

Down-modulation of nucleoporin RanBP2/Nup358 impaired chromosomal alignment and induced mitotic catastrophe

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Chromosomal missegregation is a common feature of many human tumors. Recent studies have indicated a link between nucleoporin RanBP2/Nup358 and chromosomal segregation during mitosis; however, the molecular details have yet to be fully established. Observed through live cell imaging and flow cytometry, here we show that RNA interference-mediated knockdown of RanBP2 induced G2/M phase arrest, metaphase catastrophe and mitotic cell death. Furthermore, RanBP2 down-modulation disrupted importin/karyopherin β 1 as well as the expression and localization of the Ran GTPase activating protein 1. We found that N-terminal of RanBP2 interacted with the N-terminal of importin β 1. Moreover, at least a portion of RanBP2 partially localizes at the centrosome during mitosis. Notably, we also found that GTPase Ran is also involved in the regulation of RanBP2–importin β 1 interaction. Overall, our results suggest that mitotic arrest and the following cell death were caused by depletion of RanBP2. Our findings point to a crucial role for RanBP2 in proper mitotic progression and faithful chromosomal segregation.

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The nuclear pore complex (NPC) is the major gatekeeper of macromolecular traffic between the nucleus and cytoplasm. It is composed of multiple subunits comprising ≈ 30 proteins called nucleoporins. During cell division, transmembrane nucleoporins and nuclear envelope membrane fragments move to the endoplasmic reticulum, while other nucleoporins are disassembled into subcomplexes that are distributed throughout the mitotic cytoplasm to kinetochore, spindle and centrosome regions.^{1–3} As mitosis nears completion, the NPC is sequentially reassembled from the dispersed cytosolic components.⁴ Our previous findings demonstrated that several NPC proteins, such as Rae1,^{5–8} Nup98,⁹ Tpr¹⁰ and Nup88,¹¹ do not simply disperse into the mitotic cytoplasm but instead become preferentially associated with kinetochores and/or spindle pole regions, where they are crucial in maintaining spindle bipolarity and thus prevent aneuploidy and carcinogenesis.^{4,12}

RanBP2, a Ran binding protein with four Ran binding domains, is a major cytosolic component of filaments that are derived from the cytoplasmic ring of the NPC^{13,14} during interphase. RanBP2 is composed of 3224 residues in humans. It can be divided into several domains: an N-terminal TPR domain, an α -helical region, four Ran-binding domains, eight tandem zinc fingers, a SUMO E3 ligase domain, and a C-terminal domain that displays sequence homology to cyclophilins.^{15,16} RanBP2 was initially described as a regulator of nucleocytoplasmic transport because of its link with the small G protein, Ran, and because it possesses

docking motifs for nuclear transport receptors.^{17–19} However, the multi-domain configuration of RanBP2 indicates a more pleiotropic function for this huge nucleoporin; indeed, RanBP2 is now recognized as a regulator of numerous cellular activities.^{20–27} Besides, RanBP2-RanGAP1*SUMO1/Ubc9 was identified as a multi-subunit SUMO E3 ligase.²⁸ Interestingly, RanBP2 has also been implicated in the delivery and integration of the genomic material of HIV-1.^{29–32} RanBP2 was also shown to potentiate the translation of a subset of mRNA encoding secretory proteins.³³

RanBP2 aids proper nuclear envelope assembly³⁴ and is involved in mitosis.^{35–37} Notably, together with Ran GTPase activating protein 1 (RanGAP1), RanBP2 relocates to kinetochores and mitotic spindle microtubules during mitosis.^{38,39} Whether normal mitosis also proceeds in the presence of altered RanBP2 expression is unclear.

To illustrate the participation of RanBP2 in cell-cycle regulation and to understand the underlying molecular mechanism thereof, we examined the effects of modulating RanBP2 expression on cell-cycle progression. Here, we show that knockdown of RanBP2 causes an aberrant metaphase and mitotic arrest in G2/M phase, leading to abnormal chromosome segregation, aneuploidy and metaphase catastrophe. We also found that at least a portion of RanBP2 localizes at the spindle pole/centrosome regions. Using RNAi-depletion assays, flow cytometry and live cell imaging studies, we were able to document a novel regulatory mechanism involving RanBP2 and importin β 1/karyopherin- β 1.

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Our biochemical characterization further confirms their direct interaction, while ectopic expression of the interaction domain revealed the role of the RanBP2-RanGAP1-importin β subcomplex in maintaining proper bipolarity segregation, thus preventing mitotic catastrophe. In addition to its activity as a GTPase, Ran might also regulate RanBP2-importin β interactions. These findings suggest a crucial role for RanBP2 in proper mitotic progression and chromosomal segregation, both of which prevent apoptosis.

Results

RanBP2 partially localizes at the centrosome and interacts with importin β to form a transient complex during mitosis. To understand the physiological role of

RanBP2 in cell-cycle regulation, we first examined the kinetics of endogenous RanBP2 expression during cell-cycle progression by using α -tubulin, a mitotic spindle marker. We found that RanBP2 transiently co-localized with α -tubulin during the metaphase-anaphase transition (Figure 1a). In contrast to the results of previous RanBP2 mitotic studies,^{38,39} in metaphase mitotic HeLa cells RanBP2 was visible along the spindle microtubules, and signals were also detected, to a lesser extent, at the kinetochores (Supplementary Figure S1) and even at the centrosome regions (Figure 1a). Confocal microscopy confirmed the RanBP2 centrosome staining pattern, which showed the partial association of γ -tubulin (spindle pole/centrosome marker) with RanBP2 from prophase to anaphase. In γ -tubulin immunoprecipitates from mitotic HeLa cells, low

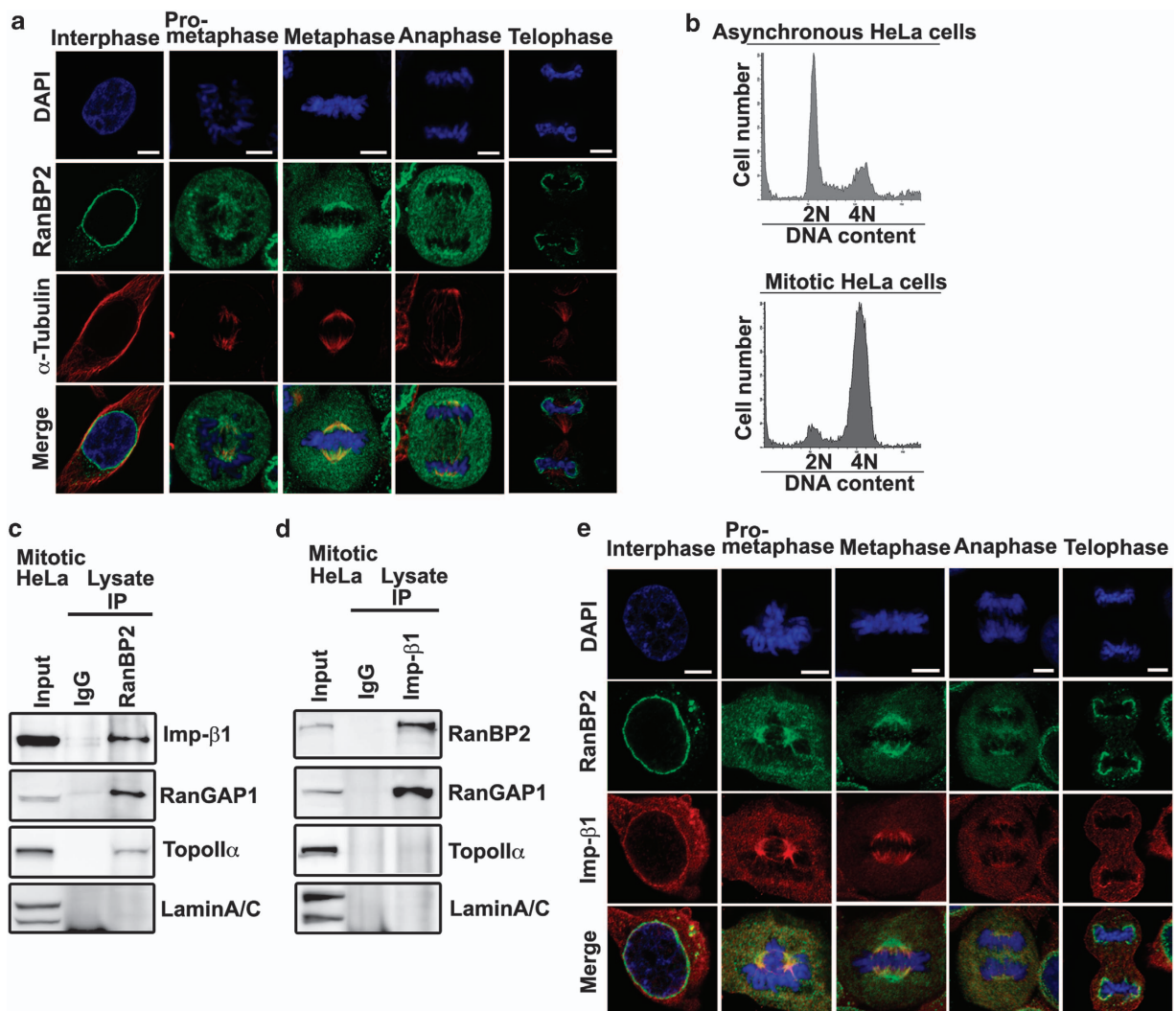


Figure 1 RanBP2 is associated with importin β during mitosis. **(a)** Confocal microscopic images of HeLa cells at different mitotic stages, stained with anti-RanBP2 (green) and anti- α -tubulin (red) antibodies. Chromatin was stained with 4',6-diamidino-2-phenylindole (DAPI; blue). Bars = 5 μ m. In lanes marked 'Input,' 20 μ l of the 500 μ l of extract used per IP were analyzed directly. **(b)** Asynchronous (upper panel) and synchronized mitotic (lower panel) HeLa cells (G2/M) were prepared via double thymidine block and confirmed by flow cytometry. **(c)** IP of mitotic HeLa cell extracts incubated with anti-RanBP2 or nonspecific rabbit antibodies (IgG) were analyzed by IB with anti-Imp- β 1, anti-RanGAP1, anti-Topoll α and anti-Lamin A/C antibodies. **(d)** IP of mitotic HeLa cell extracts incubated with anti-Imp- β 1 or nonspecific rabbit antibodies (IgG) were analyzed by IB with the indicated antibodies. **(e)** Confocal microscopic images of HeLa cells at different mitotic stages, stained with anti-RanBP2 (green) and anti-Imp- β 1 (red) antibodies. Chromatin was stained with DAPI (blue). Bars = 5 μ m

levels of RanBP2 and importin β 1 were detected (Supplementary Figure S1). Our RanBP2 centrosome localization data are also supported by recent proteomic studies involving liquid chromatography–MS of the mitotic *Drosophila* centrosome, which identified RanBP2 in the *Drosophila* centrosome.⁴⁰

Given that the regulatory functions of Ran are mediated by the dissociation of importin β 1 complexes⁴¹ and that importin β 1 also localizes at the mitotic spindle³⁹ and spindle pole region,⁴² we hypothesized that the contribution of RanBP2 (a Ran binding protein) is to spatiotemporally regulate the recruitment of importin β 1 during mitosis. We therefore examined whether RanBP2 was associated with the importin β 1 complex in mitotic HeLa cells. HeLa cells were synchronized and released from the double thymidine block treatment, the mitotic HeLa cells were harvested (~88%) and then confirmed by flow cytometry (Figure 1b). Immunoblotting (IB) of anti-RanBP2 immunoprecipitates showed the coprecipitation of importin β 1, TopoII α and RanGAP1 but not Lamin A/C (Figure 1c and Supplementary Figure S2). Using anti-importin β 1 antibodies, we immunoprecipitated RanBP2 and RanGAP1 but not TopoII α or Lamin A/C (Figure 1d). This interaction was also found in a lysate prepared from asynchronous HeLa cells, and it was not regulated by phosphorylation (Supplementary Figures S2a–c). Furthermore, RanBP2 and importin β 1 colocalized in mitotic HeLa cells (Figure 1e). Taken together, these observations suggest dynamic changes in the cellular localization of RanBP2 during mitosis and the potential role of this protein in the regulation of mitotic progression.

Knockdown of importin β 1 in vertebrate cells reduces kinetochore localization of RanBP2. To further analyze whether importin β 1 regulates the function of RanBP2 or vice versa, we used an importin β 1 siRNA knockdown approach (Figure 2a). IB analysis of HeLa cells subjected to importin β 1 siRNA treatment for 3 days revealed an 85% reduction in the level of importin β 1 compared with controls (Figure 2b and Supplementary Figure S3). The same IB membrane was re-probed with α -tubulin and β -actin to ensure equivalent loading (Figure 2b). Moreover, there were no significant changes in the levels of TopoII α (another reported binding partner of RanBP2) or the membrane nucleoporin Pom121 or LaminA/C (Figure 2c). Notably, RanBP2 levels were reduced significantly and RanGAP1 proteins were non-sumoylated, in contrast to mock-treated (control siRNA) cells (Figure 2c and Supplementary Figure S3). Moreover, we found that impaired chromosome alignments were increased fourfold compared with control siRNA mitotic cells ($n=300$ mitotic cells) (Figures 2d–e). We next assessed chromosomal defects and RanBP2 localization in importin- β 1-depleted HeLa cells. The depletion of importin β 1 reduced RanBP2 localization at the kinetochores, mitotic spindles and spindle poles (Figure 2e). RanGAP1 localization was also examined in importin- β 1-depleted cells. Consistent with our prediction, the knockdown of importin β 1 also reduced the RanGAP1 kinetochore and spindle localization pattern (Figures 2f–h). These effects were also observed in cells treated with another siRNA against importin β 1 (Supplementary Figures S3c–e). Taken together, these observations demonstrated

that the knockdown of importin β 1 reduces the mitotic localization of RanBP2 and RanGAP1 in vertebrate cells.

To determine whether soluble full-length importin β 1 could rescue the observed chromosomal defects, FLAG-importin β 1-RNAi-resistant plasmids (Supplementary Tables S1 and S2) were co-transfected together with importin β 1 siRNA, which partially rescued the chromosomal defects (Figure 2d). As predicted, there were significantly more defects in siRNA importin β 1 cells than in cells co-transfected with siRNA + FLAG-importin β 1-RNAi-resistant plasmid (14 *versus* 9%; $n=300$ mitotic cells) (Figure 2d).

To rule out the possibility that the importin- β 1-depleted phenotypes are indirect consequences of interphase, RanBP2 and Pom121 were followed with confocal microscopy, and their localization in control siRNA and in importin- β 1-depleted cells was determined (Supplementary Figure S4). Although 35% of the latter cells had an abnormal nuclear morphology (Supplementary Figure S4), triplicate experiments did not establish statistical significance ($P>0.1$; $n=150$) (Supplementary Figure S4). Why was the amount of RanBP2 protein controlled by importin β 1 depletion? Our results suggest that importin β 1 depletion leads to a reduction of RanBP2 and RanGAP1 sumoylation, such that importin β 1 and RanBP2 no longer localize on the kinetochores, mitotic spindles and centrosomes, thus leading to aneuploidy (Figures 2d–h).

RanBP2 down-modulation induces chromosome missegregation and aneuploidy. The above-described results prompted us to examine the consequences of RanBP2 depletion on importin β 1. In HeLa cells subjected to RanBP2 siRNA treatment for 3 days and analyzed by IB, RanBP2 levels were reduced by almost 90% compared with control cells (Figure 3a). Again, there were no significant changes in TopoII α expression. By contrast, the level of importin β 1 was reduced, and RanGAP1 sumoylation was significantly reduced (Figure 3b). With the nearly complete knockdown of RanBP2, less RanGAP1 and importin β 1 were localized in the kinetochores and mitotic spindles, and the frequency of chromosomal separation defects increased compared with control siRNA (Figures 3c and d). Consistent with previous findings, we found that mitotic RanBP2 siRNA treatment caused chromosomal misalignments in 25% of the mitotic cells, whereas these defects were not seen in mock-treated cells ($n=300$ mitotic cells) (Figure 3e).

RanBP2 down-modulation-induced metaphase catastrophe, G2/M phase arrest and mitotic cell death. To firmly establish that mitosis is disturbed (at different stages) in down-modulated RanBP2 cells, HeLa cells stably expressing histone H2B fused to green fluorescent protein (H2B-GFP) were examined using live cell imaging. This assay was under real-time, simultaneous detection of mitosis, ploidy and mitotic catastrophe.⁴³ Down-modulation of RanBP2 resulted in a prolonged metaphase (>40 min). We found that chromosomal segregation defects and impaired chromosome alignment were rarely observed in control cells (Figure 4a Supplementary Video S3). By contrast, in the live cell images, which showed chromosomal segregation defect phenotypes (Figure 4b and

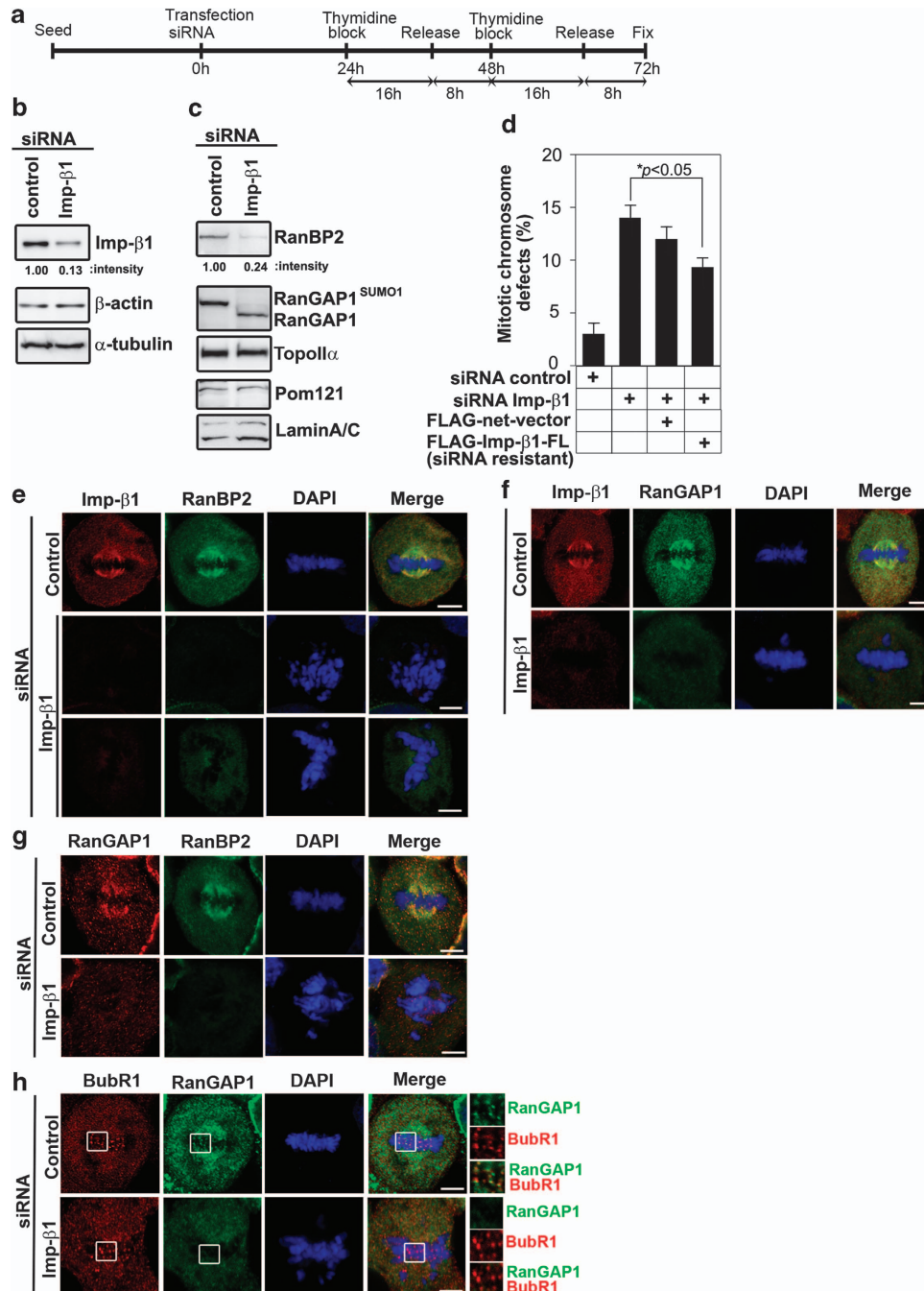


Figure 2 Importin β 1 siRNA treatment reduces RanBP2 and causes abnormal chromosomal congression and defective mitosis. (a) Schedule of fixing or collecting mitotic HeLa cells after siRNA Imp- β 1 depletion. (b) Lysates of mitotic HeLa cells transfected with control or Imp- β 1 siRNAs were analyzed 72 h later by IB with anti-Imp- β 1 antibody. The same membrane was stripped and reprobed with anti- β -actin and anti- α -tubulin (as loading control). (c) Effects of Imp- β 1 depletion on the protein levels of RanBP2-associated proteins. The same membranes as in (b) were analyzed by IB with the indicated antibodies. RanBP2 was significantly reduced in the Imp- β 1-depleted lysates, whereas Topoll α , Pom121 and Lamin A/C were not affected. (d) Quantification (relative percentage) of mitotic chromosomal defect phenotypes for the indicated siRNA and/or plasmid. Values are based on three independent experiments, counting 100 mitotic cells in each experiment. Mean values \pm S.D. (error bars) are shown. All cells were treated with double thymidine block, stained with 4',6-diamidino-2-phenylindole (DAPI) and visualized by confocal microscopy. (e) Confocal microscopic images of mitotic HeLa cells transfected with control or Imp- β 1 siRNA and analyzed 72 h post-transfection; the cells were stained with anti-RanBP2 (green) and anti-Imp- β 1 (red) antibodies and DAPI (blue). (f–h) Representative confocal microscopic images were stained with various antibodies as indicated. White boxes show the enlarged region. Bar = 5 μ m

Supplementary Video S1) together with impaired chromosome alignment (Figure 4c and Supplementary Video S2) in RanBP2 siRNA H2B-GFP cells. In response to RanBP2 depletion, cell division was halted during metaphase,

a phenomenon we refer to as metaphase catastrophe. Insufficient mitotic arrest was previously shown to result in abnormal cell-cycle progression via chromosomal abnormalities.^{44–46} We also investigated the rate of cell death under

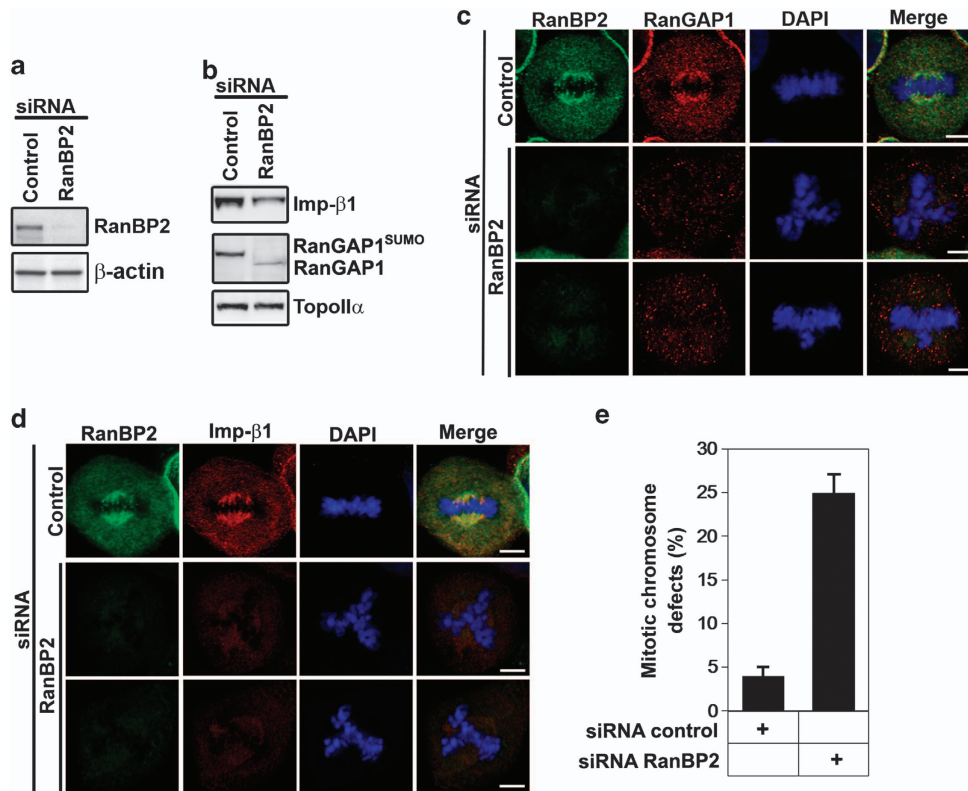


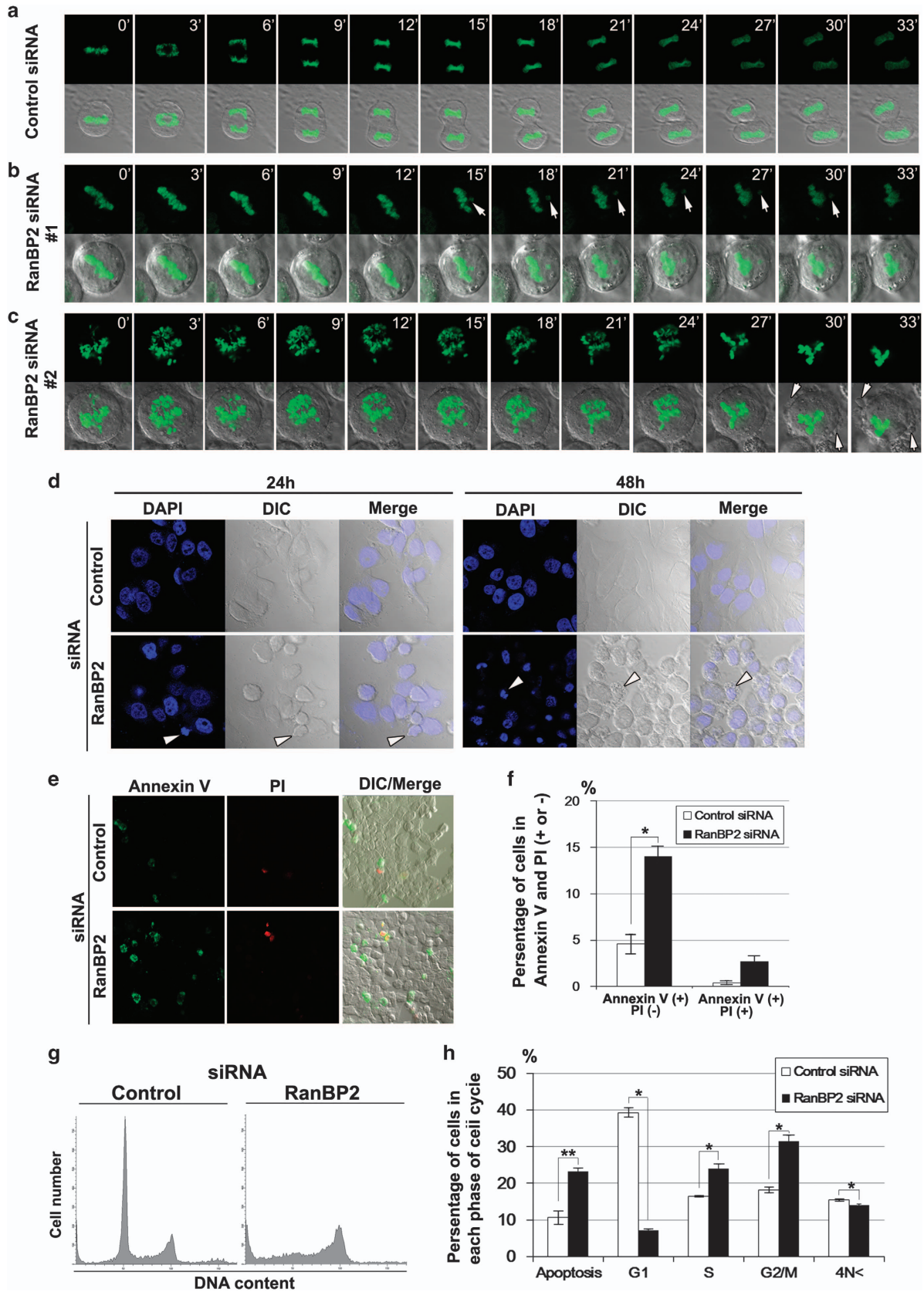
Figure 3 RanBP2 siRNA treatment reduces importin β 1 and causes abnormal chromosomal congression and defective mitosis. (a) Lysates of mitotic HeLa cells transfected with control or RanBP2 siRNAs were analyzed 72 h post transfection by IB with anti-RanBP2 antibody. The same membrane was stripped and reprobed with anti- β -actin (as a loading control). (b) Effects of RanBP2 depletion on the protein levels of RanBP2-associated proteins. The same membranes as in panel (a) were analyzed by IB with the indicated antibodies. (c and d) Confocal microscopic images of mitotic HeLa cells transfected with control or RanBP2 siRNAs. At 72 h post transfection, the cells were stained with RanBP2 (green) antibody and anti-RanGAP1 (red) or Imp- β 1 (red) antibodies and 4',6-diamidino-2-phenylindole (DAPI; blue). Bar = 5 μ m. (e) Quantification (relative percentage) of mitotic chromosomal defect phenotypes in cells transfected with control siRNA or RanBP2 siRNA. Values are based on three independent experiments, counting 100 mitotic cells in each experiment. Mean values \pm S.D. (error bars) are shown. All cells were treated with double thymidine block, stained with DAPI and visualized by confocal microscopy

these conditions. In cells transfected with siRNAs against RanBP2 but not control, the death cells gradually increased (Figure 4d). Next we plan to confirm whether this cell death is accompanied by apoptotic features. We monitored the percentage of Annexin V-positive cells. Forty-eight hours after transfection, we stained siRNA-transfected HeLa cells with FITC-labeled Annexin V and propidium iodide (PI).⁴⁷ We found that the percentage of Annexin V-positive and Annexin V/PI double-positive cells considerably increased following transfection with RanBP2 depletion (Figures 4e and f, $*P < 0.05$; $n = 300$ cells). To examine the effects of RanBP2 depletion on the cell-cycle profile, cells transfected with siRNA and incubated for 3 days thereafter were analyzed by flow cytometry (Figure 4g), which showed a twofold increase in mitotic cell death/apoptotic cells in the transfectants compared with the controls (Figures 4g and h, apoptotic cells were indicated as $**P < 0.01$). Moreover, depletion of RanBP2 resulted in an increase in the percentage of cells at the G2/M transition compared with control siRNA-transfected cells in triplicate experiments (32 *versus* 18%, respectively) (Figures 4g and h). These data indicate that cells were arrested in G2/M phase after RanBP2 depletion. Together, these data suggested that RanBP2 depletion induces metaphase catastrophe, impaired chromosome alignment,

G2/M phase arrested and mitotic cell death, indicating a crucial role for RanBP2 in faithful chromosomal segregation.

RanBP2-depletion-induced mitotic catastrophe is not a side effect of nuclear import.

Next, to rule out the possibility that the observed RanBP2-depletion phenotypes were secondary effects or byproducts of disrupted nuclear import during interphase, we generated a fluorescent import construct using a nuclear protein fragment, NuMA2/NuMA₃₂₅₋₈₂₉,⁸ fused to a nuclear localization signal and GFP, yielding GFP-NuMA2-NLS. This construct is imported into the nucleus whereas a mutant version of the same protein, GFP-NuMA2- Δ NLS, in which the NLS signal is mutated, is not. HeLa cells were transfected with either of the two constructs 24 h after treatment with RanBP2 siRNA. A significant reduction in RanBP2 localization at the nuclear membrane (nuclear rim) was observed 72 h after siRNA treatment, as was cytoplasmic staining of RanBP2 during interphase (Supplementary Figure S5). GFP-NuMA2-NLS protein accumulated in the nucleus of RanBP2-depleted cells, in contrast to GFP-NuMA2- Δ NLS, which remained cytoplasmic (Supplementary Figure S5). Consistent with these results, RanBP2 was previously shown to have little, if any, effect on NPC numbers, which during interphase



remained substantially intact after RanBP2 depletion.³⁵ Together, these data strongly suggest that RanBP2 depletion does not cause a major defect in nuclear import but is instead needed for chromosomal alignment and segregation.

Overexpression of RanBP2 does not rescue importin β 1 mitotic defects. To map the RanBP2 domain that physically binds to importin β 1, full-length GFP-RanBP2-FL (1–3224 aa), two deletion mutants, GFP-RanBP2-N1 (1–900 aa) and GFP-RanBP2-N2 (1–1826 aa), and the GFP net vector (Figure 5a) were transiently transfected into HeLa cells, followed by immunoprecipitation (IP) of the cell extracts with an anti-GFP antibody. As shown in Figure 5b, importin β 1 interacted with the RanBP2-FL and RanBP2-N2 fragments but not with RanBP2-N1 or the GFP net vector (Figure 5b) during mitosis, which is consistent with another recent report of this interaction during interphase.²⁷ To rule out that increased RanBP2 expression mimics the importin β 1 siRNA phenotypes through sequestration of endogenous importin β 1, different GFP-RanBP2 fragments were transiently overexpressed in HeLa cells. The results showed that chromosomal missegregation defects were not significantly altered by any of the GFP-RanBP2 constructs compared with GFP net vector in mitotic cells, even when transfected by various methods (Supplementary Figure S6). A careful examination of the relative intensity of RanBP2 in those GFP-RanBP2-transfected cells showed that total RanBP2 was only slightly increased through our various transient transfection methods (Supplementary Figure S6), which might explain why we did not observe enhancement of the phenotypes resulting from the chromosomal defects.

Ran is also required for the mitotic interaction between RanBP2 and importin β 1. A thorough understanding of the RanBP2–importin β 1 interaction must take into account the possible role of the GTPase Ran, as both RanBP2 and importin β 1 contain Ran-interacting regions. Ran might be part of the complex formed by RanBP2 and importin β 1, perhaps having a stabilizing role in their interaction. To test this hypothesis, GST-RanBP2-M1_{900–1826} and His-Imp- β 1-N2_{1–650} were expressed and the two proteins were then mixed *in vitro* with or without Ran-GTP (Figures 5c and d). The addition of His-Ran-GTP proteins significantly enhanced the binding between GST-RanBP2-M1_{900–1826} and His-Imp- β 1-N2_{1–650} (Figure 5d). Next, we generated constructs containing mutagenized Ran-binding domains of RanBP2²⁶ and importin β 1⁴⁸ (RanBP2-mt and Imp- β 1-mt, respectively) to determine whether the lack of Ran binding in one partner affected the mitotic localization of the other partner (Figure 5d). The absence of Ran-binding domains in RanBP2-mt and Imp- β 1-mt significantly reduced the

interaction between RanBP2 and importin β 1 compared with the wild type (Figures 5e–g). Taken together, these data suggest that Ran regulates the binding between RanBP2 and importin β 1.

The N-terminal of importin β 1 interacts with RanBP2 by sequestering endogenous RanBP2 and thereby induces mitotic catastrophe defects. Finally, we biochemically mapped the *in vivo* interaction domain between RanBP2 and importin β 1. Five cDNA fragments covering the entire length of importin β 1 (including the full length and four fragments of the protein), named Imp- β 1-FL (1–876 aa), Imp- β 1-N1 (1–250 aa), Imp- β 1-N2 (1–650 aa), Imp- β 1-M (250–650 aa) and Imp- β 1-C (650–876 aa), were generated. Each cDNA was coupled to a 6 × His tag and then expressed in an *Escherichia coli* system (Figure 6a). Pull-down assays with mitotic HeLa cell lysates showed that RanBP2 interacted with Imp- β 1-N2 (1–650 aa) but not with the other fragments nor with the vector alone. Consistent with this result, Imp- β 1-N2 (1–650 aa) also localized to the nuclear envelope, but other fragments did not (Supplementary Figure S7). Additionally, we found that Imp- β 1-N1 (1–250 aa), which contained HEAT repeats 1–6, and Imp- β 1-M (250–650aa), which contained HEAT repeats 7–14, were also required for binding; Imp- β 1-N1, Imp- β 1-M and Imp- β 1-C failed to bind to RanBP2 (Figure 6b). These mapping results suggested that overexpression of FLAG-Imp- β 1-N2 (1–650 aa) mimics the RanBP2 siRNA phenotype through the sequestration of endogenous RanBP2, making it unavailable to interact productively with full-length importin β 1, thus causing the re-appearance of the enhanced chromosome segregation abnormalities. This scenario is analogous to the mitotic catastrophe phenotype observed after the reduction of RanBP2 levels by RNAi. Indeed, we observed chromosomal segregation defects and mitotic catastrophe in 30% of FLAG-Imp- β 1-N2 (1–650 aa) mitotic cells but not in cells transfected with the other fragments (Figure 6c and Supplementary Figure S8). Taken together, our findings indicate that the N-terminal of importin β 1_{1–650} interacts with RanBP2 by sequestering endogenous RanBP2 and thereby induces mitotic catastrophe defects.

Discussion

During mitosis, chromosomal segregation is carried out by a complex cellular machinery⁴⁵ in which abnormalities in the cell cycle and mitosis eventually lead to cell death. There is increasing evidence that nucleoporins control chromosomal instability and tumor development.^{1,4,49} In the present work, we demonstrated that down-modulation of RanBP2 induces

Figure 4 RanBP2 down-modulation induces G2/M phase arrest, metaphase catastrophe and cell death. (a–c) Time-lapse microscopy of HeLa cells stably expressing histone H2B-GFP. At 72 h post transfection with (a) control or (b and c) RanBP2 siRNAs, the cells were processed for time-lapse microscopic analysis. Images were acquired every 3 min. Arrow indicates a fragmented chromosome. White arrowhead indicates an apoptotic bleb. (d) siRNAs against RanBP2 and control were transfected, respectively. Cell images were acquired using a phase-contrast microscope 24 or 48 h after siRNA transfection. White arrowhead indicates an apoptotic cell. (e) Cell death induced by knockdown of RanBP2. HeLa cells 48 h after siRNA transfection were stained with FITC-labeled Annexin V and PI. (f) Annexin V and PI-positive HeLa cells 48 h after siRNA transfection. HeLa cells randomly selected ($n=300$) were counted. Values are an average of three independent experiments. Error bars represent S.D. (g) At 48 h post transfection with control or RanBP2 siRNAs, HeLa cells were examined for their cell-cycle profiles by flow cytometry. (h) Percentage of cell-cycle phases in cells from panel (g). Asterisks indicates the significant P values (* $P<0.05$; ** $P<0.01$)

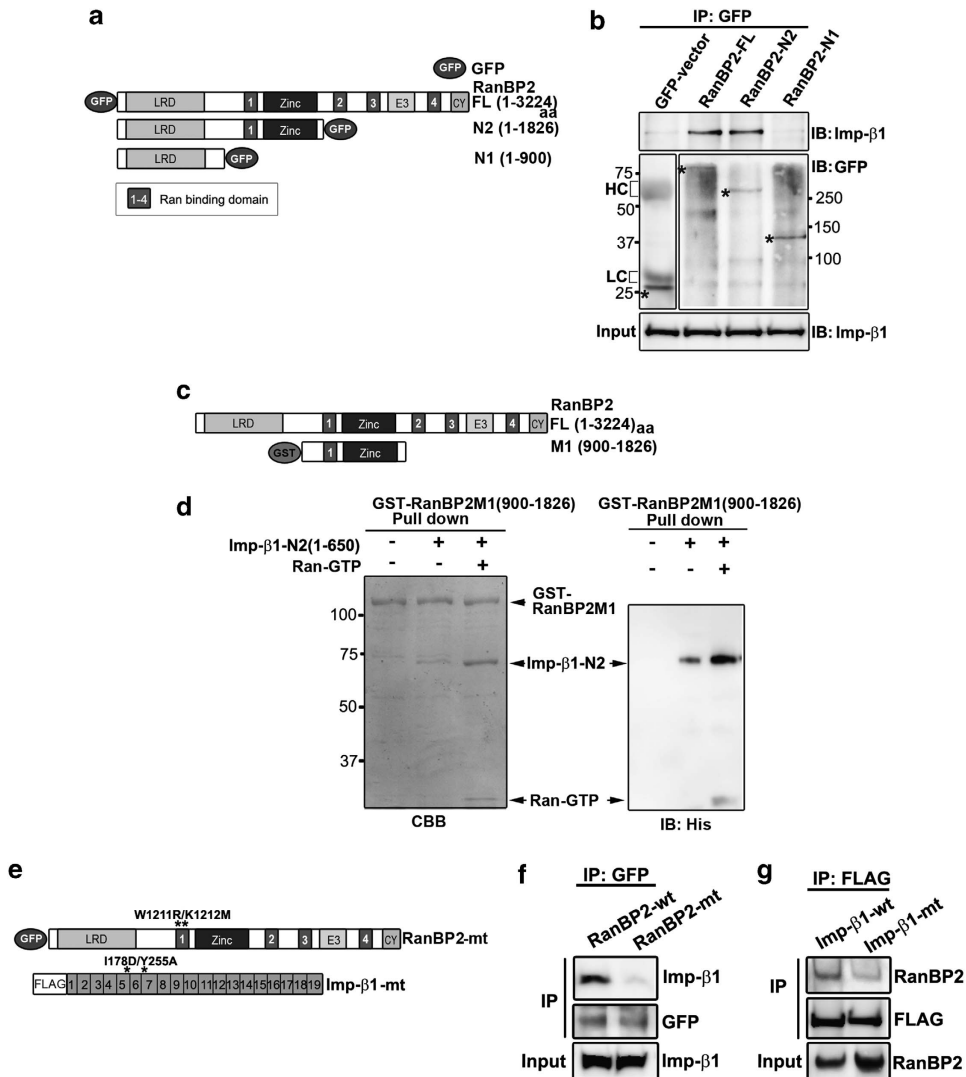


Figure 5 Ran binding is important for the interaction between the N-terminal of RanBP2 and importin β 1. (a) Scheme of full-length RanBP2 and two C-terminal deletion mutants with GFP tags. Numbers on the right refer to amino acids. LRD, leucine-rich domain; Zinc, zinc finger motifs; E3, SUMO E3 ligase domain; CY, cyclophilin homology domain. (b) HeLa cells were transfected with a GFP, GFP-RanBP2-FL, RanBP2-N2 or RanBP2-N1 expression plasmid. After double thymidine block, lysates prepared from the cells were immunoprecipitated with anti-GFP antibody and then analyzed by IB with anti-Imp- β 1 or anti-GFP antibodies. Asterisks indicate the GFP fusion proteins. Numbers indicate molecular mass markers in kilodaltons. HC, IgG heavy chain; LC, IgG light chain. In lanes marked 'Input,' 20 μ l of the 500 μ l of extract used per IP were analyzed directly. (c) Scheme of full-length RanBP2 and the N- and C-terminal deletion mutant (M1) with the GST tag. Numbers on the right refer to amino acids. (d) *In vitro* expressed and purified GST-RanBP2-M1, Imp- β 1-N2-His and His-Ran-GTP were mixed and pulled-down with glutathione-Sepharose beads. Pulled-down lysates were separated by SDS-PAGE and then either stained with Coomassie brilliant blue (CBB) or subjected to IB with anti-6 \times His antibody. Numbers indicate molecular mass markers in kilodaltons. (e) Scheme of GFP-tagged full-length RanBP2 carrying the W1211R/K1212M mutation and FLAG-tagged Imp- β 1-FL carrying the I178D/Y255A mutation. (f and g) HeLa cells were transfected with a GFP-RanBP2-FL (wt), GFP-RanBP2-FL_W1211R/K1212M (mt), FLAG-Imp- β 1-FL (wt) or FLAG-Imp- β 1-FL_I178D/Y255A (mt) expression plasmid. After double thymidine block, lysates prepared from the cells were immunoprecipitated with anti-GFP or anti-FLAG antibody and then analyzed by IB with anti-Imp- β 1, anti-GFP, anti-RanBP2 or anti-FLAG antibodies. In lanes marked 'Input,' 20 μ l of the 500 μ l of extract used per IP were analyzed directly

abnormal chromosomal segregation, improper mitotic progression and mitotic cell death, evidence of the crucial role of RanBP2 in proper mitotic progression and faithful chromosomal segregation. Mitotic defects following RanBP2 siRNA treatment were previously described only in fixed cells.^{35,36,38,39} Here, through live cell imaging, chromosomal segregation defects and metaphase catastrophe could, for the first time, be recorded in live cells undergoing mitosis (especially during metaphase-anaphase transitions) following RanBP2 siRNA treatment (Figure 4 and Supplementary Videos S1 and S2).

Mitotic catastrophe is defined in morphological terms as a mechanism of cell death occurring during or after aberrant mitosis.^{45,50} In this study, we also identified a novel link between the RanBP2-importin β 1 complex and the Ran GTPase that is required for proper chromosomal segregation. Consistent with this finding, Roscioli *et al.*⁵¹ reported the interaction of importin β 1 with RanGAP1 (the bona fide binding partner of RanBP2) in mitotic cells. In addition, previous findings suggested that importin β 1, RanBP2 and SUMO-RanGAP1 co-localize along microtubules.^{38,39} Our immunofluorescence results provide the first evidence

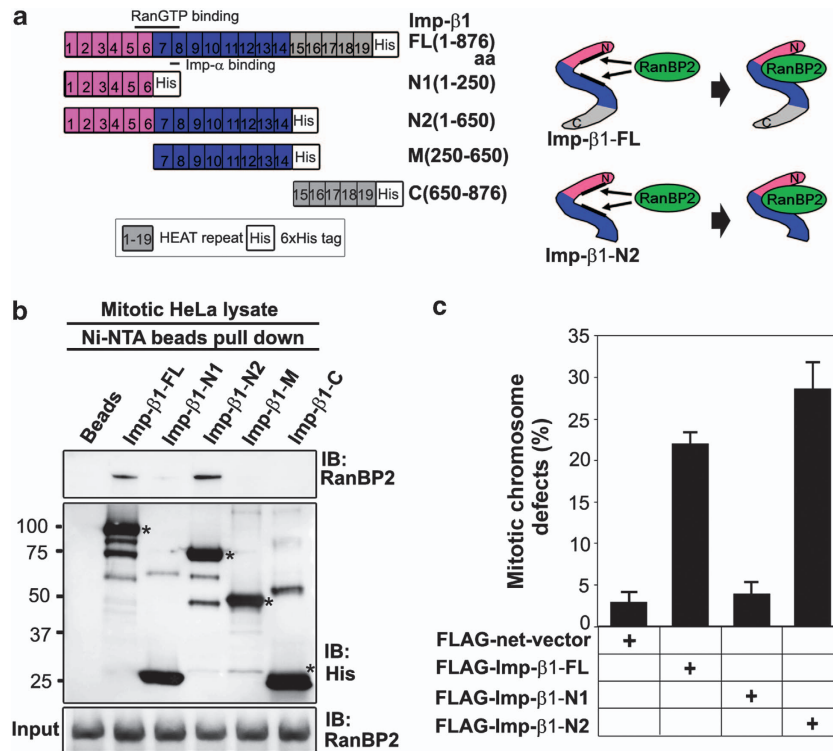


Figure 6 N-terminal of importin β 1 interacts with RanBP2. (a) Scheme of full-length Imp- β 1 and four mutants with 6 \times His tags. Numbers on the right refer to amino acids (left). An illustration of an interaction model between importin β 1 and RanBP2 (right). (b) *In vitro* expressed and purified 6 \times His tagged Imp- β 1 fragments were mixed with mitotic HeLa lysates and pulled-down with Ni-NTA beads. Pulled-down lysates were analyzed by IB with anti-RanBP2 or anti-6 \times His antibodies. Asterisks indicate the 6 \times His fusion proteins. Numbers indicate molecular mass markers in kilodaltons. In lanes marked 'Input,' 20 μ l of the 500 μ l of lysate used per pull-down were analyzed directly. (c) Quantification (relative percentage) of mitotic chromosome defect phenotypes in FLAG-net-vector, FLAG-Imp- β 1-FL, FLAG-Imp- β 1-N1 and FLAG-Imp- β 1-N2 transfected cells. Values are based on three independent experiments, counting 100 mitotic cells in each experiment. Mean values \pm S.D. (error bars) are shown. All cells were treated with double thymidine block, stained with DAPI (4',6-diamidino-2-phenylindole) and visualized by confocal microscopy

that at least a small portion of mitotic RanBP2 interacts with γ -tubulin at the centrosome region (Figure 1 and Supplementary Figure S2). They are also consistent with recent proteomic studies of the mitotic drosophila centrosome, in which RanBP2 was localized.⁴⁰ It is unclear exactly how RanBP2 is recruited to the centrosome/spindle pole, but the mechanism seems to involve a cascade of events. Our results indicated that RanBP2 is not the only factor involved in the regulation of chromosomal segregation and that it has other roles in mitotic progression in addition to mediating metaphase catastrophe (Figure 7). Thus, several issues, including the exact mechanism by which RanBP2 contributes to mitotic cell death and the roles of importin β 1 and Ran in mitotic catastrophe, are subjects for future research.

In mammals, aneuploidy has been linked to cancer progression,⁵² which, like cancer development, is a complex process involving functional and genetic abnormalities. Our recent findings of the interaction of nucleoporin Nup98 with Galectin-3⁵³ and the association of nucleoporin Tpr with p53⁵⁴ further strengthen the relationship between nucleoporins and carcinogenesis.⁴⁹

Collectively, this study sheds light on a novel physiological function of RanBP2 in the regulation of mitotic progression and chromosomal segregation, which may facilitate the discovery of new treatment

regimens for cancer by targeting mitosis. It may also provide mechanistic insights into previously reported phenotypes associated with altered RanBP2 expression, including tumor progression and the embryonic lethality of RanBP2-null mice.³⁶ Future clinical investigations should also explore how RanBP2 contributes to carcinogenesis.

Materials and Methods

Plasmids. HeLa cell cDNA was synthesized using a SuperScript III CellsDirect cDNA synthesis system kit (Life Technologies Corporation, Carlsbad, CA, USA). The full-length RanBP2/Nup358 coding region was PCR-amplified from cDNA and subcloned into pEGFP-C3 (Clontech Laboratories, Mountain View, CA, USA) by introducing a *NotI* site using a site-directed mutagenesis kit (TaKaRa, Bio Inc., Shiga, Japan). Two RanBP2 fragments (N1 and N2) were subcloned by PCR from pEGFP-C3-RanBP2 into pEGFP-N2. The RanBP2-M1 fragment was subcloned by PCR from pEGFP-C3-RanBP2 into pET28a-GST. Both a plasmid encoding the full-length human importin β 1 and RanQ69L, tagged with 6 \times His, in the pQE60 vector, were kind gifts from Dr. Ulrike Kutay (ETH Zurich, Switzerland). Four importin β 1 fragments (N1, N2, M and C) were subcloned by PCR from pQE60-importin β 1 into pQE60 or into pCMV-3 \times -FLAG. pEGFP-C2-NuMA2/NuMA₃₂₅₋₈₂₉ and pEGFP-C2-NuMA₃₂₅₋₈₂₉ Δ NLS plasmids encoding the NLS oligonucleotide were inserted into pEGFP-C2 (Clontech Laboratories). GFP-RanBP2-FL_W1211R/K1212M, FLAG-importin β 1_1178D/Y255A, wild-type Ran-6 \times His and importin β 1 (full-length) siRNA-resistant plasmids were synthesized from each full-length construct using a site-directed mutagenesis kit. The GFP-H2B plasmid was obtained from Addgene (Cambridge, MA, USA). Details of the expression constructs and cloning primers are listed in Supplementary Tables S1 and S2. All constructs were confirmed by DNA sequencing.

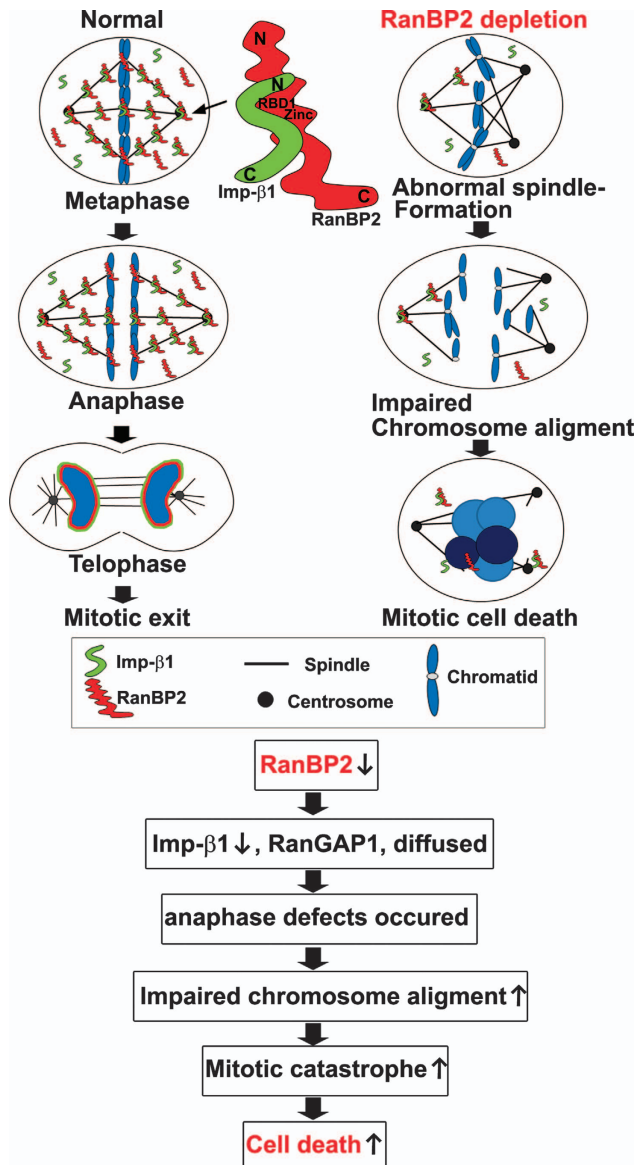


Figure 7 Model for RanBP2 in proper mitotic progression and faithful chromosomal segregation. Absence of RanBP2 during mitosis caused abnormal mitotic spindles, supernumerary centrosomes, impaired chromosomal alignment, induced mitotic catastrophe and cell death

Mammalian cell culture, transfections and cell synchronization.

HeLa cells were obtained from the American Type Culture Collection (ATCC) and were synchronized in S phase by double thymidine block using 2 mM thymidine.^{13–16} The cells were transfected with GFP-RanBP2-fragment plasmids using Lipofectamine 2000 (Life Technologies Corporation), TurboFect (Thermo Fishes Scientific Inc., Waltham, MA, USA) and Neon (Life Technologies Corporation) following the manufacturers' protocols. The HeLa EGFP-H2B cell line, used in the analysis of mitotic progression, was generated by transfecting HeLa cells with EGFP-H2B cDNA and maintained in medium containing G418 (600 µg/ml) as previously described.¹⁰

RNA interference. siRNA duplexes targeting RanBP2/Nup358 (sequence 5'-CCGUUUUGGAGUCAACAtt-3', siRNA ID s11773, cat no. 4390824) were purchased from Life Technologies Corporation. Importin β1 (sc-35736) and control siRNAs (sc-37007) were purchased from Santa Cruz Biotechnology (Dallas, TX, USA). Another independent siRNA duplexes targeting Importin β1 #2 (sequence 5'-GAAAAUUUGGGAAUGAGAACdTdT-3')⁵⁵ were purchased from Sigma-

Aldrich (St. Louis, MO, USA) for phenotypes double check. siRNA transfections were performed using Lipofectamine 2000 (Life Technologies Corporation) as previously described.¹⁰

Antibodies, immunocytochemistry (ICC) and confocal microscopy.

Anti-Nup358/RanBP2 rabbit polyclonal antibody (for ICC) was a kind gift from Dr. Elias Coutavas and Dr. Blobel (The Rockefeller University). Anti-Nup358/RanBP2 (sc-74518) (IB, IP), anti-importin/Karyopherin β1 (sc-137016) (IB, IP) and anti-β-actin (sc-47778) mouse monoclonal antibodies and anti-RanGAP1 (sc-25630) (ICC, IP) rabbit polyclonal antibody were from Santa Cruz Biotechnology. Anti-importin/Karyopherin β1 mouse monoclonal antibody (ab2811) (ICC) and anti-Pom121 rabbit polyclonal antibody (ab111610) were from Abcam (Cambridge, England). Anti-SUMO1 and anti-RanGAP1 (33-0800) (ICC, IB) mouse monoclonal antibodies were from Life Technologies Corporation. Anti-Topollα mouse monoclonal antibody (MO42-3S) was from MBL (Nagoya, Japan). Anti-Lamin A/C rabbit polyclonal antibody (2032) was from Cell Signaling (Danvers, MA, USA). Anti-GFP (A6455) (IP, IB, ICC) rabbit polyclonal antibody was from Life Technologies Corporation. Anti-GFP (012-20461) (ICC) mouse monoclonal antibody was from Wako (Osaka, Japan). Anti-α-tubulin (DM1A), anti-γ-tubulin (GTU-88) and anti-FLAG (M2) monoclonal antibodies were from Sigma-Aldrich. Anti-BubR1 (612502) mouse monoclonal antibody was from BD Transduction Laboratories (Franklin Lakes, NJ, USA). Anti-6 × His mouse monoclonal antibody was from Qiagen (KJ Venlo, Netherlands). Secondary antibodies were from Molecular Probes (Life Technologies Corporation). For immunofluorescence, synchronized HeLa cells were washed in phosphate-buffered saline (PBS) and fixed for 10 min in 4% paraformaldehyde in PBS. Cells were then permeabilized with 0.3% Triton X-100 in PBS for 10 min at room temperature. Annexin V FLUOS staining kit (Roche, Rotkreuz, Switzerland) was used for phosphatidylserine detection. Samples were mounted onto coverslips using Pro-Long Gold Antifade reagent (Life Technologies Corporation) and were examined on a Zeiss Carl Zeiss Microscopy GmbH (Jena, Germany) LSM5 EXCITER confocal microscope from Carl Zeiss Microscopy GmbH (Jena, Germany) with four laser beams, and all images were acquired using a plan-Apochromat from Carl Zeiss Microscopy GmbH at × 63 with a 1.4-N.A. objective or at × 100 with a 1.4-N.A. objective.

Immunoprecipitation.

The IP procedures were described previously.^{10,53} Briefly, mitotic HeLa cells were collected, washed with PBS, spun at 400 × g for 10 min and lysed in 1 ml of cold lysis buffer (50 mM Tris-HCl (pH 7.2), 250 mM NaCl, 0.1% Nonidet P-40, 2 mM EDTA, 10% glycerol) containing 1 × protease inhibitor mixture (Roche) and 1 mM phenylmethylsulfonyl fluoride. The lysates were centrifuged for 30 min at 4 °C at 14 000 × g. The resulting supernatants were pre-cleared with 50 µl of protein A/G bead slurry (Santa Cruz Biotechnology), mixed with 10 µl of various antibodies as specified and incubated for 1 h at 4 °C with rocking. The beads were then washed five times with 500 µl of lysis buffer. After the last wash, 50 µl of 1 × sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) blue loading buffer (New England Biolabs, Ipswich, MA, USA) was added to the bead pellet before loading. Signals were detected with an enhanced chemiluminescence system (GE Healthcare, Buckinghamshire, England) and quantified using an LAS-4000 image analyzer (Fuji Film, Tokyo, Japan) according to the manufacturer's specifications.

Expression of recombinant proteins.

To express 6 × His-tagged importin β1 fragments and Ran in *E. coli* XL-1 Blue cells (Agelent Technologies, Inc., Santa Clara, CA, USA) and GST-tagged RanBP2-M1 in *E. coli* BL21(DE3) Codon Plus (Agelent Technologies, Inc.) cells, both strains of bacteria were grown at 37 °C to an absorbance at 600 nm (A_{600}) of 0.6 and induced with 0.5 mM isopropyl-β-D-thiogalactopyranoside at 18 °C overnight. The cells were harvested by centrifugation and lysed in buffer containing 50 mM Tris-HCl (pH 7.7), 150 mM KCl, 0.1% Triton-X100 and Complete EDTA-free protease inhibitor mixture tablets (Roche Applied Science). The cells were lysed using a cell sonicator (SMT), and the lysate was clarified by centrifugation at 15 000 × g for 60 min. Proteins containing the 6 × His tag were purified by nickel-affinity chromatography (Qiagen). The GST-tagged protein was purified by glutathione-Sepharose and Superdex 200 10/300 GL size exclusion chromatography (GE Healthcare).

In vitro binding assays. *In vitro* binding assays were performed as previously described.¹⁹ Briefly, 10 µl of glutathione-Sepharose beads were

suspended in Tween 20 binding buffer (20 mM HEPES, pH 7.4, 110 mM potassium acetate, 2 mM magnesium chloride and 0.1% Tween 20) containing 1 × protease inhibitor mixture (Roche), 0.25 mM GST-RanBP2-M1, 0.25 mM Kap-β1-His and 0.8 mM Ran-GTP in a total reaction volume of 40 μl. The reactions were allowed to proceed for 2 h at 4 °C with rotation, after which the beads were washed twice in cold TB-0.1% Tween 20. After the last wash, 20 μl of 1 × SDS-PAGE blue loading buffer was added to the bead pellet before loading.

Time-lapse microscopy. Time-lapse analysis of cellular histone dynamics during metaphase and anaphase transitions in live cells was recorded in GFP-H2B stable HeLa cell lines as described previously.¹⁸ The cells were placed in a microincubation chamber (7136; Corning, Corning, NY, USA) on a Zeiss LSM5 confocal microscope stage, which was heated to 37 °C and equipped with a CO₂ supply (electric CO₂ microscope stage incubator; Okolab, Ottaviano, Italy). Time-lapse series were generated by collecting photographs every 3 min; the photographs were then converted to eight-bit images and processed using Adobe Photoshop CS5 from Adobe Systems Incorporated (San Jose, CA, USA) and NIH Image J from National Institutes of Health (Bethesda, MD, USA).

Flow cytometry. HeLa cells transfected with siRNA were trypsinized, washed twice with PBS and fixed in 70% ethanol at −20 °C overnight. The fixed cells were resuspended in PBS containing 50 μg RNase A (Nacalai Tesque, Kyoto, Japan)/ml and 50 μg PI (Sigma-Aldrich)/ml. Cellular DNA content was analyzed using a FACSCanto II (BD Biosciences, Franklin Lakes, NJ, USA) with FACS Diva software (BD Biosciences).⁵⁶

Conflict of Interest

The authors declare no conflict of interest.

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