend, see page 2847.

appeared slightly elongated after incubation in MLN8054 (compare insets on left and right panels of Fig. 1B-E; Fig. 1F).

The centriole/CLRs complexes were found in control prometaphase cells at the focus of large asters that will organize the forthcoming metaphase spindle. By contrast, distinct meiotic spindles never were found upon MLN8054 incubation and the cytoplasmic microtubules were loosely organized around the nuclear region. However, the chromatin condensed in 2–3 discrete masses as found in control prometaphase cells and compacted further in a single cluster, like that observed in metaphase or later control meiotic stages.

At the onset of the meiotic divisions the centriole/CLRs complexes retracted within a membrane pocket to become the poles of the meiotic spindles,^{16,37,38} whereas in treated cells they remained at the cell periphery far from the nuclear region, suggesting a delay in their inward movement. The length of the CLRs slightly decreased in untreated cells during metaphase, whereas this reduction was usually less evident after incubation in MLN8054 (Fig. 1E).

Since, 1 μ M concentration of MNL8054 also slightly affects Aurora B in vertebrate cells,³³ we cannot exclude off target activities of the drug against Aurora B. Therefore, the interpretation of the results involving Aurora A inhibition by MLN8054 may be complicated by low levels of Aurora B inhibition. However, the same phenotypes were also found upon incubation in 0.5 μ M of MLN8054, a concentration that does not inhibit Aurora B activity.

The tripartite distribution of Unc-GFP was usually found in both control and treated cells. However, a low but significant amount of spermatocytes showed a changed pattern of Unc-GFP upon MLN8054 treatment. 8% of the centriole/CLR complexes examined (46, n = 576) displayed abnormally short CLRs (Fig. 1G) and maintained the intermediate dot-like Unc-GFP labeling. 10% of the centriole/CLR complexes (58, n = 576) lost a distinct CLR and the Unc-GFP was localized on the centriole only, lacking the typical intermediate dot (Fig. 1H).

Since the intermediate Unc-GFP has been correlated with the transition region between the centriole and the ciliary axoneme,³⁹ we would ascertain whether the abnormal CLRs observed with conventional immunofluorescence upon Aurora A depletion might reflect ultrastructural defects of the centriole/CLRs.

Both control and treated spermatogonia had one pair of short centrioles composed of 9 triplet microtubules and a central cartwheel (not shown). During spermatocyte grow the centriole pairs

moved to the cell periphery and docked to the plasma membrane. The A- and B-tubules then elongate at the distal end of the centriole and pushed against the plasma membrane to form the ciliary axoneme.

As prophase proceeded both centrioles and CLRs elongated and during mid prophase the CLRs of untreated spermatocytes reached about half length of the centrioles (Fig. 2A). The basal region of the centriole still contained the cartwheel (Fig. 2A₁), whereas the distal lumen was empty (Fig. 2A₂). The hallmark of the axoneme formation in *Drosophila* primary spermatocytes was the reduction at the distal end of the centriole of the C-tubules to hook-like projections that represent cross-sectioned curved longitudinal blades (Fig. 2A₃). This region was also marked by the emergence from the B-tubule of thin radial projections that persisted for all the ciliary length (Fig. 2A₄₋₆). Distinct links usually connected the A and B tubules of adjacent doublets (Fig. 2A₄₋₆), reminiscent of the nexin links found in motile cilia.

79% (n = 56) of the growing primary spermatocytes scored by electron microscopy (n = 71) showed the typical organization of the CLRs seen in controls and characterized by 9 microtubule doublets, lateral C-blades and radial links. However, in some cases (21%; n = 15) we found spermatocytes in which the centrioles reached a length comparable to controls but the CLRs appeared abnormal. The CLRs may be, indeed, present but highly disorganized with doublets incomplete (Fig. 2B) or absent (Fig. 2C). These features were consistent with the observations in immunofluorescence of short (Fig. 1G, inset Fig. 2B) or lacking (Fig. 1H, inset Fig. 2C) CLRs.

The organization of the ciliary axoneme usually reflects the architecture of the centriole. However, while the ciliary axoneme was disrupted, the architecture of the centriole was unchanged in treated cells and a distinct cartwheel was always present (Fig. B_{1,2}; Fig. C_{1,2}). Thus, we asked whether the conversion of the centriole triplets to the axonemal doublets that occurs in the transition region might present eventual defects in Aurora A depleted spermatocytes.

Cross sections thought the transition regions of the abnormal centriole/CLRs complexes showed that these structures had an apparently normal organization compared to untreated controls and also consist of 9 doublets. The reduction of the C-tubules in hook-like projections was only found, however, when the ciliary axoneme was present (Fig. 2B₃). Conversely, the hook-like projections lack (Fig. 2C₃) when the transition region did not continue with the ciliary structures (Fig. 2C₄). Only the few

Figure 1. (See previous page). Aurora-A dependent centriole elongation. Control (left panel) and treated (right panel) primary spermatocytes expressing Unc-GFP were counterstained for acetylated tubulin (red) and DNA (blue). The microtubule cytoskeleton does not show significant differences in control and treated primary prophase spermatocytes (A–C). By contrast, the organization of the cytoplasmic microtubules substantially diverges during prometaphase (D) and metaphase (E) in control and treated spermatocytes: the formers display large asters and bipolar spindles, whereas the others lack both these structures. Magnifications of the centriole/CLR complexes as recognized by Unc-GFP are shown in insets. During early prophase (A) Unc-GFP recognizes small spots in both control and treated spermatocytes. As prophase progressed (B, C) Unc-GFP shows 3 distinct localization: the centriole (arrows, B), the transition region (asterisks, B), the CLR (arrowheads, B). Although the centrioles and the CLRs concurrently elongate during prophase, the centrioles appear slightly longer in late prophase (C), prometaphase (D) and metaphase (E) treated spermatocytes. (F) Quantification of centriole length at different stages of spermatogenesis following 24 hr DMSO (control) or MLN8054 (MLN8054) incubation. Error bars represent SEM. p value from Student's t-test, ***P < 0.0001. Details of abnormally short CLRs (arrowhead, G) or centrioles lacking CLRs (arrows, H); note that the intermediate Unc-GFP dot (asterisk) is only present when the centriole nucleates the ciliary axoneme. Scale bars: (A–E) = 2.5 μ m; (A–E) insets, (G, H) = 1 μ m.

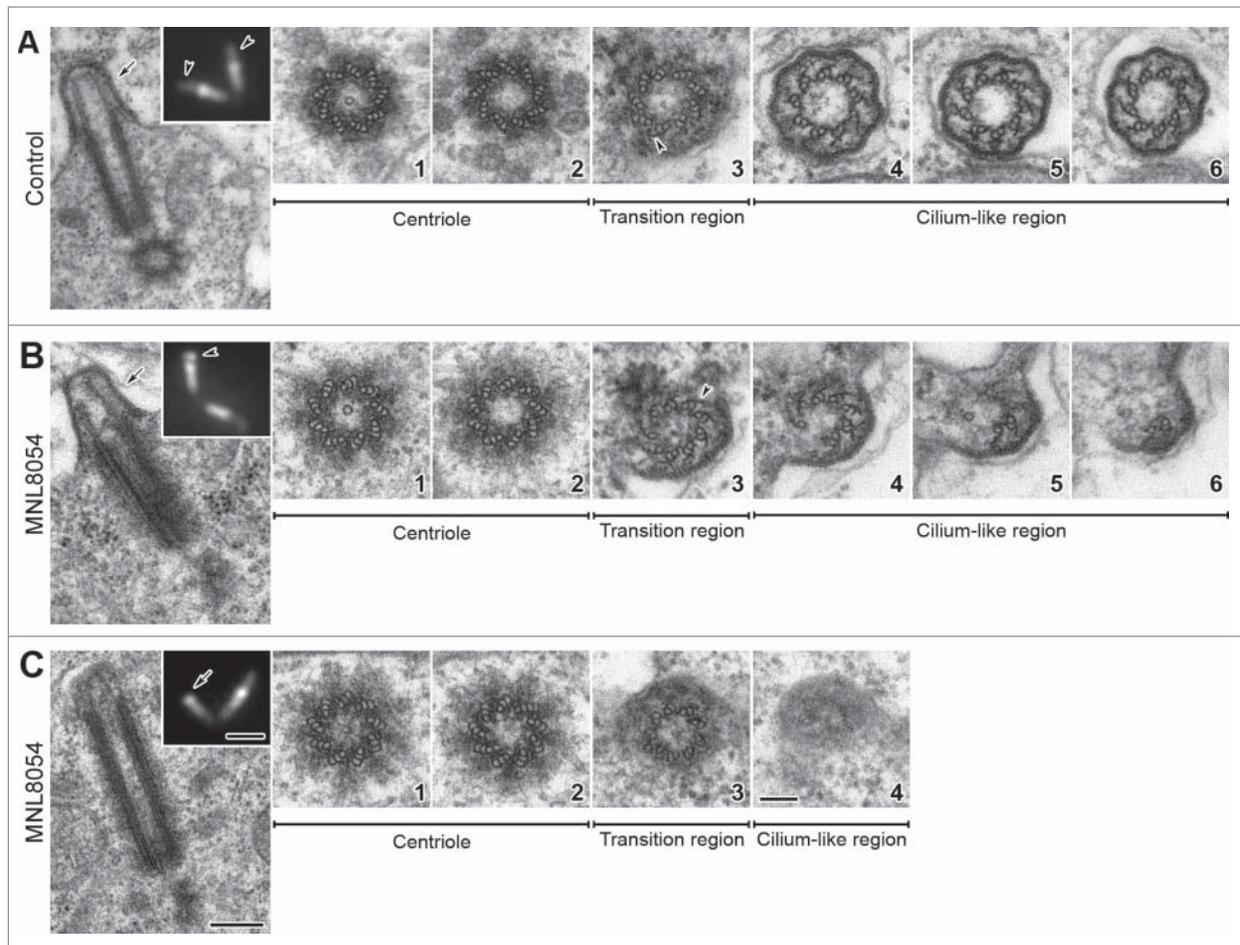


Figure 2. Defects in CLR organization upon MNL8054 incubation. Longitudinal and cross sections of centriole/CLR complexes in control (**A**) and treated (**B, C**) mid prophase primary spermatocytes; insets represent Unc-GFP localization. Control spermatocytes display distinct CLR organization evidenced by Unc-GFP (inset **A**, arrowheads) that protrude from the cell surface (**A**, arrow). Occasionally, treated spermatocytes display abnormal CLR organization (inset **B**, arrowhead) with tubules of different length (arrow, **B**) or elongated centrioles that contact the plasma membrane (**C**), without nucleating the ciliary microtubules (inset **C**, arrow). The centrioles associated with normal (**A**_{1,2}) and abnormal (**B**_{1,2}, **C**_{1,2}) CLR organization display the same architecture; as usual the cartwheel is present in the basal region only. The transition regions display distinct C-blades (**A**₃, **B**₃, arrowheads) when normal (**A**₄₋₆) or reduced (**B**₄₋₆) axonemes are present; C-blades are missing (**C**₃) when the axoneme is lacking (**C**₄). Scale bars: (**A-C**) insets = 1 μ m; (**A-C**) = 250 nm; **A**₁₋₆, **B**₁₋₆, **C**₁₋₄ = 100 nm.

doublets that extended within the abnormal CLR organization maintained, indeed, the remnants of the C-tubules in form of longitudinal blades (Fig. 2B₄₋₆). The links between adjacent doublets were barely detectable only in the proximal region of the short axoneme (Fig. 2B₄).

Centrioles and CLR organization of both control and treated spermatocytes elongated further and reached their full size in late prophase (Fig. 3A). According to Unc-GFP localization (insets in Fig. 3A, B), EM analysis of late prophase (Fig. 3A) and prometaphase (Fig. 3B) spermatocytes confirmed that the centrioles were more elongated in treated than in control cells.

Discussion

Our findings reveal that the inhibition of the Aurora A activity by the small molecule MNL8054 in *Drosophila* primary

spermatocytes results in more elongated centrioles and, to minor extent, in abnormal CLR organization. Therefore, the Aurora A kinase might play a novel role in regulating the centriole biogenesis. However, we find that male gametogenesis is normal in heterozygous *aurora*²⁰⁹/*aurora*²⁸⁷ pupae, whereas the brains of third-instar mutant larvae showed monopolar spindles focused on unseparated centrosomes. The normal meiotic phenotype contrasts with the defects of centrosome separations described in syncytial embryos obtained from homozygous females and in larval neuroblasts.¹⁹ Why should be easier to observe defects in centrosome dynamics during mitotic divisions? One possible explanation could be that there are different requirements upon *aurora* function in mitosis and meiosis. One hypothesis could be that a smaller amount of the kinase *aurora* is needed for progression through male gametogenesis. This may be accentuated by the hypomorphic nature of the *aurora* mutations in which residual function may be sufficient to allow the correct meiotic divisions

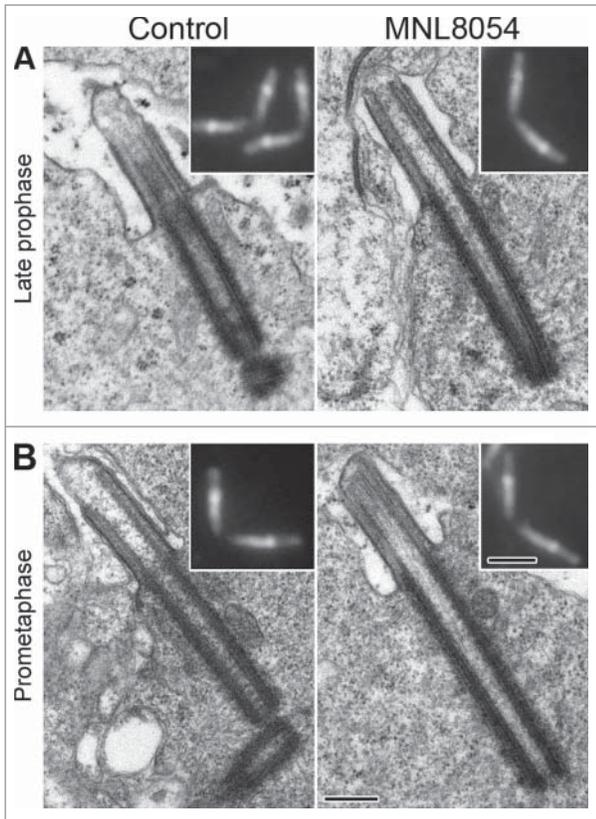


Figure 3. MLN8054 promotes centriole elongation at the onset of meiosis. Longitudinal sections of the centriole/CLR complexes in control (left panel) and treated (right panel) primary spermatocytes during late prophase (A) and prometaphase (B). Insets represent centriole/CLR complexes as recognized by Unc-GFP. Note that centrioles are longer in treated spermatocytes than in controls. Scale bars: insets = 1 μm ; A-B = 250 nm.

of the male germ cells. Treatment with MLN8054 might reduce the residual dose of the protein revealing a meiotic phenotype. However, we cannot appreciate feeble variations in centriole length during the early syncytial mitoses or in larval brains due to the reduced dimensions of the centrioles themselves.

A non-mitotic role of Aurora A kinase in controlling the dynamics of the primary cilia has been recently reported in mammalian cells. Aurora A promotes, indeed, the destabilization of the axonemal microtubules and the resorption of the primary cilium. Depletion of this kinase leads to the inactivation of the tubulin deacetylase HDAC6 resulting in stable primary cilia.^{26–28} However, the centriole/CLRs phenotype observed in *Drosophila* spermatocytes seems to be incompatible with this function. The MLN8054 treatment leaves unchanged the length of the CLRs during prophase-prometaphase, as it also occurs in control spermatocytes in which the Aurora A kinase activity is present. Rather, we observed abnormally reduced CLRs. This inconsistency is likely to stem from the fact that vertebrate primary cilia and *Drosophila* CLRs are similar but not homologous structures. *Drosophila* spermatocyte CLRs diverge, indeed, from

conventional primary cilia by several structural aspects.^{40,41} Moreover, the CLRs persisted during the meiotic divisions until the onset of spermiogenesis when they give origin to the sperm axonemes.^{16,40} By contrast, true primary cilia disassemble when G0 growth-arrested vertebrate cells re-enter the cell cycle.^{42,43} This is a critical process needed to enter cell division.^{12,18,44–47} During cilia disassembly, indeed, the centrioles detach from the plasma membrane to organize the centrosomes that will make the mitotic spindle poles.¹¹

How we can explain that the inhibition of Aurora A in the *Drosophila* primary spermatocytes results in more elongated centrioles? The axoneme of the mammalian primary cilia is nucleate by a basal body which does not change in length through ciliogenesis, when the ciliary microtubules elongate at their distal plus end by an IFT-mediated process. By contrast, both the centrioles and the associate CLRs grow in length concurrently in *Drosophila* primary spermatocytes in the absence of IFT. This raises the question of how the centrioles can elongate while at the same time they templates the assembly of the ciliary axoneme and points to novel dynamics at the distal end of these centrioles. The elongation of the A and B tubules and the conversion of the C-tubules in C-blades in the apical region of the centriole could build the proximal part of the growing axoneme. Since, the centrioles do not elongate in *Unc* mutant spermatocytes in which the aberrant CLRs lack distinct C-blades,⁴⁰ we speculate that the reverse transformation of the C-blades in complete C-tubules may lead to the elongation of the centriole. Such transformation would require a dynamic exchange between microtubule polymerization and depolymerisation. Alteration of this balance by microtubule stabilization may lead to longer centrioles. The CLRs of *Drosophila* spermatocyte are, indeed, directly affected by drugs that target microtubule polymerization. In the presence of nocodazole, a microtubule destabilizer, the axonemes of CLRs failed to assemble, whereas Taxol, a stabilizer, leads to unusually long centrioles and axonemes.⁴⁸ These findings implicate the requirement of factors that control the dynamics of the transition from centriolar triplet microtubules to the axonemal doublets of the CLRs.

Given that Aurora A accumulates at the centrosomes in most of organisms,^{21,40–53} including *Drosophila*,⁵⁴ it is tempting to speculate that this kinase might also regulate the dynamics of the centriole microtubules. Suppression of Aurora-A by small interfering RNA causes an incorrect separation of the centriole pairs in cultured cells, indicating that Aurora-A is essential for the proper execution of this process.⁵⁵ This mechanism might require the phosphorylation of microtubule-related proteins including some involved in microtubule stabilization/depolymerisation.^{56–57} Most known Aurora A substrates are associated with centrosomes,³⁰ and the Aurora A depletion results in disconnection of centrosomes from mitotic spindle poles in *Drosophila*.⁵⁹ Moreover, the Aurora A kinase activity is required to target factors involved in microtubule stabilization at the centrosome.⁶⁰ Aurora A has been reported to play a main function in protecting centrosomal microtubules against depolymerisation by its interaction with TACC complexes in both vertebrate,⁶¹ and

Drosophila cells.³² Aurora A negatively regulates the mitotic centromere-associated kinesin (MCAK),⁶² and supports the assembly of the central spindle at anaphase by promoting microtubule stabilization.⁶³ Aurora A also regulates the Kinesin-13 microtubule depolymerase Kif2A at the spindle poles of mammalian cells during prometaphase.⁶⁴

Several members of the microtubule-depolymerising kinesin-13 family have been shown to regulate the length of cilia in mammalian cells,⁶⁵ and flagella in protozoans.^{66–68} However, the only kinesin-13 demonstrated until now to play a role in centriole elongation is the *Drosophila* Klp10A.⁶⁹ *Drosophila* mutant spermatocytes for *Klp10A* had elongated centrioles that organize irregular CLRs.⁴⁰ Since the centriole phenotypes found in MLN8054 treated cells is strikingly similar to that seen in *Klp10A* mutant spermatocytes, it is conceivable that Klp10A may represent a logical target of the Aurora kinase to regulate the transition of centriolar triplet microtubules to doublets during the formation of the ciliary axoneme.

However, Aurora A promotes microtubule stabilization in mammalian cells by negatively regulating the activity of the microtubule depolymerases.^{20,57,62,70} Thus we should find shorter centrioles in *Drosophila* spermatocytes when the Klp10A activity is up-regulated in the absence of Aurora A, whereas we observed centrioles longer than usual. This discrepancy may be explained with a different function of Aurora A that during *Drosophila* spermatogenesis may activate rather than inhibit the Kinesin-13 microtubule depolymerase Klp10A. Alternatively, since many functions attributed to Aurora A are considered to be in part regulated by the interaction with multiple partner proteins,^{23,52,71,72} additional actors may be work together to Aurora A in regulating the Klp10A activity at the transition between the centriole and the axoneme. This possibility may be consistent with the observation that some centrioles of treated spermatocytes lack the intermediate Unc-GFP labeling and are unable to nucleate axonemal microtubules. The dot-like Unc-GFP has been correlated, indeed, with the transition region. Mutations in *chibby* and *dilatatory* that disrupt this region lead to defects in the organization of the ciliary axoneme.^{39,73}

Further studies are, therefore, required to identify key substrates that may interact with Aurora A kinase to control and regulate the centriole dynamics during male meiosis of *Drosophila*.

Materials and Methods

Drosophila strains

The stock containing the Unc-GFP transgene was described previously.³⁴ The *aur*²⁰⁹ and *aur*²⁸⁷ alleles were reported in Glover et al.¹⁹ Flies were raised on standard *Drosophila* medium at 24°C.

Antibodies and reagents

Mouse anti-acetylated tubulin (1:100) was from Sigma-Aldrich. Alexa Fluor 555 secondary antibody (1:800) was purchased from Invitrogen. The chemical inhibitor to Aurora A (MLN8054) was purchased by Selleck. Dimethyl sulfoxide

(DMSO) and Sang M3 Insect Medium were purchased from Sigma-Aldrich. MLN8054 was dissolved in DMSO at stock concentration of 1000 μ M and stored frozen at 20°C. The stock solution was diluted to the desired concentration in culture medium prior to incubation with testes.

Culture and drug treatment experiments

Testes were dissected from pupae between 5–7 d in M3 medium. To inhibit Aurora A, testes were incubated 24 hours in M3 medium containing 0.5 μ M or 1 μ M MLN8054 for 24 hours into a 24-well plate at 24°C.

Incubation of testes in M3 medium containing DMSO but lacking MLN8054 had no effect on the structure of the centrioles.

Indirect immunofluorescence staining

After incubation, the testes were washed in M3 medium for 10 minutes and then in phosphate buffered saline (PBS) for 5 minutes. Then testes were fixed as previously reported.⁴¹ To visualize microtubules testes were incubated with anti-acetylated tubulin antibody for 4–5 h at room temperature. After washing in PBS–BSA the samples were incubated for 1 h at room temperature with the appropriate secondary antibodies. DNA was visualized with incubation of 3–4 min in Hoechst. Samples mounted in small drops of 90% glycerol in PBS were observed by using an Axio Imager Z1 (Carl Zeiss) microscope equipped with an Axio-Cam HR cooled charge-coupled camera (Carl Zeiss). Grayscale digital images were collected separately and then pseudocolored and merged using Adobe Photoshop 7.0 software (Adobe Systems).

Transmission electron microscopy

Both drug-treated and control testes were carefully rinsed first in M3 medium and then in phosphate-buffered saline (PBS) for 5 minutes. Samples were pre-fixed in 2.5% glutaraldehyde in PBS overnight at 4°C. After washing in PBS, the testes were post-fixed in 1% osmium tetroxide in PBS for 1h at 4°C. Subsequently the material was rinsed again in PBS, dehydrated through a graded series of Ethanol, and embedded in a mixture of Epon-Araldite resin. Serial ultrathin sections (65–75 nm) were prepared with a Reichert ultramicrotome equipped with a diamond knife, collected with formvar-coated copper grids, and routinely stained with uranyl acetate and lead citrate. TEM observations were performed with a FEI Tecnai G2 Spirit transmission electron microscope operating at an accelerating voltage of 100 kV and equipped with a Morada CCD camera (Olympus).

Statistics

Centrioles from individual cysts of both control and treated spermatocytes were scored based on the Unc-GFP labeling. Only the longitudinal centrioles were measured. Centrioles scored were: control prophase (n = 103), control prometaphase (n = 85), control meiotic divisions (n = 84); treated prophase (n = 202); treated prometaphase (n = 138), treated meiotic divisions (n = 235). The error was measured as standard error of the

means (SEM). Significance was measured using the Student's *t* test. For significance ranking values ****P* < 0.0001.

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Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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