Supporting information for Green *et al.* (2003) *Proc. Natl. Acad. Sci. USA*, 10.1073/pnas.0336594100

Table 1. Yeast strains and plasmids used in this study

Strain or plasmid	Description
EGY48	MATα *ura3 his3 trp1 leu2*::*6LexAop-LEU2* plus pSH18-34 (1)
ACY192	MATa *ura3 leu2 his3 trp1* (FY23) (2)
ACY194	MATα *ura3 leu2 his3 rat7-1ts* (3)
ACY683	MATa *leu2 trp1 can1 ade2 MLP1-proteinA*::*HIS3* (CSCD15a) (4)
pSH18-34	8 ops-*lacZ, 2μ*, *URA3*
pEG202	pADH, *LexA-DBD, 2μ*, *HIS3* (1)
pJG4-5	pGAL1, *B42-AD-HA, 2μ*, *TRP1* (1)
pGEX-4T3	GST bacterial expression vector, *amp*^{r} (Amersham Pharmacia)
pAC2	pRS314, *CEN, TRP* (5)
pAC19	pGAL, *2μ*, *LEU2*
pAC213	*NLS-NES-GFP, 2μ*, *URA3* (6, 7)
pAC719	*NAB2-GFP, 2μ*, *URA3* (8)
pAC943	*NPL3-c-myc, CEN, LEU2* (9)
pAC963	*SRP1-c-myc (3X), CEN, TRP*
pAC980	Δ*RGG*(Δ600-793) -*NAB2-GFP, 2μ*, *URA3*
pAC1024	pGAL, *NLS-GFP* reporter, *2μ*, *URA3*
pAC1101	*NAB2* ORF in pEG202 to create *DBD-NAB2, 2μ*, *HIS3*
pAC1126	*NAB2-c-myc (3X), CEN, TRP*
pAC1195	*AD-MLP1*(4470-stop) in pJG4-5, *2μ*, *TRP1* (Origene)
pAC1196	pGAL, *CT-MLP1* (4470-stop), *2μ*, *LEU2*
pAC1197	pGAL, *CT-MLP1-GFP* (4470-stop), *2μ*, *URA3*
pAC1313	pGAL, *FL-MLP1* in modified pYES2 (Invitrogen), *2μ*, *URA3* (4)
pAC1315	pGAL, *ΔCT-MLP1* (1-4466) in modified pYES2, *2μ*, *URA3* , C-terminal domain was deleted by removing *Xho*I–*Xho*I fragment from pAC1313
pAC1340	*GST-CT-MLP1* (4470-stop) in pGEX-4T3, *amp*^{r}, *CT-MLP1* was created by PCR and cloned into pGEX-4T3 via *Bam*HI and *Xho*I sites

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