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Emi1 switches from being a substrate to an inhibitor of APC/ C^{Cdh1} to start the cell cycle

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Abstract

Mammalian cells integrate mitogen and stress signaling prior to the end of G1 phase to decide whether or not to enter the cell cycle^{1–4}. Before cells can replicate their DNA in S phase, they have to activate cyclin-dependent kinases (CDKs), induce an E2F transcription program, and inactivate an E3 ubiquitin ligase, the anaphase promoting complex/cyclosome (APC/C^{Cdh1}). It was recently shown that stress can return cells to quiescence after CDK2 activation and E2F induction but cannot after inactivation of APC/C^{Cdh1}, arguing that APC/C^{Cdh1} inactivation is the point-of-no-return for cell cycle entry³. While rapid inactivation of APC/C^{Cdh1} requires early mitotic inhibitor 1 (Emi1)^{3,5}, the molecular mechanism controlling this cell cycle commitment step is unknown. Here we show that cell cycle commitment is mediated by an Emi1-APC/C^{Cdh1} dual-negative feedback switch, in which Emi1 is both a substrate and an inhibitor of APC/C^{Cdh1}. The inactivation switch triggers a transition between a state with low Emi1 levels and high APC/C^{Cdh1} activity during G1 to a state with high Emi1 levels and low APC/C^{Cdh1} activity during S and G2. Cell-based analysis, in vitro reconstitution, and modeling data show that the underlying dual-negative feedback is bistable and represents a robust irreversible switch. Together, our study argues that mammalian cells commit to the cell cycle by increasing CDK2 activity and Emi1 mRNA expression to trigger a one-way APC/C^{Cdh1} inactivation switch mediated by Emi1 transitioning from a substrate to an inhibitor of APC/C^{Cdh1}.

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Contributions

S.D.C designed and carried out most experiments. K.G.M performed in vitro ubiquitination assays. D.G. performed synchronization and western blot experiments. L.R.P. made lentivirus and stable cell lines. M.R. supervised the in vitro ubiquitination studies. S.D.C and T.M. conceived the project, analyzed the data, and wrote the manuscript. All authors discussed the results and the manuscript. T.M. supervised the study.

Competing financial interests

M.R. is a co-founder and consultant to Nurix, a biotech company in the ubiquitin space.

To gain insights into the molecular control of APC/C^{Cdh1} inactivation, we used a live-cell reporter for APC/C^{Cdh1} activity³ and tested in non-transformed human MCF10A breast epithelial cells whether APC/C^{Cdh1} inactivation has the hysteresis characteristic required for an irreversible cell cycle commitment decision. As outlined in Fig. 1a, bistable decisions in cell signaling require hysteresis, which means that only weak inhibition of the trigger activity should keep APC/C^{Cdh1} “On” (solid line) while strong inhibition of the same trigger activity should keep the inactivated APC/C^{Cdh1} switch “Off” (dashed line) (Extended Data Fig. 1a-c). When we titrated a CDK1/2 inhibitor during G1 phase when APC/C^{Cdh1} was “On”, or during S or G2 phase when APC/C^{Cdh1} was “Off”, we found that the EC50 to maintain APC/C^{Cdh1} in the “On” state was 1.68 μ M, while the EC50 to turn inactive APC/C^{Cdh1} back to the “On” state was higher than 30 μ M (Fig. 1b and Extended Data Fig. 1e). Thus, cells stay in their respective “On” or “Off” APC/C^{Cdh1} state over a greater than 20-fold concentration window of the CDK1/2 inhibitor, demonstrating robust hysteresis. When we measured the fraction of cells that failed to turn APC/C^{Cdh1} “Off” as a function of APC/C^{Cdh1} activity at the time of the drug spike (Extended Data Fig. 1f,g), we found that ~70% of inactivation reflects a threshold APC/C^{Cdh1} activity when APC/C^{Cdh1} inactivation becomes irreversible. Together, the CDK2-regulated trigger mechanism, the marked hysteresis, and threshold argue that APC/C^{Cdh1} inactivation is a robust bistable switch.

For a signaling system to generate a bistable switch, it requires in addition to hysteresis a positive or dual-negative feedback⁶ (Fig. 1a). We first investigated known APC/C^{Cdh1} substrates that may also negatively regulate APC/C^{Cdh1} to generate dual-negative feedback. The cullin E3 ligases SCF^{Skp2} and SCF^{Cyclin F} have both been reported to degrade APC/C^{Cdh1} components^{7,8}, and Cyclin A2/CDK2 can mediate APC/C^{Cdh1} inhibition by phosphorylating Cdh1^{9,10}. Knockdown of Cyclin A2, Skp2, or Cyclin F (Extended Data Fig. 2a-c), did not affect the inactivation kinetics of APC/C^{Cdh1} in three cell types (HeLa, MCF10A, and U2OS; Fig. 1c and Extended Data Fig. 3a-c), suggesting that these substrates may tune APC/C activity in other phases of the cell cycle but do not control the rapid APC/C^{Cdh1} inactivation at the G1/S transition. In contrast, knockdown of the APC/C^{Cdh1} inhibitor Emi1 (alias: Fbxo5)^{5,11}, resulted in a significant decrease in the rate of APC/C^{Cdh1} inactivation (Fig. 1c and Extended Data Fig. 2d and 3a-c). Strikingly, hysteresis in APC/C^{Cdh1} inactivation was completely lost when Emi1 was knocked down (Fig. 1d and Extended Data Fig. 1e and 3d). Thus, Emi1 is responsible for the fast kinetics, bistability, as well as hysteresis observed in APC/C^{Cdh1} inactivation.

We measured Emi1 protein levels during the cell cycle to determine whether Emi1 may not only inhibit APC/C^{Cdh1}, but may also be part of a dual negative feedback by being degraded by APC/C^{Cdh1} in G1 (Extended Data Fig. 3e). Indeed, Emi1 protein levels were low in G1 phase when APC/C^{Cdh1} is active and first started to accumulate about 6 hours after mitosis coincident with the inactivation of APC/C^{Cdh1} (Fig. 2a and Extended Data Fig. 4a,b). Low Emi1 protein levels in G1 phase are a consequence of Emi1 degradation during G1 as treatment with the proteasome inhibitor MG132 increased Emi1 levels (Extended Data Fig. 4c). This degradation of Emi1 is unlikely to be mediated by Polo-like kinase and SCF ^{β -TrCP}¹² as these regulators of Emi1 degradation are active in pro-metaphase and not in G1 phase¹³. Inhibition with the small molecule APC/C inhibitor proTAME¹⁴ increased

Emi1 levels as well as the level of the APC/C substrate Cyclin A2 (Fig. 2b), consistent with Emi1 being degraded in G1 by APC/C^{Cdh1}.

In a second strategy to test whether Emi1 is a substrate of APC/C^{Cdh1} in G1, we used single-cell analysis of both Emi1 mRNA copy number and protein levels during G1 (Extended Data Fig. 4d). We confirmed that the mRNAs of the known E2F targets Emi1, Geminin, E2F1, and Cyclin E1 all immediately start to increase after anaphase (Fig. 2c and Extended Data Fig. 4e,f), and that Cyclin E1 protein levels closely follow Cyclin E1 mRNA levels (Fig. 2d and Extended Data Fig. 5a). In contrast, the increase of both Geminin and Emi1 protein levels were significantly delayed compared to their respective mRNAs (Fig. 2c,d and Extended Data Fig. 5b,c). Both Geminin, a well-characterized APC/C^{Cdh1} substrate, and Emi1 proteins levels started accumulating about 5-6 hours after anaphase, precisely when APC/C^{Cdh1} is inactivated and EdU starts to be incorporated at the start of S phase (single cell analysis in Extended Data Fig. 5d,e). We also tested for a role of APC/C^{Cdh1} in degrading Emi1 by knocking down the APC/C co-activator Cdh1 with siRNA, and found that both Geminin and Emi1 protein levels accumulated much earlier compared to control (Extended Data Fig. 5f,g), further confirming that APC/C^{Cdh1} is required for Emi1 degradation in G1.

To determine whether Emi1 is a direct substrate of APC/C^{Cdh1}, we used recombinant Emi1 and immuno-purified APC/C^{Cdh1} along with ubiquitin, E1, E2s (Ube2c and Ube2s), and ATP to test for Emi1 ubiquitination *in vitro*. Markedly, for relatively low concentrations of Emi1 in this assay (23 nM), APC/C^{Cdh1} efficiently ubiquitinated recombinant Emi1, providing evidence that Emi1 is a direct substrate for APC/C^{Cdh1} (Fig. 2e). A kinetic analysis of Emi1 ubiquitination revealed that Emi1 is a distributive substrate¹⁵, meaning that Emi1 poly-ubiquitination requires multiple successive encounters of Emi1 with APC/C^{Cdh1} (Extended Data Fig. 5h,i). Using mutant ubiquitins that lack one or all lysines, we also found that Emi1 is primarily poly-ubiquitinated via K11 and K48 ubiquitin linkages, similar to other APC/C^{Cdh1} substrates¹⁶ (Extended Data Fig. 5j).

Having established that Emi1 is an APC/C^{Cdh1} substrate that can be degraded by APC/C^{Cdh1} in G1 phase, the question arises if and how Emi1 can also be an inhibitor of APC/C^{Cdh1} in S/G2 phase. While it is conceivable that this transition is regulated by post-transcriptional modifications on Emi1¹⁷, previous structural data identified binding interactions between two domains of Emi1 and two different regions of APC/C^{Cdh1}^{18,19} (Fig. 3a and Extended Data Fig. 6a,b), which suggested that Emi1 may switch from being a substrate to an inhibitor by using two separate binding sites on APC/C^{Cdh1}. First, Emi1 binds to the substrate adapter proteins Cdh1 and APC10 through an RxxL D-Box motif, similar to typical APC/C^{Cdh1} substrates. Second, Emi1 binds to APC1, APC2, and APC11 through its Zinc Binding Region and to APC2 via its C-terminal LRRL sequence, blocking the binding of the E2 ubiquitin conjugating enzymes. We tested mutants of Emi1 deficient in each of the two binding interactions and used MCF10A cells where addition of a CDK4/6 inhibitor can block APC/C^{Cdh1} inactivation that normally occurs in most cells within 5 hours from mitosis (Fig. 3b and Extended Data Fig. 6c-e). We first confirmed that over-expression of Emi1^{WT} potently inhibited APC/C^{Cdh1} immediately after mitosis in the presence of the CDK4/6 inhibitor, showing that Emi1 can turn APC/C^{Cdh1} “Off” on its own (Fig 3c and Extended

Data Fig. 6f,g). We made a point mutation (Emi1^{C401S}; Extended Data Fig. 6d) that was previously shown to disrupt the ZBR domain of Emi1 and thereby prevent Emi1^{C401S} from inhibiting the binding of E2 enzymes to APC/C^{Cdh1}¹¹. Markedly, expressed Emi1^{C401S} failed to inactivate APC/C^{Cdh1} in this system (Fig. 3d and Extended Data Fig. 6f,h) and was instead degraded in the absence of CDK4/6 inhibitor in parallel with other APC/C^{Cdh1} substrates (Extended Data Fig. 6j-k), arguing that an interaction of Emi1 with the substrate region alone can lead to Emi1 ubiquitination and degradation without inhibiting APC/C^{Cdh1}.

To determine whether interaction of Emi1 with only the region where the E2 enzymes bind is sufficient to inhibit APC/C^{Cdh1}, we mutated the APC/C degron motif in Emi1 to prevent its interaction with this substrate docking site (Emi1^{AxxA}; Extended Data Fig. 6d). Expression of the Emi1^{AxxA} mutant rescued the effects of CDK4/6 inhibition to control levels and almost as well as overexpressing Emi1^{WT} (Fig. 3b,c,e and Extended Data Fig. 6l), arguing that Emi1 binding to the E2 enzyme interaction region can be sufficient to inhibit APC/C^{Cdh1} without requiring the interaction with the substrate region. Independent interactions of Emi1 with the substrate and inhibitory regions of APC/C^{Cdh1} are further supported by the previous finding that Emi1 and the APC/C^{Cdh1} substrate Geminin can both bind to APC/C^{Cdh1} at the same time as part of a ternary complex²⁰. Together with the previous structural and biochemical results^{18–20}, our data suggests that Emi1 is functioning independently as a substrate of APC/C^{Cdh1} by binding to the substrate region but is also an inhibitor of APC/C^{Cdh1} by binding to the E2 enzyme interaction region or at the same time to both regions (Fig. 3f).

We developed a computational model to determine how an inhibitor of an E3-ligase that is also a substrate may generate a robust bistable switch (Extended Data Fig. 7a). In the model, Emi1 reversibly binds to a low affinity substrate binding site of APC/C^{Cdh1} where it can be mono- and poly-ubiquitinated, in a non-processive manner. Emi1 also reversibly binds with higher affinity to the inhibitory site(s) to suppress poly-ubiquitination and subsequent degradation of substrates by inhibiting E2 enzyme binding and ubiquitin chain elongation²⁰ (Fig. 3f). At low Emi1 concentrations, Emi1 and other substrates can be degraded as Emi1 levels are initially lower than APC/C^{Cdh1} levels and the fraction of inhibited APC/C^{Cdh1} is therefore insufficient to block overall APC/C^{Cdh1} activity. As Emi1 levels increase, the increasing occupancy of the inhibitory site of APC/C^{Cdh1} suppresses poly-ubiquitination of unbound Emi1 and other substrates. This system has the cooperativity required for a robust switch mechanism (Fig. 1a and Extended Data Fig. 3e) since multiple ubiquitins have to be added to enable substrate degradation^{21,22}. The model includes a term for E2F-regulated synthesis of Emi1 mRNA after mitosis and also for a slow increase in CDK2 activity that partially inhibits APC/C^{Cdh1} to initiate the switch mechanism which cells trigger at a threshold of ~70% APC/C^{Cdh1} inhibition (Extended Data Fig. 1f,g).

A steady-state analysis of the model at constant Emi1 mRNA levels and CDK2 activity (Fig. 3g) shows the relationship between Emi1 levels and synthesis rate represents a bistable system with an APC/C^{Cdh1} “On”, low Emi1 state (Fig. 3g, left panel), and an APC/C^{Cdh1} “Off”, high Emi1 state (Fig. 3g, middle panel). Once the system reaches the “Off” state, APC/C^{Cdh1} activity becomes independent of CDK2 activity (Fig. 3g, right panel), a

prediction that closely matches the hysteresis characteristic we observed experimentally (Fig. 1b and Extended Data Fig. 7b).

A second key prediction of the model is that at concentrations of Emi1 below APC/C^{Cdh1}, Emi1 should be efficiently ubiquitinated by APC/C^{Cdh1}, but at concentrations higher than APC/C^{Cdh1}, Emi1 should be poorly ubiquitinated. To test this prediction, we conducted *in vitro* ubiquitination assays with both low and high concentrations of Emi1. Emi1 was again efficiently poly-ubiquitinated when we added 23 nM of recombinant Emi1 to an *in vitro* ubiquitination assay, but poly-ubiquitination of Emi1 was completely inhibited at 2.9 μ M Emi1, consistent with robust APC/C^{Cdh1} inhibition (Fig. 4a). This is also true for other substrates as Geminin was poly-ubiquitinated by APC/C^{Cdh1} in the presence of 23 nM Emi1 but Geminin poly-ubiquitination by APC/C^{Cdh1} was completely inhibited when we added 2.9 μ M Emi1 (Extended Data Fig. 8a). Emi1 added at the higher concentration could still be mono-ubiquitinated by APC/C^{Cdh1} as was Geminin (Fig. 4b and Extended Data Fig. 8b), consistent with a previous observation that Emi1 may primarily inhibit ubiquitin chain elongation by interfering with Ube2S binding^{18,20}. We also observed the same concentration-dependent ubiquitination of Emi1 when we used APC/C^{Cdc20} in the *in vitro* ubiquitination assays, indicating that Emi1 is also a substrate of APC/C^{Cdc20} (Extended Data Fig. 8c). Using densitometry of the ubiquitinated bands, we observed a nonlinear transition between being a substrate with 5 or more ubiquitins attached and an inhibitor (Extended Data Fig. 8d). Taken together, our data supports a model that Emi1 is a substrate of APC/C^{Cdh1} when present at low concentrations and an inhibitor of APC/C^{Cdh1} when it exceeds the concentration of APC/C^{Cdh1} (Fig. 4c).

Our model also predicted that cells should trigger bistable APC/C^{Cdh1} inactivation even when Emi1 mRNA increases alone without an increase in CDK2 activity (Fig. 4d-e and Extended Data Fig. 8e). Indeed, we found that HeLa and U2OS cells treated with siRNA against both Cyclin E1 and E2 (see Extended Data Fig. 2e) both rapidly inactivate APC/C^{Cdh1} after a delay, indicating that APC/C^{Cdh1} can indeed be inactivated independently of CDK2²³ activity if Emi1 synthesis is constitutively active and Emi1 mRNA can reach high enough levels (Fig. 4f and Extended Data Fig. 9a-f). Thus, cells can bypass the requirement for G1 Cyclin E/CDK2 activity by upregulating Emi1 mRNA levels to inactivate APC/C^{Cdh1}, providing a redundant mechanism to start the cell cycle (Fig. 4g and Extended Data Fig. 10a-d). This alternative path to S phase is likely important in cancer cells that often have mis-regulated E2F, Rb, or upstream regulators that constitutively activate E2F²⁴ (Extended Data Fig. 10d). Notably, we observed that knockdown of Emi1 in U2OS cells completely prevented APC/C^{Cdh1} inactivation (Extended Data Fig. 3c and 10c), suggesting that Cyclin E/CDK2 activity alone does not reach sufficiently high levels in U2OS cells to inactivate APC/C^{Cdh1} independent of Emi1. While CDK2 activity alone is capable of inactivating APC/C^{Cdh1} in some but not all cell lines, the dual-negative feedback loop between Emi1 and APC/C^{Cdh1} is required for rapid and irreversible APC/C^{Cdh1} in every cell line tested, indicating it is likely a general feature of cell cycle entry control.

Together, our results provide a mechanistic model for mammalian cell cycle commitment. We previously demonstrated that the activation of Rb-E2F and Cyclin E/CDK2, which is thought to underlie the mammalian restriction point, remains reversible when cells are

exposed to stress until APC/C^{Cdh1} is inactivated several hours later³. Thus, cells pass through two sequential steps in cell cycle entry: Rb inactivation followed by APC/C^{Cdh1} inactivation. Only after this second step can cells no longer return to quiescence. We now show that Emi1 mRNA and Cyclin E/CDK2 are synergistic triggers of a bistable switch mechanism that is driven by dual-negative feedback between Emi1 and APC/C^{Cdh1}. Figure 4h positions this Emi1-APC/C^{Cdh1} inactivation switch within the overall cell cycle, showing that cells turn APC/C^{Cdh1} “On” when they exit mitosis, keep APC/C^{Cdh1} “On” during quiescence and G1, before the Emi1-APC/C^{Cdh1} inactivation switch is triggered at the G1/S transition when cells irreversibly commit to the cell cycle.

Methods

Cell Culture

MCF10A cells were obtained from ATCC (CRL-10317) and were cultured in phenol red-free DMEM/F12 (Invitrogen) supplemented with 5% horse serum, 20,000 pg/mL EGF, 10 µg/mL insulin, 0.5 µg/mL hydrocortisone, 100 ng/mL cholera toxin, 50 U/mL penicillin, and 50 µg/mL streptomycin. HeLa (ATCC, CCL-2) and U2OS cells were cultured in DMEM (Invitrogen) plus 10% fetal bovine serum (FBS) and penicillin-streptomycin-glutamine (PSG). HeLa and U2OS cells were used because they both have constitutively active E2F-mediated transcription, which is necessary to test our hypothesis that Emi1 can inactivate APC/C independently of CDK2 activity. Where indicated, cells were treated with the small molecule inhibitors CDK1/2i III (3 µM, EMD Biosciences #217714), CDK4/6i (1 µM, PD-0332991), MG132 (10 µM), proTAME (30 µM, Boston Biochem).

Synchronization

HeLa cells were treated with 2 mM thymidine for 24 hours, washed, and allowed to grow in regular growth media for 3 hours. Cells were then treated with 100 ng/mL nocodazole for 9-12 hours. Cells synchronized in mitosis were then isolated by mitotic shake-off and replated after washing with regular growth media.

Constructs

CSII-pEF-H2B-mTurquoise and CSII-pEF-DHB(aa994-1087)-mVenus were described previously²³. CSII-pEF-Geminin(aa1-110)-mCherry was described previously³. Emi1 fused to mCitrine was cloned into the CSII-pEF lentiviral vector. Emi1 C355S mutation was made in the CSII-pEF-Emi1-mCitrine construct using the Gibson assembly method with the mutation encoded in the primers. Coding sequences for human His-Emi1^{aa299-447} was cloned into pET28 for purification from *E. coli*. Transduced cells were sorted on a Becton Dickinson Influx to obtain pure populations expressing the desired fluorescent reporters.

Time-Lapse Microscopy and Image Analysis

Cells were plated at least 24 hr prior to imaging in full growth media in a 96-well dish (Costar #3904) such that the density would remain sub-confluent until the end of the imaging period. Time-lapse imaging was performed in 290 µL full growth media. Images were taken in CFP, YFP, and RFP channels every 12 min on an IXMicro microscope (Molecular Devices) with a 10× objective, 0.3 NA. Total light exposure time was kept under

600 msec for each time point. Cells were imaged in a humidified, 37°C chamber in 5% CO₂. Image processing, cell tracking, and APC/C activity quantification was done using custom Matlab scripts described previously³.

Immunofluorescence

Cells were fixed in 4% paraformaldehyde, washed three times in PBS, permeabilized with 0.2% triton, and stained overnight at 4°C with anti-Skp2 (Cell Signaling Technology, #4358), anti-Geminin (Sigma-Aldrich, HPA049977), anti-Emi1 (Invitrogen, 37-6600), anti-Cyclin E1 (Santa Cruz Biotechnology, sc-247), or anti-Cyclin F (Santa Cruz Biotechnology, sc-952). Cells stained with anti-Cyclin E1 were treated with ice cold 100% methanol for 5 min after initially fixing with 4% paraformaldehyde. Primary antibodies were visualized using a secondary antibody conjugated to Alexa Fluor-647 and imaged with a Far Red filter. Where indicated, cells were treated with 10μM EdU for 15 min unless otherwise specified and fixed and processed according to manufacturer's instructions (Invitrogen, #C10356).

RNA FISH

RNA *in situ* hybridization was carried out using the Affymetrix Quantigene ViewRNA ISH cell assay kit and mRNA quantification was carried out using custom Matlab scripts as described previously³. Probes were custom designed (Affymetrix) to target Emi1, Geminin, Cyclin E1 or E2F1. After FISH staining, cells were blocked in PBS containing 10% FBS, 1% BSA, 0.1% TX-100, and 0.01% NaN₃ for 1 hour at room temperature, and then stained overnight with antibodies for immunofluorescence.

Live-cell imaging combined with fixed-cell analysis

To generate time course measurements of mRNA, protein, and EdU levels we combined live-cell imaging and fixed cell analysis. HeLa cells were used to simplify the analysis since HeLa cells do not enter quiescence after mitosis. We tracked unsynchronized cells using H2B-mTurquoise and imaged them for approximately 20 hours. At the end of the live-cell imaging, cells were grown in 10μM EdU for 15 minutes. The time since the previous anaphase before fixing was recorded for each individual cell. The cells were then stained for mRNA using RNA FISH (see above), protein using immunofluorescence, and EdU and imaged on the same microscope used for the live-cell imaging. Custom MATLAB scripts were used to re-align the fixed cell image with the last frame of the time-lapse imaging and then quantify the mRNA, protein, and EdU levels for each cell. Thus for each cell the time since anaphase, mRNA, protein, and EdU levels was recorded. Cells were binned in 1 hour intervals and the median value of each bin was calculated and plotted on an axis. Lines were normalized by the mRNA values at the onset of S phase (as measured by EdU incorporation), which was 6 hours after anaphase or to the maximum protein levels.

siRNA transfection

MCF10A and HeLa cells were transfected using Dharmafect 1 (Thermo Scientific) according to the manufacturer's instructions and U2OS cells were transfected with Lipofectamine RNAiMAX (Thermo Fisher Scientific) according to the manufacturer's instructions. The following siRNAs were used: control siRNA (nontargeting #2,

Dharmacon), siGenome pooled set of four siRNAs for cyclin A2, siGenome pooled set of four siRNAs for Skp2, siGenome pooled set of four siRNAs for cyclin F, siGenome pooled set of four for Emi1, siGenome pooled set of four for Fzr1 (Cdh1), siGenome pooled set of eight for cyclin E1 and E2 (Dharmacon) at final concentrations of 20nM unless noted. Six hours post-transfection, cells were washed with full growth medium and then imaging was immediately started. Cells were only considered if they went through one mitosis within the imaging period. This method works efficiently since all the genes targeted in this study are known to be degraded during or prior to the onset of mitosis. Thus, the siRNA would prevent subsequent accumulation in the next cell cycle.

***In vitro* ubiquitination Assay**

Human APC/C was purified from extracts of prometaphase or G1 HeLa cells¹⁶. For G1 extracts, prometaphase cells were washed twice in PBS and released from nocodazole-arrest for 2 hours in drug-free media. His-Emi1^{aa299-447} was purified as previously described^{16,25}. We used the C-terminal domain of Emi1 which contains all of the APC/C-interacting domains (amino acids 299-447) and is commonly used for *in vitro* ubiquitination assays involving APC/C^{18,20}. In an 8- μ l reaction volume, 0.5 μ l of 10 μ M E1 (625 nM final), 1 μ l of 10 μ M UBE2C (1.25 μ M final), 1 μ l of 10 μ M UBE2S (1.25 μ M final), 1 μ l of 10 mg/ml ubiquitin (1.25 mg/ml final), 1 μ l of 8 $\times$ ubiquitylation assay buffer (250 mM Tris 7.5, 500 mM NaCl, and 100 mM MgCl₂), 1 μ l of 100 mM DTT, 1.5 μ l of energy mix (150 mM creatine phosphate, 20 mM ATP, 20 mM MgCl₂, 2 mM EGTA, pH to 7.5 with KOH), and 1 μ l of recombinant human EMI1 or Geminin (ab86447, Abcam) was mixed with 5 μ l of APC/C resin. Reactions were carried out at 30°C for 1 hour unless otherwise indicated and shaking at 1,000 RPM. Reactions were stopped by the addition of SDS sample loading buffer, resolved on a 12% SDS-acrylamide gel. For experiments comparing *in vitro* reactions using different concentrations of Emi1 on the same gel (Fig. 4a,b and Extended Data Fig. 8a,c), equal amounts of Emi1 were loaded onto the gel to allow for direct comparisons of band intensity using a single exposure.

Immunoblotting

Cells were plated in equal number on 6cm dishes. At the time of harvesting, cells were washed in ice cold PBS. Protein extracts were produced by lysing cells directly in boiling 1 $\times$ sample buffer and resolved by 4-20% gradient SDS-PAGE. Samples were subjected to immunoblotting with Emi1 antibodies (Fig. 2a,b and Extended Data Fig. 4c: 3D2D6, Invitrogen or Fig 2e, 4a,b, Extended Data Fig. 5h,j and 8a-c: #38-5000, Zymed) at 1:1000, cyclin A2 antibodies (sc-751, Santa Cruz Biotechnology) at 1:500, Geminin antibodies (FL-209, Santa Cruz Biotechnology), and GAPDH antibodies (D16H11, Cell Signaling Technology) at 1:1000. Immunoreactive species were visualized by chemiluminescent detection of horseradish peroxidase-conjugated secondary antibodies (7074 or 7076, Cell Signaling Technology).

APC/C-Emi1 modeling

Model calculation of the dual Emi1-APC/C^{Cdh1} substrate and inhibitor switch were performed using a MATLAB program using the ODE45 solver for ordinary differential equations. The model calculates changes of the concentrations of non-ubiquitinated Emi1

and of Emi1 species with 1 to 5 ubiquitins conjugated. The model uses the following parameters and assumptions: First, since at least 4 ubiquitins have to be conjugated to a substrate for its effective degradation²², and since substrates with 5 or more ubiquitins are more effectively degraded, we made a simplifying assumption that all Emi1 species that have more than 5 conjugated ubiquitins are degraded in cells. We did not distinguish possible rate variations if more than one mono-ubiquitins might be added or if different chains might be elongated in Emi1. Second, we made the simplifying assumption that only unbound Emi1 is ubiquitinated and that ubiquitinated Emi1 does not bind to the inhibitory site. This leads to a slower mono-ubiquitination of Emi1²⁶ since most Emi1 is bound to APC/C^{Cdh1} in G1. Third, inhibition of APC/C^{Cdh1} by Emi1 is modeled by using as APC/C^{Cdh1} activity the concentration of APC/C^{Cdh1} without bound inhibitory Emi1 (Total APC/C^{Cdh1} concentration was assumed to be 500 nM, the affinity of the Emi1- APC/C^{Cdh1} inhibitory binding interaction was assumed to be 5 nM and the maximal synthesis of Emi1 from Emi1 mRNA was assumed to be 10 nM/min. The latter rate increases with a time constant of 500 min from 0 nM/min). Fourth, based on our *in vitro* results, we included in our model that ubiquitination of Emi1 is not processive, assuming faster binding and dissociation of Emi1 substrates ($\gamma+$ and $\gamma-$ terms) compared to ubiquitin chain elongation of the bound Emi1 substrate. When simulated, this model recapitulates the fast inactivation of APC/C that we observed in cells (< 1 hour) when we selected an overall rate at which the chain of ubiquitinated Emi1 is elongated as 3.7/min (binding plus elongation) and a term, β , the rate at which individual ubiquitins are removed from poly-ubiquitinated Emi1, is selected to be 2/min (most relevant for the model is the ratio of the two parameters). Finally, we modeled inhibition by CDK2 as a normalized factor, r , that is 1 at the start and inactivates APC/C^{Cdh1} using a time constant of 600 min and cooperativity of 3. The Matlab models that run the simulations are publically available on Github (see code availability).

Code availability

The code for the image analysis pipeline is available at https://github.com/scappell/Cell_tracking. The code for the model of APC/C^{Cdh1} inactivation is available at https://github.com/scappell/APC_model. Additional modified scripts can be accessed upon request.

Data availability

Source Data are available from the corresponding authors upon reasonable request.

Extended Data

Extended Data Figure 1. Single-cell APC/C^{Cdh1} data used to determine hysteresis curves
a, Experimental setup for Fig. 1b,d. and Extended Data Fig. 1b. Asynchronous MCF10A cells were tracked and imaged for approximately 18 hours and then treated with either DMSO or the indicated concentration of CDK1/2 inhibitor. Two subpopulations were analyzed: cells in G1 phase (drug added while APC/C^{Cdh1} On) or cells in S/G2 phase (Drug added while APC/C^{Cdh1} Off). **b**, Single-cell traces of APC/C activity in MCF10A cells

treated with DMSO or 30 μ M CDK1/2 either in G1 phase (top and middle) or S phase (bottom). Traces were colored red if APC/C^{Cdh1} ended up inactive and traces were colored black if APC/C^{Cdh1} ended up active 5 hours after addition of the drug. ($n=200$, 135, and 197 cells). Right: histogram of APC/C activity at the last frame of the movie for each cell. Inset: Pie chart of the percent of cells with APC/C still active at the end of the movie. **c**, Scatter plot of CDK2 activity at the time of drug spike plotted against the CDK2 activity from the same cell 1 hour later. Cells at all phases of the cell cycle were included. **d**, Top: representative APC/C activity traces for MCF10A cells in G1 phase at time of the indicated treatment. Bottom: representative APC/C activity traces for MCF10A cells in S/G2 phase at time of the indicated treatment. Traces were colored red if APC/C^{Cdh1} was inactive and traces were colored black if APC/C^{Cdh1} was still active 5 hours after addition of the drug. Pie chart shows the percent of cells with APC/C^{Cdh1} still active, which was used to make the dose-response curve in Fig. 1b. **e**, Table of the EC50 for the dose response curves in Fig. 1b,d $\pm$ SEM. $n=3$ independent experiments. Note: doses above 30 μ M CDK1/2 inhibitor killed the cells making it impossible to calculate an EC50 above 30 μ M. **f**, MCF10A Cells were treated with 10 μ M CDK1/2 inhibitor and binned by the APC/C^{Cdh1} activity at the time of drug addition. The percent of cells in each bin with APC/C^{Cdh1} activity on 2.5 hours after drug addition was calculated. Data were analyzed by nonlinear regression (sigmoidal dose-response, variable slope). Error bars are standard deviation. from 2 independent experiments. **g**, Scatter plot of APC/C activity at the time of drug spike plotted against the APC/C activity from the same cell 2.5 hours later. Vertical dashed line represents the 70% inactive threshold. Horizontal line indicates when APC/C activity is below 0.1 and considered to be “Off”.

Extended Data Figure 2. siRNA validation experiments

HeLa (**a,b,e**) or U2OS (**c,d**) cells were treated with the indicated siRNA for 24 hrs prior to fixation and immunostaining. Scale bar is 20 μ m. Single-cell quantification of immunofluorescence data was used to generate the histograms. **a**, n=7,152 cells for siControl and n=4,051 cells for siCyclin A2. **b**, n=7,772 cells for siControl and n=3,916 cells for siSkp2. **c**, n=7,003 cells for siControl and n=8,589 cells for siCyclin F. **d**, n=9,393 cells for siControl and n=7,871 cells for siEmi1. **e**, n=64,574 cells for siControl and n=53,338 cells for siCyclin E1,2.

Extended Data Figure 3. Emi1 is responsible for rapid APC/C^{Cdh1} inactivation in multiple cell lines

a,b, Single-cell traces of APC/C activity computationally aligned to when APC/C^{Cdh1} first starts to inactivate in cells treated with the indicated siRNA 6 hours prior to the start of imaging. Black line is median APC/C activity trace. Right: Median APC/C activity trace $\pm$ SD. HeLa cells, n=129/602, 126/384, 124/399, 120/228, 138/400 cells per condition respectively (representative traces plotted/all traces used to calculate median) (**a**), MCF10A cells, n=139/400, 130/400, 134/400 cells per condition respectively (**b**). **c**, Single-cell traces of APC/C activity computationally aligned to mitosis in U2OS cells treated with the indicated siRNA 6 hours prior to the start of imaging. Black line is median APC/C activity trace. Traces in U2OS cells had to be aligned by mitosis since APC/C^{Cdh1} did not inactivate when cells were treated with siRNA targeting Emi1. n=200 representative cells per condition out of a total of 2068, 1531, 875 cells respectively. **d**, Single-cell traces of APC/C activity in

MCF10A cells treated Emi1 siRNA. At the indicated time in S phase, 1 μ M CDK1/2 inhibitor was spiked in. Traces were colored red if APC/C^{Cdh1} stayed inactive and traces were colored black if APC/C^{Cdh1} turned back on 5 hours after addition of the drug. n=80. Inset: Pie chart of the percent of cells that turned APC/C back on at the end of the movie. **e**, Schematic diagram of a proposed bistable switch involving Cyclin E1,2/CDK2, APC/C, and Emi1.

Extended Data Figure 4. Measuring protein and mRNA dynamics from fixed cells

a, HeLa cells were synchronized with thymidine followed by nocodazole and mitotic shake-off (see Fig. 2a). After shake-off, cells were immediately plated in a 96-well plate for live-cell imaging. Top: single-cell traces of APC/C-degron levels. Black line is the median APC/C-degron level. Bottom: single-cell APC/C activity traces. Dashed-line indicates time when APC/C^{Cdh1} first starts to inactivate. n=200 representative cells out of a total of 3190 cells used to calculate the median. **b**, Asynchronous HeLa cells were exposed to 10 μ M EdU for 15 minutes, fixed, and then stained for both Emi1 and EdU incorporation. Top: density scatter plot of DNA content vs EdU levels, used to gate cells into specific cell cycle phases. (e.g. G1 cells have 2n DNA and low EdU, S phase cells have greater than 2n DNA and high EdU, etc). Cells were designated as Early S phase if they had 2n DNA and sub-maximal EdU, indicating they were in S phase for less than 15 minutes at the time of fixation. Bottom: density scatter plot of DNA content vs Emi1 levels. Since we measured EdU and Emi1 in the exact same cells, cell cycle phase gates were applied to the Emi1 scatter plot. Each cell was colored based on the cell cycle phase that cell was in. Large dot with black boarder marks the median values for each gated population. The single-component histograms for Emi1 levels in each cell cycle phase are shown on the right. **c**, HeLa Cells were treated as in Fig. 2a, but 2 hours after release, cells were treated with either DMSO or 10 μ M MG132. Lysates were collected 2 hours later for a total of 4 hours after release. Image is representative of n=3 independent experiments. **d**, Schematic diagram of the regulation of 3 key G1 and S phase genes, Geminin, Emi1, and Cyclin E1 and E2, by both transcription and ubiquitination-mediated degradation. **e**, HeLa cells were imaged and tracked for 20 hrs, and the time each cell went through anaphase relative to the end of the movie was recorded. Cells were then fixed and stained with single molecule mRNA FISH probes for the indicated genes (top), or with antibodies against the indicated proteins (bottom). Data are a scatter plot of the single-cell mRNA or protein levels as a function as the time relative to mitosis. Solid line represents the median levels from 1 hour bins. n=2 independent experiments. **f**, cells treated as in (e), but prior to fixation cells were incubated with 100 μ M EdU for 5 min. Dashed line indicates the first bin when EdU incorporation goes above background, indicating cells have entered S phase. Data is from one representative experiment Right: Single-cell data was binned in 1 hour intervals and the median $\pm$ S.E.M. was plotted. n=6 replicates.

Extended Data Figure 5. Emi1 accumulates coincident with APC^{Cdh1} inactivation

a-c, Normalized mRNA and protein levels for the indicated genes. Data from mRNA and protein are plotted on the same x-axis to allow comparison of the kinetics. S phase start determined by EdU incorporation (see Extended Data Fig. 4e). Data is median value for each bin. n=2 independent experiments. **d**, Density scatter plot of Emi1 protein levels and Geminin protein levels measured in the same single cells. Single-component histograms shown as insets. **e**, Density scatter plot of Cyclin E1 protein levels and Geminin protein levels measured in the same single cells. Single-component histograms shown as insets. **f**,

HeLa cells were treated with either control siRNA (black line) or Cdh1 siRNA (colored line) and then imaged and tracked for 20 hrs. The time each cell went through anaphase relative to the end of the movie was recorded. Cells were then fixed and immunofluorescence was performed with antibodies to the indicated proteins. The time to reach half-maximal is indicated with a dashed line ($t_{1/2}$). Data are median values $\pm$ S.E.M and best fit line was calculated using nonlinear regression. Data was normalized to maximum value. $n=8$ independent experiments. **g**, Bar graph of the difference in $t_{1/2}$ times in cells treated with Control siRNA and Cdh1 siRNA from (**j**). Individual data points are shown as dots. Error bars are S.E.M. p values from one-way ANOVA with Tukey multiple comparisons test. $n=8$ independent experiments. **h,i**, *in vitro* ubiquitination time-course using immuno-purified APC/C and either recombinant Emi1 (**h**) or Geminin (**i**). Lower panel is a lighter exposure showing the un-modified substrate. $n=1$ experiment. **j**, *in vitro* ubiquitination assay using immuno-purified APC/C^{Cdh1}, recombinant Emi1, and the indicated ubiquitin mutant. E2C/S, E2 enzymes Ube2C and Ube2S. $n=1$ experiment.

Extended Data Figure 6. Emi1 and Geminin protein levels are tightly correlated during the cell cycle

a, Model diagram of APC/C^{Cdh1} architecture. For simplification, the substrate binding region comprised primarily of Cdh1 and APC10 is designated as site 1, and the E2 ubiquitin conjugating enzymes Ubch10 and Ube2s are designated as site 2. **c-d**, Regulation of APC/C by two Emi1 mutants. Substrates are recruited to APC/C through a substrate binding region (site 1). Ubiquitin is transferred to the substrate by UBCH10 and UBE2S (site 2). Sub, substrate; Ub, ubiquitin. **e**, Signaling diagram of the G1 cell-cycle program. Shows inhibiting CDK4/6 activity with a small molecule inhibitor will prevent APC/C^{Cdh1} inactivation. **f**, MCF10A cells expressing either empty vector or various Emi1 variants were treated with DMSO. Single-cell APC/C activity traces were computationally aligned to the time of mitosis. DMSO treatment administered before mitosis as indicated by arrow. $N=120$ cells for each condition. **g**, Median levels of APC/C degraon (red) or wild-type Emi1

(Emi1^{WT}; green) stably expressed in the same MCF10A cells. n=330 cells. **h**, Median expression levels of ectopically expressed Emi1 mutants tagged with mCitrine. Measurements for each single-cell were the maximum fluorescence detected during each cell cycle. Single-cell data was then averaged for each replicate. Individual data points are shown as dots. Data are median values $\pm$ S.D. n=3 independent experiments. **i**, Left: median APC/C degron levels measured over 24 hours in MCF10A cells. Arrows indicate mitosis (mit). n=88,125 cells. Right: median levels of APC/C degron (red) or Emi1^{C401S} (green) stably expressed in the same cells. n=3,618 cells. Inset diagram shows that Emi1^{C401S} is a substrate of APC/C^{Cdh1} and not an inhibitor. **j**, Single-cell traces of APC/C degron (red) or Emi1^{C401S} (green) levels in MCF10A cells. Six representative cells from (**i**) are shown. **k**, Density scatter plot of APC/C degron levels versus Emi1^{C401S} levels measured in the same cells. **l**, Percent of cells from Fig. 3b-e that inactivated their APC/C^{Cdh1} at least 5 hours after mitosis. n=3 biological repeats. Individual data points are shown as dots. Error bars are S.D.

Extended Data Figure 7. Mathematical model for an APC/C-Emi1 dual negative feedback loop
a, Differential equation of the regulation of Emi1 protein concentration. Constant synthesis (S) is assumed for the steady state model. The model includes reversible low affinity binding of Emi1 to the substrate binding site ($\gamma+/-$) and inhibitory binding that blocks ubiquitin chain elongation, but not mono-ubiquitination. Emi1 ubiquitination is assumed to be distributive, requiring 5 ubiquitin elongations to become fully ubiquitinated for effective degradation. APC/C^{Cdh1} activity is inhibited by CDK2 activity or by Emi1 binding to the inhibitory site, allowing Emi1 to be subsequently de-ubiquitinated and stabilized. **b**, A

single-cell APC/C activity and CDK2 activity trace. Each steady-state from Fig. 3g is marked. At the indicated time 3 μ M CDK1/2 inhibitor was added, but APC/C activity remains off, as predicted in Fig. 3g.

Extended Data Figure 8. Emi1 is an APC/C substrate at low concentrations but an inhibitor of ubiquitin chain elongation at high concentrations

a, *in vitro* ubiquitination reaction of recombinant Geminin using immuno-purified APC/C and a range of concentrations of recombinant Emi1. Non-specific bands are marked with (*). n=1 experiment. **b**, *in vitro* ubiquitination time-course using immuno-purified APC/C and 2944 nM recombinant Emi1. Only mono-ubiquitinated Emi1 was observed after 30 minutes. n=1 experiment. **c**, *in vitro* ubiquitination assay using immuno-purified APC/C^{Cdc20} from cells arrested in mitosis and either 23 nM or 2944 nM recombinant Emi1. After the *in vitro*

reaction, equal amounts of Emi1 were loaded onto the gel for western blotting. $n=1$ experiment. **d**, Quantification of fully-ubiquitinated Emi1 from Fig. 4b using densitometry. Data is plotted as the percent of total Emi1. Best-fit line was determined using the Hill equation. Dashed lines indicate an EC50 of ~ 175 nM. **e**, Steady-state analysis of Emi1 levels as a function of the degradation and synthesis rate. Assuming constant Emi1 synthesis (dashed lines), there are two stable steady states (red and blue dots). The system can transition to a mono-stable state with APC/C “Off” and high Emi1 levels by increasing the Emi1 synthesis rate above a threshold level (red arrow). Conversely, the system can transition to a different mono-stable state with APC/C “On” and low Emi1 levels by decreasing the Emi1 synthesis rate (blue arrow). This system also exhibits hysteresis, since the occupied steady-state depends on the previously occupied state (red dots vs blue dots).

Extended Data Figure 9. Cyclin E is dispensable for APC/C^{Cdh1} inactivation in cells with constitutive active E2F

a,b, Single-cell traces of APC/C activity computationally aligned to mitosis in either HeLa (n=200 cells per condition out of a total of 1284, 434, 462 cells respectively used to calculate the median and standard deviation) (**a**) or U2OS (n=200 cells per condition out of a total of 551, 300, 271 cells respectively used to calculate the median and standard deviation) (**b**) cells treated with the indicated siRNA 6 hours prior to the start of imaging. Black line is median APC/C activity trace. Bottom: Median APC/C activity trace $\pm$ SD. **c**, U2OS cells

were treated with either Control, Cyclin E1 and E2, or Emi1 siRNA for 4 hours prior to the start of time-lapse imaging. APC/C and CDK2 activity were measured in the same cells, and traces were aligned to mitosis. Data is median activity trace $\pm$ SEM. $n=200$ cells. **d**, Time between anaphase and when relative APC/C activity reaches below 0.2 was calculated for each cell from Fig. 4f. Times from all cells were averaged for each replicate. Individual data points are shown as dots. Data are median times $\pm$ S.E.M. p values from one-way ANOVA with Tukey multiple comparisons test. $n=3$ independent experiments. **e**, Single-cell APC/ C^{Cdh1} inactivation slope at the G1/S transition for each cell from Fig. 4f. Slopes from all cells were averaged for each replicate. Data are median APC/ C^{Cdh1} inactivation slopes $\pm$ S.E.M. p values from one-way ANOVA with Tukey multiple comparisons test. Individual data points are shown as dots. $n=3$ independent experiments. **f**, Histogram of the time between CDK2 activation and APC/ C^{Cdh1} inactivation in HeLa cells treated with the indicated siRNA. Values above zero indicate CDK2 turns On before APC/ C^{Cdh1} turns Off, and values below zero indicate APC/ C^{Cdh1} turns Off before CDK2 turns On.

Extended Data Figure 10. APC/C^{Cdh1} inactivates prior to CDK2 activation in cells with Cyclin E1,2 knocked down

a,b, Single-cell traces of APC/C (red) and CDK2 (green) activity in HeLa cells treated with either control siRNA (**a**) or cyclin E1 and E2 siRNA (**b**). Ten representative cells are shown for each condition. Black arrow, time of mitosis; green dashed line, time CDK2 activity first starts to rise; red dashed line, time APC/C activity first starts to inactivate. **c**, Phase-plane diagram of median CDK2 and APC/C activity measured simultaneously in the same U2OS cells over time, starting in Mitosis or G0 phase and progressing to S phase. n=270 cells per

condition. **d**, Scheme of the regulation of APC/C^{Cdh1} inactivation in the presence (Wild-type) or absence of Cyclin E. Cells can bypass the requirement for Cyclin E1,2/CDK2 in G1 phase by upregulating Emi1 mRNA. HeLa and USO2 cells bypass Cyclin E1,2/CDK2 by having constitutively active E2F1-3.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 1. Emi1 conveys hysteresis to APC/C^{Cdh1} inactivation

a, Requirements for a bistable switch. **b**, Dose response curve for the two subpopulations of cells treated with CDK1/2 inhibitor. Data were analyzed by nonlinear regression (sigmoidal dose-response, variable slope). $n=3$ independent experiments, errorbars are S.E.M. **c**, APC/C activity traces aligned to when APC/C^{Cdh1} inactivates in HeLa cells. Top: Median and single-cell traces of APC/C activity in control cells. Bottom: Median APC/C activity traces. Error bars are SD ($n=602, 384, 399, 228, 400$ cells respectively). **d**, Same experimental setup as **(b)** but MCF10A cells were first treated with Emi1 siRNA. Data were analyzed by nonlinear regression (sigmoidal dose-response, variable slope). $n=3$ independent experiments, errorbars are S.E.M.

Figure 2. The APC/C^{Cdh1} inhibitor Emi1 is also an APC/C^{Cdh1} substrate degraded during G1 and building up at G1/S

a, HeLa cells were synchronized with nocodazole and mitotic shake-off. Image is representative of 3 independent experiments. **b**, HeLa Cells were treated as in (**a**), but 1.5hrs after release, cells were treated with either DMSO or 30 μ M proTAME, a small-molecule APC/C inhibitor. Lysates were collected 4 hours after release. Image is representative of 2 independent experiments. **c,d**, Live-cell imaging was combined with fixed cell analysis to measure mRNA (**c**) or protein and EdU (**d**) levels over time since mitosis. Median levels from 1 hour bins were plotted. $n=2$ independent experiments. **e**, *in vitro* ubiquitination assay using APC/C from cells in G1 phase and recombinant Emi1 (amino acids 299-447). E2C/S, E2 enzymes Ube2C and Ube2S. Image is representative of 2 independent experiments.

Figure 3. APC/C^{Cdh1} inactivation is a bistable switch driven by dual-negative feedback
a, Emi1 structural elements. D-Box, APC/C degen sequence (RxxL); L, linker; ZBR, zinc-binding region; T, tail. **b-e**, MCF10A cells expressing either empty vector or various Emi1 mutants were treated with DMSO (black lines; **b**) or 1 μ M CDK4/6 inhibitor (red lines; **b-e**). Drug treatment administered before mitosis as indicated by arrow. $n=120$ cells for each condition. Data representative of $n=3$ independent experiments. **f**, Two-binding site model for regulation of the APC/C by Emi1. Emi1 can bind either the substrate binding region (1) via its APC/C degen motif or the E2 binding site (2) via its ZBR and C-terminal tail. **g**, Three steady-state solutions for the mathematical model (see Extended Data Fig. 7a).

Figure 4. Cyclin E/CDK2 activity and Emi1 mRNA synthesis synergistically trigger the Emi1-APC/C^{Cdh1} inactivation switch and cell cycle commitment

a, *in vitro* ubiquitination assay using immuno-purified APC/C^{Cdh1} and recombinant Emi1. After the *in vitro* reaction, equal amounts of Emi1 were loaded onto the gel for western blotting. Image representative of *n*=2 independent experiments. **b**, same as (**a**) but using a range of concentrations of recombinant Emi1. Image representative of *n*=2 independent experiments. **c**, Schematic showing at low Emi1 concentrations, Emi1 can be poly-ubiquitinated but at high Emi1 concentrations, Emi1 inhibits E2 activity and chain elongation. **d**, Schematic diagram of a bistable switch involving Cyclin E1,2/CDK2 and a dual-negative feedback loop between APC/C^{Cdh1} and Emi1. **e**, Simulated changes in APC/C^{Cdh1} activity in G1 and S phase. **f**, HeLa cells treated with the indicated siRNA. APC/C and CDK2 activity were measured in the same cells, and traces were aligned to mitosis. Data is median activity trace $\pm$ SEM. **g**, Phase-plane diagram of median CDK2 and APC/C activity measured simultaneously in the same HeLa cells. *n*=270 cells per condition. **h**, APC/C^{Cdh1} inactivation at the G1/S transition is a one way-switch, triggered by CDK2 activity and made irreversible by a dual-negative feedback mechanism with Emi1.