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The Noc proteins involved in ribosome synthesis and export contain divergent HEAT repeats

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ABSTRACT
The Noc1–4p proteins were previously reported to be involved in intranuclear and nucleocytoplasmic transport of pre-ribosomes. Using fold recognition and structural modeling, we show that Noc1–4p are largely comprised of α-helical repeats similar to HEAT repeats. Because other HEAT-repeat proteins play key roles in transport processes, this finding provides a plausible mechanistic explanation for the function of the Noc proteins.

Keywords: rRNA; ribosome formation; ribonucleoproteins; molecular models; yeast

Eukaryotic ribosomes are assembled in the nucleolus in a series of highly coordinated events (for review, see Fatica and Tollervey 2002; Tschochner and Hurt 2003). In recent years, several reports have described around 140 nonribosomal factors involved in this multistep process (Harnpicharnchai et al. 2001; Dragon et al. 2002; Fatica et al. 2002; Grandi et al. 2002; Nissan et al. 2002). An important aspect of ribosome subunit synthesis is their transport from the nucleolus to nucleoplasm and then to the cytoplasm. Two recent reports identified a family of pre-ribosome-associated transport factors termed Noc proteins (Milkereit et al. 2001, 2003). These proteins are involved in intranuclear transport and export of the pre-60S subunit (Noc1/2/3p) and nuclear export of the pre-40S subunit (Noc4p). Moreover, Noc3p also plays a key role in the initiation of DNA replication (Zhang et al. 2002). The biochemical and genetic characterization of the Noc proteins (Milkereit et al. 2001, 2003) did not, however, reveal the mechanism(s) by which they mediate ribosomal subunit transport.

Limited sequence similarity between the Noc1/3/4p proteins over a short region of ~45 residues has been noted previously (Milkereit et al. 2001, 2003). Using profile consistency analysis (Pei et al. 2003), we have extended this alignment into a larger Noc domain (Fig. 1). This extended similarity provides further evidence that these proteins have related functions despite their nonredundancy. Orthologs of Noc proteins are present in all higher eukaryotes and were used as starting queries for PSI-BLAST searches (Altschul et al. 1997). We could not identify convincing sequence similarity to proteins of known structure or function. The search of the hidden Markov model (HMM) database of protein families (Bateman et al. 2000) resulted in Noc1p and Noc3p matching CBF/Mak21 domain (PF03914). The annotation for this domain, however, contains no additional information beyond what is already known about Noc proteins. Multiple alignments of Noc proteins were then used to train HMMs and scan the protein database, but again, no informative matches were found. We therefore resorted to fold recognition using the 3D-PSSM server (http://www.sbg.bio.ic.ac.uk/~3dpssm/; Kelley et al. 2000). With confidence in the range 70%–90%, the server returned predictions that all Noc proteins share structural similarity with proteins containing HEAT/Armadillo repeats (Andrade and Bork 1995; Andrade et al. 2001). This prediction was confirmed using the consensus of multiple fold recognition methods at the 3D-Jury metaserver (Ginalski et al. 2003). In a follow-up to this prediction, we compared Noc proteins to HMMs trained on all major classes of HEAT-repeat proteins (Andrade et al. 2001). Noc2p and Noc4p had no sequence similarity to known HEAT repeats with E < 10, while Noc1p and Noc3p had five and two HEAT repeats, respectively, with statistically insignificant E-values (E = 6.9 for Noc1p and E = 1 for Noc3p). We conclude that at the sequence level Noc proteins do not have convincing sequence similarity to any of the major classes of HEAT repeats previously described (Andrade et al. 2001).
The plausibility of the fold recognition prediction and the degree of structural similarity with known HEAT-repeat proteins were further tested by building 3D models of all Noc proteins (Sali and Blundell 1993). Shown in Figure 2 are models for Noc1p (residues 349–771) and Noc2p (residues 146–699). The models were evaluated using quality criteria for comparative modeling (Sanchez and Sali 1998). A probability ($pG$) that estimates the reliability of the overall fold of protein models was calculated for each Noc protein. Models with $pG > 0.7$ are considered to have a correct overall fold (Sanchez and Sali 1998), although they will not be correct in all details. According to these criteria, models of Noc proteins are very reliable ($pG_{\text{Noc1p}} = 0.97; pG_{\text{Noc2p}} = 0.96; pG_{\text{Noc3p}} = 0.94; pG_{\text{Noc4p}} = 0.78$). This is particularly convincing because the Noc proteins show only 8–13% identity with the templates used for modeling, and in this range of sequence identity high $pG$ values are unlikely unless the structural relationship between model and the template is genuine. The locations of the predicted HEAT repeat elements in the primary sequence of Noc1p are shown in Supplementary Figure S1 (http://www.homepage.montana.edu/~mdlakic/heat_Noc1p_suppl_FIG1.html).

In addition to the Noc proteins, we have identified four other essential HEAT-repeat proteins that are associated with yeast pre-ribosomes: Rrp12p, Sda1p, Utp10p, and Utp20p (Oeffinger et al. 2004). It was recently estimated that at least 0.2% of eukaryotic proteins have HEAT or Armadillo repeats (Andrade et al. 2001). This abundance may reflect the functional versatility of pro-
proteins with HEAT repeats. The PR65/A subunit of protein phosphatase 2A (PP2A) functions as a scaffold for assembly of the catalytic and regulatory subunits (Groves et al. 1999), while the importin-β/karyopherin-β (imp-β/kap-β) family act as molecular transporters across the nuclear envelope (Gorlich et al. 1997; Malik et al. 1997; Chook and Blobel 1999; Cingolani et al. 1999; Kobe et al. 1999; Vetter et al. 1999). Many assembly and transport steps are critical for ribosome biogenesis, potentially involving multiple HEAT-repeat proteins.

Ribosome synthesis dominates nucleocytoplasmic transport in yeast, with each nuclear pore complex (NPC) importing ∼1000 ribosomal proteins and exporting ∼25 ribosomal subunits per minute (for review, see Jorgensen et al. 2004). Efficient import of ribosomal proteins relies on multiple, partially redundant members of the imp-β/kap-β family, the founding member of which has a HEAT-repeat structure (Chook and Blobel 1999; Cingolani et al. 1999; Vetter et al. 1999). Ribosome export is also known to require a member of the imp-β/kap-β family, Crm1p/Xpo1p (for review, see Johnson et al. 2002; Tschochner and Hurt 2003), but it is unlikely that single extrinsic factor mediates the export of the very large ribosomal subunits. We therefore predict that efficient subunit export will require multiple transport factors. At least one of the other HEAT-repeat proteins we have identified, Rrp12p, is required for ribosomal subunit export (Oeffinger et al. 2004).

Here we have reported that pre-ribosomes are associated with a family of divergent HEAT-repeat proteins, which are required for ribosomal subunit transport. The Noc proteins are structurally, and potentially functionally, related to general transport factors, despite lacking detectable sequence similarity to known HEAT-repeat proteins. Future studies will determine the relative contributions of these versatile proteins to ribosome assembly and subunit transport.

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REFERENCES


