A Network of Interdependent Molecular Interactions Describes a Higher Order Nrd1-Nab3 Complex Involved in Yeast Transcription Termination*

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**Background:** How the yeast proteins Nrd1 and Nab3 provoke transcription termination is poorly understood.

**Results:** An essential part of Nab3 contains a self-assembly domain that appears unstructured. Nrd1/Nab3 double mutants disrupt the function of this higher order complex, causing lethality.

**Conclusion:** A large network of molecular interactions is needed for termination.

**Significance:** A new essential function of Nab3 has been identified.

Nab3 and Nrd1 are yeast heterogeneous nuclear ribonucleoprotein (hnRNP)-like proteins that heterodimerize and bind RNA. Genetic and biochemical evidence reveals that they are integral to the termination of transcription of short non-coding RNAs by RNA polymerase II. Here we define a Nab3 mutation (nab3Δ134) that removes an essential part of the protein’s C-terminus but nevertheless can rescue, in *trans*, the phenotype resulting from a mutation in the RNA recognition motif of Nab3. This low complexity region of Nab3 appears intrinsically unstructured and can form a hydrogel *in vitro*. These data support a model in which multiple Nrd1-Nab3 heterodimers polymerize onto substrate RNA to effect termination, allowing complementation of one mutant Nab3 molecule by another lacking a different function. The self-association property of Nab3 adds to the previously documented interactions between these hnRNP-like proteins, RNA polymerase II, and the nascent transcript, leading to a network of nucleoprotein interactions that define a higher order Nrd1-Nab3 complex. This was underscored from the synthetic phenotypes of yeast strains with pairwise combinations of Nrd1 and Nab3 mutations known to affect their distinct biochemical activities. The mutations included a Nab3 self-association defect, a Nab3-Nrd1 heterodimerization defect, a Nrd1-polymerase II binding defect, and an Nab3-RNA recognition motif mutation. Although no single mutation was lethal, cells with any two mutations were not viable for four such pairings, and a fifth displayed a synthetic growth defect. These data strengthen the idea that a multiplicity of interactions is needed to assemble a higher order Nrd1-Nab3 complex that coats specific nascent RNAs in preparation for termination.

The Nrd1 and Nab3 proteins of *Saccharomyces cerevisiae* heterodimerize, bind RNA, and are important for transcription termination during the synthesis of small RNAs, such as snRNAs, small nucleolar RNAs, and cryptic unstable RNAs (1–3, 5). Both of these essential proteins possess a conserved RNA recognition motif (RRM),

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1 The abbreviations used are: RRM, RNA recognition motif; hnRNP, heterogeneous nuclear ribonucleoprotein; CTD, C-terminal domain; SC, synthetic complete; FOA, 5-fluoroorotic acid.
binant proteins and cell lysates (1, 7, 12, 13, 16, 18, 19). The physiological significance of each is underscored by mutations in the relevant domains that affect cell growth and RNA metabolism. Corden and co-workers (1) proposed that short, non-coding, RNAs destined to be terminated by this system, collect multiple Nrd1-Nab3 dimers to assemble a large nucleoprotein complex.

This assemblage serves as the core of a machinery that contacts appropriately phosphorylated RNA polymerase II and recruits termination factor Sen1 and RNA processing enzymes, such as the TRAMP complex (19–21). A complex may resemble mammalian hnRNP-C, which packages RNA into a higher order ribonucleoprotein substrate prior to further processing (22–24).

Nab3 carries interest as well because many RNA binding proteins across phyla have unusual sequences similar to the Nab3 aspartate/glutamate- and proline/glutamine-rich regions (25–27). A variety of yeast proteins contain such low complexity sequences, some of which are intrinsically unfolded and some of which form amyloidogenic, detergent-resistant complexes (25, 28). In mammals, an expansion of polyglutamine tracts due to mutation is associated with protein aggregation and neuro-pathology (e.g. in amyotrophic lateral sclerosis and Huntington disease) (29, 30). The polyglutamine regions appear to be structurally dynamic, having the ability to adopt many types of secondary structure, including extended conformations, α-helices, and β-sheets (31). The protein sequence adjacent to polyglutamine stretches also plays a role in conformation because polyglutamine tracts tend to be non-randomly distributed and frequently neighbor helical coiled coil elements (32–34). Some RNA-binding proteins with low complexity sequences form cytoplasmic foci, such as stress granules that harbor enzymes and substrates of RNA metabolism. Some of these can reversibly form dynamic fibers, including amyloid-like polymers and hydrogels (26, 27).

Here we have tested the extent of the many interactions found in this higher order complex by making compound mutations affecting more than one function of the network. Crippling two functions leads to lethality in almost all cases. We also studied the unusual and undefined structure and function of the C terminus of Nab3. We found that the last 134 residues were required for cell viability. Biochemical evidence suggests that it is intrinsically unstructured up to the α-helical hnRNP-C homology region at its very end. Although the domain is essential, co-expression of Nab3 without it rescued the mutant phenotype of another Nab3 allele with an RRM mutation, consistent with the model that multiple copies of this protein function in a single complex. These results provide new evidence that there is a constellation of protein-protein and protein-RNA interactions needed to compile a termination-competent assembly and highlight the function of the Nab3 C terminus (Fig. 1A).

**MATERIALS AND METHODS**

**Plasmid and Strain Construction**—Plasmid pET32-Nab3-134 was constructed by inserting an Xhol- and BamHI-cut PCR product (made from BY4741 genomic DNA using 5′-caaggtatccatggtagttgtgcagga-3′ and 5′-atattcggagagcaaaaattggtggctttc-3′) into similarly cut pET32a. pRS315-Nab3 was made by inserting an Xhol- and BamHI-cut PCR product (made from BY4741 genomic DNA using 5′-caaggtatccatggtagttgtgcagga-3′ and 5′-atattcggagagcaaaaattggtggctttc-3′) into similarly cut pRS315. The 34 C-terminal amino acids were deleted from Nab3 by in vitro replication of pRS315-Nab3 with Phusion DNA polymerase (New England Biolabs) and the mutagenic oligonucleotides 5′-aggtggcggaggttggtgac-3′ and 5′-tagactcccccccccttttaccttttttcttg-3′. Replicated DNA was used to transform *Escherichia coli*, and the deletion was confirmed by sequencing the resulting plasmid pRS315Nab3Δ34. Similarly, the 134 C-terminal amino acids of Nab3 were deleted by in vitro replication of pRS315NAB3 using mutagenic oligonucleotides 5′-aggttgtgagggccggacc-3′ and 5′-tagactcccccccccttttaccttttttcttg-3′ to yield pRS315Nab3Δ134.

The LEU2-marked plasmids pRS315 REF GFP, pRS315-Nab3, pRS315-Nab3Δ34, and pRS315-Nab3Δ134 were independently introduced into DY30229, a strain that lacks chromosomal NAB3 and contains the plasmid pRS316-nab3-11. The resulting strains (DY3031, DY3133, DY3131, and DY3130, respectively) were then tested for growth at 22 or 30 °C on SCura ‘leu’ plates and SC-FOA plates. The lithium acetate method of transformation was used throughout. Diploid strains were generated by mating the nab3ΔCΔ19-containing strain 1A1F (15, 16) or its otherwise isogenic wild type strain BY4742 to the *nrd1Δ151–214* containing strain, YSB2078 (13). These diploids, DY2098 and DY2097, respectively, were then transformed with pRS316-NRD1 and sporulated. Tetradis were dissected, and the resulting spores were grown on SCura– or SC-FOA. Similarly, BY4742 or 1A1F were mated to YSB2064, which bears the *nrd1Δ6–150* allele, and diploids were transformed with pRS316-NRD1 to yield strains DY2113 and DY2114, respectively. Tetradis were dissected, and spores were grown on selective media as indicated. The nab3-11-containing strain ACY1224 was mated to YSB2064 and YSB2078 to generate diploids that were each transformed with pRS316-NRD1 to yield the strains DY1627 and DY1628, respectively. These diploids were sporulated and dissected to yield haploid spores. The strains used in this study are shown in Table 1.

**Protein Purification and Analysis**—Plasmids were introduced into BL21(DE3) *E. coli*, and expression was induced with isopropylthiogalactoside for 3 h at 37 °C. Lysozyme-treated cells were broken by sonication, and the lysate was clarified by centrifugal filtration as needed. The Nab3 134-amino acid poly-gel. Flow-through protein was collected and concentrated by immobilized nickel, dialyzed into 20 mM Tris, pH 7.5, 50 mM NaCl, 1 mM EDTA, and chromatographed onto QAE-Sepha-dex. Flow-through protein was collected and concentrated by centrifugal filtration as needed. The Nab3 134-amino acid poly-peptide was cleaved from the C terminus of the thioredoxin fusion protein using thrombin. A second round of nickel chro-matography was used to isolate the His₆-tagged thioredoxin from the free Nab3 polypeptide.

Purified thioredoxin-Nab3 134-amino acid fusion protein (0.7 mg in 100 μl) was applied to a Superdex 200 10/300 column in 20 mM Tris, pH 7.5, 200 mM NaCl, using an AKTA chromatography system at a flow rate of 0.5 ml/min, and 0.5-ml fractions were collected and analyzed by SDS-PAGE. The column was calibrated with bovine thyroglobulin (670 kDa), bovine...
Higher Order Nrd1-Nab3 Complex

RESULTS

A Conserved Region of Nab3 Is Biologically Essential and Biochemically Intriguing—A spontaneous mutation that removes 19 residues from the S. cerevisiae Nab3 C terminus compensates for the deletion and provides the biologically essential function of the 5′-3′ exonuclease activity (Table 1). This conserved region contains a stretch of 41 glutamines (Gln residues) that is not conserved among the other yeast Nab3 orthologs (16). The N-terminal 19 residues of Nab3 were not conserved among the yeast Nab3 orthologs. These results suggest that the shortened Nab3 proteins accumulated and compensated for the biological function defined by the nab3-11 mutation.

Table 1: Yeast strains used in this study

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<th>Strain</th>
<th>Description</th>
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<td>Loya et al. (15)</td>
</tr>
<tr>
<td>ACY1224</td>
<td>MATa ade2 can1-100 his3-11,15 leu-3,12 trp-1 ura-3-1</td>
<td>A. Corbett (Emory University)</td>
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<td>This work</td>
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γ-globulin (158 kDa), chicken ovalbumin (44 kDa), horse myoglobin (17 kDa), and vitamin B12 (1.4 kDa). To test heat stability, 40 μg of protein in 100 μl was incubated at 22 or 100 °C for 45 min. Denatured protein was pelleted for 10 min at 13,000 × g. Supernatants were removed, concentrated by precipitation with 10% v/v trichloroacetic acid, dissolved in SDS-sample buffer, and resolved on an 11% polyacrylamide gel. Pellets were neutralized with sodium hydroxide for separation on SDS-PAGE.

Protein cross-linking was carried out as described previously (16). Briefly, protein was mixed with the indicated concentration of bis(sulfosuccinimidyl) suberate (ThermoFisher Pierce) and incubated for 30 min at 22 °C. Reactions were terminated with the addition of glycine to 50 mM, and samples were precipitated with 10% trichloroacetic acid, dissolved in sample load buffer, and neutralized with sodium hydroxide for separation on SDS-PAGE.

To test the biological role of this region, either a plasmid encoding Nab3 lacking its terminal 34 residues (the Gln16 stretch and terminal α-helix) or one encoding Nab3 lacking its terminal 134 residues (Δ134) was introduced into a yeast strain in which the chromosomal copy of Nab3 was deleted. Nab3 function was provided via a second plasmid-borne allele called nab3-11. The nab3-11 allele is a well characterized and experimentally useful pair of point mutations in the RRM domain that result in temperature-sensitive growth (12). Cells with both a control plasmid (marked with LEU2) and the nab3-11 plasmid (marked with URA3) grew at 22 but not 30 °C, confirming the conditional phenotype (Fig. 2, left panels). Cells with a plasmid encoding wild type NAB3 covering the nab3-11 allele grew at the non-permissive temperature of 30 °C (Fig. 2, bottom left). Strains with either the nab3Δ34 or the nab3Δ134 allele in addition to the nab3-11 allele also enabled growth at the non-permissive temperature (Fig. 2, bottom left). Hence, the Nab3 protein lacking its C-terminal tail and polyglutamine stretches rescued the loss of function of nab3-11. These results suggest that the shortened Nab3 proteins accumulated and compensated for the biological function defined by the nab3-11 mutation.
The 134-amino acid Nab3 peptide (17 kDa) terminus was sufficient for self-assembly, it was produced in an orthogonal system, presumably by polymerizing on RNAs to be terminated. We conclude that the terminal 134 amino acids of Nab3 allow us to try to cure the nab3-11 missense allele, these findings support the idea that multiple copies of the Nab3 protein cooperate to provide a function important for normal yeast physiology, presumably by polymerizing on RNAs to be terminated.

The development of this strain with both nab3 alleles allowed us to try to cure the nab3-11-containing plasmid marked with URA3, by using the counterselection agent 5-fluoroorotic acid to ask if yeast with either nab3Δ34 or nabΔ134 as the only version of NAB3 were viable. The strain with nab3Δ34 as its sole version of NAB3 grew, albeit slowly, at both the permissive and non-permissive temperatures, whereas cells with nab3Δ134 as their sole source of Nab3 protein could not grow at either the permissive or the restrictive temperatures (Fig. 2, right panels). We conclude that the terminal 134 amino acids of NAB3 are essential for yeast viability. Taken together with the ability of nab3Δ134 to rescue the nab3-11 allele, these findings support the idea that multiple copies of the Nab3 protein cooperate to provide a function important for normal yeast physiology, presumably by polymerizing on RNAs to be terminated.

To examine if the 134-amino acid peptide from the Nab3 C terminus was sufficient for self-assembly, it was produced in E. coli as a recombinant protein fused to the C terminus of thioredoxin-His6. The 134-amino acid Nab3 peptide (17 kDa) was proteolytically removed from the fusion protein, purified, and subjected to gel filtration. Its mobility was calculated to be 52 kDa relative to globular reference proteins. This apparent hydrodynamic size could be explained if it exists as an anomalously large dimer resulting from a relatively unfolded pair of partners (Fig. 3B). Alternatively, the molecules could represent a trimer.

Chemical cross-linking was previously used to show that a short coiled-coil segment of hnRNP-C and the homologous region of Nab3 could multimerize (16, 17). The 134-amino acid peptide was treated with increasing concentrations of bis(sulfosuccinimidyl) suberate, an amine-directed, bifunctional cross-linker, and separated on denaturing SDS-PAGE. Dimers were readily trapped by cross-linking, and small amounts of trimer could be observed at higher levels of cross-linking agent (Fig. 3C).

Perhaps due to its glutamine richness (53%), the C-terminal 34-amino acid peptide of Nab3 demonstrated unusual behaviors, such as migrating unusually slowly in SDS-PAGE (16) and staining poorly with Coomassie Brilliant Blue (data not shown). Full-length Nab3 in yeast lysates and a number of human hnRNP proteins also migrate anomalously slowly, which could be due in part to their low complexity regions, which may be unstructured and extended (6). Because the essential 134-amino acid piece of Nab3 contains multiple stretches of consecutive glutamines (two Gln9 stretches, two Gln8 stretches, and the Gln14 stretch) and 16 prolines and is poor in aromatic amino acids (two tyrosines, no phenylalanine or tryptophan), we pursued the hypothesis that this region is relatively unstructured. If so, the 134-amino acid Nab3 peptide might remain soluble following heating (37). The isolated peptide was heated

(Fig. 3D, right panel).
to 100 °C for 45 min. Soluble protein (S) was separated from denatured material (P) by centrifugation and analyzed by SDS-PAGE. The 134-amino acid Nab3 peptide remained entirely soluble following this treatment (Fig. 4A). In contrast, the identical treatment of a control protein, deoxyribonuclease I, rendered it completely insoluble. This is consistent with the proposal that this part of Nab3 lacks a complex packing of secondary structural elements.

We also used limited proteolysis to probe the unfolded nature of this region of Nab3 because unstructured domains show more susceptibility to hydrolysis than stably folded secondary structures (28, 38, 39). A recombinant fusion protein bearing the 134-amino acid Nab3 peptide was expressed and purified from E. coli, treated with chymotrypsin, and analyzed by SDS-PAGE. The Nab3 extension was digested off of the thioredoxin tag (110 kDa) with a half-life of 5 min, whereas the thioredoxin tag and a control protein (DNase I) were stable to digestion for at least 45 min (Fig. 4B) (data not shown). Similarly, the isolated Nab3134 peptide was hydrolyzed to completion more rapidly than its purified fusion partner thioredoxin when separately exposed to chymotrypsin (data not shown).

Regions of mammalian RNA binding proteins with low sequence complexity, such as one found in hnRNPA2, have been shown to undergo a phase transition at high concentration to a hydrogel-like state (27). The purified Nab3134 protein demonstrated this phenomenon when concentrated and incubated at 4 °C (Fig. 4C). This provided additional evidence for self-assembly and the hnRNP-like character of Nab3.

New Genetic Evidence for a Network of Interactions between Nrd1, Nab3, and RNA Polymerase II—A number of interactions are important for Nrd1-Nab3-based termination, including the interactions between each of their RRM domains and the nascent transcript (Fig. 1A, circled number 1), the dimerization of these two hnRNP-like proteins through mutual contact surfaces (Fig. 1A, circled number 2), the interaction of Nrd1 with the phosphorylated C-terminal domain (CTD) of RNA polymerase II (Fig. 1A, circled number 3), and the self-assembly of Nab3 through its C-terminal tail (Fig. 1A, circled number 4). Mutations in these functions have been characterized, almost all of which lead to growth defects on their own (12, 13, 16). Apparently, the network of interactions needed for termination is redundant enough to sustain mutagenic insult. We reasoned that introducing more than one of these mutations in the same cell would be additive and potentially lethal; thus, we combined them pairwise via genetic crosses to test for synthetic lethality. First, a strain that we isolated through a screen for terminator override was examined. It has a chromosomal NAB3 mutation that encodes a termination-defective version of Nab3 lacking its 19 C-terminal amino acids (Nab3C19), which is part of its self-association and hnRNP-C-like domain (Fig. 1B). It was mated to a pair of otherwise isogenic strains containing either 1) a small deletion of the Nab3-binding domain of Nrd1 (151–214) or a small deletion of the RNA polymerase II CTD-binding domain of Nrd1 (Δ6–150). Both nrd1 mutations result in defects in the biogenesis of yeast snRNA as described previously by Buratowski and co-workers (13).

The diploids NAB3/NAB3 NRD1/nrd1 151–214 and NAB3/nab3C19 NRD1/nrd1Δ151–214 were sporulated to obtain haploid cells with either the nrd1 deletion alone or both the nrd1 and nab3 mutations. Results from more than 20 tetrads showed that spores with the nrd1Δ151–214 allele were unable to germinate to form colonies even when NAB3 was wild type.
(data not shown). To overcome this technical difficulty, diploids were transformed with a plasmid-borne copy of NRD1 to cover the nrd1Δ151–214 defect. Such a strain could yield spores (Fig. 5A, top panels). The resulting haploids were subsequently plated on FOA-containing medium that selects for cells lacking the URA3-marked plasmid containing NRD1 used to rescue germination. In the absence of wild type NRD1, doubly mutant nrd1Δnab3Δ strains were inviable (Fig. 5A, bottom middle panel), whereas haploid spores with only the nrd1 mutation were able to grow (Fig. 5A, bottom left panel). This showed that retention of the NRD1-containing plasmid is essential for these double mutants and that the combination of these nrd1 and nab3 mutations was lethal.

Similarly, when the strain with the deletion that removes the Nrd1 CTD interaction domain (Δ6−150) was mated to the nab3ΔCΔ19 mutant, only the spore that had both mutations failed to grow without plasmid-borne NRD1 (Fig. 5B), whereas sister spores that were wild type for NRD1 and NAB3 or those with mutations in either NRD1 or NAB3 could grow on FOA. These results suggest that the loss of either of two different functions of Nrd1, in conjunction with a version of Nab3 lacking its self-association domain, cripples the complex to the point of inviability.

A similar pair of matings were performed between strains with the NRD1 alleles above mated to a different nab3 mutant (i.e. a previously characterized strain nab3−11) that contains a temperature-sensitive mutation in the Nab3 RRM domain (12). This resulted in another set of nrd1Δ−nab3Δ strains in which the nab3 mutation targets a functional region of the Nab3 protein distinct from the self-assembly domain. Diploids were again transformed with a plasmid bearing NRD1 in order to facilitate germination of nrd1Δ151–214-containing spores. The resulting haploids were grown at the permissive temperature of 22 °C, which supports growth of cells with the nab3−11 allele. Spores with either the nrd1 or the nab3 mutation grew on FOA-containing medium, which selects against plasmid-borne wild type NRD1, but cells with both nab3−11 and nrd1Δ151–214 were inviable (Fig. 6, right). Cells with the nab3−11 and nrd1Δ6−150 combination were very sick, taking a number of days to yield colonies (Fig. 6, left). These synthetic genetic effects emphasize the importance of both Nab3 and Nrd1 function for an effective termination complex.

The above genetic analyses tested the viability of doubly mutant cells when two different genes in the interaction network were defective. We next tested a mutant strain that lost two different functions in a single gene. It was previously shown that when two mutations in NRD1 (the aforementioned deletion alleles, Δ6−150 and Δ151–214) were present in a single copy of the gene, the double mutant could not support viability (13), suggesting that loss of both the CTD interaction and Nab3 interaction activities of Nrd1 was lethal. In this vein, we made a double mutant of NAB3 from the originally identified nab3
allele that lost its self-association domain (nab3C/H900419) and the temperature-sensitive RRM mutation (nab3-11). A LEU2-marked plasmid with these changes was transformed into a strain lacking the chromosomal copy of NAB3 but which contained a plasmid with a rescuing version of the NAB3 on a URA3-marked plasmid. At the permissive temperature, this strain’s growth was comparable with that bearing only the nab3-11 mutation (data not shown). This was the only double mutant of the set that failed to show a synthetic phenotype. The results of the double mutant crosses are summarized in Table 2.

**FIGURE 4.** Characterization of the 134-amino acid Nab3 peptide. A, heat stability. Forty μg of isolated Nab3 134-amino acid peptide or deoxyribonuclease I (Stratagene) were incubated at 22 or 100 °C for 45 min and separated into insoluble (P) and soluble (S) fractions by centrifugation. Samples were resolved on 11% SDS-polyacrylamide gels and stained with Coomassie Blue. B, time course of chymotryptic digestion of the Nab3 134-amino acid peptide fused to thioredoxin (TRX-Nab3134). Chymotrypsin was incubated with the target protein at a molar ratio of 1:400 for the indicated times before reactions were stopped and resolved on SDS-PAGE. Graphed below is a quantification of the stained bands using ImageJ software, showing loss of the full-length fusion protein (TRX-Nab3134) and accumulation of the thioredoxin tag (TRX) over time. C, gel formation. After incubation of purified Nab3134 at 4 °C, a hydrogel could be detected with the naked eye, and it is shown here after pelleting in a microcentrifuge.

**FIGURE 5.** Synthetic lethality of a nab3 tov allele (nab3C/H900419) and mutant nrd1 alleles. A, a yeast strain with the nrd1Δ151–214 (Nab3-binding domain) deletion was mated to the control strain BY4742 or an otherwise isogenic strain with the nab3C/H900419 mutation. The resulting diploids were transformed with a URA3-marked plasmid bearing wild type NRD1, sporulated, and dissected. The resulting four sister spores from a tetrad were genotyped and plated on SCura or FOA-containing medium at 30 °C. The result from the NRD1/nrd1Δ151–214 NAB3/nab3C19 diploid’s dissection are shown in the leftmost top and bottom panels. The result from the NRD1/nrd1Δ151–214 NAB3/nab3C19 diploid’s dissection are shown in the right top and bottom panels of A. B, a yeast strain with the nrd1Δ6–150 (polymerase II CTD-binding domain) deletion was mated to an otherwise isogenic strain with the nab3C/H900419 mutation. Diploids were transformed with a URA3-marked plasmid bearing wild type NRD1, sporulated, and dissected. Four sister spores from the NRD1/nrd1Δ6–150 NAB3/nab3C19 diploid were genotyped and plated on FOA-containing medium. A spore with both mutations failed to grow on FOA (top right quadrant). Sister spores with either mutant allele alone are viable.
Based upon the cooperative interaction of the Nrd1-Nab3 heterodimer with target RNA as well as the known association of Nrd1 with RNA polymerase II, Corden and co-workers (1) postulated that termination involves assembly of a nucleoprotein complex on nascent RNA in order to achieve efficiency and specificity. We previously expanded this set of interactions by showing the functional importance of a small C-terminal Nab3 domain with self-assembly properties (15, 16). Here we elaborate on the Nab3 self-assembly domain’s biological and physical functions and test the network of interactions that stabilize the complex by generating yeast strains with compound pairwise mutations in the complex’s constellation of linkages.

Our findings emphasize that there are multiple ways in which the Nrd1-Nab3 dimer can be brought to the elongation complex in preparation for termination. These include the independent recognition of RNA by each of the proteins, recognition of polymerase by Nrd1, association of the proteins with each other, and self-association of Nab3 through its C-terminal tail (Fig. 1A). This complex network of interactions could help explain why the number, spacing, and density of Nab3 or Nrd1 recognition sites on nascent transcripts is variable between different substrate RNAs (1). Prior genetic studies have shown that crippling any one of these steps negatively impacts cell growth. Three of the nrd1-nab3 double mutant strains constructed here were not viable, and one yielded viable but very slow growing cells (Table 2). The one compound mutant that failed to show a synthetic phenotype was that harboring a doubly mutant nab3 gene containing the independently isolated RRM and self-assembly region mutations. Because the RRM mutation (nab3-11) is a tight temperature-sensitive allele, any synthetic effect could only be observed at the permissive temperature. Failure to find an exacerbation of this phenotype in the double mutant could be due to the mild impact of the nab3-11 mutation at the permissive temperature. Nevertheless, other mutations paired with nab3-11 did reveal synthetic effects, indicating that the simultaneous loss of the self-assembly and RNA binding capabilities of Nab3 has a reduced consequence in the spectrum of paired defects made to the Nrd1-Nab3 system. A more quantitative dissection of the interaction network’s function must await a biochemical reconstitution of a nucleoprotein complex with full-length proteins and target RNA. It should be noted that the mutations studied here did not result in substantial losses in steady state levels of either the
Nrd1 or Nab3 proteins, excluding the possibility that their cellular concentrations were impaired by these mutations.

Loss of the Nrd1 domain that interacts with Nab3 (Δ151–214; generated by Buratowski and co-workers (13)) resulted in a particularly severe phenotype. Cells with the mutation grew slowly, and pairing it with the RRM mutation or self-assembly mutation of Nab3 was lethal. In the strain used here, diploids bearing the mutation could form an ascus, but the resulting spores that received this allele failed to germinate (data not shown). This is notable because cells lacking TRF4, which encodes a component of the TRAMP complex involved in termination and degradation of the IMD2 cryptic unstable RNA as well as other cryptic unstable RNAs (15), are also defective in spore germination (40), suggesting a role for the Nab3-Nrd1 interaction in the emergence from meiosis. High copy suppression of a nrd1 allele by NAB3 provided early genetic evidence in support of the biological importance of the Nrd1-Nab3 interaction (12).

Both mutations in NRD1 studied here were lethal when combined with the nab3CΔ19 allele that removes the Nab3 self-assembly domain, adding to the evidence that this part of Nab3 is important. Interestingly, Nrd1 also contains a polyglutamine stretch (Gln_{8}) at its C terminus (residues 567–574 of the 575-amino acid protein), and it is predicted to lie within an aggregation-prone region, similar to the Gln_{16} stretch of Nab3 studied previously and in this report (15, 16, 25). Loss of the terminal 60 amino acids of Nrd1, including this Gln_{8} tract, however, did not reveal a termination defect at the IMD2 cryptic unstable RNA, so this polyglutamine stretch does not appear to play a significant role in termination at this locus (data not shown). It is worth noting that appending a TAP tag to the C terminus of Nrd1 or Nab3 partially compromised the termination activity of each, suggesting that the addition of this relatively large extension to either protein results in steric interference that compromises their function (data not shown).

The important biological role of the curious low complexity sequences found in many RNA-binding proteins, including Nab3, has gained recent recognition (26, 27). One function of the low complexity regions is their potential to form higher order self-assemblies while escorting RNA around the cell. Indeed, low complexity sequences are associated with intrinsically disordered regions that have the propensity to form amyloid-like aggregates, including those with the potential to cause disease (25, 29). Polyglutamine stretches are often found adjacent to α-helices in domains that can multimerize, and this juxtaposition is associated with the protein’s solubility, aggregation, and function (32) We found that such a region of Nab3 has properties of a class of domains/proteins categorized as intrinsically disordered, including 1) protease sensitivity, 2) solubility after heating, 3) anomalous hydrodynamic mobility, 4) anomalous migration on denaturing gels, and 5) a high score when analyzed by an algorithm trained to distinguish order from disorder in polypeptides. Intrinsically disordered proteins can undergo a transition to an ordered state upon binding to other subunits of a complex (41). Thus, we propose that this region of Nab3 in isolation is significantly unstructured, with the notable exception of the last 18 residues, which bear structural homology to a coiled coil-forming α-helix of hnRNP-C (Fig. 1, circled number 4), but it has the potential to become structured. The ability of this domain to form a hydrogel, as seen for a number of other unstructured low complexity domains, is thought to reflect such a potential to acquire structure. In this case, it would be in the context of Nrd1, RNA polymerase II, and nascent transcript in preparation for termination. This essential segment of Nab3 looks to be important for the protein’s ability to achieve an assembly state with a specific three-dimensional configuration. Such a geometry appears important for termination and downstream RNA processing steps, including the recruitment of the Sen1 helicase, the TRAMP complex, and the nuclear exosome.

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Higher Order Nrd1-Nab3 Complex


A Network of Interdependent Molecular Interactions Describes a Higher Order Nrd1-Nab3 Complex Involved in Yeast Transcription Termination
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