Different Phosphorylation States of the Anaphase Promoting Complex in Response to Anti-mitotic Drugs: A Quantitative Proteomic Analysis

Judith A. J. Steen;1 Hanno Steen;2 Ann Georgi;3 Kenneth Parker;4 Michael Springer;3 Marc Kirchner;5 Fred Hamprecht5 and Marc W. Kirschner3*

1Harvard Medical School and Children’s Hospital Boston, Department of Neurology, Boston, MA 02115, USA

2Harvard Medical School and Children’s Hospital Boston, Department of Pathology, Boston, MA 02115, USA

3Harvard Medical School, Dept. of Systems Biology, Boston MA 02115, USA

4Brigham and Women’s Hospital, Harvard Medical School, Department of Neurology, Boston, MA 02115, USA

5Interdisciplinary Center for Scientific Computing, University of Heidelberg, Heidelberg, Germany

*Corresponding author: Marc W. Kirschner, Department of Systems Biology, Harvard Medical School, 200 Longwood Ave, Boston, MA 02115, USA; e-mail marc@hms.harvard.edu
ABSTRACT

The anaphase promoting complex (APC) controls the degradation of proteins during exit from mitosis and entry into S-phase. The activity of the APC is regulated by phosphorylation during mitosis. As the phosphorylation pattern provides insights into the complexity of regulation of the APC, we studied in detail the phosphorylation patterns at a single mitotic state of arrest generated by various anti-mitotic drugs. We examined the phosphorylation patterns of the APC in HeLa S3 cells after they were arrested in prometaphase with taxol, nocodazole, vincristine or monastrol. There were 71 phosphorylation sites on 9 of the APC subunits. Despite the common state of arrest, the various anti-mitotic drug treatments resulted in differences in the phosphorylation patterns and phosphorylation stoichiometries. The relative phosphorylation stoichiometries were determined using a method adapted from the isotope free Quantitation of the Extent of Modification iQEM. We could show that during drug arrest the phosphorylation state of the APC changes, indicating that the mitotic arrest is not a static condition. We discuss these findings in terms of the variable efficacy of anti-mitotic drugs in cancer chemotherapy.
Introduction

The Anaphase Promoting Complex (APC) is an essential ubiquitin ligase that coordinates events in the mitotic and G1 phases of the cell cycle (1). The activity of the APC is regulated by two cofactors, CDC20 and CDH1 (2, 3) as well as by protein modifications including phosphorylation (4, 5). The APC is the target of the spindle checkpoint (6), a complex process that senses the fidelity of microtubule attachment to chromosomes before allowing chromatid separation and cell division to proceed. Improper microtubule attachment or chromosome misalignment on the spindle activates the formation of an inhibitory “supercomplex,” consisting of the APC and a second complex, the mitotic checkpoint complex (MCC), which consists of a kinase BubR1, an inhibitor MAD2, and the APC activator CDC20 (1).

Phosphorylation is thought to be important in regulating the APC and several kinases are known to affect the spindle checkpoint machinery, including CDK1, PLK1 (7-9), BUB1, MPS1, BUBR1 and Aurora B (10). Furthermore, i) the activation state of the APC is directly affected by phosphorylation (11-13); ii) phosphorylation of the APC affects the binding of interacting partners such as CDC20 (2, 14-16); and iii) the phosphorylation of CDC20 by BUB1 is also important for full inhibition of the APC by MAD2 (14).

The checkpoint machinery or related activities may be the indirect target of several anti-cancer therapies, including vinca alkaloids and taxanes, which are inhibitors of microtubule dynamics and are among the most effective anti-cancer agents (17-21). Although much is known about drug interactions with microtubules, why only some microtubule inhibitors are useful anti-cancer drugs is a puzzle: e.g. neither colchicine nor nocodazole finds use in tumor treatment. Therefore, a detailed understanding of the effects of anti-spindle drugs on the checkpoint and APC may provide clues to the development of even more effective therapies (22). In particular, since phosphorylation appears to be an important regulator of the checkpoint, a detailed monitoring of the phosphorylation state of the APC might prove informative (23).

Mass spectrometry has been used to map phosphorylation sites on the APC (4) from cells arrested with nocodazole. A total of 43 phosphorylation sites were found. Although this
study highlighted the complexity and potential importance of the phosphorylation control of the APC, it provided little functional information: the sheer number of sites suggested that functional studies would be very difficult. Even a comprehensive set of point mutations at each phosphorylation site might not be enough to gain full functional understanding. Sites might act redundantly or dependently, and it would be prohibitively difficult to study all possible combinations. Furthermore, some sites might be quantitatively unimportant, while other sites might serve as unstable intermediates in important pathways. Although a complete analysis by mutation may be impossible, a quantitative and kinetic analysis might reveal important clues to APC’s regulatory circuitry and provide new information about the dynamics of the arrested state and the action of anti-mitotic drugs.

Quantitative measurements of phosphorylation of a complex target are far from routine. Antibody-based approaches, using phosphorylation site specific antibodies/Western blotting and phosphopeptide specific enrichment strategies in combination with stable isotope labeling and mass spectrometric approaches cannot readily distinguish between changes in the degree of phosphorylation or changes in the level of a protein, which is a serious complication in the cell cycle where protein synthesis and degradation rates change rapidly. However, information about the stoichiometry of phosphorylation is critical. A 5-fold increase in signal could correspond to changes from 1 % to 5 % or to changes from 20 % to 100 % in phosphorylation. The former might be significant for an activating phosphorylation, but is probably insignificant for an inhibiting phosphorylation. Furthermore, low stoichiometries may be important if turnover is rapid, but that could only be evaluated in kinetic studies and such studies generally require an absolute measurement of stoichiometry. We recently developed a mass spectrometry technique, iQEM (isotope free Quantitation of the Extent of Modification) (24), that addresses some of these problems. It allows analysis of phosphorylation stoichiometries without isotopic labeling and can be used to study the dynamics of modifications irrespective of the source of the protein material, even from whole tissue/biopsies.
In the work reported here, we comprehensively analyzed phosphorylation sites on the APC in response to a prometaphase arrest using various anti-mitotic drugs, several of them used in anti-cancer therapies. Subsequently, we applied concepts from iQEM to quantitatively determine the differences in phosphorylation stoichiometries of some of the sites upon treatment with the various anti-mitotic drugs. In addition we used iQEM to study the kinetics/temporal order of phosphorylation of the APC. Apart from identifying characteristic differences in phosphorylation profiles for the different spindle poisons we determined kinetics for several phosphorylation sites. This demonstrates that mitotic arrest is not a unitary state; it is both dynamic and diverse.

Results

**APC and associated proteins are heavily phosphorylated in the cell cycle**

Several strategies were tested to optimize identification of the APC phosphopeptides, a scheme of the strategy used is shown in Supplementary Information (SI) Figure 1. Following immunoprecipitation the APC was analyzed in several conditions: (M) double thymidine arrest followed by an 8 hour release into mitosis, (G1) double thymidine arrest released into G1; (n) nocodazole mitotic arrest; (mn) monastrol mitotic arrest; (S) double thymidine arrest; (tl) 9 nM taxol mitotic arrest; (tm) 20 nM taxol mitotic arrest; (th) 40 nM taxol mitotic arrest; (vc) vincristine mitotic arrest. The drug treatments were selected to provide cells with similar distributions of cell cycle states. Not surprisingly, members of the spindle checkpoint complex were purified with the APC in cells treated with spindle poisons. Core APC and MCC proteins that were identified reproducibly in several experiments are listed in the SI Table 1 (incl. obtained sequence coverage). Kinases, such as PLK1, Aurora kinase B and CDK1 also co-immunoprecipitated with the APC complex (data not shown). An ubiquitin conjugating E2 protein, E2S (UBE2S_HUMAN), that was not previously known to associate with the APC was found predominantly in G1 and in the “mitosis” sample. Several other known interactors and substrates such as EMI1 and geminin, were identified in the appropriate cell cycle phase. In addition several other proteins of interest also co-purified with the APC.
Undertaking a comprehensive phosphorylation analysis we were able to map 71 distinct phosphorylation sites under 9 different conditions. Table 1 shows a list of phosphorylation sites for the APC under the conditions described. The spectral evidence for the phosphorylation sites is shown in the SI Figure 6, 12-65. In four cases the phosphorylation sites could not be unambiguously assigned to one amino acid residue. In these cases, two possible sites are indicated in the SI Figures12-65. As there were differences in the sequence coverage from treatment to treatment, we could not assume that a phosphorylation that was not observed in a given condition was in fact not present. Unobserved sites are listed in the Table 1 as absent (white) only if we saw the corresponding unphosphorylated peptide in at least one of the samples.

The sites for cells undergoing mitosis are different from those found in nocodazole arrest

The greatest number of phosphorylation sites appeared in drug-induced mitotic arrests. After nocodazole treatment, we found 43 phosphorylation sites, whereas in S- and G1-phase we found only 2 and 5 sites, respectively. The data for S-phase and nocodazole treatment were in general agreement with the work of Kraft et al. (4). In a few cases we differed with respect to the exact location of the phosphorylation site.

In most cell cycle studies nocodazole arrest is used as a surrogate for the mitotic state. To test the validity of that assumption, we compared phosphorylation in the nocodazole arrested cells with that in a set of cells undergoing mitosis, but not arrested in mitosis. We compared cells from double thymidine arrest released for 6-8 hours. These cells had a robust mitotic index of 70%, compared to 90% for nocodazole arrest. However, the spectrum of phosphorylation was very different for some APC subunits but similar for others. In the mitosis samples APC1 was phosphorylated at 5 sites, as compared to 11 for nocodazole arrest. Only 3 of the phosphorylation sites are common between the two conditions. The phosphorylation state for CDC27, APC4, and CDC16 were more similar between treatments. It is interesting to note that the subunits that showed the most similarity in phosphorylation between drug treatments contain protein interaction domains such as TPRs (tetratrico peptide repeats) and WD40 repeats.
Different spindle poisons give different qualitative phosphorylation patterns

We then extended our analysis to other mitotic poisons. As shown in Figure 1 a compilation of phosphorylation site data for the largest APC subunit, APC1, shows very different patterns of phosphorylation for different spindle poisons. While several phosphorylation sites are shared among the treatments, such as Ser202, Ser377 and Ser688, there are sites that appear to be specific to certain drugs. For example Ser233 is seen to be phosphorylated in nocodazole and taxol but not vincristine or monastrol, while Ser1552 is specific for taxol arrest. As samples were analyzed more than once, we were able to establish reproducibility. All the phosphorylation sites listed were found in more than one analysis of the same treatment.

Though these differences were reproducible, there is still some concern about the threshold of detection. Phosphorylation sites that are basally phosphorylated at a very low stoichiometry are sometimes detected by mass spectrometry, while phosphorylations with high stoichiometries may not be detected, perhaps because the phosphopeptide co-elutes with other peptides, resulting in undersampling. Although the samples were handled identically and we monitored the total amount of protein injected for each LC/MS analysis, we could not rule out relative differences in protein concentration, a crucial factor in determining the number of phosphorylation sites observed. Thus, to determine whether the differences we observed are real, we needed to provide a quantitative measure of phosphorylation that is comparable among drug treatments.

Quantitative studies confirm that phosphorylation patterns resulting from different drug treatments are distinct

Determining the degree of phosphorylation by mass spectrometry would be easy if an unmodified peptide and its phosphorylated cognate had the same flyabilities, as one could directly compare the signal intensities of these two species. However, we have recently found that using the same experimental conditions, the flyability ratio for a given peptide/phosphopeptide pair remains constant from experiment to experiment when the same peptide pair is examined (24, 25). Thus reliable information about relative changes
can be derived when the signals from identical peptides/phosphopeptide pairs are compared for the different drug treatments. In order to generate the data shown in Table 2 we determined which slices contained the protein of interest SI Figure 2-6 then we measured relative intensities of the phosphorylated peptides vs. their unphosphorylated complements. Not all of the 71 peptides mapped qualitatively provided good quantitative data. There are several explanations for this. Most frequently, a single phosphorylation site was spread over multiple peptide species, complicating quantitation. This occurs when: 1) there are multiple tryptic cleavage sites close together, resulting in a “ragged peptide”; 2) the peptide contains e.g. a methionine residue which is easily partially oxidized; or 3) the peptide is multiply modified. For 33 of the phosphorylation sites, we obtained reliable quantitative information for both the modified and unmodified peptides to determine the ratio of phosphorylated vs. unphosphorylated peptide.

The phosphopeptide vs. peptide signal intensity ratio (Equation 1) allows for an easily accessible quantitative measure that can be compared between treatments. Here \( I_{pP1} \) is the measured intensity of the phosphopeptide and \( I_{P1} \) is the intensity of the unphosphorylated cognate peptide, \( \mu_{pP} \) is the flyability of the phosphopeptide and \( \mu_P \) the flyability of the unphosphorylated cognate peptide. \([pP_1]\) is the concentration of the phosphopeptide and \([P_1]\) is the concentration of the unphosphorylated cognate peptide. The same equation was applied to another drug treatment for the same phosphopeptide/peptide pair in Equation 2. Thus, two drug arrest conditions can easily be compared as a ratio (Equation 3). It is obvious that the flyability ratios cancel out if the mass spectrometry conditions are kept similar such that the ratio of phosphopeptide vs. peptide signal intensity ratios is equivalent to the ratio of phosphopeptide vs. peptide concentration ratios (Equation 4).

\[
\frac{I_{pP1}}{I_{P1}} = \frac{\mu_{pP}[pP1]}{\mu_P[P1]} \quad (1)
\]

\[
\frac{I_{pP2}}{I_{P2}} = \frac{\mu_{pP}[pP2]}{\mu_P[P2]} \quad (2)
\]
This simple expression of the phosphorylation state facilitates the comparison between two or more drug treatments. The values for Equation 1 are provided and the qualitative “heat map” shown in Table 2. The heat map shows the ratio of the signal intensity of the phosphopeptide to that of the unphosphorylated cognate. The color coding is as follows: red indicates a ratio of over 1, orange 1 to 0.1, dark yellow 0.1 to 0.02 and light yellow detectable levels below 0.02.

Table 2 shows clearly that the different qualitative patterns of phosphorylation for different drugs cannot simply be explained by differences in the sensitivity of the analysis between samples. For example taxol treatment gives high levels of phosphorylation (ratio > 1) on 6 sites: APC1: Ser377 (# 8; ratio: 2), Ser688 (# 14; ratio: 1), Ser731 (# 15; ratio: ∞); APC2: Ser218 (# 18; ratio: 2.5), Ser314 (# 19; ratio: 75), and CDC23: Ser582 (# 62; ratio: 9). By comparison, vincristine treatment gives high levels of phosphorylation on only 4 sites. Although 3 of these sites are identical to those found in taxol arrest, one site is remarkably different: CDC16 (50) Ser559 is phosphorylated at a level of 1.45 in vincristine but only at 0.04 in taxol; CDC23 (62) Ser582 is phosphorylated at a level of 3.9 in the vincristine treatment and 0.8 in the taxol treatment. An even stronger contrast to vincristine and taxol is nocodazole vs. taxol. In nocodazole, 7 sites are phosphorylated to a high degree and these sites are found on APC1, APC2 and CDC27 and CDC16. In general CDC27 is more highly phosphorylated in the nocodazole and vincristine treatments than it is in the taxol treatment. A few sites are highly phosphorylated among all or many of the treatments but not in the normal mitotic progression. These sites
include APC2: (19) Ser314; CDC27: (25) Thr205, (37) Ser426; CDC16: (50) Ser559. Interestingly APC2, a core catalytic subunit which contains the cullin protein motif shows the highest phosphorylation ratio of all the sites observed. A remaining question is whether the patterns are stable or change over time.

**Phosphorylation patterns in nocodazole-arrested cells change over time**

To study dynamic changes in phosphorylation patterns, quantitative measurements of phosphorylation that are accurate to < 20% are needed. The error level of estimating the recovery of different peptides in multiple gel slices is too high for this kind of study. We therefore processed the entire sample after brief electrophoresis as a means of purification and immobilization without further fractionation. The complexity of the mixture reduced the number of peptides that could be identified; only 25 phosphopeptides were found using either Mascot or Protein Pilot.

On inspecting the initial data from a number of time points, several of the phosphopeptides appeared not to be at steady state. We used iQEM to look more closely at the quantitative kinetics of phosphorylation. As peptides and their phosphorylated cognates have differential ionization/detection efficiencies (here also called ‘flyability’), the phosphorylation stoichiometry cannot directly be derived from the relative peak intensities. However, although the ionization/detection efficiencies of a peptide/phosphopeptide pair are not known, their changes are correlated. This enables one to calculate the ratio of ‘flyabilities’ which, in turn, allows one to derive the degree of phosphorylation based on the signal intensities when data for several samples with significantly different degrees of phosphorylation are available. We provided the proof of concept in two recent papers (24, 25) and introduce the term iQEM (isotope-free Quantitation of the Extent of Modification) for this approach. iQEM allows us to determine the flyability ratio for a peptide/phosphopeptide pair which in turn enables the calculation of the phosphorylation stoichiometry, not just a ratio of stoichiometries, which we used in the previous example; this estimate does not require the use of isotope-labeled standards. Instead of isotope-labeled standards, iQEM uses a set of peptides derived from the protein of interest, chosen because they do not show sample-to-sample
variation and are therefore presumably unmodified, as omnipresent internal standards. By characterizing the properties of these peptides and then detecting their presence in our samples, we can normalize/correct for variations of the absolute amount of the target protein present in the extract. As we can then assume conservation of mass we can calculate the ionization/response factors as the decrease of the one signal e.g. the peptide is correlated with the increase of the cognate signal of the phosphopeptide (24).

In this case 4 or 5 standard peptides were used per protein for normalization. We chose our normalization peptides by looking at peptides that gave ion currents variations among the samples that reflected the general variations of the majority of the peptides. To illustrate the method, the expected behavior of peptides from a single protein measured over several time points is shown in a cartoon (SI Figure 7). Blue lines represent peptides that are not modified in the biological condition under study; these show a similar pattern over the time course, with the intensities measured depending largely on the amount of sample injected onto the HPLC column. Black lines show peptides that are modified during the time course; these start out following the same pattern as the blue lines, but sharply diverge at some point. The green and yellow lines show the concomitant increase of the corresponding modified peptides. In every LC/MS time course there will be some peptides that show an aberrant intensity pattern (red lines). These peptides often contain e.g. methionine, which undergoes unpredictable oxidation.

Using well-behaved peptides (blue lines) we can accurately normalize phosphorylated and unmodified peptides through a simple calculation. The phosphorylation stoichiometry can be determined based on the relative ion signal intensities of the two species after correcting for the different response factors by knowing the flyability ratio of a peptide/phosphopeptides pair calculated based on the conservation of mass (for details, see Ref. (24)). The flyabilities can be very different for the phosphorylated and unphosphorylated forms. Hence the uncorrected ion currents are unreliable for determining the precise amount of peptide present. In Figure 2A, for example, CDC16 peptide NIIs559PPWDFR displays a remarkable 60-fold increase in the intensity of the phosphopeptide and only a 3-fold decrease for the unphosphorylated cognate. Without
iQEM normalization this result is uninterpretable, but using iQEM we find that the stoichiometry of increase and decrease is exactly complementary (Figure 2B). Phosphorylation at that site increases from 0% to 80% and the cognate unphosphorylated peptide decreases from 100% to 20%. We give additional examples of such peptides in SI Figures 8 and 9.

Using this method we can show that in nocodazole arrest, APC phosphorylation changes significantly over time. Dynamic changes in phosphorylation levels for 4 different phosphorylation sites, 1 on APC2 and 3 on CDC27, are shown in Figure 2C. Each site shows different kinetics. The rate of phosphorylation at Ser426 on CDC27 steadily increases until the 9 hour time point, while that at Ser435 increases more rapidly from 2 to 8 hours and then the rate of phosphorylation decreases. This decrease in the rate of Ser435 phosphorylation is coincident with the detection of a second phosphorylation site on Ser438, manifested by the detection of the doubly phosphorylated peptide species. Phosphorylation at the Ser534 site on APC2 is undetectable until the 12 hour time point. We show other peptides quantitated in a similar fashion in SI Figures 10 and 11. Thus, quantitative measurements show that nocodazole “arrest” is not a static state: APC phosphorylation patterns change significantly depending on the duration of the arrest.

Discussion

We studied three aspects of the phosphorylation patterns on the 11 core proteins of the APC plus interactors: i) qualitative phosphorylation mapping, ii) relative quantitation to detect differences and iii) where necessary, quantitation in actual stoichiometries by iQEM. We used standard methods to identify 71 sites of phosphorylation, under a variety of conditions, and we were also able to use non-normalized ion currents to compare phosphorylation levels for a subset of these sites at different points in the cell cycle and in mitotic arrest in response to anti-mitotic drugs. Neither of these approaches was sufficiently quantitative to measure kinetic features of phosphorylation. For that we used a newly developed iQEM technique, which provides accurate absolute stoichiometry without using isotope-labeled standards.
Using these methods, we were able to uncover differences between the mitotic arrest states imposed by different spindle poisons and distinguish between these states and the normal metaphase condition. We also found that mitotic arrest in response to the spindle poison nocodazole is a dynamic, not a static, state.

**Drug-arrested mitotic states are not a faithful reflection of normal mitosis**

Our analysis of the phosphorylation state of the APC at different cell cycle stages allowed us to show that the pattern of phosphorylation in prometaphase arrest by nocodazole was significantly different from that of cells synchronized in mitosis after timed release from double thymidine block. We considered two different explanations for this disparity. The first possibility is that nocodazole might arrest cells at a specific short-lived point in the mitotic cycle, which we might have missed in our studies of the synchronized but unarrested cell cycle. The semi-quantitative data on nocodazole arrest, suggest that this is not likely to be the case. For example Thr205 (25) on CDC27 showed a phosphorylation level in nocodazole of 1.2 while that same site in the mitotic state was 0.09. Such a high phosphorylation level could only be masked if the state stabilized by nocodazole represents less than 4% of the time spent in normal mitosis (90 minutes); such extremely rapid changes in phosphorylation seem unlikely. Similarly Ser221 (47) in APC5 had a ratio of 0.2 but was undetectable in the normal mitotic cells.

A second explanation might be that the nocodazole state of arrest is not a subset of mitosis but a specific response to microtubule depolymerizing drugs. If this were true then a similar response would be expected for vincristine, which is also a microtubule depolymerizing agent. Vincristine arrest does show six of the same phosphorylation sites on APC1 as nocodazole, but we also found a number of unique sites (five for nocodazole and four for vincristine). We cannot attribute the difference between nocodazole and vincristine to peptides that were systematically missed in the different analyses, since we observed each corresponding unphosphorylated peptide in both sets of samples and the analyses were repeated.
It is clear that the nocodazole-arrested state, at least as reflected in APC phosphorylation, does not represent a well populated state found in normal mitosis. Although all the phosphorylations found in mitosis are also observable in the drug arrest, we find many specific to the drug-arrested state. Further work will be required to determine the causes and effects of these extra phosphorylations.

_mass spectrometry offers a new way of examining the differences between different spindle poisons_

Some drugs that bind to microtubules and block mitosis are ineffective in cancer treatment; others show inexplicable focal efficacy. For example, the vinca alkaloids are useful for treating lymphoma, neuroblastoma and nephroblastomas, while taxol is useful for advanced breast cancer and ovarian cancer. We do not know why these drugs are not all equally effective, nor why they have different therapeutic value against different cancers. It is possible, perhaps likely, that our observations of distinct phosphorylation states of the APC in response to different anti-mitotic drugs may point to an explanation of some of these differences. Furthermore, it is possible that cells from different tissues, or cells harboring different mutations, or cells under different physiological stresses, such as hypoxia, may differ in their response to spindle poisons and would reflect those differences in different sites of phosphorylation. In that respect differences in spindle checkpoint phosphorylation may reveal new features of the mitotic state.

It is also possible that anti-mitotic agents differentially affect the activity of kinases, such as CDK1 and aurora B, leading to variations in spindle checkpoint responses. The ability to characterize drug candidates based on the spectrum of APC phosphorylations may facilitate the categorization of drugs, the discrimination of the response of tumors to drugs, and the identification of new means of checkpoint control.

**Dynamics of Phosphorylation.**

Our results emphasize the fact that mitotic arrest is a misnomer: arrest is a dynamic state in which some cells enter apoptosis and other cells revert to interphase. The ability to observe biochemical events during arrest could be very important for understanding anti-
proliferative treatments. However, exploring the dynamics of phosphorylation makes great demands on the accuracy of quantitation. Most mass spectrometric based quantitative approaches, including SILAC and iTRAQ, give relative data, meaning that one state of phosphorylation is determined relative to another phosphorylation state (26-28); these data are insufficient to establish the kinetics of a pathway. iQEM offers a significant advance over earlier techniques. It has allowed us to measure specific quantitative changes in APC phosphorylation in cells arrested in nocodazole for varying periods. Although we do not yet understand the specific dynamics that we have discovered here, they are intriguing. If these can be correlated with the process by which the arrested state is resolved, they may provide us with new tools to understand the mitotic process and to find more effective drug targets in cancer.

Materials and Methods

Chemicals
All chemicals, if not otherwise stated, were purchased from Sigma-Aldrich (St. Louis, MO). HPLC-grade solvents for mass spectrometric analyses were acquired from Burdick and Jackson (Morristown, NJ).

Preparation of synchronized HeLa cell extracts
HeLa S3 cells were obtained from ATCC. Cells were grown in spinner flasks in Dulbecco’s Modified Eagle Medium (DMEM) supplemented with 10% FBS and 100 µg/ml penicillin/streptomycin. Cells were synchronized at the G1/S transition by double thymidine block as previously described. Details of the anti-mitotic drug treatments are described in the supplementary materials. All cultures were analyzed by FACS to ensure the correct cell cycle state. Cell extracts were prepared by lysing the cell pellet in 1 volume of PBS buffer supplemented with 0.1% Triton X-100, 1 nM okadaic acid, 1 µg/ml microcystine-LR (Alexis Biochemicals), 10 mM sodium fluoride, and 10 µg/ml each of chymostatin, pepstatin and leupeptin. The cells were passed through a 27 gauge needle to shear the DNA. The extract was precleared by centrifugation for 20 min at 20,000 xg at 4°C.
**Immunoprecipitation of APC from extracts**
The APC was immunoprecipitated using the anti-CDC27 antibody sc-6392 (Santa Cruz). Details of the procedure are described in the supplementary materials.

**Time course experiments**
Cells were synchronized using a double thymidine arrest in G1/S phase as previously described and fresh medium was added after the cells were washed. After 4 hours nocodazole was added to the cells. Time points were taken at 0, 3, 6, 8, 10 and 12 hours. This particular experiment was repeated 3 times with varying time points in order to gauge the time windows for appropriate measurements. Phosphorylation sites on CDC27 were used to check the reproducibility of the kinetics between experiments. Immunoprecipitations were performed as described above and the samples were run into a 10% gel but not separated. The protein was then in-gel digested and analyzed using LC/MS (see below).

**Mass spectrometry**
All experiments were performed by using a QSTAR XL mass spectrometer (Applied Biosystems/MDS Sciex, Concord, Canada) coupled to a microscale capillary HPLC (Famos microautosampler (LC Packings, Sunnyvale, CA), Agilent 1100 HPLC pump (Agilent, Andover, MA)). Columns (length: 15 cm, inner diameter: 75µm) were packed in-house by using Magic C18 beads (Michrom BioResources, Auburn, CA). Buffer A was 2.5% acetonitrile/0.2% formic acid; buffer B was 97.3% acetonitrile/0.2% formic acid.

**ACKNOWLEDGEMENTS**
We would like to thank Ron Bonner, Lyle Burton and Keith Ashman, for helpful discussions. We thank the National Institutes of General Medical Sciences (GM039023) for support of this work.
**Figure and Table Legends**

**Table 1.** Phosphorylation sites mapped for the different APC subunits and related spindle checkpoint proteins. The green shading shows the presence of the phosphorylation site and the white indicates the absence of a phosphorylation site.

**Table 2.** A heat map of the quantitative measurements of phosphorylation stoichiometries of the sites found for each treatment (mn: monasterol; n: nocodazole; t: taxol 20mM; vc: vincristine; M: double thymidine followed by a 8 hour release into mitosis).

**Figure 1.** A cartoon depicting the phosphorylation sites identified on APC1 during drug-induced mitotic arrest. The arrows are color coded according to the drug treatment and the number above the site indicates the site of phosphorylation.

**Figure 2. A)** Mass spectra of the CDC16 peptide NIISPPWDFR and its phosphorylated counterpart at the 0, 6 and at the 12 hour time point. The given numbers correspond to the peak intensities after averaging over the elution time window. B) Graphical representation of the in-/decrease of NIIpSPPWDFR and NIISPPWDFR. C) A plot of the phosphorylation stoichiometries of 4 different phosphorylation sites: APC2: S534; CDC27: S426, S435 and S438 as determined using the iQEM approach. Clearly different kinetics of phosphorylation are observed.
REFERENCES