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Normal Cell Cycle and Checkpoint Responses in Mice and Cells Lacking Cdc25B and Cdc25C Protein Phosphatases

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The Cdc25 family of protein phosphatases positively regulates cell division by activating cyclin-dependent protein kinases (CDKs). In humans and rodents, there are three Cdc25 family members—denoted Cdc25A, Cdc25B, and Cdc25C—that can be distinguished based on their subcellular compartmentalizations, their abundances and/or activities throughout the cell cycle, the CDKs that they target for activation, and whether they are overexpressed in human cancers. In addition, murine forms of Cdc25 exhibit distinct patterns of expression throughout development and in adult tissues. These properties suggest that individual Cdc25 family members contribute distinct biological functions in embryonic and adult cell cycles of mammals. Interestingly, mice with *Cdc25C* disrupted are healthy, and cells derived from these mice exhibit normal cell cycles and checkpoint responses. *Cdc25B*^{-/-} mice are also generally normal (although females are sterile), and cells derived from *Cdc25B*^{-/-} mice have normal cell cycles. Here we report that mice lacking both *Cdc25B* and *Cdc25C* are obtained at the expected Mendelian ratios, indicating that *Cdc25B* and *Cdc25C* are not required for mouse development or mitotic entry. Furthermore, cell cycles, DNA damage responses, and Cdc25A regulation are normal in cells lacking *Cdc25B* and *Cdc25C*. These findings indicate that *Cdc25A*, or possibly other phosphatases, is able to functionally compensate for the loss of *Cdc25B* and *Cdc25C* in mice.

Cell cycle advancement is regulated, in part, by the dephosphorylation and activation of cyclin-dependent protein kinases. Dephosphorylation and activation of cyclin-dependent protein kinases are, in turn, catalyzed by the Cdc25 family of protein phosphatases. Budding and fission yeasts encode a single member of this family, whereas the mammalian genome encodes three family members, designated Cdc25A, Cdc25B, and Cdc25C (19, 43, 45, 51, 52). The unique functional contributions made by individual Cdc25 family members to mammalian cell cycle control and checkpoint control have not yet been defined.

Experiments performed using mammalian tissue culture cells have revealed several distinguishing characteristics of Cdc25 family members. Although cells simultaneously express all three family members, Cdc25A is nuclear, whereas Cdc25B and Cdc25C shuttle in and out of the nucleus throughout interphase, and this activity is dependent on their interactions with 14-3-3 proteins (9–11, 22, 23, 26, 49). Early cell cycle studies indicated that the G₁-to-S-phase transition was regulated by Cdc25A (5, 26, 31, 55), whereas the G₂-to-M-phase transition was regulated by Cdc25B and Cdc25C. For example, the microinjection of antibodies specific for either Cdc25B or Cdc25C arrests cells in G₂, suggesting roles for these proteins at the G₂-to-M-phase transition (38, 43). In certain cell types, Cdc25B has been shown to be an unstable protein that accumulates during the S and G₂ phases of the cell cycle. In other cell lines, the activity of Cdc25B is regulated such that it is most

active during the S and G₂ phases of the cell cycle (18, 38, 47). Furthermore, Cdc25B has a Cdk binding site within its C terminus that enables the efficient dephosphorylation and activation of cyclin A/Cdk2 and cyclin B1/Cdk1 complexes in vitro (17, 27, 54). Cdc25B has been proposed to regulate centrosomal microtubule nucleation during mitosis (18). Unlike Cdc25B, the intrinsic phosphatase activity of Cdc25C is low during the S and G₂ phases of the cell cycle. Cdc25C is activated in mitosis due to phosphorylation by Cdk1/cyclin B1 and the Polo-like kinase Plk1 (1, 13, 25, 28, 29, 34–36, 38, 50, 60). Cdk1/cyclin B complexes have been proposed to be the primary targets for activation by Cdc25C (14, 20, 25, 39, 58).

Microinjection of antibodies against Cdc25A arrests cells in G₁, and the overexpression of Cdc25A accelerates entry of cells into S phase, implicating a role for Cdc25A in regulating the G₁-to-S-phase transition (5, 26, 31, 55). In addition, *Cdc25A* is an E2F target gene, and Cdc25A is required for the efficient induction of S-phase entry by E2F-1 (59). However, recent evidence suggests that Cdc25A also plays a role in regulating the G₂-to-M-phase transition (8, 42, 63, 66). Cdc25A is present and active in all stages of the cell cycle, and Cdc25A levels actually rise as cells progress from S phase to mitosis (4, 5, 26, 31, 42, 44, 55). Importantly, Cdc25A has a docking site for cyclin B1/Cdk1 within its C terminus, which is masked by 14-3-3 proteins during interphase but exposed during mitosis (8). Additional evidence includes the observation that Cdc25A overexpression accelerates mitotic entry (44) and the overproduction of phosphatase-dead Cdc25A delays mitotic entry (33).

Cdc25A and Cdc25C are targets of negative regulation by checkpoints that respond to various forms of genotoxic stress. Checkpoint activation maintains Cdc25C in a 14-3-3 protein-

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bound form, and the overproduction of a mutant of Cdc25C that cannot bind to 14-3-3 proteins causes a partial bypass of both the DNA replication and G₂ DNA damage checkpoints (49). Cdc25A stability is regulated as a function of the cell cycle, and Cdc25A is rapidly degraded in a proteasome-dependent manner in cells exposed to UV light or ionizing radiation (IR) (4, 6, 12, 16, 21, 24, 30, 41, 42, 44, 66). Chk1 phosphorylates Cdc25A to target it for proteolysis during an unperturbed cell cycle, and the integrity of the Chk1/Cdc25A pathway is required for cells to delay in the S and G₂ phases of the cell cycle following checkpoint activation (57, 63, 66). In addition, the overexpression of Cdc25A bypasses the G₁ DNA damage checkpoint and the intra-S-phase checkpoint, resulting in enhanced DNA damage and decreased cell survival (42, 44).

In mice, *Cdc25A*, *Cdc25B*, and *Cdc25C* exhibit overlapping but distinct patterns of expression during development and are expressed in tissue-specific patterns in adult mice (32, 46, 61, 62). This fact suggests that these genes have distinct biological functions in embryonic and adult mice. Mice lacking *Cdc25C* are viable, develop normally, and do not display any obvious abnormalities (7). Furthermore, the phosphorylation status of Cdk1, the timing of entry into mitosis, and the cellular responses to DNA damage are unperturbed in mouse embryo fibroblasts (MEFs) lacking *Cdc25C* (7). Mice with *Cdc25B* disrupted are also viable and healthy, although females are sterile due to a meiotic defect during oogenesis (40). One explanation for the lack of a cell cycle phenotype in mice with either *Cdc25B* or *Cdc25C* disrupted is compensation by the other family members and/or other phosphatases. To address this possibility, we generated mice lacking both *Cdc25B* and *Cdc25C*. Surprisingly, these mice are obtained at the expected Mendelian ratios and are healthy, demonstrating that *Cdc25B* and *Cdc25C* are dispensable for murine embryonic development and for mitotic entry.

MATERIALS AND METHODS

Generation and genotyping of mice with *Cdc25B* and *Cdc25C* disrupted. Mice with both *Cdc25B* and *Cdc25C* disrupted were generated by first mating *Cdc25B*^{-/-} (40) males with *Cdc25C*^{-/-} (7) females. F₂ double-heterozygous littermates of the strain background C57BL/6;129X1SvJ were then interbred to generate *Cdc25B*^{-/-} *Cdc25C*^{-/-} double knockout mice. PCR analysis was carried out on genomic tail DNA of wild-type (WT) mice, *Cdc25B*^{-/-} knockout mice, *Cdc25C*^{-/-} knockout mice, and the *Cdc25B*^{-/-} *Cdc25C*^{-/-} double knockout mice (BCKO) by using specific primers to confirm the genotypes of the mice. PCR analysis for *Cdc25C* was achieved by using a three-primer PCR with primer 5'-GGTTCCTTGGATTCATCTGGACC from exon 3, primer 5'-CCTCGTGCTTACGGTATCGCCG from the neomycin phosphotransferase gene cassette, and primer 5'-CCTACCATGAGTGCAGGGCACC from the intronic sequence between exons 3 and 4 as described previously (7). Three PCRs were carried out to identify mice lacking *Cdc25B*. Primers 5'-GCCTGTGAGGCCA CCTAC and 5'-CACTGAACACAACAGACTTC amplify a region in the 3' untranslated region. Primers 5'-TGGAGAGGCTATTCGGCTATGAC and 5'-GGCATCGCCATGGGTACAGCAGA amplify a region within the neomycin phosphotransferase gene cassette. Primers 5'-GACTGATCGGAGATTACTCT and 5'-TGGAGAGGCTATTCGGCTATGAC amplify a region between exon 10 and the neomycin phosphotransferase gene cassette.

Histology. Tissues were fixed in 10% neutral buffered formalin, rinsed in phosphate-buffered saline (PBS), and stored in 70% ethanol. Fixed tissues were embedded in paraffin by using standard procedures. Blocks were sectioned (5 mm) and stained with hematoxylin and eosin.

Animal growth measurements. Heterozygous crosses were used to generate mice, and postnatal growth curves were obtained by weighing pups at the indicated times by using a Mettler AE 50 scale. Genotyping was performed by PCR analysis of tail cuttings. All recorded weight averages shown in the growth curves were compiled from at least 30 mice per genotype per time point.

Northern blotting. RNA was isolated from cells by using the QuickPrep total RNA extraction kit (Amersham Pharmacia Biotech) according to the manufacturer's suggestions. RNA was resolved on a 1.2% agarose gel and then transferred to a GeneScreen Plus membrane (NEN). The probes used for screening mouse tissues and cells include a 440-bp HincII/DraIII restriction fragment of the murine *Cdc25C* (*mCdc25C*) cDNA, a 427-bp XmnI/PvuII restriction fragment of the *mCdc25A* cDNA, a 466-bp BglIII/SacII restriction fragment of the *mCdc25B* cDNA, a 600-bp HindIII/EcoRI restriction fragment of *GAPDH*, and an ~1.2-kb XbaI/HindIII restriction fragment of the neomycin phosphotransferase gene. Probes were labeled with dCTP (α -³²P; NEN) by using the Megaprime DNA labeling system (Amersham). Blots were prehybridized with ExpressHyb solution (Clontech) containing 100 μ g of sonicated salmon sperm DNA/ml for 2 to 3 h at 68°C with shaking. Labeled probe was added to 10⁶ cpm/ml, and the blot was hybridized in ExpressHyb solution for 2 h at 68°C. Blots were washed with 2 \times SSC (1 \times SSC is 0.15 M NaCl plus 0.015 M sodium citrate)-0.5% sodium dodecyl sulfate (SDS) at room temperature twice and then washed once with 2 \times SSC-0.5% SDS for 15 min with shaking at 50°C.

Generation of mouse embryo fibroblasts. Mouse embryonic fibroblasts were derived from 13.5-day-old embryos. Following the removal of the head and organs, each embryo was rinsed with PBS, minced, and digested with trypsin-EDTA (0.5% trypsin, 0.53 mM EDTA) for 10 min at 37°C, using 1 ml per embryo. Trypsin was inactivated by the addition of Dulbecco's modified Eagle's media (DMEM; Gibco-BRL) containing 10% fetal bovine serum (FBS), 2 mM L-glutamine, a 0.1 mM concentration of nonessential amino acids, 140 mM 2-mercaptoethanol, 100 U of penicillin G/ml, and 100 mg of streptomycin/ml. Cells from single embryos were plated into one 100-mm-diameter tissue culture dish and incubated at 37°C in a 10% CO₂-humidified chamber for 3 days and then expanded. Each trypsinization and replating represented one passage, and the split ratio of each passage was 1:3. Early-passage cells (P2 to P4, prior to crisis) were analyzed for cell cycle progression and for cellular responses to DNA damage.

Antibodies. Fluorescein isothiocyanate (FITC)-conjugated mouse anti-BrdU monoclonal antibody was purchased from BD Biosciences. Mouse monoclonal antibodies specific for Cdc25A were raised against baculovirus-produced glutathione S-transferase (GST)-mCdc25A. Cdk1 was detected with Cdk1-specific rabbit polyclonal antibody (48). β -Catenin mouse monoclonal antibodies were purchased from BD Transduction Laboratories. Phosphohistone H3 antibodies were purchased from Upstate Biotechnology (Lake Placid, N.Y.) and then visualized with FITC-conjugated goat anti-rabbit antibody (Jackson ImmunoResearch, Inc.). Bound primary antibodies were detected with horseradish peroxidase-conjugated rat anti-rabbit antibody (Zymed) or horseradish peroxidase-conjugated goat anti-mouse immunoglobulin G (IgG) plus IgM antibody (Jackson ImmunoResearch, Inc.), and proteins were visualized by chemiluminescence with the ECL reagent (Amersham Pharmacia Biotech).

S-to-G₁-phase progression in absence and presence of IR. A total of 10⁶ MEFs were seeded onto 100-mm-diameter tissue culture dishes 36 h prior to bromodeoxyuridine (BrdU) labeling. Cells were incubated in 5 ml of culture medium (DMEM, 10% FBS, 2 mM L-glutamine, a 0.1 mM concentration of nonessential amino acids, 140 mM 2-mercaptoethanol, 100 U of penicillin G/ml, and 100 mg of streptomycin/ml) containing 20 mM bromodeoxyuridine (Amersham Pharmacia Biotech) at 37°C for 1 h. Medium was removed and replaced with 10 ml of culture medium, and cells were incubated for the indicated times. Cells were harvested by trypsinization and collected by centrifugation. After the removal of the supernatant by aspiration, cells were washed once in PBS and were then suspended in 0.5 ml of PBS. Cells were fixed by the addition of 5 ml of 70% ethanol at 4°C in the dark. Pelleted cells were resuspended in 1 ml of 0.4 mg of pepsin (Sigma Chemical Co.)/ml in 0.1 N HCl and incubated with rocking for 30 min. Nuclear pellets were suspended in 1 ml of 2 N HCl-0.5% NP-40, and incubated with rocking for 1 h. After neutralization by incubation with 1 ml of 0.1 M sodium borate (pH 8.5) for 5 min, nuclear pellets were suspended in 100 μ l of PBS-TB (PBS, 0.5% Tween 20, 1% bovine serum albumin [BSA]) and stained with 5 μ l of FITC-conjugated anti-BrdU monoclonal antibody for 1 h in the dark. Nuclei were washed once with 1 ml of PBS-TB and then incubated with 1 ml of PBS-TB containing 30 μ g of propidium iodide (PI)/ml and 250 μ g of RNase A/ml for 30 min in the dark. Cells were analyzed for DNA content by flow cytometry using a FACSCalibur instrument (Becton Dickinson Instruments). The data were analyzed using CELLQUEST analysis software (Becton Dickinson).

G₀-to-S-phase progression. A total of 10⁶ MEFs were seeded onto 100-mm-diameter tissue culture dishes. The following day, cells were incubated in 10 ml of medium (DMEM, 2 mM L-glutamine, a 0.1 mM concentration of nonessential amino acids, 100 U of penicillin G/ml, and 100 μ g of streptomycin/ml) containing 0.1% FBS for 96 h. Cells were reseeded at 10⁶ in medium containing 10% FBS

and pulsed with 20 μ M BrdU for 1 h prior to harvest. Harvesting, fixing, staining, and analysis of cells were carried out as described above.

IR-induced G₁-phase checkpoint in MEFs. Cells were plated and serum starved as for G₁ analysis with the following changes. After serum starvation, 2×10^6 cells were seeded onto 100-mm-diameter tissue culture dishes and either mock irradiated or exposed to 20 Gy of IR from a ⁶⁰Co source. Medium containing 10% FBS and 20 μ M BrdU was added, and cells were harvested 24 h later. Cells were fixed, stained, and analyzed by flow cytometry as described above.

IR-induced G₁-phase checkpoint in thymocytes. Eight- to 10-week-old mice were irradiated with 10 Gy of IR, and injected intraperitoneally with 1 ml of 10 mM BrdU–1 mM fluorodeoxyuridine (Amersham Pharmacia Biotech) at 2 h postirradiation (3). Mice were sacrificed 1 h after the injection, and thymocytes were isolated by crushing thymi between glass slides. The prepared thymocytes were fixed, stained, and analyzed by flow cytometry as outlined above.

IR-induced S-phase checkpoint. MEFs were labeled with 10 nCi of ¹⁴C-thymidine (NEN Life Science Products)/ml for 24 h to control for total DNA content between samples. Cells were then washed and incubated in culture medium for 24 h. Cells were either mock irradiated or exposed to 5, 10, or 20 Gy of IR, incubated for the indicated times, and then pulse labeled with 2.5 μ Ci of [³H]thymidine (NEN Life Science Products)/ml for 15 min. Cells were harvested, washed twice in PBS, and then incubated in 100% methanol for 5 min, followed by incubation in 10% trichloroacetic acid for 5 min and then 0.3 N NaOH. After neutralization with HCl, radioactivity was determined by liquid scintillation counting. The resulting ratios of ³H counts per minute to ¹⁴C counts per minute, corrected for counts per minute that resulted from channel crossover, were measures of DNA synthesis.

IR-induced G₂-phase checkpoint. MEFs were either mock irradiated or exposed to 6 Gy of IR. After 40 min of incubation, 100 ng of nocodazole/ml was added to trap cells in mitosis, and samples were harvested by trypsinization 1 h 20 min later. A total of 5×10^5 cells were fixed by the addition of 5 ml of 70% ethanol at 4°C for as long as 24 h. After fixation, cells were washed twice with PBS, suspended in 1 ml of 0.25% Triton X-100 in PBS, and incubated at 4°C with rocking for 15 min. After centrifugation, cell pellets were suspended in 100 μ l of PBS containing 1% BSA and 0.75 μ g of anti-histone H3 antibody and incubated for 1.5 h at room temperature. Cells were rinsed with PBS containing 1% BSA and incubated with FITC-conjugated goat anti-rabbit IgG antibody diluted at a ratio of 1:1,200 in PBS containing 1% BSA. After a 30-min incubation at room temperature in the dark, cells were washed again and incubated with 500 μ l of PBS with 1% BSA containing 30 μ g of PI/ml and 250 μ g of RNase A/ml for 30 min, also in the dark. Cells were analyzed by flow cytometry as described above.

Synchronization of MEFs and analysis of Cdk1 phosphorylation. A total of 5×10^5 MEFs were seeded onto 100-mm-diameter tissue culture dishes. The following day, cells were incubated in 10 ml of culture medium containing 0.1% FBS for 48 h. Cells were then incubated in culture medium containing 15% FBS and 1 μ g of aphidicolin/ml (Calbiochem) for 20 h. Cells were released from the aphidicolin block by being washed twice with 10 ml of PBS and then being incubated in culture medium containing 15% FBS. Cells were harvested prior to release (time zero) or at 3, 6, 9, or 12 h after release by trypsinization and collected by centrifugation. After the removal of the supernatant by aspiration, cells were washed once in PBS and then were suspended in 0.5 ml of PBS. Cells were fixed by the addition of 5 ml of 70% ethanol at 4°C. Cells were washed once with 1 ml of PBS–1% FBS and then incubated with 1 ml of PBS–1% FBS containing 30 μ g of PI/ml and 250 μ g of RNase A/ml for 30 min. Cells were analyzed for DNA content by flow cytometry using a FACSCalibur instrument (Becton Dickinson). The data were analyzed by using CELLQUEST analysis software (Becton Dickinson). For monitoring Cdk1 phosphorylation, cells were lysed in MCLB {50 mM Tris-HCl, pH 8.0, 2 mM dithiothreitol, 5 mM EDTA, 0.5% Nonidet P-40, 100 mM NaCl, 1 μ M sodium orthovanadate, phosphatase inhibitor cocktail [containing 25 μ M (–)-*p*-bromotetramisole oxalate, 5 mM cantharidin, and 5 mM microcystin LR; Calbiochem] and protease inhibitor cocktail [containing 1.04 mM 4-(2-aminoethyl) benzenesulfonyl fluoride (AEBSF), 15 μ M pepstatin A, 14 μ M E-64, 40 μ M bestatin, 20 μ M leupeptin, and 800 nM aprotinin; Sigma] } for 15 min. at 4°C. Cell lysates containing 2 mg of total cellular protein were incubated with 50 μ l of packed p13^{suc1} agarose (Upstate Biotechnology). After incubation at 4°C for 3 h, precipitates were washed four times with 1 ml of MCLB. Proteins were resolved by SDS-polyacrylamide gel electrophoresis on a 12% SDS gel. Cdk1 phosphorylation status was monitored by immunoblotting.

Monitoring Cdc25A levels. Asynchronously growing populations of MEFs were lysed in MCLB. Lysates were resolved by SDS-polyacrylamide gel electrophoresis on an 8% SDS gel, transferred to nitrocellulose, and incubated with mouse ascites specific for mCdc25A. For synchronization studies, 6×10^5 MEFs

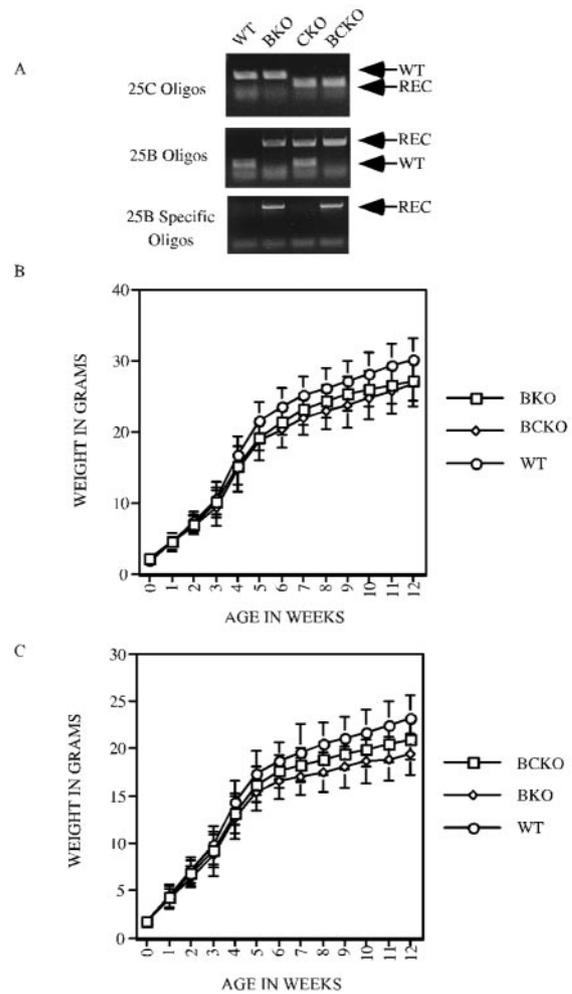


FIG. 1. Genotyping and growth rates of mice with *Cdc25B* and *Cdc25C* disrupted. (A) PCR analysis of mouse tail DNA from WT, *Cdc25B*^{-/-} (BKO), *Cdc25C*^{-/-} (CKO), and *Cdc25B*^{-/-}*C*^{-/-} (BCKO) mice, using the indicated oligonucleotides. REC, recombinant. (B and C) WT, *Cdc25B*^{-/-} (BKO), and *Cdc25B*^{-/-}*C*^{-/-} (BCKO) male (B) and female (C) mice were weighed beginning at birth and at weekly intervals up to 12 weeks. Weights (in grams) represent averages from at least 30 mice per genotype per time point. Standard deviations are shown as error bars along the y axis.

were seeded onto 100-mm-diameter tissue culture dishes and serum starved in the same manner as for the Cdk1 studies. Cells were harvested prior to release (time zero) or at 3, 6, 9, or 12 h after release, analyzed for DNA content by flow cytometry, and processed to monitor Cdc25A levels by Western blotting.

RESULTS AND DISCUSSION

Mice heterozygous for both *Cdc25B* (40) and *Cdc25C* (7) were bred to generate mice lacking both phosphatases. PCR analysis was carried out on genomic tail DNA by using specific primers to confirm the genotypes of all mice (Fig. 1A). The cumulative genotyping of heterozygous crosses resulted in the expected ratio of each genotype (Table 1). Given that mice lacking both *Cdc25B* and *Cdc25C* were obtained at the expected Mendelian ratios, this outcome indicates that *Cdc25B* and *Cdc25C* are not required for mouse development or mitotic entry. All major organs from BCKO mice were examined histologically after hematoxylin-eosin staining and appeared

TABLE 1. Actual and expected genotypes of double-heterozygous mating

Parameter (<i>n</i>)	Result (%) for indicated genotype ^a								
	W/W	W/H	W/K	H/W	H/H	H/K	K/W	K/H	K/K
Expected ratio	1 (6.25)	2 (13)	1 (6.25)	2 (13)	4 (26)	2 (13)	1 (6.25)	2 (13)	1 (6.25)
No. of mice									
Males (398)	32 (8.04)	47 (11.81)	29 (7.29)	46 (11.56)	93 (23.37)	48 (12.06)	25 (6.28)	53 (13.32)	25 (6.28)
Females (412)	25 (6.07)	68 (16.50)	33 (8.01)	51 (12.38)	95 (23.06)	44 (10.68)	25 (6.07)	40 (9.71)	31 (7.52)
No. of MEFs (236)	19 (8.05)	31 (13.14)	14 (5.93)	24 (10.17)	80 (33.89)	22 (9.32)	7 (2.97)	24 (10.17)	15 (6.36)
Total no. of mice and MEFs (1,046)	76 (7.26)	146 (13.96)	76 (7.27)	121 (11.57)	268 (25.62)	114 (10.89)	57 (5.45)	117 (11.19)	71 (6.79)

^a The first genotype represents *Cdc25B*, and the second genotype represents *Cdc25C*. W, wild type; H, heterozygous; K, knockout.

normal (data not shown). Fluorescence-activated cell-sorting (FACS) analysis using the T-cell markers CD4 and CD8 revealed the expected percentages of single-positive (CD4⁺CD8⁻ and CD4⁻CD8⁺) and double-negative (CD4⁻CD8⁻) T cells in the peripheral blood of BCKO animals (data not shown).

The weights of individual mice from birth to 3 months of age were also determined (Fig. 1B and C). As previously reported, weights of WT and *Cdc25C*^{-/-} mice were indistinguishable (data not shown) (7). In contrast, *Cdc25B*^{-/-} and BCKO mice were 8 to 14% and 6 to 10% smaller than their WT littermates, respectively, beginning at 4 weeks of age. The basis for the differences in weights is unknown at this time.

Cell cycle progression is normal in cells lacking *Cdc25B* and *Cdc25C*. Fibroblasts derived from wild-type and knockout mouse embryos were analyzed for their ability to traverse from G₀ into S phase. MEFs were synchronized by serum starvation, and at various times after the addition of complete media, cells were incubated with BrdU for 1 h to specifically label cells that were undergoing DNA replication. Cells were stained for DNA content with PI and for replicative DNA synthesis with anti-BrdU antibody and analyzed by flow cytometry. By gating on BrdU-positive cells, the timing of S-phase entry from G₀ could be monitored. As seen in Fig. 2A, similar percentages of WT and BCKO MEFs were undergoing DNA replication at each time point tested, indicating that the progression from G₀ into S phase is not perturbed in the absence of *Cdc25B* and *Cdc25C*.

Wild-type and BCKO MEFs were also analyzed for their ability to traverse from S phase through G₂ and mitosis and into G₁. Cells were pulse labeled with BrdU to mark S-phase cells. Cells were harvested at various times after the incubation and processed for analysis by flow cytometry. By gating on BrdU-positive cells, cell cycle progression from S phase through G₂/M phase and into the G₁ phase of the cell cycle could be monitored. At the time of the pulse, 17% of WT cells and 19% of BCKO cells incorporated BrdU. As seen in Fig. 2B, the percentages of BrdU-positive wild-type and BCKO cells in each phase of the cell cycle were similar. By 3 h, 42.3 and 42.9% of BrdU-positive wild-type and BCKO cells, respectively, had progressed to the G₂/M phase of the cell cycle, and by 6 h, 31.9 and 25.2% of BrdU-positive wild-type and BCKO cells, respectively, had progressed to the G₁ phase of the cell cycle. These findings indicate that the time required for cells to progress from S to G₁ phases of the cell cycle is unaltered in murine cells lacking *Cdc25B* and *Cdc25C*.

Next, the phosphorylation status of Cdk1 was monitored in

early passage fibroblasts derived from wild-type and BCKO mouse embryos. Cdk1 is proposed to be a key target of *Cdc25B* and *Cdc25C* in vivo. The phosphorylation of Cdk1 varies as a function of the cell cycle, and changes in the electrophoretic mobility of Cdk1 can be used as a specific indicator of Cdk1 phosphorylation status and cell cycle position (2, 56). Wild-type and BCKO MEFs were arrested in early S phase, by first being cultured in medium containing low serum and then being released into complete medium containing aphidicolin. Cells released from G₁/S arrest were then analyzed for DNA content by flow cytometry and were processed to monitor Cdk1 phosphorylation status (Fig. 2C). Wild-type and BCKO cells were in S phase by 3 h after release, in G₂ phase by 6 h after release, and in the M and G₁ phases of the cell cycle by 9 and 12 h after release, respectively. As seen in Fig. 2C, two electrophoretic forms of Cdk1 were present in cells arrested at the G₁/S border (lane 1), as well as cells in the S (lane 2) and G₂ (lane 3) phases of the cell cycle. As expected, a loss of the slower electrophoretic form of Cdk1 with a concomitant increase in the fastest electrophoretic form of Cdk1 was observed as cells entered mitosis (lane 4) and moved into G₁ (lane 5). The slight delay in the dephosphorylation of Cdk1 seen in BCKO cells at the 9-h time point is a consistent finding. However, the cell cycle analysis did not reveal significant differences between wild-type and BCKO cells in their abilities to traverse through the cell cycle after release from the G₁/S-phase arrest. These findings demonstrate that sufficient quantities of active Cdk1 were generated in BCKO cells to drive cells into mitosis.

Cells lacking *Cdc25B* and *Cdc25C* respond normally to ionizing radiation. Ionizing radiation activates the IR-induced DNA damage checkpoint to induce cell cycle delays. This checkpoint monitors the integrity of the genome and arrests cells in G₁ before DNA replication (termed the IR-induced G₁-phase checkpoint), in S phase (the IR-induced S-phase checkpoint), or in G₂ before mitosis (the IR-induced G₂-phase checkpoint). Checkpoints target the *Cdc25A* and *Cdc25C* regulatory pathways to elicit cell cycle delays following ionizing radiation exposure (15, 37, 41, 42, 44, 49, 66). To monitor the IR-induced G₁-phase checkpoint in wild-type and BCKO cells, early passage MEFs were synchronized by serum starvation, released into complete medium containing BrdU, and immediately mock irradiated or gamma irradiated. Cells harvested 24 h after release were stained for DNA content with PI and for replicative DNA synthesis with anti-BrdU antibody. As seen in Fig. 3A, wild-type and BCKO cells showed 57 and 48% reductions, respectively, in the numbers of S-phase cells rela-

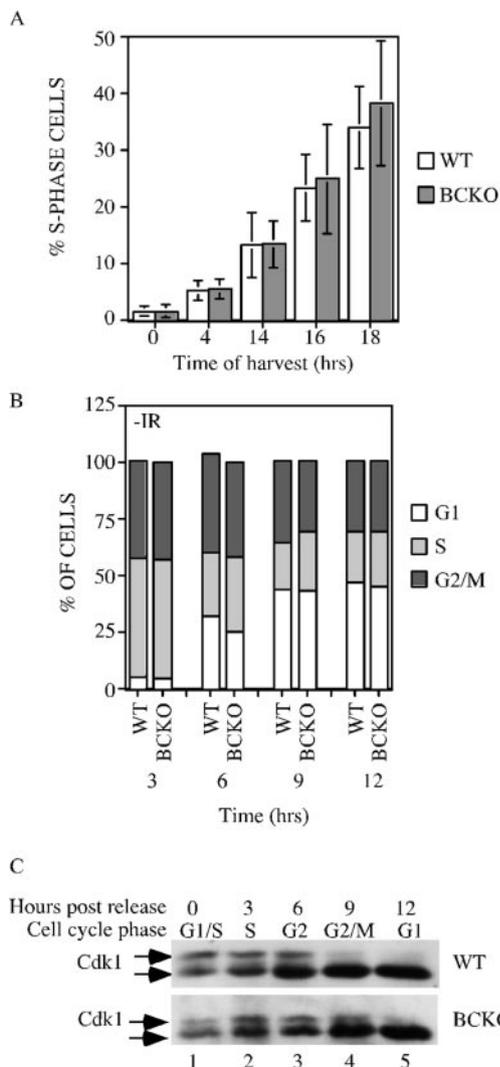


FIG. 2. Cell cycle analysis of cells lacking *Cdc25B* and *Cdc25C*. (A) WT and BCKO MEFs (passages 4 to 6) were serum starved for 96 h to synchronize cells in G₀. After the addition of serum, cells were incubated with BrdU for 1 h prior to harvest. Cells were stained with PI and for BrdU and were analyzed by flow cytometry. Each experiment was performed six times in duplicate with independent MEF strains, and standard deviations are shown as error bars along the y axis. Differences between WT and BCKO MEFs were not statistically different at any of the time points (0 h [$P = 0.936$], 4 h [$P = 0.7374$], 14 h [$P = 0.9441$], 16 h [$P = 0.6278$], and 18 h [$P = 0.2667$]). (B) MEFs at passages 4 to 6 were pulse labeled with BrdU for 1 h. Cells were harvested at the indicated times (hours) and stained with PI and for BrdU. The cellular DNA content of BrdU-positive cells was analyzed by flow cytometry. Graphs show the averages from six independent experiments. (C) Early-passage MEFs prepared from wild-type and BCKO mice were synchronized in early S phase. Cells were harvested prior to release (time zero) or at various times (hours) after release. Cdk1 precipitates were resolved on a 12% SDS gel, and Cdk1 was detected by Western blotting. The arrows indicate two electrophoretic forms of Cdk1.

tive to the numbers of mock-irradiated control cells at the 24 h time point. The difference was not statistically significant ($P = 0.2909$), indicating that the ability of cells to delay entry into S phase is intact in cells lacking *Cdc25B* and *Cdc25C*.

The IR-induced G₁-phase checkpoint in mice was also mon-

itored (3). Animals were mock irradiated or irradiated with 10 Gy of IR and injected with BrdU 2 h postirradiation. At 1 h postinjection, mice were sacrificed and their thymi were dissected. Thymocytes were analyzed by flow cytometry. As seen in Fig. 3B, irradiated wild-type and BCKO mice exhibited 48 and 51% reductions, respectively, in the numbers of thymocytes undergoing DNA replication compared with the numbers of their mock-irradiated counterparts. The difference was not statistically significant ($P = 0.1638$), indicating that the ability of thymocytes to delay entry into S phase is intact in animals lacking *Cdc25B* and *Cdc25C*.

To determine whether the absence of *Cdc25B* and *Cdc25C* perturbed the IR-induced S-phase checkpoint, [³H]thymidine incorporation into DNA was monitored 1 h after exposure of cells to various doses of IR (Fig. 3C). Both wild-type and BCKO cells had an intact S-phase checkpoint, as indicated by

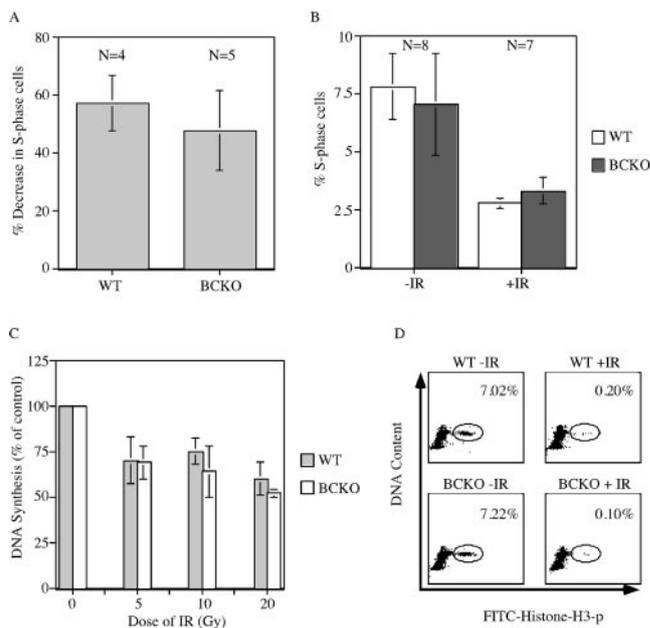


FIG. 3. The IR-induced DNA damage checkpoint is intact in BCKO cells. (A) Early-passage MEFs were serum starved for 96 h. Arrested cells were mock irradiated or gamma irradiated and then incubated in BrdU-containing complete medium. Cells were stained with PI and for BrdU and analyzed by flow cytometry 24 h later. The percent decreases in S-phase cells following irradiation are indicated. Differences between WT and BCKO cells were not significant ($P = 0.2909$) (B) WT and BCKO mice were mock irradiated (-IR) or gamma irradiated (+IR) with 10 Gy of IR and injected intraperitoneally with BrdU 2 h later. One hour after the BrdU injection, thymocytes were prepared and analyzed by flow cytometry. The percentages of S-phase cells are indicated. Standard deviations are shown as error bars along the y axis. (C) Radioresistant DNA synthesis was assessed 1 h after the exposure of early-passage MEFs to various doses of IR. Experiments were repeated four times. Differences between WT and BCKO MEFs were not statistically different at any dose of IR (5 Gy [$P = 0.9035$], 10 Gy [$P = 0.2095$], and 20 Gy [$P = 0.1333$]). Standard deviations are shown as error bars along the y axis. (D) Early-passage MEFs were mock irradiated or gamma irradiated and incubated in nocodazole 40 min later. Cells were costained for DNA content and histone H3 phosphorylation (p) and were analyzed by flow cytometry 2 h after IR. The percentages of mitotic cells are indicated. The panels are representative results obtained using WT ($n = 2$) and BCKO ($n = 3$) MEFs.

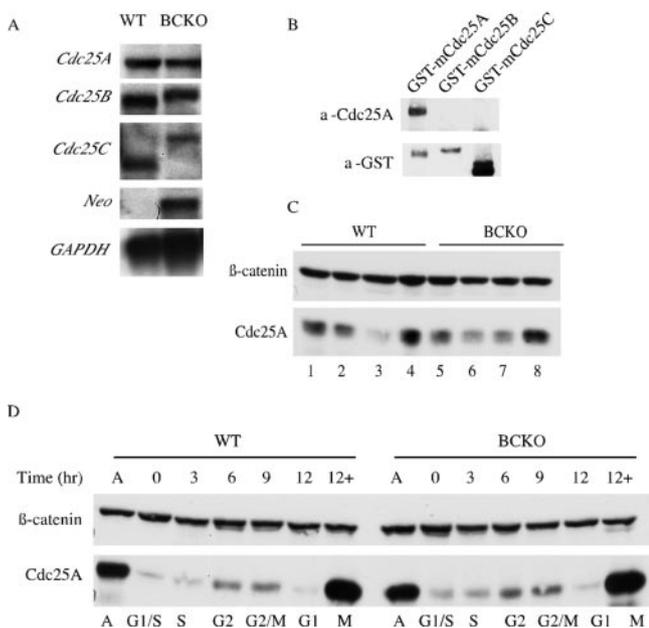


FIG. 4. Regulation of *Cdc25A* is normal in cells lacking *Cdc25B* and *Cdc25C*. (A) RNA isolated from WT and BCKO MEFs was processed for Northern blot analysis using probes specific for *mCdc25A*, *mCdc25B*, *mCdc25C*, neomycin phosphotransferase (*Neo*), and *GAPDH*. (B) GST-tagged mouse *Cdc25* proteins purified from overproducing insect cells (7) were analyzed for reactivity with the mouse *Cdc25A* (a-Cdc25A) monoclonal antibody and GST (a-GST) antibody. (C) Lysates derived from early-passage WT and BCKO MEFs were analyzed for *Cdc25A* and β -catenin levels by Western blotting. (D) Asynchronous early-passage MEFs (A) were synchronized at the G_1/S border. G_1/S -arrested cells (0) were released from the block and processed for Western blotting and flow cytometry at the indicated time points. +, samples that were incubated with nocodazole at the time of release.

the dose-dependent decrease in [3 H]thymidine incorporation into newly synthesized DNA. Finally, the ability of G_2 cells in cultures of wild-type and BCKO MEFs to delay in G_2 within 2 h post-IR was monitored (Fig. 3D). Cells were costained with PI to assess DNA content and with anti-phosphohistone H3 antibody to identify mitotic cells (64, 65). This experimental paradigm determines the percentages of G_2 cells that move into mitosis in the absence and presence of double-strand DNA breaks (64, 65). There was a 96% decrease in the number of wild-type MEFs that moved from G_2 into mitosis following exposure to IR. A similar decrease in the percentage of mitotic cells was measured for cells lacking both *Cdc25B* and *Cdc25C*, which is indicative of an intact IR-induced G_2 -phase checkpoint.

Analysis of *Cdc25A* in MEFs lacking *Cdc25B* and *Cdc25C*. One potential explanation for the lack of an apparent phenotype in mice and cells with *Cdc25B* and *Cdc25C* disrupted is compensation by the remaining *Cdc25* family member, *Cdc25A*. The levels of *Cdc25A* mRNA in wild-type and BCKO MEFs were monitored by Northern blotting, and the levels of *Cdc25A* protein were monitored by Western blotting (Fig. 4). As seen in Fig. 4A, the levels of *Cdc25A* mRNA in BCKO cells were not detectably altered. The larger RNA species detected in Northern blots probed for *Cdc25B*- and *Cdc25C*-specific mRNAs are due to the insertion of the neomycin phospho-

transferase gene after homologous recombination. Importantly, *Cdc25B* and *Cdc25C* proteins are not produced in BCKO cells (7, 40). A mouse monoclonal antibody raised against recombinant GST-mCdc25A was used to examine the levels of *Cdc25A* protein in various primary MEF strains. To test the specificity of the monoclonal antibody, GST-tagged *Cdc25A*, *Cdc25B*, and *Cdc25C* were blotted with the monoclonal antibody and an antibody against GST (Fig. 4B). Only GST-mCdc25A was recognized by the monoclonal antibody, indicating that it does not cross-react with mCdc25B or mCdc25C. Differences in total cellular levels of *Cdc25A* proteins of various MEF strains were observed, but a consistent pattern of elevated levels of *Cdc25A* in BCKO-derived MEFs was not seen (Fig. 4C). Next, *Cdc25A* regulation as a function of the cell cycle was monitored because the abundance of *Cdc25A* is regulated in a cell cycle-specific manner by ubiquitin-mediated proteolysis (4, 6, 12, 26, 30, 41, 42, 44). As seen in Fig. 4D, *Cdc25A* levels rose as wild-type and BCKO MEFs progressed from S phase through G_2 and into mitosis and then fell as cells exited mitosis and entered G_1 . A substantial increase in *Cdc25A* was observed in cells arrested with nocodazole in mitosis. Thus, the regulation of *Cdc25A* turnover during the cell cycle was normal in cells lacking *Cdc25B* and *Cdc25C*.

Concluding remarks. Mice lacking two members of the *Cdc25* family of protein phosphatases, *Cdc25B* and *Cdc25C*, are viable, develop normally, and do not display any obvious abnormalities with the exception that females lacking both phosphatases are sterile, as are females lacking *Cdc25B* alone. Importantly, these findings challenge the paradigm that entry into mitosis requires *Cdc25B* and *Cdc25C*. It is unknown at this time whether *Cdc25A*, the remaining member of the *Cdc25* family, functionally compensates for the lack of *Cdc25B* and *Cdc25C* or whether other compensatory pathways are involved. Importantly, *Cdc25A* has a docking site for Cdk1/cyclin B1 within its C terminus, indicating that *Cdc25A* is fully capable of activating Cdk1 to regulate mitotic entry (8). Mice that can have *Cdc25A*, *Cdc25B*, and *Cdc25C* conditionally deleted are currently being generated. In this way, *Cdc25* phosphatases can be disrupted individually and in combination to determine if "adaptation" during development accounts for the lack of observed phenotype in the BCKO animals, as has been seen in other germ line knockout models (53).

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