Decreased mechanotransduction prevents nuclear collapse in a Caenorhabditis elegans laminopathy

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The function of the nucleus depends on the integrity of the nuclear lamina, an intermediate filament network associated with the linker of nucleoskeleton and cytoskeleton (LINC) complex. The LINC complex spans the nuclear envelope and mediates nuclear mechanotransduction, the process by which mechanical signals and forces are transmitted across the nuclear envelope. In turn, the AAA+ ATPase torsinA is thought to regulate force transmission from the cytoskeleton to the nucleus. In humans, mutations affecting nuclear envelope-associated proteins cause laminopathies, including progeria, myopathy, and dystonia, though the extent to which endogenous mechanical stresses contribute to these pathologies is unclear. Here, we use the Caenorhabditis elegans germline as a model to investigate mechanisms that maintain nuclear integrity as germ cell nuclei progress through meiotic development and migrate for gametogenesis—processes that require LINC complex function. We report that decreasing the function of the C. elegans torsinA homolog, OOC-5, rescues the sterility and premature aging caused by a null mutation in the single worm lamin homolog. We show that decreasing OOC-5/torsinA activity prevents nuclear collapse in lamin mutants by disrupting the function of the LINC complex. At a mechanistic level, OOC-5/torsinA promotes the assembly or maintenance of the lamin-associated LINC complex and this activity is also important for interphase nuclear pore complex insertion into growing germline nuclei. These results demonstrate that LINC complex-transmitted forces damage nuclei with a compromised nuclear lamina. Thus, the torsinA–LINC complex nexus might comprise a therapeutic target for certain laminopathies by preventing damage from endogenous cellular forces.

nuclear lamina | torsinA | nuclear mechanotransduction | laminopathy | LINC complex

Nuclear mechanotransduction enables the nucleus to adapt its rigidity in response to applied forces and is necessary for transmitting mechanical signals across the nuclear envelope (NE) and for maintaining the integrity of the genome (1, 2). Nuclear mechanotransduction requires an intact nuclear lamina, which comprises an intermediate filament network of lamins and associated proteins that line the nuclear-facing surface of the NE. The LINC (linker of nucleoskeleton and cytoskeleton) complex is a core NE component required for nuclear mechanotransduction, as it acts as a molecular bridge coupling the inside of the nucleus with the cytoplasm. The LINC complex consists of two interacting protein components: 1) SUN (Sad1 and UNC-84) domain proteins in the inner nuclear membrane; and 2) KASH (Klarsicht, ANC-1, Syne homology) domain proteins in the outer nuclear membrane (2). The LINC complex is tethered to the cytoplasmic cytoskeleton and nuclear lamins and transduces forces across the NE.

Mutations in many NE components are associated with a range of seemingly unrelated tissue-specific pathologies, collectively known as nuclear envelopathies (3). The apparent tissue specificity of these diseases is somewhat surprising because most of these NE components are widely expressed proteins. For example, mutations in genes encoding nuclear lamins cause diseases, including Hutchinson-Gilford progeria, dyslipidemias, and myopathies (3). By contrast, mutations affecting the NE-associated AAA+ ATPase (ATPases associated with diverse cellular activities) torsinA cause early-onset DYT1 dystonia and congenital joint contractures (4–6). An attractive hypothesis is that endogenous mechanical forces contribute to the pathology of nuclear envelopathies by causing damage to cells with defective NE components, yet few studies have directly tested this possibility. In many cases, experimental investigations are hampered by the fact that the mechanistic functions of many key NE components remain incompletely understood—such is the case of torsinA.

In this study, we set out to elucidate molecular interactions and mechanisms by which torsinA functions in the NE. Although the cellular functions of torsinA have remained enigmatic, they appear to be conserved across cell types and species because distantly related organisms lacking torsinA function display similar ultrastructural NE defects (7, 8). A plausible hypothesis is that torsinA plays a role in regulating nuclear mechanotransduction, as its loss disrupts cell migration, nuclear positioning, and polarization (8–10). Yet, torsinA has been implicated in other cellular processes, including endoplasmic reticulum (ER) stress (11–14), lipid metabolism (15), and membrane remodeling (16). In this study, we took advantage of the fact that mutations in the Caenorhabditis elegans torsinA ortholog, OOC-5 (abnormal oocyte formation, 5), cause oocyte growth and embryonic polarity defects.
(17, 18). Further, we used the *C. elegans* germline as a model system to investigate how cytoskeletal forces are transduced as germ cell nuclei progress through meiosis and migrate for gametogenesis. Our results reveal that OOC-5/torsinA promotes the activity of the LINC complex in transducing forces to the nuclear lamina. Remarkably, decreasing the function of OOC-5/torsinA or the LINC complex prevents nuclear collapse in the absence of a functional nuclear lamina. These results highlight a critical role for torsinA in nuclear mechanotransduction and suggest that an inability to cope with endogenous and exogenous forces may underlie the etiology of nuclear envelopathies.

**Results**

**OOC-5/TorsinA Associates with the LINC Complex Protein SUN-1 and Lamin.** To address the mechanism of OOC-5 function, we purified OOC-5-associated proteins from soluble and membrane protein fractions using tandem affinity purification (SI Appendix, Fig. S1). Both purifications identified highly conserved regulators of ER and NE homeostasis (SI Appendix, Table S1). The OOC-5-associated proteins include OOC-3, a protein required for OOC-5 localization and function (17, 18), ER chaperone CNX-1/calnexin, which associates with torsinA in mammalian cells (19), and OEE-15 and T09B4.2, the divergent *C. elegans* homologs of the torsinA-interacting proteins LAP1 and LUL1 (20).

Also, among the OOC-5-associated proteins was SUN-1, the SUN domain inner nuclear membrane protein component of LINC complexes in the *C. elegans* germline (21). SUN domain proteins interact with nuclear lamins (22, 23), and the sole *C. elegans* lamin, LMN-1 (24), was well represented among OOC-5-associated proteins (42 to 43% protein coverage; SI Appendix, Table S1). The genetic interaction studies described below suggest that the association between OOC-5, which resides in the lumen of the NE and ER, and nucleoplasmic LMN-1 is likely mediated through SUN-1.

**Decreasing OOC-5/TorsinA Function Prevents Nuclear Collapse and Extends Lifespan in Lamin Mutants.** To address the significance of the protein associations with OOC-5, we conducted genetic analyses. The most striking result was an interaction between ooc-5 and lmn-1. Most *lmn-1* null mutants are agametic, sterile, and have small gonads (Figs. 1A and 2A). However, 12.3 ± 7.8% of animals (“escapers”) develop larger gonads and produce dead embryos (Figs. 1A and 2A). Surprisingly, compromising ooc-5 function rescued the sterility of *lmn-1* mutants such that all animals produced gametes and dead embryos (Figs. 1A and 2A and SI Appendix, Fig. S2A). Importantly, reducing the dosage of ooc-5 was sufficient to suppress *lmn-1* sterility (Figs. 1A and 2A). These and the following genetic analyses were performed using a deletion of the *ooc-5* locus that we generated by CRISPR-Cas9 genome editing (SI Appendix, Fig. S3A). Yet, similar results were observed for other ooc-5 alleles (n = 69 for the *tm1709* deletion allele, and n = 57 for the *it145* R9stop nonsense allele). Because prior work established that *lmn-1* mutants exhibit a reduced lifespan (25), we tested whether decreasing ooc-5 function could suppress the premature aging observed in *lmn-1* mutants. Indeed, reducing or eliminating ooc-5 function suppressed the decreased lifespan of *lmn-1* mutants (Fig. 1B and SI Appendix, Table S2). Importantly, reducing or eliminating ooc-5 function did not increase lifespan in an otherwise wild-type genetic background (Fig. 1B and SI Appendix, Table S2).

The germ cells of the *lmn-1* escapers appeared to deteriorate as they progressed through meiotic prophase; they assembled a synaptosomal complex (SC) but their nuclei appeared to collapse soon after the pachytene stage (SI Appendix, Figs. S3D and S4). We assessed the dynamic nature of this apparent nuclear collapse using time-lapse imaging of *lmn-1* mutant gonads (Fig. 1C and Movies S1–S3). Proximal germ cell nuclei that appeared normal in size and shape at the beginning of the recording were observed to rapidly condense and even fragment. We observed nuclear collapse in *lmn-1* mutants in at least three situations: 1) simultaneously with the ovulation of the most proximal oocyte (Fig. 1C, Top and Movie S1); 2) spontaneously in a cellularized oocyte (Fig. 1C, Middle and Movie S2); and 3) while migrating in the proximal direction with the cytoplasmic flows, which also resulted in nuclear fragmentation (Fig. 1C, Bottom and Movie S3). In all cases, nuclear collapse occurred within the span of a few minutes. We also observed that SUN-1 lost its normal NE localization and appeared diffuse in most collapsing nuclei (SI Appendix, Fig. S5 C and D). Remarkably, reducing the dosage of ooc-5 prevented the nuclear collapse in *lmn-1* mutants (Fig. 1A and SI Appendix, Fig. S3C). Whereas diakinesis-stage chromosomes were rare in *lmn-1* mutant oocytes, they were observed in most or all *lmn-1* mutants after reducing ooc-5 function (Fig. 1A and SI Appendix, Fig. S3C). Further, reducing or eliminating ooc-5 function enabled SUN-1 to retain its NE association in *lmn-1* mutants (SI Appendix, Fig. S5 E and F). Additionally, we observed increased levels of DNA damage in late-stage pachytene female germ cells of *lmn-1* mutants (SI Appendix, Fig. S6). Nevertheless, decreasing ooc-5 function did not appear to completely rescue this defect (SI Appendix, Fig. S6 A and B).

**OOC-5/TorsinA Acts through the LINC Complex.** Studies in mammalian cells suggest that torsinA regulates force transmission from the cytoplasm to the nucleus (9, 10). Thus, we examined a model in which OOC-5 promotes LINC complex function, which damages nuclei with a compromised nuclear lamina. We tested for genetic interactions between *lmn-1* mutants and genes encoding components of the LINC complex, utilizing mutations in *sun-1* and zyg-12, which encode the KASH domain outer nuclear membrane LINC complex component in the germline (21). As observed for ooc-5, the proportion of fertile *lmn-1* mutant animals increased when LINC complex function was compromised, and nuclear collapse was substantially ameliorated (Fig. 2A and SI Appendix, Fig. S2B). These results suggest that OOC-5 and LINC complex-transmitted forces contribute to nuclear damage in *lmn-1* mutants.

**OOC-5/TorsinA Promotes LINC Assembly or Maintenance in Growing Late Prophase Meiotic Nuclei.** TorsinA function is required for the proper localization of KASH domain proteins ZYG-12 and nesprin3 in *C. elegans* germ cells and mammalian fibroblasts, respectively (8, 9). To assess whether OOC-5 promotes LINC complex integrity, we used superresolution microscopy to examine the colocalization of SUN-1 and ZYG-12 in the wild type and ooc-5 mutants. Prior work showed discontinuities in ZYG-12 localization in ooc-5 mutants commencing in pachytene germ cells (8). In addition to these discontinuities, we observed a reduction of ZYG-12 NE localization as germ cells progressed from the pachytene stage to the diakinesis stage of meiotic prophase (Fig. 2B). By contrast, SUN-1 remained in the NE at high levels in ooc-5 mutants (Fig. 2B). We quantified SUN-1 and ZYG-12 colocalization as a function of nuclear diameter by measuring Pearson’s correlation and Mander’s overlap as a function of nuclear diameter. We observed that as the ooc-5(-) germ cells grew, colocalization declined (Fig. 2C). These results suggest that OOC-5 promotes the assembly or maintenance of the LINC complex, which appears particularly important for growing meiotic prophase nuclei.

**OOC-5/TorsinA Promotes the Activity of the LINC Complex in Early Meiotic Prophase Nuclei.** The LINC complex is essential for the pairing of homologous chromosomes in *C. elegans* meiosis (26–28). OOC-5 is not required for SC formation (SI Appendix, Figs. S3B and S4) and appears dispensable for meiotic chromosome segregation (17, 18). To examine whether OOC-5 promotes...
the efficiency of meiotic pairing, we examined the localization of ZIM-3 and HIM-8, which bind the pairing center regions on chromosomes I and IV and the X chromosome, respectively (29). The spatial distribution of unpaired meiotic chromosomes across the distal gonad suggests that pairing of chromosomes I and IV is delayed in ooc-5 mutants (Fig. 3 A and B, Upper). By contrast, ooc-5 appears dispensable for the timely pairing of the X chromosome (Fig. 3 A and B, Lower). Persistent SUN-1 phosphorylation correlates with synaptic or recombination defects (30). We observed that phosphorylated SUN-1 (S8Pi) persisted in an extended zone in ooc-5 mutants (Fig. 3 C and D), and that the number of SUN-1 foci and their speed of motion increased (Fig. 3 E and F and Movies S4–S6). The increased movement of SUN-1 foci in ooc-5 mutants may reflect a partial release from cytoskeletal tension. By contrast, we observed lower numbers and slower movement of SUN-1 foci in lmn-1 mutants (Fig. 3 E and F). We hypothesize that this observation reflects a lower level of force resistance exerted by the weakened nuclear lamina in the absence of LMN-1 such that the SUN-1 foci are subjected to enhanced cytoskeletal tension and thus move more slowly and coalesce into fewer foci. Taken together, these results suggest that the function of the LINC complex is perturbed in ooc-5 mutants even in early prophase germ cells, which exhibit normal SUN-1 and ZYG-12 colocalization.

OOC-5/TorsinA and the LINC Complex Promote Interphase Nuclear Pore Insertion. C. elegans OOC-5 and mammalian torsinA are required for the proper localization of nuclear pore complex (NPC) proteins (8, 31). Mislocalization of nucleoporins (Nups) may be sufficient to explain the embryonic polarity defects in ooc-5 mutants (32). The two transmembrane Nups, NPP-12 and NPP-22, were identified among OOC-5-associated proteins (SI Appendix, Table S1). A prior study concluded that an NPP-22::GFP-expressing transgene localized more normally in ooc-5 mutants than other Nups, but the localization of NPP-12 was not examined (8). Thus, we used genome editing to generate GFP fusions to the endogenous transmembrane Nups and analyzed their localization. By examining the localization of these two fusion proteins we confirmed that they localize to patches within the NE of ooc-5 mutant germ cells, consistent with the previous report (8). However, we observed further localization changes in that both NPP-12::GFP and GFP::NPP-22 exhibited substantial mislocalization in ooc-5 mutants, accumulating largely outside the NE and in the cytoplasm of proximal oocytes (Fig. 4 A and B). We tested the possibility that this mislocalization of transmembrane Nups in ooc-5 mutants might be a consequence of the impaired function of the LINC complex. Thus, we used a temperature-sensitive zyg-12 mutant allele to examine the localization of...
We observed that both NPP-12::GFP and GFP::NPP-22 exhibited clear mislocalization in the germlines zyg-12 mutants (Fig. 4A and B). We also examined the effect of a sun-1 reduction-of-function (rf) allele on the localization of these transmembrane Nups. We found that NPP-12::GFP and GFP::NPP-22 abnormally aggregate in a patchy pattern on the NE of sun-1(rf) nuclei, but they only subtly changed their subcellular localization (Fig. 4A and B). The subtle effect on the localization of Nups might be a reflection of the reduction-of-function nature of this sun-1 allele.

**Discussion**

Decreased function of torsinA, the LINC complex, lamins, and other NE-associated proteins, result in altered nuclear deformability (1, 2, 33). Prior studies suggest that torsinA might function as a regulator of the LINC complex (9, 10). Four of our findings provide direct support for this model: 1) OOC-5 associates with SUN-1, as detected using tandem affinity purification (SI Appendix, Table S1); 2) mutations in OOC-5 and LINC complex components can suppress nuclear collapse in lamin null mutants (Fig. 2A); 3) OOC-5 is required for LINC complex assembly or maintenance in growing oocyte nuclei (Fig. 2B and C); and 4) LINC complex-dependent activities, including the pairing of meiotic chromosomes and interphase nuclear pore insertion, are misregulated in the germline of ooc-5 mutants (Figs. 3 and 4 and Movies S4–S6). Consistent with these findings, observations in mouse embryos suggest that torsinA acts upstream of the LINC complex (34). However, decreasing the dosage of the LINC complex protein Sun2 prevented abnormal brain morphogenesis in torsinA-deficient mouse embryos (34). Taken together with our results, this latter finding suggests that torsinA might promote or inhibit LINC complex-dependent activities depending
on cellular context or the specific SUN-KASH pairs composing the particular LINC complexes. Further work will be needed to determine the precise molecular mechanisms by which torsinA regulates the activity of LINC complexes.

When torsinA is not functional, cells accumulate abnormal vesicles within the perinuclear space, which are often referred to as “blebs” (7, 8, 35, 36). One possibility is that the NE blebs observed in torsinA mutant cells might originate from defects in interphase NPC assembly or insertion. Consistent with this possibility, budding yeast mutants that disrupt the process of NPC insertion or assembly accumulate similar NE blebs (37, 38).

In multiple cell types lacking torsinA function, several Nups localize to abnormal perinuclear plaque-like structures (8, 31). In this study, we found that OOC-5/torsinA associates with the two C. elegans transmembrane Nups, NPP-12 and NPP-22, and that these proteins lose their strict NE localization in the growing nuclei of ooc-5 mutant oocytes (Fig. 4 A and B and SI Appendix, Table S1). The mislocalization of transmembrane Nups in ooc-5 mutants might be a consequence of the impaired function of the LINC complex because NPP-12::GFP and GFP::NPP-22 were mislocalized in sun-1 and zyg-12 mutants (Fig. 4 A and B). In support of this idea, prior work found that Sun1 disruption in mammalian cells leads to NPC mislocalization (39).

The physical association between the LINC complex and nuclear lamins is critical for nuclei to prevent deformation under mechanical stress (1, 2). Our findings suggest that in the absence of a functional nuclear lamina, LINC complex-dependent forces, promoted by OOC-5, result in nuclear damage (Fig. 4C). Using time-lapse microscopy, we were able to directly observe rapid nuclear collapse occurring in oocyte nuclei in the absence of LMN-1 (Movies S1–S3). Our work here capitalized on the fact that C. elegans has but a single nuclear lamin. Nonetheless, we speculate that rapid nuclear collapse might contribute to the pathology of laminopathies caused by milder perturbations to the nuclear lamina. Consistent with our model derived from studies of the C. elegans germline (Fig. 4C), disruptions of the LINC complex can ameliorate defects in cultured cells, mouse models of laminopathies, and fibroblasts derived from Hutchinson-Gilford progeria patients (40, 41). Age-related alterations in the ability to cope with mechanotransduction across the NE may contribute to progeroid syndromes and reduce normal healthspan.

Nuclear envelopopathies exhibit a broad range of symptoms and are characterized by a complex pathophysiology. Understanding the normal cellular functions and interactions of NE components and how their function is perturbed in nuclear envelopopathies is vital for advancing treatments. This study in the C. elegans germline system provides insights into the normal functions and interactions of torsinA with conserved NE components and elucidates how this normal function becomes deleterious when the structure of the nuclear lamina is compromised. Our findings indicate that OOC-5/torsinA promotes or modulates vital LINC complex functions needed for maintaining NE homeostasis. This activity of torsinA might be relevant to the pathophysiology of DYT1 dystonia because Sun1 deficiency leads to cerebellar ataxia in mice (42). Thus torsinA-dependent LINC complex activities might influence the pathophysiology of multiple nuclear envelopopathies.
Materials and Methods

Detailed materials and methods are available in SI Appendix, Materials and Methods.

Strains. C. elegans strains were maintained at 20 °C on Nematode Growth Medium (NGM) and fed with the Escherichia coli strain OP50-1, unless otherwise noted (for the genotypes of all strains used see SI Appendix, Table S3). Genes and mutations are described in WormBase (https://wormbase.org/). The following mutations were used: LGI—lmn-1(tm1502), npp-12(tn1780 [npp-12::gfp::tev::3x-flag]); LGII— ooc-5(tn1757), ooc-5(tn1758 [ooc-5::gfp::tev::3xflag]), zyg-12(ts), zyg-12(ts578), and LGV—sun-1(jf18), sun-1(jf199), npp-22(tm1794[tev::3xflag::npp-22]), and ooc-5(tn1709), ooc-5(tn1758 [ooc-5::gfp::tev::3xflag]), zyg-12(ts578), and LGV—sun-1(jf18), sun-1(jf199), npp-22(tm1794[tev::3xflag::npp-22]). The following rearrangements were used: tmC18[dpy-5(tmIs1236)], L mn1[dpy-10e128[mls14] II, and tmC12[egl-9[mls1194]] V. The following transgenes and transgene insertions were used: ieSi21[sun-1p-sun-1::mRuby::sun-13′ UTR + Cbr-unc-119] IV, ith37[pie-1p::mCherry::H2B::pie-1 3′ UTR + unc-119] IV, qj158[jzyg-12::gfp + unc-119(+) II, and ith38[pie-1p::GFP::PH(PLC1delta1) + unc-119(+) II]. Phenotypic analyses were performed on well fed animals to avoid the effects of starvation in germline development.

Genome Editing. CRISPR-Cas9 genome editing used standard methods (43–46) as described in detail in SI Appendix. All new alleles were verified by Sanger sequencing and outcrossed to wild-type males at least three times. The sequence of oligonucleotides used to construct single-guide RNAs, repair templates, and sequencing primers are listed in Dataset S1.

OOC-5 Immunopurifications. OOC-5::GFP::TEV::3xFLAG and interacting proteins were purified from soluble protein extracts and membrane protein fractions prepared from the DG4377 strain using modifications of methods we previously described (47–49) as detailed in SI Appendix. Purifications from the wild-type strain (N2) were run in parallel as a control. Small aliquots of protein fractions from every step of the purifications were analyzed by Western blotting (Fig. 1 and SI Appendix, Materials and Methods).

Mass Spectrometry. Immunopurified proteins were concentrated on a centrifugal evaporator (Jouan RC 10-10 Vacuum Concentrator) for 5 h, briefly separated on a 12% Bis-Tris NuPAGE gel, and stained using the Colloidal Blue Staining Kit (Invitrogen) (SI Appendix, Fig. S1G). Lanes were subdivided into six slices and mass spectrometry was performed at the Taplin Biological Mass Spectrometry Facility (Harvard Medical School) using an LTQ ion-trap mass spectrometer. In addition to control immunopurifications from the N2 strain, to account for abundant contaminant proteins that bound nonspecifically to the immunopurification matrix, we utilized data from multiple purifications of GFP fusion proteins, which we previously reported (47–49; see SI Appendix, Materials and Methods for the details of the filtering criteria used).

Fig. 4. OOC-5 and the LINC complex promote the localization of transmembrane Nups. (A) Confocal images of the expression of GFP::NPP-22 or NPP-12::GFP in the gonad. Animals were grown at 20 °C except zyg-12(ts) mutants, which were grown at 25 °C. (Scale bars, 20 μm.) (B) Quantification of the fluorescence intensity of GFP::NPP-22 or NPP-12::GFP in subcellular compartments of proximal oocytes (mean ± SD). Alleles used: tn1757 as ooc-5(–), jf18 as sun-1(+)I, and ct350 as zyg-12(ts). (C) Model. OOC-5 promotes LINC complex function, which damages nuclei with a compromised nuclear lamina.
Lifespan Analysis. All of the strains were maintained at 20 °C. Ten mid-L4 hermaphrodites were transferred onto NGM plates freshly seeded with OP50. The plates were scored for viability every 1 to 3 d, either by movement or tapping them with a platinum wire. The wild-type animals were transferred to NGM plates from the same batch every other day for a week to avoid overcrowding by their progeny. The animals that crawled on the plate sides were censored (SI Appendix, Table S2).

Immunostaining. The gonads of day 1 adult hermaphrodites were dissected in egg salt buffer fixed in 1% paraformaldehyde for 5 min and postfixed in cold methanol (−20 °C) for 1 min. Then, the samples were blocked with 1 mg/mL bovine serum albumin (BSA) in 1× phosphate-buffered saline containing 0.1% Tween 20 (PBT) overnight at 4 °C. Primary antibody incubation was performed in PBT + BSA overnight at 4 °C, while secondary antibody incubation was performed in PBT + BSA for 1.5 to 2 h at room temperature. Gonads were stained with the following antibodies provided by Abby Dernburg, University of California, Berkeley, CA: goat anti-SYP-1 (1:4,000), rabbit anti-ZIM-3 (1:1,000), guinea pig anti-HIM-8 (1:250), and guinea pig anti-HTP-3 (1:300). The guinea pig anti-SUN-1(SBP1) (1:700) was provided by Verena Jantsch, Max Pertuz Labs, Vienna, Austria, and the rabbit anti-RAD-51 (1:30,000) was provided by Sarit Smolickov, University of Iowa, Iowa City, IA. Secondary antibodies were Cy3-conjugated goat anti-mouse (1:500, Jackson Immunoresearch), Cy3-conjugated donkey anti-chicken IgY (1:1,000, Jackson Immunoresearch), and DyLight 488-conjugated donkey anti-guinea pig (1:800, Thermo Fisher Scientific). DAPI was used to detect DNA, and Vectashield (Vector Laboratories) was used as a mounting medium.

Image Acquisition and Processing. Images of day 1 adult hermaphrodites, anesthetized with 0.1% levamisole on 2% agarose pads, were acquired on a Nikon T2 inverted microscope equipped with an epifluorescence, high-performance, achromat objective correction (Apo) infrared (IR) 60× and high power (HP) Apo total internal reflection fluorescence 100× high-magnification oil immersion objective lenses, numerical aperture (NA) 1.27 and 1.49 respectively. Additional details of image acquisition and processing are detailed in SI Appendix. To examine the colocalization of SUN-1::mRuby and ZYG-12::GFP, we used Spinning Disk Super Resolution by Optical Pixel Reassignment (SoRA). The 0.30-μm step z-series images were acquired in a Nikon T2 inverted microscope equipped with a super-resolution HP plan Apo lambda S 100× silicon oil immersion objective lens, NA 1.35. Further details of image acquisition and processing are described in SI Appendix. The brightness and contrast of the images shown in Fig. 4 were adjusted for presentation identically across all of the strains.

Time-Lapse Acquisition and Processing. Hermaphrodites were selected as mid-L4 larvae and incubated at 20 °C. About 24 h later, young adults were anesthetized in 10 mM levamisole and mounted on 2% agarose pads. Nuclear collapse in the oocytes of lmn-1 mutants was documented by acquiring every 0.4-μm step z-series every 20 s. Extended depth of field (EDF) projections were created with the NIS Elements software. The movement of SUN-1 foci was documented as previously reported with the following modifications (50): The 0.25-μm step z-series were acquired approximately every 15 s for 6 to 7 min. Maximum intensity projections were made with ImageJ (Fiji), the images were converted to grayscale and their Lookup Tables were adjusted. Individual SUN-1 foci were followed using StackReg (51) and manual tracking plug-ins. The speed was determined for 84 to 390 individual foci per animal, and the number of animals assessed was equal to or greater than three per strain. All time-lapse images were acquired with the Apo IR 60× water immersion, 1.27 NA objective.

Quantitative and Statistical Analyses. The statistical analyses described below were performed using JMP Pro v14 software (Statistical Discovery, SAS) or Prism 8 (GraphPad Software, LLC). The statistical analysis details (n value, P values, confidence intervals, and measures of central tendency) for each experiment are either stated below or in the respective figures, supplemental materials, and figure legends. Kaplan-Meier survival curves followed by Wilcoxon tests were performed for the lifespan data. The Wilcoxon tests of individual pair comparisons for all of the genotypes had a P < 0.001, except the lmn-1(m1502) vs. lmn-1(m1502); +/+mIn1 pair, which had a P = 0.119. For the suppression of sterility phenotype data, a contingency analysis of outcome by genotype was performed (SI Appendix, Fig. S2A). Standard chi square tests were conducted to compare the data from each mutant to those of their respective control [e.g., lmn-1(–);ooc-5(–);mIn1 was compared to lmn-1(–);mIn1+]. SUN-1(SBP1), SUN-1 foci number and speed of movement, and the position of unpaired chromosomes data were tested for equality of group variance. Since they all had unequal variances, nonparametric comparisons were made for each pair using the Wilcoxon method. A one-way ANOVA, followed by Dunnett’s comparisons, was performed to analyze the start of the SUN-1(SBP1) region, using the N2 data as control. The comparison of fluorescence intensity of NPP-12::GFP or GFP::NPP-22 in different subcellular compartments between genetic backgrounds was performed using Prism 8 (GraphPad Software, LLC). A two-way ANOVA showed that the effect of the different genetic backgrounds was significant, P < 0.001. Bonferroni multiple comparison tests were performed using the fluorescent intensity of the nucleoporins in the otherwise wild-type background as controls. Fig. 4B shows the P values for these comparisons in the NE.

Data Availability. All data generated in this study are found in the main text or SI Appendix. Further information and requests for resources and reagents should be directed to and will be fulfilled by D.G. (greend59@umn.edu). We gladly share all C. elegans strains generated in this study.

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