The epigenetic regulator SIRT7 guards against mammalian cellular senescence induced by ribosomal DNA instability

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Edited by John M. Denu

In the yeast Saccharomyces cerevisiae, genomic instability in rDNA repeat sequences is an underlying cause of cell aging and is suppressed by the chromatin-silencing factor Sir2. In humans, rDNA instability is observed in cancers and premature aging syndromes, but its underlying mechanisms and functional consequences remain unclear. Here, we uncovered a pivotal role of sirtuin 7 (SIRT7), a mammalian Sir2 homolog, in guarding against rDNA instability and show that this function of SIRT7 protects against senescence in primary human cells. We found that, mechanistically, SIRT7 is required for association of SNF2H (also called SMARCA5, SWI/SNF-related matrix-associated actin-dependent regulator of chromatin, subfamily A, member 5), a component of the nucleolar heterochromatin-silencing complex NoRC, with rDNA sequences. Defective rDNA–heterochromatin silencing in SIRT7-deficient cells unleashed rDNA instability, with excision and loss of rDNA gene copies, which in turn induced acute senescence. Mounting evidence indicates that accumulation of senescent cells significantly contributes to tissue dysfunction in aging-related pathologies. Our findings identify rDNA instability as a driver of mammalian cellular senescence and implicate SIRT7-dependent heterochromatin silencing in protecting against this process.

Cellular senescence is a state of permanent cell cycle arrest that is induced by diverse types of stress (1–3). Replicative senescence of primary mammalian cells occurs as a result of exhaustive cell division, during which erosion of telomere sequences eventually triggers senescence as a DNA damage response (4). Senescence can also be induced by cellular stress associated with oncogene activation, DNA damage, or chromatin deregulation and can have tumor-suppressive effects (4). However, senescent cells also have profound deleterious effects that enhance tumor malignancy or contribute to tissue dysfunction in aging and disease. Indeed, senescent cells undergo dramatic alterations in metabolic and gene expression profiles with acquisition of a “senescence-associated secretory phenotype” (SASP) (1, 5). Through the SASP, even relatively low levels of senescent cells (<20%) can have far-ranging noncell-autonomous effects that influence tissue function (2).

The evolutionarily conserved sirtuin family of proteins is an important group of lysine deacetylase and deacetylases enzymes (6, 7). The founding member of this family, Saccharomyces cerevisiae silent information regulator-2 (Sir2), promotes chromatin silencing through histone deacetylation at telomeres, mating-type loci, and ribosomal DNA (rDNA) genes (8, 9). Interest in Sir2 greatly increased with the finding that it protects against yeast cell senescence by preventing genomic instability at rDNA genes, a large family of gene repeats that is highly prone to recombination (9, 10). As the most abundant and highly transcribed gene family in eukaryotes, rDNA sequences pose a significant challenge for the replication machinery (11). They are particularly susceptible to replication stress, and stalled replication forks at rDNA give rise to DNA double strand breaks (DSBs). Recombinational repair of rDNA DSBs can lead to deletions or expansions via unequal sister chromatid exchanges (12).

In the human genome, rDNA genes comprise ~350 copies distributed in large clusters (13). As in yeast, mammalian rDNA genes are prone to instability, and recombination among repeats can lead to expansions, contractions, or translocations (11). Thus, maintaining rDNA stability is a serious challenge for genome integrity, and rDNA instability is a potential driving force of genomic instability in cancer. Indeed, rDNA sequences are “hot spots” for DNA DSBs and recombinational instability in adult solid tumors (14, 15). rDNA instability is also observed in cells from patients with Bloom syndrome and ataxia telangi...
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Results

SIRT7 protects against cellular senescence and is localized at silent rDNA repeats

Given the aging-related phenotypes observed in SIRT7-deficient mice, we investigated a potential role of SIRT7 in protecting against senescence of primary human cells. We found that shRNA depletion of SIRT7 from WI38 primary human fibroblasts led to substantial accumulation of senescent cells within days after SIRT7 depletion (Fig. 1A). Levels of the senescence marker p16 were also increased in the SIRT7-deficient cell cultures (data not shown). Notably, the acute onset of senescence occurred much earlier than replicative senescence induced by telomere shortening, and a similar increase in senescent cells was observed when the WI38 cells were first immortalized by expression of hTERT telomerase subunit, which counteracts telomere shortening (Fig. 1B). These data indicate that SIRT7 protects against a novel trigger of cellular senescence that is independent of telomere shortening.

Because SIRT7 is enriched in nucleoli where rDNA genes are located, we hypothesized that SIRT7 might function like yeast Sir2 in guarding against senescence by preventing rDNA instability. To test this model, we first explored SIRT7 association with rDNA regulatory sequences using chromatin immunoprecipitation (ChIP) assays in U2OS cells (which allowed more robust ChIP DNA purification than primary cells). These analyses revealed SIRT7 binding to multiple rDNA promoter elements, consistent with previous reports (Fig. 1C) (20, 32). The association of SIRT7 with rDNA promoters was previously linked to the function of SIRT7 in stimulating RNA PolII activity, which occurs at transcriptionally active rDNA genes (20, 32). By contrast, we sought to determine whether a fraction of the rDNA-bound SIRT7 protein might occur in the context of rDNA silencing at heterochromatic rDNA gene clusters.

Because traditional ChIP assays cannot distinguish between silent versus active rDNA, we used a modified protocol, termed “ChIP–CHOP” (32), to quantify SIRT7 occupancy specifically at silent rDNA (Fig. 1D). Digestion of ChIP DNA within rDNA genes by the SmaI methylation–sensitive restriction enzyme selectively cleaves within active, but not silent, rDNA copies, and subsequent PCR with primers that bridge the SmaI site amplifies only silent rDNA, whereas downstream primers detect total rDNA. This assay revealed that over 60% of rDNA-bound SIRT7 is present at silent rDNA (Fig. 1D). As a control, the NoRC protein SNF2H was almost exclusively detected at silent rDNA, as previously shown, consistent with its known function in maintaining rDNA heterochromatin structure (19). Together, these data suggested that, in addition to its activity in stimulating PolII transcription at active rDNA genes, SIRT7 has a novel function at silent rDNA clusters.

SIRT7 acts as a scaffold to stabilize SNF2H protein at rDNA promoters for chromatin silencing

Previous proteomic studies have identified components of the NoRC complex as potential SIRT7-interacting proteins, but the physiologic relevance of these interactions was not known (33, 34). To confirm these interactions, we carried out coimmunoprecipitation assays of both FLAG-tagged and endogenous...
SIRT7 proteins. This analysis revealed specific binding of SIRT7 to both the SNF2H and TIP5 subunits of NoRC (Fig. 1, E and F, and data not shown). Intriguingly, these experiments also revealed that overexpression of SIRT7 leads to increased SNF2H protein levels (Fig. 1, E and G). To verify the generality of this finding, we also confirmed this effect in two primary cell lines (RPE-I and WI38) (Fig. 1G). These observations suggested that SIRT7 might function as a scaffold that stabilizes SNF2H protein. To test this possibility, we asked whether the catalytic activity of SIRT7 is required for increasing SNF2H levels. Overexpression of the catalytically inactive H187Y SIRT7 protein (22) had the same effect on SNF2H levels as wildtype (WT) SIRT7 (Fig. 1G). Moreover, cycloheximide chase experiments revealed a longer half-life of SNF2H protein in SIRT7-overexpressing cells (Fig. S1). Together, these data suggest that interaction of SIRT7 with SNF2H enhances SNF2H stability, and increasing SIRT7 expression is sufficient to increase cellular levels of SNF2H protein.

We next asked whether, conversely, a decrease in SIRT7 can reduce SNF2H levels. We found that depletion of SIRT7 (by shRNA depletion in human cells or gene-targeted knockout in mouse embryonic fibroblasts) led to modest decreases in SNF2H protein (Fig. 1H). However, the magnitude of this effect was quite variable, possibly due to cell type–specific compensatory mechanisms. Alternatively, it is possible that the stabilizing effect of SIRT7 on SNF2H protein is only limiting in nucleoli, which could be masked by nonnucleolar SNF2H complexes. We therefore asked whether SIRT7-deficient cells show decreased SNF2H levels specifically in nucleoli or rDNA loci. First, quantitative SILAC analysis of nucleolar protein extracts from SIRT7-deficient or control cells revealed that SIRT7 depletion leads to decreased nucleolar levels of SNF2H (data not shown), suggesting that SIRT7 stabilizes SNF2H in nucleoli. To further support this notion, we next investigated the effects of SIRT7 depletion on rDNA heterochromatin formation and rDNA transcription rates. These analyses revealed that SIRT7 depletion led to decreased rDNA heterochromatin formation and increased rDNA transcription rates (Fig. 1I and J). Together, these data suggest that SIRT7 regulates rDNA instability–induced senescence by maintaining rDNA heterochromatin and guarding against cellular senescence.
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not shown). We then examined SNF2H levels specifically at rDNA promoters by ChiP analysis, which revealed a dramatic loss of SNF2H occupancy in SIRT7-depleted cells (Fig. 1I). Similar results were observed for TIP5 (data not shown). Together, these observations identify a novel, nonenzymatic scaffolding function of SIRT7 that is required for maintaining SNF2H levels at silent rDNA genes.

SNF2H functions as a component of NoRC, which promotes heterochromatin silencing at rDNA by recruiting histone-modifying and DNA methylation enzymes. Thus, inactivation of NoRC by RNAi leads to changes in histone marks and DNA methylation patterns that reflect heterochromatin loss (19). We asked whether previously reported NoRC-dependent changes in DNA methylation and histone modifications are also observed in SIRT7-deficient cells. shRNA depletion of either SNF2H or SIRT7 led to a significant decrease in DNA methylation levels at rDNA promoters (Fig. 1J). Curiously, rDNA promoter methylation levels were lower in SIRT7-depleted cells than in SNF2H-depleted cells. This result could reflect different thresholds required for SIRT7 versus SNF2H protein or, alternatively, could indicate that SIRT7 influences rDNA methylation through additional mechanisms. SIRT7-deficient cells also showed an increase in multiple histone acetylation marks associated with active rDNA loci (Fig. 1K). Conversely, analysis of histone methylation marks associated with silent heterochromatin revealed a substantial decrease in rDNA promoter levels of H4K20me3, but not H3K9me3 (Fig. 1K), a pattern that is also observed in NoRC-deficient cells (19). The mechanistic underpinnings that distinguish H4K20me3 from H3K9me3 in this setting are still unclear, but our findings indicate that SIRT7 knockdown phenocopies multiple aspects of chromatin deregulation in NoRC-deficient cells.

SIRT7 protects against endogenous DSBs, rDNA instability, and nucleolar fragmentation

Chromatin silencing by yeast Sir2 represses both rDNA transcription and instability (35). In mammalian systems, much work on rDNA chromatin silencing has focused on rDNA transcription, but relatively little is understood about the factors that govern rDNA stability. Recently, inactivation of NoRC by RNAi depletion of its TIP5 subunit was shown to cause a loss of rDNA copies, implicating NoRC as one of the first known factors to control rDNA instability in human cells (19). When we assayed for rDNA copy number loss in SIRT7-depleted cells (36), we detected dramatically increased rDNA instability with an ~50% reduction in rDNA copy number, levels comparable with what we observed in NoRC-deficient cells (generated by shRNA depletion of SNF2H) (Fig. 2A). One consequence of rDNA recombination is thought to be an increase in nucleolar fragmentation, which has been associated with rDNA silencing defects from yeast to humans (37–39). Accordingly, SIRT7-deficient mouse embryonic fibroblasts (MEFs) and WI38 cells exhibited significant disruption in nucleolar integrity (Fig. 2B and data not shown). These data suggest that, like Sir2 in yeast, SIRT7 links maintenance of rDNA heterochromatin to rDNA stabilization.

A large body of work in yeast has provided evidence that rDNA instability results from recombinational repair of DNA DSBs that arise in rDNA due to replication stress (35). Moreover, rDNA sequences are thought to be the most fragile part of the genome in both yeast and mammalian cells (11, 14). In addition to being a hot spot for translocations in human cancers, recent work indicates that rDNA sequences are hot spots for phosphorylated γ-H2AX, a chromatin marker of DNA DSBs (40). These DSB hot spots colocalize with signs of active transcription, consistent with the model that an open chromatin state may be permissive for rDNA damage. We hypothesized that SIRT7-dependent rDNA silencing may be important for protecting against baseline rDNA damage that could otherwise arise as a result of replication stress. Consistent with this model, we found that, in SIRT7-deficient cells, γ-H2AX levels at rDNA are significantly increased (Fig. 2C). In biochemical fractionation studies, baseline γ-H2AX levels were considerably higher in nucleoli than throughout the nucleoplasm, consistent with the increased fragility of rDNA sequences, and overexpression of SIRT7 in these cells preferentially reduced nucleolar γ-H2AX levels (Fig. 2D). Notably, this analysis considerably underestimates the relative concentration of endogenous DSBs at rDNA because rDNA sequences comprise less than 0.5% of the human genome (15). Together, our results identify a novel function of SIRT7 in protecting against baseline endogenous rDNA damage and rDNA instability in mammalian cells.

dDNA instability is a trigger of mammalian cellular senescence and is regulated by SIRT7

Our data suggest that SIRT7 functions like yeast Sir2 in protecting against cellular senescence by preventing rDNA instability. However, although rDNA instability is a key driver of senescence in yeast, it has not yet been implicated in mammalian cellular senescence. We therefore set up a system to induce rDNA instability and directly test its effects on senescence. We used CRISPR/Cas9 technology to generate DNA DSBs specifically at rDNA sequences using guide RNAs homologous to 28S rDNA sequences or negative control luciferase sequence guide RNAs (which lack genomic targets) (Fig. 2E). rDNA DSB induction led to increased rDNA sequence loss and cellular senescence, similar to what we observed in cells lacking SIRT7 (Fig. 2, F and G).

Next, we sought to test directly whether rDNA silencing defects and instability are underlying causes of the senescence observed in SIRT7-deficient cells. Strikingly, we found that overexpression of SNF2H in SIRT7-depleted cells is sufficient to completely reverse both the rDNA instability and increased senescence in these cells (Fig. 2, H and I). This observation provides direct evidence that cellular senescence induced by SIRT7 loss is due to defective SNF2H-dependent rDNA silencing rather than other functions of SIRT7. We note that in these experiments SIRT7 levels were depleted transiently by siRNA knockdown, which led to a more modest change in rDNA copy number than was observed with stable shRNA knockdown of SIRT7 (Fig. 1A). Nonetheless, this level of instability was sufficient to induce acute senescence. Together, our data demonstrate that rDNA instability can be a driving force in provoking human cellular senescence and that SIRT7 plays a crucial role in preventing rDNA instability and senescence through rDNA chromatin silencing.
Discussion

Senescent cells accumulate in aging organisms and in the context of numerous aging-related chronic disease processes from osteoarthritis to diabetes (5). A growing body of work has shown that genetic or pharmacologic clearance of senescent cells in mice can extend lifespan and prevent or reverse a wide range of aging-related pathologies (41, 42). Thus, elucidating the upstream triggers and regulatory factors that control cellular senescence should provide important insights into basic mechanisms of aging and aging-related pathology. In this study, we have identified rDNA instability as an underlying trigger of senescence in mammalian cells and implicated SIRT7 as an important regulator of this process. Notably, by using CRISPR/Cas9 targeted cleavage to generate rDNA deletions, we also provide the first demonstration that rDNA instability has a direct causal role in provoking mammalian cellular senescence.

These findings fit well with models of aging mechanisms that have been proposed based on rDNA-driven senescence in yeast, collectively called the “rDNA theory of aging” (35). In this theory, rDNA sequences are more susceptible to DSBs and instability than other genomic regions and thus can serve as a sensor that induces senescence to prevent accumulation of whole-genome damage. Our data support such models, showing that a majority of baseline endogenous DSBs that accumulate in cells occur at rDNA in nucleoli, and SIRT7 is both necessary and sufficient to protect against accumulation of such rDNA DSBs.

Our findings show that, in human cells, rDNA-driven senescence occurs acutely, within days of SIRT7 depletion or targeted induction of rDNA DSBs, and is observed at levels (~25%) that are sufficient to induce tissue dysfunction through the SASP (2). These observations suggest that even transient
changes that can lead to senescence. Studies in rDNA copy number adds another layer to the epigenetic specific epigenetic landscape. SIRT7-dependent regulation of senescence in such contexts. Concentrations at rDNA could contribute to induction of response to stress (21), and a resulting decrease in local SIRT7 been shown to translocate dynamically out of nucleoli in increasing senescent cell burden. Indeed, SIRT7 has previously been suggested to interact with SNF2H at rDNA sequences. However, we cannot exclude that SIRT7 might also influence rDNA chromatin silencing by additional mechanisms. For example, SIRT7 inactivation leads to hyperacetylation of histone acetylation sites, including the known SIRT7 substrate H3K18Ac, and it is possible that SIRT7 also directly deacetylates this residue at rDNA. In addition, during preparation of our manuscript, a study from Bober and co-workers (45) suggested that SIRT7 interacts with and recruits DNA methyltransferase DNMT1 to rDNA promoters. Because NoRC recruits DNMT1 (and histone-modifying enzymes) to rDNA, our data are consistent with the model that the effects of SIRT7 on DNMT1 are mediated by SNF2H in the context of NoRC. However, our observation that SIRT7-deficient cells exhibit a greater loss of rDNA methylation than SNF2H-deficient cells is consistent with the possibility that SIRT7 interacts independently with SNF2H and DNMT1 and that the coordinated interactions provide stronger effects than each interaction alone.

rDNA stability may contribute to a number of human pathologies associated with aging. In a study of adult solid tumors, rDNA rearrangements were the most common chromosomal alterations detected and were present in over 50% of the samples analyzed (15). In addition, growing evidence suggests that rDNA instability may be linked to aging-associated disorders. Cells from patients with Werner syndrome progeria exhibit an abnormal number of palindromic rDNA repeats (46), and cells from Bloom syndrome and ataxia telangiectasia patients exhibit rDNA copy number variability and instability (16). Furthermore, age-related disorders such as Hodgkin’s, Parkinson’s, and Alzheimer’s diseases show rDNA rearrangements, nucleolar disruption, and changes in rDNA chromatin marks, respectively (47–49). Thus, our data implicating SIRT7 in rDNA silencing and stabilization suggest potential roles for SIRT7 in the pathology of these and other human disease processes.

**Experimental procedures**

**Cell culture and transfections**

Human U2OS, 293T, WI38, hTERT-WI38, and hTERT-RPE-1 were cultured in Eagle’s minimum essential medium supplemented with 10% FBS and penicillin/streptomycin. Stable retroviral and lentiviral knockdown or expression was performed as described previously (50). shRNA target sequences for SIRT7 knockdown (KD) were described previously (22). SNF2H shRNA target sequence was 5’-gtaatactctttagc4caaa-3’. rDNA guide RNA targets were caaagctcggaaggcgc (28S1) and atgaagcgcgggtaaggcgc (28S2). siRNA transfections were performed with Dharmafect reagent (Dharmacon) according to the manufacturer’s instructions. SIRT7 and control double-stranded siRNAs were purchased from Dharmacon as described previously (22).

**ChIP and quantitative real-time PCR**

ChIPs were performed in U2OS cells as described previously (51) with the following modifications for SIRT7 and SNF2H ChIPs. Briefly, cells were cross-linked with 1% formaldehyde for 25 min at room temperature, quenched with 1.25 M glycine, and washed in ice-cold PBS, and pellets were resuspended in lysis buffer (20 mM Tris, pH 8.0, 85 mM KCl, 0.5% Nonidet P-40). Nuclear pellets were obtained by centrifugation and resuspended in 50 mM Tris, pH 8.0, 10 mM EDTA, 1% SDS. Cellular lysates were sonicated and diluted in dilution buffer (10 mM Tris, pH 7.5, 140 mM NaCl, 1 mM EDTA, 0.5 mM EGTA, 1.1% Triton, 0.01% SDS). Magnetic bead–antibody complexes (coupled for >2 h at 4 °C) were added to sonicated lysates and rotated overnight at 4 °C. Rabbit anti-mouse IgG was used as a negative control. Beads were washed three times in radioimmuno precipitation assay buffer (10 mM Tris, pH 7.5, 140 mM NaCl, 1 mM EDTA, 0.5 mM EGTA, 1% Triton, 0.1% SDS, 0.1% sodium deoxycholate), and DNA was eluted and reverse cross-linked at 65 °C for 2 h using elution buffer (20 mM Tris, pH 7.5, 5 mM EDTA, 50 mM NaCl, 1% SDS, 5 μg/ml proteinase K). DNA was purified using a PCR purification kit (Qiagen).

ChIP-associated DNA was analyzed by qPCR on a LightCycler 480 II (Roche Applied Science) using SYBR Green Master Mix (Roche Applied Science). For γ-H2AX ChIPs, -fold enrichment over control was calculated as the percentage of input normalized to total H2AX. Purified DNA was diluted 10× before PCR to avoid saturation of repetitive sequences.

**qPCR rDNA copy number quantification**

DNA was isolated from U2OS cells 5–10 days after lentiviral knockdown of SIRT7 or SNF2H using a DNeasy Blood and Tissue kit (Qiagen). Purified DNA was quantified, and 10 ng was used for PCR. PCR was performed on a LightCycler 480 II using SYBR Green Master Mix. rDNA copy number was quantified by amplification of the 18S gene normalized to the rDNA13S gene. Primer sequences are listed on Table S1.

**Coimmunoprecipitation**

Cells pellets were resuspended and sonicated in Buffer A (50 mM Tris–HCl, pH 7.4, 250 mM NaCl, 0.5% Triton X-100, 10% glycerol), and lysates were incubated overnight at 4 °C with
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anti-FLAG-agarose beads (Sigma). Following five washes in Buffer A, the FLAG IPs were pelleted and eluted in Laemmli buffer.

Endogenous IP was performed similarly but under higher salt concentration (50 mM Tris-HCl, pH 7.4, 300 mM NaCl, 0.5% Triton X-100, 10% glycerol). Lysates were incubated with primary antibodies overnight, and Protein A/G-agarose beads (Thermo Scientific) were added postincubation for 2 h.

Nucleolar fractionation

Nucleolar fractionation was performed as described previously (52).

Mass spectrometry

Light and heavy labeled cell extracts (53) were combined, and nucleolar material was isolated using a sucrose gradient. Sucrose was removed by SDS-PAGE, and proteins were processed by in-gel digestion with trypsin prior to analysis by LC and tandem MS (LC-MS/MS). Briefly, samples were electrophoresed 1 cm on an SDS-polyacrylamide gel, and lanes were diced into small pieces, treated with 10 mM DTT for 30 min at 60 °C and then with 55 mM iodoacetamide for 60 min in darkness at room temperature, dehydrated with acetonitrile (ACN), and rehydrated in 50 mM ammonium bicarbonate containing 12 ng/μl trypsin (Promega). Samples were digested overnight at room temperature and stopped with 50% ACN, 5% formic acid. Peptides were extracted by dehydration in ACN, and salts were removed using a C18 Stage Tip (Thermo Scientific). LC-MS/MS was performed on an Orbitrap Fusion mass spectrometer (Thermo Scientific) using data-dependent acquisition with a 4-h HPLC gradient. Protein identification and SILAC quantification were performed using MaxQuant version 1.3.0.5 with default settings (54). Proteins quantified with a minimum of four peptides were considered for further analysis.

Senescence-associated β-gal assay

SA-β-gal activity was detected with a Senescence Cells Histochemical Staining kit (Sigma-Aldrich) according to the manufacturer’s instructions. Cells were imaged with a Leica DM5000B microscope.

DNA methylation

DNA was isolated from hTERT-RPE-1 cells 5–10 days following transduction of SIRT7 or SNF2H shRNA using a DNeasy Blood and Tissue kit. Purified DNA (250 ng) was digested overnight with Smal and purified using a PCR Purification kit. Smal-resistant rDNA was normalized to total rDNA calculated by amplification of undigested DNA.

ChIP–CHOP

ChIP was done as described above, and CHOP was done as described previously (32) with some modifications. Briefly, 10 μl of chromatin immunoprecipitated or input DNA and 2 ng of pBlueScript plasmid were mixed and digested overnight with Smal. The restriction enzyme was heat-inactivated, and DNA was purified using a PCR Purification kit. DNA methylation was detected by qPCR quantification using 2 μl of purified Smal-digested DNA on a LightCycler 480 II using SYBR Green Master Mix.

Immunofluorescence

Cells were grown in glass coverslips, fixed with 4% paraformaldehyde, washed with 1 × PBS, permeabilized with 0.1% Triton X-100 for 10 min, blocked with 2% BSA in PBS, and immunostained with primary antibodies (Table S2). Coverslips were mounted with ProLong Gold Antifade reagent with DAPI (Invitrogen) and imaged with a Zeiss LSM700 confocal laser-scanning microscope.

Author contributions—S. P. and K. F. C. conceptualization; S. P., L. T., S. M. C., M. A.-I., and W. Z. data curation; S. P. formal analysis; S. P. and K. F. C. funding acquisition; S. P., M. A.-I., L. T., S. M. C., W. Z., and T.-M. L. investigation; S. P., M. A.-I., T.-M. L., and K. F. C. methodology; S. P. writing—original draft; S. P. and K. F. C. project administration; S. P. and K. F. C. writing—review and editing.

Acknowledgments—We thank members of the laboratories of K. F. C. and O. Gozani for useful discussions.

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