

**Table S1. Plasmids and Strains**

<b>Plasmid</b>	<b>Fluorescent Protein<sup>1</sup></b>	<b>Selective Marker<sup>2</sup></b>	<b>Genbank Accession</b>	<b>Strain<sup>3</sup></b>
<b>pBS41</b>	CFP	URA3	KF177452	EMY200
<b>pBS43</b>	YFP	URA3	KF177455	EMY201
<b>pBS49</b>	YFP	LEU2	KF177459	EMY202
<b>pBS42BN<sup>4</sup></b>	YFP-Pro0-CFP	URA3	KF177454	EMY203
<b>pBS46</b>	YFP-Pro5-CFP	URA3	KF177456	EMY204
<b>pBS47</b>	YFP-Pro10-CFP	URA3	KF177457	EMY205
<b>pBS48</b>	YFP-Pro15-CFP	URA3	KF177458	EMY206
<b>pBS50</b>	YFP-Pro20-CFP	URA3	KF177460	EMY207
<b>pFTR74-YFP</b>	FTR74-YFP	LEU2	KF177461	EMY208
<b>pFTR117-CFP</b>	FTR117-CFP	URA3	KF177462	EMY208
<b>pYFP-TAF28-CFP</b>	YFP-TAF28-CFP	URA3	KF177465	EMY209
<b>pTAF18<sup>5</sup></b>		LEU2	KF177463	EMY209
<b>pYFP-Lim4-CFP</b>	YFP-Lim4-CFP	URA3	KF177464	EMY210
<b>pYFP-THP112-CFP</b>	YFP-THP112-	URA3	KF177466	EMY211

<sup>1</sup> Expression of all proteins is driven by the TEF promoter. All proteins have an N-terminal nuclear localization signal.

<sup>2</sup> Plasmids were integrated into strain BSY9 (MATa/MAT $\alpha$  ade2-1oc/ade2-1oc ADE3/ade3?100 can1-100/can1-100 CYH2s/cyh2r his3-11,15/his3-11,15 leu2-3,112/leu2-3,112 trp1-1/ trp1-1 ura3-1/ura3-1 ) at URA3 by digesting the plasmids with *Stu*I, and integrated at LEU2 by digestion with *Bst*EII. Since the Lim4 gene contains a *Stu*I site, pYFP-Lim4-CFP was digested with *Apa*I instead.

<sup>3</sup> Plasmids were integrated at the selective marker.

<sup>4</sup> Contains *Bam*HI and *Nhe*I sites between YFP and CFP genes.

<sup>5</sup> TAF18 is not tagged, but co-expressed with YFP-TAF28-CFP.