

pAGH1	Ycplac33-RPL16B	URA3	product of a PCR with genomic DNA as template and primers AG9/O1235 was SpeI/PstI digested and cloned into Ycplac33
pAGH2	Ycplac33-RPL16BΔInt	URA3	product of a PCR with pAGH1 as template and primers AG9/AG10 was SpeI/Bpu10i digested and cloned into SpeI/Bpu10i digested pAGH1
pAGH3	Ycplac33- <i>rpl16b</i> Δ51	URA3	Site-directed mutagenesis using QuickChangeII Site-Directed Mutagenesis kit from Agilent with the oligos AG11/AG12 and pAGH2 as template. * product of a PCR with mutagenized pAGH3 as template and primers AG9/O1235 was subcloned into Ycplac33 to avoid other mutations in the vector backbone
pAGH4	Ycplac33- <i>rpl16b</i> Δ28	URA3	Site-directed mutagenesis using QuickChangeII Site-Directed Mutagenesis kit from Agilent with the oligos AG13/AG14 and pAGH2 as template. * product of a PCR with mutagenized pAGH4 as template and primers AG9/O1235 was subcloned into Ycplac33 to avoid other mutations in the vector backbone
Yeplac195	Yeplac195	URA3	Gietz and Sugino, Gene 74 (2), p. 527-534
TK487	Yeplac195-pRPS28-RPS24-FLAG	URA3	Ferreira-Cerca et al., 2007 (suppl. data)
pAGH5	Yeplac195-pRPS28-RPL16B-FLAG	URA3	product of a PCR with pAGH2 as template and primers AG15/AG16 was BamHI/PstI cloned into TK487
pAGH6	Yeplac195-pRPS28- <i>rpl16b</i> Δ51 -FLAG	URA3	product of a PCR with pAGH3 as template and primers AG15/AG17 was BamHI/PstI cloned into TK487
pAGH7	Yeplac195-pRPS28- <i>rpl16b</i> Δ28 -FLAG	URA3	product of a PCR with pAGH4 as template and primers AG15/AG18 was BamHI/PstI cloned into TK487
pAGH8	Ycplac33-pRPS28-RPL16B-FLAG	URA3	pvull cassette from pAGH5 was cloned into Ycplac33
pAGH9	Ycplac33-pRPS28- <i>rpl16b</i> Δ51 -FLAG	URA3	pvull cassette from pAGH6 was cloned into Ycplac33
pAGH10	Ycplac33-pRPS28- <i>rpl16b</i> Δ28 -FLAG	URA3	pvull cassette from pAGH7 was cloned into Ycplac33