Disruption of Nuclear Lamin Organization Alters the Distribution of Replication Factors and Inhibits DNA Synthesis

Timothy P. Spann,* Robert D. Moir,* Anne E. Goldman,* Reimer Stick,‡ and Robert D. Goldman*

*Department of Cell and Molecular Biology, Northwestern University Medical School, Chicago, Illinois 60611; and ‡ Institut fuer Biochemie und Molekulare Zellbiologie der Universitat Goettingen, D-37073 Goettingen, Germany

Abstract. The nuclear lamina is a fibrous structure that lies at the interface between the nuclear envelope and the nucleoplasm. The major proteins comprising the lamina, the nuclear lamins, are also found in foci in the nucleoplasm, distinct from the peripheral lamina. The nuclear lamins have been associated with a number of processes in the nucleus, including DNA replication. To further characterize the specific role of lamins in DNA replication, we have used a truncated human lamin as a dominant negative mutant to perturb lamin organization. This protein disrupts the lamin organization of nuclei when microinjected into mammalian cells and also disrupts the lamin organization of in vitro assembled nuclei when added to Xenopus laevis interphase egg extracts. In both cases, the lamina appears to be completely absent, and instead the endogenous lamins and the mutant lamin protein are found in nucleoplasmic aggregates. Coincident with the disruption of lamin organization, there is a dramatic reduction in DNA replication. As a consequence of this disruption, the distributions of PCNA and the large subunit of the RFC complex, proteins required for the elongation phase of DNA replication, are altered such that they are found within the intranucleoplasmic lamin aggregates. In contrast, the distribution of XMCM3, XORC2, and DNA polymerase α, proteins required for the initiation stage of DNA replication, remains unaltered. The data presented demonstrate that the nuclear lamins may be required for the elongation phase of DNA replication.

The nuclear lamin proteins form a fibrous structure, termed the nuclear lamina, which is concentrated at the nucleoplasmic face of the nuclear envelope (40). The lamins are also found in nucleoplasmic foci, the distribution of which is related to the cell cycle (7, 18, 32, 39, 48). The lamins are highly conserved proteins that are closely related to cytoplasmic intermediate filament (IF) proteins and, as such, are classified as type V IF (51). In vertebrates, as many as five lamin proteins have been reported. These are divided into two types, A and B, based on criteria such as expression patterns and exon positions (53). B-type lamins are expressed in all cells, while the A-type lamins are expressed in differentiated cells (40). In Xenopus laevis, there are five or more lamins, also showing cell type-specific expression patterns (3, 40, 54). As is the case with cytoplasmic IF, the lamins have a central rod domain that forms an α helix, composed of heptad repeats. The rod domains are primarily responsible for the higher order lamin–lamin interactions that govern lamin assembly (20). The two non-α-helical end domains are also involved in assembly and may interact with other nuclear structures (see 40, for detailed discussion).

In addition to providing mechanical support to the nucleus and influencing its shape and volume, the lamins appear to interact with other nuclear components and thereby may influence a number of nuclear processes (40). For example, the lamins interact with chromatin in vitro and probably in vivo (2, 16, 22, 47). Through this interaction, the lamins may be involved in DNA replication. For example, lamin B is associated with replicating chromatin in mammalian cells (39). During S phase, lamin B appears not only at the nuclear periphery, but also in nucleoplasmic foci that frequently coincide with sites of bromodeoxyuridine incorporation and the location of the DNA replication factor, PCNA (39). It has also been suggested on morphological grounds that a filamentous network of lamins acts as a scaffold for “DNA replication factories” (24, 25).

Further evidence for a role of the nuclear lamins in DNA replication comes from the nuclear assembly system prepared from Xenopus laevis eggs (3, 4, 30). In this sys-
when DNA or chromatin is added to interphase egg extracts. These nuclei carry out processes such as nuclear import, lamin assembly, and DNA replication. The presence of highly concentrated, soluble, nuclear components in the extract makes the Xenopus system particularly useful for biochemical manipulations. Proteins from a wide range of species, including yeast and humans, have been added to interphase and mitotic extracts to examine their function in the regulation of the cell cycle (41). In addition, immunodepletion of specific proteins from these extracts has been used to determine their involvement in nuclear functions. For example, when the major endogenous lamin (lamin B3) is immunodepleted from interphase extracts, nuclei form, but they cannot replicate their DNA (34, 45). Furthermore, when the eluted lamin B3 is added back to depleted extracts, nuclear DNA replication is restored (17). These results suggest that nuclear lamins play a role in DNA replication, although it is unclear how or at what stage DNA synthesis is blocked under these experimental conditions.

The Xenopus nuclear assembly system has also been useful in characterizing other proteins involved in regulating DNA replication. Immunodepletion experiments involving the removal of XMCM3 and XORC2, as well as a number of other factors, have demonstrated that they are essential for DNA replication. XMCM3, for example, is a putative component of the licensing factor that is thought to limit replication to one round for each cell cycle. It binds to chromatin early in the process of nuclear assembly, before the nuclear membrane forms (10, 29, 33). XORC2 is the Xenopus homologue of the yeast protein, ORC2 (9). In yeast, this protein is required to initiate DNA synthesis and is part of a complex that binds to origins of replication (55). In Xenopus, XORC2 binds to chromatin before nuclear envelope formation and appears to be involved in the initiation of DNA synthesis (9). Other proteins involved in replication have also been identified as constituents of the Xenopus system, including PCNA (26), a required cofactor of DNA polymerase δ. This polymerase is responsible for the elongation phase of DNA replication (59).

The immunodepletion approach has been extremely valuable in defining roles for the lamins as well as for other proteins in DNA synthesis, but this method has several limitations. For example, in the case of the nuclear lamins, it is difficult to completely immunodeplete lamin proteins (31). In addition, any effects seen after immunodepletion may not be due to the removal of targeted antigenic components, but rather to the immunoprecipitation of bound, associated proteins. Immunodepletions of XMCM3, XORC2, and lamin B3 all result in the specific removal of other proteins, in addition to the antigen targeted by the antibody (9, 10, 17, 33).

To define more precisely the functions of nuclear lamins in nuclear assembly and DNA replication, we have developed a method that avoids some of the pitfalls inherent in the immunodepletion techniques. Our approach uses a human lamin mutant as a dominant negative disrupter of the nuclear lamin organization in two experimental systems. When this mutant protein is microinjected into mammalian cells or added to the Xenopus nuclear assembly extract, the lamin organization is disrupted. The mutant protein as well as the endogenous lamins appear to colocalize in nucleoplasmic aggregates, with little or no detectable lamin protein at the nuclear periphery. We have concentrated our efforts on defining the effects of this disruption in nuclei assembled in Xenopus extracts, since more nuclei can be studied under conditions permitting coordinated biochemical and morphological assays. As in the case for Xenopus nuclei assembled after lamin B3 immunodepletion (34, 45), the disrupted nuclei described in this study cannot complete DNA replication. However, because the lamins are retained within the nucleus, we are able to examine their distribution relative to other DNA replication markers. We find that, as a consequence of the disruption, there is an altered distribution of proteins specifically involved in the elongation phase, but not in the initiation phase of DNA replication.

**Materials and Methods**

**Expression of Human Lamins in Escherichia coli**

Full-length human lamin A (LA) and ΔNLA were cloned in a pET-derived vector and expressed in the NovaBlue (DE3) strain of E. coli (Novagen, Madison, WI). ΔNLA lacks the first 33 amino acids of human lamin A. Both of the expressed proteins were purified by ion-exchange chromatography as described previously (38). The protein in column buffer (6 M urea, 25 mM Tris, 2 mM EDTA, and 1 mM DTT) was dialyzed against PB (300 mM NaCl, 25 mM Tris Base, pH 9, and 1 mM DTT). After dialysis, SDS-PAGE analysis of the resulting protein solution showed the presence of one major band of protein. In the case of ΔNLA, the protein had a molecular mass of 69 kD (Fig. 1 a). This 69-kD protein reacted with a rabbit polyclonal antibody directed against human lamins A/C as demonstrated by immunoblotting (Fig. 1 b). SDS-PAGE and blotting analyses were carried out as described in (39). The relatively minor bands seen in the immunoblot (Fig. 1 b) are due to a small amount of proteolysis that is present in all preparations of nuclear lamins (38). The same gel profiles are seen for the wild-type LA protein, but the apparent molecular mass is 72 kD (not shown). The protein solutions were aliquoted and stored at −80°C at a final concentration of 2 mg/ml. Before use in microinjection experiments and nuclear assembly assays, samples were centrifuged at 20,000 g for 10 min at room temperature to remove insoluble material.

**Microinjection of Mammalian Cells and Analysis by Immunofluorescence**

BHK-21 cells were cultured as described elsewhere (18). Single cells were injected with ΔNLA at a concentration of 1 mg/ml in PB (18). Controls consisted of the injection of cells with PB. Cells were fixed in methanol 2 h after microinjection and processed for immunofluorescence as described previously (18). A rat polyclonal antibody directed against human lamins A and C (39) or a rabbit polyclonal antibody directed against human lamin B

**Figure 1.** (a) SDS-PAGE of purified ΔNLA and (b) corresponding Western blot using a polyclonal lamin A/C antibody (39). The major band runs at 69 kD, the predicted molecular mass of ΔNLA. There are also minor bands, representing the proteolytic fragments seen in nuclear lamin preparations expressed in E. coli (38). Numbers on the left represent molecular mass size markers in kD.
 Xenopus Intercalase Extracts and In Vitro Nuclear Assembly Reactions

Xenopus laevis egg interphase extracts were prepared as described in (43). The extract was frozen in liquid nitrogen in 70-μl aliquots. Demembranated chromatin from Xenopus sperm was prepared as described in (42).

For all nuclear assembly reactions, aliquots of interphase extract were thawed rapidly and brought to 15 mM Hepes, pH 7.4. For ATP generation, the extract was made 1 mM ATP (Sigma Chemical Co., St. Louis, MO), 10 mM phosphocreatine (Sigma Chemical Co.), and 50 μg/ml creatine phosphokinase (Sigma Chemical Co.) (see 43). Stock solutions of bacterially expressed human LA or NLA in PB were added to the assembly reaction to a final concentration of 200 μg/ml. Controls consisted of the addition of an equal volume of PB to parallel assembly reactions. The volume of protein solution or PB alone added to an assembly reaction was maintained at <10% of the final volume of the reaction mixture. 15 min after the addition of protein or PB to an assembly reaction, demembranated sperm chromatin was added to initiate nuclear assembly. Sufficient sperm chromatin was added to achieve a final concentration of 1,000 nuclei per μl. Unless otherwise specified, nuclei were either fixed for immunofluorescence or prepared for electrophoretic analysis at 90–120 min after initiating the assembly reaction as described below.

In some experiments, assembled nuclei were removed from one assembly reaction and transferred to another. To accomplish this, the assembly reaction mixture was diluted 50-fold with NWB (200 mM sucrose, 15 mM Hepes, pH 7.4, 50 mM NaCl, 2.5 mM MgCl₂). The nuclei were recovered by centrifugation for 5 min at 1,600 g (5). The resulting pellets were then resuspended in fresh nuclear assembly reaction mixture.

The effects of ΔNLA on assembled nuclei were studied by adding ΔNLA (200 μg/ml final concentration) at a time interval of 90 min after the addition of sperm chromatin to a nuclear assembly reaction. After an additional 45 min, nuclei were fixed and processed for immunofluorescence as described below.

In Vitro DNA Replication Assays

DNA replication in the in vitro assembled nuclei was assayed with fluoroscence microscopy by adding 1 mM bio-11-dUTP (Enzo Diagnostics, Farmingdale, NY) to the nuclear assembly reaction to achieve a final concentration of 10 μM (6). The incorporated nucleotides were detected with fluochrome-tagged streptavidin as described below.

Preparation of Nuclear Matrices

Nuclear matrices were prepared from the nuclei assembled in vitro as described in (11). Nuclei were assembled in a 100-μl nuclear assembly reaction, and the reaction mixture was diluted with 650 μl of NWB containing 0.5% Triton X-100. DNase I (DPRF; Worthington Biochemical Corp., Freehold, NJ) was added to a final concentration of 8.3 μg/ml, and, after a 10-min incubation at 20°C, an additional 750 μl of 4 M NaCl, 20 mM Hepes, pH 7.4, 20 mM EDTA, and 1 mM DTT was added. After 10 min at 20°C, the matrices were fixed at the same temperature by adding 160 μl of 10 mM ethylene glycol bis [succinimidyl succinate] (Pierce Chemical Co., Rockford, IL). The fixation was stopped after 7 min by the addition of 42 μl of 1 M Tris HCl, pH 7.4.

Microscopic Analyses of Xenopus Nuclei Assembled In Vitro

Nuclei assembled in vitro were fixed for 10 min at 20°C by diluting the assembly reaction mixture 10-fold in NWB and adding 0.1 vol of 100 mM ethylene glycol bis [succinimidyl succinate] in DMSO (35). Fixation was stopped by adding 1 M Tris HCl, pH 7.4, to achieve a final concentration of 25 mM. Alternatively, nuclei were fixed with 4% paraformaldehyde in NWB for staining with XMCM3 (33). Subsequently, nuclei were pelleted onto coverslips as described elsewhere (37). After fixation, coverslips were placed on 0.1% NP-40 or 0.1% Triton X-100 in PBS for 2 min, and then rinsed twice for 2 min in PBS at room temperature. 30-μl aliquots of primary antibodies, diluted 1:200 in PBS, were overlaid on the coverslips. After a 30-min incubation at 37°C, coverslips were washed four times with PBS and incubated for an additional 30 min at 37°C with a 1:50 dilution of the appropriate fluorochrome-labeled secondary antibody. The coverslips were then washed five times in PBS and mounted in 50 mM Tris Base, pH 9.0, 50% glycerol, and 2 mg/ml of p-phenylenediamine (Sigma Chemical Co.).

The rabbit polyclonal sera used for these studies were directed against human lamins A and C (39), XORC2 (9; a gift from William Dunphy, California Institute of Technology, Pasadena), and XMCM3 (33; a gift from Ronald Laskey, Cambridge University, UK). Monoclonal asces and supernatants used included L6-5D5 directed against Xenopus lamin B3 (52), CRL 1640 directed against DNA polymerase α (57; American Type Culture Collection, Rockville, MD), a RFC 11 directed against the large subunit of replication factor C (8; a gift from Bruce Stillman, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY), and PC10 directed against PCNA (Boehringer Mannheim Biochemicals, Indianapolis, IN). The myc 9E10 epitope antibody was also used (13; American Type Culture Collection). The antibodies directed against PCNA, DNA polymerase α, XMCM3, and XORC2 have all been shown to react only with their targeted antigen in Xenopus extracts (9, 26, 27, 33). The antibody directed against human lamin A does not cross-react with Xenopus lamin B3 in immunofluorescence assays (data not shown). The secondary antibodies used were FITC-labeled donkey anti-mouse IgG, tetramethyl rhodamine isothiocyanate-labeled donkey anti-rabbit IgG, and lissamine rhodamine-labeled donkey anti-rabbit IgG (Jackson Immunoresearch Laboratories, Inc.). Detection of bio-11-dUTP incorporation involved the incubation of fixed nuclei for 30 min at 37°C in Texas red- or FITC-labeled streptavidin (Amersham Corp.) at a 1:50 dilution in the presence of 12.5 μg/ml of RNase A (4; Sigma Chemical Co.). Nuclear membranes were stained with the lipophilic dye dihexyloxacarbocyanine (DIOC₆) (Molecular Probes, Eugene, OR) at 2.5 μg/ml during the secondary antibody incubation and at 0.25 μg/ml in the mounting medium (44). Nuclear pore proteins were stained by adding FITC-labeled WGA (Sigma Chemical Co.) at 50 μg/ml during secondary antibody incubation (14). DNA was visualized by adding Hoechst dye (Molecular Probes) at 1 μg/ml to the mounting medium.

Microscopic observations were carried out on an Axiohot (Carl Zeiss, Inc., Thornwood, NY) equipped with a 35-mm camera or an LSM 410 confocal microscope (Carl Zeiss, Inc.) equipped with an argon/krypton laser. Confocal micrographs were stored on optical disks, and micrographs were printed on an UP-D8800 video printer (Sony Corp., Park Ridge, NJ).

Results

ΔNLA Disrupts Nuclear Lamin Organization In Mammalian Nuclei

To determine the function of nuclear lamins, we sought a method to perturb lamin organization. In previous studies, we found that microinjection of bacterially expressed human LA into mammalian cells resulted in its incorporation into the endogenous nuclear lamin structures (18). This technique allowed us to follow the pathway of nuclear lamin assembly in situ (18). We used this same technique to introduce a mutant lamin protein into BHK-21 cells. The mutant, ΔNLA, lacks the NH₂-terminal nonhelical domain (33 amino acids) of LA, but it contains the entire α-helical rod and COOH terminus. Bacterially expressed ΔNLA cannot form typical LA paracrystals in vitro, showing that the NH₂ terminus is required for higher order lamin assemblies (21, 38). To determine the effects of ΔNLA in vivo, a solution of this mutant protein (1 mg/ml; see Materials and Methods) was injected into the cyto-
plasm of BHK cells. Injected cells were fixed after 2 h and then processed for immunofluorescence. As can be seen in Fig. 2, there is a dramatic alteration in the distribution and organization of the nuclear lamins. Instead of producing the typical rim pattern associated with the nuclear envelope as well as nucleoplasmic foci (Fig. 2, a and b), lamins A/C and B colocalize almost exclusively in large nucleoplasmic aggregates in injected cells (Fig. 2, c and d). These results show that ∆NLA acts as a dominant negative mutant that can induce the rapid disruption of the endogenous lamin structures that are composed of both A- and B-type lamins.

**Human Lamin A Is Incorporated into the Endogenous Lamin Structures of Xenopus Nuclei Assembled In Vitro**

While microinjection of ∆NLA can disrupt the lamina of individual BHK-21 cells, such studies are not amenable to detailed biochemical and structural analyses, as only a limited number of cells can be injected, and it is difficult to control the amount of protein injected (for a discussion of this latter point, see 19). Similarly, others have shown that transfection of lamin cDNA mutants into mammalian cells can also result in similar nucleoplasmic structures and apparent disruption of the endogenous lamin organization, but the amount of protein expressed per cell and the percentage of cells expressing the protein are quite variable (23). In light of these limitations, we have developed a hybrid system in which controlled concentrations of mutated and wild-type human lamins are added to nuclear assembly extracts prepared from *Xenopus laevis* eggs (4, 30, 43). One of the major advantages of this in vitro nuclear assembly system includes the ability to precisely control the amount of protein added to the extract. In addition, large numbers of nuclei can be prepared for morphological and biochemical studies. We have used the human lamins in this system because species-specific antibodies allow us to distinguish between the endogenous *Xenopus* lamin B3 and the mutant human lamin.

As a control for the use of the *Xenopus* system, wild-type LA was added to the nuclear assembly reactions. Nuclei assembled under these conditions contain LA, which colocalizes with the endogenous *Xenopus* lamin B3 (LB3) (Fig. 3, a and b). Staining is present at the nuclear periphery and elsewhere through the nucleus when viewed by conventional immunofluorescence. The LB3 staining pattern is similar to that seen in nuclei assembled without LA (compare Fig. 3 b with Fig. 6 a). Furthermore, nuclei assembled in the presence of LA contain a normal DNA staining pattern as indicated with Hoechst dye (Fig. 3 f). They also possess a normal distribution of nuclear membrane, as shown by staining with the membrane intercalatiny dye DIOC4 (data not shown), and a nuclear envelope with nuclear pore complexes, as suggested by the presence of a typical WGA staining pattern (Fig. 4, c and d). This organization of human LA in a heterologous system is consistent with results obtained in other laboratories. For example, when human lamin RNA is expressed in *Xenopus* oocytes, the expressed protein is localized to the lamina of the germinal vesicle (28). Similarly, *Xenopus* lamins expressed in mammalian cells integrate normally into the lamina (15).

To determine if LA is stably incorporated into the endogenous nuclear lamina, nuclear matrices were prepared from nuclei assembled in the presence of LA. Nuclei were first treated with DNase I, and subsequently extracted with 2 M NaCl and 0.1% Triton X-100 (see Materials and Methods). The nuclear lamina is resistant to this digestion and extraction procedure that has been shown to remove ~90% of the nuclear proteins and DNA (11). Indirect immunofluorescence shows that the LA within the lamina is resistant to extraction and is therefore incorporated into the lamina of these nuclei (Fig. 3 e).

*Xenopus* nuclei assembled in vitro replicate their DNA once per cell cycle in a semiconservative manner (4, 30, 43). To determine whether the incorporation of LA affects DNA replication, biotinylated dUTP was added to nuclear assembly reactions (see Materials and Methods) containing LA. After fixation and staining with Texas red–streptavidin, fluorescence microscopic observations show that this nucleotide is incorporated in a fashion indistinguishable from control nuclei (compare Figs. 3, d and e, and 6, a and b).

**∆NLA Disrupts Nuclear Lamin Organization in Nuclei Assembled In Vitro and Inhibits DNA Replication**

When the mutant human lamin ∆NLA is added to the nuclear assembly reaction at the same concentration as that used for the wild-type LA (200 μg/ml final concentration; see Materials and Methods), normal lamin assembly is al-

---

**Figure 2.** Double label immunofluorescence showing nuclear lamin patterns in a normal interphase BHK cell and in a cell injected with ∆NLA. Nuclei were stained for lamins A/C (LA) (a and c) or lamin B (LB) (b and d). In the un.injected cell, there is a distinctive lamin rim as well as less intense nucleoplasmic foci (a and b), as previously described (39). In cells fixed 2 h after injection, the lamin rim is no longer obvious, and the lamin staining for both lamins A/C and B appears mainly in the same foci (c and d). Confocal optics showing sections through the mid-region of the nucleus. Bar, 5 μm.
tered in >90% of the nuclei. Instead of producing a typical nuclear lamin staining pattern, antibodies directed against Xenopus LB3 and LA stain large nucleoplasmic spheroidal bodies (Fig. 5, a and b). Confocal microscopic analysis of nuclei assembled with ΔNLA shows that both Xenopus LB3 and ΔNLA colocalize within these nucleoplasmic aggregates and that there is no obvious staining of the nuclear periphery (Fig. 5, d and e). However, the DNA of these lamin-disrupted nuclei appears to be distributed normally as indicated by Hoechst dye (Fig. 5 c). Furthermore, nuclear pore complexes, as indicated by WGA staining (Fig. 4, e and f), and the nuclear membranes, as indicated by DIOC₆ staining, (Fig. 4, g and h) appear to be normal.

We also determined if nuclear protein import could take place in disrupted nuclei. This involved the use of nuclei that were assembled in the presence of ΔNLA for 90 min (see Materials and Methods). Under these conditions, 100% (n = 84), of the in vitro assembled nuclei were disrupted. Subsequently myc-tagged wild-type LA was added to the same extract at a concentration equivalent to 25% of the mutant protein. 30 min later, the reactions were stopped and immunofluorescence assays demonstrated that the wild-type lamin A was imported into 100% (n = 80) of the disrupted nuclei (Fig. 4, i and j). These observations demonstrate that the disrupted nuclei are able to import nuclear proteins. Furthermore, under the experimental conditions used, the nuclei maintained their disrupted phenotype throughout the transport process. In contrast with the control nuclei, the myc-tagged lamin A colocalized with ΔNLA in the nucleoplasmic aggregates in disrupted nuclei (Fig. 4, i and j).

Interestingly, the disrupted nuclei are smaller than those formed in control assembly reactions or in reactions containing LA (e.g., compare Fig. 6 a and 6 b with 6 c and 6 d). Nuclei assembled in the presence of ΔNLA also seem more fragile than nuclei formed in either control or LA-containing assembly reactions, as indicated by their increased tendency to break open when centrifuged at low forces (see Materials and Methods; data not shown).

In contrast with the results obtained with the buffer control (Fig. 6, a and b) or with LA (see Fig. 3 e), the addition of ΔNLA to nuclear assembly reactions greatly inhibits DNA replication. When examined with conventional fluorescence optics, the disrupted nuclei show greatly reduced or no detectable incorporation of biotinylated dUTP after 90- and 180-min incubations (Fig. 6 d). However, confocal microscopy demonstrates that the majority of these nuclei do exhibit a faint punctate nucleoplasmic pattern of biotinylated dUTP incorporation (Fig. 6 f). To obtain a more quantitative measure of the inhibition of DNA synthesis, 32P-labeled dCTP was added to nuclear assembly reactions, in the presence or absence of ΔNLA. After a 90-min incubation, the DNA was isolated and resolved by gel electrophoresis (see Materials and Methods). Autoradiographic (Fig. 7 A) and phosphoimage analysis (Fig. 7 B) demonstrates that ΔNLA reduces the level of 32P-incorporation by 94% in disrupted nuclei.

**Disruption of Lamin Organization Alters the Distribution of Factors Required for the Elongation Phase of DNA Synthesis**

To examine the effects of altered lamin organization on the major steps of DNA replication, the distributions of
five proteins known to be involved in either the initiation or the elongation phases of DNA replication were examined. The organization of factors involved in the initiation phase of DNA replication was studied with antibodies directed against DNA polymerase α, XORC2, and XMCM3. DNA polymerase α is believed to catalyze the formation of primers at origins of replication (55). XORC2 has been shown to be essential for DNA replication in the Xenopus nuclear assembly extracts, and, as a result of its sequence homology to the yeast ORC2, it is most likely involved in the initiation of DNA replication (9). XMCM3 has been characterized as a component of the licensing factor for DNA replication in Xenopus nuclear assembly extracts, and it is thought to be required for the initiation of DNA replication (10, 29, 33).

The DNA polymerase α and XORC2 staining patterns were unaffected in ΔNLA-disrupted nuclei (Fig. 8, a–d), compared with nuclei formed in the presence of buffer (data not shown). In both disrupted and control nuclei, staining with these antibodies indicated that they colocalized with chromatin (not shown). This is in agreement with previous studies (9, 26). These results suggest that the disruption of the lamin network does not affect the early stages of replication. The XMCM3 staining pattern of ΔNLA-disrupted nuclei was also coincident with chromatin (Fig. 8, e and f). This was true of 45-, 90-, and 180-min nuclear assembly reactions. In addition, we noticed that, in nuclei assembled in the presence of buffer, the XMCM3 staining colocalizes with chromatin at 45 min, but the fluorescence intensity decreases over time so that at 90 min it is not detectable (data not shown). This loss of XMCM3 signal is identical to the results obtained by other groups and is believed to represent a displacement of the protein from chromatin as replication proceeds (10, 29, 33). The disruption of lamin organization apparently prevents this XMCM3 displacement.

In contrast, the staining patterns produced by the antibodies directed against the elongation factors, PCNA and the large subunit of RFC, were dramatically altered in nuclei assembled in the presence of ΔNLA. Both of these proteins are required cofactors for DNA polymerase δ, the polymerase known to be responsible for chain elongation during DNA synthesis (55, 59). As assayed by indirect immunofluorescence, these two factors are associated with chromatin in nuclei assembled in control reactions (Fig. 9, a and d) or in assembly reactions containing LA (data not shown). However, when nuclei are assembled in the presence of ΔNLA, PCNA and RFC colocalize with the lamin aggregates (Fig. 9, b, c, e, and f). These observations sug-
suggest that the inhibition of DNA replication resulting from the disruption of nuclear lamin organization may be caused by alterations in the localization and/or targeting of the components of the DNA replication machinery responsible for chain elongation.

**Disruption of Nuclear Lamin Assembly Is Reversible**

To determine if the ∆NLA-induced alterations of nuclear structure and function are reversible, nuclei were assembled in the presence of ∆NLA for 90 min. The disrupted nuclei were then removed from nuclear assembly reactions containing ∆NLA by centrifugation. These nuclei were resuspended in fresh extract to which no ∆NLA was added and were assayed for DNA replication. The nuclei were fixed 90 min later for immunofluorescence or processed for autoradiography (see Materials and Methods). As seen by both biotinylated dUTP incorporation (Fig. 10, a and b) and [32P]dCTP incorporation (data not shown), nuclear DNA synthesis was “rescued” when the disrupted nuclei were transferred to normal nuclear assembly reactions. Interestingly, although a few of the lamin aggregates remained in these nuclei, apparently normal nuclear lamina staining was reestablished (Fig. 10 a).

**Disruption of Nuclear Lamin Organization after Nuclear Assembly In Vitro**

The microinjection of ∆NLA into cultured mammalian cells

the lamin-disrupted nuclei do contain a faint punctate nucleoplasmic pattern of labeled nucleotide incorporation that is readily resolved by confocal microscopy. (a–d) Conventional optics (see f). (e and f) Confocal optics. Bar, 5 μm.
demonstrates the capacity of this truncated protein to disrupt endogenous lamin organization in fully formed interphase nuclei. At 90 min, nuclei formed in the *Xenopus* nuclear assembly reactions under normal conditions contain a normal lamin organization as indicated by lamin antibody staining (see, e.g., Fig. 4 a). DNA replication is complete or nearly complete, as indicated by the low level of incorporation of a 5-min pulse of biotinylated dUTP at this time interval (data not shown). To determine if ΔNLA can disrupt the lamin organization once it is established, ΔNLA was added to the nuclear assembly reaction 90 min after the initiation of nuclear assembly. The reaction was allowed to continue for an additional 45 min (see Materials and Methods). We found that the addition of ΔNLA under these conditions induced a dramatic disruption of the endogenous lamina in >90% of the nuclei observed. This resulted in the formation of nucleoplasmic aggregates containing LB3 (Fig. 10 c) and ΔNLA (data not shown). The addition of ΔNLA at both earlier and later time points (45, 120, and 150 min) after the initiation of nuclear assembly also resulted in the formation of nucleoplasmic aggregates indistinguishable from those seen in Fig. 10 c. In all cases, these aggregates appear to be structurally identical to those formed when nuclei are assembled in the presence of ΔNLA (see Figs. 3–6). However, it should be noted that nuclei disrupted 90 min after the initiation of assembly are larger than nuclei formed in the presence of ΔNLA.

**Discussion**

In this study we describe the use of a dominant negative mutant human lamin to disrupt the organization of the nuclear lamina assemblies both in vivo by microinjection into mammalian cells and in vitro assembled *Xenopus* nuclei. We further demonstrate that a normal distribution of nuclear lamins is required for DNA synthesis. Specifically, the addition of ΔNLA, a mutant human lamin, to the *Xenopus laevis* nuclear assembly system blocks the formation of a normal lamina at the nuclear periphery. Instead, the endogenous LB3 and ΔNLA are found as constituents of the same large aggregates dispersed throughout the nucleoplasm. Under these conditions, DNA synthesis is dramatically reduced to ~5% of its normal level. These results are consistent with those obtained from immunodepletion studies demonstrating that LB3 is required for DNA synthesis in in vitro assembled nuclei (17, 34, 45). The nuclei

Figure 7. The addition of ΔNLA to nuclear assembly reactions reduces [³²P]dCTP incorporation by ~95%. (A) Autoradiogram of an agarose gel showing the incorporation of [³²P]-labeled dCTP into the DNA of nuclei formed in the presence of buffer control or ΔNLA. After nuclear assembly, the samples were treated as described in Materials and Methods and resolved on an 0.8% agarose gel. The upper band is at the origin of the gel. (B) Quantitation of replication assays shown in A. The radioactive signal of the dried gel was quantitated with a FUJIX BAS 2000 phosphoimager. The sum of the signal intensity/area value for both bands in each lane was used to measure the total incorporation of radioactivity into DNA. The average value for four replicate assays was plotted in a bar graph, where the vertical axis represents the signal/area values (in thousands) determined by the imager. The average value for samples containing ΔNLA was 2,292, with values ranging from 2,035–2,513. The average value for the control samples was 39,030, with samples ranging from 34,514–47,443. The addition of ΔNLA to the nuclear assembly reaction reduced the incorporation of [³²P]-labeled dCTP to ~5% of that found in control reactions.

Figure 8. Nuclei formed in the presence of ΔNLA, and then subsequently stained for ΔNLA or lamin B3 and one of several early DNA replication markers. (a and b) Nucleus stained for (a) ΔNLA and (b) DNA polymerase α. (c and d) Nucleus stained for (c) LB3 and (d) XORC2. (e and f) Nucleus stained for (e) LB3 and (f) XMCM3. The distribution of DNA polymerase α, XORC2, and XMCM3 is not altered by the disruption of the lamin structure. Confocal microscopic images showing sections through the middle of the nuclei. Bar, 5 μm.
formed in LB3-immunodepleted extracts or ΔNLAs-containing extracts described in this study have other features in common, including the fact that nuclear membranes and pores appear to assemble in a relatively normal fashion (34, 45). However, the dominant negative approach introduced in this study avoids one of the major problems inherent in the immunodepletion experiments: that the block in DNA replication could be caused by the removal of other proteins associated with lamin B3 in the assembly extract. In the experiments presented in this study, no components are removed from the nuclear assembly system.

The disruption of lamin organization alters the distribution of both PCNA and the large subunit of RFC, two essential cofactors for DNA polymerase δ during the elongation phase of replication (55). Normally both the large subunit of RFC (see Fig. 9) and PCNA (26) are distributed along chromatin. In ΔNLAs-disrupted nuclei, these cofactors are reorganized, along with the nuclear lamins, to form nucleoplasmic aggregates. These aggregates are not obviously associated with chromatin, and this in turn may have an inhibitory effect(s) on the assembly and function of the elongation machinery. Alternatively, disruption of lamin organization may also alter an aspect of the initiation process itself that we have been unable to detect.

At the present time, we believe that our results support an effect on the elongation phase of DNA replication for several reasons. Evidence suggesting that the initiation of DNA synthesis does not rely on an intact lamin organization comes from our immunofluorescence studies of XORC2, XMCM3, and DNA polymerase α. These three proteins are thought to be involved in the initiation and primer formation steps of DNA replication (9, 10, 29, 33, 55). The disruption of the nuclear lamin structure does not detectably alter the distribution of DNA polymerase α and XORC2 or the initial distribution of XMCM3, and all three of these factors remain associated with chromatin. Interestingly, during nuclear assembly, it appears that XORC2 and XMCM3 along with two other initiation factors, RFA and FFA, bind to chromatin before the assembly of either higher order lamin structures or the nuclear membrane (1, 9, 10, 29, 33, 55).

Figure 9. Staining patterns of lamin and DNA replication factors involved in elongation in nuclei assembled in the presence of (a and d) buffer or (b, c, e, and f) ΔNLAs: (a) Control nucleus stained for PCNA. (b and c) Nucleus assembled in the presence of ΔNLAs stained for PCNA. (d) Control nucleus stained for RFC. (e and f) Nucleus assembled in the presence of ΔNLAs stained for RFC. (e and f) ΔNLAs. PCNA and RFC distributions are altered from the control as a consequence of lamin disruption such that PCNA and RFC colocalize with lamin aggregates in these nuclei. Confocal microscope showing sections through the middle of the nuclei. Bar, 5 μm.

Figure 10. (a and b) Lamin and biotinylated dUTP incorporation in a nucleus formed in the presence of ΔNLAs, and subsequently transferred to an interphase extract containing biotinylated dUTP but lacking ΔNLAs (see text). Confocal micrographs showing sections through the middle of the nucleus. (a) Nucleus stained for Xenopus LB3. (b) Pattern of biotinylated dUTP incorporation as shown by binding of Texas red–conjugated streptavidin. The disrupted nuclei were transferred to a nuclear assembly reaction lacking ΔNLAs, where they form a lamin rim and replicate DNA. However, some lamin foci remain. (c) Postassembly disruption of the lamin structure of an in vitro assembled nucleus. ΔNLAs was added 90 min after the onset of nuclear formation, a point at which the nuclei have normal lamin organization and have largely completed DNA replication. The addition of ΔNLAs disrupts the assembled LB3 staining pattern. Bar, 5 μm.

Spann et al. Disruption of Nuclear Lamins and DNA Replication
These findings suggest that both the sites of initiation of DNA replication, as well as the organization of initiation cofactors at these sites, take place early in the process of nuclear assembly. In addition, the results reported here indicate that the location, organization, and function of these origins of replication may be independent of the presence of normal lamin organization.

Evidence indicating that the initiation of DNA synthesis occurs in disrupted nuclei comes from the observations that ∆NLA dramatically reduces but does not eliminate incorporation of biotinylated dUTP or [32P]dCTP. The very faint punctate pattern of biotinylated nucleotide incorporation in disrupted nuclei (Fig. 6 f) is reminiscent of the centers of DNA synthesis observed in normal nuclei at the onset of replication (24, 25, 37). Therefore, it is possible that the pattern and low levels of nucleotide incorporation seen in the disrupted nuclei described in this study may result from sites of primer and initial strand synthesis. This is also consistent with the unaltered distribution of DNA polymerase α in the lamin-disrupted nuclei. However, at this point, we cannot eliminate the possibility that the low level of incorporation of biotinylated nucleotide detected in disrupted nuclei may be due to other processes such as DNA repair.

Further clues for the role of the nuclear lamins are derived from the immunofluorescence pattern of XMCM3 in ∆NLA-disrupted nuclei. Normally, in control nuclei, XMCM3 is displaced from chromatin as replication progresses and a dramatic decrease in fluorescence intensity is observed (10, 29, 33). However, in the ∆NLA-disrupted nuclei reported in this study, the fluorescence intensity of XMCM3 staining appeared unchanged throughout the entire assembly reaction. This suggests that the disruption of the lamin structure prevents the dissociation of XMCM3 from chromatin, presumably by arresting replication before the normal displacement of this factor. Interestingly, it has been proposed that XMCM3 may be involved in the switch between the initiation and elongation phases of replication (60). Taken together, these results suggest that disruption of lamin organization blocks replication after the initiation of DNA synthesis and prevents the switch from the initiation to the elongation phase of DNA replication. However, the precise elucidation of the point at which lamin disruption blocks DNA synthesis requires a more complete understanding of the steps involved in DNA replication.

Most of our understanding of the biochemistry of DNA replication has come from the coupling of genetic studies in yeast with in vitro studies of SV-40 replication (55). The mechanisms involved in the regulation of DNA replication in more complex genomes remain largely unknown. However, it appears from the data presented in this study and elsewhere (17, 34, 44, 55) that, in higher eukaryotic cells, the nuclear lamins play a vital role in this process. We have previously reported that lamin B colocalizes with PCNA in mammalian cells during S phase (39). In this report we find that the disruption of nuclear lamin organization also alters the normal organization of RFC and PCNA, such that all three proteins are found in the same aggregates. These results suggest that lamins interact with the components of the strand elongation complexes. This interaction could be direct, with lamins binding to a component of the replication machinery, or indirect through unknown nuclear proteins or structural entities. This proposed interaction of PCNA with nuclear lamins is also supported by the finding that PCNA is readily extracted from nuclei formed in lamin B3-depleted assembly reactions, but is resistant to extraction in nuclei assembled in control reactions (27).

Traditionally, lamins have been thought to be located exclusively at the nuclear periphery, which makes it difficult to model lamin involvement in DNA synthesis, since much of the replication process takes place deep in the nucleoplasm. However, the presence of lamins in the nucleoplasm is supported by a number of studies (7, 18, 39), and we have found lamin B3 within the nucleoplasm of control Xenopus nuclei (Fig. 3 a). This lamin staining may be part of a dynamic nucleoplasmic lamin network. Such a network could form a scaffold (24, 25) upon which replication factors are assembled into functional units that facilitate the formation of active elongation complexes and/or stabilize such complexes once they are formed. Such an organization would explain why perturbations in nuclear lamin organization can block DNA replication and cause the abnormal distribution of RFC and PCNA.

The reduced size of nuclei assembled in the presence of ∆NLA provides evidence that the nuclear lamins are involved in the growth of the nucleus after its initial assembly. In addition, the increased fragility of the nuclei assembled in the presence of ∆NLA supports the idea that the nuclear lamin structure also provides a mechanical support system for the nucleus. These findings are consistent with previous reports of the small size and increased fragility of nuclei assembled in the absence of LB3 (34, 45).

The ∆NLA-induced disruption of lamin structure is most likely related to the dynamic characteristics of lamins in vivo. These properties of nuclear lamins are very similar to those found for other types of intermediate filament systems (36, 46, 50, 56, 58). Specifically, it has been shown that during interphase the nuclear lamins do not form a static polymer in vivo, but rather they are in a state of dynamic equilibrium between subunits and polymer. For example, microinjected nuclear lamins are rapidly incorporated into endogenous lamin polymers (18). Similarly, the results of fluorescence recovery after photobleaching experiments demonstrate that lamin assemblies undergo continuous subunit exchange in living cells (49).

Since it is known from in vitro studies that the central α-helical rod domain is required for normal lamin–lamin interactions, it is probable that ∆NLA and LB3 interact through their highly conserved rod domains to form heterostructures. However, the in vitro assembly of higher order lamin structures such as tetramers and larger oligomeric complexes requires the NH2-terminal domain that is missing in ∆NLA (20, 38). Therefore, the ∆NLA/LB3 heterostructures most likely cannot be incorporated into higher order lamin complexes. In the presence of excess ∆NLA, the normal process of subunit exchange could produce a large pool of heterostructures. In turn, this could drive the equilibrium in the direction of disassembly, ultimately resulting in the disruption of the endogenous lamin structure described in this study.

In summary, the addition of exogenous normal and mutated lamins to the Xenopus nuclear assembly system has provided evidence that a normal nuclear lamin organiza-

The Journal of Cell Biology, Volume 136, 1997 1210
tion is required to proceed from the initiation to the elongation phase of DNA replication. The assays used are relatively simple and should continue to provide further structural and biochemical information about the role of nuclear lamins in DNA replication. Furthermore, the availability of a soluble pool of nuclear components should allow us to fractionate the interphase extract and to determine whether lamin proteins are interacting directly with elongation factors at replication forks or indirectly through other unidentified nuclear components. This use of Xenopus extracts has already proven to be very important in identifying and characterizing the interactions of factors involved in the initiation of DNA replication within nuclei (1, 9, 10, 29, 33, 61). We believe such an approach will also help to elucidate the role of nuclear lamins in other processes such as postmitotic nuclear assembly, nuclear growth, and the maintenance of the overall shape and structural integrity of the nucleus.

We thank Ms. Satya Khuon for help in preparing some of the micrographs and Ms. Laura Davis for help in manuscript preparation.

This work is supported by grant CA31760 from the National Cancer Institute.

Received for publication 24 October 1996 and in revised form 29 January 1997.

References


