Primary rat embryo fibroblasts were transformed by a p53 mutant (alanine to valine change at amino acid 135) plus ras. This p53$^c^135$ mutant is temperature sensitive for a conformational change detected by the binding of a monoclonal antibody, PAb246, which recognizes the wild-type protein or the great majority of p53$^c^135$ at 32.5°C. At 37°C, both mutant and wild-type p53 conformational forms coexist in the cells, while at 39.5°C, the majority of the p53$^c^135$ in the cell is in a mutant conformation not recognized by PAb246 antibody. At 39.5°C, the mutant p53 is localized in the cytoplasm of the cell. At 32.5°C, the p53 protein enters the nucleus and stops the growth of these cells. At 37°C where there is a mixture of mutant and wild-type p53, the wild-type p53 protein is in a complex with hsc70 and mutant p53 protein in the cytoplasm of the cell during G$_1$. This wild-type protein enters the nucleus as the cells enter the S-phase of the cell cycle. At 32.5°C, the cells stop replication and arrest at the G$_1$/S border. After 48 hr at 32.5°C, 91% of the cells are in the G$_1$ fraction of the cell cycle. The S-phase cells appear to be immune to the p53 negative regulation of growth until they enter the next G$_1$ period. These data strongly suggest that mutant p53 proteins in transformed cells act to sequester the wild-type p53 protein in an hsc70–p53 complex, which resides in the cytoplasm during the stage of the cell cycle, G$_1$, when nuclear wild-type p53 would normally act to regulate cell growth and progression through the cycle. In this way, mutant p53 proteins can act in a trans-dominant fashion to overcome growth regulation by the wild-type p53 allele and protein in a cell.

[Key Words: p53 mutant; rat embryo fibroblasts; transformed cells]

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mune how mutant p53 protein in a cell can overcome the regulation of endogenous wild-type p53 proteins and transform a cell. Two murine p53 mutants, p53val135 [an alanine to valine change at residue 135] [Eliyahu et al. 1985; Hinds et al. 1989] and p53H215 [an insertion of four amino acids at residue 215] [Tan et al. 1986] were employed to transform primary rat embryo fibroblasts in cooperation with the ras oncogene. High levels of these murine mutant p53 proteins were expressed in these cells along with lesser amounts of the rat p53 protein [Finlay et al. 1989]. A polymerase chain reaction [PCR] amplification of the reverse transcriptase copy of the rat p53 mRNA from both p53val135 and p53H215 plus ras-transformed cells was sequenced. This showed that the endogenous rat p53 mRNA and protein was a wild-type copy of p53. The murine mutant p53 protein, the wild-type rat p53 protein, and the cellular heat shock protein hsc70 were found in an oligomeric protein complex in these cells [Finlay et al. 1989]. It has been suggested that the murine mutant p53 protein could sequester the wild-type rat p53 protein and inactivate its ability to negatively regulate growth in these cells [Baker et al. 1989; Finlay et al. 1989].

In agreement with Michalovitz, et al. [1990], the p53val135 mutant is shown to be temperature sensitive. At 32.5°C, the great majority of this p53 protein is in the wild-type conformation (PAb246⁺, PAb240⁻), whereas at 39.5°C, most of this p53 protein is in the mutant conformation (PAb246⁻, PAb240⁺). At 39.5°C, during the G1 phase of the cell cycle, the transformed cells replicate rapidly, whereas at 32.5°C, cell growth stops and the cells are blocked at the G1/S border of the cell cycle. Those cells in S-phase continue through the cell cycle to the next G1/S border. At 39.5°C, the p53 mutant protein is localized in the cytoplasm, while at 32.5°C, the p53 protein is in the nucleus. At 37°C where both mutant and wild-type p53 coexist in these transformed cells, the mutant p53–wild-type p53–hsc70 complex [Hinds et al. 1987] is in the cytoplasm during the G1 phase of the cell cycle. The wild-type p53 proteins move into the nucleus at the start of the S phase. Thus, wild-type p53 appears to be sequestered in the cytoplasm during a critical time, G1, when it could act in the nucleus to block entry into the S phase. As such, mutant p53 acts dominantly to sequester the wild-type p53 protein in a place where it cannot function. The p53H215 mutant protein, which is not temperature sensitive, remains in the cell cytoplasm throughout the cell cycle [PAb⁻, PAb240⁺, hsc70⁺ complex] [Hinds et al. 1987].

**Table 1.** Detection of wild-type and mutant forms of p53 in p53 plus ras-transformed cells at 39.5°C, 37°C, and 32.5°C

| Temperature—percent reactive | Monoclonal antibody | A1-5 cells (p53val135) | T101-4 cells (p53H215) |
|------------------------------|---------------------|------------------------|------------------------|
|                              | detect | 39.5°C | 37°C | 32.5°C | 39.5°C | 37°C | 32.5°C |
| PAb246 | 27%   | 36%   | 82%  | PAb246 | 0%    | 0%   | 0%    |
| PAb240 | 73%   | 65%   | 18%  | PAb240 | 100%  | 100% | 100%  |

Soluble protein extracts were prepared from p53val135 plus ras (A1-5 cells) or p53H215 plus ras (T101-4 cells) labeled with [35S]methionine for 1 hr at the indicated temperature. The p53 protein was immunoprecipitated with PAb246 (wild type), PAb240 (mutant), or PAb421 (total p53), and the level of p53 was quantitated by densitometer tracings of autoradiograms from SDS-polyacrylamide gels of the immunoprecipitates [Finlay et al. 1989]. In replicate experiments, scintillation counting was employed to quantitate these results with the levels remaining essentially the same.
suggest that the \(p53^{\text{val135}}\) allele is temperature sensitive, in agreement with Michalovitz et al. (1990).

When Al-5 cells (\(p53^{\text{val135}}\)) and T101-4 cells (\(p53^{\text{KH215}}\)), grown at 37°C, were stained with fluorescent antibodies that detect the wild-type p53 (PAb246) or both mutant and wild-type p53 (PAb421), the PAB246 antibody detected nuclear p53 in a punctate pattern in Al-5 cells (at 37°C, 35% of the p53 is wild type). PAB246 failed to detect any wild-type p53 in \(p53^{\text{KH215}}\) transformed cells (T101-4; Fig. 1). The PAB421 antibody detects p53 predominantly in the cytoplasm of T101-4 cells and both nuclear and cytoplasmic p53 in Al-5 cells.

Because the great majority of p53 in these cells derives from the murine p53 transgene (Finlay et al. 1988), it is unlikely that these fluorescent probes detect the rat endogenous p53 protein. To determine whether the rat p53 protein was a mutant or wild-type p53, we amplified a cDNA copy of the rat p53 by PCR and sequenced it directly (see Materials and methods). Both the Al-5 and the T101-4 cells had wild-type rat p53 mRNAs. Thus, these transformed cells have a mixture of mutant p53 and wild-type p53, whereas the mutant form is usually in the highest concentration because it has a longer half-life (Reich and Levine 1984).

**Intracellular localization of \(p53^{\text{val135}}\) changes with incubation temperature**

The murine \(p53^{\text{val135}}\) mutant has recently been shown to be temperature-sensitive for transforming activity. At 32.5°C, it is unable to cooperate with ras in transformation assays, yields no foci, and behaves in a wild-type manner (Michalovitz et al. 1990). Al-5 cells contain this temperature-sensitive mutant, whereas T101-4 cells express wild-type rat p53 mRNAs. Thus, these transformed cells have a mixture of mutant p53 and wild-type p53, whereas the mutant form is usually in the highest concentration because it has a longer half-life (Reich and Levine 1984).

The Al-5 cells at 37°C contain a mixture of murine mutant p53 and wild-type murine and rat wild-type p53 in the cell. The nuclear and cytoplasmic localization of these p53 molecules (Figs. 1 and 2) suggested the possibility that different stages of the cell cycle might affect the location of p53 in the cell. To test this, Al-5 cells were synchronized by selective mitotic detachment. These cultures were pulse-labeled with \[^{3}H\]thymidine to determine the time of entry into S phase or stained by indirect immunofluorescence using PAB426 to examine the intracellular location of the wild-type p53 (conformation) in these cells. The results of this experiment are presented in Fig. 3. \[^{3}H\]Thymidine incorporation (i.e., entry into S phase) begins 3 hr after mitosis (the generation time of these cells was 13 hr) and reaches a maximum by 4 hr. Mitotic cells all have wild-type (but not mutant) p53 attached to the chromosomes in the cell (zero time). At 1–2 hr, virtually all of the cells contained p53 (wild-type p53) in the cytoplasm (Fig. 3). As the cells entered S phase, the percentage of cells with wild-type p53 in the nucleus increased in parallel with the increase in \[^{3}H\]thymidine incorporation. By 4 hr, most cells had the punctate staining pattern of nuclear p53, which continued through S phase. Throughout G\(_{1}\), Al-5 cells do not show any nuclear staining for p53 with PAB426 (Fig. 3, wild-type) or PAB421 (total p53, not shown). The wild-type conformation of p53 is excluded from the cell nucleus during the G\(_{1}\) phase of the cell cycle. It enters the nucleus as the cells enter S phase. Interestingly, this is
Figure 3. Determination of initiation of DNA replication and intracellular distribution of p53\textsuperscript{val135} in synchronized A1-5 cells. (A) A1-5 cells were synchronized by mitotic detachment, incubated at 37°C, and analyzed for TCA-precipitable \([\text{H}]\)-thymidine incorporation (\(\bullet\)). The percent of p53 positive nuclei was determined after indirect immunofluorescence using PAb246 (\(\circ\)). (B) Synchronized A1-5 cells grown on coverslips were stained with PAb246 at hourly intervals. A representative field for each sample time point is presented. Phase-contrast photomicrographs are included for reference.

the same pattern of cell localization reported for hsp70 in a cell [Milarski and Morimoto 1986]. TlOl-4 cells contain no wild-type murine p53 and has all of its p53 (PAb421) in the cytoplasm throughout the cell cycle (results not shown).

**p53 localization affects cell growth**

p53 is a nuclear protein and has been implicated in growth control of the cell. In A1-5 cells, the subcellular localization of p53\textsuperscript{val135} can be varied by changing the incubation temperature. On this basis, the effect of a p53 nuclear versus cytoplasmic compartmentalization on cell growth was investigated. Doubling times of six different cell lines were determined at 32.5°C, 37°C, and 39.5°C (Table 2). The following cell lines were studied: A1-5; B4, an independently isolated cell line also expressing the p53\textsuperscript{val135} mutant; TlOl-4; TlOl-1, an independently isolated clone also containing the p53\textsuperscript{KH215} mutant; Rat-1, an immortalized rat embryo fibroblast cell line; and secondary rat embryo fibroblasts, which are not immortalized. Growth of A1-5 (Fig. 4) and B4 cells [Table 2] that express p53\textsuperscript{val135} was arrested at 32.5°C. All other cell lines slow their growth considerably at 32.5°C but continue to grow and divide. Viability of A1-5 cells maintained at 32.5°C, as assayed by trypan blue exclusion, is 100% even when incubated for periods of up to 1 week at the low temperature. These cells continue to synthesize proteins and are capable of incorporating \([\text{H}]\)methionine at rates nearly equal to that of cells grown at 37°C (results not shown). Finally, the rate of \([\text{H}]\)thymidine incorporation for A1-5 cells grown at

| Cell line | Transforming p53 mutant\(^a\) | Doubling time [hr] |
|-----------|-------------------------------|--------------------|
|           |                               | 39.5°C 37°C 32.5°C\(^b\) |
| A1-5      | p53\textsuperscript{val135}   | 13 13 *            |
| B4        | p53\textsuperscript{val135}   | 15 18 *            |
| TlOl-1    | p53\textsuperscript{KH215}    | 18 20 54           |
| TlOl-4    | p53\textsuperscript{KH215}    | 14 22 41           |
| Rat-1     | NA                            | 13 14 44           |
| REF       | NA                            | 18 19 31           |

\(^a\) (NA) Not applicable.  
\(^b\) * Cells remained viable but did not divide.

Figure 4. Growth curves of A1-5 cells at 39.5°C (\(\triangle\)), 37°C (\(\bullet\)), and 32.5°C (\(\square\)).
32.5°C drops sevenfold when shifted from 37°C to 32.5°C, whereas Tl01-4 cells continue at only slightly reduced levels (data not shown). Therefore, growth inhibition of A1-5 cells by incubation at 32.5°C does not appear to result from nonspecific toxicity of p53野生型. Since [3H]thymidine incorporation was reduced at the low temperature, it was expected that DNA synthesis might be impaired.

Inhibitory effects of p53 on cell growth are restricted to $G_1$

In A1-5 cells grown at 37°C, p53野生型 had a cytoplasmic location during $G_1$ (Fig. 2). Incubation at 32.5°C resulted in nuclear localization of p53野生型 accompanied by cell growth inhibition (Figs. 2 and 4). As a next step, growth-inhibited A1-5 cells at 32.5°C were tested to determine whether they were blocked at a specific point in the cell cycle. A1-5 cells maintained for 48 hr at 39.5°C, 37°C, and 32.5°C were lysed and the nuclei were treated with ethidium bromide and analyzed by flow cytometry (Fig. 5, Table 3). At 32.5°C, 91% of the cells were in $G_1$, less than one-tenth of all the cells are present in $S$ and $G_2$. In contrast, the profiles for cells growing at 37°C and at 39.5°C are indicative of normal exponential growth. Therefore, it appears likely that the progression of A1-5 cells incubated at 32.5°C is impeded at the $G_1$ phase of the cell cycle.

The results in Figure 5 provide a static profile of the cells maintained at the three temperatures. It would be informative to explore the changes that occur when a rapidly growing population of A1-5 cells maintained at high temperature are shifted to low temperature. Log-phase cultures of A1-5 cells growing at 39.5°C were shifted to 32.5°C and analyzed by flow cytometry (Fig. 6). This demonstrated that cells in $G_1$ did not advance through the cell cycle. However, the shift from 39.5°C to 32.5°C did not retard the progression of cells already in $S$ phase through the remainder of the cycle (Fig. 6). The block was coincident with a rapid shift in p53野生型 localization from cytoplasm to nucleus, as seen by immunofluorescent staining. Interruption of cell cycle progression by shifting to low temperature appears to affect only those cells that have not yet entered $S$ phase. These data are most consistent with the possibility that p53野生型 functions at 32.5°C during $G_1$ as a growth inhibitor and

### Table 3. Percent of A1-5 cells in $G_1$, $S$, and $G_2$ at three incubation temperatures

| Incubation temperature (°C) | $G_1$ | $S$ | $G_2$ |
|-----------------------------|------|-----|------|
| 39.5                        | 51   | 31  | 18   |
| 37                          | 35   | 42  | 23   |
| 32.5                        | 91   | 4   | 5    |

Figure 5. Histograms of A1-5 cells incubated for 48 hr at 39.5°C, 37°C, or 32.5°C. Nuclei were prepared and analyzed by flow cytometry. Relative DNA content is represented on the x-axis; cell numbers are on the y-axis. Cell cycle fractions $G_1$, $S$, and $G_2$ are indicated; the percent of total cells in each fraction is presented in Table 1.

Figure 6. Histograms of A1-5 cells shifted down from 39.5°C to 32.5°C. Samples were analyzed by flow cytometry. Histograms are aligned relative to their $G_1$ peaks. The time point of each sample is indicated.
DNA replication. Once replication has begun, the cell cycle continues unhindered until G\textsubscript{i} is encountered once again.

Discussion

Normal or untransformed cells contain very low levels of p53 [Linzer and Levine 1979], which has made it very difficult to study this protein in these cells. Expressing higher levels of wild-type p53 appears to suppress cell growth [Baker et al. 1990; Diller et al. 1990; Mercer et al. 1990; Michalovitz et al. 1990], further complicating the study of p53 function in these cells. Mutant p53, on the other hand, cooperates with the ras oncogene to transform cells [Eliyahu et al. 1984; Parada et al. 1984], and high levels of mutant p53 proteins are found in these cells. Several biochemical and immunological criteria have been used to distinguish between the mutant and wild-type p53 proteins [Levine 1990]: (1) PAb246 binds to wild-type but not most mutant p53 proteins [Tan et al. 1986; Yewdell et al. 1986]; (2) PAb240 binds to many mutant proteins but not to wild-type p53 proteins [Bartek et al. 1990]; (3) many mutant p53 proteins form a complex with hsp70 in the cell [Hinds et al. 1987]; (4) the half-life is longer and the cellular concentrations of mutant p53 proteins are higher than wild-type p53 [Finlay et al. 1989]. Recently, Michalovitz et al. [1990], have shown that the p53\textsuperscript{val1355} mutant is temperature sensitive. In this study we confirmed that observation with each of the criteria stated above. At 32.5°C the great majority of the p53\textsuperscript{val1355} protein is wild type, at 39.5°C most of it is mutant, and in A1-5 cells at 37°C a mixture of these two forms has been shown previously to coexist [Hinds et al. 1987; Finlay et al. 1988]. This mixture of mutant and wild-type forms of p53 in A1-5 cells may be a good model system to ask the question, how can mutant p53 have a trans-dominant phenotype, eliminating the negative regulatory functions of wild-type p53 [Finlay et al. 1989]? The hypothesis put forth to explain this observation was that murine mutant p53–hsp70 and murine and rat wild-type p53 in these cells was in a complex that inactivated the p53 wild-type function. The results presented in this report provide a mechanism to explain the dominant loss of function mediated by mutant p53. First, the results of a shift of A1-5 cells from 39.5°C to 32.5°C strongly suggests that wild-type p53 acts in G\textsubscript{i} to block cells from entering the S phase of the cell cycle. Second, in A1-5 cells at 37°C wild-type p53 is sequestered, likely by its association with mutant p53–hsp70 [Hinds et al. 1987] in the cytoplasm in G\textsubscript{i} and cannot act in the nucleus at that time. When wild-type p53 returns to the nucleus in S phase, possibly as a result of the movement of hsp70 into the nucleus at that time [Miaarski and Morimoto 1986], it is too late in the cell cycle to function and block progression through the cell cycle.

In addition to providing a mechanism for mutant p53 to act in a dominant-negative (loss of function) fashion, the results reported here suggest that wild-type p53 acts during the G\textsubscript{i} phase of the cell cycle. This result is in good agreement with Diller et al. [1990] and Michalovitz et al. [1990]. The only caution in all of these observations is that the wild-type p53 levels required to observe these effects are higher than levels normally seen in cells. This is really an overexpression phenotype. This communication is the first report of the genotype of the p53 alleles from rat cells transformed by mutant murine p53 plus ras. In both p53\textsuperscript{val1355} and p53\textsuperscript{K4215} plus ras-transformed...
cell lines, the endogenous rat p53 is wild type. It remained possible that the endogenous rat cell p53 gene had sustained a mutation to produce a permanent transformed cell line. Many permanent cell lines in culture have been shown to have mutant p53 genes and proteins (Finlay et al. 1988; Levine 1990). On the contrary, the two different mutant p53 plus ras rat embryo fibroblast cell lines examined here each had wild-type endogenous rat p53 mRNAs, consistent with the trans-dominant inactivation of a negative regulator by mutant murine p53 proteins. This is based on the assumption that p53<sup>val135</sup> in its wild-type conformation is a good model for studying the behavior of the wild-type rat p53 protein.

Finally, there are some suggestions in this study that p53 does not act (in a mutant or wild-type form) only as a negative regulator of cell growth. First, the punctate pattern observed in the nuclei of A1-5 cells grown at 37°C and stained with PAb246 (wild-type p53, Fig. 3) is not observed in these same cells at 32.5°C (Fig. 2). The wild-type p53 protein in cells that are actively replicating their DNA [having allowed cells to enter S phase] behaves differently [has a different nuclear distribution] in S phase than when it is in the nucleus in G1 [at 32.5°C]. Similarly, the wild-type p53 protein [and not the mutant form] binds tightly to metaphase chromosomes [Fig. 3B]. This has also been observed with wild-type p53 in nontransformed cells [rat embryo fibroblast cells; results not presented]. Third, the results presented here would predict that wild-type p53 proteins that failed to enter the nucleus [because they have lost their nuclear localization signals] would also fail to suppress transformation of cells by p53 plus ras or E1A plus ras. That is, p53 must mediate its suppressive effects by acting in the nucleus (because they have lost their nuclear localization signals) would predict that wild-type p53 proteins that failed to enter the nucleus [because they have lost their nuclear localization signals] would also fail to suppress transformation of cells by p53 plus ras or E1A plus ras. That is, p53 must mediate its suppressive effects by acting in the nucleus of a cell. This has recently been tested (G. Shaulsky, N. Goldfinger, M. Tosky, A. Levine, and V. Rotter, in press) and is correct. Wild-type p53 proteins without a nuclear localization signal fail to suppress transformation of cells in culture. On the other hand, one might predict that mutant p53 proteins that are activated for transformation, but lack a nuclear localization signal, would transform cells in culture by cooperating with ras. In this case, being sequestered in the cytoplasm seems to constitute a mechanism to effect a dominant loss of function for the endogenous wild-type p53. This hypothesis is being tested.

**Materials and methods**

**Cells and antibodies**

A1 and B4 cells are primary rat embryo fibroblasts transformed by transfection with activated ras [T24] (Goldfarb et al. 1982) and plasmid LTR p53CG, which directs overexpression of p53<sup>val135</sup> (Finlay et al. 1988). These two cell lines were derived from separate transformation experiments. A1-5 cells were derived from the A1 line by limiting dilution and were used throughout these studies, except where indicated. Cell lines T101-1 and T101-4 are rat embryo fibroblasts transformed by transfection with activated ras [T24] and plasmid MSUKH215, which directs the overexpression of p53<sup>val135</sup>. These two cell lines are independent clones from the same transformation experiment and have been described in detail elsewhere (Finlay et al. 1988). Rat-1 cells were derived from methylcholanthrene-treated Fisher rat embryo fibroblasts (Freeman et al. 1973). Rat embryo fibroblasts are secondary or tertiary cultures. Culture medium for all cells was Dulbecco's modified Eagle medium (DMEM) supplemented with 10% heat-inactivated fetal calf serum (FCS) at 37°C, except where stated otherwise.

PAb246 is a pan-specific monoclonal antibody that detects p53 from all species (Harlow et al. 1981). PAb246 recognizes a specific conformational epitope present only on wild-type murine p53 (Yewdell et al. 1986). PAb240 is specific for mutant murine p53 proteins (Gannon et al. 1990).

**Indirect immunofluorescent staining**

Coverslips with attached cells were rinsed three times in ice cold PBS [137 mM NaCl, 2.7 mM KCl, 4.3 mM Na,HPO<sub>4</sub> • H<sub>2</sub>O, 1.4 mM KH<sub>2</sub>P0<sub>4</sub> • pH 7.3]. The cells were fixed in methanol/acetone [50 : 50] for 2 min at −20°C and rehydrated with cold PBS. Undiluted monoclonal antibody cell culture supernatants were applied to the coverslips and incubated in a humidified atmosphere for 2 hr at 37°C. After washing three times with cold PBS, FITC-conjugated sheep [Fab']2 fragment anti-mouse IgG (heavy and light chains) (Cooper Biomedical, Inc., Malvern, PA) diluted 1 : 50 was added. The coverslips were incubated for 30 min at 37°C and washed three times with cold PBS. Samples were mounted on glass slides with mounting solution (1 x PBS, 90% glycerol, 0.1% p-phenylendiamine). A Zeiss model IIIRS microscope was used for all observations and photographs.

**PCR amplification and DNA sequencing**

Total RNA was isolated as described by Maniatis et al. (1982). cDNA was generated essentially as described by Kawaski (1990), using 5 μg of total RNA, except that oligo[d(T)]<sub>18</sub> was used to prime the reverse transcriptase reaction. On completion of the reverse transcriptase reaction, the enzyme was inactivated by heating to 94°C for 56 min. The PCR was performed by adding 80 μl of 1 × PCR buffer [50 mM KCl, 2.5 mM MgCl<sub>2</sub>, 10 mM Tris-Cl (pH 8.3), 200 μM dNTPs, 0.1 mg/ml BSA], 10 pmol of 5′ primer [5′-CTGGAAGACTCGTATACTGTCCGAGAT-3′], and 5 pmol of 3′ primer [5′-AGACCCCGAGTGATCTCTCAAGGTA-3′], and 5 units of Taq polymerase (Cetus Emeryville, CA) directly to the cDNA reaction mixture. Rat p53 cDNA was amplified by 30 cycles (1 min at 94°C, 2 min at 45°C, and 3 min at 72°C). Single-stranded DNA from this reaction was amplified by asymmetric PCR as described by McCabe (1990) and sequenced directly by the dideoxy chain-termination method (Sanger et al. 1977).

**Growth curves**

Log-phase cells grown to no more than 80% confluence were fed fresh media 12–24 hr before they were replated at a density of 5 × 10<sup>4</sup> cells/60-mm plate. The cells were allowed to attach at 37°C for 8–12 hr and were left at 37°C or shifted to either 32.5°C or 39.5°C. The initial sample was taken 12–24 hr later (approximately one doubling time). Cell numbers were determined using an Electrozone model 112LTSNBCD/ADC (Particle Data, Inc). All time points were performed in duplicate. Doubling times were calculated by fitting data points to an exponential curve using regression analysis.

**Mitotic detachment**

Cells for mitotic detachments were fed fresh culture media 15
hr prior to the experiment. All cells were grown at 37°C to 90% confluency in Corning T150 flasks (Corning, Chicago, IL). Miotic cells were harvested by replacing culture media with 6 ml of fresh suspension-modified minimum essential medium (SMEM) and 10% FCS, prewarmed to 37°C. After continued incubation at 37°C for 30 min, the flasks were shaken rapidly for 15 sec. The media now containing mitotic cells were pooled and centrifuged at low speed (4°C). Cell pellets were resuspended in culture medium (Terasima and Tolmach 1963). This procedure regularly yielded populations that consisted of 70-80% mitotic cells.

Cell cycle progressions

For determination of S phase, synchronized cells obtained by mitotic detachment were aliquoted into 35-mm plates and incubated at 37°C. At hourly intervals, cells were labeled by replacing culture medium with prewarmed [37°C] medium containing 50 μCi/ml [3H]-thymidine (73.7 Ci/mmol, Dupont/ New England Nuclear, Boston, MA). Incubation was continued for 15 min at 37°C. Samples were harvested by rinsing cell monolayer with ice-cold PBS and applying 1 ml of lysis buffer [10% SDS, 10 mM EDTA (pH 8.0)] directly to the plate. TCA-precipitable radioactivity was determined for each sample by scintillation counting in a Beckman LS5000TD.

Synchronized cells for indirect immunofluorescence studies were resuspended in 5 ml of culture medium and applied to poly-L-lysine-coated coverslips for 10 min on ice. The coverslips were placed in 35-mm plates with medium and incubated at 37°C. Coverslips were removed at hourly intervals and prepared for immunofluorescence as described.

For analysis of cell cycle progression using flow cytometry were trypsinized from 60-mm plates and lysed in FACS buffer [3.4 mM Na citrate, 10 mM NaCl, 0.1% NP-40, 75 mM ethidium bromide]. The cell nuclei were either analyzed immediately, or pelleted, resuspended in freezing buffer [10 mM Tris-HCl (pH 7.5), 10% sucrose], and stored at -70°C for analysis at a later date (Sherley and Kelly 1988).

Temperature shift assays and flow cytometry analysis

Cells were plated at a density of 5 × 10^3 cells/10-cm plate and incubated at 37°C overnight to allow attachment. For shift-down experiments, plates were incubated at 39.5°C for 24 hr and shifted to 32.5°C. Samples were taken at appropriate intervals and prepared for analysis by flow cytometry as described (Sherley and Kelley 1988). Cells for indirect immunofluorescent staining were grown on coverslips and subjected to the same temperature-shift protocols.

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