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A pathway in quiescent cells that controls p27<sup>Kip1</sup> stability, subcellular localization, and tumor suppression

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We have created two knock-in mouse models to study the mechanisms that regulate p27 in normal cells and cause misregulation of p27 in tumors: p27<sup>S10A</sup>, in which Ser10 is mutated to Ala; and p27<sup>CK−</sup>, in which point mutations abrogate the ability of p27 to bind cyclins and CDKs. These two mutant alleles identify steps in a pathway that controls the proteasomal degradation of p27 uniquely in quiescent cells: Dephosphorylation of p27 on Ser10 inhibits p27 nuclear export and promotes its assembly into cyclin–CDK complexes, which is, in turn, necessary for p27 turnover. We further show that Ras-dependent lung tumorigenesis is associated with increased phosphorylation on Ser10 and cytoplasmic mislocalization of p27. Indeed, we find that p27<sup>S10A</sup> is refractory to Ras-induced cytoplasmic translocation and that p27<sup>S10A</sup> mice are tumor resistant. Thus, phosphorylation of p27 on Ser10 is an important event in the regulation of the tumor suppressor function of p27.

[Keywords: p27<sup>Kip1</sup>; cyclin; CDK; tumor suppressor; tumorigenesis; Ras; lung cancer]

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The transitions from one phase of the cell cycle to the next are driven by the activities of various cyclin/cyclin-dependent kinase [CDK] complexes (Sherr and Roberts 2004). One level of regulation of cyclin–CDK complexes is provided by cyclin–CDK inhibitors [CKIs], which modulate their activity [Sherr and Roberts 2001]. The CKI p27<sup>Kip1</sup> (p27) regulates the transition from G0, through G1, into S phase [Nourse et al. 1994; Polyak et al. 1994a,b; Toyoshima and Hunter 1994; Coats et al. 1996]. The importance of p27 as a regulator of cell proliferation is illustrated by the phenotype of the p27-null mice, which display a marked increase in body size and multiorgan hyperplasia [Fero et al. 1996; Kiyokawa et al. 1996; Nakayama et al. 1996]. Moreover, p27 is also a tumor suppressor in mice, as animals lacking p27 are predisposed to both spontaneous and induced tumorigenesis [Fero et al. 1996, 1998; Kiyokawa et al. 1996; Nakayama et al. 1996]. p27 is haploinsufficient for tumor suppression, and loss of a single p27 allele is sufficient to increase tumor incidence [Fero et al. 1998]. p27 may also act as a tumor suppressor in humans, as decreased expression of nuclear p27 is commonly observed in various human cancers and is a significant prognostic marker for clinical outcome [Slingerland and Pagano 2000; Philipp-Staheli et al. 2001].

p27 protein levels are elevated in quiescent cells and decrease following mitogen stimulation and entry into G1 [Coats et al. 1996]. p27 levels can be affected by transcriptional and translational pathways, but the major mechanisms for p27 regulation are thought to be post-translational proteolytic degradation [Pagano et al. 1995; Hengst and Reed 1996]. Best understood is the proteolytic pathway that causes p27 degradation in late G1- and S-phase cells. Cdk2 first becomes active in late G1 and phosphorylates p27 on Thr187. This creates a recognition site for the Skp2–SCF E3 ubiquitin ligase, which ubiquitinates p27, thereby promoting its degradation by the proteasome [Sheaff et al. 1997; Vlach et al. 1997; Carrano et al. 1999; Montagnoli et al. 1999, Sutterluty et al. 1999; Tsvetkov et al. 1999]. In addition, a stable interaction with cyclin–CDK complexes is important for p27 turnover. Thus, a p27<sup>T187A</sup> mutant that cannot bind to cyclins and CDKs was still a substrate for phosphorylation [Vlach et al. 1997; Montagnoli et al. 1999]. Supporting this idea, others have found a noncatalytic requirement for cyclin A–CDK2 complexes for p27 turnover [Zhu et al. 2004].

However, analysis of a p27<sup>T187A</sup> knock-in mouse model revealed that this phosphorylation-resistant form
of p27 was stabilized in S phase but was still degraded normally during G0 and G1, suggesting that other proteolytic pathways must exist [Malek et al. 2001]. In fact, a second ubiquitin–proteasome pathway was recently described that causes the degradation of p27 in the cytoplasm during the G1 phase, mediated by the KPC1 and KPC2 E3 ubiquitin ligases [Kamura et al. 2004]. CRM1–KPC2–mediated nuclear export of p27 was necessary for KPC1–KPC2–mediated degradation, but it remains unknown whether a direct modification of p27, such as phosphorylation, is also required [Kamura et al. 2004]. Moreover, it is unlikely that all p27 translocates into the cytosol in G1 cells [Rodier et al. 2001; Boehm et al. 2002; Ishida et al. 2002; Connor et al. 2003; McAllister et al. 2003; Kamura et al. 2004; Shin et al. 2005]. Indeed, when p27 is confined to the nucleus either by pharmacological or genetic inhibition of its nuclear export, its stability decreases [Rodier et al. 2001]. It is therefore probable that another turnover pathway acts during G1 in the nucleus. Whether it involves a Skp2–containing ubiquitin ligase that functions independently of Thr187 phosphorylation remains to be clarified [Malek et al. 2001].

The main phosphorylation site on p27 is Ser10, which has been found to regulate both its subcellular localization and stability [Ishida et al. 2000; Rodier et al. 2001]. Phosphorylation at this site, or mutation of Ser10 to a phospho-mimetic residue [Asp or Glu], correlates with increased stability of p27, which is consistent with this being a modification primarily detected in nondividing cells [Ishida et al. 2000; Rodier et al. 2001; Boehm et al. 2002; Deng et al. 2004]. The mechanism by which Ser10 phosphorylation stabilizes p27 in the G0 nucleus is not understood. Although Ser10 phosphorylation is associated with p27 stabilization in quiescent cells, it has also been linked to p27 degradation in G1. Several groups reported that Ser10 phosphorylation of p27 was required for its export from the nucleus, possibly by mediating the binding of p27 to the exportin CRM1 [Rodier et al. 2001; Boehm et al. 2002; Ishida et al. 2002; Connor et al. 2003; McAllister et al. 2003; Shin et al. 2005]. This may trigger p27 degradation in the cytoplasm of G1 cells, possibly by the KPC pathway. Several kinases are capable of phosphorylating p27 on Ser10: Mirk/dyrk1B phosphorylates and stabilizes p27 in quiescent cells [Deng et al. 2004], while hKIS phosphorylates this site at the G0–G1 transition following mitogen stimulation, and this is accompanied by translocation of p27 in the cytoplasm [Boehm et al. 2002]. Other kinases, such as PKB/Akt and ERK2, were also reported to phosphorylate p27 on Ser10, at least in vitro [Ishida et al. 2000; Fujita et al. 2002].

The regulation of p27 stability and subcellular localization, which are critical for the control of p27 function, are complex phenomena modulated by phosphorylation and protein–protein interactions. In order to dissect the importance of these events in the regulation of p27 stability, localization, and function in vivo, both in physiological and pathological settings, we constructed knock-in mice harboring specific point mutations in the p27 gene. In the experiments described below, we focused on the unexpected interactions between two pathways that have been proposed to regulate p27 stability: its phosphorylation on Ser10 and its assembly into complexes with cyclins and CDKs. In quiescent cells, phosphorylation of p27 on Ser10 regulated its subcellular localization and its association with cyclin–CDK complexes, and this was critical for controlling p27 stability. Not only do we show that this constitutes a new turnover pathway for p27 that is uniquely active in quiescent cells but also that it represents a key mechanism for regulating the tumor suppressor function of p27.

Results

Generation of p27S10A and p27CK− mice

To investigate the pathways that control p27 turnover, we engineered knock-in mice in which the p27 gene was mutated at specific sites [Fig. 1A]: In one strain, the p27S10A allele carried a mutation of Ser10, the main phosphorylation site on p27, to Ala [Ishida et al. 2000]; in the other strain, the p27CK− allele carried the mutations Arg30 to Ala and Leu32 to Ala in the cyclin-binding domain of p27, and Phe62 to Ala and Phe64 to Ala in the CDK-binding domain of p27. The p27CK− mutation has previously been reported to abolish p27 binding to cyclins and CDKs [Vlach et al. 1997], and this was confirmed by us [see below]. Precise replacement of the endogenous p27 gene with the mutant alleles was confirmed by Southern blot (data not shown), genomic PCR [Fig. 1B], and DNA sequencing [data not shown]. The progeny from p27+/S10A × p27+/S10A or p27+/− × p27+/− crosses were recovered with approximately the expected Mendelian ratios [Table 1]. Below, we describe how mutation of either Ser10 or the cyclin–CDK-binding domain affects p27 function and regulation both in vivo and in cell culture, and then consider how these elements interact with each other to control p27 stability during the cell cycle.

The p27CK− mutation

Mice lacking p27 have an increased growth rate and mature body size because of the occurrence of extra cell divisions in essentially all lineages, presumably a consequence of the lack of cyclin–CDK regulation by p27 [Fero et al. 1996; Kiyokawa et al. 1996; Nakayama et al. 1996]. This is supported by the phenotype of the p27CK− mice, which were similarly affected. We weighed cohorts of 30 animals per genotype weekly for 3 mo after weaning. Mice homozygous for p27+ or p27− were larger than their wild-type littermates, and p27−/+ had an intermediate phenotype [Fig. 1C, upper panels]. Moreover, certain organs such as the spleen and thymus are differentially enlarged in p27null mice, and those organs were also disproportionately larger in the p27CK− background [Fig. 1D].

We confirmed that the p27CK− protein did not associate with cyclins or CDKs. The presence of p27 bound to cyclins A, E1, or D1 was determined by immunoprecipi-
tation (IP) from primary mouse embryonic fibroblast (MEF) extracts. Whereas wild-type p27 associated with all three cyclins, the p27\textsuperscript{CK\textendash} protein was associated with none (Figs. 1E, 5D–F; Supplementary Fig. 5). Thus, both phenotypic and molecular observations confirm that the p27\textsuperscript{CK\textendash} mutation prevents the interaction of p27 with cyclins and CDKs. We are continuing to characterize the phenotype of p27 CK\textsuperscript{\textendash} mice in order to address whether...
there are cyclin–CDK-independent functions of p27 (A. Besson and J.M. Roberts, in prep.).

We next examined the effect of the p27<sup>CK−</sup> mutation on the abundance of the p27 protein in vivo. In all tissues tested (spleen, uterus, brain, liver, thymus, and lung), p27<sup>CK−</sup> levels were increased [Fig. 2A, data not shown]. Similar results were obtained by immunohistochemistry, which revealed a stronger signal for p27<sup>CK−</sup> than wild-type p27 in the brain (Supplementary Fig. 1A). p27 levels in primary MEFs derived from wild-type and p27<sup>CK−</sup> embryos were similar to those observed in tissues, with p27<sup>CK−</sup> being elevated in all conditions (exponentially growing, serum starved, or confluent) (Fig. 2C). To further explore the effect of the p27<sup>CK−</sup> mutation on p27 abundance, we monitored primary MEFs following serum stimulation over a 24-h time course both for p27 levels (Fig. 2D) and progression through the cell cycle by flow cytometry (Fig. 2E). As previously reported, wild-type p27 levels were elevated in serum-starved cells, progressively decreased during G1 [between 3 h and 15 h], and remained low in S phase [between 18 h and 24 h]. In contrast, p27<sup>CK−</sup> protein levels were strikingly different from wild-type p27. As expected, the relative abundance of the p27<sup>CK−</sup> protein was increased in S-phase cells, consistent with the requirement for phosphorylation of p27 by cyclin E–CDK2 and binding to cyclin–CDK for ubiquitination by the SCF-Skp2 E3 ligase (Sheaff et al. 1997; Vlach et al. 1997; Montagnoli et al. 1999; Zhu et al. 2004). Surprisingly, the p27<sup>CK−</sup> protein was also elevated in quiescence, suggesting that cyclin–CDK–p27 heterotrimer assembly also sets the level of p27 in nongrowing cells. The p27<sup>CK−</sup> protein behaved similarly to wild-type p27 only during entry into G1 following serum stimulation, where both proteins were effectively downregulated (Fig. 2D). Thus, p27 regulation during G1 was independent of its binding to cyclins and CDKs and therefore proceeded by a pathway that was distinct from those that operated in quiescence and S phases.

We directly tested whether the p27<sup>CK−</sup> mutation affected p27 steady-state abundance by changing its rate of degradation. The rate of p27 turnover was measured in exponentially growing (Fig. 3A), serum-starved (Fig. 3B), G1-phase (Fig. 3C), and S-phase (Fig. 3D) primary MEFs using cycloheximide to block protein synthesis. The graph in Figure 3E represents the means and standard deviations for p27 half-lives obtained in multiple experiments at each phase of the cell cycle. In every case, the effect of the p27<sup>CK−</sup> mutation on p27 steady-state protein abundance could be explained by its effect on p27 turnover. The p27<sup>CK−</sup> protein was more stable than wild-type p27 in exponentially growing cells, in S-phase cells, and in quiescent cells [Fig. 3]. Importantly, the rate of p27<sup>CK−</sup> degradation was significantly increased during G1 compared with other phases of the cell cycle [Fig. 3C,E], with the small difference between it and wild-type p27 probably reflecting the fact that not all cells entered the cell cycle following serum stimulation and that this was therefore only an enriched population for G1 cells [Fig. 3E]. We concluded that the formation of heterotrimeric p27–cyclin–CDK complexes is required for p27 degradation both in quiescent cells and during S phase.

**The p27<sup>S10A</sup> mutation**

In confirmation of recent reports, we observed that the abundance of the p27<sup>S10A</sup> protein was decreased in quiescent cells compared with wild-type p27, both in vivo and in vitro (Deng et al. 2004; Kotake et al. 2005). p27<sup>S10A</sup> protein levels were decreased in extracts from all tissues examined [Fig. 2A,B], and this was consistent with p27 immunohistochemistry in the brain of p27<sup>S10A</sup> mice (Supplementary Fig. 1B). As expected, no signal was detected when extracts from the brain or MEFs from p27<sup>S10A</sup> mice were probed with a phospho-Ser10 p27-specific antibody, while both wild-type p27 and p27<sup>CK−</sup> displayed abundant Ser10 phosphorylation [Fig. 2B,C]. Similarly to what was observed in tissues, the p27<sup>S10A</sup> protein was decreased in abundance relative to wild-type p27 in primary MEFs independently of culture conditions [Fig. 2C].

Analyses of synchronized MEFs demonstrated that the p27<sup>S10A</sup> mutation specifically affected p27 abundance and stability in quiescent cells. p27<sup>S10A</sup> protein levels were low in quiescent MEFs, and in response to serum stimulation they were rapidly down-regulated to lower levels than wild-type p27 in G1 and S phases [Fig. 2D].

Effect of the p27<sup>CK−</sup> mutation on p27 abundance could be explained by its effect on p27 turnover. The p27<sup>CK−</sup> protein was more stable than wild-type p27 in exponentially growing cells, in S-phase cells, and in quiescent cells [Fig. 3]. Importantly, the rate of p27<sup>CK−</sup> degradation was significantly increased during G1 compared with other phases of the cell cycle [Fig. 3C,E], with the small difference between it and wild-type p27 probably reflecting the fact that not all cells entered the cell cycle following serum stimulation and that this was therefore only an enriched population for G1 cells [Fig. 3E]. We concluded that the formation of heterotrimeric p27–cyclin–CDK complexes is required for p27 degradation both in quiescent cells and during S phase.

### Table 1. Ratios of p27<sup>+/CK−</sup> × p27<sup>+/CK−</sup> and p27<sup>−/S10A ×</sup> p27<sup>−/S10A</sup> progeny

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Number of animals</th>
<th>Percentage</th>
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<tr>
<td>+/S10A × +/S10A</td>
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<td>100%</td>
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<tr>
<td>+/S10A</td>
<td>73</td>
<td>23.2%</td>
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<tr>
<td>S10A/S10A</td>
<td>163</td>
<td>51.9%</td>
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<tr>
<td>S10A/S10A</td>
<td>78</td>
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<tr>
<td>+/CK− × +/CK−</td>
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<tr>
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<tr>
<td>CK−/CK−</td>
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junction with cycloheximide efficiently blocked wild-type p27, p27\textsuperscript{CK}, and p27\textsuperscript{S10A} turnover (Fig. 4B).

**Mechanism of stabilization of p27\textsuperscript{CK}**

Our data indicate that p27 turnover in G0 requires binding to cyclin/CDK complexes. However, which cyclins and/or CDKs are involved and whether CDK activity is required are unclear. To determine the importance of a direct interaction with cyclin or CDK or both proteins in p27 turnover in G0, we measured the half-lives of p27\textsuperscript{CK} (Arg30 to Ala and Leu32 to Ala) and p27\textsuperscript{S10A} (Phe62 to Ala and Phe64 to Ala). The interaction with cyclin appeared critical for turnover in quiescent cells, as p27\textsuperscript{CK} had a
half-life similar to that of p27 CK−; however, the interaction with CDK may also be important since the half-life of p27 CK− was intermediate between wild-type p27 and p27C− (Fig. 4C). This was consistent with previous data showing that the cyclin-binding domain is more important for the stable assembly of the p27–cyclin–CDK complex than is the CDK-binding domain (Vlach et al. 1997), and was also in agreement with a study on the requirements for p27 ubiquitination in vitro (Ungermannova et al. 2005). Pharmacological CDK inhibitors, roscovitine or olomoucine, had no significant effect on p27 half-life in serum-starved wild-type or p27 S10A primary MEFs (Supplementary Fig. 2), indicating that CDK activity was not required for p27 turnover during quiescence. Next we monitored p27 half-life in MEFs lacking specific cyclins or CDKs (cyclin D1−/−, cyclin D1/D2−/−, cyclin E1/E2−/−, CDK2−/−, and CDK4−/−). The absence of one type of cyclin or CDK did not significantly affect p27 half-life in G0 (Supplementary Fig. 3). Therefore, the cyclin/CDK complexes that are present in quiescent cells may be functionally interchangeable with respect to promoting p27 turnover.

One of the amino acids mutated in p27 CK−, Leu32, is part of a putative nuclear export signal in p27 (between amino acids 32 and 45) and has been proposed to mediate p27 CK’s interaction with CRM1 (Connor et al. 2003). How-
ever, we found that endogenous p27<sup>CK−</sup> could be coimmunoprecipitated with endogenous CRM1, similarly to wild-type p27 (Fig. 4D). This ruled out the possibility that the p27<sup>CK−</sup> mutation increased p27 stability due to a defect in CRM1-mediated export. The increased stability of the p27<sup>K−</sup> protein, which is mutated outside of the proposed CRM1-binding domain, also indicated that the affected turnover pathway did not involve CRM1.

**A pathway for p27 turnover in quiescent cells**

Phosphorylation of p27 on Ser10 and the ability of p27 to interact with cyclins and CDKs both regulate p27 stability in quiescent cells. To determine if these events were part of a single pathway, we tested how the conjunction of the Ser10 and CK− mutations affected p27 stability. We found that the p27<sup>S10A/CK−</sup> double mutant was as stable as the p27<sup>CK−</sup> mutant in quiescent (Fig. 4E) or exponentially growing cells (Supplementary Fig. 4), suggesting that cyclin–CDK assembly functions downstream of, or coincidentally with, Ser10 phosphorylation to control p27 stability.

An attractive hypothesis to explain the above result is that Ser10 phosphorylation may determine the rate of p27 turnover by inhibiting the ability of p27 to associate with cyclins and CDKs. Indeed, we found that the p27<sup>S10A</sup> mutation had a dramatic effect on p27 assembly into cyclin–CDK complexes. We tested this hypothesis by performing IPs using a monoclonal p27 antibody (F8) that recognizes free p27, but not p27 bound to cyclin–CDK complexes (see below). When this antibody was used to immunoprecipitate p27 in serum-starved or exponentially growing primary MEF extracts (Fig. 5A, B, respectively), hardly any p27<sup>S10A</sup> was detected, while wild-type p27 and p27<sup>CK−</sup> were readily detected, consistent with the fact that the latter cannot bind cyclins and CDKs. In addition, no cyclin D was detected in these IPs (Fig. 5A, C). To rule out the possibility that the S10A mutation disrupted the epitope recognized by this antibody, we used p27<sup>−/−</sup> MEFs retrovirally infected with wild-type p27, p27<sup>S10A</sup> (p27<sup>−/−</sup>/pQP-p27<sup>S10A</sup>), and p27<sup>CK−</sup> (p27<sup>−/−</sup>/pQP-p27<sup>CK−</sup>). After selection of infected MEFs, cells were serum starved, treated with CHX, and processed as described in Figure 3. (B) p27 degradation is proteasome-dependent in quiescent cells. Wild-type (WT), p27<sup>S10A</sup> (S10A), and p27<sup>CK−</sup> (CK−) primary MEFs were serum starved for 72 h and treated with 25 µg/mL CHX or with both CHX and 10 µM MG132 for the indicated time. Samples were processed as in Figure 3. (C) Cyclin interaction is critical for p27 turnover in G0. The half-lives of wild-type p27, p27<sup>S10A</sup> (p27<sup>−/−</sup>/pQP-p27<sup>S10A</sup>), and p27<sup>CK−</sup> (p27<sup>−/−</sup>/pQP-p27<sup>CK−</sup>) were determined as in A. (D) p27<sup>CK−</sup> can still interact with CRM1. CRM1 was immunoprecipitated from extracts prepared from primary wild-type or p27<sup>CK−</sup> MEFs either serum starved for 72 h, or starved and stimulated for 4 h with 10% serum. Immunoprecipitates were resolved on SDS-PAGE and probed for p27. A fraction of the extracts was used for Western blotting to determine the amount of CRM1, p27, and Grb2 (loading control) present in these cells. (E) Stabilization of a p27<sup>S10A</sup> mutant that cannot bind cyclins and CDKs (p27<sup>S10A/CK−</sup>) in quiescent cells. p27-null MEFs were retrovirally infected with either wild-type p27 (p27<sup>−/−</sup>/LNSX-p27), p27<sup>CK−</sup> (p27<sup>−/−</sup>/pQP-CK−), or p27<sup>S10A/CK−</sup> (p27<sup>−/−</sup>/LNSX-S10A/CK−), and treated as in A.
rally infected with the p27S10A/CK− double mutant. This mutant was immunoprecipitated very efficiently by the p27 F8 antibody, while very little p27 S10A could be detected (Fig. 5C). In contrast, coimmunoprecipitation of cyclin D1 with either wild-type p27 or p27 S10A was readily detected with a polyclonal anti-p27 antibody (Fig. 5D), while no cyclin D1 could be detected in association with p27CK− or p27S10A/CK−. Thus, most p27 S10A protein is bound to cyclin–CDK complexes, whereas the wild-type p27 protein is found in both the bound and free compartments.

In further support of an effect of the p27S10A mutation on p27’s assembly into cyclin–CDK complexes, coimmunoprecipitation experiments were performed to measure the amount of p27 associated with various cyclins. In serum-starved primary MEFs, the amount of p27S10A coimmunoprecipitated with cyclin D1 was similar to the amount of wild-type p27 associated with cyclin D1, despite the fact that the total amount of p27S10A in the cell was reduced compared with wild-type p27 [Fig. 5E]. Similar results were obtained in exponentially growing MEFs [Fig. 5F]. Likewise, Cyclin A and Cyclin E coimmunoprecipitations yielded similar or higher amounts of p27S10A compared with wild-type p27 [Supplementary Fig. 5], despite the lower overall abundance of p27S10A.

The increased ability of the p27S10A protein to associate with cyclins and CDKs was also supported by the phenotype of the knock-in mice. p27 S10A/S10A homozygous mice expressed the p27 protein at a level similar to mice heterozygous for p27 (p27+/−; i.e., 50% of normal wild-type levels). However, whereas p27+/− mice showed an increased body size and generalized organomegaly (Fero et al. 1996), the p27S10A/S10A homozygous male mice were indistinguishable from wild-type mice in these respects, and the p27S10A/S10A homozygous females had an increased size of only borderline significance [Fig. 1C]. Together, the molecular and phenotypic data show that the p27S10A mutation promotes the association of p27 with cyclin–CDK complexes, and suggest that Ser10 phosphorylation and heterotrimer assembly

Figure 5. Most p27S10A is bound to cyclin–CDK complexes. IPs in primary MEF protein extracts. (A,B) IP with a monoclonal p27 antibody [p27 F8] that immunoprecipitates only free p27 in serum-starved [A] or exponentially growing [B] cells. The amounts of p27 present in the cell extracts used for IP are also indicated. Immunoprecipitates were resolved on SDS-PAGE, and membranes were probed with a polyclonal p27 antibody (C19). In A, the membrane was stripped and reprobed with a polyclonal antibody to cyclin D. (C) Same as A, except that an additional extract from p27−/− cells expressing a p27S10A/CK− mutant was used for IP. (D) Same as C with IPs performed with a polyclonal p27 antibody [C19] that immunoprecipitates all forms of p27. (E,F) IPs with a monoclonal cyclin D1 antibody in serum-starved (E) or exponentially growing (F) primary MEFs. Membranes were probed successively with polyclonal antibodies to phospho-Ser10 p27, p27, CDK4, and cyclin D. For F, the cell extracts used were the same as in B. In B and F, the control lane (C) corresponds to an IP of wild-type extract performed with an irrelevant antibody.

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operate in a single p27 turnover pathway in quiescent cells.

One possible explanation for these data is that Ser10 phosphorylation directly affects the binding of p27 to cyclins and CDKs or that it modulates its ability to inhibit cyclin/CDK complexes. To test this, the ability of recombinant p27, p27S10A, and the phospho-mimetic mutant p27S10E to bind to and inhibit cyclin E–CDK2 in vitro was measured. We found that the amounts of recombinant p27, p27S10A, and p27S10E bound to cyclin E–CDK2 were similar [Fig. 6A]. Likewise, the Ki of these three forms of p27 for inhibition of recombinant cyclin E–CDK2 were almost indistinguishable [Fig. 6B], indicating that the abilities of these proteins to bind to and to inhibit cyclin E–CDK2 complexes in vitro were not substantially different from one another.

An alternative hypothesis to explain the sequestration of p27S10A into cyclin–CDK complexes was that Ser10 phosphorylation promoted p27 export to the cytoplasm, which would create a pool of unbound p27 by segregating it from its cyclin and CDK partners (Rodier et al. 2001; Boehm et al. 2002; Ishida et al. 2002; Connor et al. 2003). To address this, the subcellular localization of the p27S10A and wild-type p27 proteins in MEFs was compared. In these experiments we previously found it crucial to use p27-null MEFs as a control, because all available p27 antibodies detect a modest nonspecific background signal in the cytoplasm of fibroblasts (Besson et al. 2004b). Wild-type p27 MEFs contained a prominent nuclear pool of the p27 protein, and additionally a weak cytoplasmic signal that was reproducibly stronger than the background detected in the p27-null cells [Fig. 7; Supplementary Fig. 6; Besson et al. 2004b]. p27S10A cells also contained a clear nuclear pool of p27 protein, but the cytoplasmic signal did not surpass the nonspecific background signal detected in the p27-null cells [Fig. 7; Supplementary Fig. 6]. Nevertheless, since the overall amount of p27S10A was lower than the wild-type p27 protein, it was possible that some cytoplasmic p27S10A protein was below the limit of detection. To overcome this limitation, similar experiments were performed in the presence of the proteasome inhibitor MG132, so that if p27 were unstable in the cytoplasm, its abundance would be increased by proteasome inhibition, making it readily detectable. Indeed, in the presence of MG132, a clear pool of cytoplasmic p27 was visible in wild-type MEFs [Fig. 7; Supplementary Fig. 6]. This pool was more abundant following serum stimulation, consistent with previous reports that p27 nuclear export is increased in proliferating cells [Rodier et al. 2001; Boehm et al. 2002; Connor et al. 2003; Shin et al. 2005]. In contrast, although proteasome inhibition increased the abundance of nuclear p27S10A, hardly any p27S10A could be detected in the cytoplasm [Fig. 7; Supplementary Fig. 6]. Further evidence in support of the hypothesis that the p27S10A mutation alters the subcellular localization of the protein is provided below, as is evidence that misregulation of this pathway is critical during tumor progression. Thus, our results suggest that the increased association of p27S10A with cyclin–CDK complexes is a likely consequence of its sequestration in the nucleus.

Phosphorylation on Ser10 controls tumor suppression by p27

p27 is a dosage-dependent tumor suppressor [Fero et al. 1998]. Since the p27S10A/S10A homozygous mouse ex-
pressed reduced amounts of p27, it might have been predicted that it would demonstrate increased tumor susceptibility, similar to the p27−/− mouse. However, the p27S10A mutation increases its association with cyclin–CDK complexes, probably by enforcing the nuclear retention of p27, and thus may enhance the ability of p27 to inhibit cell proliferation. Therefore, the overall effect of the Ser10 pathway on tumor suppression was not easily predicted.

To address the importance of Ser10 phosphorylation in controlling tumor suppression by p27, we compared the susceptibility of wild-type, p27+/−, p27−/−, and p27S10A/S10A mice to urethane-induced tumorigenesis. In this model, mice develop primarily lung tumors that carry an activating mutation at codon 61 of the K-Ras oncogene (Barbin 2000; Meuwissen and Berns 2005). Loss of one allele of p27 was sufficient to significantly increase the number of lung tumors compared with wild-type animals, but the loss of both p27 alleles did not further increase tumor number [Fig. 8A]. Even though p27S10A protein levels in the lungs were similar to the p27 levels found in the p27 heterozygous mice [Fig. 8B], p27S10A/S10A mice developed less tumors than all the other groups. Thus, despite equivalent levels of p27 protein expression, p27S10A/S10A mice were substantially less susceptible to lung tumor development than p27+/− animals (∼6.7 tumors per animal compared with 12.6, respectively).

When lung tumors were scored according to their size, the number of small tumors (0–2 mm) mirrored the total number of tumors [Fig. 8C]. However, the number of large tumors (2–4 mm) was directly proportional to the number of p27 alleles: p27−/− mice had more large tumors than p27+/− animals, and the latter had more large tumors than wild-type mice (Fig. 8D). These results suggested that the growth rate of tumors was related to the dosage of the p27 protein, with the absence of p27 allowing for faster growth. In contrast, p27S10A/S10A had a greatly reduced number of large tumors compared with p27+/−, suggesting that tumors arising in the former had reduced growth rate [Fig. 8D]. Figure 8E shows representative lungs from urethane-treated animals of each genotype. In sum, p27S10A/S10A mice were partially resistant to lung tumor development, and the growth rate of tumors arising in p27S10A/S10A animals was reduced.

Although our studies focused on lung tumorigenesis, urethane treatment for 20 wk resulted in pituitary adenomas and harderian gland tumors exclusively in the

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**Figure 7.** Nuclear sequestration of p27S10A. Immunofluorescence with a monoclonal p27 antibody (clone 57, BD-Transduction Laboratories) in the presence or absence of 10 µM MG132 in wild-type and p27S10A MEFs either serum starved for 48 h (0.1% FCS), or starved and stimulated with 10% FCS for 4 h or 8 h. All images were taken with identical settings using a 100× lens, with the exposure time calculated on p27−/− MEFs processed identically so that the nonspecific staining of the antibody was not visible.
p27-null mice with a 100% penetrance (Table 2). Liver hemangiomas were observed in animals of all genotypes, with incidence increasing with the reduction of the number of p27 alleles; and the lowest incidence observed in p27S10A/S10A animals (Table 2). Uterine tumors were observed with increasing penetrance in p27+/− and p27−/− mice, but not in wild-type or p27S10A/S10A animals. Ovarian granulosa cell tumors were seen in all genotypes, but were more prominent in p27−/− mice (Table 2).

Histological analyses revealed that all tumors developing in the lungs, regardless of the genotype of the mice, were a mix of bronchioalveolar adenomas and papillary adenomas (Fig. 9A). Cytoplasmic localization of p27 is observed in many types of human cancers and is considered a negative factor for prognosis (Slingerland and Pagano 2000; Philipp-Staheli et al. 2001; Liang et al. 2002). In normal lung tissue, p27 was mostly localized in the nuclei, however, in the tumors from wild-type and p27+/− mice, p27 was largely excluded from the nuclei of

Figure 8. Decreased incidence and growth rate of urethane-induced lung tumors in p27S10A mice. (A) Total number of lung tumors in wild-type, p27+/−, p27−/−, and p27S10A mice 20 wk following a single dose of urethane. Lungs were dissected and macroscopic tumors were counted by visual inspection. The results were analyzed using a one-way analysis of variance with the Tukey-Kramer multiple comparison test. [***] p < 0.001; [**] p < 0.01; [*] p < 0.05. (B) p27S10A protein levels in the lung are similar to p27 levels in p27+/− mice. Lung extracts from wild-type (WT), p27+/−, p27−/−, p27S10A/S10A (S10A), and p27CK−/CK− (CK−) mice were resolved on SDS-PAGE, and the membrane was probed with a polyclonal p27 [C19] antibody, stripped, and reprobed with a monoclonal Grb2 antibody to evaluate protein loading. (C,D) Lung tumors were counted according to their size: small tumors (between 0 and 2 mm in diameter) (C), and large tumors (between 2 and 4 mm in diameter) (D). Statistical analyses were performed as in A. (E) Representative lungs for each mouse genotype following dissection. Tumors are indicated by arrows.
tumor cells (Fig. 9B). In contrast, in the tumors arising in the p27 promin A mice, the p27 protein remained predominantly nuclear [Fig. 9B]. In these tumors, the complete loss of the p27 protein was never observed, consistent with our immunohistochemistry data [Fig. 9C]. However, Ser10 phosphorylation was abundant in tumors from wild-type and p27−/− animals, despite high levels of proliferation as indicated by PCNA levels [Fig. 9C]. This is unlike normal tissue, in which Ser10 phosphorylation is abundant only in nonproliferating cells. These data suggest that tumor cells activated a Ser10-dependent pathway that relocated much of the cellular p27 to the cytoplasm.

Urethane-induced lung tumorigenesis has been shown to be caused by mutational activation of K-Ras [Barbin 2000; Meuwissen and Berns 2005], and tumors harvested from our cohort all showed evidence of activation of the Ras pathway, as indicated by increased phospho-ERK1/2 levels [Fig. 9C]. It has been reported that active Ras may promote p27 mislocalization to the cytoplasm. However, the importance of Ser10 phosphorylation in the pathway by which Ras induces p27 nuclear export has not been addressed [Liu et al. 2000; Kfir et al. 2005]. We therefore compared the subcellular localization of wild-type p27 and p27 promin A in MEFs expressing activated K-Ras [Fig. 10]. Overexpression of K-Ras V12 caused an increase in cytoplasmic wild-type p27, while it had little or no effect on the localization of p27 promin A. Most dramatically, Ras-expressing cells treated with the proteasome inhibitor MG-132 accumulated large amounts of wild-type p27 protein in the cell cytoplasm, whereas p27 promin A accumulated almost exclusively in the cell nucleus. Thus, Ras-induced cytoplasmic mislocalization of p27 occurred via the Ser10-dependent nuclear export pathway. In sum, our data suggest that the mutation of Ser10 to Ala impairs the Ras-induced nuclear export of p27, thus providing an explanation for the lower tumor burden in p27 promin A mice and for the reduced growth rate of these tumors. Therefore, phosphorylation of Ser10 may constitute an important pathway by which tumor cells modulate the tumor suppressor function of p27.

Discussion

The CDK inhibitor p27Kip1 plays a major role in regulating the transition between proliferating and quiescent cell states. In quiescent cells, p27 levels are elevated and then decrease when cells enter the cell cycle, remaining low through the S and G2/M phases. Remarkably, p27 expression is regulated by at least three pathways that mediate its ubiquitin-dependent proteolysis, each of which operates during a unique window of the cell cycle or in a specific subcellular compartment: a nuclear Skp2-dependent pathway that operates in S/G2 and M, and is dependent on phosphorylation of p27 on T187 [Malek et al. 2001; Nakayama et al. 2004], a KPC-dependent pathway that acts in the cytoplasm of G1 cells [Kamura et al. 2004], and a less well-understood pathway that regulates p27 degradation in quiescent cells. There may also be a fourth p27 degradation pathway that acts in the nucleus of G1 cells.

In the present study, we have used two genetically engineered mouse strains to characterize the pathway for p27 degradation that operates exclusively in quiescent cells, and have further shown that misregulation of this pathway is a key mechanism for disrupting p27’s tumor suppressor function. p27CK−/− mice carry point mutations in the protein interaction domains that are necessary for p27 to bind to cyclins and CDKs [Vlach et al. 1997]. In p27 promin A mice, the main phosphorylation site on p27 has been mutated to Ala [Ishida et al. 2000]. Unexpectedly, we discovered that these mutations together define a single pathway for p27 degradation in quiescent cells. The p27CK protein did not associate with, and therefore did not regulate, cyclins or CDKs, as evidenced by its absence from cyclin and CDK immunoprecipitated complexes, and by the increased growth rate and generalized organomegaly of the p27CK−/− mice. p27CK protein levels were increased in tissues and MEFs, probably because the degradation of p27CK− was impaired in both G0 and S phases. The increased stability of p27CK− in S phase was largely expected since transfection experiments had shown that p27CK− was resistant to cyclin E-CDK2-induced degradation, although it was still a target for phosphorylation on Thr187 by cyclin E-CDK2, indicating a requirement for a stable interaction with cyclin–CDK complexes for efficient turnover in S phase [Vlach et al. 1997; Montagnoli et al. 1999]. On the other hand, the dramatically increased stability of p27CK− in quiescent cells demonstrated that cyclin–CDK binding to p27 is also required for p27 turnover in G0. Thus, the degradation of p27 both in S phase and G0 requires its interaction with cyclin–CDK complexes. In S phase this involves the Skp2–SCF E3-ubiquitin ligase. However,

### Table 2. Reduced incidence of urethane-induced tumors in p27 promin A mice

<table>
<thead>
<tr>
<th>Tumor type</th>
<th>p27+/−</th>
<th>p27+/−</th>
<th>p27−/−</th>
<th>p27 promin A/p27 promin A</th>
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</thead>
<tbody>
<tr>
<td>Pituitary tumor</td>
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<td>0 (0/16)</td>
<td>100 (17/17)</td>
<td>0 (0/15)</td>
</tr>
<tr>
<td>Harderian gland tumor</td>
<td>0 (0/20)</td>
<td>0 (0/16)</td>
<td>100 (17/17)</td>
<td>0 (0/15)</td>
</tr>
<tr>
<td>Uterine hyperplasia</td>
<td>0 (0/8)</td>
<td>30 (8/10)</td>
<td>73 (8/11)</td>
<td>0 (0/3)</td>
</tr>
<tr>
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<td>30 (8/10)</td>
<td>82 (9/11)</td>
<td>33.3 (1/3)</td>
</tr>
<tr>
<td>Liver hemangiomia</td>
<td>25 (5/20)</td>
<td>50 (8/16)</td>
<td>59 (10/17)</td>
<td>20 (3/15)</td>
</tr>
<tr>
<td>Thymic lymphoma</td>
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<td>12.5 (2/16)</td>
<td>0 (0/17)</td>
<td>6.7 (1/15)</td>
</tr>
<tr>
<td>Lung tumors</td>
<td>100 (20/20)</td>
<td>100 (16/16)</td>
<td>100 (17/17)</td>
<td>100 (15/15)</td>
</tr>
</tbody>
</table>

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the ubiquitin ligase responsible for p27 turnover in G0 is unlikely to use Skp2. Indeed, not only is the recognition motif for Skp2 (phospho-T187) unnecessary for p27 turnover in G0, but also quiescent Skp2-null cells do not express increased amounts of p27 [Nakayama et al. 2000; Hara et al. 2001; Malek et al. 2001]. Studies using MEFs lacking specific cyclins or CDKs did not implicate one specific cyclin/CDK complex in p27 degradation in G0. This is not surprising given the very high level of redundancy among different cyclins and CDKs [Pagano and Jackson 2004; Sherr and Roberts 2004], and may thus suggest that any cyclin–CDK complex that can bind p27 can mediate this function. Our data also indicate that these cyclin–CDK complexes do not need to be enzymatically active to promote p27 degradation in G0, possibly providing a structural scaffold for p27 recognition by ubiquitin ligases as reported by others [Zhu et al. 2004]. Interestingly, cyclin–CDK binding was dispensable for p27 turnover only during the G1 phase, which may reflect p27 degradation by the cytoplasmic KPC pathway, recently found to mediate the degradation of free p27 [Kamura et al. 2004; Kotoshiba et al. 2005].

Ser10 phosphorylation stabilizes p27 in quiescent cells

p27 stability in quiescent cells is also controlled by phosphorylation on Ser10. The levels of the p27S10A protein were reduced in tissues and MEFs, consistent with its accelerated turnover in G0 compared with wild-type p27. This was in agreement with previously reported transfection experiments, which showed that p27 phosphorylation on Ser10 by the kinase Mirk/Dirk stabilizes p27 in quiescent cells [Deng et al. 2004], and that mutation of Ser10 to Ala or Glu destabilizes or stabilizes p27, respectively [Ishida et al. 2000; Rodier et al. 2001]. During the course of this work, Kotake et al. [2005] reported similar findings with regards to p27S10A turnover in a p27S10A/S10A mouse knock-in model.

A central conclusion of our experiments was that dephosphorylation of p27 on Ser10 and assembly of p27

Figure 9. Nuclear localization of p27S10A is maintained in lung tumors. (A) Lung tumor histology. Five-micron-thick slices of paraffin-embedded lungs tissues from wild-type, p27+/−, p27−/−, and p27S10A/S10A mice were stained with eosin and hematoxylin. Tumors were classified as bronchioalveolar adenomas and papillary adenomas. (B) p27 immunohistochemistry of lung tumors and normal surrounding tissue. Five-micron-thick slices of paraffin-embedded lung tissues from wild-type, p27+/−, and p27S10A/S10A mice were stained using the mouse monoclonal p27 antibody [clone 57, BD-Transduction Laboratories]. (C) Western blot analysis of normal lung tissue and individual tumors from wild-type, p27+/−, and p27S10A/S10A mice. Forty micrograms of proteins were loaded per well. Membranes were probed successively with polyclonal antibodies to phospho-Ser10 p27, p27, PCNA, phospho-ERK (activated), and with a monoclonal anti-Grb2 antibody for protein loading.
into cyclin–CDK complexes coordinately regulated p27 stability in quiescent cells, participating in a single turnover pathway. One hypothesis consistent with our data was that dephosphorylation of Ser10 promoted the assembly of heterotrimeric p27–cyclin–CDK complexes, thereby promoting p27 degradation. Indeed, the p27 S10A mutation increased the proportion of p27 that was assembled into cyclin–CDK complexes. In accord with this, p27 S10A/S10A mice had a normal growth rate and size, and a partial resistance to urethane-induced tumorigenesis, despite having overall reduced amounts of p27 protein. Also, the p27 S10A/CK− double mutant protein was stabilized relative to p27 S10A, suggesting that p27–cyclin–CDK assembly functioned downstream of Ser10 phosphorylation to control p27 stability.

Nevertheless, if Ser10 phosphorylation controls the interaction of p27 with cyclins and CDKs, it probably does so indirectly. Thus, substitution of Ser10 by the phospho-mimetic amino acid Glu (p27 S10E) did not inhibit the interaction of recombinant p27 with cyclin–CDK complexes in vitro, even though this same mutation stabilized p27 in vivo. In proliferating cells, Ser10 phosphorylation promotes the interaction of p27 with CRM1, and thereby stimulates its nuclear export (Rodier et al.

Figure 10. Unlike wild-type p27, p27 S10A is resistant to activated Ras-induced cytoplasmic translocation. p27-null, wild-type, and p27 S10A MEFs either expressing K-Ras V12 or not were serum starved for 72 h and stimulated with 10% serum in the presence or absence of MG132, fixed, and stained with a monoclonal antibody to p27. As in Figure 7, the exposure time was set on the p27−/− cells [left column] and kept constant to acquire all the other images, to take into account the nonspecific signal generated by the antibody.
p27 stability and tumor suppression

2001; Boehm et al. 2002; Ishida et al. 2002; Connor et al. 2003; McAllister et al. 2003; Shin et al. 2005). Export of p27 into the cytoplasm could represent one pathway by which Ser10 phosphorylation indirectly modulates the interaction of p27 with CDKs. Alternatively, Ser10 phosphorylation may control the interaction of p27 with other proteins in quiescent cells that could modulate its interactions with cyclin–CDK complexes.

An alternative hypothesis is that Ser10 phosphorylation directly affects the rate of degradation of p27 within cyclin–CDK complexes. For example, phosphorylation of Ser10 could inhibit p27 ubiquitination by interfering with its recognition by an E3 ubiquitin ligase. Kinetic modeling showed that both hypotheses—an effect of Ser10 phosphorylation either on heterotrimer assembly or degradation of CDK-bound p27—were equally consistent with the observed redistribution of p27S10A into cyclin–CDK complexes, as well as the relative stabilities, abundance, and phosphorylation states of p27S10A, p27C, and wild-type p27 [F. Cross, pers. comm.]. Interestingly, these models showed, in some scenarios, that the free pool of p27S10A becomes depleted because p27 levels are reduced to the point where cyclin–CDK complexes are in excess of p27. Thus, the Ser10 kinase system may regulate p27 through this critical threshold, and thereby ensure that p27 levels in quiescent cells remain in excess of cyclin–CDK complexes.

Ser10 phosphorylation in the regulation of p27 subcellular localization

Mitogenic signals activate a kinase that phosphorylates p27 on Ser10, promotes its interaction with CRM1, and thereby induces its export from the nucleus [Rodier et al. 2001; Boehm et al. 2002; Ishida et al. 2002; Connor et al. 2003; McAllister et al. 2003; Kamura et al. 2004; Shin et al. 2005]. In contrast to previous reports, Kotake et al. (2005) concluded that the p27 S10A mutation did not in- fluence the subcellular distribution of p27. However, mice expressing p27T187A (which cannot be degraded by this pathway) are not resistant to lung tumorigenesis, and lung tumors arising in those mice express reduced amounts of p27 similarly to mice expressing wild-type p27 [I. Timmerbeul, C.M. Garret-Engle, U. Kossatz, X. Chen, E. Firpo, V. Grunwald, K. Kamino, S. Kubicka, M.P. Manns, J.M. Roberts, et al., in prep.]. Cytoplasmic localization of p27 in tumor cells can be promoted by Akt-mediated phosphorylation of Thr157 (Liang et al. 2002), but this phosphorylation site is not evolutionarily conserved, and is absent from murine p27.

p27S10A mice are partially resistant to urethane-induced tumorigenesis compared with mice expressing wild-type p27 protein. Even more dramatic is their resistance to tumorigenesis compared with p27null mice, which express an amount of p27 protein that closely approximates that in the p27S10A homozygote. In our tumorigenesis experiment, p27null mice developed more tumors than p27-null animals. Others also reported an increased susceptibility to tumorigenesis of p27−/− mice compared with p27+/− mice in breast and prostate tumor models, suggesting that the maintenance of reduced amounts of p27 may, in fact, actively promote tumorigenesis [Muraoka et al. 2002; Gao et al. 2004]. In contrast, despite its reduced stoichiometry, p27S10A was a better tumor suppressor than wild-type p27. The enhanced tumor suppression by p27S10A correlates with a higher proportion of p27S10A associated with cyclin–CDK complexes than wild-type p27. This may be due to the constitutive nuclear localization of p27S10A, which increases its availability to bind to and inhibit cyclin–CDKs. Our data suggest that increased activity of the pathway that phosphorylates p27 on Ser10 may be important during tumor progression, and interfering with this pathway may be a useful therapeutic strategy. The tumor resistance of the p27S10A mice is the first example of a p27 mutation that enhances its tumor suppressor function, and thus constitutes the first direct demonstration of a pathway that is important for misregulation of p27 in cancer.
Materials and methods

Constructs, antibodies, and reagents

The 4317G9 targeting vector was kindly provided by Richard Palmitter (University of Washington, Seattle, WA). The Mox2Cre mice were kindly provided by Philippe Soriano (Fred Hutchinson Cancer Research Center, Seattle, WA). The polyclonal anti-CRM1 antibody was kindly provided by Larry Gerace (Scripps Research Institute, La Jolla, CA). pBabe-K-Ras V12 was a gift from Channing Der (University of North Carolina, Chapel Hill, NC). Reagents were purchased from the following sources: puromycin and hygromycin (Calbiochem); polyclonal antibodies to p27 [C19], CDK4 [H303], Cyclin D [H295], phospho-Ser10 p27, and monomeric antibodies to Cyclin D1 [A12], CDK2 [D12], Cyclin E [HE111], PCNA, and p27 [F8] (Santa Cruz Biotechnology); monoclonal antibodies to p27 and Grb2 (BD-Transduction Laboratories); polyclonal antibodies to phospho-p42/p44 MAPK (Thr202/Tyr204) (Cell Signaling); pQCXIP and pQCXIH retroviral vectors (BD-Clontech); cycloheximide and diphtheria toxin (DTA) were both used as negative selection markers. Targeting vectors were electroporated into AK7.1 ES cells and after positive and negative selections, Southern blot hybridization using a KpnI genomic fragment containing the two coding exons of the p27 gene was inserted into a pcDNA3.1 vector. Mutagenesis of Ser10 to Ala for the p27S10A gene was inserted on the p27+/-EcoRI genomic fragment containing the two coding exons flanked by LoxP sites, and a 2800-bp KpnI genomic fragment containing the SV40-neo cassette (used as a positive selection marker) was applied, and resistant clones were screened by PCR and direct DNA sequencing. The mutant genomic fragment was then electroporated into AK7.1 ES cells and after positive and negative selections, western blotting. Kinase assays were performed as described previously (Besson et al. 2004b). Western blotting. Kinase assays were performed as described previously (Besson et al. 2004b), with a preincubation of the indicated amount of recombinant p27 with a fixed volume of kinase reactions were incubated for 15 min at 30°C, with a preincubation of the indicated amount of recombinant p27 with a fixed volume of kinase reactions were incubated for 15 min at 30°C.

Generation of the knock-in mice

To generate the p27S10A and p27CK alleles, a 4700-base-pair (bp) KpnI–EcoRI genomic fragment containing the two coding exons of the p27 gene was inserted into a pcDNA3.1 vector. Mutagenesis of Ser10 to Ala for the p27S10A allele, and Arg30 to Ala, Leu32 to Ala, Phe62 to Ala, and Phe64 to Ala for the p27CK allele was performed using a Quick-change site-directed mutagenesis kit (Stratagene) according to the manufacturer’s instructions. Correct mutagenesis of the genomic DNA was verified by direct DNA sequencing. The mutant genomic fragment was then inserted into the 4317G9 targeting vector on one side of the SV40-neo cassette [used as a positive selection marker] flanked by LoxP sites, and a 2800-bp KpnI–Xhol genomic fragment containing the 3’-UTR of the p27 gene was inserted on the other side of the SV40-neo cassette (Fig. 1A). The coding sequences of the mutant p27 alleles were sequenced in the final targeting vectors. The herpes simplex virus thymidine kinase [HSV-TK] and diphtheria toxin [DTA] were both used as negative selection markers. Targeting vectors were electroporated into AK7.1 ES cells and after positive and negative selections were applied, and resistant clones were screened by PCR and Southern blot hybridization using a KpnI–HindIII probe containing the 3′-UTR of the p27 gene was inserted on the other side of the SV40-neo cassette [Fig. 1A]. The coding sequences of the mutant p27 alleles were sequenced in the final targeting vectors. The herpes simplex virus thymidine kinase [HSV-TK] and diphtheria toxin [DTA] were both used as negative selection markers. Targeting vectors were electroporated into AK7.1 ES cells and after positive and negative selections were applied, and resistant clones were screened by PCR and Southern blot hybridization using a KpnI–HindIII probe containing the 3′-UTR of the p27 gene was inserted on the other side of the SV40-neo cassette [Fig. 1A]. The coding sequences of the mutant p27 alleles were sequenced in the final targeting vectors.

Tissue culture and retroviral infections

Tissue culture conditions, primary and immortalized MEFs preparation, and retroviral infections were performed as described previously [Besson et al. 2004b].

Immunofluorescence, Western blotting, and flow cytometry

Immunofluorescence and Western blots were performed as described previously [Besson et al. 2004b], except for the phospho-specific antibodies, which were diluted in PBS, 0.1% Tween 20, and 5% bovine serum albumin. Cell cycle profiles by flow cytometry were performed as described previously [Besson and Yong 2000].

Protein half-life determination

For half-life determination in exponentially growing cells, cells were seeded at ~60% confluency in 100-mm plates, grown over night, and treated with 25 µg/mL cycloheximide for the indicated time. Cells were lysed in 200 µL of 2× sample buffer, sonicated for 10 sec, and boiled. Equal volumes of each lysate were resolved on SDS-PAGE. For half-life determination of other conditions, cells were seeded as above, grown overnight, and serum starved in 0.1% FCS DMEM for 72 h. For half-lives in serum-starved cells, cycloheximide was then added. For G1-phase half-lives, serum-starved cells were trypanospinized and replated for 4 h in 10% FCS DMEM prior to cycloheximide treatment. For S-phase half-lives, serum-starved cells were replated in 10% FCS DMEM for 15 h prior to cycloheximide treatment.

Coimmunoprecipitations

Primary MEFs were lysed in IP buffer [50 mM HEPES at pH 7.5, 150 mM NaCl, 1 mM EDTA, 2.5 mM EGTA, 10% glycerol, 0.1% Tween 20, complemented with 1 mM dithiotreitol, 10 mM β-glycerophosphate, 10 mM NaF, 1 mM sodium orthovanadate, 10 µg/mL leupeptin, 10 µg/mL aprotinin, 10 µg/mL pepstatin-A, and 1 mM phenylmethylsulfonylfluoride]. Lysates were syringed several times with a 26 g/2-gauge needle. Cellular debris were eliminated by centrifugation for 5 min at 14,000 rpm. Protein concentration was determined using the Bio-Rad protein assay. For each immuno precipitate, 4 µg of the indicated antibody, 500 µg of lysate, 16 µL of protein A sepharose beads (Amersham), and 500 µL of IP buffer were incubated for 3 h at 4°C. Immunoprecipitates were rinsed three times in IP buffer, and 10 µL of 4× sample buffer were added to the bead pellet. The immunoprecipitates were subjected to Western blotting as described previously [Besson et al. 2004b].

In vitro binding and kinase assays

Cyclin E–CDK2 complexes were produced in SF21 cells, and an equal amount of lysate was used in each assay. For the binding assays, 2.5 or 5 ng of recombinant p27 were mixed with 5 µL of cyclin E–CDK2-containing lysate in IP buffer. Cyclin E was then immunoprecipitated with a polyclonal antibody, and the amount of p27 associated with cyclin E was determined by Western blotting. Kinase assays were performed as described previously [Besson and Yong 2000], with a preincubation of the indicated amount of recombinant p27 with a fixed volume of lysate containing the kinase complexes for 15 min at 4°C, and the kinase reactions were incubated for 15 min at 30°C.

Urethane tumor study

p27+/- mice in the 129S4 background were bred to generate the p27+/-, p27+/-, and p27+/- mice used in the study. Mixed 129S4/C57BL6 p27+S10A mice backcrossed twice into 129S4 were bred to generate the p27+/-, p27+S10A, and p27+S10A/S10A animals. A single intraperitoneal injection of urethane (1 g/kg) in pups between 21 and 28 d old was used as a carcinogenic insult. Mice were sacrificed after 20 wk, dissected, and examined for gross lesions of internal organs. Lungs were dissected and photographed with a Nikon Coolpix 3700 digital camera, and lung tumors were counted and their size measured. Lungs were fixed...
overnight in formalin, embedded in paraffin, and sectioned for histochemistry (5 µm thickness).

**Immunohistochemistry**

Lung tissue from wild-type, p27−/−, p27+/−, and p27S10A/S10A mice was fixed in 4% paraformaldehyde in PBS, processed as paraffin sections, and deparaffinized. p27 immunostaining was performed according to the methods provided in the Immunocruz Staining System Kit (Santa Cruz Biotechnology). Briefly, hydrated tissue sections were steamed for 20 min in 0.1% citric acid and blocked in serum for 20 min at room temperature. Samples were subsequently incubated with anti-p27 antibody [Rb p27 C19, 1:100 dilution] for 1 h and biotinylated secondary antibody for 30 min, and visualized using the chromogen 3′,3′-diaminobenzidine [DAB].

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