Table 2. Ste11p target genes expressed in both cell types

Common name	Systematic name	Sexual diff. function	Known Ste11p target	Description			
Regulation of meiosis							
mei2	SPAC27D7.03c	Meiotic defect	Yes	RNA-binding protein that regulates meiosis progression			
sme2	SPNCRNA.103	Meiotic defect	Yes	Noncoding RNA that binds to mei2			
ran1;pat1	SPBC19C2.05	Meiotic defect	Not studied	Protein kinase that negatively regulates meiosis			
Regulation of ma	Regulation of mating/pheromone response						
spk1	SPAC31G5.09c	Abnormal mating	Predicted	MAP kinase of the pheromone pathway			
pmp1;dsp1	SPBC1685.01	Not studied	Not studied	Protein phosphatase, may dephosphorylate spk1			
рур2	SPAC19D5.01	Not studied	Not studied	Protein phosphatase, may dephosphorylate spk1			
gap1;src1;sar1	SPBC646.12c	Abnormal mating	Reported not regulated by starvation or pheromone	GTPase-activating protein, negative regulator of mating			
rgs1	SPAC22F3.12c	Abnormal mating	Yes	Regulator of G protein signalling (RGS) family member			
ste6	SPCC1442.01	Abnormal mating	Yes	Guanyl-nucleotide exchange factor, regulates conjugation			
ste7	SPAC23E2.03c	Abnormal mating	Yes	Regulator of mating			
gpa1	SPBC24C6.06	Abnormal mating	Induced by starvation, Ste11p-dependency not tested	G-alpha family			
ste11;steX;aff1	SPBC32C12.02	Abnormal mating	Yes	Transcription factor, contains an HMG box			
map1	SPAC11E3.06	Abnormal mating	Yes	MADS box transcription factor, regulates of mating-type specific genes			
ste4	SPAC1565.04c	Abnormal mating	Yes	Contains SAM and RAS association domains, regulates mating			
Nuclear moveme	nt						
dhc1	SPAC1093.06c	Meiotic defect	Induced by starvation, Ste11p-dependency not tested	Cytoplasmic dynein heavy chain, required for nuclear movement			
Cell fusion							
fus1	SPAC20G4.02c	Abnormal cell fusion	Yes	Required for cell fusion			
	SPAP7G5.03	Not studied	Not studied	Similar to Sa. cerevisiae Prm1p, which is required for mating			
	SPAC31G5.07	Not studied	Not studied	Similar to Sa. cerevisiae Fig1p, which is required for mating			
Nuclear fusion							
tht1	SPAC13C5.03		Not studied	Required for nuclear fusion after mating			

Cell cycle regu	lators			
cig2;cyc17	SPAPB2B4.03	No	Not studied	G₁/S-phase cyclin
rum1	SPBC32F12.09	Abnormal mating	Not studied	cdk inhibitor
cdc10	SPBC336.12c	No	Not studied	Subunit of the MBF transcription factor complex, required for S phase
арс2	SPBP23A10.04	Not studied	Not studied	Component of the Anaphase Promoting Complex (APC)
Others		<u>. </u>	•	
ubp11	SPBC19C2.04c	Not studied	Not studied	Ubiquitin C-terminal hydrolase activity
shk2;pak2	SPAC1F5.09c	Normal mating	Not studied	PAK-related serine/threonine protein kinase
	SPCC162.10	Not studied	Not studied	Serine/threonine protein kinase
ehs1;yam8	SPAC1F5.08c	Abnormal mating	Not studied	Predicted glycoprotein, required for cell wall integrity
itr2	SPAC20G8.03	Abnormal mating	No, induced by glucose starvation	Myo-inositol transporter, essential gene
	SPBP4H10.11c	Not studied	Not studied	Long-chain-fatty-acid-CoA ligase activity (predicted)
срр1	SPAC17G6.04c	Abnormal mating	Not studied	Beta subunit of farnesyltransferase (FTase)
rlc1	SPAC926.03	Not studied	Not studied	Myosin II regulatory light chain, involved in cytokinesis
rad54;rhp54	SPAC15A10.03c	Not studied	Not studied	DEAD/DEAH box helicase involved in DNA repair
	SPBC3F6.01c	Not studied	Not studied	Serine/threonine protein phosphatase
rep1;rec16	SPBC2D10.06	Normal mating	Yes	Transcription factor regulator of pre-meiotic DNA synthesis
	SPBC3D6.13c	Not studied	Not studied	Member of the protein disulfide isomerase oxidoreductase family
	SPAC4G9.14	Not studied	Not studied	Member of the Mpv17/PMP22 family of peroxisomal membrane proteins
etr1	SPAC26F1.04c	Not studied	Not studied	Enoyl-[acyl-carrier protein] reductase (predicted)
тср5	SPBC216.02	Not studied	Not studied	Similarity to nuclear migration protein 1 (Sa. cerevisiae Num1p),
 	SPAC7D4.09c	Not studied	Not studied	Member of the 3-oxo-5-alpha-steroid 4-dehydrogenase family
	SPBC13G1.02	Not studied	Not studied	Mannose-1-phosphate guanyltransferase (predicted)
rad17	SPAC14C4.13	Not studied	Not studied	Involved in DNA repair and replication checkpoint control
	SPBC4.01	Not studied	Not studied	Sequence orphan, unknown function
	SPBC146.02	Not studied	Not studied	Sequence orphan, unknown function
	SPCC338.18	Not studied	Not studied	Sequence orphan, unknown function
	SPAC29B12.11c	Not studied	Not studied	Orthologue of human WW domain binding protein-2, unknown function
	SPAC18G6.01c	Not studied	Not studied	Conserved hypothetical protein, unknown function
	SPAC8C9.16c	Not studied	Not studied	Unknown function, contains TLD domain
	SPAP27G11.14c	Not studied	Not studied	Sequence orphan, unknown function
	SPBC1604.01	Not studied	Not studied	Contains a domain of unknown function (DUF323)
	SPBC359.06	Not studied	Not studied	Contains a Class II aldolase domain and adducin N terminal domain
	SPBC902.06	Not studied	Not studied	Sequence orphan, predicted coiled-coil, localised to spindle pole body

	SPCC1393.07c	Not studied	Not studied	Sequence orphan, unknown function
eta2	SPAC31G5.10	Not studied	Not studied	Contains two Myb DNA-binding domains
	SPAC22A12.06c	Not studied	Not studied	Contains a domain of unknown function (DUF341)
	SPAC2C4.17c	Not studied	Not studied	Possible ion channel
	SPAC32A11.01	Not studied	Not studied	Conserved hypothetical protein, unknown function
pvg2	SPAC27E2.07	Normal mating	Not studied	Involved in cell wall biosynthesis
	SPBC19G7.14c	Not studied	Not studied	Sequence orphan, unknown function
elg1	SPBC947.11c	Not studied	Not studied	Replication factor C complex
omt2	SPAC27D7.04	Abnormal sporulation	Not studied	Required for spore wall formation
	SPBC354.08c	Not studied	Not studied	Contains a domain of unknown function (DUF221)
	SPAC227.06	Not studied	Not studied	Member of the YIP1 family, predicted to be involved in vesicle traffic

Stellp targets were identified as described in the text. Column three indicates whether the genes are known to have a function in sexual differentiation and column four indicates whether their Stellp-dependency has been studied. Mutant phenotypes and descriptions were compiled by using information from GeneDB (www.genedb.org), PombePD (www.proteome.com), and our own literature searches.