Supplemental data



Figure S1. Colony and cellular images of the white phenotype of the WT and *phr2/phr2* **mutant under different pH conditions.** The parent strain GH1013 served as the control. White cells were plated onto Lee's glucose medium plates and incubated at 25°C for five days.



Figure S2. Colony and cellular images of the opaque phenotype of the WT and *phr2/phr2* **mutant under different pH conditions.** The parent strain GH1013 served as the control. Opaque cells were plated onto Lee's glucose medium plates and incubated at 25°C for five days.

Strain name	Parent	Genotype	Reference
	strain		
GH1349	WUM5A	MTLα/α ura3::FRT/ura3::FRT arg4::dpl200/	(1)
		arg4::dpl200-URA3-dpl200	
P37005		Clinical isolate, MTLa/a	(2)
GH1012	CAI4	As CAI4, but <i>MTLa</i> /a	(3)
GH1013	BWP17	As BWP17, but <i>MTL</i> a/a	(3)
WO-1		Clinical isolate, <i>MTLα/α</i>	(3, 4)
(FC4)			
SN250α	SN250	As SN250, but <i>MTLα/α</i>	(5)
12C		Clinical isolate, MTLa/a	(2)
GH1120	CAN52	As CAN52, but MTLa/mtla::dpl200	(3)
GH1352	GH1349	As GH1349, but arg4::dpl200/ arg4::dpl200	This study
cyr1/cyr1	GH1352	As GH1352, but cyr1:arg4/cyr1::URA3	This study
pde2/pde2	WUM5A	As WUM5A, but pde2::dpl200/pde2::dpl200	(6)
WUM5A	WO-1	As WO-1, but ura3::FRT/ura3::FRT	(7)
CAR2	CAI4	As CAI4, rim101::hisG/	(8)
		rim101::hisG-URA3-hisG	
CAR2a	CAR2	As CAR2, but <i>MTLa/ mtl</i> α:: <i>FRT-SAT1-FRT</i>	This study
phr1/phr1-2	GH1013	As GH1013, but mfa1::ARG4/mfa1::HIS1	This study
phr2/phr2-2	GH1013	As GH1013, but mfa1::ARG4/mfa1::HIS1	This study
SZ306u-a	SZ306u	As SZ306u, but <i>MTL</i> a / <i>mtl</i> α:: <i>FRT-SAT1-FRT</i>	(9)
GH1600	SZ306a	As SZ306a, but MFA1/MFA1::MFA1p-GFP	(9)
SZ306u	SZ306	As SZ306, but ura3::FRT/ura3::FRT	(9)
GH1349a	GH1349	As GH1349 but arg4::dpl200/ arg4::dpl200	This study
		-URA3-dpl200::CaARG4	
ste11/ste11	SC5314	As SC5314, but ste11::FRT/ ste11::FRT	(9)
		MTL a /mtla::FRT	
GH1613	JKC131	As JKC131, but <i>MTLa/mtlα::FRT</i>	(9)

Table S1. Strains used in this study

References

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Table S2. Primers used in this study

Name	Sequence (5' to 3')	Purpose	
Marker-F	CCGCTGCTAGGCGCGCCGTGACCAGTGTGATG GATATCTGC	Amplification for ARG4, HIS1, LEU2,	
Marker-R	GCAGGGATGCGGCCGCTGACAGCTCGGATCCA CTAGTAACG	URA3 fragments used in fusion PCR	
PHR1-L FWD	CATAACTCTAACGATTCCA		
PHR1-L REV	GTCAGCGGCCGCATCCCTGC-TATCTAACTGGAG TGTATAAG	Fusion PCR for	
PHR1-R FWD	CACGGCGCGCCTAGCAGCGG-GTTCTGTTGTATA GGAC	PHR1 KO	
PHR1-R REV	TACTTGGGATAAGGAGAG		
PHR1-CHF	TCGGCTAAGTATTGTAATGGC	<i>PHR1</i> KO check (first copy)	
PHR1-CHR	GGACAATGTTCGTGATGATGA		
PHR1-orf-F	TACATTGTTGCTGATTTATC	PHR1 KO check	
PHR1-orf-R	GCAATCGATATAACCAC	(second copy)	
PHR2-L FWD	AGTCAATGAGATGGGAAG		
PHR2-L REV	GTCAGCGGCCGCATCCCTGC-TACAATTCTGCTA CTGTTG	Fusion PCR for	
PHR2-R FWD	CACGGCGCGCCTAGCAGCGG-	PHR2 KO	
	AGTATTATGCCACCACTAGT		
PHR2-CHF		PHR2 KO check	
PHR2-CHR	CCTTGGTGGAAATCCTTGAT	(first copy)	
PHR2-orf-F	AGATACACTTCTGTTGTCG	PHR2 KO check	
PHR2-orf-R	TACTACCACTTGAACCAGA	(second copy)	
RAS1-L FWD	CAGAATTGCATTTCTCCGATC		
RAS1-L REV	CACGGCGCGCCTAGCAGCGGTGAAGAAGGAAA AACAGACGAG	Fusion PCR for	
RAS1-R FWD	GTCAGCGGCCGCATCCCTGCACATTCAACATCCT CTTCATACC	RAS1 KO	
RAS1-R REV	ATACTATATCAGCTCAACCTCG		
RAS1-CHF	тсстсстссататттсастттатс	RAS1 KO check (first copy)	
RAS1-CHR	TTCTTCTTCCTCCACAACAAC		

RAS1-orf-F	AAGAATATCTGGCCATGAGAG	RAS1 KO check			
RAS1-orf-R	ACCACCATTAACAGCACTAG	(second copy)			
MFA1-RT-F	ATGGCTGCTCAACAACAATC				
MFA1-RT-R	AACAGAACAAGTGGAACAGC				
MFα1-RT-F	TGACAGTAACCAAGTTGTTG				
MFα1-RT-R	AGCACCAGAGGTAAGAGTAG	Quantitative real time			
STE2-RT-F	TACTGGTTGGTATGATGGATC	PCR			
STE2-RT-R	AAGGCAACAACAATCAATCC				
GFP-RT-F	AGACACAACATTGAAGATGG				
GFP-RT-R	AGCAGCTGTTACAAACTCAAG				
For the construction of CYR1-reconstituted strain					
LWH105	ACAAGCGGAAGAGTATGAATG	CYR1 promoter			
LWH106	CTTCCTAACATTCTCAACTTCATTTAAATTGAGAATGAA TGGT	region amplification			
LWH107	ACCATTCATTCTCAATTTAA <u>ATG</u> AAGTTGAGAATGTTA GGAAG	CYR1 catalyzed			
LWH108	AACGACAACTATATATGTATCTATCATTGTATGAAAAA TTCAA	domain sequence amplification			
LWH109	TTGAATTTTTCATACAATGATAGATACATATATAGTTGT				
LWH110	CACGGCGCGCCTAGCAGCGGATGCAGTTGTCATTGAAG	<i>CYR1</i> terminator amplification <i>CYR1</i> 3-UTR amplification			
LWH111	GTCAGCGGCCGCATCCCTGCATGATGATGGTTATGTTG GA				
LWH112	CGAACTATTTGTGACGCAT				
	For the construction of <i>PHR1</i> -reconstituted strain				
LWH117	TCACTTATACACTCCAGTTAG	<i>PHR1</i> gene amplification			
LWH118	CACGGCGCGCCTAGCAGCGGGAAACTTTGCTGAGTACT TG				
LWH119	GTCAGCGGCCGCATCCCTGCTCATCATCACGAACATTG TCC	PHR1 3-UTR amplification			
LWH120	GGTGTCCCATGCATTAACTA				
For the construction of <i>PHR2</i> -reconstituted strain					
LWH121	ATACCACGACAATAGGAGAG	PHR2 gene			

LWH122	CACGGCGCGCCTAGCAGCGGCACCACTAGTAACATTAG	amplification			
	AGT				
LWH123	GTCAGCGGCCGCATCCCTGCAGAGTTGTGGAGACTAGT				
	TG	PHR2 3-UTR			
LWH124	CTGAGTGAGATGAGGTCATT	amplification			
For the construction of <i>RIM101</i> -reconstituted strain					
LWH129	TAGTAGGTAGTGTGATGTTG	RIM101 gene			
LWH130	CACGGCGCGCCTAGCAGCGGCTGTGGAATTGACGCAGT	amplification			
	ТА				
LWH131	GTCAGCGGCCGCATCCCTGCTATCGCTGACAAACCAAA				
	TG	<i>RIM101</i> 3-UTR			
LWH132	GTTACTATCTTTCGTCGAT	amplification			
LT391	CCGCTGCTAGGCGCGCCGTGCCATCATAAAATGTCGAG	CaSAT1 cassette			
	CGTC	amplification			
LT392	GCAGGGATGCGGCCGCTGACTGCAGGACCACCTTTGAT				
	TG				