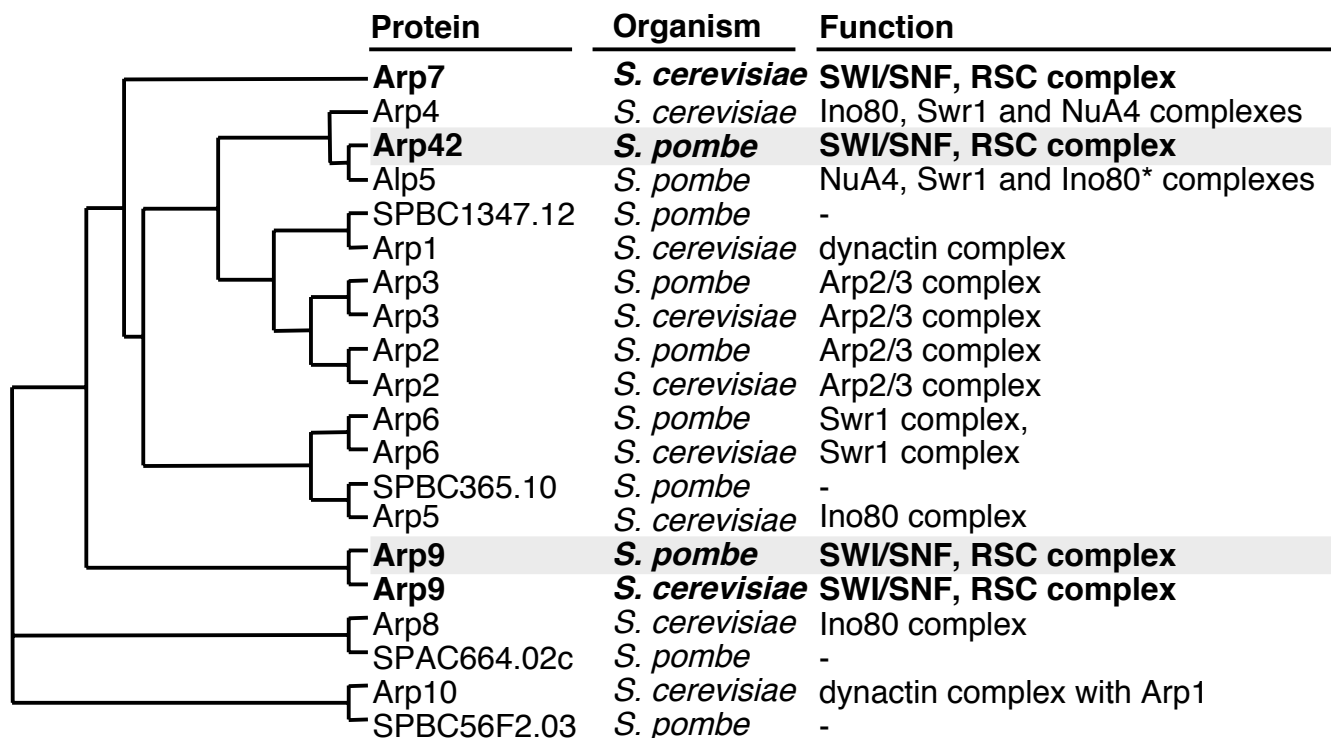
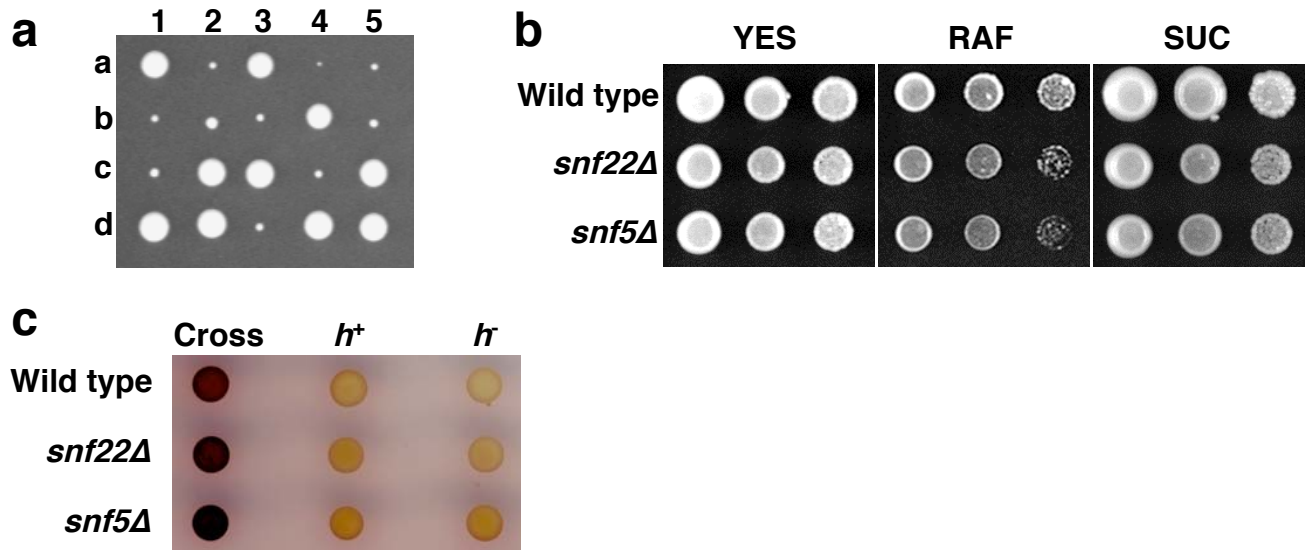

SUPPLEMENTARY MATERIAL.

Fission yeast SWI/SNF and RSC complexes show compositional and functional differences from budding yeast

B.J. Monahan, J. Villén, S. Marguerat, J. Bähler, S.P. Gygi, and F. Winston.



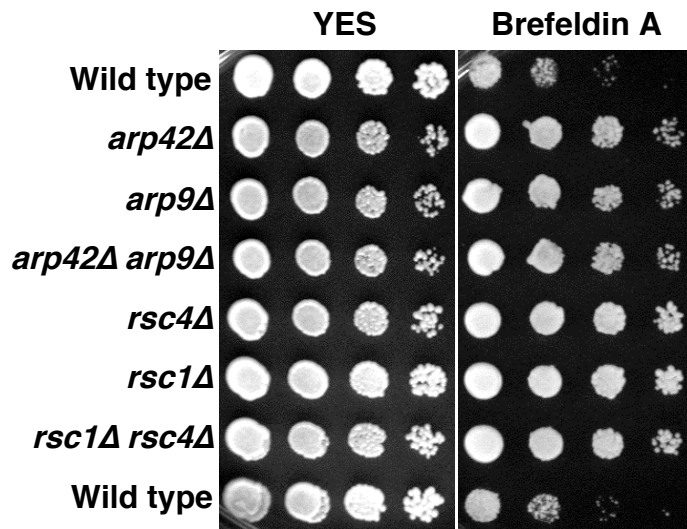
Supplementary Figure 1. *S. pombe* actin related protein (ARP) family. A dendrogram showing relatedness of the ten *S. pombe* and *S. cerevisiae* ARPs. *S. pombe* does not contain a gene encoding an Arp7 homolog but does contain two apparent Arp4 paralogs, named Alp5 and Arp42. ARPs present in SWI/SNF and RSC are highlighted in bold with the *S. pombe* complexes shaded in grey boxes. Putative function or association of each ARP is indicated where applicable. The tree was constructed by the neighbor-joining method with ClustalX⁴.



Supplementary Figure 2. Deletion analysis of *S. pombe* SWI/SNF and RSC genes (a) The *S. pombe rsc58Δ* mutant displays a severe growth phenotype. Shown are progeny after sporulation and tetrad dissection of a *rsc58*⁺/*rsc58Δ* diploid. The four progeny of each tetrad are labeled a-d. The small colonies are *rsc58Δ* and the large colonies are *rsc58*⁺. (b and c) *S. pombe* SWI/SNF mutants do not display phenotypes characteristic of *S. cerevisiae* SWI/SNF mutants. In contrast to phenotypes noted for *S. cerevisiae* SWI/SNF mutants *S. pombe snf22Δ* and *snf5Δ* mutants display wildtype growth on media containing sucrose or raffinose as the sole carbon source (b). (c) *S. pombe snf22Δ/snf22Δ* and *snf5Δ/snf5Δ* mutants display normal sporulation although a small proportion of asci have incorrect spore number (data not shown). Iodine staining of mating plates is shown. As a negative control, staining of the *h*⁺ and *h*⁻ parent strains is also shown.

SYSTEMATIC	Protein	Arp42:TAP		Arp9:TAP		no tag	
		Exp 1	Exp 2	Exp 1	Exp 2		
RSC	SPAC1250.01	Snf21	116	93	108	97	0
	SPBC4B4.03	Rsc1	52	43	54	41	0
	SPBC1703.02	Rsc9	55	49	46	44	0
	SPBC1734.15	Rsc4	44	34	46	31	0
	SPCC16A11.14	Sfh1	22	15	20	17	0
	SPAC1F3.07c	Rsc58	22	21	19	21	0
	SPCC1281.05	Rsc7	24	20	25	19	0
Shared	SPAC17G6.10	Ssr1	45	37	49	35	0
	SPAC23H3.10	Ssr2	37	35	36	31	0
	SPAC23G3.10c	Ssr3	47	44	48	38	0
	SPBP23A10.05	Ssr4	30	29	31	28	0
	SPAC1071.06	Arp9	37	25	37	34	0
	SPAC23D3.09	Arp42	31	30	33	30	0
Swi/Snf	SPCC1620.14c	Snf22	54	41	39	41	0
	SPBC30B4.04c	Sol1	28	27	24	28	0
	SPAC2F7.08c	Snf5	22	16	20	18	0
	SPBC26H8.09c	Snf59	29	29	22	25	0
	SPAC23G3.07c	Snf30	9	8	6	7	0
	SPAC22H12.02	Tfg3	1	2	0	2	0
Ino80	SPAC29B12.01	Ino80	0	0	0	0	0
	SPBC365.10	Arp5	0	0	0	0	0
	SPAC664.02C	Arp8	0	0	0	0	0
	SPBC28F2.11	(Nhp10)	0	0	0	0	0
Swr1	SPAPB8E5.09	Rvp1	1	0	0	0	1
	SPBC83.08	Rvp2	0	0	1	0	0
	SPBP23A10.08*	Alp5	0	0	0	0	0
	SPBC32H8.12c*	actin	19	9	21	10	11
	SPAC11E3.01c	Swr1	0	0	0	0	0
	SPCC550.12	Arp6	0	0	0	0	0
	SPBP35G2.13C	Swc2	0	0	0	0	0
	SPAC9G1.13C*	Swc4	0	0	0	0	0
	SPCC576.13	Swc5	0	0	0	0	0
	SPAC17G8.07*	(Yaf9)	0	0	0	0	0
NuA4	SPBC29A3.05	(Vps71)	0	0	0	0	0
	SPAC637.12C	Mst1	0	0	0	0	0
	SPCC1795.08C	(Eaf1)	0	0	0	0	0
	SPAC23H4.12	Alp13	0	0	0	0	0
	SPAC6F6.09	(Eaf6)	0	0	0	0	0
	SPBC16A3.19	Eaf7	0	0	0	0	0
	SPCC830.05C	Epl1	0	0	0	0	0
	SPBC1709.11C	Png2	0	0	0	0	0
	SPBP16F5.03C	Tra1	0	0	0	0	0

Supplementary Figure 3. The *S. pombe* actin related proteins Arp42 and Arp9 are specific to the SWI/SNF and RSC chromatin remodeling complexes. Mass-spectrometry analysis of duplicate Arp42-TAP and Arp9-TAP purifications are shown. The number of unique peptides identified in the mass-spectra for the respective protein for each purification is shown. To illustrate specificity of Arp42 and Arp9, various proteins predicted to be members of the *S. pombe* chromatin remodeling or modifying complexes known to contain ARPs, namely NuA4, Swr1 and Ino80, are shown. The name of the respective complex is indicated on the left. Asterisks represent proteins that are also putative components of the NuA4 complex.



Supplementary Figure 4. *S. pombe* RSC mutants show brefeldin A resistance. Wild type (FWP52), *arp42Δ* (FWP239), *arp9Δ* (FWP240), *arp42Δ arp9Δ* (FWP249), *rsc1Δ* (FWP234), *rsc4Δ* (FWP242), and *rsc1Δ rsc4Δ* (FWP250) strains were grown in liquid YES medium to stationary phase, subjected to 10-fold serial dilutions, and spotted onto solid YES medium containing 10 $\mu\text{g ml}^{-1}$ brefeldin A.

Supplementary Table 1. Summary of BLASTP analysis of the *S. pombe* protein set searching for RSC (a) and SWI/SNF (b).

(a)

<i>S. cerevisiae</i> RSC complex			<i>S. pombe</i> BLASTP top hit(s) ¹			
protein	systematic	size (aa)	protein	systematic	size (aa)	e-value
Sth1	YIL126W	1359	Snf21	SPAC1250.01	1199	7E-262
			Snf22	SPCC1620.14c	1680	3E-253
Rsc1	YGR056W	928	Rsc1	SPBC4B4.03	803	1E-56
Rsc2	YLR357W	889	Rsc1	SPBC4B4.03	803	1E-58
Rsc3	YDR303C	885		SPAPB1A11.04c	697	2E-06
Rsc30	YHR056C	832	Moc3	SPAC821.07c	497	2E-05
Rsc4	YKR008W	625	Rsc4	SPBC1734.15	542	3E-17
Rsc9	YML127W	581	Rsc9	SPBC1703.02	780	1E-23
Rsc8	YFR037C	557	Ssr2	SPAC23H3.10	503	4E-73
			Ssr1	SPAC17G6.10	527	5E-64
Rsc58	YLR033W	502		SPBC215.07c	568	6E-02
Rsc6	YCR052W	483	Ssr3	SPAC23G3.10c	425	5E-15
Arp7	YPR034W	477	Arp42	SPAC23D3.09	430	4E-27
Arp9	YMR033W	467	Arp9	SPAC1071.06	523	3E-21
Rsc7	YMR091C	435	Rsc7	SPCC1281.05	390	2E-46
Sfh1	YLR321C	426	Sfh1	SPCC16A11.14	418	1E-33
Rtt102	YGR275W	186	Swr1	SPAC2H10.03c	1288	7E-02
Ldb7	YBL006C	145		No hits, E < 0.01		
Htl1	YCR020W-B	78	Mde1	SPBC31F10.08	209	3E-02

(b)

<i>S. cerevisiae</i> SWI/SNF complex			<i>S. pombe</i> BLASTP top hit(s) ¹			
protein	systematic	size (aa)	protein	systematic	size (aa)	e-value
Snf2	YOR290C	1703	Snf22	SPCC1620.14c	1680	6E-279
			Snf21	SPAC1250.01	1199	2E-231
Swi1	YPL016W	1314	Sol1	SPBC30B4.04c	865	7E-24
Snf5	YBR289W	905	Snf5	SPAC2F7.08c	632	8E-45
Swi3	YJL176C	825	Ssr2	SPAC23H3.10	503	9E-60
			Ssr1	SPAC17G6.10	527	3E-51
Swp82	YFL049W	623	Rsc7	SPCC1281.05	390	1E-11
Snf12	YNR023W	566	Ssr3	SPAC23G3.10c	425	1E-22
Arp7	YPR034W	477	Arp42	SPAC23D3.09	430	4E-27
Arp9	YMR033W	467	Arp9	SPAC1071.06	523	3E-21
Snf6	YHL025W	332		No hits E < 0.01		
Taf14	YPL129W	244	Tfg3	SPAC22H12.02	241	3E-48
Rtt102	YGR275W	186	Swr1	SPAC2H10.03c	1288	7E-02
Snf11	YDR073W	169	-	SPBC660.06	273	1E-03

¹ Bold indicates that the protein was subsequently identified as a component of the respective complex.

Supplementary Table 2. Deletion analysis of *S. pombe* SWI/SNF and RSC genes and comparison to *S. cerevisiae*

<i>S. pombe</i>		<i>S. cerevisiae</i>	
<u>subunit^a</u>	<u>deletion phenotype</u>	<u>ortholog</u>	<u>deletion phenotype</u>
Snf22	viable	Snf2	viable
Snf5	viable	Snf5	viable
Sol1	viable	Swi1	viable
Tfg3	viable	Taf14	viable
Snf59	viable	Swp82	viable
Snf30	viable		
Ssr1	inviable	Swi3 ^b , Rsc8 ^c	<i>swi3Δ</i> viable, <i>rsc8Δ</i> inviable
Ssr2	inviable	Swi3 ^b , Rsc8 ^c	<i>swi3Δ</i> viable, <i>rsc8Δ</i> inviable
Ssr3	inviable	Snf12 ^b , Rsc6 ^c	<i>snf12Δ</i> viable, <i>rsc6Δ</i> inviable
Ssr4	inviable		
Arp42	viable		
Arp9	viable	Arp9	inviable or sick ^d
Snf21	inviable	Sth1	inviable
Sfh1	inviable	Sfh1	inviable
Rsc9	inviable	Rsc9	inviable
Rsc7	inviable	Rsc7	viable
Rsc1	viable	Rsc1, Rsc2	double mutant inviable
Rsc4	viable	Rsc4	inviable
Rsc58	viable	Rsc58	inviable

^aThe top group of subunits in the unshaded boxes is in *S. pombe* SWI/SNF, the middle shaded group is shared between SWI/SNF and RSC, and the bottom unshaded group is in *S. pombe* RSC.

^bPart of *S. cerevisiae* SWI/SNF.

^cPart of *S. cerevisiae* RSC.

^dThe null phenotype depends upon the genetic background¹.

Supplementary Table 3. Mass-spectrometry analysis of RSC and SWI/SNF purifications in the presence or absence of Arp42 and Arp9. Shared components are within grey box with RSC-specific subunits above and SWI/SNF-specific subunits below. This analysis also indicated that no other *S. pombe* ARP or actin substitute for either *arp42Δ* or *arp9Δ*. The 10 *S. pombe* ARPs and actin(Act1) are shown in red.

protein	RSC (Snf21-TAP)				SWI/SNF (Snf21-TAP)				controls	
	350 mM NaCl		150 mM NaCl		350 mM NaCl		150 mM NaCl		350	150
	<i>arp+</i>	<i>arpΔ</i>	<i>arp+</i>	<i>arpΔ</i>	<i>arp+</i>	<i>arpΔ</i>	<i>arp+</i>	<i>arpΔ</i>		
Snf21	103 ^a	123	130	109	10	7	9	9	2	1
Rsc1	44	45	53	47	0	0	0	1	0	0
Rsc9	50	47	60	45	2	3	0	1	0	0
Rsc4	35	36	46	36	0	0	0	0	0	0
Sfh1	16	15	20	20	1	0	0	0	1	0
Rsc58	17	18	23	22	1	0	0	0	0	1
Rsc7	21	24	27	21	0	0	0	0	1	1
Ssr1	39	46	44	38	42	44	38	40	1	0
Ssr2	37	38	39	31	32	33	29	38	0	2
Ssr3	50	50	52	46	40	40	40	41	0	0
Ssr4	27	27	32	26	20	20	24	26	0	1
Arp9	24	0	31	0	25	1	23	3	2	1
Arp42	29	0	30	0	30	0	23	0	0	1
Snf22	0	1	0	0	96	90	99	97	0	0
Sol1	0	0	0	0	38	40	40	50	0	0
Snf5	0	0	0	0	47	35	39	33	0	0
Snf59	0	0	0	0	62	60	52	54	0	0
Snf30	0	0	0	0	20	18	17	12	0	0
Tfg3	0	0	0	0	9	9	5	11	0	0
<i>Alp5</i>	0	0	0	0	0	0	0	0	0	0
<i>Arp1</i>	0	0	0	0	0	0	0	0	0	0
<i>Arp2</i>	0	0	0	0	0	0	0	0	0	0
<i>Arp3</i>	0	0	0	0	0	0	0	0	0	0
<i>Arp5</i>	0	0	0	0	0	0	0	0	0	0
<i>Arp6</i>	0	0	0	0	0	0	0	0	0	0
<i>Arp8</i>	0	0	0	0	0	0	0	0	0	0
<i>Arp10</i>	0	0	0	0	0	0	0	0	0	0
<i>Act1</i>	25	19	21	22	28	25	34	23	25	9

^aNumber of unique peptides for the respective protein.

Supplementary Table 4. Microarray data of SWI/SNF mutants showing the fold-change in mRNA levels for the hexose and iron transporter genes and their known repressors (in blue).

gene	<i>snf22</i>Δ	<i>snf5</i>Δ	Description
<i>fip1</i> ⁺	4.3	3.2	Iron permease FTR1 family
<i>frp1</i> ⁺	15.0	10.4	Ferric reductase transmembrane component
<i>fio1</i> ⁺	2.5	2.2	iron transport multicopper oxidase
<i>str1</i> ⁺	2.2	2.1	Siderophore iron transporter
<i>str2</i> ⁺	1.7	1.5	Siderophore iron transporter
<i>str3</i> ⁺	8.1	14.1	Siderophore iron transporter
<i>ght1</i> ⁺	2.5	2.9	hexose transporter
<i>ght2</i> ⁺	2.0	1.6	hexose transporter
<i>ght3</i> ⁺	5.3	4.4	hexose transporter
<i>ght4</i> ⁺	11.4	10.3	hexose transporter
<i>ght5</i> ⁺	3.6	3.7	hexose transporter
<i>ght6</i> ⁺	4.3	3.0	hexose transporter
<i>ght7</i> ⁺	0.9	1.3	hexose transporter
<i>ght8</i> ⁺	3.4	2.5	hexose transporter
<i>fep1</i> ⁺	0.9	1.0	GATA transcription factor iron repressor
<i>scr1</i> ⁺	1.4	1.3	Mig1/CreA-like carbon catabolite repressor
<i>tup11</i> ⁺	1.1	1.1	transcriptional corepressor
<i>tup12</i> ⁺	1.2	1.2	transcriptional corepressor

Supplementary Table 5. *S. pombe* strains used in this study

Strain	Genotype	Source
FWP52	<i>h⁺ leu1-32 ade6-m216</i>	Lab stock
FWP165	<i>h⁺ ura4-D18 leu1-32 ade6-m216</i>	Lab stock
FWP172	<i>h⁻ ura4-D18 leu1-32 ade6-m210</i>	Lab stock
FWP253	<i>h⁺ fep1::ura4⁺ ura4-D18 leu1-32 ade6-M210 his7-366</i>	Pelletier <i>et al</i> ²
FWP197	<i>h⁻ tup11::ura4⁺ tup12::ura4⁺ ura4-D18 leu1-32</i>	Janoo <i>et al</i> ³
FWP218	<i>h⁺ snf21⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP219	<i>h⁺ snf22⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP220	<i>h⁺ snf5⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP221	<i>h⁺ sfh1⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP222	<i>h⁺ snf59⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP223	<i>h⁺ snf30⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP224	<i>h⁺ rsc7⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP225	<i>h⁺ rsc1⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP226	<i>h⁺ arp42⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP227	<i>h⁺ arp9⁺-TAP::KANMX6 ura4-D18 leu1-32 ade6-m216</i>	This study
FWP287	<i>h⁺</i>	This study
FWP289	<i>h⁺ snf22⁺-TAP::KANMX6</i>	This study
FWP228	<i>h⁺/h⁻ snf21::ura4⁺ ura4-D18 /ura4-D18 leu1-32/leu1-32 ade6m216/ade6-m210</i>	This study
FWP229	<i>h⁺ snf22::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP230	<i>h⁻ snf22::ura4⁺ ura4-D18 leu1-32 ade6-m210</i>	This study
FWP231	<i>h⁺ snf5::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP232	<i>h⁻ snf5::ura4⁺ ura4-D18 leu1-32 ade6-m210</i>	This study
FWP233	<i>h⁺/h⁻ sfh1::ura4⁺ ura4-D18 /ura4-D18 leu1-32/leu1-32 ade6-m216/ade6-m210</i>	This study
FWP234	<i>h⁺ rsc1::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP235	<i>h⁺/h⁻ rsc7::ura4⁺ ura4-D18 /ura4-D18 leu1-32/leu1-32 ade6-m216/ade6-m210</i>	This study
FWP236	<i>h⁺/h⁻ ssr1::ura4⁺ ura4-D18 /ura4-D18 leu1-32/leu1-32 ade6-m216/ade6-m210</i>	This study
FWP237	<i>h⁺/h⁻ ssr2::ura4⁺ ura4-D18 /ura4-D18 leu1-32/leu1-32 ade6-m216/ade6-m210</i>	This study
FWP238	<i>h⁺/h⁻ ssr3::ura4⁺ ura4-D18 /ura4-D18 leu1-32/leu1-32 ade6-m216/ade6-m210</i>	This study
FWP239	<i>h⁺ arp42::ura4⁺ ura4-D18 leu1-32 ade6-m210</i>	This study
FWP240	<i>h⁺ arp9::ura4⁺ ura4-D18 leu1-32 ade6-m210</i>	This study
FWP241	<i>h⁺/h⁻ ssr4::ura4⁺ ura4-D18 /ura4-D18 leu1-32/leu1-32 ade6-m216/ade6-m210</i>	This study
FWP242	<i>h⁺ rsc4::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP243	<i>h⁺/h⁻ rsc9::ura4⁺ ura4-D18 /ura4-D18 leu1-32/leu1-32 ade6-m216/ade6-m210</i>	This study
FWP244	<i>h⁺ rsc58::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP245	<i>h⁺ sol1::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP246	<i>h⁺ tfg3::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP247	<i>h⁺ snf59::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP248	<i>h⁺ snf30::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP249	<i>h⁺ arp42::ura4⁺ arp9::ura4⁺ ura4-D18 leu1-32 ade6-m210</i>	This study
FWP250	<i>h⁺ rsc1::ura4⁺ rsc4::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study
FWP251	<i>h⁺ snf21⁺-TAP::KANMX6 arp42::ura4⁺ arp9::ura4⁺ ura4-D18 leu1-32 ade6-m210</i>	This study
FWP252	<i>h⁺ snf22⁺-TAP::KANMX6 arp42::ura4⁺ arp9::ura4⁺ ura4-D18 leu1-32 ade6-m216</i>	This study

Supplementary Table 6. Oligonucleotide primers used in this study.

Gene	Name	Sequence	Description
<i>snf21⁺</i>	FO3577	TGTTTGTTTTTATTTCAATAATAACGACCTGGAAATCGTCTA TTTATGTGGTAATTTGGATTTTTGATCACAAATCGCCAGGGT TTTCCCAGTCACGAC	Deletion-For
	FO3578	AACAAAAAAAAAGACAATCTGTGTTTTTATGAGACTTCAGCAC TTGCATCTATAGTAGTTCAAAAAAGCAATTAATAGCGGATA ACAATTTACACAGGA	Deletion-Rev
	FO3569	AGGATGGCACATTAGCAACGCTTCGCGGAATGGAGGCGGA GGCTACATCGCAATTGGAAGACAGAATTGAAAATGAGGCTC GGATCCCCGGGTAAATTA	TAP-For
	FO3570	AAAAACAAAAAAAAAGACAATCTGTGTTTTTATGAGACTTCA GCACTTGCATCTATAGTAGTTCAAAAAAGCAATTAATGAAT TCGAGCTCGTTAAAC	TAP-Rev
<i>snf22⁺</i>	FO3579	TCCATTTGTACGTTGAGGGTTTTTCGGTCATTGTTGTTGTAAT ATTGATTTGCGATTTACTAAAACACATTTCTCCGCGCCAGGG TTTTCCCAGTCACGAC	Deletion-For
	FO3580	CCTCACCTAACAAAATGTACCAAAATATTATAAACAAGGCAT TAAATAAAACGACAAAAGGTAAAACGTTAGTCCAGCGGATA ACAATTTACACAGGA	Deletion-Rev
	FO3571	TACCGTTGGATTCTGGTATAGTAAGCGCCGAAGATGACAAA GTTATTACTTATGAAGATTCTTCTTCTTATTTCGGAGCGG ATCCCCGGGTAAATTA	TAP-For
	FO3572	TAGTCCTCACCTAACAAAATGTACCAAAATATTATAAACAAG GCATTAAAATAAAACGACAAAAGGTAAAACGTTAGTCCGAAT TCGAGCTCGTTAAAC	TAP-Rev
<i>snf5⁺</i>	FO3581	AAGAATGACACCGATATATGATATTCTTTCTAATAACATTTGC ACTTAAACGAATCTTTACTTTTTTTTTAAATCTCGCCAGGGTT TTCCCAGTCACGAC	Deletion-For
	FO3582	CTGGTCCGTCGAGCTATCCATAACGTGAGTCAAATCTTGATT ATGGTCTTTGATAAAGATTCTTCCAAAATTGTATAGCGGATA ACAATTTACACAGGA	Deletion-Rev
	FO3573	ATTTCCATATCCACTTGTTGTCAAGCTCGCATCAACAAAAA AGGAGGTGAGAATGAATACGGTTTTGGATAGAAATACTCGG ATCCCCGGGTAAATTA	TAP-For
	FO3574	GAAACTGGTCCGTCGAGCTATCCATAACGTGAGTCAAATCT TGATTATGGTCTTTGATAAAGATTCTTCCAAAATTGTATGAAT TCGAGCTCGTTAAAC	TAP-Rev
<i>sfh1⁺</i>	FO3583	CCAAGTAAATTGACTTGCGGCTTACATAGTTTTTTTGGC ATACTTGTGCAAACACGACTTACAAATTTACACGCCAGG GTTTTCCCAGTCACGAC	Deletion-For
	FO3584	TTCTCATTAAATTCGTCTCCTTAAAACACGACTAATGAACAT CAAATACATCAGTCCATTAGAGTAAAACATCTTAGCGGATAA CAATTTACACAGGA	Deletion-Rev
	FO3575	ACAAATCATTGTGCAATGCTTGTGGTGTTCATATGCTAAAA CAGGACAATTGCCATATTGGAGAAAATCTCTATACACTCGGA TCCCCGGGTAAATTA	TAP-For
	FO3576	GACCTTCTCCATTAAATTCGTCTCCTTAAAACACGACTAATG AACATCAAATACATCAGTCCATTAGAGTAAAACATCTTGAAT TCGAGCTCGTTAAAC	TAP-Rev
<i>rsc1⁺</i>	FO4016	TGATATACCCGTTAAAAATCTTACGAACAAGGTACGGGTAC GTCTTCCACGAAAACACCAAGAAAATCCATCAGTGAGACGC CAGGGTTTTCCCAGTCA	Deletion-For

	FO4017	ATATACATTATTAGGTTCTAAAAATCTTCAAATCTATCACAAT ATTTGTTTTCTAAAAATCACAGTGAGTAATTTTCACTAGCGG ATAACAATTTTCACAC	Deletion-Rev
	FO3575	ACAAATCATTGTGCAATGCTTGTGGTGTTCATATGCTAAAA CAGGACAATTGCCATATTGGAGAAAATCTCTATACACTCGGA TCCCCGGGTTAATTAA	TAP-For
	FO3576	GACCTTCTCCATTAATTCGTCTCCTTAAAACACGACTAATG AACATCAAATACATCAGTCCATTAGAGTAAAACATCTTGAAT TCGAGCTCGTTTAAAC	TAP-Rev
<i>rsc7⁺</i>	FO4018	AAAAAGCAATTTTTAACACAGTGCTATTCAGTACGATAAGAT TGAAGTGTATGGAATAAACACCTTGCAATTAAGTACGCC AGGGTTTTCCCAGTCA	Deletion-For
	FO4019	TGATAAATGACTGTATACAATCTGAAAATTAATATTATAAAAC ATCCAAAACAAAATAGGATTAAGAAAACGAGCACTAGCG GATAACAATTTTCACAC	Deletion-Rev
	FO3575	ACAAATCATTGTGCAATGCTTGTGGTGTTCATATGCTAAAA CAGGACAATTGCCATATTGGAGAAAATCTCTATACACTCGGA TCCCCGGGTTAATTAA	TAP-For
	FO3576	GACCTTCTCCATTAATTCGTCTCCTTAAAACACGACTAATG AACATCAAATACATCAGTCCATTAGAGTAAAACATCTTGAAT TCGAGCTCGTTTAAAC	TAP-Rev
<i>ssr1⁺</i>	FO4020	TGCATTTTATTGACGTGAAGCTTTTAAAGCCCTTGGCGTTGA GGATTTTGAAATATTATACTAGACTCTAACACCACAACGCC AGGGTTTTCCCAGTCA	Deletion-For
	FO4021	ATTAAGATAAGATAAAAAAAAAAATTCATTTAAATAAAAGCCTA ATAAATTTTTATCCGATAAGCATATTAGTTGAGATTAAGCGG ATAACAATTTTCACAC	Deletion-Rev
<i>ssr2⁺</i>	FO4022	TTACAACTTAAAAACAACCTAATTTTGTGGGTATTACATTTTT TTCGTGAGTGGACTATTCAAAAATCAGACCTTCACGACGCC AGGGTTTTCCCAGTCA	Deletion-For
	FO4023	TACATATATATATGTCTATAAAAAGTAGATAGCCAAAAGAACT TCATGCAGAATACAAATTTGAATTTGGTGAACCTAACAGCGG ATAACAATTTTCACAC	Deletion-Rev
<i>ssr3⁺</i>	FO4024	ACGTCCACCAATATCGAAGATTATTGATCGGTTCCGGATTGC CGCATATTTGAGAGGAGCTTGAATAAGGACAGCTTATCCGC CAGGGTTTTCCCAGTCA	Deletion-For
	FO4025	AATTTGTAATAATATACACGTTTGTGACACACGCAGGAATA TACCGTTGCTTTCTACTGGTAACGTAGTAATAGTCTATAGCG GATAACAATTTTCACAC	Deletion-Rev
<i>arp42⁺</i>	FO4031	ATACTAACTATTTACATAGACATGGGTAAGTGCGGTTAATC AACCATAATATGTGTTGAGACGAAGACAATGGCATGAACGC CAGGGTTTTCCCAGTCA	Deletion-For
	FO4026	TGCGAGATTTTCGTAAATTTATTTTATACTCTCTTGGGTTTTGA AGACTTTATTTTTGTGAGCAAATAACACAGGCGGATAGCG GATAACAATTTTCACAC	Deletion-Rev
	FO5236	TTCAACATTTGTGGGTATCTAAGCAAGAGTACGATGAAGTAG GTGTTGACAGAGCTTTGTTTGTGAGAAAAGATGCAAGCGG ATCCCCGGGTTAATTAA	TAP-For
	FO5237	ATGCGAGATTTTCGTAAATTTATTTTATACTCTCTTGGGTTTTG AAGACTTTATTTTTGTGAGCAAATAACACAGGCGGAGAATT CGAGCTCGTTTAAAC	TAP-Rev
<i>arp9⁺</i>	FO4027	AGATTAGTGAGTAGTATGACTTGTATCGTAAACCACAATTCA TCCATAGCACAAACAAAGCAATCGGTTTGGGAAAACGACGC CAGGGTTTTCCCAGTCA	Deletion-For

	FO4028	AAGTACCTTTTCGCACGAGTCAATTGATTTCTTAAAATACCGT GACAGGCAATTAACCTTTTATGTGATGAACGAGAGTAAAGC GGATAACAATTTACACAC	Deletion-Rev
	FO5238	TTAATGAATCTGTATCTTCCCATTATGTTACACTTGAGGAATA TGCCCAACATGGACCCACAGCGATTCATACAAAGCAACGGA TCCCCGGGTTAATTAA	TAP-For
	FO5239	AAGTACCTTTTCGCACGAGTCAATTGATTTCTTAAAATACCGT GACAGGCAATTAACCTTTTATGTGATGAACGAGAGTAAGAAT TCGAGCTCGTTTAAAC	TAP-Rev
<i>ssr4⁺</i>	FO4029	AACAAAGCCAAAACCTTCTGCATCTTGATAGATATGAGTAAGT TGTACCATCCAAAACCTTACATATAATTAATAAATCAACAACGCC AGGGTTTTCCCAGTCA	Deletion-For
	FO4030	TAAAGTAGGACGTACGTACTTTTTCAAATCTTCTTTCTGATTA GCCTCAACATGTTTTTCTATACTACACTTTAAATCTTAGCGG ATAACAATTTACACAC	Deletion-Rev
<i>rsc4⁺</i>	FO4285	CAGTCGCAAGCGGACCTAATTAAGTTGATATACTCTATTCA TTGTACATAAACGTTGCAATATAATAAATTATTTAAAACGCCA GGTTTTCCCAGTCACGAC	Deletion-For
	FO4286	TGTACATATTTCAAACGAATGATACCATTTATTTTAATTCCTT GTATTAATACATTTTGTCTCTATCCAAAAGTGTAGCAGCGG ATAACAATTTACACACAGGA	Deletion-Rev
<i>rsc9⁺</i>	FO4911	CCACTTAGTATTTTCGAAAAGTTGTGAAAGGTTTCGTTATACT ACTTTTGACCACCTACTTACTTTACGACGCGCGCTTCACGC CAGGGTTTTCCCAGTCA	Deletion-For
	FO4912	ATATGAAAAGTGATGTCGAGATGCTTGACCTTAGCAATGAG ATTTAAAAGACATGTTATGACTATCGAGCCATCCAAAATAGC GGATAACAATTTACACAC	Deletion-Rev
<i>rsc58⁺</i>	FO4913	TCTGGGCATATTTTGAGTATTTGAAACGCATTACCGTTATCC GGCCTTTTTTGATAGCAATTCGTAAAACAACTCGGCACGC CAGGGTTTTCCCAGTCA	Deletion-For
	FO4914	TATATATATGTATGTATATACATCTATTAATTAGTTAACTAAA ATCTTTAAGTAACACCATGGAAATAGAGAACGAATTAGCGGA TAACAATTTACACAC	Deletion-Rev
<i>sol1⁺</i>	FO4915	CCAAGCGCCTTAGACGTTAATATATTGAAACGTGTATTTTAA GGTCCCTTACTCTTTGTTTGATTTAGTATGAGGAATCCGCC AGGGTTTTCCCAGTCA	Deletion-For
	FO4916	GACAATATACATACAGAAACGCATCAGATTAAGTAACTAGAAATGAA GACAAAAATATTAACGTTTGGAAAATTATCTACAAATAGCG GATAACAATTTACACAC	Deletion-Rev
<i>taf14⁺</i>	FO4917	GCCCGATATCTTCTCACGACTCGCTAAGCGTCTTCTATCC ATATATTTTTCGTCCTCATATACATTGATTTAAGAATACGCCA GGTTTTCCCAGTCA	Deletion-For
	FO4918	CAGTCTTCAGGAAAAAAGTAGAATGGGGAACTAACGCTTC TCCCTTAAAGTAGGCTGCACAAAACGCAATGCTCTTAAAGC GGATAACAATTTACACAC	Deletion-Rev
<i>snf59⁺</i>	FO4919	AATTAGGTACAAGTCTATAAAATCACGCTTGGTTTTGGAATT GATATTCGTAACATTTTCGTTTGACACCAACTTTTCTTCGCC AGGGTTTTCCCAGTCA	Deletion-For
	FO4920	ACCAATAGCAACTAGCCTAAATCCAATTGATCGAAAATATGA TAGCTCAGCAATGCAATTAATAATTTTAAATATAAAGTAGCG GATAACAATTTACACAC	Deletion-Rev
	FO3968	TCATGAGGACAAGAAATCTAAGGAAAGAAGCAAGGTTAAGT TATTATACAAAATTACGAGGTGTCAACAGATCTGTTTCACGG ATCCCCGGGTTAATTAA	TAP-For

	FO3969	ACCAATAGCAACTAGCCTAAATCCACTTGATCGAAAATATGA TAGCTCAGCAATGCAATTAATAATTTTTAATATAAAAGTGAATT CGAGCTCGTTTTAAAC	TAP-Rev
<i>snf30⁺</i>	FO4921	ACAAGAAGTTACAACAAATTTTTTAAAAAACTGGTTATTTAA GTTATATCTCCTTTTTAAGTTAACTACGTTTGTAATACGCCAG GGTTTTCCCAGTCA	Deletion-For
	FO4922	TTACGTTTAGATAAATGATCAATTA AACATAAAAATTACAGGA CATAGCTTTTCTATATCATAAACTTCAAAAACACCACAGCGG ATAACAATTTACAC	Deletion-Rev
	FO3970	GTTTTTTTGATTACAACAAGCTCAAGTTCAGGCTCGAGCAT TGATGCAAAGATATCAACAGGGGATGGAATTTAACAAACGG ATCCCCGGGTTAATTA	TAP-For
	FO3971	TTACGTTTAGATAAATGATCAATTA AACATAAAAATTACAGGA CATAGCTTTTCTATATCATAAACTTCAAAAACACCACGAATTC GAGCTCGTTTTAAAC	TAP-Rev
<i>fip1⁺</i>	FO6125	GTCGGAGTTGGTGTCCACTT	ChIP
	FO6126	GCCAATGTCCTTTGTCTCGT	ChIP
<i>str3⁺</i>	FO6127	GCTAATCAGTAACCAGGGTTGT	ChIP
	FO6128	CACCGCGGTTAAGAAGCTC	ChIP
<i>frp1⁺ 3'</i>	FO5994	CTTCCAGACTTCGGAGTGAA	ChIP
	FO5995	TTTCCAAAAGCAAACGGAGT	ChIP
<i>frp1⁺</i>	FO5990	TGTACAAAGCCTCCCCTCTG	ChIP
	FO5991	GCCGCTTCGCTATACACTA	ChIP
<i>pex7⁺</i>	FO6270	TCCCACCCTTATCGAAGCTA	ChIP
	FO6271	GCGAAGCTTGTTTCCTCAAC	ChIP
<i>obr1⁺</i>	FO6272	ACCCAAAGTCCCCACTTAG	ChIP
	FO6273	CAAACCGATTTTGTGCTTGA	ChIP
<i>pho1⁺</i>	FO6282	TGGTTCGGGACACATTTACA	ChIP
	FO6283	CACAAACGAAACCAACAACG	ChIP
<i>spo3⁺</i>	FO4013	GTGCGCTTCAATGAGTTAACACTC	ChIP
	FO4014	AGTGTGGGGAAACTTTTTATTAATC	ChIP
<i>hcs1⁺</i>	FO6129	GCGAATTGTTCACTTGTTCCA	ChIP
	FO6130	TCATACGTTACGTTTTCTCG	ChIP
	FO5758	CTCGTGTTGTACTCCCAAT	RT-PCR
	FO5759	CTTCCAAACGACCAATCTTGC	RT-PCR
<i>ght3⁺</i>	FO5113	GAAGAATTCATTGAAAATGCT	RT-PCR
	FO5764	TCTAGAAGTGCCTCGCTATCA	RT-PCR
<i>ght4⁺</i>	FO5115	ACGTTTTATCGAAAATGCAGA	RT-PCR
	FO5765	CCTCATGAAACTAGTCACTTCC	RT-PCR
<i>ght5⁺</i>	FO5117	TCTAGCAATGACATTAGTTCC	RT-PCR
	FO5766	AAATGCAGCGAGTGAGACAAG	RT-PCR

Supplementary references.

1. Cairns, B.R., Erdjument-Bromage, H., Tempst, P., Winston, F. & Kornberg, R.D. Two actin-related proteins are shared functional components of the chromatin-remodeling complexes RSC and SWI/SNF. *Mol Cell* **2**, 639-51 (1998).
2. Pelletier, B., Beaudoin, J., Mukai, Y. & Labbe, S. Fep1, an iron sensor regulating iron transporter gene expression in *Schizosaccharomyces pombe*. *J Biol Chem* **277**, 22950-8 (2002).
3. Janoo, R.T., Neely, L.A., Braun, B.R., Whitehall, S.K. & Hoffman, C.S. Transcriptional regulators of the *Schizosaccharomyces pombe fbp1* gene include two redundant Tup1p-like corepressors and the CCAAT binding factor activation complex. *Genetics* **157**, 1205-15 (2001).
4. Jeanmougin, F., Thompson, J.D., Gouy, M., Higgins, D.G. & Gibson, T.J. Multiple sequence alignment with Clustal X. *Trends Biochem Sci* **23**, 403-5 (1998).