

## **PLK4 is a microtubule-associated protein that self assembles promoting *de novo* MTOC formation**

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PLK4, MTOCs, *in vitro* reconstitution, microtubule nucleation, PCM, centrosomes, *de novo*, supramolecular assemblies.

### **Summary statement**

PLK4 binds to microtubules and self-assembles into condensates that recruit tubulin and trigger *de novo* MTOC formation *in vitro*.

## Abstract

The centrosome is an important microtubule-organizing centre (MTOC) in animal cells. It consists of two barrel-shaped structures, the centrioles, surrounded by the pericentriolar material (PCM), which nucleates microtubules. Centrosomes can form close to an existing structure (canonical duplication) or *de novo*. How centrosomes form *de novo* is not known. The master driver of centrosome biogenesis, PLK4, is critical to recruit several centriole components. Here, we investigate the beginning of centrosome biogenesis, taking advantage of *Xenopus* egg extracts, where PLK4 can induce *de novo* MTOC formation (Eckerdt et al., 2011; Zitouni et al., 2016). Surprisingly, we observe that *in vitro*, PLK4 can self-assemble into condensates that recruit  $\alpha/\beta$ -tubulin. In *Xenopus* extracts, PLK4 assemblies additionally recruit PLK4's substrate, STIL, and the microtubule nucleator,  $\gamma$ -tubulin, forming acentriolar MTOCs *de novo*. The assembly of these robust microtubule asters is independent of dynein, similarly to centrosomes. We suggest a new mechanism of action for PLK4, where it forms a self-organizing catalytic scaffold that recruits centriole components, PCM factors and  $\alpha/\beta$ -tubulin, leading to MTOC formation.

## Introduction

Centrosomes are important microtubule organizing centres (MTOCs) in animal cells involved in a variety of processes, including cell motility, division and polarity (Sanchez and Feldman, 2017). They are composed of a core structure, centrioles, surrounded by pericentriolar material (PCM), which nucleates and anchors microtubules (MTs) (Paz and Luders, 2017). PCM proteins also associate with other cellular structures to assemble non-centrosomal MTOCs, a less characterised process (Sanchez and Feldman, 2017).

PLK4, a serine-threonine kinase, triggers procentriole formation close to a pre-existing centriole (the canonical pathway), or induces centriole *de novo* formation when centrioles are absent (*de novo* pathway) (Bettencourt-Dias et al., 2005; Habedanck et al., 2005; Rodrigues-Martins et al., 2007). Recent evidence showed that both pathways use a common but highly conserved set of proteins: PLK4, STIL and SAS6, that are vital players in centriole assembly (Rodrigues-Martins et al., 2007; Loncarek and Khodjakov, 2009; Loncarek and Bettencourt-Dias, 2018) Upon achieving a critical PLK4 concentration threshold at the centriole or in the cytoplasm, PLK4 becomes active (Loncarek and Bettencourt-Dias, 2018; Lopes et al., 2015) determining the place of biogenesis. Then, PLK4 recruits and phosphorylates STIL, which in turn recruits SAS6 to start forming the cartwheel (Zitouni et al., 2016).

Recently, it was demonstrated that PLK4 promotes MT nucleation in acentriolar mouse embryo, being essential for spindle assembly, suggesting it also contributes to acentriolar MTOC formation (Bury et al., 2017; Coelho et al., 2013).

How PLK4 drives *de novo* MTOC formation is not understood. To study PLK4 function in acentriolar systems, we used both *in vitro* systems and acentriolar *Xenopus* extracts, where PLK4 is sufficient to generate *de novo* MTOCs (Eckerdt et al., 2011; Zitouni et al., 2016). We show that *in vitro* PLK4 self-organizes into supramolecular assemblies that recruit tubulin and promote nucleation. In *Xenopus* extracts, PLK4 assemblies recruit STIL,  $\gamma$ -tubulin and  $\alpha/\beta$ -tubulin, forming acentrosomal MTOCs *de novo*. Thus, PLK4 plays an important role in forming both centriole-containing and acentriolar MTOCs. It is important to explore the non-canonical functions of PLK4 to better understand the first events of *de novo* MTOC formation.

## Results

### PLK4 self-assembles into condensates that concentrate soluble tubulin *in vitro*

We wanted to understand how PLK4 drives *de novo* MTOC formation. Recently, Woodruff and colleagues used a minimal set of *C. elegans* proteins to reconstitute a functional MTOC *in vitro*. The PCM scaffold protein, SPD-5, self-assembles into spherical assemblies named condensates, which concentrate homologs of XMAP215 and TPX2, resulting in tubulin recruitment and acentrosomal MTOCs formation (Woodruff et al., 2017).

To explore a minimal system to study PLK4 function, we expressed GFP-tagged *Xenopus* PLK4 in SF9 insect cells using recombinant baculovirus (Fig. S1A). Purified PLK4 is active as shown by western blot using an antibody against threonine 170 in its activation loop, within the kinase domain (Fig S1B) (Lopes et al., 2015). We were surprised to observe that purified active GFP-PLK4 self-assembles *in vitro* into abundant sphere-like structures, when lowering the concentration of NaCl in the buffer (Fig. 1A). The PLK4 sphere-like assemblies were dense at the electron microscope (Fig. 1B and Fig. S1C,D), and recovered very little after photo-bleaching (~ 10% recovery; Fig. 1C), suggesting that, similarly to the SPD-5 spherical assemblies (Woodruff et al., 2017), PLK4 assemblies coarsen and become less dynamic with time, with no internal rearrangement. It is possible that PLK4 assemblies form through multimerization, as different regions of PLK4 can dimerise (Jana et al., 2014). We observed that PLK4 retains its activity while forming these assemblies (Fig. S1D-E). From hereafter, for their similarity with SPD5 assemblies, we will call PLK4 assemblies, condensates.

Given that PLK4's ability to form centrioles in cells and MTOCs in *Xenopus* extract requires its kinase activity (Rodrigues-Martins et al., 2007; Zitouni et al., 2016), we asked whether this was also the case for PLK4 condensates. We found that recombinant GFP-PLK4<sup>AS</sup> (L89A/H188Y), which can specifically fit bulky ATP-analogues, making it sensitive to ATP-analogue inhibitors *e.g.* NAPP1 (Bishop et al., 2000; Zitouni et al., 2016), also self-assembles (Fig. 1D and Fig. S1F). Importantly, as a control, NAPP1 impairs formation of these structures as observed by confocal and electron microscopy (Fig. 1D-F), but not the ones generated by PLK4<sup>WT</sup>, which were only impaired by the PLK4 inhibitor, centrinone (Wong et al., 2015) (Fig. S1F). Instead of robust spherical structures, inactivated PLK4 assembles into an amorphous network with no regular shape (Fig. 1D-F), comparable to PLK4<sup>WT</sup> treated with centrinone or with  $\lambda$ -phosphatase for 1 hour at room temperature (RT) (Fig. S1F-G). Together, this data suggests that PLK4 needs to be in its active conformation to self-assemble into sphere-like structures *in vitro* (Fig. 1D-F and S1).

We next asked whether PLK4 condensates could form an MTOC, by incubating it with  $\alpha/\beta$ -tubulin. We observed that PLK4 condensates can recruit and concentrate  $\alpha/\beta$ -tubulin (Fig. 1G). A full tubulin bleach around PLK4 condensates revealed that tubulin does not recover ( $\sim 10\%$  recovery; Fig. 1H), suggesting they provide a solid platform for immobile tubulin dimers (Fig. 1H), as observed for tau droplets that enhance tubulin assembly (Hernandez-Vega et al., 2017).

### **PLK4 binds microtubules *in vitro***

Given our intriguing observation that PLK4 condensates can recruit  $\alpha/\beta$ -tubulin, we inquired whether PLK4 could bind polymerised  $\alpha/\beta$ -tubulin and promote MT stabilization, which could contribute for its ability to promote MTOC formation. We observed that GFP-PLK4 condensates associate with stable MTs *in vitro* (Fig. 2A). The great majority of PLK4 condensates *in vitro* are associated to MTs ( $\sim 95.4\%$ ) (Fig. 2A-B). We then asked if PLK4 binds directly to MTs. Polymerized MTs pelleted (P) with bound PLK4 whereas the unbound fraction remained suspended in solution (S). We observed that purified PLK4 co-pellets with the MT fraction *in vitro*. To calculate the binding dissociation constant (Kd), we performed pelleting assays with constant PLK4 concentration (0.7  $\mu\text{M}$ ) and increasing MTs concentrations (0 to 4  $\mu\text{M}$ ) (Fig. 2C). Reciprocally, we performed the same assay using constant amount of MTs (10  $\mu\text{M}$ ) and increasing amounts of PLK4 (0 to 4  $\mu\text{M}$ ) until the saturation point (Fig. 2C). We plotted PLK4 bound to MTs versus MT concentration, from three independent experiments. Calculated Kd is the concentration of MTs that is required to sediment half of PLK4 (Fig 2D). These data strongly indicate that PLK4 is a MT-associated protein that binds MTs directly with high-affinity (Kd= 0.62  $\mu\text{M} \pm 0.071$ ). PLK4's presence led to an increase of MT bundles (Fig. 2E-F), as many other MT-binding proteins do (Walczak and Shaw, 2010). As MT bundles stabilize MT dynamics, perhaps PLK4 promotes MT stabilization and nucleation (Brandt and Lee, 1994; Umeyama et al., 1993).

Importantly, PLK4 kinase activity was not required for both  $\alpha/\beta$ -tubulin recruitment (Fig. S2A) and MT-binding (Fig. S2B-C), suggesting they rely on a scaffold but not on the catalytic function of PLK4. A time course using PLK4 and MTs showed that PLK4 binds to MTs independently of PLK4 sphere-like, condensate formation (Fig. 2G). These results show a new role for PLK4 as a MT associated protein involved in MT stabilization and possibly in MT nucleation.

### **PLK4 condensates function as MTOCs in *Xenopus* extracts**

We asked whether PLK4 condensates could promote MT nucleation in the presence of a more complex environment. It was previously shown that PLK4 induces *de novo* MTOC formation after M-phase exit in

*Xenopus* extracts (Eckerdt et al., 2011; Zitouni et al., 2016). We observed that PLK4 condensates, initially formed *in vitro*, can nucleate MTs when *Xenopus* extracts were added, suggesting that they can act as a scaffold forming an active MTOC, given the right factors supplied (Fig. 3A).

Next, we investigated whether GFP-PLK4 would also form assemblies in the more physiological context of *Xenopus* extracts. We used rhodamine-tubulin and EB3-mCherry to visualise GFP-PLK4 induced MT-nucleation (Fig. 3B, Movie 1). PLK4 assemblies were also formed in this context, bearing variable sizes (Fig. 3C), which are akin to centrosome size (300 to 1000 nm; average size ~650 nm). These MTOCs contained GFP-PLK4 at their core, surrounded by tubulin (Fig. 3B), similarly to what we observed *in vitro* (Fig. 1G), suggesting that the same type of assemblies are present *in vitro* and in extracts.

We thus investigated whether PLK4 MTOCs are dynein dependent. Purified centrioles and DMSO asters were used as controls. While DMSO asters are expected to be destroyed in the presence of dynein inhibitors such as vanadate and ciliobrevin, centrioles, as expected, remained capable of nucleating MTs (Fig. S3A). In the case of PLK4-driven MTOCs', asters formed in a dynein-independent manner (Fig. 3D, S3A), suggesting they are indeed self-assembled, and more similar to centrosomes.

Importantly, even in the extract, we observed PLK4's affinity for tubulin. We used interphase extracts to obtain stable asters, using GFP-centrioles, and supplemented with soluble PLK4. We observed some PLK4 associating with MTs (Fig. S3B and Movie 2). However, the majority of PLK4 was localised at the MTOCs centre, as seen *in vitro*, suggesting a core role for PLK4 in MT organisation.

We next examined whether PLK4 condensates could form centrioles in *Xenopus* extracts, using correlative light-electron microscopy (CLEM) (Fig. 3E). Unexpectedly, we observed no centrioles. Instead, PLK4-GFP assemblies corresponded to hollow or sometimes filled spheres of variable sizes, 100 – 600 nm (Fig. 3E-F). These structures agreed with the appearance of PLK4 MTOCs observed by confocal microscopy. We speculate that the formation of PLK4 assemblies that appear hollow in the *Xenopus* extracts, which were not observed *in vitro*, could be due to the presence of additional factors present in this system. Notwithstanding this difference, we conclude that PLK4-MTOCs nucleate MTs, analogous to bona-fide centrosomes.

### **PLK4 MTOCs recruit STIL and $\gamma$ -tubulin in *Xenopus* extracts, leading to MT nucleation**

To further understand PLK4's condensates capacity to form MTOCs, we asked whether their function depends on their ability to recruit other PLK4 partners and/or modulators of  $\alpha/\beta$ -tubulin function. First, we used structured illumination microscopy (SIM) to characterise PLK4-MTOCs in *Xenopus* extracts. We

could observe GFP-PLK4 ring-structure in the crosscuts (Fig. 4A) and sphere-structure after 3D rendering (Fig. 4B, Movie 3).

STIL is PLK4's best characterised substrate and formation of PLK4-STIL's complex triggers centriole biogenesis (Arquint et al., 2015; Kratz et al., 2015; Moyer et al., 2015; Ohta et al., 2014).  $\gamma$ -tubulin is a highly conserved protein, which is associated to most characterised MTOCs (O'Toole et al., 2012; Teixido-Travesa et al., 2012). Most  $\gamma$ -tubulin in animal cells appears to exist as  $\gamma$ -TuRCs and to be important for centrosome-dependent MT nucleation (Wiese and Zheng, 1999). Interestingly, GCP6, a  $\gamma$ -TuRC member, is also a PLK4's substrate (Bahtz et al., 2012; Martin et al., 2014). Thus, we investigated whether STIL and  $\gamma$ -tubulin associate with PLK4-MTOCs by immunofluorescence. We found STIL and  $\gamma$ -tubulin to associate with PLK4 MTOCs in a spatially ordered manner (Fig. 4C-D) so that PLK4 formed an inner layer, followed by STIL, and  $\gamma$ -tubulin (Fig. 4C-D and S4A, Movie 4). Depletion of STIL prevented the formation of PLK4-induced MTOCs in extracts, suggesting that the same pathway triggers the formation of centrioles and acentriolar MTOCs (Fig. 4E-F) (Zitouni et al., 2016). Due to the lack of specific antibodies against *Xenopus* proteins, we were unable to analyse whether other centrosomal proteins associate with PLK4 MTOCs.

#### **PLK4 leads to the formation of stronger centrosomal MTOCs**

PLK4 could promote MT nucleation in different ways also in centrosomal MTOCs. It was recently proposed that MT stabilizers can promote MT nucleation in cells (Roostalu et al., 2015; Roostalu and Surrey, 2017; Wieczorek et al., 2015). The authors proposed that MT stabilizers control the nucleation efficiency by stabilizing the MT centre or "nucleus", either by providing a template for assembly or by promoting longitudinal or lateral tubulin-tubulin interactions. Additionally, by locally recruiting  $\alpha/\beta$ -tubulin, PLK4 assemblies could promote high  $\alpha/\beta$ -tubulin concentrations around the MTOC centre, which is known to lead to enhanced-MT assembly, as it is observed in BuGZ and SPD-5 condensates (Jiang et al., 2015; Woodruff, 2018; Woodruff et al., 2017).

To address this hypothesis, we used GFP-centrin purified centrioles from HeLa cells and incubated them with *Xenopus* extracts. Shortly after addition to interphasic *Xenopus* extract, purified centrioles could recruit PCM components and nucleate MTs. However, when we added GFP-PLK4, we observed a very significant increase in MT nucleation capacity, MT elongation and a decrease in MT dynamics, suggesting their stabilization (Fig. 4G-H; Movie 5 and 6). We observed the same effect in *Xenopus* extracts upon M-phase release (Fig. S4B-C).

## Discussion

The centrosome is an important MTOC in animal cells. Centrosomes can form close to an existing structure (canonical duplication) or *de novo* (Bettencourt-Dias et al., 2005; Habedanck et al., 2005; Rodrigues-Martins et al., 2007). However, the mechanisms underlying *de novo* centrosome formation remain unclear.

In this study, we show that PLK4 self-organizes into condensates that can recruit  $\alpha/\beta$  tubulin. However, this did not lead to MTOC formation, suggesting that other molecules are needed for the observed PLK4's ability to form centrioles and MTOCs in extracts and *in vivo* (Arquint et al., 2015; Baumbach et al., 2015; van Breugel et al., 2011). When PLK4 is added to *Xenopus* extracts it forms condensates, which can further recruit STIL and  $\gamma$ -tubulin, and possibly other components, forming a layered MTOC, similar to centriole-containing centrosomes. Since GCP6 (PLK4's substrate) is required to recruit  $\gamma$ -TuRC to centrosomes, perhaps it can recruit  $\gamma$ -TuRC to PLK4 assemblies (Bahtz et al., 2012; Oriolo et al., 2007; Teixido-Travesa et al., 2012). Additionally, CPAP, a binding partner of STIL involved in MT stabilization, could indirectly enhance MT nucleation from PLK4- MTOCs (Sharma et al., 2016). The ability to mimic the layered centrosome *in vitro* opens new ways of understanding PCM assembly.

In addition, we observed that PLK4 binds to MTs and induces bundling formation. PLK4 kinase activity is not needed for its binding to MTs and recruitment of  $\alpha/\beta$  tubulin but is needed for PLK4 self-assembly into sphere-like structures and for forming an MTOC in *Xenopus* extracts. These observations suggest that PLK4 may have both scaffold and catalytic functions in MTOC formation. Further work is needed to understand those two functions and how they are coordinated. Together, these results suggest a new role for PLK4, in which it binds MTs and serves as a scaffold to concentrate  $\alpha/\beta$ -tubulin, increasing the local tubulin critical concentration, which in turn could promote MT nucleation. Similar conclusions were drawn by a recent study showing similar structures into which tubulin can accumulate, enhancing MT polymerization (Jiang et al., 2015; Woodruff, 2018; Woodruff et al., 2017).

PLK4 forms condensates that recruit several important components in MT nucleation, including  $\gamma$ - and  $\alpha/\beta$ -tubulin. We suggest that this lowers the critical concentration of spontaneous MT nucleation leading to MTOC formation (Fig. 4I). Our data strongly suggests that in the extract, PLK4-driven MTOCs start from PLK4 condensates, and contrary to acentrosomal MTOCs, do not depend on motor based self-assembly of MT minus-end bound material (acentrosomal MTOCs) (Compton, 1998; Mitchison, 1992; Sanchez and Feldman, 2017), where motor proteins, e.g. dynein, play a crucial role in forming acentrosomal MTOCs (Gaglio et al., 1997; Gaglio et al., 1996).



Furthermore, PLK4 MTOCs exhibit a layered organisation, as it was shown to exist in the interphasic centrosome in animal cells (Lawo et al., 2012; Mennella et al., 2012; Sonnen et al., 2012). It is thus possible that, even upon the presence of centrioles, PLK4 could promote the formation of assemblies that concentrate components critical to form centrioles: including STIL,  $\gamma$ -tubulin and  $\alpha/\beta$  tubulin. PLK4 itself, given that it binds and stabilises MTs, can also promote further MT nucleation (Roostalu and Surrey, 2017), hence contributing to daughter centriole assembly. Indeed, we observed that PLK4 enhances centrosomal MTOCs nucleation. These observations are very similar to the ones from Popov and colleagues with XMAP215, a processed MT polymerase that plays an important role in MT-nucleation, in addition to  $\gamma$ -tubulin (Brouhard et al., 2008; Popov et al., 2002). Unlike XMAP215 and like Aurora B (Tsai and Zheng, 2005), PLK4-coated beads do not nucleate MTs in the presence of tubulin (data not shown) below critical concentration ( $< 20\mu\text{M}$ ) (Wieczorek et al., 2015). One explanation could be that PLK4-coated beads alter PLK4 conformation, preventing PLK4-assemblies' formation and blocking MTOC formation. In addition, Aurora B kinase activity is essential for MTOCs formation, identical to what happens with PLK4 (Tsai and Zheng, 2005). Altogether, our observations suggest a mechanism by which PLK4 promotes MT nucleation and stabilization in centrosomal and acentrosomal systems. Further work is needed to understand how PLK4 condensates are formed and whether they are formed at the site of centriole birth, on mother centrioles.

In conclusion, our work shows that PLK4 assembles into condensates, which recruit downstream centrosomal components that are important for both centriolar duplication and microtubule nucleation. These condensates form only in the presence of active PLK4. However, the recruitment of downstream components is independent of the kinase activity. This suggests that PLK4 has both catalytic and scaffold roles in MTOC formation.

Although further work is needed to clarify these mechanisms, this study provides new ways to understand PCM assembly and *de novo* centrosome formation.

## Materials and Methods

### PLK4 protein purification

Full-length *Xenopus* PLK4 gene lacking a stop codon was amplified by PCR and inserted into in-house-designed baculoviral expression plasmids (pOCC series) to generate the following construct: MBP-PreScission::PLK4::mEGFP::PreScission-6xHis; The protein was expressed in SF9 insect cells and harvested 72 hr post infection. Cells were collected, washed, and resuspended in harvest buffer (50 mM Tris HCl, pH 7.4, 150 mM NaCl, 30 mM imidazole, 1% glycerol) + protease inhibitors (1 mM PMSF, 100 mM AEBSF, 0.08 mM Aprotinin, 5 mM Bestatin, 1.5 mM E-64, 2 mM Leupeptin, 1 mM Pepstatin A)(Calbiochem) and frozen in liquid nitrogen. The protein was purified using a two-step purification protocol described in Woodruff, J. B., & Hyman, A. A. (2015)\*. PLK4 clarified lysate was incubated first with Ni-NTA agarose beads followed by a second incubation with amylose resin. The MBP and 6xHis tags were cleaved and PLK4 was eluted by overnight incubation with PreScission protease. PLK4 was concentrated with a 50K Amicon Ultra centrifugal concentrator units (Millipore), aliquoted and flash frozen in liquid nitrogen. The lysis and final elution buffer used (called “PLK4 buffer”) contained 50 mM Tris-HCl, pH 7.4, 500 mM NaCl, 0.5 mM DTT, 1% glycerol, 0.1% CHAPS. The intermediate elution buffer from the Ni-NTA beads contained additional 250 mM imidazole (Woodruff and Hyman, 2015). As a control, we used the construct mEGFP::6-His::Z-tag::TEV::SBP to express GFP, which is purified as previously mentioned.

### *In vitro* PLK4 self-assembly

PLK4 assemblies were formed by adding purified GFP-PLK4 (1  $\mu$ M) to the condensate buffer (150 mM NaCl, 25 mM Hepes (pH 7.4) and 1 mM DTT). PLK4 was incubated for 5 to 10 min at room temperature (or at different temperatures when checking their effect on PLK4 assemblies (Fig. S2C)) and then imaged using a spinning disk CSU-X1 (Yokogawa) confocal scan head coupled to a Nikon Eclipse Ti-E and controlled using MetaMorph 7.5 (Molecular Devices). For tubulin recruitment, tubulin-labelled rhodamines TRITC (Cytoskeleton) (500 nM) were added to the condensate buffer. We used BRB80 as a control. For the experiments using inactive PLK4, NAPP1 (100  $\mu$ M) was used to inactivate PLK4<sup>AS</sup>; DMSO was used as a control. PLK4's dephosphorylation was performed using the typical reaction protocol for  $\lambda$ -phosphatase, incubated for 1 hour at room temperature and processed either for confocal imaging or for EM. A sample was taken to check PLK4's activity by western blot using a Threonine-170 rabbit antibody raised in our laboratory.

### **Microtubule pelleting assay**

Tubulin was polymerized into MTs stabilized with 20  $\mu$ M taxol in BRB80 buffer (25 mM HEPES, pH 6.8, 2 mM MgCl<sub>2</sub>, 1 mM EGTA, 0.02% Tween 20 (v/v)), and quantified by absorbance measurements at 280 nm. Various concentrations of MTs were mixed with constant concentrations of PLK4 in BRB80 Buffer or vice-versa. Samples (final volume=40  $\mu$ l) were equilibrated at 37°C for 25 minutes, centrifuged in Airfuge at 90 000 rpm for 30 minutes, and both the supernatant (S) and pellet (P) collected and resuspended in SDS sample buffer, and equal amounts of supernatant and pellet were run on 4–20% Tris-HCl gradient gels (Bio-Rad). Gels were stained by Coomassie Blue or blotted for western blot to detect MTs and PLK4. Quantification of the relative amounts of PLK4 in supernatants and pellets was performed using ImageJ (National Institutes of Health, Bethesda, MD). The dissociation constants measured by MT co-sedimentation represent the average and propagated error from three independent experiments.

### ***In vitro* PLK4 and microtubule bundling on confocal assay**

Taxol-stabilized MTs seeds were incubated with GFP-PLK4 (1  $\mu$ M) for 15 minutes at 37°C and then mounted in a slide and observed by confocal microscope. Taxol MT seeds were done as previously described (Honnappa et al., 2009). Briefly, lyophilized 1mg of tubulin (Cytoskeleton) was resuspended in 100  $\mu$ l of BRB80 buffer, 1mM GTP and 4 mM MgCl<sub>2</sub> and incubated for 30 minutes at 37°C. After 20 minutes, taxol was added to a final concentration of 20  $\mu$ M. Finally, taxol-MTs were stored at room temperature.

### **Fluorescence recovery after photo bleaching (FRAP)**

Tubulin recruitment by PLK4 assemblies was performed as described before (Woodruff et al., 2017). For the FRAP analysis of the *in vitro* PLK4 assemblies (alone or coated with tubulin), samples were mounted in an imaging chamber after 10 min of incubation at RT. FRAP experiments were performed on the same spinning disk confocal microscope mentioned above using a 100x 1.49 NA oil immersion objective. Laser micro-irradiation was performed with a 491 nm or 561 nm laser with a 100 ms exposure time. Tubulin coating PLK4 was bleached for 15 ms and images were taken at 0.4 s intervals. Analysis of the recovery curves and the half-time recovery were carried out with the FIJI/ImageJ macro ([http://imagej.net/Analyze\\_FRAP\\_movies\\_with\\_a\\_Python\\_script](http://imagej.net/Analyze_FRAP_movies_with_a_Python_script)). The normalized data was parsed and plotted using R.

### **Electron microscopy negative staining assay**

For the electron microscopy assays, GFP-PLK4<sup>WT</sup> or GFP-PLK4<sup>AS</sup> was used to visualize the structures when the proteins were incubated in different buffers and at different conditions (PLK4's buffer, BRB80, with or without DMSO, in the presence or absence of microtubules and incubated at different temperatures). A time-course (0 to 10 min) was also performed to observe the evolution of the assemblies. For acute inactivation of PLK4, NAPP1 or centrinone (PLK4 inhibitor) was added simultaneously with GFP-PLK4<sup>AS</sup> or GFP-PLK4<sup>WT</sup>, respectively. When PLK4 was mixed with microtubules, the incubation was performed at 37°C for 30 minutes. After the reaction, samples were adhered to glow discharged copper 150 mesh grids coated with 1% (w/v) formvar (®Agar Scientific) in chloroform (®VWR) and carbon. Following attachment, samples were rinsed with distilled water and stained with 2% (w/v) uranyl acetate. Electron microscopy images were acquired on a Hitachi H-7650 operating at 100 keV equipped with a XR41M mid mount AMT digital camera.

### **Preparation of *Xenopus Laevis* egg extracts and MTOC formation assay**

MII-arrested and interphase egg extracts were prepared as previously described (Lorca et al., 2010; Zitouni et al., 2016). Purified PLK4 (0.675 nM) was added to 20 µl of CSF extracts and released into interphase using calcium (20 mM). MTOCs were analysed by using rhodamine-labelled porcine tubulin (Cytoskeleton). Depletion of STIL in *Xenopus* extracts was performed as described in (Zitouni et al., 2016).

**Assays using GFP-centrin labelled centrosomes:** centrosomes were added to interphasic extract or to MII-arrested extract containing rhodamine-labelled tubulin in the presence or the absence of GFP-PLK4 (1 µM). These extracts are incubated for 30 min (interphasic extract; Fig 4H) and 15 min (MII released extract; Fig S5B) at 18°C and visualised by confocal microscope. When GFP-centrin centrosomes were added in the extract in the presence or absence of PLK4, we measured the length of MTs (Fig. 4I) or the intensity of MT nucleation via rhodamine-tubulin intensity (Fig. S5C). Prism (version 5.0c; GraphPad) was used for statistical analysis and plotting when was needed. To visualize PLK4 binding to MTs, as previously done, centrioles labelled with GFP-centrin were added to interphasic extracts and supplemented with more soluble GFP-PLK4 (1 µM).

## Antibodies

Antibodies against STIL (Human SIL Antibody, A302-441A, Rabbit anti-SIL Antibody, Affinity Purified; dilution 1/1000) and  $\gamma$ -tub (GTU-88, T5326 Sigma; dilution 1:500) were purchased from Bethyl Laboratory and Sigma, respectively. Anti-xCEP192 (dilution 1:500) polyclonal antibodies were generated by immunizing rabbits with the purified fragments of *Xenopus* Cep192 [aa 521–1000 ( $\alpha$ Cep192-N)]. Antibodies were affinity-purified by using the relevant antigens cross-linked to AminoLink Coupling Resin (Thermo Scientific) according to the manufacturer's instructions) was a gift from V. Joukov (Department of Dana-Farber Cancer institute, Boston) and XPLK4 was used in (Zitouni et al., 2016). Rabbit Phospho-Thr170 antibody (dilution 1:500) was produced by immunizing rabbits with a synthetic peptide against the activation T-loop ((K)HYpTLCGTPNY) of the human sequence. Antibodies against Histidine (dilution 1:1000) (Novagen Material Number: 70796-3, Kit Batch Number: D00106784) and GFP (dilution 1:1000) (Roche Applied Science; catalog: 11814460001, Clone name 7.1, 13.1) were purchased from Novagen and Roche, respectively.

## Super-resolution of PLK4 assemblies in *Xenopus* extracts

Structured Illumination Microscopy (SIM) of PLK4-GFP assemblies was performed on N-SIM, Nikon Inc., equipped with Apo TIRF 100x NA 1.49 Plan Apo oil objective, back-illuminated EMCCD camera (Andor, DU897), and 405, 488, 561 and 640 nm excitation lasers. 100 nm Z sections were acquired in 3D SIM mode generating 15 images per plan, and reconstructed. XYZ corrections were applied using the signals of 100 nm multi-spectral fluorescent spheres (Invitrogen) included in the sample.

## Correlative light and electron microscopy

To correlate light and electron microscopy images of PLK4-GFP assemblies, *in vitro* PLK4-GFP self-assembly reaction mixture was overlaid directly to the coverslips mounted in Attofluor Cell Chambers (Invitrogen; A7816) and kept at 37°. The coverslips contained previously sparsely seeded and fixed HeLa cells. HeLa cells served as landmarks for subsequent identification of target PLK4-GFP assemblies during trimming, sectioning, and imaging on electron microscope. PLK4-GFP MTOCs were fixed in 2.5% glutaraldehyde, and immediately imaged on an inverted microscope (Eclipse Ti; Nikon, Tokyo, Japan) equipped with a spinning-disk confocal (CSUX Spinning Disk; Yokogawa Electric Corporation, Tokyo, Japan), back-illuminated 13  $\mu$ m pixel EMCCD camera (Andor, DU888), 100x NA 1.42 Plan Apo objective lens with 1.5x magnifying tube lens, and a 2x lens in front of the confocal head. The positions of GFP-

PLK4 assemblies were recorded by acquiring a stack of 200 nm-thick Z sections in fluorescent mode and then in DIC (using Nikon DS-U3 camera). The position of the target PLK4-GFP assemblies and fiducial cells on the coverslip was marked by a diamond scribe, as described earlier (Kong and Loncarek, 2015). After fixation, the samples were washed in PBS for 30 min, pre-stained with osmium tetroxide and uranyl acetate, dehydrated in ethanol, and then embedded in Embed 812 resin. 80 nm thick serial sections were sectioned, transferred onto the formvar coated copper slot grids, stained with uranyl acetate and lead citrate, and imaged using a transmission electron microscope (H-7650; Hitachi, Tokyo, Japan) operating at 80 kV. For the alignment of serial sections and for image analysis we used Photoshop (Adobe) and Fiji (NIH).

## Author Contributions

S.M.G purified the proteins, performed the microtubule assays, the EM and the SIM experiments with the participation of P.D, E.T., A.L, D.K and J.L. The CLEM experiments were performed by D.K and J.L. S.Z. performed the aster formation assays in extracts, the EM, the FRAP experiments and the PLK4 assembly formation assays with the participation of B.R.G. and T.H. B.R.G. analyzed the FRAP data. S.M.G and S.Z wrote the manuscript and made the figures. All authors read, discussed and approved the manuscript. S.M.G. and M.B.D. conceived the study and were in charge of overall direction and planning.

## Competing interest statement

The authors declare no competing interests.

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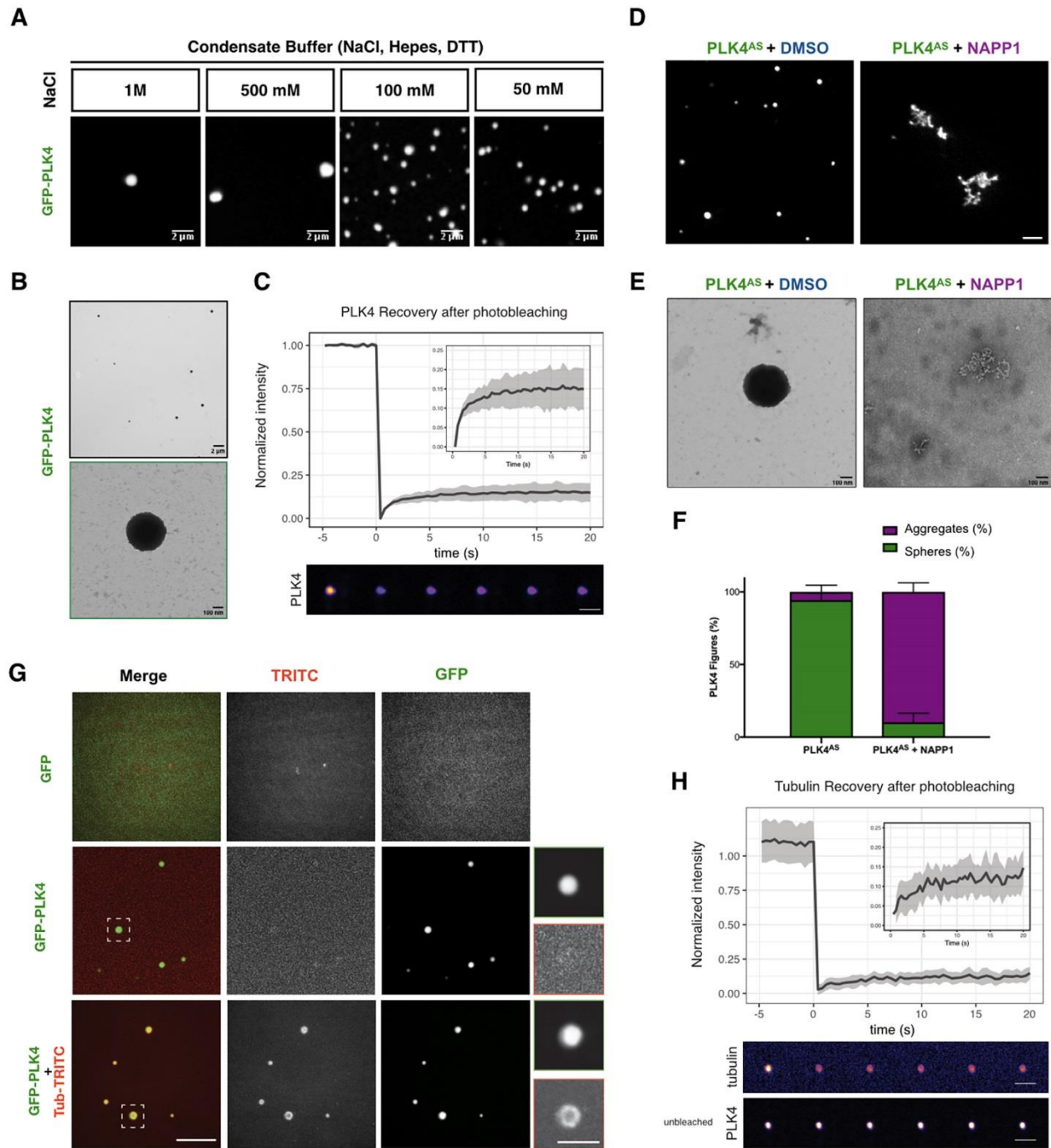
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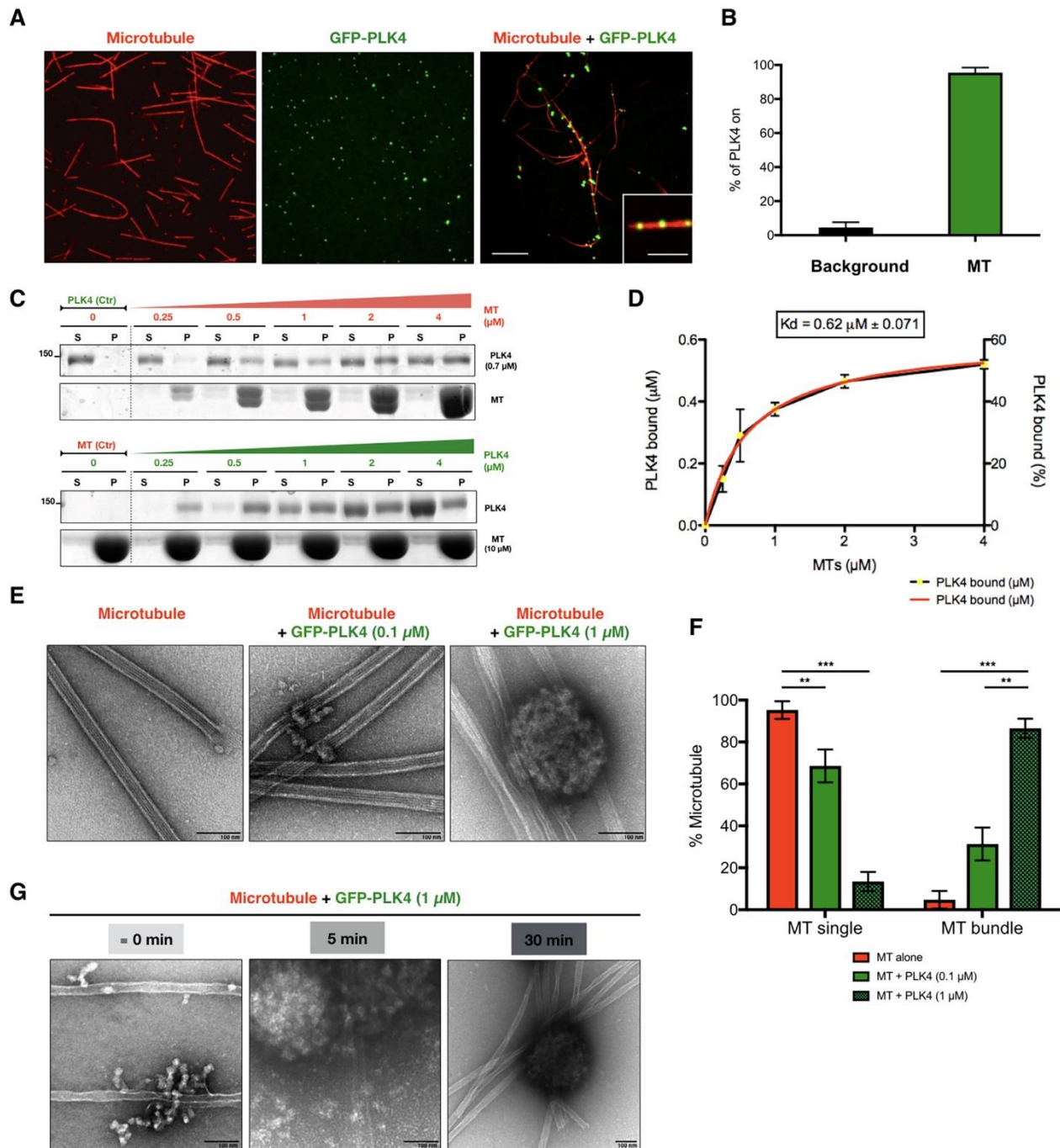
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# Figures



**Fig. 1. PLK4 self-assembly is dependent on its kinase activity and PLK4 assemblies concentrate tubulin *in vitro*.**

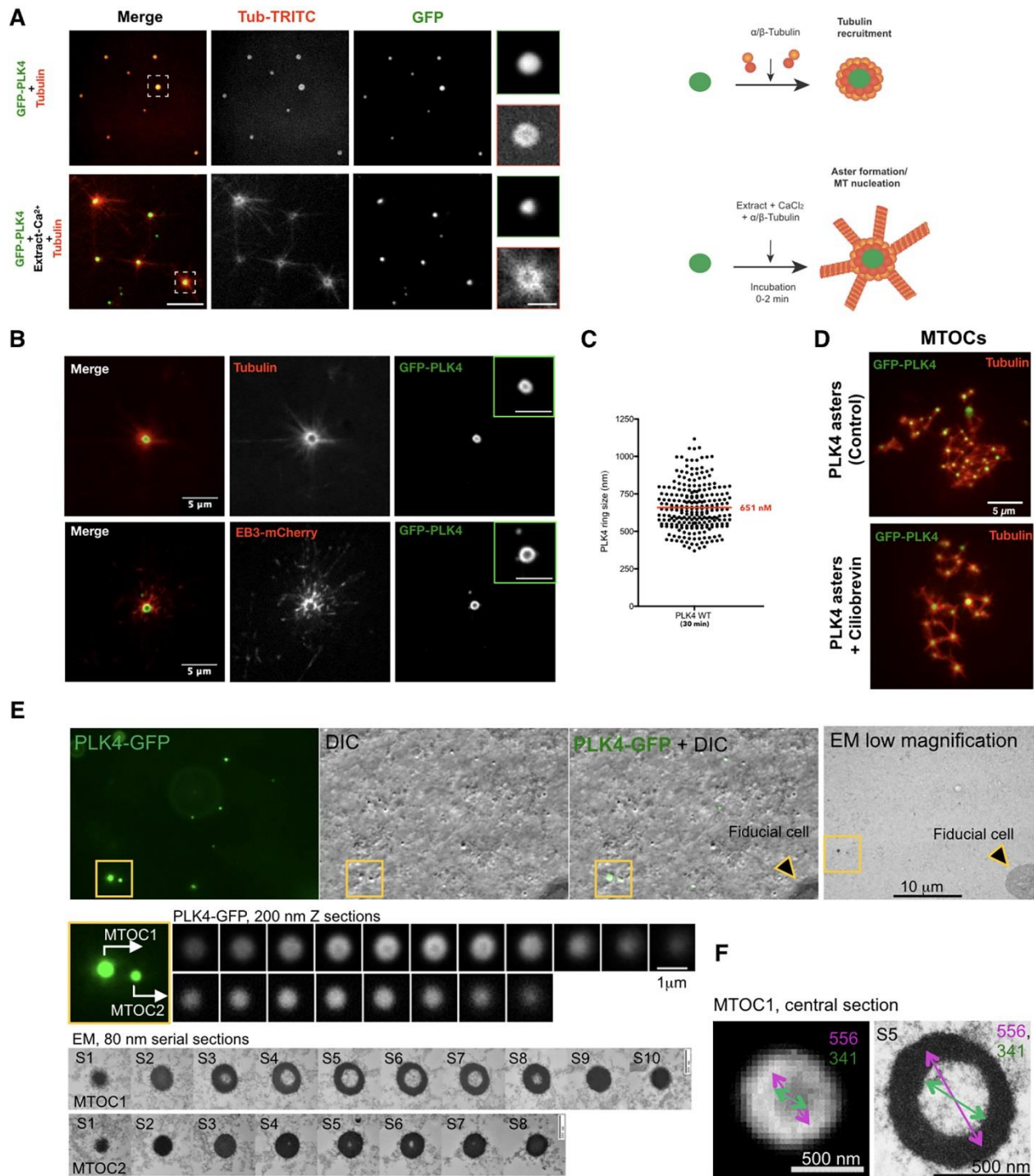
(A) Representative images of GFP-PLK4 assemblies formed at different concentrations of NaCl. (B) Electron Microscopy (EM) images showing PLK4 assemblies *in vitro*. Scale bars: 2  $\mu\text{m}$  and 100 nm. (C) FRAP Analysis of PLK4 *in vitro*. Fluorescence intensity recovery after photobleaching of PLK4 assemblies. Scale bar: 2  $\mu\text{m}$ . (D) Confocal images representing GFP-PLK4<sup>AS</sup> in the absence or presence of NAPP1. DMSO was used as a control for NAPP1. Scale bars: 5  $\mu\text{m}$ . Note that, in presence of NAPP1, GFP-PLK4 forms disorganized structures. (E) EM images of GFP-PLK4<sup>AS</sup> in the presence or absence of NAPP1. Scale bars: 100 nm. (F) Quantification (%) of sphere-like assemblies versus aggregates obtained from EM data. Three independent experiments were counted. (G) Confocal images of GFP-PLK4 assemblies formation in the absence or presence of rhodamine-labelled tubulin (500 nM). GFP was used as a control. Scale bars: 5  $\mu\text{m}$ ; Insets: 2  $\mu\text{m}$ . (H) FRAP Analysis of tubulin coating PLK4 spheres *in vitro*. Fluorescence intensity recovery after photobleaching of the tubulin coating PLK4 spheres, showing little dynamicity Scale bar: 2  $\mu\text{m}$ .



**Fig. 2. PLK4 is a microtubule-associated protein that promotes microtubule bundling *in vitro*.**

(A) Confocal images of Taxol-stabilized MTs alone (rhodamine-labelled tubulin, red), recombinant purified GFP-PLK4 alone (green) and the mixture of both, showing association of PLK4 condensates to MTs. Scale bar: 5  $\mu\text{M}$ ; Inset: 2  $\mu\text{m}$ . (B) Quantification of PLK4 assemblies associated to MTs compared to

free PLK4 in the background. (N=3, n=100 spot/conditions). (C) MT-pelleting assays. The two assays are showing either a constant PLK4 concentration (0.7  $\mu\text{M}$ ) mixed and incubated with different MTs concentrations (0 to 4  $\mu\text{M}$ ) or increasing amounts of GFP-PLK4 (0 to 4  $\mu\text{M}$ ) in the presence of constant MTs concentration (10  $\mu\text{M}$ ). Coomassie gel is showing supernatant (S) and pellet (P) for each condition. (D) Quantitative analysis of binding properties between PLK4 and MTs. Note that the dissociation constant ( $K_d$ ) for PLK4, determined by best fit to the data (red curve), is  $0.62 \pm 0.071 \mu\text{M}$ . Note that the dotted line is the real data and the red line is the fitted curve to derive constants. The data were collected from three independent experiments. (E) EM images showing MTs alone or MTs incubated with two concentrations of PLK4 (0.1  $\mu\text{M}$  and 1  $\mu\text{M}$ ). Scale bars: 100 nm. (F) Percentage of single or bundled MTs quantified from EM data in presence of PLK4 (0.1  $\mu\text{M}$  or 1  $\mu\text{M}$ ); MTs alone are used as a control. Results were scored using 30 images per condition obtained from 3 independent experiments each; (\*\* $P < 0.01$ ; \*\*\* $P < 0.001$ ). (G) Time course of PLK4 (1  $\mu\text{M}$ ) incubated with MTs. Note that PLK4 binds to MTs before PLK4 condensates are formed; (“ $\equiv$ ” means approximately 0 min, as feasible experimentally). Scale bars: 100 nm.

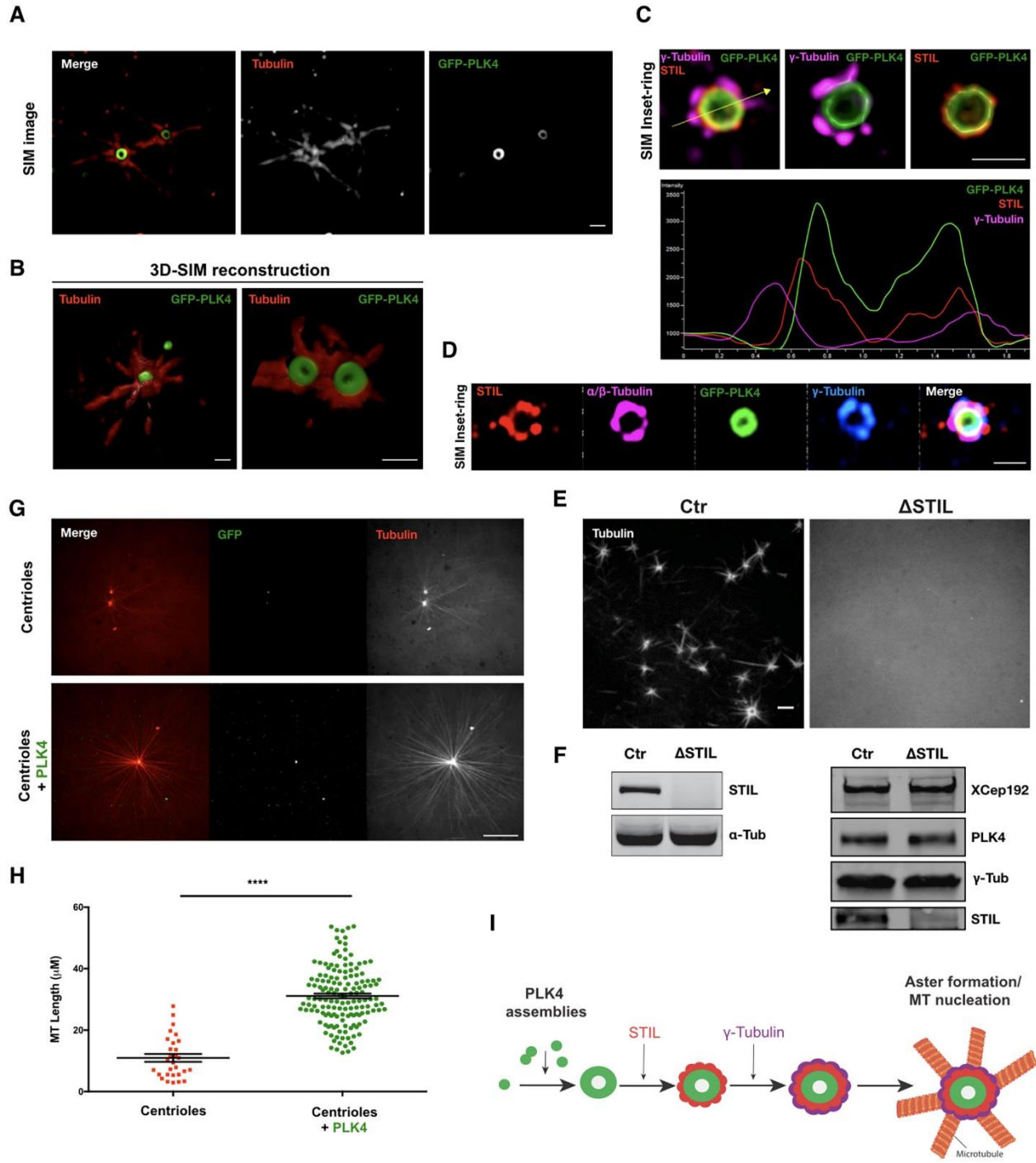


**Fig. 3. PLK4 condensates form *de novo* MTOCs in *Xenopus* extracts that are independent of motor proteins and mimic centrosomes *in vivo*.**

(A) GFP-PLK4 was mixed concomitantly with rhodamine-labelled tubulin and condensates were formed

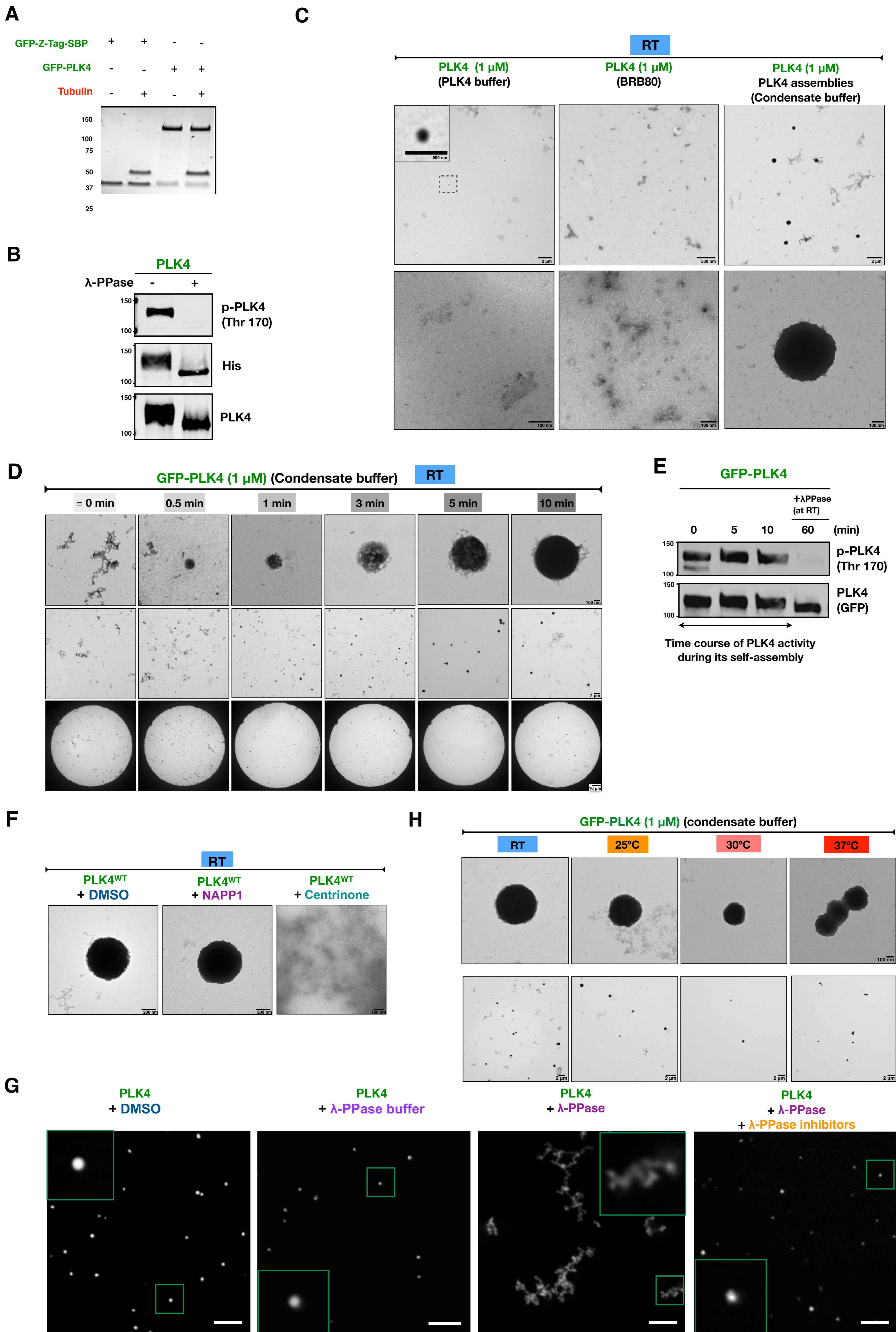
(step I) and then extract released to interphase with calcium containing rhodamine-labelled tubulin were added to these assemblies (step II). Note that nucleation was observed instantly after the addition of the mixture (0-2 min). Scale bars: 5  $\mu\text{m}$ , inset= 2  $\mu\text{m}$ . (B) Confocal images showing MTOC formation in *Xenopus* MII-calcium-released extracts in the presence of recombinant GFP-PLK4 (green). MTs are visualised using rhodamine-labelled tubulin (upper panel) and EB3-mCherry (lower panel). MT plus ends visualised by EB3-mCherry point out to the edge of the aster. Insets show PLK4 as a ring-like structure (Movie 1). (C) Quantification of the size (nm) of GFP-PLK4 ring-like structure after 30 min of incubation. GFP-PLK4 rings were measured from 3 independent experiments. (D) PLK4 asters are independent of dynein. Confocal images of PLK4 asters are shown in the control and in the presence of ciliobrevin (dynein inhibitor). (E) Correlative light/electron microscopy analysis of PLK4's MTOCs. PLK4-GFP signals were first visualised by fluorescence and DIC, and then by EM. A series of 200 nm sections (confocal) and 80 nm EM sections are presented for two MTOCs (yellow box, MTOC1 and MTOC2) (F) Measurements of the central sections of MTOC1 (section S5 in E). Scale bars: 500 and 1  $\mu\text{m}$ .





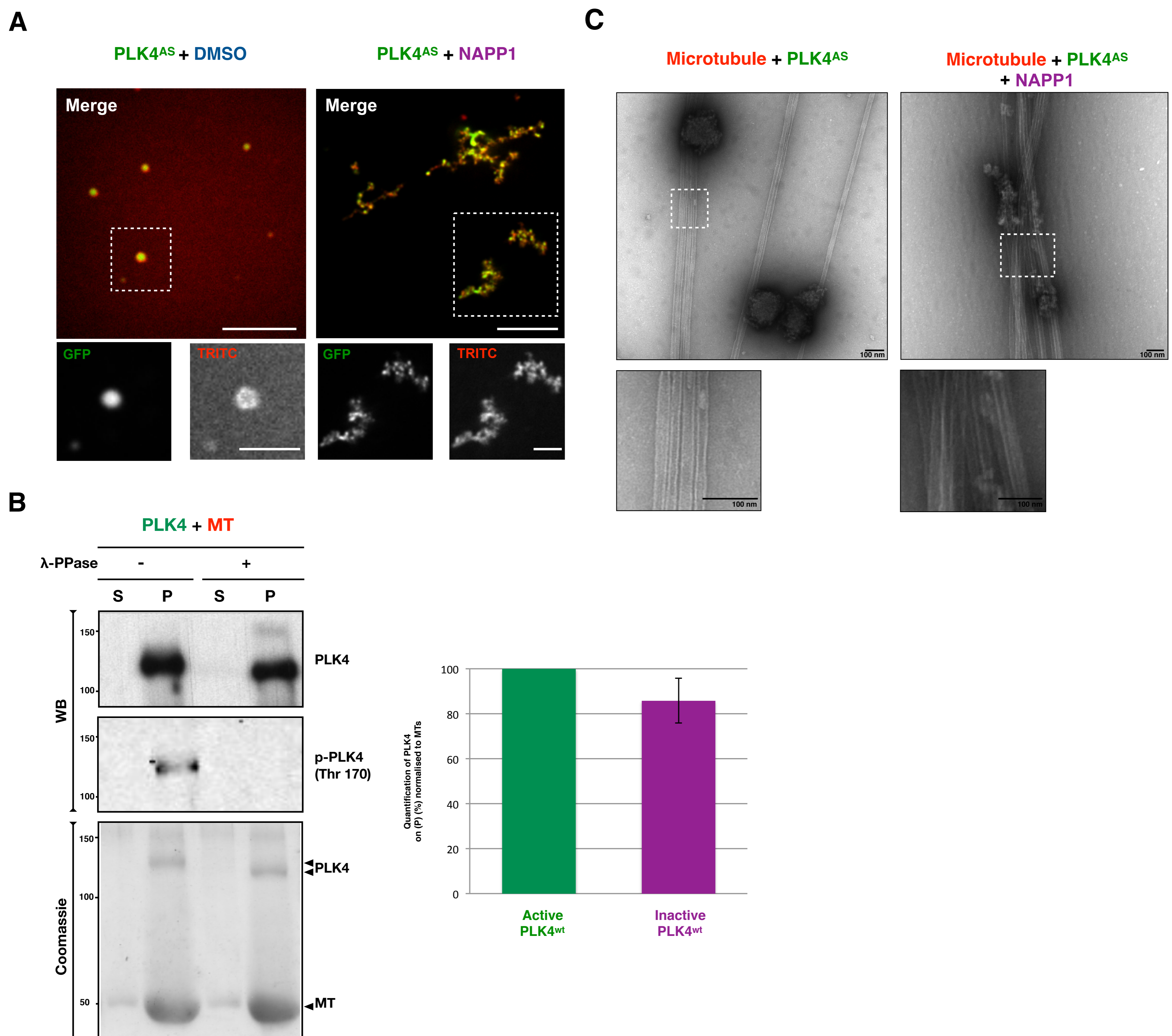
**Fig. 4. PLK4 MTOCs can recruit STIL and  $\gamma$ -tubulin in *Xenopus* released extracts and are able to enhance centrosomal MT nucleation.**

(A) Images of 3D-SIM showing a ring-like structure of PLK4 MTOCs formed in calcium-released *Xenopus* extracts.  $\alpha$ -tubulin and GFP-PLK4 are presented in red and green, respectively. Scale bars: 1  $\mu$ m. (B) 3D-reconstitution of PLK4 asters (Movie 3). Scale bars: 1  $\mu$ m. (C) SIM images showing the co-localisation of STIL (red),  $\gamma$ -tubulin (magenta) and GFP-PLK4 (green) within PLK4 MTOCs. Scale bars: 1  $\mu$ m. (Movie 4) (D) SIM images showing the co-localisation of GFP-PLK4 (green), STIL (red),  $\alpha/\beta$ -tubulin (magenta) and  $\gamma$ -tubulin (blue) that co-localise with PLK4 MTOCs. (E) Confocal images showing PLK4 induced-MTOCs in control extracts (Ctr) and STIL depleted extract ( $\Delta$ STIL) using rhodamine-tubulin. (F) Western blots showing depletion of STIL in the extracts used in (4E). The total level of proteins in these extracts is shown using antibodies against Xcep 192,  $\gamma$ -tub and PLK4. (G) PLK4 enhances MT nucleation. Confocal images showing MT nucleation using purified centrioles labelled with GFP-centrin incubated in *Xenopus* interphasic extract in the presence or absence of GFP-PLK4 (rhodamine-labelled tubulin (red); centriole and PLK4 (green)). Images were taken after 30 min incubation (Movie 5, 6). (H) Quantifications of MTs length ( $\mu$ m) visualised from the centrioles (GFP-centrin MTOCs) in the presence or absence of GFP-PLK4. MTs were measured from 2 independent experiments, where 4 different MTOCs were analysed. (N is the total number of MTs measured in the presence of GFP-PLK4, N= 225). Scale bar: 5  $\mu$ m. The statistical data are presented as  $\pm$  s.d. \*\*\*\*P < 0.0001, (Mann-Whitney U). (I) Representative scheme of PLK4 MTOCs formation in *Xenopus* extracts.



**Fig. S1. (Related to Fig. 1). Formation of PLK4 supramolecular assemblies is dependent on its activity.**

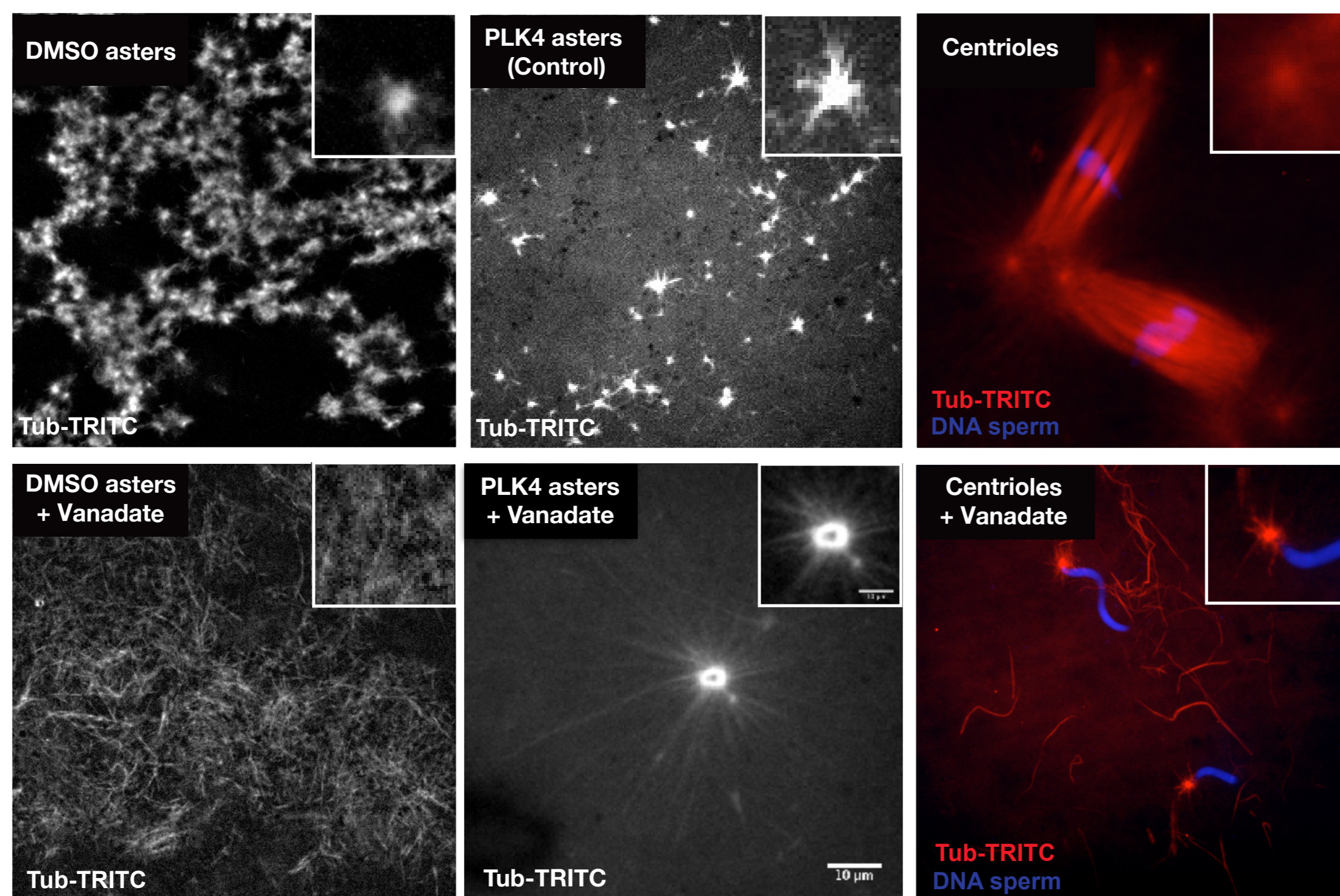
(A) Coomassie-stained gels showing the proteins used in different conditions in Figure 1. (B) Western blots showing PLK4's activity, when treated or not with lambda-phosphatase ( $\lambda$ -PPase), using an antibody against threonine-170 localised within the kinase domain. (C) EM images of PLK4 assemblies in control buffer (*PLK4 buffer* (50 mM Tris-HCl, pH 7.4, 500 mM NaCl, 0.5 mM DTT, 1% glycerol, 0.1% CHAPS) and *BRB80/MTs buffer* (25 mM HEPES, pH 6.8, 2 mM MgCl<sub>2</sub>, 1 mM EGTA, 0.02% Tween 20 (v/v)) and in the *Condensate buffer* (150 mM NaCl, 25 mM Hepes (pH 7.4); 1 mM DTT). Note that we detected few small PLK4 assemblies in PLK4 buffer similarly to what is shown in Figure 1A at 1 M and 500 mM of NaCl. (D) Time course of PLK4 assembly formation. Note that PLK4 assemblies grow and become round and dense while forming, with heterogeneous sizes (500 nm to 1  $\mu$ m). (E) WB showing the activity of PLK4 while forming the assemblies (Thr170 antibody and GFP for total PLK4 protein). Note that we used the  $\lambda$ -PPase treatment for 1 hour at room temperature (RT) to show the efficiency of the dephosphorylation compared to the typical reaction protocol of the phosphatase (30 min at 30°C; Fig. S1B). (F) EM images showing PLK4-WT assemblies in the control condition (DMSO) and when treated with NAPP1 (specifically inhibiting PLK4-AS) and centrinone (specific inhibitor of PLK4). Note that NAPP1 and centrinone were added simultaneously with the protein for acute inactivation. (G) Confocal images of GFP-PLK4 treated with  $\lambda$ -PPase used according to the typical reaction protocol, incubated for 1 hr at RT. The corresponding controls were performed to check effects of the buffers on PLK4 assemblies. To inactivate  $\lambda$ -PPase we used: 10 mM sodium orthovanadate, 50 mM EDTA, protease inhibitor cocktail 10%, leupeptin and aprotinin 10  $\mu$ g/ml, added to the buffer simultaneously with the phosphatase for acute inactivation. Scale bars: 5  $\mu$ m. (H) EM images of PLK4 assembly formation at different temperatures. Note that the structures are stable at 37°C, the temperature used for the assays with MTs.



**Fig. S2. (Related to Fig. 1 and 2). PLK4 assemblies recruit and bind to MTs independently of PLK4 activity.**

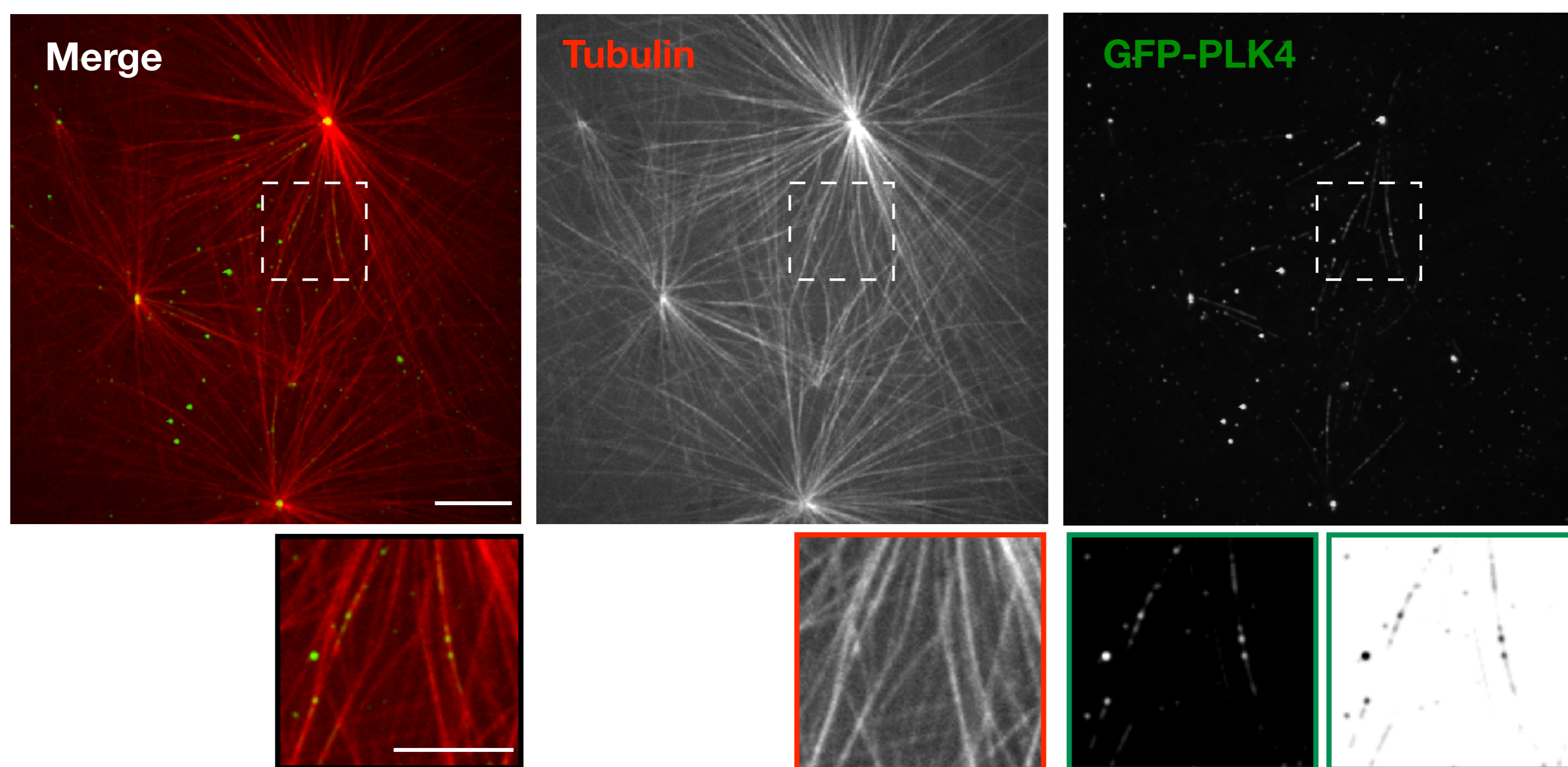
(A) Confocal images of PLK4-AS (treated or not with NAPP1) incubated with tubulin-rhodamine TritC. Either active or inactive PLK4 recruits tubulin similarly to the inactivated PLK4 in the presence of MTs. Scale bars: 10  $\mu$ m; insets: 5  $\mu$ m. (B) MT pelleting assay. MTs are incubated with PLK4 (active or inactive (treated with  $\lambda$ -PPase)). Coomassie gel and western blots (WB) show supernatant (S) and pellet (P) for each condition. We used anti-PLK4 (for total protein detection) and phospho-Thr-170 (PLK4 activity). A plot of PLK4 quantification on the pellet was performed from three independent experiments. We compared the level of PLK4 treated with  $\lambda$ -PPase to the level of active PLK4 in the pellet. Both PLK4 levels were normalised to MT level in the pellet. (C) EM pictures showing PLK4-AS binding to MTs when treated or not with NAPP1. Scale bars: 100 nm.

A



B

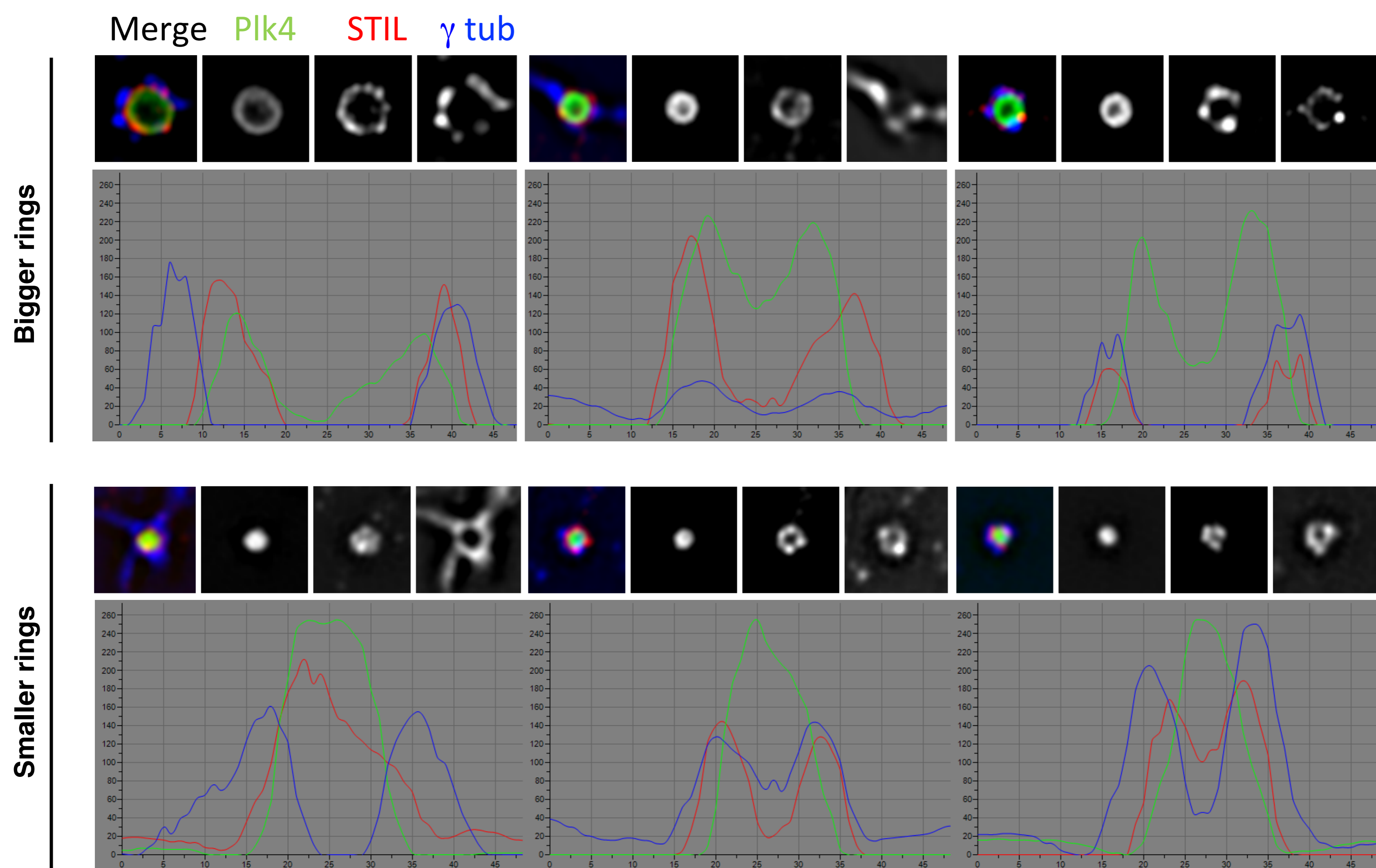
Interphase extract + GFP-Centrin-labeled centrosomes + GFP-PLK4 (1 μM)



**Fig. S3. (Related to Fig. 3). PLK4 binds to Microtubules in *Xenopus* extracts.**

(A) Images showing PLK4, DMSO and centriole asters in *Xenopus* extracts in the presence or absence of vanadate. DMSO forms asters in *Xenopus* extracts. Note that those asters are destroyed in the presence of vanadate (dynein inhibitor). Bipolar spindles assemble (red) in *Xenopus* extracts in the presence of sperm, which carries centrioles. In the presence of vanadate bipolar spindles are destroyed, but centrioles still nucleate MTs (positive control). Scale bars: 10 μm. (B) Confocal images showing GFP-PLK4 binding to MTs in interphase extract. Stable asters were obtained by adding GFP-centrin labeled purified centrosome (from HeLa cells) to interphase extracts. Soluble GFP-PLK4 was added to the extract to visualise PLK4 on MTs. Note that even though GFP-PLK4 is mainly at the MTOC centre, PLK4 is also observed on the MTs. Scale bars: 10 μm; insets: 5 μm.

A

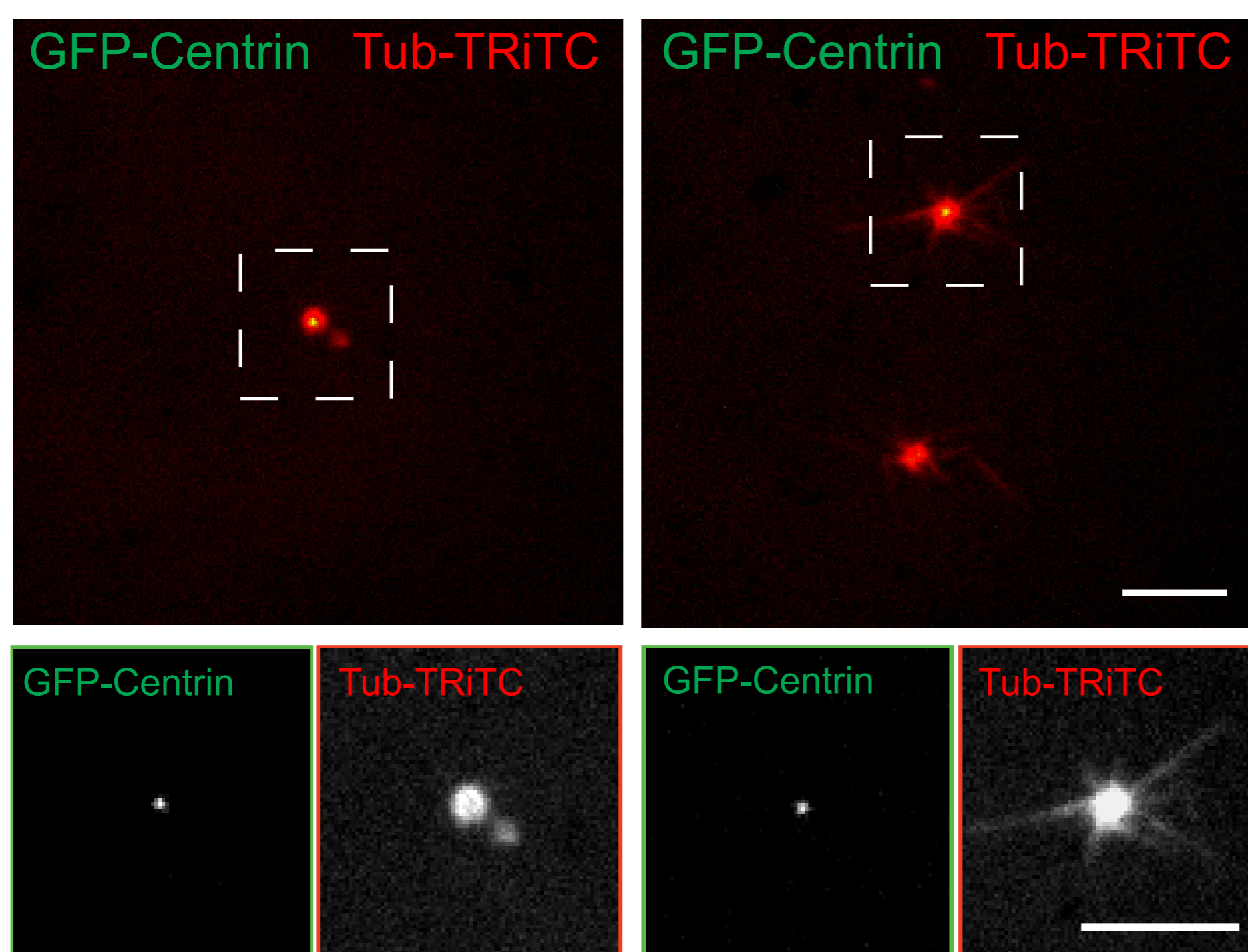


B

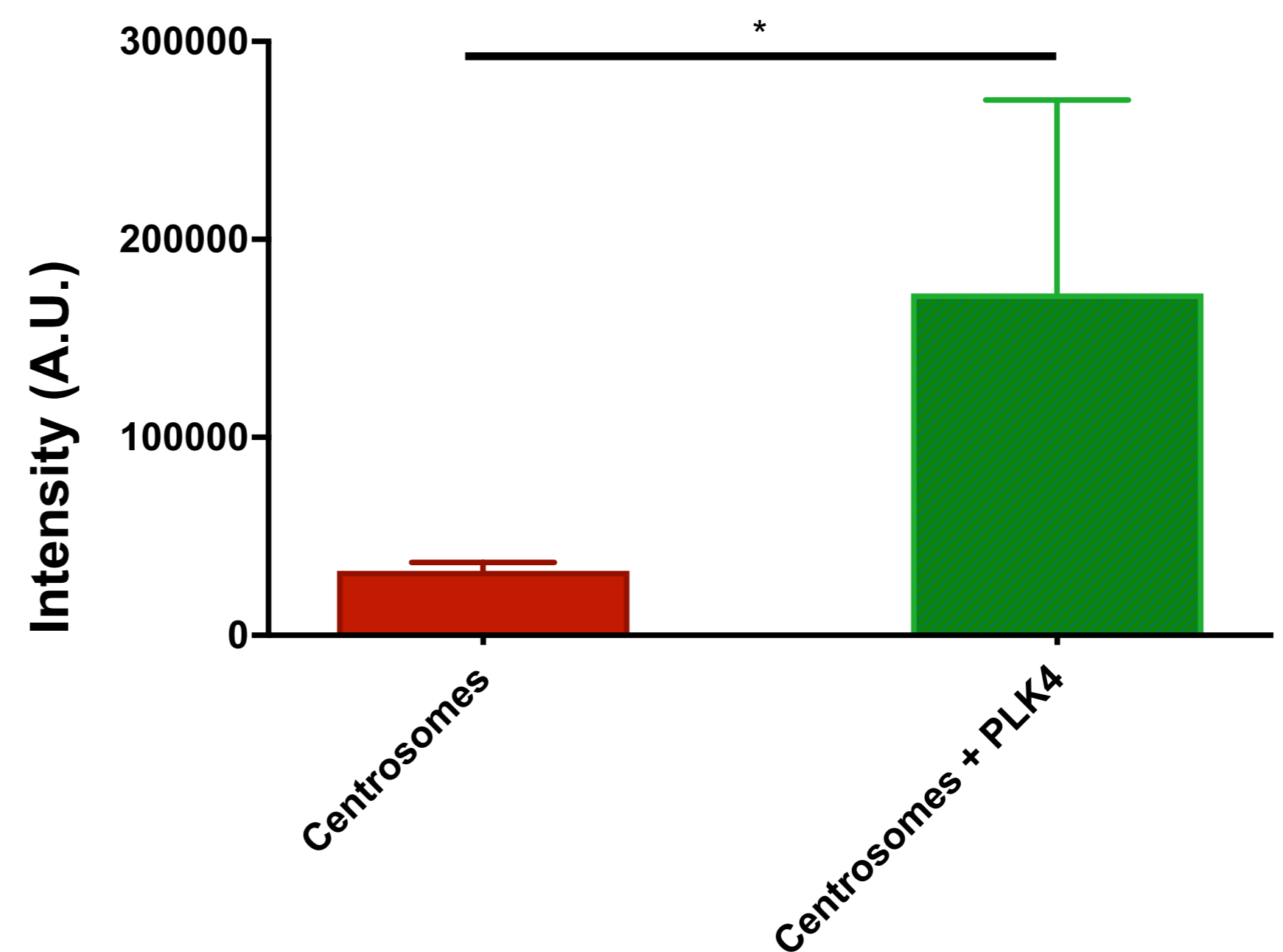
**CSF extract released with CaCl<sub>2</sub>**

**Centrosomes  
(GFP-Centrin)**

**Centrosomes  
(GFP-Centrin) + PLK4**

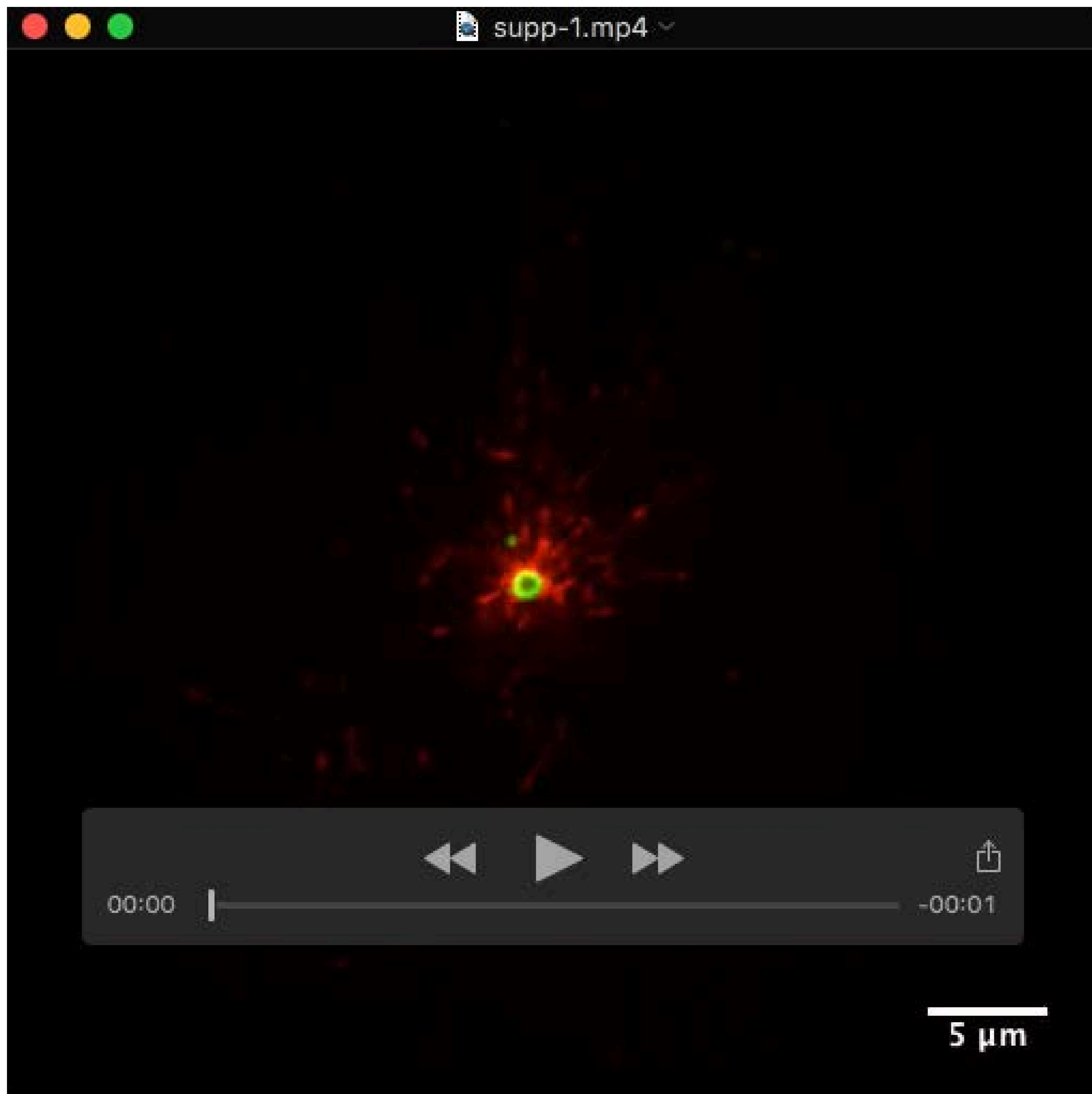


C



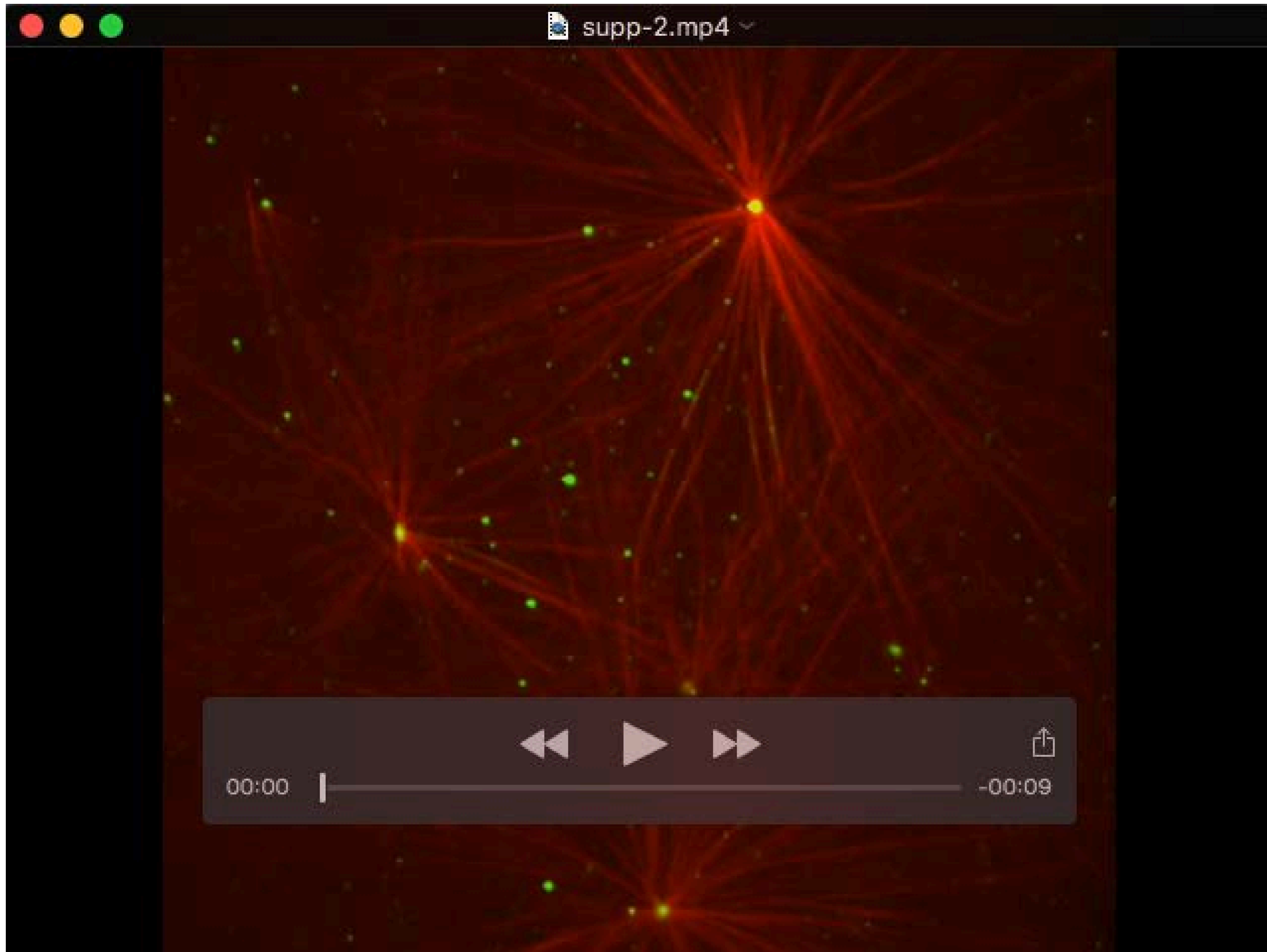
**Fig. S4. (Related to Fig 4). PLK4 enhances centrosomal MT nucleation.**

(A) Examples of spatial localisation of PLK4, STIL and  $\gamma$ -Tubulin using 3D-SIM of PLK4 assemblies at different sizes (bigger rings and smaller rings). 3D-SIM images of examples of PLK4 assemblies stained with PLK4, STIL and  $\gamma$ -Tubulin. Note that we show examples of PLK4 structures at different sizes and that STIL and  $\gamma$ -Tubulin coat PLK4 assemblies in an organised spatial localisation. (B) Purified centrosomes from HeLa cells labeled with GFP-centrin were added to MII *Xenopus* extracts (CSF) released to interphase with CaCl<sub>2</sub>. Labeled tubulin was added to visualise asters. Note that the time of incubation is 15 min to visualise only the nucleation that results from centrosomes. GFP-centrin (green) and MTs (red). Scale bars: 5  $\mu$ m. (C) Quantification of the intensity of centrosomal MT nucleation in MII-extracts released into interphase with CaCl<sub>2</sub> in the presence or absence of GFP-PLK4 (1  $\mu$ M). Results were obtained from four different independent experiments. (A.U: arbitrary units). The statistical data are presented as  $\pm$  s.d. \* $P$  < 0.05, (Student's t-test).

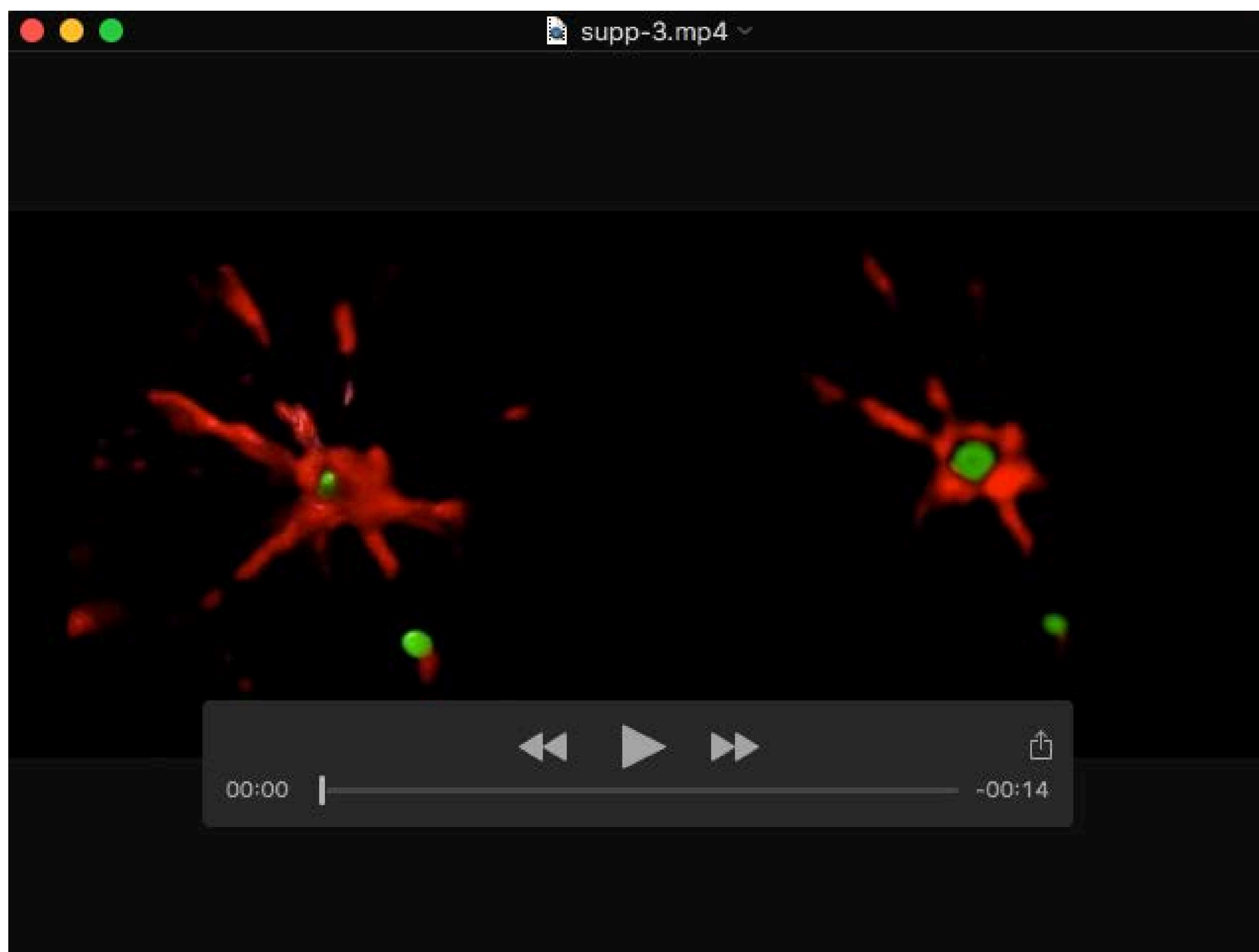


**Movie S1 (Related to Fig. 3B).** GFP-PLK4 (green) MTOC formation in *Xenopus* egg extract using EB3-mCherry (red) to visualise MT plus ends.

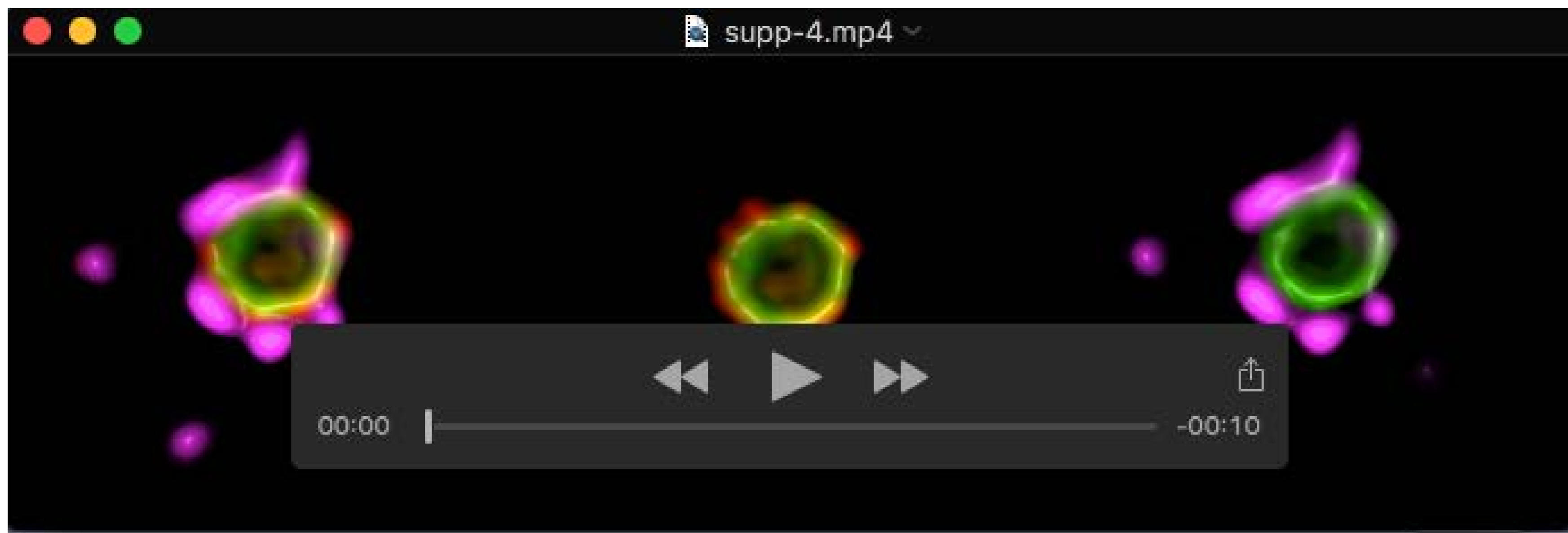




**Movie S2 (Related to Fig. 3).** GFP-PLK4 (green) binds to MTs (red) in *Xenopus* interphase extracts. We visualise PLK4 at the centre of the aster, and soluble PLK4 was observed sticking to MTs.



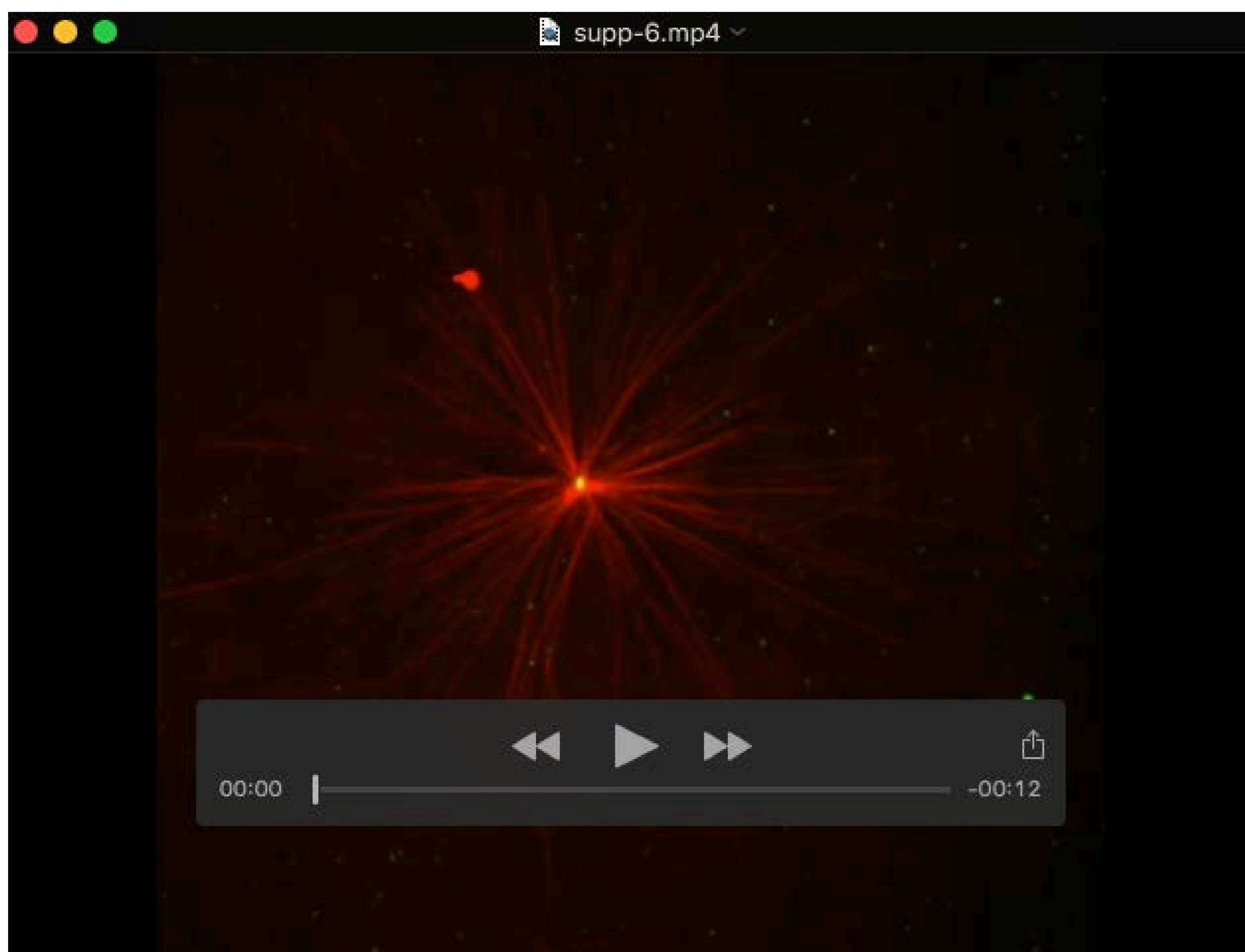
**Movie S3 (Related to Fig. 4B).** 3D-SIM reconstruction of PLK4 MTOCs in *Xenopus* egg extracts using rhodamine-tubulin. Reconstruction of the whole aster.



**Movie S4 (Related to Fig. 4C)** 3D-SIM reconstruction of PLK4 MTOCs in *Xenopus* egg extracts with STIL and  $\gamma$ -tubulin.



**Movie S5 (Related to Fig. 4G).** Purified centrioles in *Xenopus* egg extracts.



**Movie S6 (Related to Fig. 4G).** Purified centrioles in *Xenopus* egg extracts in the presence of GFP-PLK4.