

# Evidence that Myb-related CDC5 proteins are required for pre-mRNA splicing

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**The conserved CDC5 family of Myb-related proteins performs an essential function in cell cycle control at G<sub>2</sub>/M. Although c-Myb and many Myb-related proteins act as transcription factors, herein, we implicate CDC5 proteins in pre-mRNA splicing. Mammalian CDC5 colocalizes with pre-mRNA splicing factors in the nuclei of mammalian cells, associates with core components of the splicing machinery in nuclear extracts, and interacts with the spliceosome throughout the splicing reaction *in vitro*. Furthermore, genetic depletion of the homolog of CDC5 in *Saccharomyces cerevisiae*, *CEF1*, blocks the first step of pre-mRNA processing *in vivo*. These data provide evidence that eukaryotic cells require CDC5 proteins for pre-mRNA splicing.**

The *Schizosaccharomyces pombe* *cdc5-120* mutant was isolated in a screen for mutants defective in cell cycle progression (1). At the restrictive temperature, *cdc5-120* cells arrest growth in G<sub>2</sub> (1, 2), indicating that *cdc5*<sup>+</sup> function is required for G<sub>2</sub>/M progression. CDC5 has been conserved throughout evolution, and related genes have been cloned from *Saccharomyces cerevisiae* (termed *CEF1*; ref. 3), *Arabidopsis thaliana* (4), *Drosophila melanogaster* (3), *Caenorhabditis elegans* (3), *Xenopus laevis* (5), and *Homo sapiens* (3, 6, 7). We conclude that these proteins are conserved functionally, because *D. melanogaster* and human CDC5 (hCDC5) complement the *cdc5-120* mutant, *S. cerevisiae* *CEF1* is essential during G<sub>2</sub>/M in this evolutionarily distinct yeast (3), and overexpression of dominant negative forms of hCDC5 slows G<sub>2</sub> progression in mammalian cells (8).

In their N termini, CDC5 proteins are highly related to the DNA-binding domain of human c-Myb (2, 3, 9). Whereas human c-Myb contains three Myb repeats, ≈50-amino acid motifs with characteristic spacing of tryptophan residues (9), CDC5 proteins contain two Myb repeats (R1 and R2) followed by a Myb-like-repeat (MLR3) that contains some, but not all, of the hallmarks of a typical Myb repeat (3). Based on their homologies to c-Myb, CDC5 proteins were hypothesized to carry out their essential function in cell cycle control through transcriptional regulation, a notion supported by the following observations: (i) the Myb repeats of *S. pombe* *cdc5p* fused to glutathione S-transferase-bound DNA cellulose (2); (ii) the Myb repeats of *A. thaliana* *cdc5p* selected a specific DNA sequence in a cyclic amplification and selection of targets protocol (4); and (iii) the C terminus of hCDC5 fused to the GAL4-DNA binding domain activated transcription in a reporter assay (8). To date, however, no downstream transcriptional targets for any of the CDC5 proteins have been identified.

hCDC5 was identified recently in a biochemical purification of the mammalian spliceosome assembled *in vitro* (10), indicating that CDC5 proteins may be involved in pre-mRNA splicing rather than transcriptional regulation. Herein, we extend this observation by showing that mammalian CDC5 colocalizes with splicing factors in the nuclei of mammalian cells, coimmunoprecipitates with core components of the splicing machinery from nuclear extracts, and interacts with the spliceosome throughout the splicing reaction *in vitro*. Finally, we present evidence that eukaryotic cells require the CDC5 family of proteins for pre-mRNA splicing *in vivo*.

## Materials and Methods

DNA manipulations were performed by using standard techniques (11). Details of plasmid construction and oligonucleotide sequences are available on request.

**Mammalian Cell Culture Conditions and Flow Cytometric Analysis.** NIH 3T3 and HeLa cells were cultured in DMEM and 10% (vol/vol) bovine calf serum or 10% (vol/vol) FBS, respectively. TsBN2 cells (12) were cultured and shifted to the restrictive temperature for 10 h as described (13). Asynchronously growing subconfluent NIH 3T3 cells were rendered quiescent by incubation for 24 h in DMEM containing 0.1% bovine calf serum. For flow cytometric analysis, cells were trypsinized, collected by centrifugation, and fixed overnight at 4°C in 70% (vol/vol) ethyl alcohol. The next day, cells were incubated for 30 min with 0.1 mg/ml RNase and 0.05 mg/ml propidium iodide in 50 mM sodium citrate. Flow cytometric analysis was conducted with a Becton Dickinson fluorescence-activated cell sorter (FACScan).

**Generation of hCDC5 Polyclonal Antisera.** The C-terminal 440 and the N-terminal 363 amino acids of hCDC5 were produced separately in *Escherichia coli* as hexahistidine fusion proteins and used to generate polyclonal antisera in rabbits. Antisera were affinity-purified against hCDC5C or hCDC5N as described (14).

**Protein Lysates, Subcellular Fractionation, Production, and Immunoprecipitation of hCDC5 *in Vitro*.** Subconfluent NIH 3T3 and HeLa cells were lysed in 1× SDS sample buffer [0.125 M Tris-HCl, pH 6.8/10% (vol/vol) glycerol/2% (vol/vol) SDS/10% (vol/vol) β-mercaptoethanol Bromophenol blue]. Lysates were heated to 95°C for 5 min and clarified before loading. Nuclear and cytoplasmic fractions of cultured cells were prepared as described (15). hCDC5 and 6XHis/Myc-tagged hCDC5 were produced *in vitro* in the presence of TRAN<sup>35</sup>S-LABEL (ICN) by using rabbit reticulocyte lysate (Promega). Nonidet P-40 buffer (500 μl; ref. 16) was added to each 50-μl reaction, and the lysates were clarified for 30 min at 4°C. Proteins were immunoprecipitated with 3 μl of preimmune or immune hCDC5C antiserum or 5 μg of the anti-Myc (9E10) monoclonal antibody. For immunoprecipitations with 9E10, affinity-purified rabbit anti-mouse secondary antibodies were added before addition of protein A Sepharose (Amersham Pharmacia).

**Immunoblotting.** Proteins were separated by SDS/PAGE and transferred to poly(vinylidene difluoride) membranes. Primary antibodies were used at dilutions of 1:1,000 or 1:2,000 for preimmune and immune hCDC5C antisera, respectively; 1:300

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Abbreviations: hCDC5, human CDC5; snRNA, small nuclear RNA; snRNP, small nuclear ribonucleoprotein; mCDC5, mouse CDC5.

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for affinity-purified hCDC5C antiserum; and at 0.5  $\mu\text{g}/\text{ml}$  for preimmune IgG or affinity-purified hCDC5N antiserum. Monoclonal anti-actin antibodies (Amersham Pharmacia) were used at 1:2,000. Peroxidase-conjugated goat anti-rabbit or sheep anti-mouse secondary antibodies were used at 10,000-fold dilutions. Reactive proteins were detected by using enhanced chemiluminescence detection (Amersham Pharmacia).

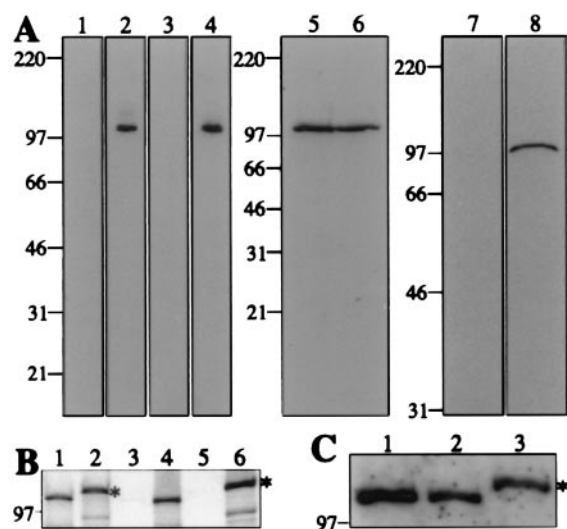
**Indirect Immunofluorescence.** Subconfluent cells attached to coverslips were fixed with 1:1 (vol/vol) methanol:acetone for 5 min at room temperature. Cells were blocked at 37°C with 3% (vol/vol) BSA, followed by incubation with 1:25 or 1:50 affinity-purified hCDC5C antiserum. For colocalization studies, cells were simultaneously incubated with 0.05 mg/ml anti-SC35 monoclonal antibody (PharMingen). Fluorescein goat anti-rabbit IgG conjugate (1:100 dilution of 2 mg/ml stock; Molecular Probes) and Texas Red goat anti-mouse IgG conjugate (1:100 dilution of 2 mg/ml; Molecular Probes) were used to detect the primary antibodies. DNA was visualized by incubating the cells in 0.5  $\mu\text{g}/\text{ml}$  Hoechst 33258 (Sigma). Cells were photographed through a Zeiss Axioskop microscope with  $\times 40$  or  $\times 100$  objectives and appropriate filters. Images were captured with a cooled CCD camera (Optronics ZVS-47DEC, Goleta, CA). Confocal microscopy and overlay analysis were performed on a Zeiss LSM410 Confocal Laser Scanning Microscope with a  $\times 63$  objective.

**Transient Transfection of Clk/Sty Kinase in NIH 3T3 Cells.** NIH 3T3 cells were transfected with a plasmid that constitutively expresses N-terminally Myc-tagged mouse Clk/Sty (pECE-M-STY; ref. 17). Transfections were performed with Lipofectamine (Life Technologies). After 24 h, cells were processed for indirect immunofluorescence with 9E10 monoclonal antibody and affinity-purified hCDC5C antiserum.

**Immunoprecipitation of Small Nuclear RNAs (snRNAs), Small Nuclear Ribonucleoproteins (snRNPs), and Spliceosomes.** Immunoprecipitations were carried out in IPP150 (10 mM Tris-HCl, pH 8.0/150 mM KCl/0.1% NP-40) buffer essentially as described (18). Briefly, 20  $\mu\text{l}$  of packed protein A Sepharose 4B fast flow resin (Sigma) was precoated with 20  $\mu\text{l}$  of hybridoma ascites containing monoclonal antibodies against maltose-binding protein (AK105; ref. 48) or 40  $\mu\text{g}$  of Sm Ab-1 (Clone Y12; Neomarkers, Fremont, CA). Beads (20  $\mu\text{l}$ ) were also precoated with 2  $\mu\text{l}$  of either preimmune or immune hCDC5C antiserum. The beads were then incubated for 1 h at 4°C with 25  $\mu\text{l}$  of HeLa-cell nuclear extract prepared as described below. snRNAs were immunoprecipitated with 100  $\mu\text{g}$  of agarose-linked 2,2,7-trimethylguanosine (Ab-1) monoclonal antibodies (Oncogene Research Products, Cambridge, MA).

*In vitro* splicing assays were carried out in HeLa-cell nuclear extracts as described (19, 20). Capped and uniformly  $^{32}\text{P}$ -labeled  $\beta$ -globin pre-mRNA substrate was produced *in vitro* from pSP64-H $\beta$ Δ6 (20, 21).  $\beta$ -Globin RNA species were visualized on 9% polyacrylamide (29:1, acrylamide/bisacrylamide)/8 M urea denaturing gels (20). Immunoprecipitation of spliceosomes was carried out after 45 or 90 min of the reaction in IP100 (50 mM Tris-HCl, pH 7.6/100 mM KCl/2 mM MgCl<sub>2</sub>/0.5 mM DTT/0.05% NP-40) buffer essentially as described, with 15  $\mu\text{l}$  of protein A beads precoated with preimmune or immune anti-hCDC5C or 30  $\mu\text{l}$  of Y12 hybridoma ascites (monoclonal anti-Sm; refs. 22 and 23).

**S. cerevisiae Strains, Preparation of Total RNA, and Northern Blot Analysis.** KGY1120 and KGY1140 strains were grown under permissive conditions and shifted to restrictive conditions exactly as described (3). *prp3-1(ts125)*; refs. 24 and 25), *prp18(ts503)*;



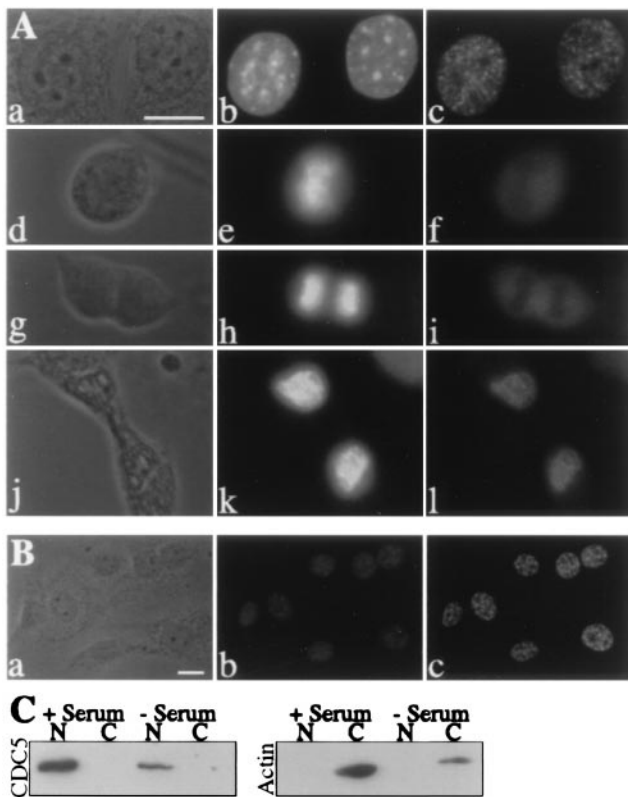
**Fig. 1.** Characterization of hCDC5 antisera. (A) Immunoblots of whole-cell protein extracts from NIH 3T3 (lanes 1, 2, and 5) and HeLa (lanes 3, 4, and 6–8) cells probed with preimmune (lanes 1 and 3), immune (lanes 2 and 4), and affinity-purified (lanes 5 and 6) antisera raised against the C terminus of hCDC5 (hCDC5C), and preimmune IgG (lane 7) and affinity-purified antisera (lane 8) raised against the N terminus of hCDC5 (hCDC5N). (B and C) Analysis of hCDC5 produced *in vitro*. hCDC5 or 6XHis/Myc-tagged hCDC5 proteins were produced *in vitro* in the presence (B) or absence (C) of TRAN<sup>35S</sup>-LABEL. In B, total products of the reactions (lanes 1 and 2), preimmune (lane 3) or anti-hCDC5C (lane 4) immunoprecipitates, and immunoprecipitates of the 6XHis/Myc-tagged hCDC5 with no primary antibody (lane 5) or 9E10 antibody (lane 6) were resolved by SDS/PAGE. Proteins were detected by fluorography. In C, an NIH 3T3 cell lysate (lane 1), hCDC5 produced *in vitro* (lane 2), and 6XHis/Myc-tagged hCDC5 produced *in vitro* (lane 3) were resolved by SDS/PAGE and immunoblotted with anti-hCDC5C serum. In both B and C, asterisks denote the 6XHis/Myc-tagged species of hCDC5. For all panels, numbers to the left indicate molecular mass (in kDa).

ref. 24), and *cdc28-1N* (26) temperature-sensitive strains were grown in yeast extract/peptone/dextrose (27).

Total RNA was prepared from cells by extraction with hot acidic phenol as described (28). Total RNA (20  $\mu\text{g}$  per sample) was electrophoresed on formaldehyde-agarose gels and capillary blotted to a Duralon-UV membrane (Stratagene). After UV crosslinking, blots were hybridized with labeled probes from *RP51a*, *DYN2*, *DBP2*, or *GLC7* ORFs or PCR fragments representing the *ACT1* or *TUB3* intron sequences. Blots were exposed to PhosphorImager screens and visualized by using MD IMAGE QUANT (version 3.3) or IMAGE QUANT for Macintosh 1.1 (Molecular Dynamics).

## Results

**Characterization of hCDC5 Antisera.** To facilitate studies of hCDC5, rabbit polyclonal antibodies were raised against the C-terminal 440 amino acids (hCDC5C) or N-terminal 363 amino acids (hCDC5N) of the protein. Crude and affinity-purified hCDC5C and hCDC5N sera recognized specifically a single protein species of 105 kDa from HeLa and NIH 3T3 whole-cell lysates (Fig. 1A; data not shown). To confirm that the 105-kDa proteins were indeed hCDC5 and mouse CDC5 (mCDC5), two full-length variants of hCDC5 differing only by the presence of a C-terminally located 6XHis/Myc epitope were produced by *in vitro* transcription/translation. Both proteins were specifically immunoprecipitated by the hCDC5C antiserum (Fig. 1B), and hCDC5 produced *in vitro* comigrated with mCDC5 from NIH 3T3 cells (Fig. 1C). Collectively, these data confirm that mammalian CDC5 migrates at 105 kDa and that our polyclonal antibodies recognize specifically mammalian CDC5 proteins.



**Fig. 2.** Subcellular localization of CDC5 throughout the cell cycle and in serum-deprived cells. (A) Phase contrast (a, d, g, and j) and fluorescence micrographs of NIH 3T3 cells stained with Hoechst (b, e, h, and k) and affinity-purified hCDC5C antiserum (c, f, i, and l). Cells in interphase (a–c), metaphase (d–f), anaphase (g–h), and telophase (j–l) are shown. (B) Localization of CDC5 in serum-deprived NIH 3T3 cells. Phase contrast (a) and fluorescence micrographs of NIH 3T3 cells deprived of serum for 24 h and stained with Hoechst (b) and affinity-purified hCDC5C antiserum (c). (C) Localization of CDC5 in nuclear and cytoplasmic fractions of asynchronously growing and serum-deprived NIH 3T3 cells. Equivalent amounts of nuclear and cytoplasmic fractions prepared from asynchronously growing NIH 3T3 cells or cells that were deprived of serum for 24 h were analyzed by immunoblotting by using either hCDC5C antiserum or a monoclonal anti-actin antibody. (Bars = 10  $\mu$ m.)

**Subcellular Localization of CDC5.** By indirect immunofluorescence, endogenous mCDC5 protein was localized to the nuclei of NIH 3T3 cells in a punctate pattern, except during metaphase and anaphase when mCDC5 became localized diffusely throughout the cell (Fig. 2A). mCDC5 was excluded from phase-dark regions of the nucleus presumed to be nucleoli. Cells in metaphase and anaphase, identified by cellular and chromosomal morphology, displayed a uniform distribution of mCDC5 with exclusion from segregating chromosomes. In telophase cells, mCDC5 localized exclusively to the reforming nucleus in a speckled pattern (Fig. 2A). Identical staining patterns were observed with affinity-purified hCDC5N antibodies, by using formaldehyde rather than methanol as fixative, in HeLa, CV-1, and COS-7 cell lines (data not shown). Cells stained with preimmune serum or secondary antibodies alone failed to produce fluorescent signals (data not shown).

To rule out the possibility that a cytoplasmic pool of mCDC5 had gone undetected by indirect immunofluorescence, NIH 3T3 cells were fractionated biochemically into nuclear and cytoplasmic pools. In agreement with the indirect immunofluorescence results, mCDC5 was detected in the nuclear but not the cytoplasmic fraction (Fig. 2C). Conversely, actin was detected in the

cytoplasmic but not the nuclear fraction, indicating that the fractionation procedure had been successful (Fig. 2C).

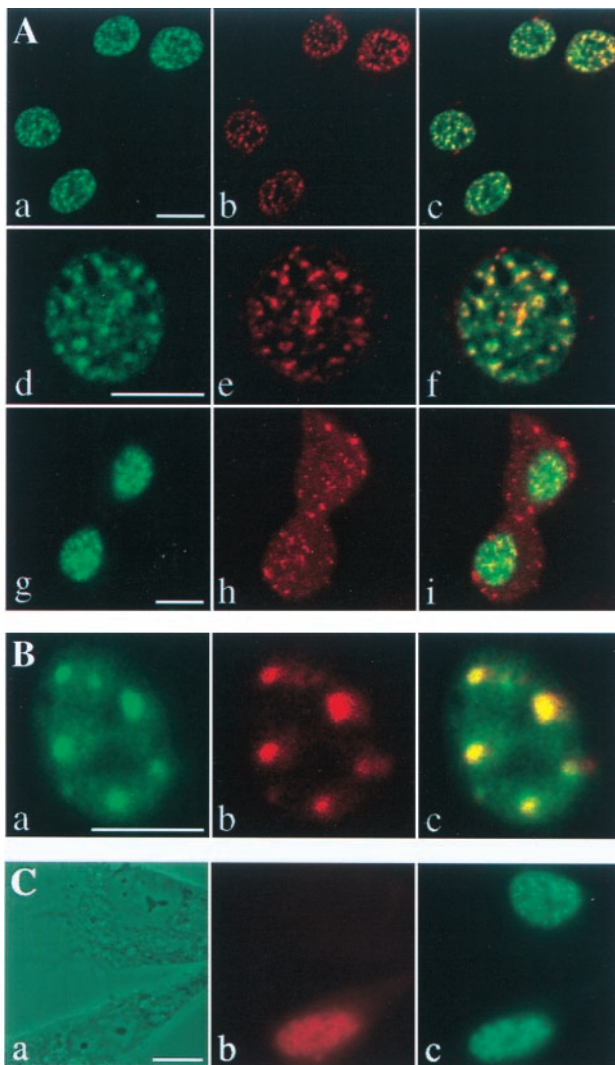
It has been reported that exogenously expressed hCDC5 is cytoplasmic after serum deprivation, and that, on serum stimulation, hCDC5 translocates into the nucleus (6). To determine whether the localization of endogenous CDC5 was altered by serum conditions, indirect immunofluorescence was performed on NIH 3T3 cells that had been deprived of serum for 24 h. A significantly greater proportion (95%) of serum-deprived cells contained a 1 N content of DNA when compared with asynchronously growing serum-fed cells (63%). Thus, these cells had been appropriately serum starved. Serum-deprived NIH 3T3 cells displayed a punctate, nuclear staining pattern of CDC5, indistinguishable from that seen in asynchronously growing cells (Fig. 2B). The nuclear localization of CDC5 was again confirmed by biochemical fractionation (Fig. 2C). CDC5 was also detected in the nucleus of serum-deprived COS-7 and CV-1 cells under the conditions used previously (data not shown; ref. 6). The pattern of punctate staining was not altered in these cells after serum addition (data not shown).

**CDC5 Localizes to the Nuclear Speckles.** The nuclear, punctate distribution of CDC5 resembled that of factors involved in pre-mRNA splicing, which are concentrated into “nuclear speckles” (for review see ref. 29). To determine whether CDC5 localized to nuclear speckles, NIH 3T3 cells were costained for a non-snRNP component of the spliceosome and a marker of the nuclear speckle (SC35; refs. 30 and 31) by using an anti-SC35 monoclonal antibody and for mCDC5 by using affinity-purified hCDC5C antiserum. In all interphase nuclei observed, the distribution of brightly staining mCDC5 foci mirrored the SC35 staining pattern (Fig. 3A), although the more diffuse pool of mCDC5 did not colocalize with SC35. Colocalization of CDC5 with SC35 was also observed in asynchronously growing and serum-deprived NIH 3T3, CV-1, and COS-7 cells (data not shown). No staining was observed when the secondary antibodies used to detect CDC5 antibodies and the SC35 monoclonal antibody were interchanged (data not shown).

During metaphase and early anaphase, SC35 staining is uniform throughout the cell body (32). In late anaphase and telophase, SC35 antigens reassociate into speckles that appear in both the cytoplasm and reforming telophase nuclei (Fig. 3A; 32). In contrast, during telophase, mCDC5 was localized completely to the reforming nuclei in foci that colocalized with the nuclear subset of SC35 speckles (Fig. 3A). These data indicate that CDC5 and SC35 translocate into the nucleus with different kinetics after chromosome segregation.

We reasoned that, if CDC5 proteins are indeed bona fide components of the nuclear speckles, then treatments that alter the spatial distribution of speckle proteins (for review see ref. 29) should similarly modulate the subnuclear localization of mammalian CDC5. In particular, tsBN2 cells (12) reorganize splicing factors into 4–10 large clusters at the restrictive temperature (13). To determine whether the CDC5 distribution is similarly reorganized, tsBN2 cells were shifted to 39.5°C for 10 h and costained for SC35 and CDC5. At 32.5°C, SC35 and CDC5 colocalized and displayed a normal speckled distribution (data not shown). In contrast, at 39.5°C, SC35 antigens and CDC5 reorganized into abnormally large clusters (Fig. 3B), indicating that CDC5 behaves like other speckle-associated proteins under these conditions.

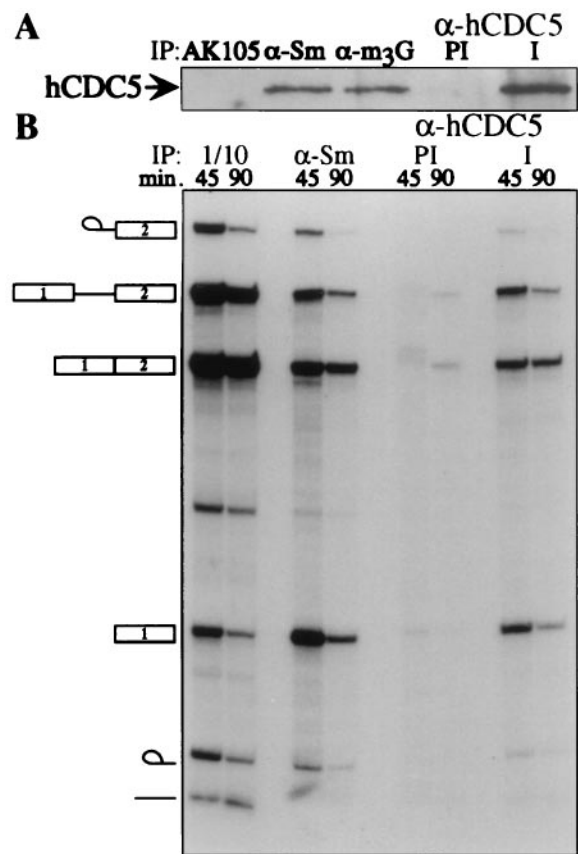
To disperse nuclear speckles, we overexpressed the SR protein kinase Clk/Sty in NIH 3T3 cells (33). A plasmid expressing Myc-tagged Clk/Sty under control of the cytomegalovirus promoter was transfected into NIH 3T3 cells, and 24 h later, cells were fixed and costained for the Myc epitope, which allowed the identification of transfected cells and mCDC5. As described previously, transfected Clk/Sty localized diffusely throughout



**Fig. 3.** CDC5 is a component of the nuclear speckles. (A) NIH 3T3 cells were costained with affinity-purified hCDC5C antiserum and a monoclonal antibody against SC35. Fluorescein and Texas Red-conjugated secondary antibodies were used to detect the distribution of CDC5 (a, d, and g) and SC35 (b, e, and h), respectively. Confocal images of cells in interphase (a–f) and telophase (g–i) were obtained and merged (c, f, and i). Regions of colocalization are yellow in merged images. (B) Asynchronously growing tsBN2 cells were shifted to the nonpermissive temperature and costained with affinity-purified hCDC5C antiserum (a) and a monoclonal antibody to SC35 (b). The confocal images were merged in c. (C) Myc-tagged Clk/Sty kinase was transfected into NIH 3T3 cells. Phase contrast images (a) of cells costained with anti-Myc epitope monoclonal antibodies (9E10; b) and affinity-purified hCDC5C antiserum (c) at 24 h after transfection. (Bars = 10  $\mu$ m.)

the nucleus (33). In transfected cells, mCDC5 was significantly more dispersed compared with untransfected cells (Fig. 3C). These data indicate that CDC5 is a bona fide component of the nuclear speckles.

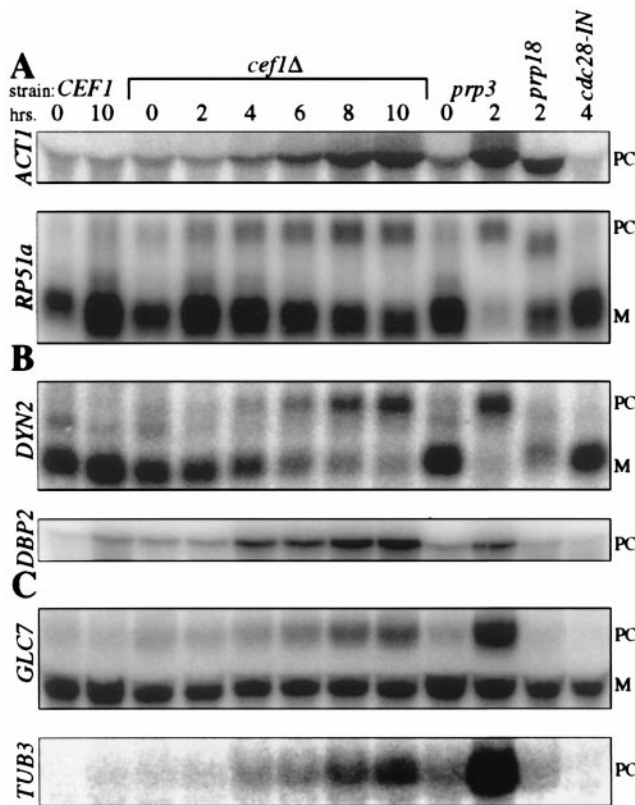
**hCDC5 Associates with Core Components of the Splicing Machinery and the Spliceosome *in Vitro*.** The observation that CDC5 colocalized with pre-mRNA splicing factors raised the possibility that CDC5 associates *in vivo* with components of the pre-mRNA splicing machinery. Indeed, hCDC5 was found in both anti-Sm and anti-m<sub>3</sub>G immunoprecipitates prepared from HeLa-cell nuclear extracts (Fig. 4A), and approximately 20% of total hCDC5 in the extract could be immunoprecipitated by using



**Fig. 4.** hCDC5 interacts with core components of the splicing machinery and the spliceosome *in vitro*. (A) Immunoprecipitations (IP) from a nuclear splicing extract were performed with three monoclonal antibodies [irrelevant monoclonal antibody (AK105), anti-Sm (Y12), and anti-snRNA cap (anti-trimethylguanosine;  $\alpha$ -m<sub>3</sub>G)] and preimmune (PI) or immune (I) hCDC5C antiserum. Immunoprecipitates were blotted with hCDC5C antiserum. (B) hCDC5C or Sm proteins were immunoprecipitated from an *in vitro* splicing reaction containing labeled  $\beta$ -globin pre-mRNA at 45 or 90 min after addition of pre-mRNA, and 1/10 of the total reactions were run as standards. The identities of the RNA species are indicated to the left.

anti-m<sub>3</sub>G antibodies (data not shown). We next investigated whether hCDC5C antiserum could immunoprecipitate spliceosomes from *in vitro* splicing reactions. After incubation of a nuclear splicing extract containing <sup>32</sup>P-labeled  $\beta$ -globin pre-mRNA for 45 or 90 min, both anti-Sm (Y12 hybridoma ascites; ref. 23) and anti-CDC5 but not preimmune antibodies immunoprecipitated approximately 5% of  $\beta$ -globin pre-mRNA, the lariat intermediate, and mature  $\beta$ -globin mRNA (Fig. 4B). By quantitating the maximal amount of  $\beta$ -globin immunoprecipitated by anti-hCDC5C serum relative to that of anti-Sm antibodies, we determined that hCDC5C was associated with approximately one-third of the spliceosomes *in vitro* (data not shown).

**S. cerevisiae Cells Lacking the Ortholog of *S. pombe cdc5*<sup>+</sup>, *CEF1*, Are Defective in pre-mRNA Splicing.** Finally, we wanted to test whether eukaryotic cells require Cdc5 proteins for pre-mRNA splicing *in vivo*. We decided to analyze *S. cerevisiae* cells genetically depleted of *CEF1* for pre-mRNA splicing defects. A *S. cerevisiae* strain (KGY1120) lacking endogenous *CEF1* and harboring plasmid-borne *CEF1* cDNA under the control of the *GAL1* promoter arrests growth during G<sub>2</sub>/M after 10 h of *CEF1* repression (3). We looked for accumulation of immature forms of several intron-containing genes by Northern blot analysis over a 10-h time course of *CEF1* repression. Cef1p is detectable until



**Fig. 5.** *S. cerevisiae* cells lacking *CEF1* are defective in pre-mRNA splicing. Strains containing (*CEF1*) or lacking the endogenous copy of *CEF1* (*cef1Δ*) harboring plasmid-borne *CEF1* cDNA under the control of the *GAL1* promoter were maintained in synthetic medium containing raffinose and galactose. *CEF1* expression was repressed by shifting the cells to synthetic medium containing glucose (SD). Aliquots of cells were collected at the number of hours indicated after the shift into SD. Total RNA was also purified from temperature-sensitive mutants *prp3-1*, *prp18* (*ts503*), and *cdc28-1N* shifted to the restrictive temperature (35.5°C) for the number of hours indicated. Total RNA (20 μg) was electrophoresed and blotted. (A) Northern blots probed with the *ACT1* intron sequence or the *RP51a* ORF. (B) Northern blots probed with the labeled ORFs for *DYN2* and *DBP2*. (C) Northern blots probed with the *GLC7* ORF or *TUB3* intron sequence. PC, precursor mRNA; M, mature mRNA.

8 h of repression (data not shown). KGY1120 mRNA was compared with that isolated from four control strains: (i) KGY1140 (3), a strain isogenic to KGY1120 but containing endogenous *CEF1* as a control for growth conditions; (ii) *prp3-1*, a positive control for a defect in the first step of splicing (24); (iii) *prp18* (*ts503*), a positive control for a defect in the second step of splicing (24); and (iv) *cdc28-1N*, a G<sub>2</sub> arrest (26) control to ensure that any observed splicing defects were not secondary to cell cycle arrest.

Initially, we assayed two intron-containing transcripts that are routinely used to analyze splicing defects in *prp* mutants *ACT1* and *RP51a* (24). In KGY1120 cells under *CEF1* repression, intron-containing forms of both transcripts steadily accumulated and peaked by 10 h (Fig. 5A). At this time, the abundance of each unspliced transcript was comparable to that observed in *prp3-1* cells (Fig. 5A). Next, we assayed two transcripts with unusual intron properties, *DYN2*, one of two transcripts in *S. cerevisiae* that contains two introns, and *DBP2*, the transcript with the largest intron in the *S. cerevisiae* genome (34). Similar to *ACT1* and *RP51a*, *DYN2* and *DBP2* unspliced transcripts accumulated progressively after *CEF1* repression (Fig. 5B). At 10 h, *DYN2* and *DBP2* pre-mRNAs had roughly equivalent or greater abundance, respectively, than the same pre-mRNAs in the *prp3-1* mutant.

Furthermore, the levels of mature *DYN2* decreased dramatically throughout the time course. Finally, we analyzed two intron-containing transcripts, *GLC7* and *TUB3*, that encode proteins required for mitotic progression (35–38). Unspliced *GLC7* accumulated throughout the time course of *CEF1* repression (Fig. 5C). Similarly, intron-containing transcripts of the α-tubulin encoding *TUB3* (36) increased throughout the time course (Fig. 5C). Taken together, these data indicate that *CEF1* is essential for pre-mRNA splicing *in vivo*.

## Discussion

CDC5 proteins are essential for G<sub>2</sub>/M progression in *S. pombe* (2), *S. cerevisiae* (3), and likely in higher eukaryotic cells (8). Despite a well documented role for these proteins in cell cycle progression, their biochemical function or functions are unknown. Consistent with CDC5 proteins containing homology to the well characterized DNA-binding factor and transcriptional activator c-Myb, we and others have demonstrated that the Myb domains of CDC5 proteins display affinity for double-stranded DNA (2, 4). Furthermore, the C terminus of hCDC5 fused to the GAL4-DNA-binding domain activated transcription in a transcriptional reporter assay (8). Beyond these properties, however, there is little evidence to suggest that CDC5 proteins function as transcription factors *in vivo*.

In fact, a recent report that hCDC5 associates with the spliceosome assembled *in vitro* (10) raised the possibility that CDC5 proteins are involved in pre-mRNA splicing. Our data confirm and extend this idea and show that mammalian CDC5 colocalizes with pre-mRNA splicing factors in mammalian cells, associates with core components of the splicing machinery in nuclear extracts, and interacts with the spliceosome throughout the splicing reaction *in vitro*. We have also shown that genetic depletion of *S. cerevisiae* *CEF1* blocks pre-mRNA splicing *in vivo*. These data provide evidence that eukaryotic cells require CDC5 proteins for pre-mRNA splicing. While an earlier version of this report was under review, similar conclusions were reached regarding the requirement for Cef1p in pre-mRNA splicing *in vivo* (39).

Although our data, as well as those of others (10, 39), are consistent with the notion that CDC5 proteins function directly in pre-mRNA splicing, we are unable to draw such a strong conclusion. To do so, it would have to be shown that immunodepletion of hCDC5 abrogated pre-mRNA splicing *in vitro*. Subsequent readdition of hCDC5 should then restore pre-mRNA splicing to the depleted extract. Although we were able to remove the majority of hCDC5 from a HeLa-cell S100 extract by immunodepletion, ≈5% of the protein remained in the extract, and the depleted extract spliced β-globin (data not shown). We did observe, however, that preincubation of S100 extract with affinity-purified hCDC5N antibodies inhibited the splicing reaction, whereas equivalent amounts of control antibodies had no effect on the reaction (data not shown). That antibodies to hCDC5 inhibited pre-mRNA splicing *in vitro* corroborates the finding that antibodies to Cef1p inhibit pre-mRNA splicing in *S. cerevisiae* extracts (39). Thus, further investigations into hCDC5/Cef1p function are likely to reveal that these proteins play a direct role in the process of pre-mRNA splicing.

We have observed that mammalian CDC5 localized to the nucleus regardless of serum conditions. This result contrasts with a previous report indicating that epitope-tagged and overexpressed hCDC5 localized exclusively to the cytoplasm in serum-deprived cells (6, 8). There are at least two simple explanations for this discrepancy. First, whereas the previous study examined the localization of epitope-tagged cytomegalovirus-expressed hCDC5, we have examined the localization of endogenous CDC5. Second, both of these studies observed the localization of CDC5 at steady state. Because some splicing factors shuttle

continuously between the cytoplasm and the nucleus (40), the difference between our observations may result from nucleocytoplasmic shuttling of mammalian CDC5. Perhaps serum starvation causes cytoplasmic retention of overexpressed hCDC5, thereby explaining the observations reported in the previous study.

Recent data call attention to the idea that pre-mRNA splicing and cell cycle control are linked (reviewed in ref. 41). As examples, most *S. pombe prp* mutants display cell cycle phenotypes (42, 43); *S. pombe cdc28<sup>+</sup>*, a putative RNA helicase, is required for both pre-mRNA splicing and G<sub>2</sub>/M progression (44); and cyclin E associates with and phosphorylates protein components of the U2 snRNP (45). Several hypotheses have been formulated to explain how a protein would function simultaneously in cell cycle control and pre-mRNA splicing (for review see ref. 41). One model predicts that a block in pre-mRNA splicing depletes a transcript or transcripts necessary for cell cycle progression. Indeed, it has been suggested that the mitotic arrest of *S. cerevisiae prp22* mutants results from a failure to splice the mRNAs for *TUB1* and *TUB3* (46). Because we have found that unprocessed *TUB3* transcripts accumulate in cells lacking *CEF1*, these cells might also arrest during G<sub>2</sub>/M (3) because of a deficiency in  $\alpha$ -tubulin protein.

Because precursor mRNAs were the most abundant species to accumulate in cells lacking Cef1p, it is likely that CDC5 proteins are required for the first biochemical step of pre-mRNA splicing. Notably, genetic depletion of *CEF1* did not affect the abundance

of several intronless transcripts, indicating that *CEF1* function is not globally required for mRNA production (3). Because the function of *S. pombe cdc5<sup>+</sup>* is structurally and functionally conserved in eukaryotes (3), these data are consistent with the notion that all eukaryotic cells require CDC5 proteins for pre-mRNA processing. Consistent with this hypothesis, we have shown recently that *S. pombe cdc5p* is also required for pre-mRNA splicing *in vivo* and associates with a snRNP-containing complex (47). The CDC5 family of proteins represents a conserved subfamily of Myb-related proteins necessary for pre-mRNA splicing.

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- Nurse, P., Thuriaux, P. & Nasmyth, K. (1976) *Mol. Gen. Genet.* **146**, 167–178.
- Ohi, R., McCollum, D., Hirani, B., Den Haese, G. J., Zhang, X., Burke, J. D., Turner, K. & Gould, K. L. (1994) *EMBO J.* **13**, 471–483.
- Ohi, R., Feoktistova, A., McCann, S., Valentine, V., Look, A. T., Lipsick, J. S. & Gould, K. L. (1998) *Mol. Cell. Biol.* **18**, 4097–4108.
- Hirayama, T. & Shinozaki, K. (1996) *Proc. Natl. Acad. Sci. USA* **93**, 13371–13376.
- Stukenberg, P. T., Lustig, K. D., McGarry, T. J., King, R. W., Kuang, J. & Kirschner, M. W. (1997) *Curr. Biol.* **7**, 338–348.
- Bernstein, H. S. & Coughlin, S. R. (1997) *J. Biol. Chem.* **272**, 5833–5837.
- Groenen, P. M., Vanderlinden, G., Devriendt, K., Fryns, J. P. & Van de Ven, W. J. (1998) *Genomics* **49**, 218–229.
- Bernstein, H. S. & Coughlin, S. R. (1998) *J. Biol. Chem.* **273**, 4666–4671.
- Lipsick, J. S. (1996) *Oncogene* **13**, 223–235.
- Neubauer, G., King, A., Rappsilber, J., Calvio, C., Watson, M., Ajuh, P., Sleeman, J., Lamond, A. & Mann, M. (1998) *Nat. Genet.* **20**, 46–50.
- Ausubel, F. M., Brent, R., Kingston, R. E., Moore, D. D., Seidman, J. G., Smith, J. A. & Struhl, K., eds. (1997) *Current Protocols in Molecular Biology* (Wiley, New York).
- Nishimoto, T., Eilen, E. & Basilico, C. (1978) *Cell* **15**, 475–483.
- Huang, S., Mayeda, A., Krainer, A. R. & Spector, D. L. (1997) *Mol. Biol. Cell* **8**, 1143–1157.
- Olmsted, J. B. (1981) *J. Biol. Chem.* **256**, 11955–11957.
- Krude, T., Jackman, M., Pines, J. & Laskey, R. A. (1997) *Cell* **88**, 109–119.
- Gould, K. L., Moreno, S., Owen, D. J., Sazer, S. & Nurse, P. (1991) *EMBO J.* **10**, 3297–3309.
- Colwill, K., Feng, L. L., Yeakley, J. M., Gish, G. D., Cáceres, J. F., Pawson, T. & Fu, X. D. (1996) *J. Biol. Chem.* **271**, 24569–24575.
- Blencowe, B. J., Carmo-Fonseca, M., Behrens, S. E., Lührmann, R. & Lamond, A. I. (1993) *J. Cell Sci.* **105**, 685–697.
- Mayeda, A. & Krainer, A. R. (1999) *Methods Mol. Biol.* **118**, 309–314.
- Mayeda, A. & Krainer, A. R. (1999) *Methods Mol. Biol.* **118**, 315–322.
- Krainer, A. R., Maniatis, T., Ruskin, B. & Green, M. R. (1984) *Cell* **36**, 993–1005.
- Blencowe, B. J., Nickerson, J. A., Issner, R., Penman, S. & Sharp, P. A. (1994) *J. Cell Biol.* **127**, 593–607.
- Lerner, E. A., Lerner, M. R., Janeway, C. A., Jr., & Steitz, J. A. (1981) *Proc. Natl. Acad. Sci. USA* **78**, 2737–2741.
- Vijayraghavan, U., Company, M. & Abelson, J. (1989) *Genes Dev.* **3**, 1206–1216.
- Blanton, S., Srinivasan, A. & Rymond, B. C. (1992) *Mol. Cell. Biol.* **12**, 3939–3947.
- Piggott, J. R., Rai, R. & Carter, B. L. (1982) *Nature (London)* **298**, 391–393.
- Guthrie, C. & Fink, G. R., eds. (1991) *Guide to Yeast Genetics and Molecular Biology* (Academic, San Diego), Vol. 194.
- Collart, M. A. & Oliviero, S. (1993) in *Current Protocols in Molecular Biology*, eds. Ausubel, F. M., Brent, R., Kingston, R. E., Moore, D. D., Seidman, J. G., Smith, J. A. & Struhl, K. (Wiley, New York), Vol. 2, pp. 13.12.1–13.12.50.
- Misteli, T. & Spector, D. L. (1998) *Curr. Opin. Cell Biol.* **10**, 323–331.
- Fu, X. D. & Maniatis, T. (1992) *Science* **256**, 535–538.
- Fu, X. D. & Maniatis, T. (1990) *Nature (London)* **343**, 437–441.
- Spector, D. L., Fu, X. D. & Maniatis, T. (1991) *EMBO J.* **10**, 3467–3481.
- Colwill, K., Pawson, T., Andrews, B., Prasad, J., Manley, J. L., Bell, J. C. & Duncan, P. I. (1996) *EMBO J.* **15**, 265–275.
- Spingola, M., Grate, L., Haussler, D. & Ares, M., Jr. (1999) *RNA* **5**, 221–234.
- Hisamoto, N., Sugimoto, K. & Matsumoto, K. (1994) *Mol. Cell. Biol.* **14**, 3158–3165.
- Schatz, P. J., Pillus, L., Grisafii, P., Solomon, F. & Botstein, D. (1986) *Mol. Cell. Biol.* **6**, 3711–3721.
- Schatz, P. J., Solomon, F. & Botstein, D. (1988) *Genetics* **120**, 681–695.
- Jacobs, C. W., Adams, A. E., Szaniszlo, P. J. & Pringle, J. R. (1988) *J. Cell Biol.* **107**, 1409–1426.
- Tsai, W. Y., Chow, Y. T., Chen, H. R., Huang, K. T., Hong, R. I., Jan, S. P., Kuo, N. Y., Tsao, T. Y., Chen, C. H. & Cheng, S. C. (1999) *J. Biol. Chem.* **274**, 9455–9462.
- Cáceres, J. F., Screaton, G. R. & Krainer, A. R. (1998) *Genes Dev.* **12**, 55–66.
- Burns, C. G. & Gould, K. L. (1999) in *Post-Transcriptional Regulation of Gene Expression and Its Importance to the Endocrine System*, ed. Chew, S. L. (Karger, Basel), Vol. 25, pp. 59–82.
- Urushiyama, S., Tani, T. & Ohshima, Y. (1996) *Mol. Gen. Genet.* **253**, 118–127.
- Potashkin, J., Kim, D., Fons, M., Humphrey, T. & Frendewey, D. (1998) *Curr. Genet.* **34**, 153–163.
- Lundgren, K., Allan, S., Urushiyama, S., Tani, T., Ohshima, Y., Frendewey, D. & Beach, D. (1996) *Mol. Biol. Cell* **7**, 1083–1094.
- Seghezzi, W., Chua, K., Shanahan, F., Gozani, O., Reed, R. & Lees, E. (1998) *Mol. Cell. Biol.* **18**, 4526–4536.
- Hwang, L. H. & Murray, A. W. (1997) *Mol. Biol. Cell* **8**, 1877–1887.
- McDonald, W. H., Ohi, R., Smelkova, N., Frendewey, D. & Gould, K. L. (1999) *Mol. Cell. Biol.* **19**, 5352–5362.
- Hanamura, A., Cáceres, J. F., Mayeda, A., Franza, B. R., Jr., & Krainer, A. R. (1998) *RNA* **4**, 430–444.