

# Chemical genetics reveals the requirement for Polo-like kinase 1 activity in positioning RhoA and triggering cytokinesis in human cells

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**Polo-like kinases (Plks) play crucial roles in mitosis and cell division. Whereas lower eukaryotes typically contain a single Plk, mammalian cells express several closely related but functionally distinct Plks. We describe here a chemical genetic system in which a single Plk family member, Plk1, can be inactivated with high selectivity and temporal resolution by using an allele-specific, small-molecule inhibitor, as well as the application of this system to dissect Plk1's role in cytokinesis. To do this, we disrupted both copies of the *PLK1* locus in human cells through homologous recombination and then reconstituted Plk1 activity by using either the wild-type kinase (Plk1<sup>WT</sup>) or a mutant version whose catalytic pocket has been enlarged to accommodate bulky purine analogs (Plk1<sup>AS</sup>). When cultured in the presence of these analogs, Plk1<sup>AS</sup> cells accumulate in prometaphase with defects that parallel those found in *PLK1*<sup>Δ/Δ</sup> cells. In addition, acute treatment of Plk1<sup>AS</sup> cells during anaphase prevents recruitment of both Plk1 itself and the Rho guanine nucleotide exchange factor (RhoGEF) Ect2 to the central spindle, abolishes RhoA GTPase localization to the equatorial cortex, and suppresses cleavage furrow formation and cell division. Our studies define and illuminate a late mitotic function of Plk1 that, although difficult or impossible to detect in Plk1-depleted cells, is readily revealed with chemical genetics.**

cell division | Ect2 | knockout | mitosis

Members of the Polo-like kinase (Plk) family are critical for the proper timing and fidelity of chromosome segregation and cell division (1, 2). Humans and other mammals possess four distinct but closely related Plks. The most extensively studied of these isoforms, Plk1, has important functions throughout late G<sub>2</sub> and early M phase (1, 2). Plk1 is also thought to be necessary for cytokinesis, but its exact contributions have been difficult to investigate, because the early mitotic defects caused by Plk1 depletion trigger the spindle assembly checkpoint (SAC), precluding direct observation of cell division. One solution would be to use small molecules that can rapidly inhibit Plk1 in anaphase, after the SAC has been satisfied. Some compounds that inhibit Plk1 activity *in vitro* have been reported (3–7), but their specificities *in vivo* are either poor or largely unknown. Although screening panels of kinases can provide one estimate of inhibitor selectivity, such *in vitro* assays are unavailable for most of the ≈500 kinases present in the human genome and would miss off-target effects on other relevant classes of enzymes, such as cytoskeletal motor proteins (8, 9). In addition, because of the strong conservation of the Plk kinase domain, none of these compounds is expected to be selective for individual Plk isoforms, complicating their use as precise surrogates of Plk1 depletion or genetic ablation.

In yeast, a powerful method for monospecific kinase inhibition involves replacing the target enzyme of interest with a variant whose catalytic pocket has been modified to accept bulky purine analogs (10). As these analogs cannot fit into the nucleotide-binding sites of endogenous kinases, they do not affect the

growth of isogenic yeast strains lacking the mutant kinase, providing strong evidence of their specificity *in vivo*. We show here that a similar approach can be used in human cells to dissect kinase function with high temporal resolution and genetic specificity. Using gene targeting and transgenic complementation, we established somatic cell lines in which Plk1 activity was provided solely by an analog-sensitive mutant (Plk1<sup>AS</sup>). Although Plk1<sup>AS</sup> cells grew comparably to Plk1 wild-type (Plk1<sup>WT</sup>) cells, they were uniquely sensitive to two different purine analogs, arresting in mitosis with characteristic defects in spindle assembly, centrosome maturation, and chromosome alignment. In addition, acute inhibition of Plk1<sup>AS</sup> during anaphase prevented cleavage furrow formation and cell division. This defect was accompanied by a failure to localize the RhoA GTPase, an essential regulator of actomyosin dynamics during cytokinesis, and its upstream guanine nucleotide exchange factor (GEF) Ect2 to the equatorial cortex and central spindle, respectively. Furthermore, Plk1's own recruitment onto the central spindle was also abolished. Our studies identify distinct functions of Plk1 throughout mitosis and cytokinesis and demonstrate the power of chemical genetics in dissecting these complex but short-lived events within human cells.

## Results

To develop a chemical genetic system for Plk1, we first needed to eliminate the endogenous kinase from human cells. We used adeno-associated virus vectors (11, 12) (Fig. 1*a*) to mutate both copies of the *PLK1* locus in telomerase-expressing human retinal pigment epithelial cells. Because *PLK1* is expected to be essential, one allele was deleted outright, whereas the other could be conditionally deleted by using Cre recombinase. After two rounds of targeting, we obtained several independent *PLK1*<sup>fllox/Δ</sup> clones (Fig. 1*b*). These cells were infected with adenoviruses expressing Cre recombinase (AdCre) or adenoviruses expressing β-galactosidase (Adβgal) as a negative control. Over the next 24–48 h, the AdCre-infected cells ceased to express Plk1 (Fig. 1*c*) and accumulated in mitosis (Fig. 1*d*). As expected, many of these cells had severe spindle and centrosome maturation defects (see Fig. 3 and Table 1). Subsequently, the proportion of mitotic

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The authors declare no conflict of interest.

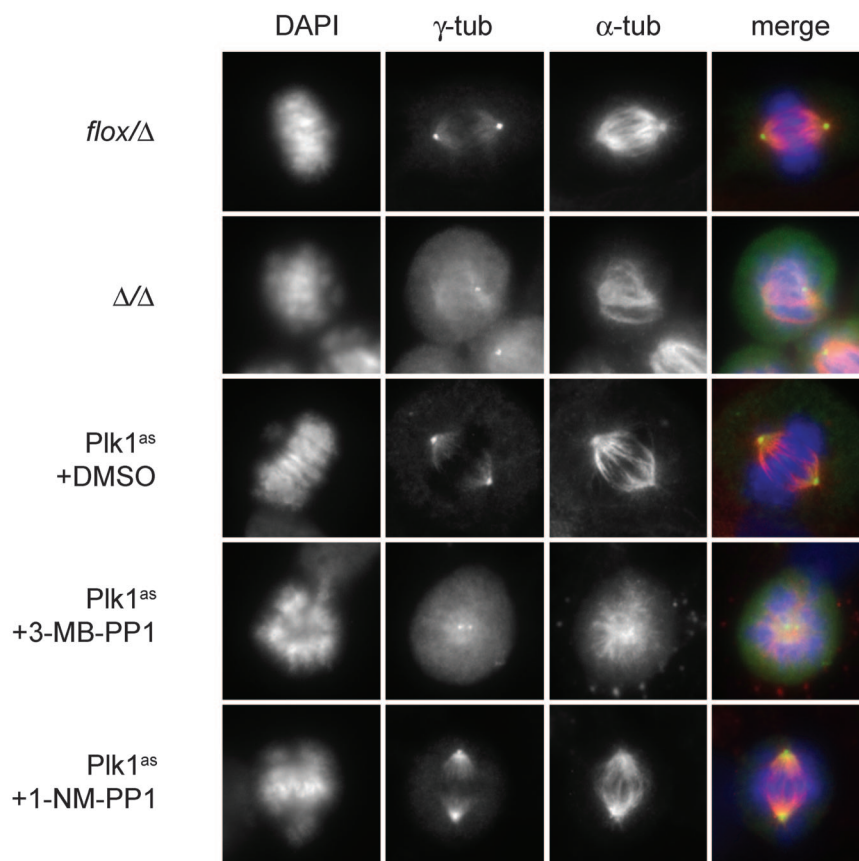
Abbreviations: Adβgal, adenoviruses expressing β-galactosidase; AdCre, adenoviruses expressing Cre recombinase; GEF, guanine nucleotide exchange factor; Plk, Polo-like kinases; Plk1<sup>WT</sup>, Plk1 wild type; Plk1<sup>AS</sup>, analog-sensitive Plk1 mutant; SAC, spindle assembly checkpoint.

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**Fig. 3.** Comparison of early mitotic defects in *PLK1*-null and *Plk1<sup>as</sup>* cells. *PLK1<sup>flox/Δ</sup>* cells were infected with Ad $\beta$ gal or AdCre for 48 h. *Plk1<sup>as</sup>* cells were incubated for 10 h with 10  $\mu$ M 3-MB-PP1, 10  $\mu$ M 1-NM-PP1, or DMSO. Cells were stained with antibodies to  $\gamma$ -tubulin ( $\gamma$ -tub) (green) and  $\alpha$ -tubulin ( $\alpha$ -tub) (red) and analyzed by immunofluorescence microscopy.

(SI Fig. 6]. Upon AdCre infection and limiting dilution, we recovered EGFP-positive clones in which both *PLK1* alleles had been deleted (i.e., a *PLK1 $\Delta/\Delta$*  genotype; SI Fig. 6), demonstrating functional rescue by both transgenes.

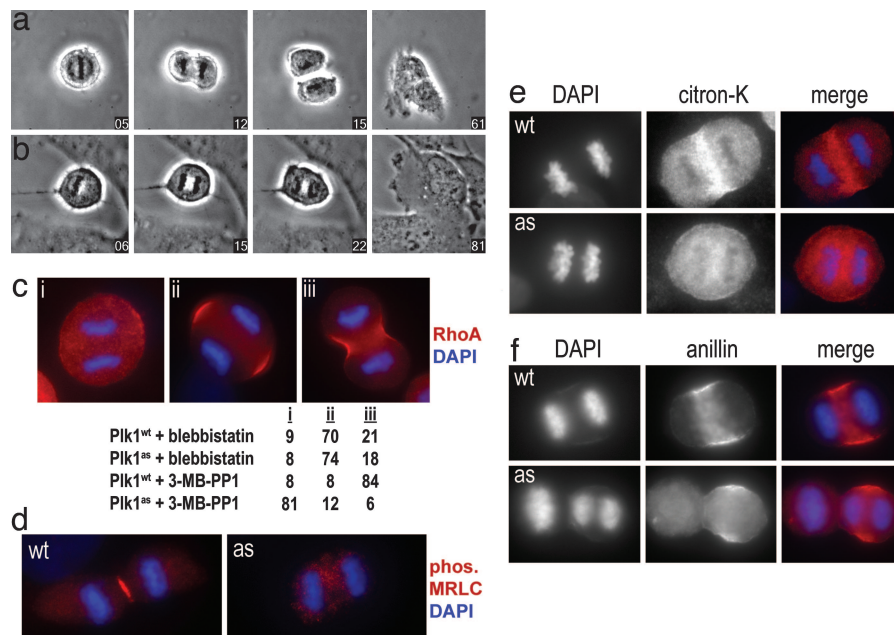
*In vitro*, *Plk1<sup>as</sup>* was able to use *N*6-(benzyl)-ATP and sensitive to the purine analogs 1-NM-PP1 (10) and 3-MB-PP1 (Fig. 2*a*), in contrast to *Plk1<sup>wt</sup>*, which was  $\approx$ 12-fold more active than *Plk1<sup>as</sup>* in the presence of ATP (Fig. 2*b–d*). In comparison, the widely used *cdc28-as1* allele behaves comparably to *CDC28* in genetic assays but encodes a kinase with a 50-fold lower  $k_{cat}/K_m$  for ATP than the wild-type enzyme (10). We therefore focused our attention on whether *Plk1<sup>as</sup>* conferred analog sensitivity *in vivo*. Both 1-NM-PP1 and 3-MB-PP1 inhibited the growth of *Plk1<sup>as</sup>* cells in a dose-dependent manner but had little effect on *Plk1<sup>wt</sup>* cells (Fig. 2*e*). Like *PLK1 $\Delta/\Delta$*  cells, inhibitor-treated *Plk1<sup>as</sup>* cells first accumulated in mitosis but then succumbed to mitotic slippage (Fig. 2*f*). In titrations, 3-MB-PP1 induced mitotic arrest

at a 10-fold lower concentration than 1-NM-PP1 (Fig. 2*g*), paralleling its greater potency as a *Plk1<sup>as</sup>* inhibitor *in vitro*. Importantly, the kinetics of this arrest were much faster than with AdCre-mediated gene deletion, as expected for an inhibitory mechanism that does not rely on mRNA or protein turnover. Moreover, *Plk1<sup>as</sup>* cells resumed cell division once the inhibitor was removed (SI Fig. 7). From these data, we conclude that these compounds rapidly, selectively, and reversibly inhibit *Plk1<sup>as</sup>* *in vivo*.

To evaluate the penetrance of *Plk1* inactivation, we examined spindle assembly and centrosome maturation in analog-treated *Plk1<sup>as</sup>* cells by using *PLK1 $\Delta/\Delta$*  cells as a qualitative and quantitative reference for the null phenotype (Fig. 3 and Table 1). Confirming earlier studies (16–18), mitotic *PLK1 $\Delta/\Delta$*  cells had monopolar or disorganized bipolar spindles with adjacent and immature centrosomes, as reflected in poor concentration of  $\gamma$ -tubulin at the poles. In agreement with our titrations, there was

**Table 1. Quantification of centrosomal and spindle defects in *PLK1*-null and *Plk1<sup>as</sup>* cells**

Cell line	Virus or compound	Separated centrosomes (%)	Unseparated centrosomes (%)	Normal bipolar spindles (%)	Disorganized bipolar spindles (%)	Monopolar spindles (%)	Multipolar spindles (%)
<i>flox/Δ</i>	Ad $\beta$ gal	49 (98)	1 (2)	49 (98)	0 (0)	1 (2)	0 (0)
<i>flox/Δ</i>	AdCre	15 (30)	35 (70)	2 (4)	19 (38)	29 (58)	0 (0)
<i>Plk1<sup>as</sup></i>	DMSO	50 (100)	0 (0)	49 (98)	0 (0)	0 (0)	1 (2)
<i>Plk1<sup>as</sup></i>	1-NM-PP1	45 (90)	5 (10)	24 (48)	21 (42)	5 (10)	0 (0)
<i>Plk1<sup>as</sup></i>	3-MB-PP1	25 (50)	25 (50)	10 (20)	15 (30)	25 (50)	0 (0)
<i>Plk1<sup>wt</sup></i>	3-MB-PP1	25 (100)	0 (0)	25 (100)	0 (0)	0 (0)	0 (0)
<i>flox/Δ</i>	3-MB-PP1	25 (100)	0 (0)	24 (96)	0 (0)	0 (0)	1 (4)

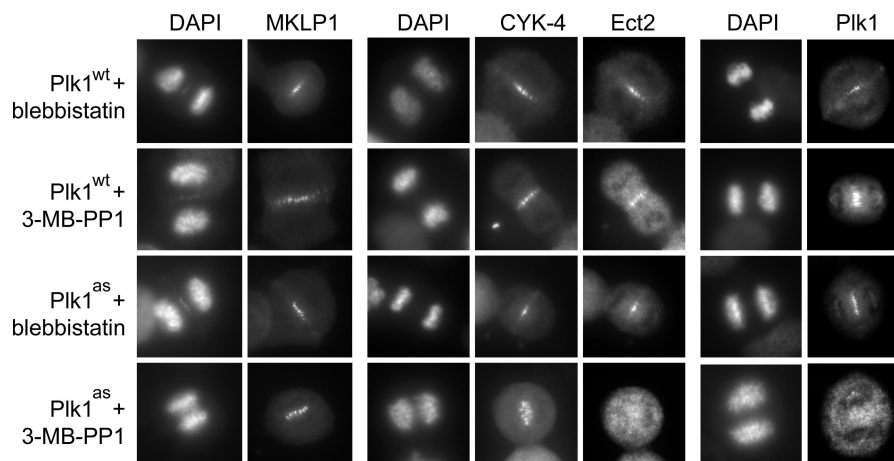


**Fig. 4.** Acute inhibition of Plk1 during anaphase disrupts RhoA localization and blocks the onset of cytokinesis. (a and b) Metaphase Plk1<sup>wt</sup> (a) and Plk1<sup>as</sup> (b) cells were treated with 10  $\mu$ M 3-MB-PP1 (time 0) and followed during progression into anaphase. Although chromatid separation occurred normally, Plk1<sup>as</sup> cells failed to divide and became binucleated (SI Movies 1 and 2). (c) Plk1 activity is required to localize RhoA at the equatorial cortex. Plk1<sup>wt</sup> and Plk1<sup>as</sup> cells were synchronized with monastrol, released for 30 min, and then treated for 20 min with either blebbistatin or 3-MB-PP1. RhoA accumulation at the equatorial cortex was visualized after trichloroacetic acid fixation (21, 27). Anaphase cells ( $n = 100$  per sample) were classified with respect to equatorial RhoA staining and cleavage furrow formation. (d–f) Plk1<sup>wt</sup> and Plk1<sup>as</sup> cells were treated with 3-MB-PP1 and stained with antibodies against phosphorylated myosin regulatory light chain (MRLC) (d), citron kinase (citron-K) (e), and anillin (f).

a clear difference between 1-NM-PP1 and 3-MB-PP1 when applied to Plk1<sup>as</sup> cells at the same concentration (10  $\mu$ M): Most 1-NM-PP1-treated cells had well separated centrosomes and bipolar spindles but incompletely aligned chromosomes (Fig. 3). By contrast, half of the 3-MB-PP1-treated cells had adjacent centrosomes and monopolar spindles, a fraction comparable to that observed in *PLK1* $\Delta\Delta$  cells (Fig. 3 and Table 1). Crucially, none of these defects was observed when Plk1<sup>wt</sup> or *PLK1*<sup>fox1 $\Delta$</sup>  cells were treated with 3-MB-PP1, ruling out the possibility that they arose from an off-target activity against other endogenous cellular enzymes, including other Plk isoforms or microtubule motor proteins required for spindle bipolarity (1, 8). We conclude that, although both analogs inhibit Plk1<sup>as</sup> sufficiently to

block cell proliferation and induce mitotic arrest, 3-MB-PP1 elicits a more penetrant phenotype that is virtually identical to *PLK1* deletion. We therefore used 3-MB-PP1 for all subsequent analyses.

To determine whether our chemical genetic system could be used to probe Plk1's role in cytokinesis, we located metaphase cells and treated them with 3-MB-PP1 shortly before they entered anaphase. Whereas Plk1<sup>wt</sup> cells divided normally ( $n = 10$  cells; Fig. 4a and SI Movie 1), furrows were either absent or highly labile in Plk1<sup>as</sup> cells, resulting in a single binucleated cell upon mitotic exit ( $n = 8$  cells; Fig. 4b and SI Movie 2). Thus, Plk1's kinase activity is needed for an early event in cytokinesis. Given the well documented role of the small GTPase RhoA at



**Fig. 5.** Plk1 activity recruits both the RhoGEF Ect2 and Plk1 itself to the central spindle. Plk1<sup>wt</sup> and Plk1<sup>as</sup> cells were treated as in Fig. 4c. Whereas MKLP1 and CYK-4 localized correctly under all conditions, Ect2 and Plk1 were both absent from the central spindle after Plk1 inhibition.

the cleavage furrow (19), as well as recent data that the yeast Plk Cdc5 controls the local activation of the RhoA ortholog Rho1 at the site of cell division (20), we asked whether Plk1 regulates RhoA localization in dividing human cells. Plk1<sup>wt</sup> and Plk1<sup>as</sup> cells were synchronized in prometaphase with monastrol (8), were released into monastrol-free medium for 30 min to allow spindle bipolarization and silencing of the SAC, and then were treated for 20 min with 3-MB-PP1 (to inhibit Plk1<sup>as</sup>) or blebbistatin (to inhibit myosin II; see ref. 9) during anaphase (Fig. 4c). We confirmed that blebbistatin inhibits furrowing without affecting the cortical localization of RhoA (21–23), consistent with myosin II being a downstream target of RhoA (19). In contrast, 3-MB-PP1 suppressed both equatorial RhoA recruitment and furrowing in Plk1<sup>as</sup> cells, implying that Plk1 acts upstream of RhoA (Fig. 4c). Consistent with the lack of RhoA at the equatorial cortex, 3-MB-PP1-treated Plk1<sup>as</sup> cells also lacked the phosphorylated form of myosin regulatory light chain, which normally accumulates at the central spindle and contractile ring in response to RhoA signaling ( $n = 46$  of 50 cells affected; Fig. 4d) (21–23). The RhoA-dependent localization of citron kinase (24, 25) was similarly lost upon Plk1 inhibition ( $n = 47$  of 50 cells; Fig. 4e). A fourth component of the contractile ring, anillin (26), was partially affected by Plk1 inhibition. In Plk1<sup>wt</sup> cells, anillin was concentrated in an equatorial band and excluded from the poles. In contrast, although some anillin reached the equatorial cortex in inhibitor-treated Plk1<sup>as</sup> cells, a large fraction was dispersed throughout the cytoplasm ( $n = 30$  cells; Fig. 4f). These results indicate that Plk1 activity is required for the equatorial localization of RhoA and its effectors during anaphase.

In animal cells, RhoA localization and furrow assembly are regulated by the RhoGEF Ect2, which is recruited to the central spindle by the Rho GTPase activating protein (GAP) CYK-4/MgcRacGAP and the kinesin MKLP1 (21–23, 27). Furthermore, Plk1 itself relocates onto the central spindle, making it an ideal candidate to regulate Ect2-mediated signaling. Intriguingly, whereas both MKLP1 and CYK-4 were recruited to the central spindle under all conditions (Fig. 5), Ect2 was dispersed throughout the cytoplasm in 3-MB-PP1-treated Plk1<sup>as</sup> cells ( $n = 48$  of 50 cells delocalized). In addition, Plk1<sup>as</sup> itself was absent from the central spindle ( $n = 44$  of 50 cells), which is consistent with a previous report that Plk1's targeting to this site requires phosphorylation of MKLP2 (28). We conclude that Plk1 activity controls not only the kinase's own recruitment at the central spindle but also that of Ect2, the critical GEF for RhoA during cytokinesis (19). Our results identify a clear requirement for Plk1 in activating the RhoA network at the start of cytokinesis, a function that, although in close temporal proximity to Plk1's early mitotic roles, could nonetheless be isolated by using chemical genetics.

## Discussion

To date, defining the precise roles of Plk1 in mammalian cytokinesis has been difficult, because inactivating Plk1 by standard techniques results in spindle defects and SAC-dependent prometaphase arrest (17, 18). Efforts to circumvent this arrest (for example, through simultaneous inactivation of the SAC and Plk1) suggested that Plk1 may be dispensable for the initiation of cytokinesis in humans (18, 29). However, we found that acute inhibition of Plk1<sup>as</sup> in cells that entered anaphase spontaneously (i.e., after satisfying rather than bypassing the SAC) suppressed cleavage furrow formation. Furthermore, Plk1 activity is necessary to target both the kinase itself and the RhoGEF Ect2 to the central spindle, a structure that has an important instructive role in contractile ring assembly (19). When Plk1 is inhibited, RhoA and its effectors fail to localize at the presumptive furrow, readily explaining why cytokinesis fails to occur. Given the similarity of these effects to RNAi depletion of Ect2 or its binding partner CYK-4 (21–23, 27), we conclude that Plk1 is a critical upstream activator of the Ect2–RhoA

network during cytokinesis. We speculate that, once localized to the central spindle, Plk1 stimulates Ect2's recruitment and/or GEF activity through phosphorylation, because Ect2 is a Plk1 substrate *in vitro*, and its interaction with CYK-4 is sensitive to phosphatase treatment (21, 30).

Current models place Ect2 and RhoA at the apex of a signal transduction pathway that stimulates the onset of cytokinesis, with additional factors, such as anillin, participating at later steps (19). Indeed, anillin is not required for either furrow formation or Ect2's central spindle recruitment (26, 31, 32), arguing that the mild delocalization of anillin in analog-treated Plk1<sup>as</sup> cells was not responsible for their early cytokinesis defects. Nevertheless, we do not exclude the possibility that regulation of anillin by Plk1 may be necessary for later stages of cytokinesis.

Our experience highlights three strengths of the chemical genetics approach. First, the bulky size of the purine analogs reduces the likelihood of off-target inhibition *a priori*, as the compounds are sterically occluded from the active sites of most if not all wild-type kinases. Second, the *in vivo* selectivity of these compounds was explicitly verified in each experiment, by ensuring that all phenotypes were linked to the analog-sensitive kinase. In principle such verification is possible with any bioactive small molecule (for example, by introducing an inhibitor-resistant version of the putative target into cells or organisms and assaying for functional rescue) but, to our knowledge, has not been conducted with existing Plk inhibitors. Third, because analog sensitivity is a genetically encoded trait, the same compounds can be used to inhibit more than one kinase in a manner equivalent to temperature shift in yeast genetics. For example, a similar strategy was recently used to replace Cdk7, the cyclin-dependent kinase-activating kinase (CAK), with an analog-sensitive version in human cells (33). Given the large number of kinases involved in mitosis and cytokinesis, as well as the short time scales on which these enzymes act, further chemical genetic analysis of the human cell division cycle should be highly informative.

## Materials and Methods

**Molecular Genetics.** Generation of adeno-associated virus particles and screening for gene-targeted clones was performed as described in refs. 11 and 12. Adenoviruses were used at a multiplicity of infection of 100. Plk1<sup>wt</sup> and Plk1<sup>as</sup> retroviruses were produced by cotransfection of retroviral and vesicular stomatitis virus glycoprotein envelope plasmids into Phoenix cells.

**Immunofluorescence and Live-Cell Microscopy.** For detection of citron kinase and RhoA, cells were fixed with 4% paraformaldehyde or 10% trichloroacetic acid and stained in accordance with published protocols (21–27). For all other antigens, cells were fixed with 100% methanol at  $-20^{\circ}\text{C}$  and stained as described in ref. 21. Live-cell imaging was conducted on a temperature- and  $\text{CO}_2$ -controlled microscope platform (11). For further details, see [SI Movies 1 and 2](#).

**Kinase Assays.** Plk1 was immunoprecipitated from whole-cell extracts, washed in lysis buffer (11) three times, once in kinase buffer [20 mM Tris (pH 7.4)/10 mM  $\text{MgCl}_2$ /50 mM KCl/1 mM DTT], and then incubated in kinase buffer plus 5  $\mu\text{g}$  of casein and [ $\gamma$ - $^{32}\text{P}$ ]ATP or [ $\gamma$ - $^{32}\text{P}$ ]N6-(benzyl)-ATP for 15–20 min at  $30^{\circ}\text{C}$  (9).  $^{32}\text{P}$  incorporation was quantified by SDS/PAGE and PhosphorImager (Fuji-Film Medical Systems USA, Stamford, CT) analysis.

**Chemicals.** 1-NM-PP1 has been described in ref. 10. 3-MB-PP1 was synthesized by using a modified procedure (11). Monastrol (8) and blebbistatin (9) were used at 100  $\mu\text{M}$ .

See [SI Text](#) for detailed protocols.

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