

Supplementary Data for:

The telomeric transcriptome of *Schizosaccharomyces pombe*

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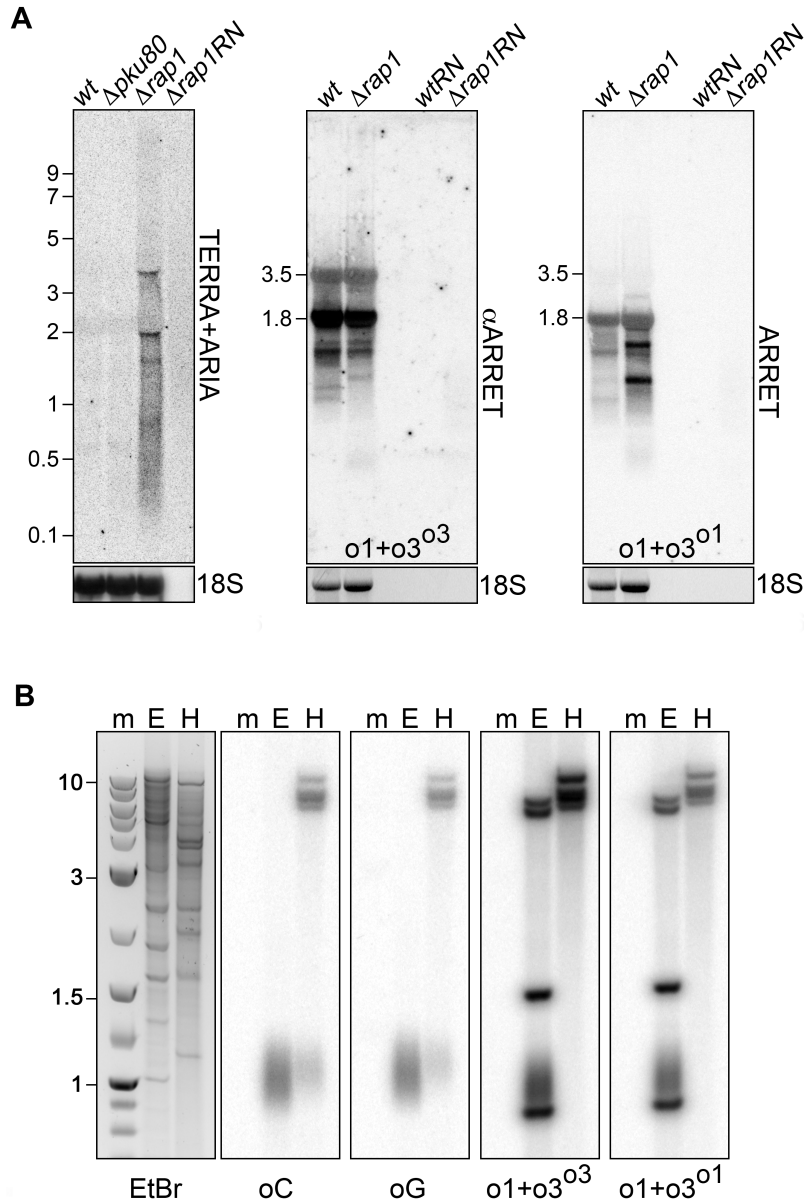


Figure S1. Control experiments for northern blot hybridizations. **(A)** Total RNA isolated from the indicated strains was treated with RNase A or left untreated prior to electrophoresis. RNA was then transferred to nylon membranes and hybridized with a double stranded telomeric probe detecting TERRA and ARIA simultaneously or with $o1+o3^{o3}$ and $o1+o3^{o1}$ probes to detect α ARRET and ARRET, respectively. To control for loading and RNase efficiency, 18S rRNA was detected by hybridization or Ethidium Bromide staining. Marker molecular weights are on the left in kb. **(B)** Genomic DNA was prepared from wt yeasts according to standard procedure. DNA was digested with *Eco*RI (E) or *Hind*III (H), electrophoresed and transferred to nylon membranes. Membranes were hybridized using the indicated probes and exposed in parallel. One typical example of Ethidium Bromide (EtBr) stained DNA is shown on the left together with marker molecular weights in kb.

Supplementary Table S1. Oligonucleotides used in this study

Name	Sequence (5'-3')	Application
o1	GAAGTTCACTCAGTCATAATTAATTGGGTAAC	PCR, RT-PCR, NB
o2	GAAGTTCACTCAGTCATAATTAATTGGGTAACGGAG	PCR, RT-PCR, NB
o3	GGGCCCAATAGTGGGGGCATTGTATTTGTG	PCR, RT-PCR, NB, 3' & 5' RACE
oC	TGTAACCGTGTAAACCACGTAACCTTGTAAACCC	PCR, RT-PCR, NB, RNA-FISH
oG	GGGTTACAAGGTTACGTGGTTACACGGTTACA	PCR, RT-PCR, NB, RNA-FISH
U6	ATGTCGCAGTGTCATCCTTG	NB
U6 ^{IN}	GTATGACTCGAACCTTGGTAAATATTGTTACTTAC	NB
35/32SA	ATTCCCAAAAAGTTAAAAGATGGAAA	NB
35/32SB	TTAGATATAATTAATTCAGACTTC	NB
18S	TCAATAAAGAACATGAGTAGTTTGCCAGAA	NB
act1	ATCCTTTTGTCCCATACCTACCATAATACC	NB
act1RT-Fw	TGCCGATCGTATGCAAAAGG	RT-PCR, CHIP
act1RT-Rv	CCGCTCTCATCATACTCTTG	RT-PCR, CHIP
Poly(T)	(T) ₂₀	RT-PCR
o5o	AGCCTCGCCTTACGGCTCGGCTGACGGGTG	5' RACE
o3i	TGAATACCGTATTTCAATTTCTATTTCTTTA	3' RACE
TSS-Fw	TATTTCTTTATTCAACTTACCGCACTTC	ChIP
TSS-Rv	CAGTAGTGCAGTGATTATGATAATTAATGATGG	ChIP

NB: northern blot