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The trypanosome Pumilio-domain protein PUF7 associates with a nuclear cyclophilin and is involved in ribosomal RNA maturation

Dorothea Droll a, Stuart Archer a, Katelyn Fenn b, Praveen Delhi a, Keith Matthews b, Christine Clayton a, * 

a Zentrum für Molekulare Biologie der Universität Heidelberg (ZMBH), DKFZ-ZMBH Alliance, Im Neuenheimer Feld 282, Heidelberg D69120, Germany 
b Centre for Immunity, Infection and Evolution, Institute of Immunology and Infection Research, School of Biological Sciences, Ashworth Laboratories, University of Edinburgh, West Mains Road, Edinburgh EH9 3JT, UK

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A B S T R A C T
Proteins with Pumilio RNA binding domains (Puf proteins) are ubiquitous in eukaryotes. Some Puf proteins bind to the 3-untranslated regions of mRNAs, acting to repress translation and promote degradation; others are involved in ribosomal RNA maturation. The genome of Trypanosoma brucei encodes eleven Puf proteins whose function cannot be predicted by sequence analysis. We show here that epitope-tagged TbPUF7 is located in the nucleolus, and associated with a nuclear cyclophilin-like protein, TbNCP1. RNAi targeting PUF7 reduced trypanosome growth and inhibited two steps in ribosomal RNA processing.

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1. Introduction
Proteins with domains of the Pumilio family (Puf) are ubiquitous in eukaryotes [1]; they are involved in various processes that require RNA binding. When several Puf domains are linked together, they form a crescent-like structure; the inner face binds to RNA while other surfaces are available for protein–protein interactions [2,3]. The genome of Saccharomyces cerevisiae encodes seven proteins with Puf domains. Five of them (ScPufs1-5) bind to, and regulate, specific subsets of mRNAs in the cytosol [4]. The remaining two, ScPuf6 and ScNop9, are predominantly nucleolar [5] (http://yeastgfp.ucsf.edu/). For Puf6, this is paradoxical because its only known function is in translational repression and localisation of the Ash1 mRNA [6,7]. The function of Nop9, in contrast, is consistent with the localisation since it plays a role in RNA maturation: it migrates with pre-ribosomes on a sucrose gradient, associates with preribosomal particles and various ribosomal assembly factors, and is required for the first steps of cleavage of the rRNA precursor [8]. Puf6 and Nop9 associate with a variety of nucleolar proteins [9,10].

Trypanosoma brucei diverged from the animal/fungal branch early in eukaryotic evolution. It has eleven Puf proteins [11]. Phylogenetic analysis for TbPUF7 (Tb11.01.6600) gave conflicting information: using only the Puf domains placed TbPUF7 together with TbPUF8 (Tb927.3.2470), ScNop9 and ScPuf6 [11]. In contrast, when we used the whole TbPUF7 sequence, instead of the Puf domains alone, the protein sequence was grouped with Saccharomyces Pufs 3, 4 and 5. Since TbPUF7 also lacks a nuclear targeting signal, we therefore suspected that it might have a function in regulation of mRNA abundance or translation. We describe here results that indicate that TbPUF7 has a function in the nucleolus.

2. Methods
2.1. Plasmid constructs
A PUF7 RNAi construct was generated using p2T7 Tablu, with the Puf7 insert and oligonucleotide primers designed using RNAit [12]. This construct was transfected into T. brucei Lister 427 single marker (‘S16’) bloodstream-form cells expressing T7 RNA polymerase and the tetracycline repressor, and transfectants selected. The PUF7 construct used for RNAi in procyclic forms was based on p2T7 Tablu but targeted a different region of the gene than that described in [11]. Primers were cz3012 (gagaagatctgcatgctAAAATGTC TCCCAGCGAC) and cz3013 (cggaattcgtcgacCGAAGAGCGCTTTAC) (restriction sites are underlined and the hybridising parts of the primers are in upper case). The NMD3 (Tb927.7.970) RNAi

* Corresponding author. 
E-mail address: cclayton@zmbh.uni-heidelberg.de (C. Clayton).
construct was made using the stem-loop strategy [13] and RNAi; this and the PUF7 RNAi plasmid were transfected into procyclic trypanosomes expressing the tet repressor and T7 polymerase [14].

To express PUF7-TAP, pHD918 [15] was modified by addition of a polylinker (Avr I–Asc I–Xma I–SpI I) to give pHD1744. The PUF7 open reading frame was amplified and cloned into the Avr I–Asc I sites. For V5 tagging (pHDI911), the plasmid used was from [16] and the primers were: ORF–cz2992 (gacctcgag TGCTCCCTTTAGTTCACTTCAA).

3.1. Sequence alignments

To investigate the phylogeny of S. cerevisiae Nop9 and Puf6, and of TbPUF7 and TbPUF8 in more detail, we used the S. cerevisiae genes to perform BLASTp searches on selected genomes from all major eukaryotic groups, then made a phylogenetic tree with these sequences and all of the major eukaryotic organisms appeared to lack Nop9. We constructed a phylogenetic tree for T. brucei and S. cerevisiae Puf proteins, and discovered that TbPUF8 was clustered with the Puf6 homologues, while TbPUF7, TbPUF10 and sometimes, TbPUF11, grouped with Nop9 proteins (not shown). A phylogenetic tree including fewer yeast and trypanosome Puf proteins is shown in Fig. 1A. Most branches were extremely poorly supported with bootstrap values below 20. The Puf6 group remained constant independent of which sequences were included, and TbPUF8 was always found within it (not shown), although the branching order was not at all robust. TbPUF7 and TbPUF10 did not branch with the Puf6’s, and their position relative to the putative Nop9 homologues changed every time the list of included proteins was altered. The Giardia protein with accession number EES98274 was consistently least related to all of the others, although the BLASTp analysis with Nop9 worked in both directions. From this analysis we could conclude that TbPUF8 could be a homologue of ScPuf6, but ScNop9 is poorly conserved in eukaryotic evolution. The status of TbPUF7 remained unresolved.

3.2. TbPUF7 is required for growth and is located in the nucleolus

For myc tagging, the TbNPC1 open reading frame (CZ2998 (gaccggatcagctcTTTCTCTTCCGCCTGGGC)) was cloned into pHDI700 [17]. Trypanosome transfection and growth analysis were as described previously [13,18].

RNA was prepared using TRIzol, denatured with formamidine and formaldehyde, and separated on denaturing formaldehyde-agarose or urea-acrylamide gels. RNA was blotted onto Nytran and hybridised using polynucleotide kinase. These were: 3′-UTR of 18S, CZ1478 (CAACACCGACACGCAACC); 7SL, CZ1478 (CAACACCGACACGCAACC). Hybridisation with oligonucleotides was as described [15] except that for the SSU rRNA, washing was at 30 °C. Probes were detected by phosphorimager.

For immunofluorescence, Cells were prepared, labeled with primary antibodies to the V5 tag (Invitrogen), the TAP tag (peroxidase-anti-peroxidase, GE Healthcare) or RNA polymerase I (kind gift from Miguel Navarro, Granada, Spain) and secondary antibodies coupled to Alexa594, 488 or 568 (Molecular Probes), as in [19].

2.2. Northern blots and immunofluorescence

PuF7-TAP was purified, and proteins identified, as described [15,20]. For immunoprecipitation, the cell lysate was obtained as for TAP, and bound to myc-(Bethyl) or V5-(Sigma) coupled beads. After washing, bound protein was eluted by boiling with reducing SDS loading buffer and analysed by Western blotting. Blots were probed with antibodies to the myc tag (Santa Cruz Biotechnology), the TAP tag (peroxidase-anti-peroxidase, GE Healthcare) or RNA polymerase I (kind gift from Miguel Navarro, Granada, Spain), and secondary antibodies coupled to Alexa594, 488 or 568 (Molecular Probes), as in [19].

2.3. Tandem affinity purification (TAP) and co-immunoprecipitation

For immunoprecipitation, the cell lysate was obtained as for TAP, and bound to myc-(Bethyl) or V5-(Sigma) coupled beads. After washing, bound protein was eluted by boiling with reducing SDS loading buffer and analysed by Western blotting. Blots were probed with antibodies to the myc tag (Santa Cruz Biotechnology), the TAP tag (peroxidase-anti-peroxidase, GE Healthcare) or RNA polymerase I (kind gift from Miguel Navarro, Granada, Spain), and secondary antibodies coupled to Alexa594, 488 or 568 (Molecular Probes), as in [19].

3. Results and discussion

3.2. TbPUF7 is required for growth and is located in the nucleolus

3.3. Depletion of TbPUF7 inhibits ribosomal RNA processing

We next analysed the pattern of rRNAs after growth-inhibitory PUF7 RNAi (Fig. 3). As a control we used cells with RNAi against NMD3. Yeast NMD3 is involved in export of large ribosomal subunits: mutations cause feedback inhibition of RNA processing [27,28]. Trypanosome NMD3 is associated with rRNA export factors [29]. A summary of rRNA processing in trypanosomes is shown in Fig. 3A [30,31]. A 9.2 kb precursor is cleaved to give a 3.4 kb small subunit (SSU) precursor, and a 5.8 kb precursor for the large
subunit (LSU) and smaller rRNAs. The 3.4 kb RNA is processed via 2.6 kb and 2.5 kb intermediates; the 5.8 kb LSU precursor is processed via 5.0 kb and 3.9 kb intermediates. The 5.8 kb large subunit precursor is visible by total RNA staining (Fig. 3B).

NMD3 RNAi inhibited growth (not shown) and large subunit processing, causing accumulation of the 5.0 kb fragment (Fig. 3B). After PUF7 RNAi, the initial processing step was inhibited such that the relative abundance of the 9.2 kb RNA increased two-to-four fold (Fig. 3B, C and D), while the 2.6 kb intermediate decreased (Fig. 3C and D). Neither RNAi affected maturation of the 5.8S rRNA, the 7SL (SRP) RNA, or the overall staining pattern of mature rRNA and small RNAs (not shown). In three bloodstream form RNAi lines, the baseline level of the 9.2 kb precursor was lower but PUF7 RNAi had the same effect as in procyclics (Fig. 3D). The results therefore indicated that PUF7 is required, either directly or indirectly, for efficient cleavage of the 9.2 kb precursor, and perhaps also for processing of the 3.4 kb pre-SSU RNA to 2.6 kb. The PUF7 RNAi did not inhibit these processes sufficiently for either to become rate-limiting in overall rRNA maturation, since neither the pre-LSU intermediates, nor steady-state SSU rRNA were affected.

Fig. 1. (A) Phylogenetic tree of possible homologues of S. cerevisiae Puf6 and Nop9. Accession numbers for all species apart from T. brucei and S. cerevisiae are shown. Each protein is designated according to the sequence sued for the BLASTp search – for example, TpPuf6 was the best match to ScPuf6. protein used for the Species abbreviations are: Tt – Tetrahymena thermophila; Pf – Plasmodium falciparum; Gi – Giardia intestinalis; Gli – Giardia lamblia; Dd – Dictyostelium discoideum; Tp – Thalassiosira pseudonana; A – Arabidopsis thaliana (official gene designations given); Hs – Homo sapiens; Tv – Trichomonas vaginalis (here there were several matches, only the best is shown). Note that the Dictyostelium Nop9 sequence gave the putative human Nop9 as the best match in the human genome, but did not give a significant match to S. cerevisiae. The tree was created in DNAStar using ClustalW; bootstrap values are from 1000 trials with a seed of 111. (B) Western blot for a procyclic trypanosome line with V5 tagged PUF7, with RNAi against PUF7. The blot was probed with antibody to the V5 tag, and to aldolase (ALD) as loading control. WT: trypanosomes expressing the tet repressor but without an RNAi construct. (C) Quantitation of PUF7 RNA and PUF7 protein after RNAi. Results are for trypanosomes without an RNAi plasmid (WT) and two experiments for the RNAi line.
In *S. cerevisiae*, the role of Nop9 in rRNA biogenesis was analysed by conditional expression under a Gal promoter. 12 h after repression, Nop9 was no longer detectable. A defect in processing of pre-rRNA to give 18S rRNA was evident from pulse-chase labelling, while production of the 25S rRNA was relatively unaffected. Steady-state levels of the 35S precursor (equivalent to the trypanosome 9.2 kb rRNA precursor) were strongly increased while the amounts of all smaller small subunit rRNA precursors, as well as the small subunit rRNA itself, were decreased [8]. The effects seen after PUF7 depletion were, in some ways, similar, but not as strong, perhaps because PUF7 down-regulation by RNAi was inefficient. Thus, although our data suggest that PUF7 has a role in rRNA maturation, we cannot conclude that it is a functional homologue of yeast Nop9.

3.4. *TbPUF7* is associated with a cyclophilin-like protein

*S. cerevisiae* Nop9 was found to be associated with a multitude of rRNA processing factors and with rRNA and pre-rRNA [8]. To find out if PUF7 was stably associated in an rRNA-processing complex, PUF7-TAP was purified. When RNA was purified from the preparations [21,22] we obtained a very low RNA yield, and reverse-transcription PCR revealed no specific association with pre-rRNA (not shown).

The gel picture of purification of PUF7-TAP was published as the control-TAP lane in Fig. 2 of [20]. Two strong bands were seen. One was tagged PUF7 (migrating at 75 kDa, MOWSE score 1323, 38 peptides, 30.7% coverage) and the other was a 34 kDa protein encoded by locus Tb927.8.2000 (MOWSE score 583, 16 peptides, 33.1% coverage).
The association appeared stoichiometric by SyproRuby staining. No other specifically-associated proteins were present. A few clusters of bands were nevertheless sequenced, and found to contain ribosomal proteins. This might be

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Fig. 3. RNAi targeting PUF7 inhibits rRNA processing. (A) Schematic overview of rRNA processing in trypanosomes. (B) Total RNA was prepared from cells with no RNAi (lane 3), RNAi targeting PUF7 (lanes 4, 5, 6, and 7) or RNAi targeting NMD3 (lanes 1 and 2). To induce RNAi the cells were grown for 48 h with tetracycline (lanes 2, 5 and 8). After denaturing agarose gel electrophoresis, RNA was blotted and stained with methylene blue. LSU: large subunit rRNA; SSU: small subunit rRNA; M: markers. (C) The blot was hybridised with a probe located 3’ of the mature SSU rRNA. (D) The intensities for the pre-rRNA, 3.4 kb, and 2.5 + 2.5 kb bands are expressed as mean ± standard deviation. Sample numbers are shown above the columns. The results shown are for procyclic cells without V5-tagged PUF7; in an experiment with the V5-tagged line, in which V5-PUF7 was decreased only two-fold, an increase in the 9.2 kb band was also seen. Using a Students t-test, the induced RNAi lines were statistically significant from the uninduced lines at the following levels: (a) <0.01; (b) <0.05; (c) <0.1.
a genuine association, but we cannot be certain because we have found ribosomal proteins as contaminants in every protein purification we have analysed so far. No proteins involved in rRNA processing or assembly were detected, although very few of these are actually known in trypanosomes. Thus our evidence so far suggests that PUF7 is not stably associated with pre-ribosomal particles.

BLASTp searches with Tb927.8.2000 yielded cytoplasmic prolyl cis–trans isomerases so we designated it nuclear cyclophilin 1 (NCP1). We confirmed the interaction between PUF7 and NCP1 by expressing NCP1 with a C-terminal myc tag in cells expressing V5-PUF7. In cells expressing both tagged proteins, precipitation with anti-V5 antibody resulted in co-precipitation of NCP1-myc (Fig. 4A, lane 6). Precipitation with anti-V5 from extracts of cells expressing NCP1-myc alone did not result in co-immunoprecipitation (Fig. 4A, lane 5). In cells expressing both tagged proteins, precipitation with anti-myc antibody resulted in co-precipitation of V5-PUF7 (Fig. 4B, lane 6), with no co-precipitation in the control (Fig. 4B, lane 5).

Inducibly expressed NCP1-myc was throughout the nucleus, and in thin inter-nuclear bridges during mitosis (Fig. 4C). In cells grown in the absence of tetracycline, a faint myc signal was still visible, again throughout the nucleus (not shown); thus the distribution was not due to over-expression. RNAi targeting NCP1 in procyclic trypanosomes had no effect on growth, but the extent of protein depletion was unknown. The distribution throughout the nucleus suggests that NCP1 may have several functions.

In other organisms, several cyclophilin-like proteins have been reported to be located in the nucleus, or to shuttle between nucleus and cytoplasm, but none is a clear NCP1 homologue. S. cerevisiae cyclophilin A has been implicated in control of meiosis [32] and of nuclear protein trafficking [33]. Arabidopsis Cyp59 interacts with SR-domain proteins and RNA polymerase II [34]; the best trypanosome match is Tb927.5.3750. Yeast Pin1 has roles in mitotic chromosome condensation [35]; Tb927.8.690 is a likely homologue.

The location of PUF7, together with the effects of RNAi, indicated that PUF7 has a role in pre-rRNA processing; but we have no evidence for stable association with pre-ribosomal particles. We speculate, therefore, that the NCP1–PUF7 complex might assist in rearrangement of the rRNA processing complex, or in reorganisation of ribosomal proteins during small subunit assembly. Inhibition of this activity could cause a feedback inhibition on earlier steps, including rRNA processing.

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References


