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ASPM and CITK regulate spindle orientation by affecting the dynamics of astral microtubules

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Abstract

Correct orientation of cell division is considered an important factor for the achievement of normal brain size, as mutations in genes that affect this process are among the leading causes of microcephaly. Abnormal spindle orientation is associated with reduction of the neuronal progenitor symmetric divisions, premature cell cycle exit and reduced neurogenesis. This mechanism has been involved in microcephaly resulting from mutation of *ASPM*, the most frequently affected gene in autosomal recessive human primary microcephaly (MCPH), but it is presently unknown how *ASPM* regulates spindle orientation. In this report, we show that *ASPM* may control spindle positioning by interacting with Citron-kinase (CITK), a protein whose loss is also responsible for severe microcephaly in mammals. We show that the absence of CITK leads to abnormal spindle orientation in mammals and insects. In mouse cortical development, this phenotype correlates with increased production of basal progenitors. *ASPM* is required to recruit CITK at the spindle and CITK overexpression rescues *ASPM* phenotype. *ASPM* and CITK affect the organization of astral microtubule (MT) and low doses of MT-stabilizing drug revert the spindle orientation phenotype produced by their knockdown. Finally, CITK regulates both astral MT nucleation and stability. Our results provide a functional link between two established microcephaly proteins.

Synopsis

CITK controls spindle orientation in neural progenitor cells in *Drosophila* and mouse.

ASPM recruits CITK to the spindle, where it promotes the nucleation and stabilization of astral microtubules, providing a link between the two microcephaly proteins.

- CITK $-/-$ mouse developing cortex is characterized by increased cell cycle exit and generation of basal progenitors.
- CITK physically interacts with ASPM at the spindle pole in HeLa cells.
- CITK and ASPM depleted HeLa cells displayed a significant reduction in the number and length of astral microtubules.

Introduction

The orientation of cell division is carefully controlled in both embryonic and adult tissues, in which it may regulate cell fate, generate tissue shape and maintain normal histological architecture [1-3]. Studies conducted over the last two decades have established that this process may play an important role to regulate the delicate balance between proliferation and differentiation that underlies normal brain development, with particular regard to the generation of a normal number of cortical neurons [2, 4, 5]. During embryogenesis, the cerebral cortex is first composed of a single layer of neuroepithelial (NE) cells, which initially expand through symmetric divisions [6-9]. As development progresses, NE progenitors give rise to radial glia (RG) cells that may further expand by dividing symmetrically or may switch to asymmetric division, producing a self-renewing progenitor and a daughter cell committed to differentiation [5, 10-12]. A characteristic feature of proliferative divisions of NE and RG cells is that cleavage occurs perpendicular to the ventricular surface of the neuroepithelium, while the switch from symmetric to asymmetric divisions is accompanied by a deviation of the cleavage plane [13-15]. Oriented cell division is achieved through the proper positioning of the mitotic spindle [16-18], which depends on the formation of molecular links between actin-rich cell cortex and astral microtubules (MT) emanating from centrosome-derived spindle poles [15, 19, 20]. Indeed either loss of Doublecortin (*DCX*), which destabilizes MT [21], or mutations of *LIS1*, *NDE1* and *NDEL1*, which disrupt dynein–dynactin function at the cell cortex [22-24], randomize the mitotic spindle in NE progenitors and lead to their early exhaustion. A similar depletion of progenitor cells, correlated to randomized spindle orientation, can be produced by disruption of proteins involved in centrosome function [25-28]. Whether the relationship between spindle orientation and cell fate choice in cortical development is causal or only correlative is currently debated, on the basis of recent studies suggesting that spindle orientation may not be essential for cortical neurogenesis [26, 29, 30]. Mutation of many

proteins contributing to centriole biogenesis, centrosome maturation and spindle organization [31, 32] has been associated to human primary microcephaly (MCPH). MCPH is characterized by reduced head circumference, accompanied by relatively preserved brain architecture, resulting in mild to moderate intellectual disability and few associated symptoms [33, 34]. At least thirteen MCPH loci (MCPH1-MCPH13) have been mapped to date [35-37] and it has been established that most of the encoded proteins are capable of localizing at the centrosome, at the spindle poles or at the spindle. Nevertheless, our understanding of how these proteins may affect spindle orientation and cell fate determination is still very limited. *ASPM* (abnormal spindle-like microcephaly associated, *MCPH5*) is the most frequently mutated gene in MCPH [28, 38]. *ASPM* is a conserved protein that associates with the MT minus ends, is recruited at the spindle poles during mitosis and controls spindle MT organization, spindle function and cytokinesis from insects to mammals [39-41]. In mammals, NE cells with reduced *ASPM* expression fail to orient the mitotic spindle perpendicular to the ventricular surface of the neuroepithelium and show increased frequency of asymmetric divisions, with a reduction of the pool of neuronal precursors [25, 42]. Recent evidence indicates that *ASPM* may control cell fate choice by regulating the activity of Cdk2/Cyclin E complex [43]. However, it is presently unknown how *ASPM* regulates spindle orientation. Previous studies [44] showed that *ASPM* physically interacts with the cytokinesis regulator Citron Kinase (CITK) [45, 46]. CITK is a conserved protein prominently localized at the cleavage furrow and at the midbody of mitotic cells [46, 47] and is involved in abscission control at the end of cytokinesis [48, 49]. In rodents, CITK inactivation leads to severe microcephaly and lethal epilepsy [45, 50]. This phenotype is believed to result from apoptosis due to cytokinesis failure. Delayed metaphase-anaphase transition in CITK deficient neuronal progenitors has also been described [51]. Although *ASPM* and CITK are both associated with microcephaly and may

form a complex in HeLa cells and in developing neural tissue [44], it remained unclear whether their interaction could have functional relevance.

In this report we show that, in addition to its role in cytokinesis, CITK plays a phylogenetically conserved role in the control of mitotic spindle orientation, by promoting the nucleation and stabilization of astral MT. Moreover, we show that CITK is recruited to the spindle by ASPM and that CITK overexpression rescues the spindle orientation defect elicited by ASPM knockdown. These results provide new insight into the mechanisms by which ASPM loss may cause microcephaly and suggest that a spindle orientation defect may contribute to the CITK microcephaly phenotype.

Results and discussion

1. CITK is a phylogenetically conserved determinant of spindle orientation.

In light of the previously reported association between CITK and ASPM [44] and of the role played by ASPM in regulating spindle orientation [25, 39], we asked whether CITK may also play a role in controlling spindle orientation, besides promoting abscission [49]. To address this question, we first examined whether CITK contributes to maintaining mitotic spindle perpendicular to the apical–basal axis of mouse NE progenitors. To do so, we immunostained fixed cryosections of E14.5 mice neocortices for spindle pole protein γ -tubulin and DNA, and we measured the angle formed by the cleavage plan of apical progenitors (AP) with the apical surface of cortex [13, 14, 25, 30] (Fig. 1A). This angle was defined as vertical when it was comprised between 75° and 90° and oblique when it was in the range from 0 to 75° . Most AP (75%) divide vertically in CitK $+/+$ cortices, while vertical divisions occurred in only 24% of AP in CitK $-/-$ mice (Fig. 1B), indicating that spindle orientation is perturbed by CITK loss.

To evaluate whether this phenotype correlates with increased commitment to differentiation, we pulse-labeled E13.5 embryos with BrdU and analyzed how many cells had left the cell cycle 24 hours later, measuring the percentage of cells positive for BrdU

but negative for the cell cycle marker Ki67 [52]. A significant increase of BrdU+/Ki67- cells was observed in the proliferative regions and in the intermediate zone (Fig. 1C-D), suggesting a shift from proliferative to differentiative divisions. Indeed, the increase of cells exited from cell cycle in these regions could not be explained by neuron migration defects, because the percentage of neurons in VZ/SVZ relative to the total neuron population is the same in WT and CITK $-/-$ mice (Fig. 1E). Interestingly, the percentage of BrdU+/Tbr2+ cells in the proliferative regions was significantly increased (Fig. 1F). In contrast, the total number of Tbr2+ cells was not different in control and knockout cortices ($55,4 \pm 4,6$ and $52,2 \pm 4,1$ per 100 μm -wide cortical column, respectively). It must be noted that the number of neurons produced in CITK $-/-$ may be strongly underestimated, because of their apoptotic death (Fig. 1E). Indeed, we observed a strong absolute reduction of BrdU-positive cells in the neuronal layers of CitK $-/-$ mice (Fig. 1C), consistent with the previous finding that in this model apoptosis occurs especially in post-mitotic neuroblasts and neurons [53]. Altogether, these results indicate that CitK $-/-$ developing cortex is characterized by increased cell cycle exit and increased generation of basal progenitors. Since the sequence of CITK is well conserved between *Drosophila* and mammals, and since the loss of CITK in *Drosophila* produces a cytokinesis-failure phenotype remarkably similar to the phenotype detected in mammalian cells [47, 54-56], we asked whether the role of CITK in spindle orientation is conserved as well. To address this question, we analyzed neuroblast (NB) divisions in larval brains from individuals homozygous for either *dck*¹ or *dck*², two presumptive null alleles at the locus encoding the *Drosophila* orthologue of CITK [47]. *Drosophila* NB are stem cells that divide asymmetrically, to give rise to another NB and to a smaller ganglion mother cell (GMC) committed to differentiation. To ensure a correct asymmetric division, NB spindle must be aligned to the cell polarity axis determined by the differential apico-basal concentration of several proteins. The basal cortex is enriched in proteins that are preferentially segregated into the GMC at the end of

division and whose localization is in turn mediated by a large multiprotein complex that concentrates at the apical cortex [57]. We immunostained wild type and *dck* mutant larval brains for Tubulin and for the basal marker Miranda (Mira) and measured the angle between a line connecting the two spindle poles and a line bisecting the crescent formed by Mira in metaphase NBs (Fig. 1G). The angle ranged between 0° and 5° in 88% of control NBs (Fig. 1H), indicating a tight coupling of the mitotic spindle with the polarity axis. Conversely, although both *dck*¹ and *dck*² mutant NBs consistently displayed well formed Mira crescents, the majority of the spindles showed more oblique orientations, ranging from 6° to 45° (n= 33 for *dck*¹, and 45 for *dck*², respectively; Fig. 1H).

These results indicate that, besides to its role in abscission control, CITK plays a phylogenetically conserved role earlier in mitosis, to ensure correct positioning of the mitotic spindle.

To better characterize this function, we resorted to HeLa cells, which are sensitive to CITK depletion [49, 58] and have extensively been used to study spindle positioning mechanisms [59]. We depleted CITK in these cells by RNAi using validated sequences [49] (Fig. EV1A), and analyzed cell division through phase-contrast time-lapse microscopy. Interestingly, divisions of cells treated with CITK-specific siRNAs were misoriented, with the two forming daughters partially overlapping rather than being adjacent and with one daughter cell dividing outside of the focal plan, thus delaying adherence to the substrate (Fig. 2A-C and Movies EV1-2). We then analyzed, in fixed samples, the angles formed by metaphase spindles with the culture dish. CITK-depleted cells displayed angles distribution skewed towards high values and significant increase in angles average (Fig. 2D-E), whereas spindle length was not affected by CITK depletion (Fig. EV4A). A similar phenotype was observed by inducing CITK depletion with a second, independent siRNA sequence (Fig. EV1B-C). The phenotype was rescued by restoring CITK levels through the expression of an RNAi-resistant construct (Fig. EV1C), further confirming that it is due

to CITK depletion rather than to off-target effects of the siRNA sequences used. The phenotype observed in CITK-depleted cells was similar to the phenotype produced by ASPM depletion [39] (Fig. 2F). Moreover, the defect on mitotic spindle orientation caused by CITK and ASPM simultaneous knock-down was the same as single knock-down (Fig. 2F), indicating that these two proteins are on the same pathway.

Altogether, these results show that CITK is a conserved determinant of spindle orientation and demonstrate that it acts in mitosis even before becoming enriched at the cleavage furrow and midbody during cytokinesis.

2. CITK is associated with mitotic spindle through ASPM.

Several proteomic studies reported that CITK is associated with the mitotic spindle [60-62], but this localization was not confirmed by other techniques. Previous immuno-localization studies have shown that CITK is localized to the nucleus in interphase cells, accumulates in the cytoplasm before anaphase and becomes enriched at the cleavage furrow and at the midbody during cytokinesis [46, 63]. A similar pattern was detected by live cell imaging in HeLa cells expressing physiological levels of GFP-tagged CITK from a stably integrated BAC transgene [64-66] (Movie EV3). As expected, during mitosis the protein appears to be evenly distributed in the cytoplasm and no enrichment is visible at the spindle or at the cell cortex (Movie EV3).

Although these data confirmed the previous reports [46, 63], they did not exclude the possibility that a pool of the protein may associate with mitotic spindle. To address this issue, we performed mild detergent extraction before fixation, which removes cytosolic proteins and facilitates the visualization of cytoskeleton-associated proteins. Under these conditions, immunofluorescence with anti-CITK antibodies detected a clear enrichment of the protein on both the spindle and the spindle poles, as it partially overlapped with the γ -tubulin signal (Fig. 3A). This signal disappears in cell treated with CITK siRNA (Fig. 3A). Similar results were obtained using anti-GFP antibodies in HeLa cells expressing GFP-

tagged CITK from a BAC transgene [66] (Fig. 3B). Specific association of CITK with spindle poles was further validated through biochemistry, as we found that CITK is enriched in centrosomal preparations obtained from HeLa cells synchronized in metaphase (Fig. 3C). In addition, we observed that the signals of both CITK endogenous protein and CITK-GFP extensively overlap with ASPM immunoreactivity at the spindle poles (Fig. 3D-E).

To confirm the physical interaction between CITK and ASPM, we performed *in situ* proximity ligation assay (PLA) which can reveal if two proteins are at a distance <40 nm [67]. In mitotic cells expressing physiological levels of ASPM-GFP [66] and immunostained for GFP and CITK, we observed a strong signal, which was completely absent in normal HeLa cells, not expressing tagged ASPM (Fig. 3F). Fluorescence dots were observed throughout the cytoplasm and in proximity of spindle and spindle poles (Fig. 3F). Since these cells were not pre-extracted, the result indicates that CITK interacts with both the cytoplasmic ASPM pool and with spindle-associated ASPM.

We then examined which CITK regions are involved in the interaction with ASPM. Based on the previously shown interaction between CITK and ASPM C-terminal region (CTR) [44] we tested the ability of ASPM-CTR to coimmunoprecipitate with different myc-CITK fusion proteins (Fig. EV2A), including kinase active (FL) or kinase dead (KD) full length CITK, Citron-N (CN), which is a brain-specific isoform of CITK completely lacking the kinase domain [68], the N-terminal and the C-terminal halves of CITK (CITK-Nt and CITK-Ct, respectively). We found that ASPM-CTR can bind all these proteins (Fig. EV2B), suggesting that it may interact, directly or indirectly, with different regions of CITK, independently from its kinase activity.

We next analyzed the localization of CITK in ASPM-depleted cells and vice versa. While the localization pattern of ASPM was not affected by CITK-RNAi (Fig. 3G), the percentage of cells with CITK-positive spindle poles and the mean intensity of the CITK at the spindle

poles of positive cells were significantly decreased by ASPM depletion (Fig. 3H-I). These results confirm the interaction between CITK and ASPM in mitotic cells and indicate that a pool of CITK is recruited through ASPM at the spindle and spindle poles, suggesting that CITK may act downstream of ASPM.

3. CITK regulates astral MT organization downstream of ASPM.

Proper orientation of mitotic spindle during mitosis requires a dynamic connection of astral MT to the cortical cytoskeleton through the MT motor complex Dynein/Dynactin, which is localized at a cortical crescent by cortical adaptor proteins such as NuMA [19, 69-71]. We therefore analyzed by immunofluorescence whether CITK absence alters the localization of these proteins at metaphase. No significant changes were observed in the localization pattern of Dynein or NuMA (Fig. EV3A-C), while the percentage of cells with a Dynactin (P150) crescent was slightly increased (Fig. EV3A-C), indicating that the phenotype induced by CITK depletion cannot be explained by defective localization of these proteins. Moreover, no differences were observed in the fluorescence intensity of these proteins at spindle poles (Fig. EV3A-C).

In contrast, if compared to controls, CITK-depleted cells displayed a significant reduction in the number and length of astral MT (Fig. 4A-B). A similar phenotype was observed in the case of ASPM depletion (Fig. 4A-B). It has been reported that asymmetric neurogenic division is associated with spindle-size asymmetry, negatively controlled by the Wnt pathway [72]. Since ASPM has been shown to act upstream of the Wnt pathway to trigger symmetric divisions of cortical progenitors [73], we assessed whether CITK loss of function impacts spindle symmetry. We found that the difference (Δ) in the number of astral MT on the two centrosomes of a mitotic cell didn't change when either CITK or ASPM were knocked down (Fig. EV4B). These data further indicate that CITK and ASPM may act through similar mechanisms and reveal that they influence astral MT organization. To functionally test the possibility that CITK may work downstream of ASPM, we transfected

ASPM-depleted cells with CITK overexpression constructs. Remarkably, the overexpression of CITK reverted the reduction of astral MT (Fig. 4C) and the average increase of mitotic angle (Fig. 4D) elicited by ASPM depletion back to control values. No rescue was observed when ASPM-depleted cells were cotransfected with a kinase-dead mutant of CITK (CKD) or with CITN (Fig. 4D), indicating that CITK catalytic activity may be required for spindle orientation. These results suggest that, when CITK is expressed at physiological levels, the presence of ASPM is a limiting factor for its recruitment to spindle poles and for its activity on spindle orientation, while CITK overexpression overcomes the requirement for ASPM. A possible explanation for the latter phenomenon could be provided by the finding that CITK can bind to microtubules [48] through many microtubule-binding proteins, including KIF14, MKLP1, PRC1 [58], KIF4A and KIF10 [65]. Since the localization of at least some of these proteins during metaphase may partially overlap with ASPM localization [74], it is conceivable that at high CITK levels the binding with these proteins, or other unknown partners, may compensate for ASPM loss.

Next, we treated HeLa cells, depleted for either proteins, with low doses of the MT-stabilizing drug Paclitaxel, which are known to specifically affect the most dynamic pool of MT [75]. In both cases, the average of mitotic spindle angles was restored to control values (Fig. 4E). Altogether, these results indicate that ASPM and CITK regulate spindle orientation through astral MT organization and that CITK is a downstream player of ASPM.

4. CITK regulates astral MT length, stability and nucleation.

To investigate which aspects of the dynamic behavior of MT are affected by CITK, we performed RNAi in HeLa cells stably expressing the MT+TIP protein EB3, fused with the fluorescent Tomato tag, to track the plus ends of growing MT by time lapse fluorescence microscopy [20] (Fig.5A). Quantitative analysis of the corresponding movies allows estimation of the different MT dynamic parameters, as it has previously been described [20, 76] (Fig. 5B). This technique revealed that CITK loss leads to decreased MT stability

and maximal distance travelled from the spindle pole towards the cell cortex, while no differences were observed in MT growth speed (Fig. 5C). Moreover, we observed a decrease of nucleation frequency (Fig. 5C).

To better address the latter phenotype we resorted to MT growth assay after cold-induced depolymerization, by incubating cells at 4°C for 30 minutes. 1 minute after switching from 4°C to 37°C, the number of cells showing detectable MT growth (aster size >1µm) was significantly decreased in CITK-depleted cells (Fig. 5D-E). Moreover, although at 2 minutes the value of this parameter was comparable to the control (Fig. 5E), the aster size was still significantly smaller in CITK-depleted cells (Fig. 5F). Altogether, these results indicate that CITK loss decreases MT nucleation and MT stability in mitotic cells.

In summary, in this study we have provided evidence that ASPM controls spindle orientation by regulating the dynamics of astral MT and that CITK is a critical downstream partner of ASPM for this activity. Moreover, although the analysis of CITK *-/-* mice is complicated by the high prevalence of cytokinesis failure and apoptosis, our *in vivo* data suggest that the two proteins may also cooperate in determining cell fate of dividing NP, a possibility consistent with the reduction of post-natal neural stem cells described in CITK *-/-* rats [77]. Thus, our study suggest that, besides apoptosis, disturbed cell fate contributes to microcephalic phenotype of CITK *-/-* mice. Therefore, it will be interesting to test whether CITK may cooperate with ASPM in controlling the activity of Cdk2/Cyclin E complex [43]. It also remains to be established how CITK may modulate astral MT dynamics. We have shown that CITK affects both MT nucleation and stability. The function of CITK in cytokinesis has already been linked to MT organization. Indeed, CITK promotes midbody maturation by recruiting at the midbody the kinesins KIF14 [48, 78] and MLKP1 [58], which in turn recruit the MT-crosslinking protein PRC1. CITK stabilizes midbody microtubules by indirectly stimulating TUBB3 phosphorylation [53]. Finally, systematic proteomic studies have found that CIT-K may interact with the kinesin KIF4A and KIF10

[65]. However, the roles of these proteins in astral MT organization and in spindle orientation have not been established. Addressing the functional significance of CITK interaction with these proteins and identifying CITK substrates will be an interesting subject for future studies. In addition, it will be very interesting to address whether other MCPH proteins associated with MT function, such as WDR62, CDK5RAP2, CENPJ and STIL [33] may regulate astral MT functionally cooperating with ASPM and CITK.

Material and Methods

Cell culture, synchronization and drug treatments

Unmodified HeLa cells were originally obtained from ATCC and a batch was frozen after 5 passages. All cells are routinely screened for mycoplasma contamination.

HeLa cells were cultured in RPMI medium supplemented with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin. HeLa cells expressing GFP-tagged CITK and ASPM from a BAC transgene were obtained by courtesy of Hyman Lab (Max Planck Institute of Molecular Cell Biology and Genetics, Dresden) cultured in DMEM 10% heat inactivated FBS, 1% penicillin/streptomycin, 0.4 mg/ml G418, Geneticin® (Life Technologies). The HeLa cell line expressing EB3–tdTomato [79], obtained by courtesy of Dr. Ann Straube (University of Warwick, Coventry, United Kingdom) was maintained in DMEM-GlutaMAX (Invitrogen) supplemented with 10% FBS, 100 Uml⁻¹ penicillin, 100µgml⁻¹ streptomycin, 200µgml⁻¹ Geneticin (Sigma) and 0.5µgml⁻¹ puromycin [20]. All cells were cultured in a humidified 5% CO₂ incubator at 37°C.

Cells were synchronized by a double- or single-thymidine block. For the double thymidine block, asynchronous cultures were treated with 2 mM thymidine (Sigma) for 16 h, then released for 4–6 h in fresh complete medium and again blocked for 16 h. Finally, cells were washed twice with fresh medium and allowed to progress through mitosis.

For drug treatments, synchronized HeLa cells were treated with 250 pM of Paclitaxel (Sigma) for 30 minutes before fixation for immunofluorescence.

Transfection, RNAi and rescue.

In this study, we used a previously validated (CK1 = AUGGAAGGCACU AUUUCUCAA) (Gai et al., 2011) and a second CITK-siRNA sequence (CK2 = CCAUUCUUGGCACCUUGCUU). The knockdown of ASPM was obtained using a previously validated sequence [39] (GUGGUGAAGGUGACCUUUC). All siRNAs were obtained from GE-Healthcare (Dharmacon, Lafayette, CO). The ON-TARGETplus non-targeting siRNA #1 (GE Healthcare, Dharmacon) was used as a negative control for potential off-target effects. The expression construct coding for CITK-Cherry was generated by inserting the Myc-CITK cDNA into the pm-Cherry-N1 plasmid (Clontech, Mountain View, CA) [49]. The same method was used to generate constructs coding for CKD-Cherry and for CITN-Cherry. HeLa cells plated on a six-well plate were transfected using 1 µg of the required plasmid DNA and 3 µl of *Trans-IT-LT1* transfection reagent (Mirus Bio, Madison, WI), or 6.25 µl of the required siRNA (20 µM) and 2.5 µl Lipofectamine 2000 (Invitrogen, Carlsbad, CA), according to the manufacturers' instructions. For rescue experiments, cells were transiently transfected with siRNAs; 24 h later cells were transfected with DNA constructs and analyzed 24 h later.

Cell staining

Fixation methods are as follows: for α -tubulin staining, 4% paraformaldehyde at room temperature for 10 min or with methanol at -20°C for 10 min; for γ -tubulin, ASPM and NuMA staining, methanol at -20°C for 10 min; for Dynactin, Dynein staining, cells were pre-extracted in 0.5% Triton X-100 in PHEM buffer (60 mM PIPES, 25 mM HEPES, 10 mM EGTA, and 4 mM MgSO₄) for 1 min, and then fixed in methanol at -20°C for 10 min; for CITK staining, cells were pre-extracted in 0.5% Triton X-100 in PHEM buffer for 1 min and fixed with 4% paraformaldehyde at room temperature for 10 min. In all cases, cells were permeabilized in 0,1% Triton X 100 in PBS for 10 minutes, saturated in 5% BSA in PBS for 30 minutes and incubated with primary antibody for 45 min at room temperature. Primary

antibodies were detected with anti-rabbit Alexa Fluor 488 or 568 (Molecular Probes, Invitrogen), anti-mouse Alexa Fluor 488 or 568 (Molecular Probes, Invitrogen) used at 1:1000 dilution for 30 min. Counterstaining was performed with the DNA dye DAPI (Sigma) at 0.5 µg/ml for 1 minute.

Antibodies

The following primary antibodies were used:

mouse anti-alpha-tubulin, 1:1000 (Sigma, Cat. T5168, clone B-5-1-2); rabbit anti-alpha-tubulin, 1:200 (Abcam, cat. ab15246); mouse anti-gamma-tubulin, 1:200 (Abcam, clone TU-30, cat. ab27074); rabbit anti-gamma-tubulin, 1:1000 (Sigma, cat. T5192); mouse anti-CITK, 1:100 (BD Transduction Laboratories, clone 6/CRIK, cat. 611376); rabbit anti-Numa, 1:2000 (Abcam, cat. ab36999); rabbit anti-Dynein Heavy Chain, 1:300 (Santa Cruz, cat. sc-9115); rabbit anti-ASPM (1:2000 [39]); mouse anti-p150, 1:1000 (BD Biosciences, cat. 610474); rabbit anti-GFP, 1:1000 (Abcam, cat. ab290); mouse anti-Ki67, 1:1000 (BD Biosciences, clone B56, cat. 550609); mouse anti-BrdU, 1:200 (Chemicon, Cat. MAB-1467); mouse anti-TUBB3, 1:1000 (clone TuJ1, Covance, cat. MMS-435P); rabbit anti-Tbr2, 1:200 (Millipore, cat. AB2283).

Mice.

CITK ^{-/-} E14.5 embryos and matched controls were obtained by crossing CITK ^{+/-} mice (mixed genetic background C57/Bl6 x Sv129). Animals were housed in the Animal facility of Molecular Biotechnology Centre, University of Torino, under license n. 343/2015 PR from Italian Ministry of Health. For estimate of sample size, CITK phenotypes are usually 100% penetrant, with minimal variability due to genetic background. Under these conditions, analysis of three biological replicates is usually sufficient to reveal the existence of significant differences.

Mouse embryonic cortex staining

Embryonic brains were dissected at E14.5 and fixed for 12-16h at 4°C in 4% PFA. For cell cycle exit analysis, pregnant females were previously injected at E13.5 with 75 mg/kg BrdU. Fixed brains were equilibrated in 30% sucrose in PBS for 12-24h at 4°C. Brains were embedded with Tissue-TEK (O.C.T, Sakura Finetek), snap frozen in liquid nitrogen and stored at -20°C. Sectioning was performed at 20µm with a cryostat. Cryosections were rehydrated five times in PBS before further processing. For measurement of spindle angle, cryosections were subjected to antigen retrieval by heating in 0.01 M citrate buffer pH 6.0 at 70°C for 1 h. Sections were then permeabilized using 0.3% Triton X-100 in PBS for 30 min and quenched with 0.1 M glycine for 30 min. Sections were then incubated with primary antibodies overnight at 4°C, followed by secondary antibody for 1 h at room temperature in a solution of 0.2% gelatin, 300 mM NaCl, and 0.3% Triton X-100 in PBS. DNA was stained in the last wash using DAPI. For cell cycle exit experiments, BrdU antigen was retrieved using 2N HCl and a 10 min in a steamer, followed by neutralization with 0.1M borate buffer pH 8.5. Slides were then processed for IF as above. We usually exclude samples that are not judged of sufficient technical quality by microscopic analysis. In this study no samples were excluded.

***Drosophila* strains**

The *Drosophila citron kinase*¹ (*ck*¹) mutant allele, also called *l(3)7m-62* or *sticky*¹ (*sti*¹), is described by [80] and by FlyBase (<http://flybase.bio.indiana.edu/>).

The *dck*² allele was isolated from a collection of 1600 ethylmethanesulfonate (EMS)-induced third chromosome late lethals, generated in Charles Zuker's laboratory (University of California, San Diego, CA) [47]. *ck*¹ and *ck*² mutant alleles were kept in stocks over the TM6 C, a chromosome balancer which carry the dominant larval marker *Tubby* (<http://flybase.bio.indiana.edu/>); homozygous mutant larvae were recognized for their non-*Tubby* phenotype. All stocks were maintained on standard *Drosophila* medium at 25°C.

Immunostaining of *Drosophila* neuroblast cells

Brains from third instar larvae were dissected and fixed according to [81]. After two rinses in phosphate buffered saline (PBS), brain preparations were incubated overnight at 4°C with a monoclonal anti- α -tubulin antibody (Sigma), diluted 1:1000 in PBS and rabbit anti-Miranda antibody (a gift of Y. N. Jan) diluted 1:100 in PBS. After two rinses in PBS, primary antibodies were detected by 1 hour incubation at room temperature with FITC-conjugated anti-mouse IgG + IgM (1:20; Jackson Laboratories) and Alexa-Fluor-555-conjugated anti-rabbit IgG (1:300; Molecular Probes), diluted in PBS. Immunostained preparations were mounted in Vectashield H-1200 (Vector Laboratories) containing DAPI (4,6 diamidino-2- phenylindole). The angle between a line connecting the two spindle poles and a line bisecting the crescent formed by Mira in metaphase NBs was measured, by hand with a goniometer, on images captured by a Zeiss Axioplan fluorescence microscope, equipped with an HBO100W mercury lamp and a cooled charged-coupled device (CCD camera; Photometrics CoolSnap HQ).

Microscopy

Imaging was performed using a Leica TCS SP5-AOBS 5-channel confocal system (Leica Microsystems) equipped with a 405nm diode, an argon ion and a 561nm DPSS laser. Fixed cells were imaged using a HCX PL APO 63x/1.4 NA oil immersion objective at a pixel resolution of 0.108 x 0.108 μm .

Time lapses were recorded overnight with an interval of 5 minutes using a 40X PlanApo N.A. 1.4 oil immersion objective on the cells kept in the microscope incubator at 37°C and 5% CO₂.

Image analysis

Spindle angle measurements were obtained using a trigonometric formula, once measured the three dimensional (3D) distance (across the x, y, and z planes) and the 2D distance (across the z planes) between the two centrosomes of the spindle.

To quantify NuMA, Dynein and p150Glued signals at the centrosome, we used Fiji (fiji.sc; [82]). After obtaining the sum of intensity of all the Z stacks from multi stacks images, we built a ROI surrounding the signal at the centrosome and a ROI around the cells. Once the ROIs were established, they were used for all the images analyzed. We evaluated the Corrected Integrated Density (CTCF), for both the centrosomes, calculating as Integrated Density (Area of selection x Mean fluorescence of the background). Analysis requiring manual counting were performed on blinded samples.

Measurement of astral-MT dynamics

Astral-MT nucleation frequencies were quantified as described [20] by counting the number of EB3 comets in the focal plane that originated from each spindle pole during the observation time. Cytoplasmic growth speeds for individual comets were measured as the total distance travelled from the spindle pole towards the cell cortex divided by the time. Kymographs of astral-MT and spindle-pole movements were obtained using Fiji.

Proximity Ligation Assay

PLA analysis [67] was performed using the kit provided by Olink Bioscience (Uppsala Science Park, Uppsala, Sweden). Metaphase HeLa cells and ASPM-GFP HeLa stable cell lines were incubated with the previously described primary antibodies and reactions were performed following manufacturer's instructions.

MT regrowth assay

48 h after siRNA transfection, MT were depolymerized at 4°C for 30 minutes. Cells were then washed and incubated in culture medium at 37°C to allow regrowth. Cells were fixed at different time intervals in methanol at -20°C for 10 min and processed for immunofluorescence microscopy to examine MT regrowth from spindle poles in metaphase cells. The MT signal was quantified using Fiji.

Centrosomal fractionation

Centrosome fractionation was performed as described previously [83]. In brief, exponentially growing cells were incubated with 10 µg/ml nocodazole (Sigma-Aldrich) and 5 µg/ml Cytochalasin B (Sigma-Aldrich) for 90 min, and then lysed by hypotonic treatment. Centrosomes were harvested by centrifugation onto a 20% Ficoll cushion and further purified by centrifugation through a discontinuous (70, 50, and 40%) sucrose gradient. Fractions of 0.3 ml were collected and analyzed.

Immunoprecipitation and Western blotting

For all immunoprecipitations, cells were extracted with lysis buffer containing 1% Triton X-100, 120 mM NaCl, 50 mM Tris-HCl (pH 7.5), protease inhibitors (Roche, Mannheim, Germany), and 1 mM phenylmethylsulfonyl fluoride (PMSF). The extracts were then clarified by centrifugation. Antibodies and protein-G–Sepharose beads (Amersham Biosciences, Uppsala, Sweden) were added to lysates and incubated overnight at 4°C. Pellets were washed four times with lysis buffer and analyzed by SDS–PAGE.

For immunoblots, immunoprecipitates or equal amounts of proteins from whole-cell lysates were resolved by reduction with SDS–PAGE and blotted to nitrocellulose filters, which were then incubated with the indicated antibodies and developed using the ECL System (Amersham Biosciences).

Statistics

For the experiments in which we can show statistically significant differences, sample size was determined on the basis of pilot studies estimating the order of magnitude of differences. For the experiments showing non significant differences, we used the same sample size of those revealing differences. Normal distribution of data is evaluated by graphical analysis of data plots. All the numbers represented by histograms were obtained by averaging all the data points contained in the replicas (with a comparable number of points per each replica). Statistical significance of the differences were assessed using a two tails Student T-test.

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Conflict of interests

The authors declare that they have no conflict of interest.

Author contributions

All experiments were designed and conceived by M.G. and F.D.C. The mouse in vivo experiments were performed by F.T.B. with the support and the advice of W.B.H. The *Drosophila* experiments were performed by F.V. and S.B. The PLA experiment was performed by L.A. and A.S. The analysis of centrosome fractions were performed by A.B. and N.L. All the other experiments were performed by M.G, with the support of C.V., S.P., G.E.B., F.S. and A.A.C. ASPM reagents were provided by J.B. The manuscript was written by M.G. and F.D.C.

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Figure Legends

Figure 1. CITK controls mitotic spindle orientation in developing mouse and fly brains.

(A) WT and CITK $-/-$ embryonic (E14.5) mouse cerebral cortex was stained for γ -tubulin (green) and DNA (gray). The ventricular plane is marked by a red dashed line, and the spindle axis of apical progenitors is indicated by a white dashed line. The angle between these two lines represents the mitotic angle. Scale bars = 5 μ m.

(B) Quantification of vertical divisions of apical progenitors in WT and CITK $-/-$ mice. $n=3$ per each genotype.

(C) Pregnant CITK $+/-$ females, crossed with CITK $+/-$ males, were injected with BrdU at E13.5; 24 hours later (E14.5) WT and CITK $-/-$ embryonic mouse cerebral cortex were

fixed in 4% PFA, cryosectioned and stained for Ki67 (red), BrdU (green), Tbr2 (blue) and DAPI (gray). The different cortical regions are indicated: CP=cortical plate; IZ= intermediate zone; SVZ= subventricular zone; VZ= ventricular zone. Scale bars = 10 μ m. White arrows in the inset indicate two Ki67- cells positive for BrdU and Tbr2.

(D) Quantification of cell cycle exit (ratio of cells BrdU+/ Ki67-) in the proliferative regions (VZ/SVZ) and in the neuronal layers (IZ and CP) of sections prepared as in panel C (n=4 per each genotype).

(E) Sections prepared as in C were stained for TUBB3 (TuJ) to reveal post-mitotic neurons and with DAPI. Red arrowheads indicate apoptotic neurons (defined by pyknotic nuclei). Quantification of neuronal distribution in VZ-SVZ versus total neurons population is showed on the right. Scale bars = 10 μ m.

(F) Quantification of the percentage of Tbr2-positive cells in the BrdU-positive population of the same sections analyzed for panel D (n=4 per each genotype).

(G) Neuroblasts of wild type and *dck* alleles immunostained for α -tubulin and Miranda. Note that whereas in wild type cells there is a tight coupling of the mitotic spindle with the polarity axis, in *dck* alleles the spindle show a more oblique orientation respect to Mira crescent. Scale bar = 5 μ m.

(H) Distribution of spindle angle amplitude ($^{\circ}$) in neuroblasts of wild type and *ck* alleles (n=58).

Data shown in histograms are means \pm SEM, statistical significance was assessed using a two tails Student T-test. ** P < 0.01; * P < 0.05.

Figure 2. CITK controls mitotic spindle orientation in HeLa cells.

(A) Selected frames from time-lapse imaging experiments (see Movies EV1 and EV2) showing two dividing cells transfected with either control (CTRL) or CITK-specific siRNAs,

respectively. Note one of the two daughter cells in the lower panel (red arrow) going out of focus as a consequence of oblique division.

(B) Graphical representation of vertical and oblique divisions. The cleavage plane is indicated by a black dashed line.

(C) Quantification of divisions showing uneven timing of daughter cell flattening onto the substrate after mitosis (oblique division) in CITK siRNA treated HeLa cells compared to control (n > 50 cells, 3 independent experiments).

(D) Control or CITK-depleted cells were immunostained for γ -tubulin (red) and DNA (blue) and imaged in z (0.3 μ m-thick sections). Upper panel: maximum intensity projections of confocal z-stacks are shown. Lower panel: cross-section (XZ) through the two poles of the same cell.

(E) Distribution of spindle angles ($^{\circ}$) in control and in CITK-depleted cells. The values represent the angles between the axis crossing the two poles of metaphase spindles and the coverslip. (n \geq 150 cells, 6 independent experiments).

(F) Quantification of spindle angles average in control cells and in cells depleted of CITK, ASPM or co-depleted of the two proteins (n \geq 150 cells, in at least 3 independent experiments).

Scale bars, 5 μ m. Data shown in histograms are means \pm SEM. Statistical significance was assessed using a two tails Student T-test. *** P < 0.001, ** P < 0.01.

Figure 3. CITK is associated with mitotic spindle poles through ASPM.

(A) HeLa cells pre-extracted for 1 min with 0.5% Triton X-100 in PHEM buffer and immunostained for CITK (green), γ -tubulin (red) and DNA (blue). Scale bars, 10 μ m

(B) HeLa cells expressing CITK-GFP (green) from a BAC transgene treated as in (A) and immunostained for γ -tubulin (red) and DNA (blue). Scale bars, 5 μ m

(C) Western blot of centrosome-containing fractions from HeLa cells, showing that CITK co-purifies with γ -tubulin.

(D) HeLa cells pre-extracted 1 min with 0.5% Triton X-100 in PHEM buffer and immunostained for CITK (green), ASPM (red) and DNA (blue).

(E) HeLa cells expressing CITK-GFP treated as in (A) and immunostained for GFP (green), ASPM (red) and DNA (blue).

(F) Close physical proximity between CITK and ASPM, revealed by PLA on ASPM-GFP-expressing HeLa cells, incubated with GFP and CITK antibodies and developed with PLA-specific secondary reagents.

(G) Control or CITK-depleted cells immunostained for ASPM (green), γ -tubulin (red) and DNA (blue).

(H) Control or ASPM-depleted cells treated as in (A) and immunostained for CITK (green), γ -tubulin (red) and DNA (blue).

(I) Quantification of CITK positive centrosomes and of the ratio of CITK spindle pole intensity versus total cell mean intensity, in control and ASPM-depleted cells. (n \geq 75 cells, 4 independent experiments).

Data presented are means \pm SEM. *** P < 0.001; ** P < 0.01. Statistical significance was assessed using a two tails Student T-test. All cell images represent maximum intensity projections of confocal z-stacks. Scale bars, 5 μ m.

Figure 4. CITK regulates astral MT and spindle orientation downstream of ASPM and through a kinase-dependent mechanism.

(A) Control, CITK or ASPM-depleted HeLa cells immunostained for α -tubulin (red) and DNA (blue). Maximum intensity projections of confocal z-stacks are shown.

(B) Quantification of astral MT number and maximal length in HeLa cells transfected with CTRL, CITK or ASPM siRNA (n > 130 cells in 3 independent experiments).

(C) Quantification of the rescue of astral MT number after CITK overexpression in ASPM-depleted HeLa cells (n > 50 cells in 3 independent experiments).

(D) Quantification of spindle orientation rescue after overexpression of CITK, CITN (lacking the kinase domain) or CKD (Citron kinase dead) mutants in ASPM-depleted HeLa cells ($n > 50$ cells in 3 independent experiments).

(E) Synchronized HeLa transfected with control, ASPM- or CITK-specific siRNAs treated with DMSO or with 250 pM Paclitaxel (taxol) 30 min before immunostaining for α -Tubulin. Spindle angles were measured as above ($n > 100$ cells in 6 independent experiments).

Data presented in (B), (C), (D) and (E) are means \pm SEM. Statistical significance was assessed using a two tails Student T-test. *** $P < 0.001$. All cell images represent maximum intensity projections of confocal z-stacks. Scale bars, 5 μ m.

Figure 5. Loss of CITK decreases astral MT length, stability and nucleation.

(A) Exemplar image of EB3–tdTomato HeLa cell used for kymograph analyses. To assess astral MT number and dynamics a line (blue) was added to images from the pole to the cortex.

(B) Kymographs of pole (bottom)-to-cortex (top) line scans of control and CITK-depleted EB3–tdTomato HeLa cells. The maximum projection of a 20-pixel-wide line (exemplified above) was plotted at 500 ms time intervals from left to right. MT growth results in diagonal lines.

(C) Quantification of astral MT dynamics. MT stability, length, growth speed ($n > 250$ MT, 30 cells) and nucleation frequency were calculated ($n > 30$ cells, 3 independent experiments).

(D) Immunofluorescence microscopy images showing MT growth at mitotic spindle poles 0, 1, and 2 min after nocodazole washout. Control and CITK-depleted cells were immunostained for α -tubulin (red) and DNA (blue).

(E) Percentage of cells showing detectable MT nucleation (aster size $> 1 \mu$ m) 0, 1, and 2 min after nocodazole washout ($n > 100$ cells, 4 independent experiments).

(F) Quantification of aster size 2 min after nocodazole washout ($n > 100$ cells, 4 independent experiments).

Data are presented as mean \pm SEM. Statistical significance was assessed using a two tails Student T-test. ** $P < 0.01$; *** $P < 0.001$. Scale bars, 5 μm .

Expanded View Figure Legends

Figure EV1. Specificity of the CITK RNAi experiments.

(A) Western blot analysis performed with the indicated antibodies on lysates of HeLa cells transfected with control or with two different CITK-specific siRNAs (CITK1 and CITK2). The results of experiments shown in the main figures were obtained with the CITK1 sequence.

(B) Distribution of spindle angles ($^{\circ}$) in HeLa cells depleted of CITK using the CITK2 sequence or transfected with control sequence. The values represent the angles between the axis crossing the two poles of metaphase spindles and the coverslip ($n \geq 150$ cells in three independent experiments).

(C) Quantification of average and distribution of spindle angles ($^{\circ}$) in HeLa cells depleted of CITK using the CITK2 ($n \geq 50$ cells in three independent experiments).

(D) Quantification of angles average in HeLa cells transfected with either control or CITK1 siRNAs and co-transfected with expression plasmids expressing mCherry alone or fused to RNAi-resistant (mouse) CITK wild type sequence ($n \geq 50$ cells in three independent experiments).

Data presented are means \pm SEM. Statistical significance was assessed using a two tails Student T-test. *** $P < 0.001$; ** $P < 0.01$; * $P < 0.05$.

Figure EV2. Analysis of CITK regions involved in the interaction with ASPM.

(A) Representation of domain structure of CITK and its fragments: Ser/Thr kinase domain (kinase), coiled-coil region (Coiled-coil), Rho-binding domain (RhoBD), type-2 zinc finger (ZF), pleckstrin homology domain (PH), CMG domain (CMG), proline-rich, putative SH3

binding domain (SH3B), PDZ binding domain (PDZB); full length CITK (FL), kinase dead full length CITK (KD), Citron-N (CN), the N-terminal (Nt) and the C-terminal halves of CITK (Ct).

B) Representation of domain structure of ASPM and CTR fragment: putative microtubule binding domain (MTB), calponin homology domain (CH), calmodulin binding domains (IQ), and the C-terminal region (CTR).

(C) The indicated Myc-CITK fusion constructs were cotransfected with ASPM-CTR-GFP in HEK293T. After 48 hours immunoprecipitations were performed from total cell lysates using anti-GFP antibody and the immunoprecipitated proteins were revealed by Western blotting for the GFP and Myc epitopes. The results shown are representative examples of at least three different experiments.

Figure EV3. Loss of CITK does not impair the localization of the dynein/dynactin motor complex and NuMA.

(A) Control and CITK depleted cells were immunostained for γ -tubulin (red), DNA (blue) and NuMA, Dynactin (p150) or Dynein (green). Images represent maximum intensity projections of confocal z-stacks. Scale bars, 5 μ m.

(B) Quantification of fluorescence intensity at spindle poles of NuMA, Dynactin (p150) or Dynein in HeLa cells transfected with either CTRL or CITK siRNA.

(C) Percentage of mitosis with cortical localization of NuMA or Dynactin (p150) in HeLa cells transfected with either CTRL or CITK siRNA. Data are presented as mean \pm SEM. Statistical significance was assessed using a two tails Student T-test. *, P < 0.05.

Figure EV4. Loss of CITK does not alter spindle length or symmetry.

(A) Quantification of spindle length in HeLa transfected with control sequence or with CITK-specific siRNA.

(B) Quantification of spindle symmetry in HeLa transfected with control sequence or with CITK-specific siRNA.

Differences in both panels were not statistically significant ($n \geq 150$ cells in three independent experiments). Statistical significance was assessed using a two tails Student T-test.

Movies EV1 and EV2. CITK depletion induces abnormal spindle orientation.

Mitotic spindles of control (Movie EV1) or CITK RNAi-treated (Movie EV2) HeLa cells were analyzed by time-lapse microscopy. Frames were taken every 5 min for 12 h.

Movie EV3. CITK localization during cell division.

HeLa cells expressing physiological levels of GFP-CITK from a stably integrated BAC transgene were analyzed by time-lapse microscopy. Frames were taken every 5 min for 12h.

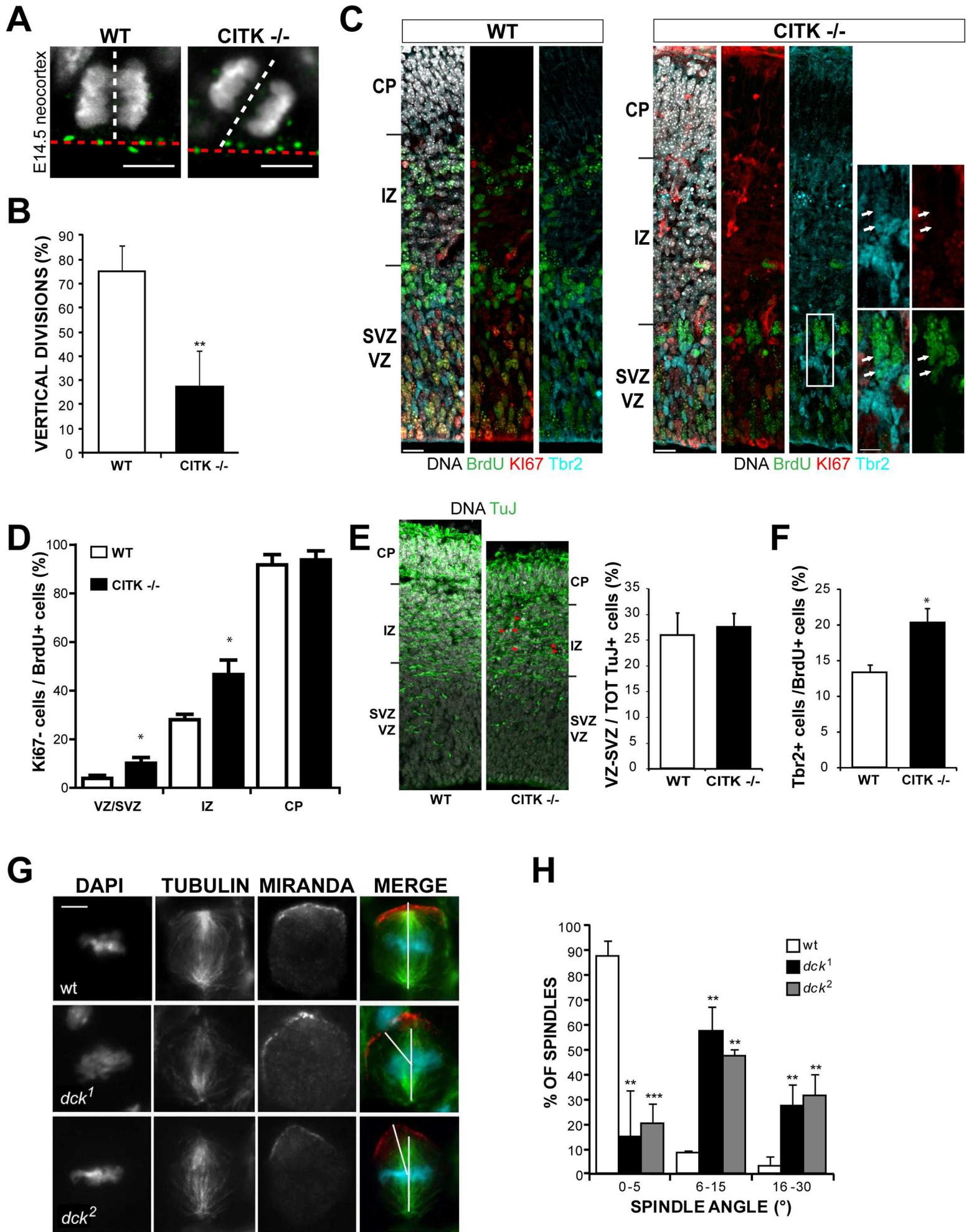
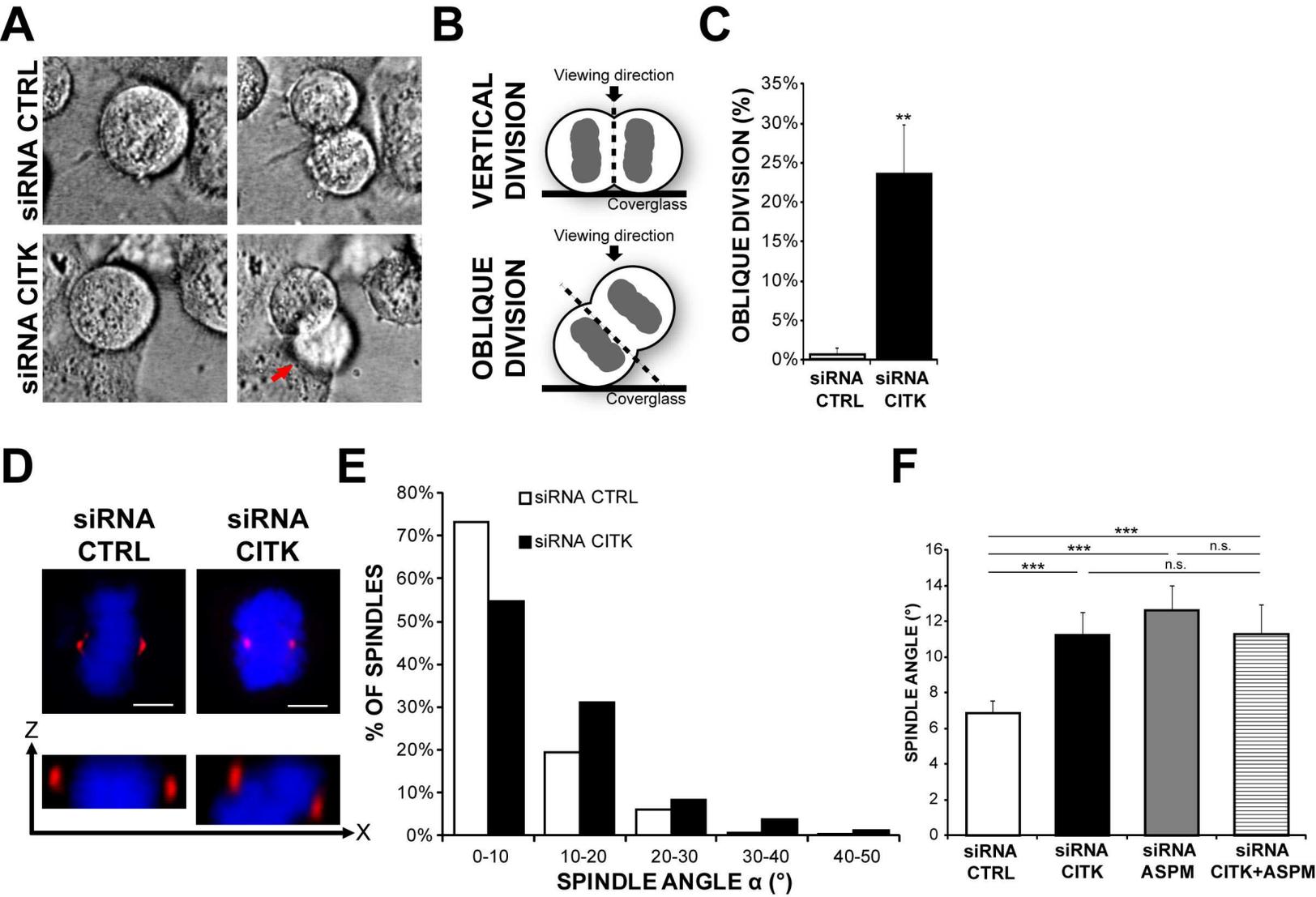


Figure 2

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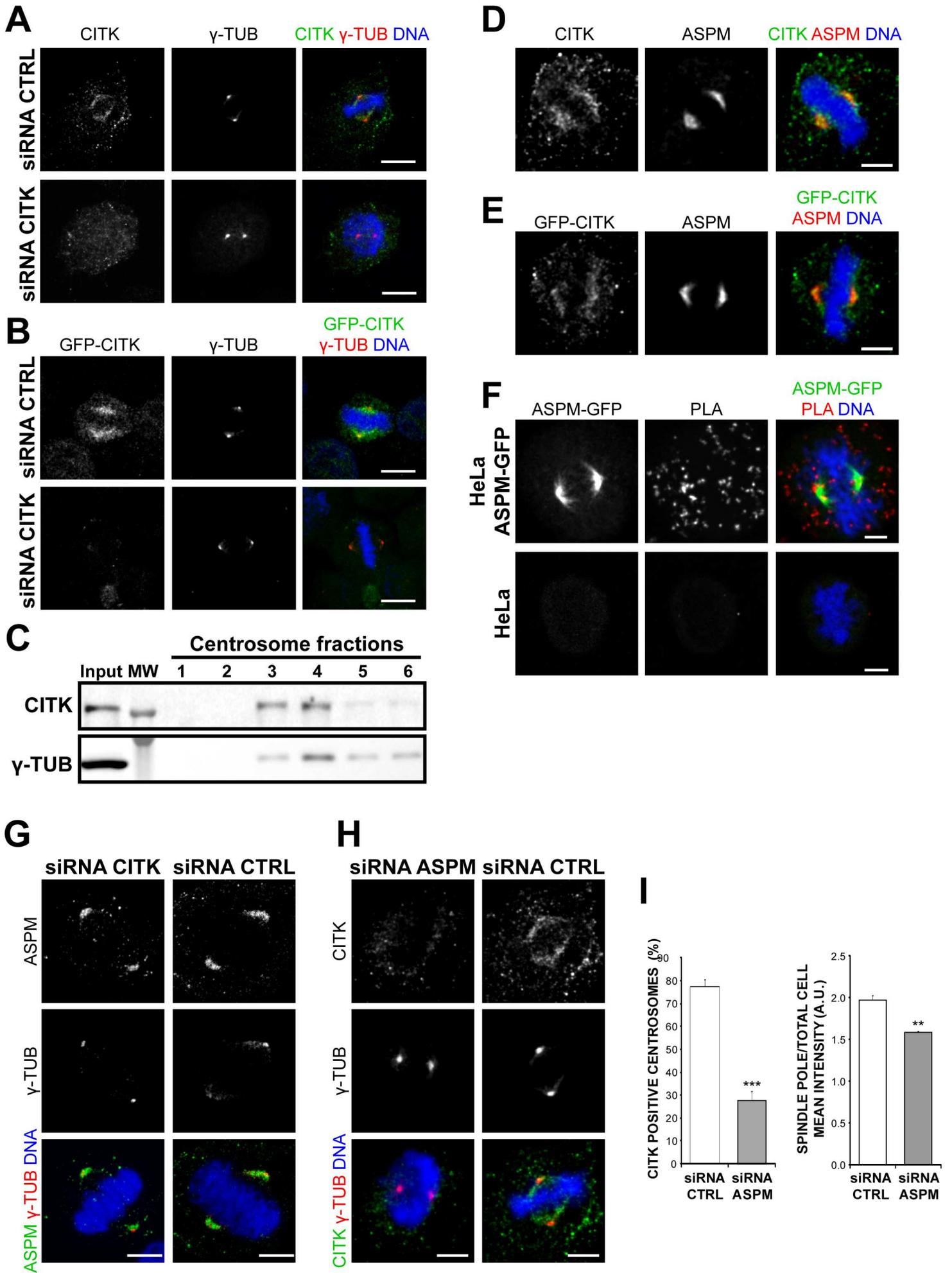
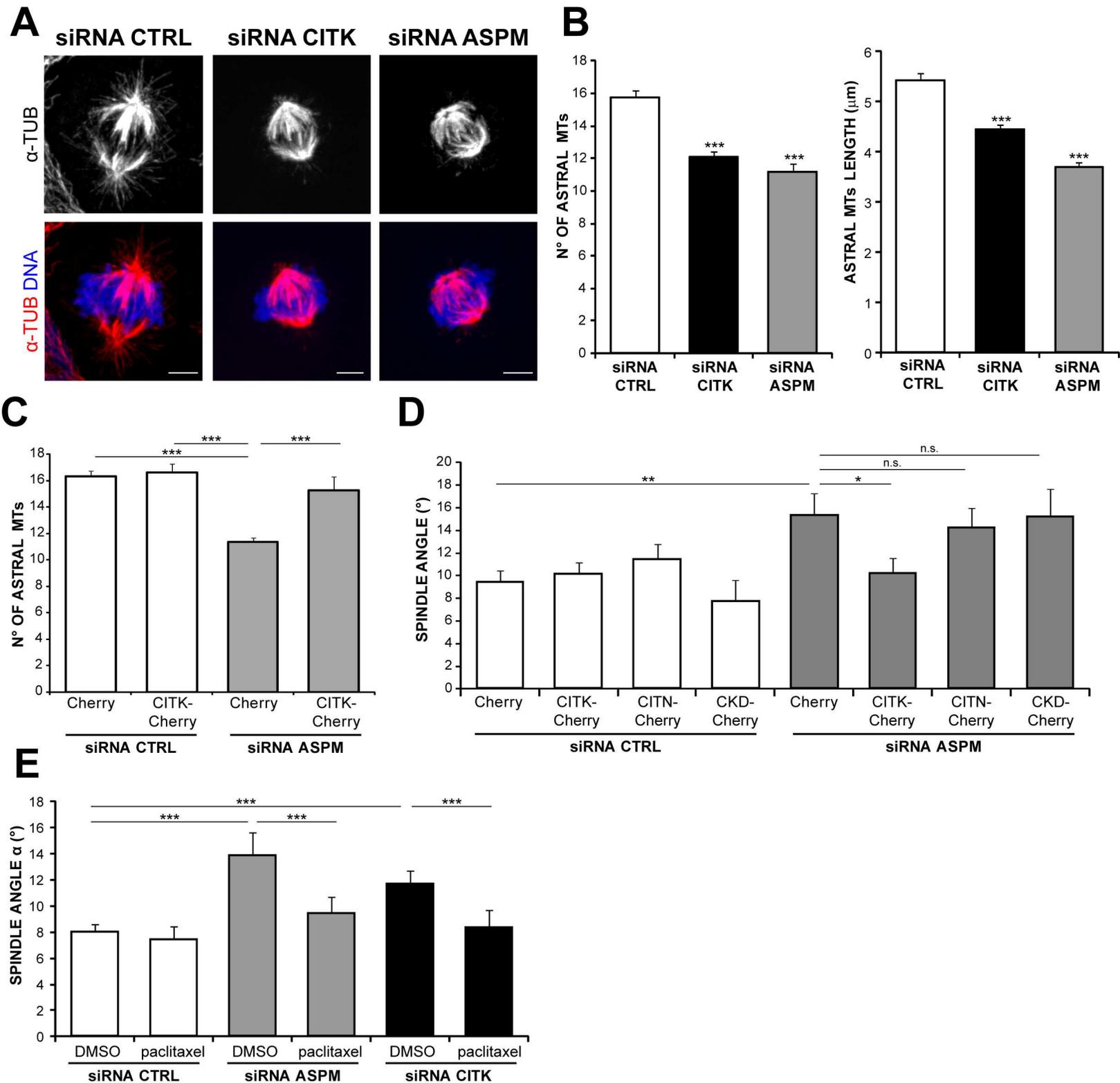


Figure 4

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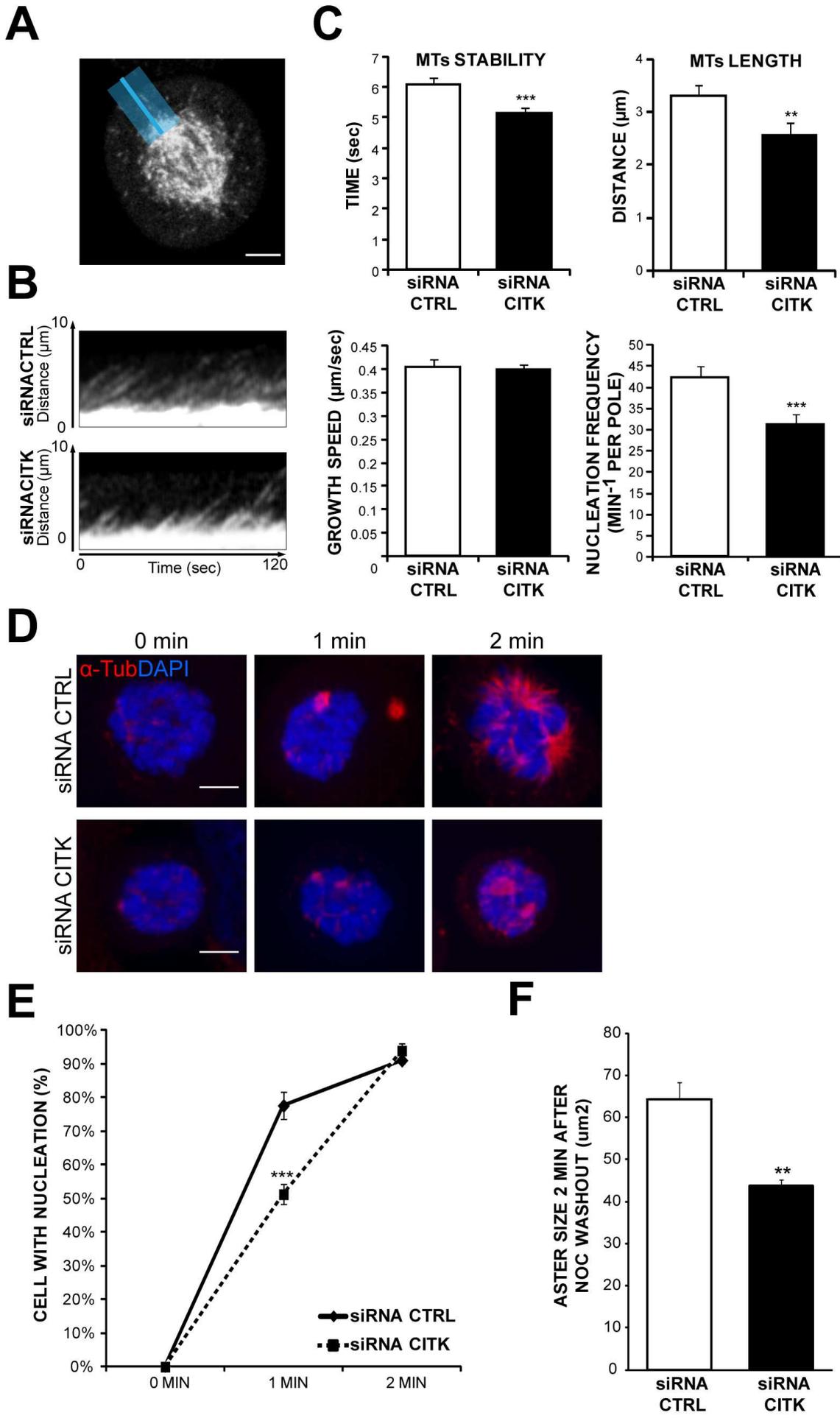
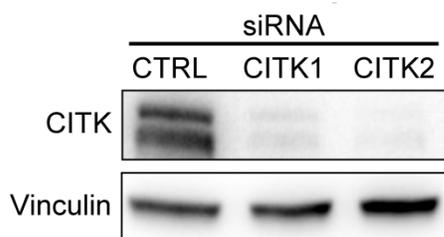


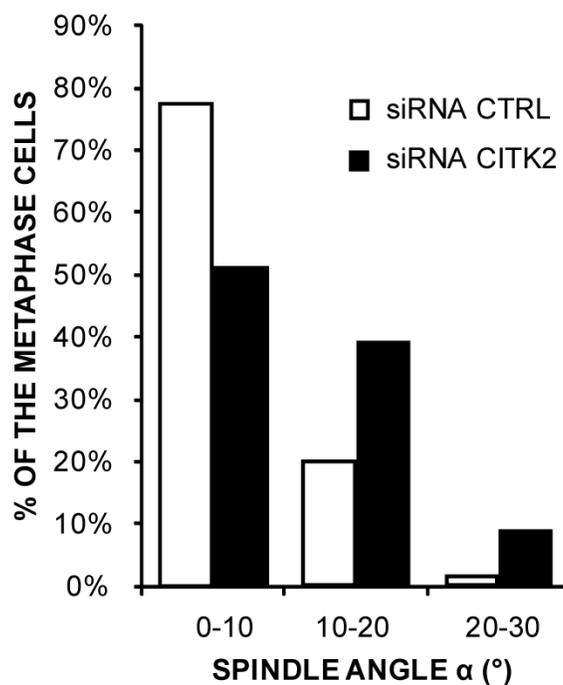
Figure EV1

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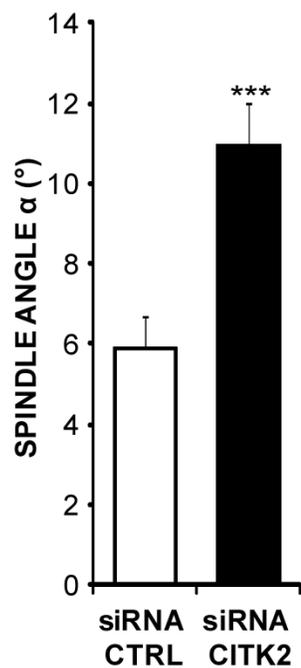
A



B



C



D

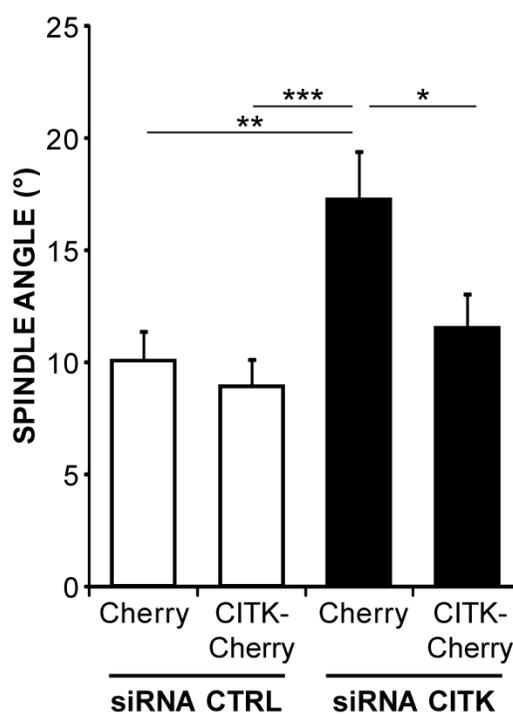
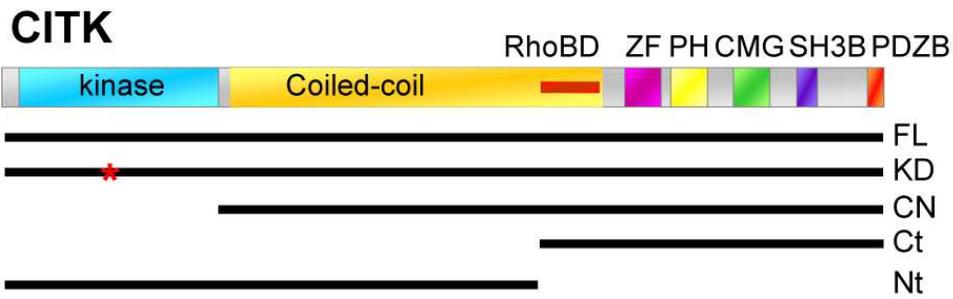


Figure EV2

A



B



C

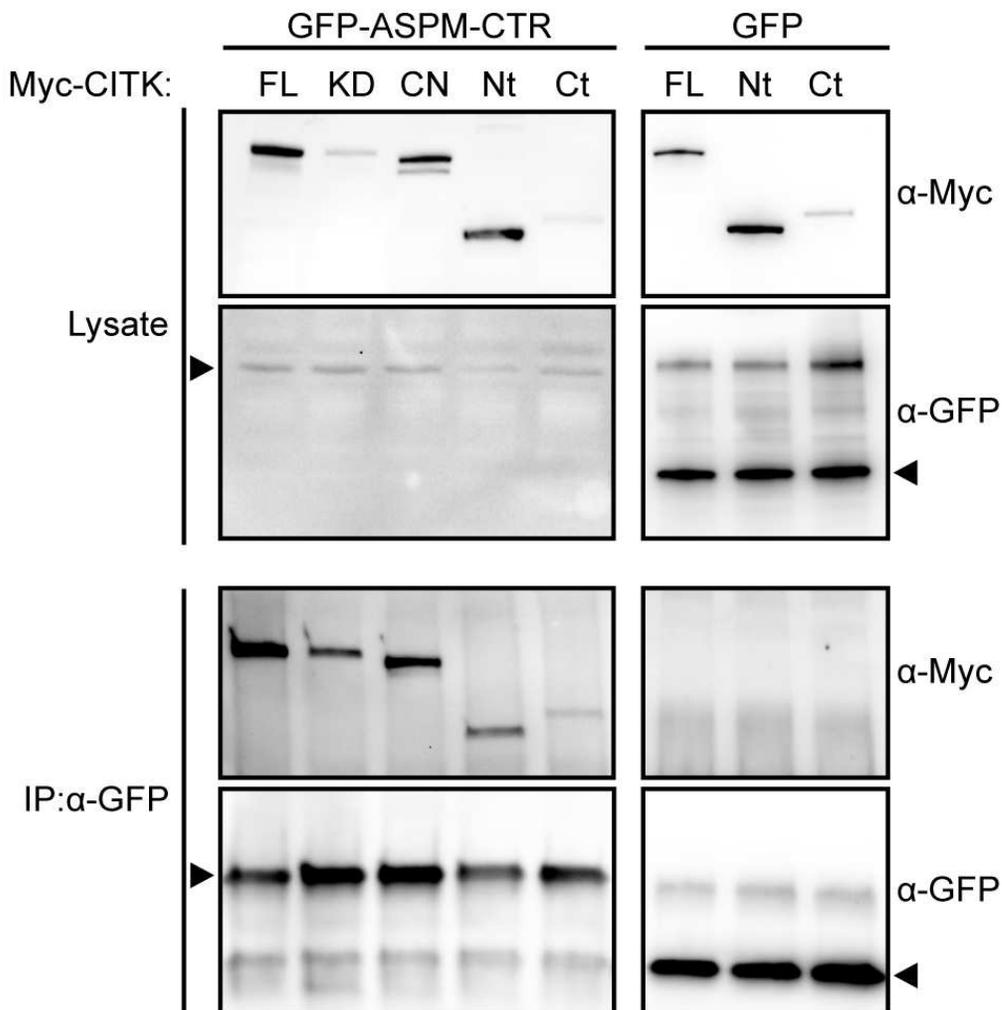


Figure EV3

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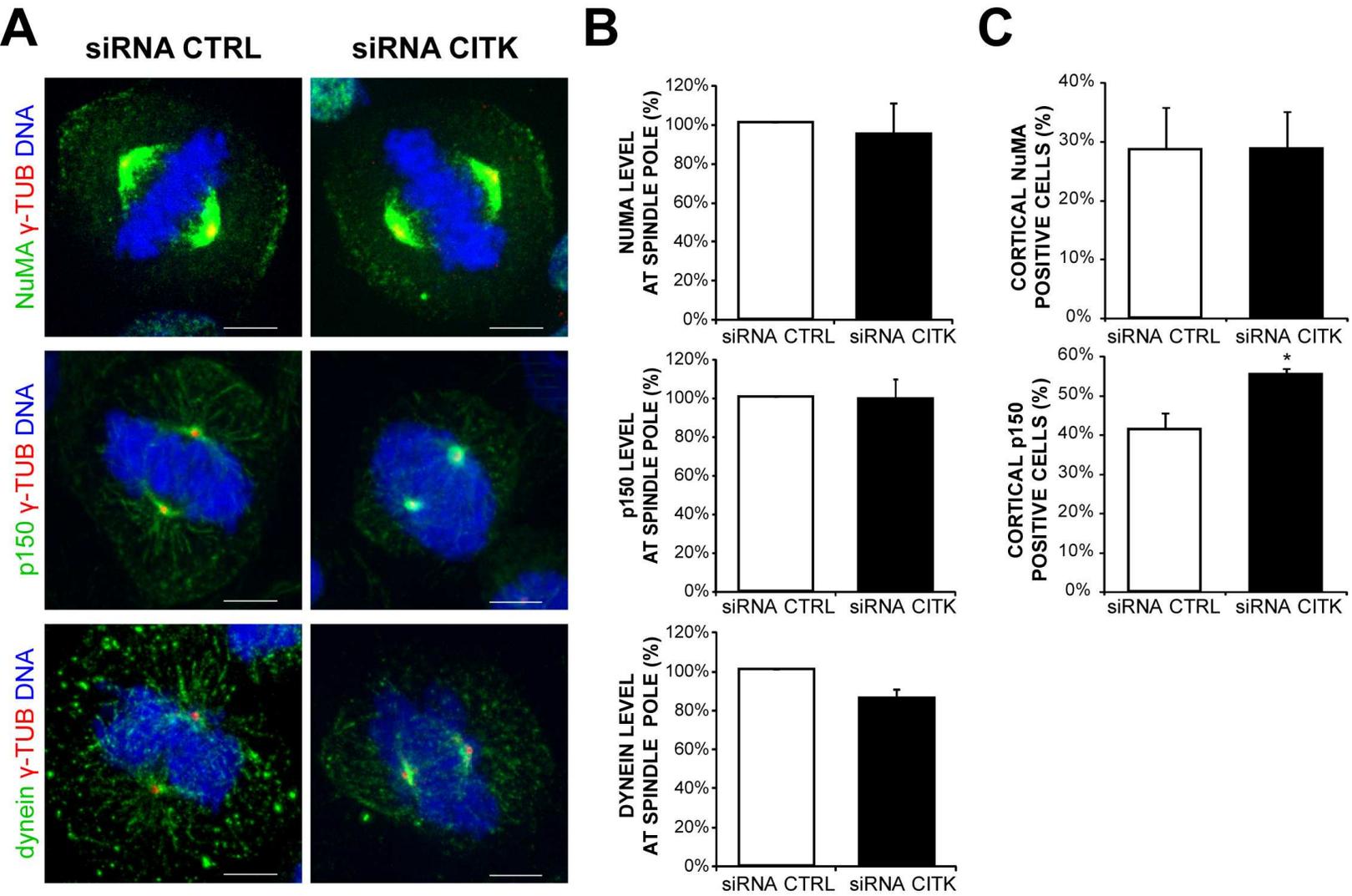
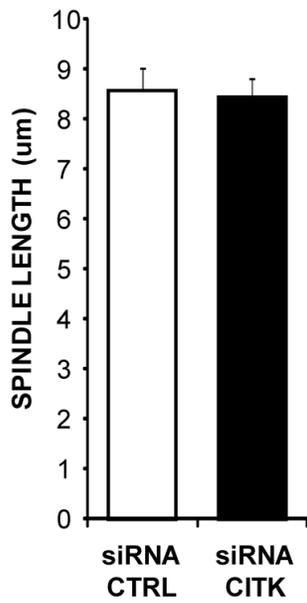


Figure EV4

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A



B

