Borealin: a novel chromosomal passenger required for stability of the bipolar mitotic spindle

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The chromosomal passenger complex of Aurora B kinase, INCENP, and Survivin has essential regulatory roles at centromeres and the central spindle in mitosis. Here, we describe Borealin, a novel member of the complex. Approximately half of Aurora B in mitotic cells is complexed with INCENP, Borealin, and Survivin; and Borealin binds Survivin and INCENP in vitro. A second complex contains Aurora B and INCENP, but no Borealin or Survivin. Depletion of Borealin by RNA interference delays mitotic progression and results in kinetochore-spindle misattachments and an increase in bipolar spindles associated with ectopic asters. The extra poles, which apparently form after chromosomes achieve a bipolar orientation, severely disrupt the partitioning of chromosomes in anaphase. Borealin depletion has little effect on histone H3 serine10 phosphorylation. These results implicate the chromosomal passenger holocomplex in the maintenance of spindle integrity and suggest that histone H3 serine10 phosphorylation is performed by an Aurora B–INCENP subcomplex.

Introduction

In recent years, it has emerged that chromosomal passengers (Earnshaw and Bernat, 1990), a conserved complex of Aurora B kinase, INCENP, and Survivin (Adams et al., 2000; Kaitna et al., 2000; Kang et al., 2001; Wheatley et al., 2001a; Bolton et al., 2002; Cheeseman et al., 2002; Leversen et al., 2002), are key regulators of mitotic events (for review see Andrews et al., 2003; Carmena and Earnshaw, 2003). In keeping with this role, chromosomal passengers show a dynamic localization pattern during mitosis, appearing on the chromosome arms and inner centromeres from prophase through metaphase and then transferring to the spindle midzone and midbody from anaphase through cytokinesis (Cooke et al., 1987). In yeast, relocalization of the chromosomal passenger complex to the central spindle in early anaphase is mediated by the dephosphorylation of Slj15 (INCENP) by Cdc14 (Pereira and Schiebel, 2003). The requirements for centromere targeting of the chromosomal passengers are still largely unknown, although Aurora A phosphorylation of CENP-A may contribute (Kunitoku et al., 2003).

Aurora B kinase activity peaks in mitosis (Bischoff et al., 1998; Terada et al., 1998), and its regulation involves both specific targeting and modulation of the kinase activity by the other chromosomal passengers. INCENP is a multi-domain microtubule-binding protein (Cooke et al., 1987; Wheatley et al., 2001b) that binds to Aurora B through the conserved COOH-terminal IN box (Adams et al., 2000; Honda et al., 2003). Binding stimulates Aurora B to phosphorylate INCENP within the IN box, resulting in further kinase activation in a feedback loop (Bishop and Schumacher, 2002; Cheeseman et al., 2002; Honda et al., 2003). Survivin has been shown to contribute to Aurora B activation in Xenopus laevis and fission yeast (Bolton et al., 2002; Leversen et al., 2002; Petersen and Hagan, 2003), although in vitro no direct stimulation was observed with the human proteins (Honda et al., 2003). Numerous chromosomal and cytoskeletal Aurora B substrates are now being discovered...
Interference with the subunits of the chromosomal passenger complex produces severe mitotic defects. A profound inhibition of chromosome congression to the metaphase plate (Adams et al., 2001; Ditchfield et al., 2003; Hauf et al., 2003) reflects an involvement of Aurora B in the correction of syntelic chromosome attachments (Tanaka et al., 2002; Hauf et al., 2003; Andrews et al., 2004; Lampson et al., 2004). A requirement for chromosomal passengers for spindle assembly checkpoint function was uncovered in yeast through analysis of Ipl1 (Aurora) and Bir1 (Survivin-like) function (Biggins and Murray, 2001; Petersen and Hagan, 2003). Interference with passenger function in vertebrate cells compromises the spindle assembly checkpoint when microtubules are perturbed using taxol or monastrol (Carvalho et al., 2003; Ditchfield et al., 2003; Hauf et al., 2003; Lens et al., 2003). Aurora B, INCENP, and Survivin are also essential for the late stages of cytokinesis (Terada et al., 1998; Kaitna et al., 2000; Severson et al., 2000; Adams et al., 2001; Giet and Glover, 2001; Carvalho et al., 2003; Lens et al., 2003).

A fourth protein, TD-60, was described as a chromosomal passenger based on its localization during mitosis (Andreasen et al., 1991). Aurora B and TD-60 are interdependent for localization (Mollinari et al., 2003), but it is presently unclear if TD-60 is part of the chromosomal passenger complex. TD-60 appears to be a guanosine nucleotide exchange factor that may act on Rac1 (Mollinari et al., 2003).

Here, we report the initial characterization of Borealin, a novel fourth subunit of the chromosomal passenger complex in vertebrates. Functional studies reveal that Borealin is required for targeting of Aurora B, INCENP, Survivin, and TD-60; correction of kinetochore attachment errors; and stability of the bipolar spindle in human cells. Our data suggest that there are at least two distinct chromosomal passenger complexes during mitosis.

Results

Borealin is a novel chromosomal passenger protein

A proteomic screen for novel human proteins associated with histone-depleted mitotic chromosomes (unpublished data) identified the uncharacterized protein FLJ10468. A GFP fusion of FLJ10468 showed a localization pattern during mitosis identical with that of chromosomal passenger proteins. It appeared on the chromosome arms and the inner centromeres in prophase (Fig. 1 A) and concentrated at the inner centromeres during metaphase (Fig. 1 A’). It then relocated to the spindle midzone at anaphase onset and remained at the midbody throughout telophase (Fig. 1 A”) and cytokinesis.
Affinity-purified rabbit polyclonal antibody R1647 raised against the full-length protein recognized a protein at the expected size (31 kD) on immunoblots of HeLa cell lysates (Fig. 1 C). Indirect immunofluorescence confirmed the dynamic localization of this protein in mitosis (Fig. 1, E–E”). Simultaneous staining for Aurora B confirmed that the two proteins colocalize throughout mitosis (Fig. 1, B and B’; and Fig. S1, available at http://www.jcb.org/cgi/content/full/jcb.200404001/DC1). These results demonstrate that the protein, which we termed Borealin, is a novel chromosomal passenger.

Chromosomal passenger proteins Aurora B, INCENP, and Survivin are expressed in a cell cycle–dependent manner (Li and Altieri, 1999; Honda et al., 2003). Immunoblotting of extracts from HeLa cells arrested in S-phase (Fig. 1 D, S) or mitosis (Fig. 1 D, M) revealed that Borealin levels are also significantly increased in mitosis. Northern blotting detected a corresponding increase in mRNA levels (unpublished data). A slower migrating form of INCENP, presumably the phosphorylated protein, was detected solely in mitotic cells.

**Borealin is conserved among vertebrates with distantly related proteins in insects and nematodes.** Multiple sequence alignment and secondary structure (A; Ginalski et al., 2003) plus tentative tree (B; Brodskii et al., 1995) for the vertebrate Borealin family. Colors indicate similarity of >60% (yellow) and >80% (green). Secondary structure predicted consistently for Borealin 1 (blue) is indicated (H, helix; E, strand), with plausible core elements underlined based on their length (H >3; E >2) and presence in all sequences. NCBI/SPTrEMBL accession codes: Human, Hs_Bor,cdc-associated protein 8, NP_060571/ Q9NVW5 [cDNA, chromosome 1]; Hs_hypBor, likely pseudogene, Hs7_34123 (5467829–5468473) [genomic DNA chromosome 7]. Mouse, Mm_Bor, BAC37258/Q8BH4Q [cDNA chromosome 4]. Chicken, Gg_Bor1, (BU137562 + BU317190) [EST]; Gg_Bor2, (BU315188 + BU328208 + BU324297 + BU225870) [EST], X. laevis, xl_Bor1 (Bu626030 + Bu642424) [EST], xl_Bor2, CA973941/CA982691 [EST]. Zebrafish, Dr_Bor1, CK683767 [cDNA]; Dr_Bor2, BM777478 + BM80944. Fugu, Fr_Bor, SIFRUP000000052671 corrected in one exon [pred. protein]. Caenorhabditis, Ce_CSC-1, O45952/ CAB07696. Mosquito, Ag_Borealin-related1, XP_309425 [pred. protein]; Ag_Borealin-related2, XP_309424 [pred. protein]. Drosophila, Dr_Borealin-related, CG4454-PB [pred. protein].
Furthermore, a manual alignment with the *Caenorhabditis elegans* chromosomal passenger protein CSC-1 revealed weak sequence similarities within two short regions of Borealin. Therefore, we tentatively include these proteins in a Borealin family tree (Fig. 2 B). We have been unable to find Borealin-like proteins in plants and yeasts.

**Borealin localization is disturbed in Survivin-depleted cells and in cells expressing a dominant INCENP mutant**

To determine whether or not Borealin exhibits functional interactions with the other chromosomal passengers, we exploited the mutual interdependence of these proteins for their localization in vivo. In cells depleted of Survivin by RNAi, Aurora B is localized diffusely in the cytoplasm in mitosis (Carvalho et al., 2003). Therefore, we transfected HeLa cells with Survivin-specific siRNAs and used Aurora B delocalization as a marker to identify Survivin-depleted cells. Borealin was delocalized in all cells in which Aurora B was delocalized (Fig. 3 B), but was localized correctly in cells transfected with control oligo (Fig. 3 A).

The chicken INCENP deletion mutant INCENP1–405 disrupts chromosomal passenger localization when overexpressed in vertebrate cells (Mackay et al., 1998). We transfected HeLa cells with the INCENP1–405 construct and identified cells overexpressing the protein by the abnormal localization of endogenous INCENP (Fig. 3 C, right). In these cells, cotransfected Borealin-GFP was diffusely localized on the chromosomes with little or no detectable enrichment at centromeres (Fig. 3 C, left). In anaphase cells expressing INCENP1–405, Borealin-GFP and INCENP remained associated with the chromosomes (Fig. 3 D). Interestingly, TD-60 also showed a diffuse chromosomal localization in the presence of INCENP1–405 (Fig. 3, E and F), as did Survivin and Aurora B (not depicted).

We conclude that Borealin localization is dependent on the correct localization of Survivin and INCENP, suggesting that Borealin is functionally linked with the chromosomal passenger complex.

**Borealin is a subunit of the chromosomal passenger complex in vivo and associates with INCENP and Survivin in vitro**

We then investigated whether or not Borealin is a subunit of the chromosomal passenger complex by immunoprecipitating the endogenous protein from mitotic cell lysates. Aurora B, INCENP, and Survivin coimmunoprecipitated with Borealin (Fig. 4 A). About half of Aurora B and most of INCENP and Survivin could thus be depleted from mitotic cell lysates. In contrast, TD-60 did not coimmunoprecipitate with Borealin. To test if the INCENP, Aurora B, and TD-60 remaining soluble after Borealin immunoprecipitation were present in a complex, we subjected the supernatant to a second round of immunoprecipitation with antibody to Aurora B. Under these conditions, INCENP was readily detected on the beads; however, Borealin, Survivin, and TD-60 were not (Fig. 4 A). The TD-60 antibody, made against a COOH-terminal peptide and affinity purified, recognized bands of 56 and 45 kD in immunoblots of mitotic cells. Pre-

...incubation of the antibody with the peptide abolishes both bands (unpublished data), and thus both may be isoforms of TD-60.

In vitro binding experiments revealed that GST-Borealin can interact directly with in vitro translated INCENP and Survivin, but not Aurora B or TD-60 (Fig. 4 B). Interestingly, Borealin bound readily to itself. Quantification of the binding experiments revealed that the interaction with Survivin is particularly efficient (Fig. 4 C). When the NH2- and COOH-terminal halves of Borealin (1–141 and 142–280 aa) were used for similar binding experiments (Fig. 4 D), neither half bound INCENP but both halves bound Borealin. Borealin1–141 bound Survivin as efficiently as full-length Borealin, whereas only a weak interaction was detected for Borealin142–280. Thus, the two halves of Borealin interact differentially with subunits of the chromosomal passenger complex.
This analysis suggests that during mitosis, cells may have at least two chromosomal passenger complexes, a holocomplex containing INCENP, Aurora B, Borealin, and Survivin and a subcomplex containing INCENP and Aurora B.

A Borealin fragment selectively perturbs chromosomal passenger targeting to centromeres in vivo

Knowing which regions of Borealin mediate interactions with other chromosomal passengers, we probed the functional significance of these interactions by transiently transfecting HeLa cells with a series of Borealin deletion mutants fused to GFP. Note that full-length Borealin (280 aa) localizes correctly with GFP at either end (Fig. 1, A and B; and Fig. 5 A).

All deletion mutants (GFP-Borealin1–141, GFP-Borealin142–280, GFP-Borealin89–280, and Borealin142–280-GFP) were nuclear in interphase (Fig. 5, B–E), confirming that the predicted nuclear localization signals, one in each half of the protein, are functional in vivo. The nucleolar localization of these GFP fusion proteins is probably an artifact, as both R1647 and another polyclonal antibody against Borealin (not depicted) fail to stain nucleoli (Fig. S1 A).

None of the deletion mutants localized to centromeres. Therefore, the NH$_2$-terminal 88 amino acids are necessary, but not sufficient, for centromere targeting (Fig. 5, D and E). GFP-Borealin1–141 transferred to the spindle midzone normally, whereas the more extensive deletion GFP-Borealin1–88 did not (Fig. 5, B and D). Both Borealin142–280-GFP and Borealin89–280-GFP failed to localize to the spindle midzone (Fig. 5, C and E).

INCENP, Survivin, Aurora B, and TD-60 were delocalized from centromeres in cells expressing GFP-Borealin1–141 (Fig. 5, G and H) and GFP-Borealin1–88 (not depicted), but not in cells expressing Borealin142–280-GFP (Fig. 5, I and J) or Borealin89–280-GFP (not depicted). Interestingly, GFP-Borealin1–141, although interfering with centromere targeting, did not prevent the other chromosomal passengers from accumulating at the spindle midzone. The chromosomal passengers also transferred normally to the spindle midzone in cells expressing Borealin142–280-GFP and Borealin89–280-GFP. In cells transfected with GFP-Borealin1–88, the few anaphase cells found expressed very low amounts of the deletion construct, suggesting that this protein is toxic. The chromosomal passengers in these cells...
localized to the midzone, albeit at reduced levels (unpublished data).

These observations suggest that the NH2-terminal half of Borealin, but not the COOH-terminal half, interacts with chromosomal passengers in vivo and can perturb their targeting to centromeres but not to the spindle midzone.

**Depletion of Borealin by RNA interference causes spindle abnormalities and failure of cytokinesis**

To examine the contribution of Borealin to mitotic regulation, we depleted the protein by RNA interference in HeLa cells. Cells were synchronized in S-phase, released, and subsequently analyzed upon entry into the first round of mitosis after exposure to siRNA. Immunoblotting showed that Borealin was significantly depleted in cells treated with Borealin siRNA (Fig. 6 A). Levels of Survivin and phosphorylated INCENP also fell, but no significant change in Aurora B, Aurora A, TD-60, or histone H3 phosphorylation at serine10 was observed. Indirect immunofluorescence showed that >95% of cells exposed to Borealin siRNA had no detectable Borealin in mitosis. INCENP, Survivin, Aurora B, and TD-60 were all delocalized in Borealin-depleted cells (Fig. 6, D–G). As observed for other chromosomal passengers (Adams et al., 2001; Carvalho et al., 2003), depletion of Borealin caused cells to accumulate in prometaphase with a corresponding decrease in later mitotic stages (Fig. 6 B).

To analyze the effect of Borealin depletion on mitotic progression, cells were grown on grid coverslips. Starting 6 h after their release from the S-phase block, phase-contrast images of several groups of cells were taken at 1-h intervals for a total of 7 h. On coverslips treated with control siRNA, 92% of cells went through mitosis and completed cytokinesis successfully (Fig. 6 C; representative cells shown in Fig. S2 and Fig. S3, available at http://www.jcb.org/cgi/content/full/jcb.200404001/DC1). In contrast, only 23% of cells treated with Borealin siRNA completed cytokinesis, whereas 41%

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**Figure 5.** Transient expression of Borealin deletion mutants identifies functional domains in vivo. (A–E) Localization of deletion mutants expressed as GFP fusions in interphase (left), prometaphase/metaphase (middle), and anaphase (right). HeLa cells were transfected with GFP constructs containing Borealin amino acids 1–280 (A), 1–141 (B), 142–280 (C), 1–88 (D), and 89–280 (E) and processed for fluorescence microscopy after 36 h. (F) Summary of Borealin deletion and endogenous chromosomal passenger localization in mitosis. Cen, centromere; mdz, anaphase spindle midzone. (G–I) Effect of Borealin deletion expression on the localization of endogenous chromosomal passengers at prometaphase/metaphase. Transfected cells were stained for Survivin (G and I) and TD-60 (H and J) 36 h after transfection. Bars, 5 μm.
failed cytokinesis, 33% underwent apoptosis, and 3% stayed in mitosis longer than 3 h. The average duration of mitosis was 1.3 ± 0.2 h for control cells and 2.6 ± 0.1 h for cells treated with Borealin siRNA (n = 3).

Closer examination of mitotic phenotypes in Borealin-depleted cells revealed defective interactions between kinetochores and the mitotic spindle. Among the many misaligned chromosomes observed (Fig. 7, B–B”), we noted both syntelic attachments (both kinetochores attached to the same pole; Fig. 7, B” and D) as well as merotelic attachments (one kinetochore attached to both poles; Fig. 7, C–C”). Merotelic attachments cause kinetochores to lag in the spindle midzone during anaphase, and 21 of 21 anaphases observed after Borealin RNAi had such lagging kinetochores (Fig. 7 E).

In addition to kinetochore misattachments, we also observed spindle abnormalities in Borealin-depleted cells. In addition to a main bipolar spindle, 40% of mitotic cells exhibited ectopic asters containing Aurora A kinase and γ-tubulin (Fig. 7, G and H; and Fig. 8, B–D). Importantly, these extra poles did not represent centrosome amplification resulting from prior failed cytokinesis, as the images shown were collected in the first mitosis following release from the S-phase block. They were also unlikely to be due to the S-phase block itself, as they were only rarely observed in cells transfected with the control oligo (Fig. 8 D).

To investigate the effects of these spindle abnormalities on mitotic chromosome dynamics, we examined cells treated with Borealin siRNA by live cell imaging. A total of 14 HeLa cells expressing histone H2B-GFP were individually followed in mitosis after exposure to siRNA by capturing a stack of DIC and FITC images every 7 min. None of these cells completed mitosis normally.

The most common phenotype, observed in 8 of 14 cells, was extremely unusual. These cells progressed through the early stages of mitosis normally and ultimately formed an apparently normal metaphase plate (Fig. 8 E, frame 2:01). The plate often appeared to become frayed at either end (Fig. 8 E, frame 3:26). Eventually, when the cells entered anaphase, the sister chromatids abruptly segregated from the apparently bipolar metaphase plate in several differing direc-
tions toward what appeared to be multiple spindle poles (Fig. 8 E, frame 3:33; and Fig. S4, A, C, and D, available at http://www.jcb.org/cgi/content/full/jcb.200404001/DC1). Other cells had lagging chromosomes in anaphase and clearly failed cytokinesis (Fig. S4 B). None of the seven control cells we followed exhibited any of these defects (unpublished data).

Thus, the chromosomal passenger complex containing Borealin has an essential role in maintaining the bipolar configuration of the mitotic spindle.

Borealin is phosphorylated by Aurora B in vitro
Aurora B kinase activity is stimulated by INCENP binding (Kang et al., 2001; Bishop and Schumacher, 2002; Honda et al., 2003). Therefore, we asked whether or not Borealin had any effect on Aurora B kinase activity in the presence of INCENP and Survivin using myelin basic protein as substrate. We expressed recombinant proteins containing either an NH$_2$-terminal GST (Aurora B and INCENP) or His (Survivin and Borealin) tag in insect cells coinfected with the corresponding baculoviral constructs. As expected, Aurora B kinase activity was greatly increased when it was copurified from insect cells with INCENP and Survivin (using glutathione sepharose beads) compared with Aurora B on its own (Fig. 9 A). We then copurified Aurora B/INCENP/Survivin/Borealin from insect cells and compared the resulting kinase activity with that exhibited by Aurora B in the presence of INCENP and Survivin. Coexpressing Borealin with the other subunits did not alter Aurora B kinase activity appreciably, nor did the addition of separately purified Borealin to Aurora B/INCENP/Survivin (Fig. 9 B).

When we added bacterially expressed Borealin to Aurora B/INCENP/Survivin, the full-length protein GST-Borealin$_{1-280}$ was readily phosphorylated (Fig. 9 C). GST-Borealin$_{142-280}$ was also phosphorylated, but GST-Borealin$_{1-141}$ was not (Fig. 9 C). A serine to alanine mutation at one predicted phosphorylation site (165 aa) resulted in a reduction, but not the abolition, of phosphorylation, suggesting that Borealin is phosphorylated at more than one site.

Discussion
Borealin is an essential subunit of the chromosomal passenger holocomplex in mitosis
Borealin is a novel 31-kD chromosomal passenger protein with homologues in all vertebrates examined, as well as more distant relatives in mosquito, D. melanogaster, and, possibly, C. elegans.

C. elegans CSC-1 was identified in a genetic screen for mutations affecting chromosome segregation and cytoki-
Borealins is a novel chromosomal passenger complex, and subsequently found to associate with the chromosomal passenger complex (Romano et al., 2003). Similarity between the vertebrate Borealins and CSC-1 is confined to short domains near the NH2 and COOH termini, and the Borealin sequence does not have the prominent imperfect repeat found in CSC-1. Furthermore, the net charge of the two proteins is very different (pI 9.9 for Borealin compared with pI 5.1 for CSC-1), and Borealin binding to Survivin does not require Zn2+
(unpublished data). Thus, it is not clear whether CSC-1 is a true orthologue of Borealin.

Like Aurora B, INCENP, and Survivin (Li et al., 1998; Honda et al., 2003), Borealin is cell-cycle regulated. This confirms the findings of a previous study, which identified eight human transcripts, including that for Borealin, on the basis of their coexpression with known cell cycle genes including CDC2, MCAK, and topoisomerase II (Walker, 2001). Borealin was also one of 50 ESTs identified in a genomic screen for mRNAs expressed in G2/M similar to Aurora A and B (Tien et al., 2004).

Borealin is a novel member of the chromosomal passenger holocomplex, together with Aurora B, INCENP, and Survivin (Adams et al., 2000; Kaitna et al., 2000; Kang et al., 2001; Bolton et al., 2002; Cheeseman et al., 2002; Leverson et al., 2002). Based on our in vitro binding data, Survivin and INCENP are likely to be the major binding partners of Borealin within the complex. Borealin associates with itself in vitro and may be a multimer in the complex. Importantly, almost all of the Survivin is complexed with Borealin in colcemid-arrested cells.

Figure 8. Live imaging of Borealin-depleted cells reveals severe defects in the partitioning of chromosomes in anaphase. (A–C) β- and γ-tubulin costaining of Borealin-transfected cells (B and C) and a control cell (A). (D) Quantification of multipolar mitotic cells (n = 3; total number of cells scored: control siRNA, 138; Borealin siRNA, 144). (E) Selected frames of a representative live cell imaging experiment performed on mitotic HeLa cells expressing histone H2B-GFP treated with Borealin siRNA. Bar, 5 μm.
Our data suggest that there are multiple chromosomal passenger complexes during mitosis and that these contain differing stoichiometries of Aurora B, INCENP, Survivin, and Borealin. For example, in addition to a holocomplex containing all four passengers, we obtained evidence for an Aurora B/INCENP subcomplex that does not contain Survivin or Borealin. This subcomplex may be responsible for serine\(^{10}\) phosphorylation of histone H3, as this modification was not significantly decreased following RNAi for Borealin or Survivin (unpublished data). In contrast, the holocomplex is required for the correction of kinetochore attachment errors and for bipolar spindle stability. We believe that complexes containing chromosomal passengers might be highly dynamic, and their composition may vary at differing times and intracellular locations during mitosis.

**Borealin may target the passenger holocomplex to centromeres**

The dramatic movements of the chromosomal passengers during mitosis are thought to reflect the spatio-temporal regulation of Aurora B kinase activity throughout the mitotic cell (Carmena and Earnshaw, 2003), and the underlying mechanisms are therefore of considerable significance. We observed that when the NH\(_2\)-terminal half of Borealin is overexpressed in HeLa cells, the endogenous chromosomal passengers are unable to accumulate at centromeres, whereas spindle midzone targeting is unaffected. This effect is unlikely to be due to a disruption of kinase activity of the complex because the ability of INCENP to stimulate Aurora B kinase activity in vitro is unaffected by the presence of excess NH\(_2\)-terminal Borealin (unpublished data). The selectivity with which the mutant interferes with the localization of the endogenous chromosomal passengers is consistent with a specific role for Borealin in centromere targeting and implies that centromere and spindle midzone targeting may involve distinct mechanisms.

We have used chromosomal passenger complexes copurified from baculovirus-infected Sf-9 cells to ask if the presence of Borealin has a stimulatory or inhibitory effect on the activity of Aurora B kinase. Like INCENP, Borealin is phosphorylated in its COOH-terminal domain by Aurora B in vitro. When comparable amounts of Aurora B, INCENP, and Survivin were used in the assays, the presence of Borealin did not result in a detectable change of kinase activity. Therefore, the major role of Borealin is unlikely to involve direct modulation of Aurora B kinase activity. However, it is still conceivable that Borealin plays a more subtle role, as we did observe a substantial decrease in the levels of a mitosis-specific slower migrating form of INCENP in Borealin-depleted cells.

**Borealin is required for kinetochore error correction and stability of the bipolar mitotic spindle**

Aurora B has a major role in the correction of kinetochore orientation defects in mitosis, and particularly in the resolution of syntelic attachments (Tanaka et al., 2002; Hauf et al., 2003; Lampson et al., 2004). Therefore, it was not surprising that syntelic chromosomes were observed in cells depleted of Borealin. More surprising was the marked increase in chromosomes exhibiting merotelic attachments, which are not thought to be readily detected by the mitotic checkpoint and have been proposed to be a major source of human aneuploidy (Cimini et al., 2001). These results imply that the chromosomal passenger complex has a significant role in the prevention and/or resolution of merotelic attachments and strengthen the notion of the chromosomal passengers as master overseers of kinetochore orientation during mitosis. The underlying mechanisms remain to be determined, but regulation of factors that influence microtubule behavior such as the KinI kinesin MCAK (Andrews et al., 2004; Kline-Smith et al., 2004; Lan et al., 2004) or its regulators (Ohi et al., 2003) are strong possibilities.

The roles of chromosomal passengers in kinetochore function and cytokinesis are widely documented, but less is known about their role in spindle structure and stability.
Borealin is a novel chromosomal passenger

Here we have shown by live imaging that deletion of Borealin results in a characteristic phenotype in which cells align their chromosomes on what appears to be a nearly normal metaphase plate, but then abruptly undergo multipolar anaphases. In fixed images, small ectopic poles can be seen beside otherwise normal appearing bipolar spindles. These poles, which bind both Aurora A and γ-tubulin, are associated with microtubules, yet the live imaging data suggest that they do not orient significant numbers of chromosomes during prometaphase, as near normal metaphase chromosome alignment is observed. Instead, ectopic asters appear to interact with the chromosomes only after a nearly normal bipolar alignment has been achieved, so that at anaphase onset the chromosomes abruptly leave the metaphase plate in multiple directions.

**TD-60: a link between Aurora B and small GTPases?**

Our data strongly suggest that there is an intimate functional link between TD-60 and the chromosomal passenger complex. TD-60 colocalizes with the chromosomal passengers (Martineau-Thuillier et al., 1998), and in a recent study, depletion of Aurora B by RNAi was shown to perturb TD-60 localization in cells (Mollnari et al., 2003). We find that whenever passenger localization is perturbed, either by transient expression of dominant-negative INCENP or Borealin mutants or by depletion of Borealin, the localization of TD-60 shows an identical perturbation. The nature of the functional link remains to be determined and may not involve direct binding of TD-60 to the complex, as we have been unable to demonstrate a physical association between TD-60 and the other chromosomal passengers by immunoprecipitation.

TD-60 is structurally homologous to RCC1, a guanine nucleotide exchange factor, and has been shown to bind Rac1 (Mollnari et al., 2003). Therefore, the available data suggest a role for the small GTPases as either regulators or effectors of Aurora B kinase.

**Conclusions**

The present study has revealed new levels of complexity and regulation of the chromosomal passengers. The discovery of Borealin has raised the number of components of the vertebrate chromosomal passenger complex to four, and the demonstration of close functional links with TD-60 has strengthened the possibility of regulation of Aurora B kinase by small GTPases. Furthermore, our data suggest that there are at least two chromosomal passenger complexes in mitotic cells. The chromosomal passenger holocomplex containing Borealin is required for a range of kinetochore functions, including the resolution of merotelic attachments. Borealin is also required for the stability of the bipolar spindle during mitosis. Thus, the chromosomal passengers are now firmly established as master regulators of mitotic events.

**Materials and methods**

**Cloning and expression of recombinant proteins**

The cDNAs and baculoviral constructs for Aurora B, Survivin, and INCENP have been described previously (Wheatley et al., 2001a; Honda et al., 2003). The cDNA of Borealin and TD-60 were obtained by RT-PCR using total RNA prepared from HeLa S3 cells. The PCR product was introduced into the appropriate vectors and fully sequenced. Recombinant baculovirus-expressing Borealin was generated using the pFastBacHTA vector and the Bac-to-Bac Expression System (Invitrogen). Insect Sf-9 cells were infected with appropriate combinations of the constructs, and complexes containing GST-Aurora B, GST-INCENP, His-Survivin, and His-Borealin were purified on glutathione sepharose beads (Amersham Biosciences) as described previously (Honda et al., 2003). Other recombinant Borealin or GST alone prebound to glutathione sepharose beads. Samples were incubated on a rotating wheel for 1 h at 4°C. The supernatant was precipitated with TCA and the beads were washed three times with binding buffer, once with binding buffer without Triton X-100, and then boiled in sample buffer. After SDS-PAGE, proteins labeled with [35S]methionine were detected using a phosphorimager (Storm 860) with ImageQuant software (Amersham Biosciences).

**Immunoprecipitations**

HeLa cells were harvested after a 20-h colcemid block by mitotic shake-off and lysed in IP buffer (50 mM Tris-HCl, pH 8.0, 0.4 M NaCl, 0.5% NP-40, 0.1% deoxycholate, 30 µg/ml RNase A, 80 U/ml micrococcal nuclease [a gift of J. Allan, University of Edinburgh, Edinburgh, UK], 1 mM PMSF, and 1 µg/ml CLAP for 20 min on ice. The cleared extract was incubated with affinity-purified anti-Borealin antibody R1647 or preimmune serum R1647 coupled to protein A beads (Dynal) for 5 h at 4°C on a rotating wheel. The beads were washed twice with IP buffer without RNase and micrococcal nuclease and once with 10 mM Tris-HCl pH 7.4, 150 mM NaCl. Supernatant and beads were processed for SDS-PAGE and the proteins were transferred to nitrocellulose membranes for immunoblotting. Immunoprecipitation of Aurora B was carried out under the same conditions using anti-Aurora B antibody (rabbit; Abcam).

**Indirect immunofluorescence microscopy and immunoblotting**

Indirect immunofluorescence was performed as described previously (Carvalho et al., 2003) using antibodies against Aurora B (mouse; Translab); Survivin (rabbit; Novus); INCENP (rabbit 1186), α-tubulin (mouse; Sigma-Aldrich), β-tubulin (Faint and Merdes, 2002), γ-tubulin (mouse; Sigma-Aldrich), TD-60 (human; a gift of R.L. Margolis, CEA-CNRS, Grenoble, France), Borealin (affinity-purified rabbit 1647), Aurora A (a gift of C. Pri gent, Université de Rennes, Rennes, France), CENP-C (rabbit 558), and appropriate secondary antibodies coupled to FITC, Texas red, or Cy5 (goat; Jackson Immunoresearch Laboratories). DNA was visualized with 0.5 µg/ml DAPI. Image stacks were captured on a microscope (model IX-70, Olympus) controlled by Delta Vision SoftWorx (Applied Precision) using a 100 or 40X objective. After deconvolution, image stacks were quick-projected and saved as TIFF files.

Immunoblotting was performed as described previously (Carvalho et al., 2003) using antibodies against Aurora B (rabbit; Abcam), TD-60 (affinity-purified rabbit R2316), phospho-H3 (rabbit; Upstate Biotechnology), or others as described for indirect immunofluorescence.

**Antibody production**

Rabbit polyclonal antibody R1647 was generated against bacterially expressed human His-Borealin by Diagnostics Scotland and affinity-purified against GST-Borealin using standard methods. A polyclonal antibody (R2316) against a COOH-terminal peptide (KEIKKILPEYNPR) of human TD-60 was raised in rabbit by Genosphere. The serum was affinity purified against the peptide using the SulfoLink coupling gel according to the manufacturer’s instructions (Pierce Chemical Co.).

**RNA interference**

HeLa cells (ACC57 and DSA42) at 50% confluence were treated with 2 mM thymidine for 16 h and released into fresh medium without antibiotics for 8 h. Cells were transfected with control or Borealin siRNAs as described previously (Carvalho et al., 2003) 4 h into the release period. A second thymidine block was then applied for 16 h in the presence of the
siRNAs. Cells were released into fresh medium and analyzed upon entry into mitosis by immunoblotting or indirect immunofluorescence.

For movies, HeLa cells stably expressing histone H2B-GFP were treated as described in the preceding paragraph and released into fresh medium without phenol red (GIBCO BRL). After 8 h, the medium was supplemented with 10 mM Hepes, pH 7.5, and the coverslips were transferred to a fixed chamber (Bioptics) and kept at 37°C. DIC and FITC three-dimensional image data sets were collected every 7 min as described for indirect immunofluorescence.

Online supplemental material
Fig S1: Aurora B and Borealin colocalize during mitosis. Fig S2: examples of cell lates during the time-lapse experiment quantitated in Fig 6 C. Fig S3: Biornucleate cells observed after depletion of Borealin by RNAi. Fig S4: selected frames of representative movies of Borealin siRNA-transfected mitotic cells. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200404001/DC1.

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