## Table A1: Plasmids

Plasmid	<b>Relevant genotype</b>	Reference
Cloning Vectors		
pRS313	Cloning replicative vector, CEN HIS3Ap <sup>R</sup>	(8)
pRS425	Cloning replicative vector, $2\mu LEU2 \text{ Ap}^{\text{R}}$	(3)
pTC3	Cloning, expression vector CEN TRP1, Ap <sup>R</sup>	(1)
YGALSET351	Cloning, expression vector CEN LEU2 Ap <sup>R</sup>	(4)
YGALSET983	Cloning, expression vector $2\mu LEU2 \text{ Ap}^{\text{R}}$	(4)
YGALSET986	Cloning, expression vector CEN HIS3 Ap <sup>R</sup>	(4)
pUC19	Cloning vector Ap <sup>R</sup>	(12)
pBSIIKS+/-	Cloning vector Ap <sup>R</sup>	Stratagene
pGAD-C1-3	Expression vector, $2\mu$ LEU2 Ap <sup>R</sup>	(5)
pGBD-C1-3	Expression vector, 2µ <i>TRP1</i> Ap <sup>R</sup>	(5)
pCgGAD-C1-2	A 1.5kb <i>Sph</i> I-fragment containing the GAL4AD/Polylinker/ Terminator cassette of pGAD-C1-3 was cloned into <i>Eco</i> RI/ <i>Sal</i> I digested, blunt ended and religated pCgACT14 digested with <i>Sph</i> I	This work
pCgACT14	Cloning, replicative vector, Cg <i>CEN</i> Cg <i>TRP1</i>	(7)
pCgACH3	Cloning, replicative vector, CgCEN CgHIS3	(7)
p112-8XM	Cloning, replicative vector, CgCEN ScURA3	(7)
pGEX-2T	Cloning, expression vector, Ap <sup>R</sup>	Novagen
pGEX-4T-2	Cloning, expression vector, Ap <sup>R</sup>	Novagen
pADNS	Cloning replicative vector,	(2)

	2μ <i>LEU2</i> Ap <sup>R</sup>	
pET16b	Cloning, expression vector, Ap <sup>R</sup>	Novagen
pET28a+	Cloning, expression vector, Km <sup>R</sup>	Novagen
Plasmid clones from genomic library		
YepTS4	Yep24 containing a ~11kb Sau3a digested genomic DNA fragment in <i>Bam</i> H1 site carrying Cg <i>CEN</i> .	This work
YepTS20.1	Yep24 containing a ~14.6kb Sau3a digested genomic DNA fragment in <i>Bam</i> H1 site carrying Cg <i>NDC10</i> .	This work
YepTS15.1	Yep24 containing a ~13.7kb Sau3a digested genomic DNA fragment in BamH1 site carrying CgCEP3.	This work
YepTS5.4	Yep24 containing a ~14kb Sau3a digested genomic DNA fragment in BamH1 site carrying CgCTF13.	This work
YepTS3.3	Yep24 containing a ~9kb Sau3a digested genomic DNA fragment in BamH1 site carrying CgCTF13.	This work
YepTS34.2	Yep24 containing a ~10.9kb Sau3a digested genomic DNA fragment in BamH1 site carrying CgMIF2.	This work
YepTS27.2	Yep24 containing a ~14kb Sau3a digested genomic DNA fragment in BamH1 site carrying CgCSE4.	This work
YepTS9.1	Yep24 containing a ~13kb Sau3a digested genomic DNA fragment in BamH1 site carrying CgCBFI.	(11)
Plasmid subclones used		
for sequencing pTS22	A 4kb <i>Sph</i> I-fragment containing Cg <i>NDC10</i> (from YepTS#20.1) was cloned into pUC19.	This work

pTS26	A 3.8kb <i>PstI-SphI</i> fragment containing Cg <i>MIF2</i> (from YepTS#34.2) was cloned into pUC19.	This work
pTS38	A 3.2kb <i>Eco</i> RI/ <i>Sph</i> I fragment containing Cg <i>CEP3</i> (from YepTS#15.1) was cloned into pUC19.	This work
pTS29	A 3kb <i>Bam</i> HI/ <i>Eco</i> RI fragment containing Cg <i>CSE4</i> (from YepTS#27.2) was cloned into pUC19.	This work
pTS30	A 4kb <i>Sma</i> I-fragment containing Cg <i>CTF13</i> (from YepTS#5.4) was cloned into pUC19.	This work
Plasmids for CBF3-		
deletions	A 470 hr Kaul/VL J DCD	This work
pTS53 (Cgndc10Δ)	A 479 bp <i>KpnI/XhoI</i> PCR fragment (Primers CgA1/1 and CgA1/2), a 467bp <i>XhoI/XbaI</i> PCR fragment (Primers CgA2/1 and CgA2/2) and a 1kb <i>XhoI</i> fragment containing Cg <i>TRP1</i> (from pCgACT14) were cloned into pUC19 digested with <i>KpnI/XbaI</i> . ApR.	
pTS54 (Cg <i>ctf13</i> Δ)	A 389 bp <i>KpnI/XhoI</i> PCR fragment (Primers CgC1/1 and CgC1/2), a 437bp <i>XhoI/XbaI</i> PCR fragment (Primers CgC2/1 and CgC2/2) and a 1kb <i>XhoI</i> fragment containing Cg <i>TRP1</i> (from pCgACT14) were cloned into pUC19 digested with <i>KpnI/XbaI</i> . ApR.	This work
pTS55 (Cg <i>cep3</i> Δ)	A 458 bp <i>KpnI/XhoI</i> PCR fragment (Primers CgB1/1 and CgB1/2), a 429bp	This work

	XhoI/XbaI PCR fragment(Primers CgB2/1 andCgB2/2) and a 1kb XhoIfragment containingCgTRP1 (from pCgACT14)were cloned into pUC19digested with KpnI/XbaI.ApR.	
Vectors with Open Reading frames		
pJL33	A 1.8kb <i>NdeI/Bam</i> HI fragment containing the Sc <i>CEP3</i> ORF was cloned into pET16B.	(9)
pJL36	A 1.4kb <i>NdeI/Bam</i> HI fragment containing the Cg <i>CTF13</i> ORF was cloned into pET16B.	(9)
pTS41	A 540bp <i>Bam</i> HI/ <i>Eco</i> RI PCR fragment of Cg <i>SKP1</i> (primer GST-3D1 and GST- 3D2) was cloned into pGEX-2T.	This work
pTS43	A 1622 bp BamHI/EcoRI PCR fragment encoding for CgMIF2 (primer GST- MIF2.1 and GST-MIF2.2) was cloned into pGEX-2T.	This work
pTS45	A 2.7 kb <i>SmaI/XhoI</i> PCR fragment encoding for the ORF of Cg <i>NDC10</i> (primer GST-p110-1 and GST- p110-2) was cloned into pGEX-4T-2.	This work
pTS52	A 1.8kb <i>SmaI/XhoI</i> PCR fragment encoding for the ORF of Cg <i>CEP3</i> (primer GST-3B-1.2 and GST-3B- 2.1) was cloned into pGEX- 4T-2 digested with <i>SmaI- XhoI</i> .	This work
pTS56	A 1.4 kb <i>Bam</i> HI/ <i>Eco</i> RI PCR fragment encoding Cg <i>CTF13</i> was cloned into pGEX2-T digested with	This work

	BamHI/EcoRI.	
pTS57	A 1.8kb <i>Bam</i> H1/ <i>Xho</i> I fragment of Cg <i>CEP3</i> (from pTS52) was cloned into pRSETA.	This work
pTS60	A 1 kb <i>Bam</i> HI/ <i>Cla</i> I- fragment from pTS45 was replaced with WT sequence from pTS22.1 to remove a point mutation in the gene.	This work
pTS62	A 2.7kb <i>Smal/XhoI</i> fragment containing the Cg <i>NDC10</i> ORF was cloned into pUC19 digested with <i>Smal/Sal</i> I.	This work
pTS66	A 1.4kb <i>Bam</i> HI/ <i>Eco</i> RI fragment containing the Cg <i>CTF13</i> ORF was cloned into pET28a+.	This work
pTS74	A 1.8kb <i>Bam</i> H1/ <i>Xho</i> I fragment encoding for Cg <i>CEP3</i> (from pTS52) was cloned into pUC19 digested with <i>Bam</i> H1/ <i>Sal</i> I.	This work
pTS75	A 1.4kb <i>Bam</i> H1/ <i>Sac</i> I fragment containing the Cg <i>CTF13</i> ORF 9from pTS66) was cloned into pUC19.	This work
pTS77	A 2.7kb <i>SacI/PstI</i> fragment containing the Cg <i>NDC10</i> ORF (from pTS62) was cloned into pBSKS+/–.	This work
pCgGAD-CgCBF1	A 1.3kb <i>Bam</i> HI PCR fragment of Cg <i>CBF1</i> (from pET-Cg1/2) was cloned into pCgGAD-C2.	This work
pCgGAD-Cg <i>SKP1</i>	A 540bp <i>NcoI/Bam</i> HI PCR fragment of Cg <i>SKP1</i> was cloned into pCgGAD-C2.	This work
pET-Cg1/2	A 1.3kb <i>Bam</i> HI PCR fragment of CgCBF1 was cloned into pET28a+. A 431bp <i>MscI/Hind</i> III fragment was replaced with	(11)

	a 1kb <i>MscI/Hind</i> III	
Plasmids for studying the species specificity in <i>S. cerevisiae</i> and <i>C. glabrata</i>	fragment from pRSCg1	
pWJ110B	A 4.2kb <i>Pvu</i> II fragment was cloned into pBSKS+/– digested with <i>Sma</i> I	(6)
pJL33		J. Lechner, personal communic.
pJL36		J. Lechner, personal communic.
pMB038 (ScP-ScMIF2)	A 1.9kb <i>Sph</i> I fragment containing the Sc <i>MIF2</i> gene was cloned into pTC3.	(1)
pYHYCbf1	PRS314 containing the Sc <i>CBF1</i> gene on a SphI fragment.	H.Y. Yoon, personal communic.
p112-Cp1	A 2.1 kb blunt ended SpeI/PstI fragment containing CgCBF1 was cloned into p112-8XM.	(11)
pTS32 (CgP-Cg <i>NDC10</i> )	A 4kb blunt ended <i>Sph</i> I- fragment from pTS22.1 containing Cg <i>NDC10</i> into pRS425 digested with <i>Sma</i> I.	This work
pTS33 (CgP-Cg <i>CSE4</i> )	A 3kb blunt ended <i>Xba</i> I- fragment containing Cg <i>CSE4</i> from pTS29 into pRS425 digested with <i>Sma</i> I.	This work
pTS34 (CgP-Cg <i>MIF2</i> )	A 3.5kb blunt ended <i>SphI/PstI</i> -fragment containing the Cg <i>MIF2</i> gene was cloned into <i>SphI</i> digested and blunt ended pMB038.	This work
pTS36	A 5.5kb <i>SphI/Sma</i> I fragment containing Cg <i>CEP3</i> was cloned into pUC19.	This work
pTS37 (CgP-Cg <i>CTF13</i> )	A 4kb <i>Sma</i> I fragment containing Cg <i>CTF13</i> from pTS30 was cloned into pRS313.	This work

pTS39 (CgP-Cg <i>CEP3</i> )	A 3.4kb blunt ended <i>Eco</i> RI- <i>Sph</i> I fragment containing Cg <i>CEP3</i> was cloned into pRS425 digested with <i>Sma</i> I.	This work
pRSCg3D (CgP-Cg <i>SKP1</i> )	A 986bp <i>Eco</i> RI/ <i>Bam</i> HI PCR fragment of Cg <i>SKP1</i> (primer CgD3 and CgD2) was cloned into pRS425.	(10)
pRSCg1 (CgP-Cg <i>CBF1</i> )	A 2.2 kb <i>SpeI/Hind</i> III fragment carrying Cg <i>CBF</i> 1 was cloned into pRS425.	(11)
pTS46 (CgP-CgNDC10)	A 4kb <i>Sph</i> I blunt ended fragment containing Cg <i>NDC10</i> (from pTS22) was cloned into p112-8XM digested with <i>Sma</i> I.	This work
pTS48 (CgP-Cg <i>CEP3</i> )	A 3.4kb blunt ended <i>Eco</i> RI/ <i>Sph</i> I fragment containing Cg <i>CEP3</i> was cloned into p112-8XM digested with <i>Sma</i> I.	This work
pTS50 (CgP-Cg <i>CTF13</i> )	A 4kb <i>Sma</i> I fragment containing Cg <i>CTF13</i> from pTS30 was cloned into p112-8XM.	This work
pTS71 (ScP-ScNDC10)	A 4kb <i>Bam</i> HI/ <i>Pst</i> I fragment containing the Sc <i>NDC10</i> gene (from pWJ110p) was cloned into pCgACH3.	This work
pTS125 (CgP-ScNDC10)	A 600bp PCR fragment (Primer CgA4 and CgA5.2) encoding for the Cg <i>NDC10</i> - Promoter was digested with <i>SmaI/Bam</i> H1 and ligated into pTS123 digested with <i>SmaI/Bam</i> H1.	This work
pTS128	A 4000bp <i>SmaI/PstI</i> fragment from pTS125 encoding for the Cg <i>NDC10</i> promoter and the Sc <i>NDC10</i> gene was ligated into pCgACH3 digested with <i>SmaI/PstI</i> .	This work
pTS72 (CgP-Sc <i>CEP3</i> )	A 480 bp <i>SmaI/NdeI</i> PCR fragment encoding for the	This work

	Cg <i>CEP3</i> promoter (primer3B1 and 3B2), a 900bp <i>Bam</i> HI/ <i>Pst</i> I PCR fragment encoding for the Cg <i>CEP3</i> terminator (primer 3B3 and 3B4), and a 1.8 kb <i>NdeI/Bam</i> HI fragment containing the Sc <i>CEP3</i> ORF were cloned into pCgACH3 digested with <i>SmaI/Pst</i> I.	
pTS73 (CgP-Sc <i>CTF13</i> )	A 480bp <i>SmaI/AseI</i> PCR fragment encoding for the Cg <i>CTF13</i> promoter (primer 3C1 and 3C2), a 500bp <i>BamHI/PstI</i> PCR fragment encoding for the Cg <i>CTF13</i> terminator (primer 3C3 and 3C4), and a 1.4 kb <i>NdeI/BamHI</i> fragment containing the Sc <i>CTF13</i> ORF were cloned into pCgACH3 digested with <i>SmaI/PstI</i> .	This work
pTS76 (GalP-Cg <i>CEP3</i> )	A 1.8 kb <i>SacI/PstI</i> -fragment containing the Cg <i>CEP3</i> ORF (from pTS74) was cloned into YGALSET983 digested with <i>SacI/PstI</i> .	This work
pTS78 (GalP-Cg <i>NDC10</i> )	A 2.7 kb <i>Kpn</i> I fragment carrying Cg <i>NDC10</i> (from pTS60) was cloned into YGALSET351 digested with <i>Kpn</i> I.	This work
pTS79 (ScP-Cg <i>CBF1</i> )	A 1.3 kb <i>Bam</i> HI/ <i>Hind</i> III fragment containing Cg <i>CBF1</i> (from pETCg1/2) was cloned into pADNS digested with <i>Hind</i> III and blunt ended.	This work
pTS81 (CgP-Sc <i>CBF1</i> )	A 339bp <i>SacI/Hind</i> III PCR fragment encoding for the Cg <i>CBF1</i> - Promoter and the first 8 aa of the Sc <i>CBF1</i> gene (Primer Cg1-P.1 and	This work

	Cg1-P.2), and a 1031bp <i>Hind</i> III/ <i>Xho</i> I-fragment	
	containing the Sc <i>CBF1</i>	
	ORF and 744bp 3'-	
	nontranslated region were	
	cloned into pCgACH3 digested with <i>SacI</i> and	
	HindIII.	
	A 1.8 kb BamHI/HindIII-	
	fragment encoding for the	
pTS85	Cg <i>CBF1</i> -ORF (from pETCg1/2) and the	This work
	3'nontranslated region was	
	cloned into pBSIIKS+/	
	A 1.6 kb <i>Bam</i> HI/ <i>Eco</i> RI	
	fragment encoding for the	
pTS86	Cg <i>MIF2</i> -ORF (from	This work
	pTS43) was cloned into	
	pBSIIKS+/	
	A 1.8kb blunt ended <i>NdeI/Bam</i> HI fragment	
	containing Sc <i>CEP3</i> (from	
pTS96 (GalP-Sc <i>CEP3</i> )	pJL33) was cloned into	This work
	YGALSET983 digested	
	with PvuII.	
	A 1.4 kb blunt ended	
	Ndel/BamHI fragment	
pTS98 (GalP-ScCTF13)	containing ScCTF13 (from	This work
	pTS36) was cloned into YGALSET986 digested	
	with <i>Pvu</i> II.	
	A 4kb <i>Bam</i> HI/ <i>Pst</i> I fragment	
pTS100 (GalP-Sc <i>NDC10</i> )	containing ScNDC10 (from	This work
	pWJ110p) was cloned into	
	YGALSET983.	
	A 1.4kb SacI fragment	
pTS109 (GalP-CgCTF13)	containing Cg <i>CTF13</i> (from pTS75) was cloned into	This work
	YGALSET983.	
	A 3.5kb blunt ended <i>SpeI</i> -	
	fragment containing	
pTS120 (ScP-ScCBF1)	ScCBF1 was cloned into	This work
	pCgACH3 digested with	
Expression vestors for	SmaI.	
<b>Expression-vectors for</b>		

two-hybrid analysis		
pTS88 (GBD-Cg <i>NDC10</i> )	A 2.7 kb <i>SmaI/XhoI</i> fragment containing Cg <i>NDC10</i> (from pTS60) was cloned into pGBD-C1 digested with <i>SmaI/SaI</i> I.	This work
pTS89 (GBD-Cg <i>CEP3</i> )	A 1.8 kb <i>SmaI/XhoI</i> fragment containing the Cg <i>CEP3</i> ORF (from pTS 52) was cloned into pGBD- C1 digested with <i>SmaI/SaI</i> I.	This work
pTS90 (GBD-Cg <i>CTF13</i> )	A 1.4kb <i>Bam</i> HI/ <i>Sal</i> I fragment containing Cg <i>CTF13</i> (from pTS66) was cloned into PGBD-C1.	This work
pTS91 (GBD-Cg <i>CBF1</i> )	A 1.8kb <i>Bam</i> HI/ <i>Sal</i> I fragment containing Cg <i>CBF1</i> (from pTS85) was cloned into pGBD-C1.	This work
pTS92 (GBD-Cg <i>MIF2</i> )	A 1.6kb <i>Bam</i> HI/ <i>Sal</i> I fragment containing the Cg <i>MIF2</i> ORF(from pTS 43) was cloned into pGBD- C1.	This work
pTS93 (GBD-Cg <i>SKP1</i> )	A 540bp <i>Eco</i> RI/ <i>Cla</i> I fragment containing Cg <i>SKP1</i> (from pCgGAD- 3D) was cloned into pGBD- C1.	This work
pTS101 (GAD-Cg <i>CEP3</i> )	A 1.8 kb <i>SmaI/XhoI</i> fragment containing the Cg <i>CEP3</i> ORF (from pTS 52) was cloned into pGAD- C1 digested with <i>SmaI/SaI</i> I.	This work
pTS102 (GAD-Cg <i>CTF13</i> )	A 1.4kb <i>Bam</i> HI/ <i>Sal</i> I fragment containing Cg <i>CTF13</i> (from pTS66) was cloned into PGAD-C1.	This work
pTS103 (GAD-Cg <i>MIF2</i> )	A 1.6kb <i>Bam</i> HI/ <i>Sal</i> I fragment containing the Cg <i>MIF2</i> ORF(from pTS 43) was cloned into pGAD- C1.	This work
pTS104 (GAD-Cg <i>CBF1</i> )	A 1.8kb <i>Bam</i> HI/ <i>Sal</i> I fragment containing	This work

	Cg <i>CBF1</i> (from pTS85) was	
	cloned into pGAD-C1.	
	A 540bp <i>Eco</i> RI/ <i>Cla</i> I	
	fragment containing	
pTS105 (GAD-Cg <i>SKP1</i> )	Cg <i>SKP1</i> (from pCgGAD-	This work
	3D) was cloned into pGAD-	
	C1.	
	A 2.7 kb SmaI/XhoI	
	fragment containing	
pTS106 (GAD-Cg <i>NDC10</i> )	Cg <i>NDC10</i> (from pTS60)	This work
	was cloned into pGAD-C1	
	digested with SmaI/SalI.	
pTS110 (GAD-CgCSE4)	A 722bp <i>EcoRI/Cla</i> I PCR-	This work
	fragment of CgCSE4 was	
	cloned into pGAD-C1.	
pTS111(GBD-Cg <i>CSE4</i> )	A 722bp <i>EcoRI/Cla</i> I PCR-	This work
F(	fragment of CgCSE4 was	
	cloned into pGBD-C1.	
pTS115 (GAD-Cg <i>CEP3-</i> aa	PTS101 was digested with	This work
1-470)	<i>Xbal/PstI</i> , blunt ended and	
1 1/0)	religated.	
pTS119 (GAD-CgCEP3-	A 1.7 kb <i>KpnI-XhoI</i>	This work
aa 38-611)	fragment of pTS101	
	containing CgCEP3 was	
	cloned into pGAD-C1	
	digested with <i>Bam</i> HI and	
	blunt ended.	
pAM5 (GAD-CgCEP3-aa	A 1600bp PCR fragment	This work
87-611)	encoding for aa 77-611 of	
	Cg <i>CEP3</i> (PrimersCg3B.1	
	/CgB5.2)was digested with	
	<i>Eco</i> RI and <i>Bam</i> H1 and	
	ligated into pGAD-C3.	
	Amino acid sequence	
	identity was confirmed by	
	sequencing the DNA.	
pAM1 (GBD-ScCTF13)	A 1437bp <i>NdeI/Bam</i> H1	This work
	fragment of pJL36 encoding	
	the Sc <i>CTF13</i> ORF was	
	blunt ended and ligated into	
	pGBD-C3 digested with	
	Smal.	
pAM2 (GAD-ScCEP3)	A 1827bp NdeI/BamH1	This work
	fragment of pJL33 encoding	
	the Sc <i>CEP3</i> ORF was blunt	
	the Sector 5 OKI' was bluilt	

	ended and ligated into	
	pGAD-C3 digested with	
	Smal.	
pAM3 (GBD-ScCEP3)	A 1827bp NdeI/BamH1	This work
	fragment of pJL33 encoding	
	the ScCEP3 ORF was blunt	
	ended and ligated into	
	pGBD-C3 digested with	
	SmaI.	
pAM4 (GAD-ScCTF13)	A 1437bp NdeI/BamH1	This work
	fragment of pJL36 encoding	
	the ScCTF13 ORF was	
	blunt ended and ligated into	
	pGAD-C3 digested with	
TC100	Smal.	
pTS122	A 3393bp <i>Cla</i> I-fragment	This work
	from pWJ110p was ligated	
	into pGAD-C3 digested	
pTS123	with <i>Cla</i> I.	This work
p13123	A 3393bp <i>Cla</i> I-fragment from pWJ110p was ligated	
	into pGBD-C3 digested	
	with <i>Cla</i> I.	
Vectors for ChIP analysis		
pTS124	A 1572bp HpaI (blunt	This work
1	ended)/HindIII fragment	
	from pTS30 encoding for	
	part of the CgCTF13 gene,	
	and a 5331bp	
	<i>Hind</i> III/ <i>Eco</i> RI (blunt	
	ended) fragment from	
	pTS126 encoding for the	
	HA-tag was ligated into	
	pCgACH3 digested with	
7010	Smal.	
pTS126	A 485bp PCR-fragment	This work
	encoding for part of the	
	Cg <i>CTF13</i> gene (Primers	
	<b>e</b> 1	
	-	
	(Primers CgC6.3 and	
	CgC6.1/CgC6.2) was ligated into pCR2.1-TOPO (Invitrogen). The resulting plasmid was digested with <i>KpnI/NdeI</i> and ligated with a 324bp PCR-fragment (Primers CgC6.3 and	

	O(O(5) = 1 = 1.0	
	CgC6.5) released from a	
	TOPO-vector with <i>Kpn</i> I,	
	encoding for the 3'nt region	
	of the CgCTF13 gene, and a	
	1100bp PCR-fragment	
	(Primers HA-F/CgHis3R on	
	template pRSCg1HA,	
	Stoyan et al., 2001)	
	released from a TOPO	
	vector with KpnI/NdeI.	
pTS129	A 996bp PCR-fragment	
r	(Primers CgA6.1.2/CgA6.2)	
	was digested with	
	<i>Bg/III/Nde</i> I, and a 160bp	
	<i>Ndel/Kpn</i> I fragment from	
	pTS124 were ligated into	
	pTS22 digested with	
	BglII/KpnI.	
pTS130	A 435bp PCR-fragment	
p15150	(Primers CgB6.1.2/CgB6.2)	
	was digested with	
	0	
	<i>XbaI/Nde</i> I, and a 160bp	
	<i>Ndel/Eco</i> RI fragment from	
	pTS124 were ligated into	
	pTS38 digested with	
T0121	Xbal/EcoRI.	
pTS131	A ~3500bp SphI/KpnI	
	fragment from pTS129	
	encoding for CgNDC10-HA	
	was blunt ended and ligated	
	into pCgACH3 digested	
	with SmaI.	
pTS132	A ~2500bp <i>SphI/Eco</i> RI	
	fragment from pTS130	
	encoding for CgCEP3-HA	
	was blunt ended and ligated	
	into pCgACH3 digested	
	with SmaI.	

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- Brown, M., Goetsch, L., and L. Hartwell. 1993. *MIF2* is required for mitotic spindle integrity during anaphase spindle elongation in *Saccharomyces cerevisiae*. J. Cell Biol. 123:387-403.
- Colicelli, J., Birchmeier, C., Michaeli, T., O'Neill, K., Riggs, .M. and M. Wigler. 1989. Isolation and characterization of a mammalian gene encoding a high-affinity camp phosphodiesterase. Proc Natl Acad Sci USA 86:3599-3603.
- Christianson, T. W., Sikorski, R. S., Dante, M., Shero, J. H., and P. Hieter. 1992. Multifunctional yeast high-copy-number shuttle vectors. Gene 110:119-122.
- Enomoto, S., Chen, G. and Berman, J. 1998. Vectors for expressing T7 epitopeand His6 affinity tagged fusion proteins in *S. cerevisiae*. Biotechniques 24:782-788.
- James, P., Halladay, J., and E. A. Craig. 1996. Genomic libraries and a host strain designed for highly efficient two-hybrid selection in yeast. Genetics 144:1425-1436.
- Jiang, W., Lechner, J., and J. Carbon. 1993. Isolation and characterization of a gene (CBF2) specifying a protein component of the budding yeast kinetochore. J. Cell Biol. 121:513-519.
- Kitada, K., Yamaguchi, E., and M. Arisawa. 1996. Isolation of a *Candida glabrata* centromere and its use in construction of plasmid vectors. Gene 175:106-108.

- Sikorski, R. S. and P. Hieter. 1989. A system of shuttle vectors and yeast host strains designed for efficient manioulation of DNA in Saccharomyces cerevisiae. Genetics 122: 19-27.
- Stemmann. O. and J. Lechner. 1996. The Saccharomyces cerevisiae kinetochore contains a cyclin-CDK complexing homologue, as identified by in vitro reconstitution. EMBO J. 15:3611-3620.
- Stoyan, T., Eck, R., Lechner, J., Hemmerich, P., Kuenkel, W., and S. Diekmann 1999. Cloning of a centromere binding factor 3D (*CBF3D*) gene from *Candida glabrata*. Yeast 15:793-798.
- 11. Stoyan, T., Gloeckner, G., Diekmann, S., and J. A. Carbon. 2001.
  Multifunctional centromere binding factor 1 (Cbf1) is essential for chromosome segregation in the human pathogenic yeast *Candida glabrata*. Mol. Cell .Biol. 21:4875-4888.
- 12. Yannish-Perron, C., Vieira, J., and J. Messing. 1985. Improved M13 phage cloning vectors and host strains: Nucleotide sequences of the M13 mp18 and pUC19 vectors. Gene 33:103-119.