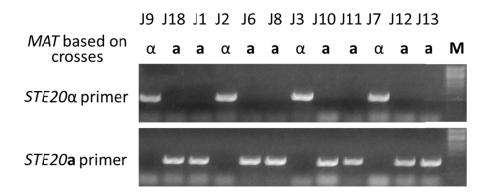
SUPPLEMENTAL MATERIAL

SUPPLEMENTAL TABLE 1 RFLP markers used in comparing the parental strains XL280p and JEC20a.

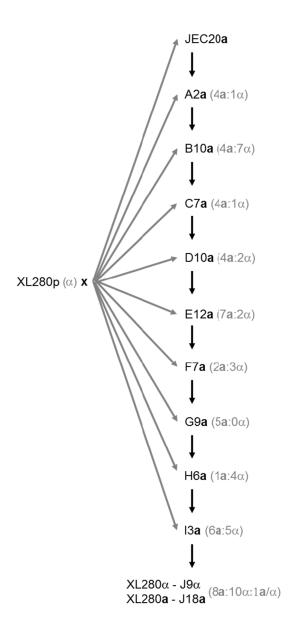
Marker name	Chromosome	Primer sequences (5'-3')	Polymorphic
Hind28	1	GGTGTGCTACATCTCTTGGTTG	No
		TCATCATGTCTCATGCAGCTTAC	
Eco23	2	TATTTAGGTATGGCCGATTGTG	No
		TGCCCAACTCTCTTCCTATTTC	
Pst19	3	AACCTCGTGGAGTCTTTGTCC	Yes
		CTGGATCATGGCTAGATGATTG	
Hind8	3	CTTGATGCTCTTTATGGGGAAG	Yes
		TGTGCCAAGGTTATGGAGATG	
Hind33	4	AGTCACTCTGACACCTCAGTCG	Yes
		CTTACTTGAAGACTCCCGTTCG	
Hind10	4	CGGTATGTCAATGCTCTCAATC	No
		TTTCTCCACCTCTGGAAACAAC	
Hind34	5	AGTCCTCCTCTCCGGGTATTC	Yes
111110.54		AGAGCAATGACCCTGTCCAC	
Hind13	6	GTGATCATGCAGAACTTGGTGT	Yes
11111013	U	CCGAGATGTCGAAGAAGAAGAT	
Hind19	7 or 8	GCCACTCTTCATTCTTCCTCTG	Yes
		TCAACGCCTTCTTCTTCTCC	
Hind25	7 or 11	TCATAGTCTCGGCGTATGTCTC	No
		ATGGGTTGGCTCTGTTTGTC	
Hind11	10	ATACGACATACAAGGAGGGTCTG	Yes
		TGATTGACCTGCACAGAGAAAC	
Hind9	12	GGATTCGGCGTTCTATACAGTC	Yes
		TTGCTGATTCAAGTTGTTGCTC	
Hind15	13	GCTATGTGCCTACTGCTACTGG	No
		CCGACTCTGCTTCTCATACTTG	

The restriction enzymes used were EcoRI (Eco), HindIII (Hind), or PstI (Pst). Polymorphic indicates whether or not there was a polymorphism between strains JEC20a and XL280p.

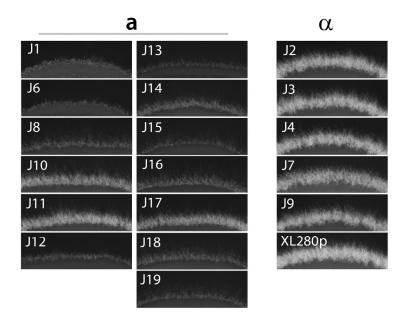
Supplemental Figures



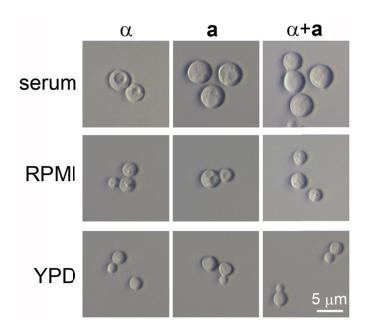
SUPPLEMENTAL FIG 1 The mating type determined by crosses is the same as determined by amplification of the MAT-specific STE20 gene. Progeny from the J generation were randomly selected (strain number indicated above). The mating type, based on crosses with reference strains JEC21 α and JEC20 α , is indicated below their strain number. Gel images of PCR amplicons specific for $STE20\alpha$ and $STE20\alpha$ of these strains are shown below.



SUPPLEMENTAL FIG 2 The pedigree of the congenic pair strains XL280α and XL280a. The progeny with the a mating type from a cross was selected at random and used for the next backcross with XL280p. The series of backcrosses gave rise to the congenic pair strains XL280α and XL280a.



SUPPLEMENTAL FIG 3 Enhanced self-filemantation is associated with the α mating type locus. All progeny generated in the ten backcrosses (generation J) were examined for self-filamentation on V8 juice agar medium at 22°C in the dark. The parental XL280p strain was also cultured under the same condition for comparison. All the α isolates examined produced filaments as robustly as the parental XL280p strain. By comparison, all the α isolates filamented relatively poorly, with strain to strain variations.



SUPPLEMENTAL FIG 4 The congenic strains maintained yeast growth when cultured under *in vitro* conditions that are relevant to host physiology. Strains XL280α, XL280a, and the mixture of equal number of XL280α and XL280a were inoculated into fetal bovine serum, RPMI medium, or YPD medium with the final cell density of 1x10⁵ cells/ml. The cells were incubated at 37°C with 5% CO₂. Photographs were taken after 3 days of incubation. All cells were in the yeast form and no filaments were observed.