# Α

#### spt16-11 histone deletion strain

Plasmid Allele	YPAD 30° 2d	YPAD 34° 2d	YPAD 35° 2d	YPAD 36° 2d	-lys 30° 3d	HU30 30° 3d
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H2B-V101F		0	*			·

#### В

#### WT histone deletion strain Plasmid Allele YPAD 30° 2d YPAD 38° 3d HU150 30° 5d -lys 30° 3d C 30° 2d wт 2. 23 ÷. 1 H2A-N39K 1 5 3 H2A-P81Q ..... **3** 25 H2A-P81G 4 142 $\mathbf{a}$ H2B-L83F . 10 žà, H2B-L83S ŵ 2 H2B-A84T A ... 3 H2B-Y86C -1. -H2B-Y86H ģi. 3 1 H2B-N87K 2 \* H2B-S93Y - 32 H2B-V101F

#### С

*pob3-Q308K* histone deletion strain; viability test

pob3-Q308K histone deletion strain; phenotypes of viable combinations YPAD 30° 2d YPAD 35° 2d YPAD 37° 2d YPAD 38° 2d



### Figure S1

Effects of *spt6-F249K* suppressors in FACT mutants or a WT strain. Tests were conducted as in Fig 1. A) Strain 9029-3-2 pTF237 (*spt16-11* with histone deletions) was transformed with plasmids carrying suppressors of *spt6-F249K* to test for overlap between the two sets of suppressors. B) Strain 9028-6-1 pTF237 (WT with histone deletions) was tested as above to determine the effects of histone mutations in an otherwise normal strain. C) Strain 9028-1-4 pTF237 (*pob3-Q308K* with histone deletions) was tested with the *spt6-F249K* suppressor plasmids and tested as above. Only a subset of the plasmids was able to support growth of the *pob3-Q308K* strain, with others failing to grow on medium containing FOA, indicating inability to lose the WT histone plasmid pTF237 (left panel).



## Figure S2

Deletion of *HTA2-HTB2* has mild effects on an *spt6-F249K* strain. Strains with the relevant genotypes listed (Table 1) were tested for phenotypes as in Fig 1.



### Figure S3

H2A-N39K has minor effects on other alleles of *spt6*. Mutations affecting the central core (*spt6-1004* and *spt6-14*) or C-terminal domain (*spt6-50*) of Spt6 do not show the strong suppression of phenotypes by *hta1-N39K hta2-N39K* that was observed with *spt6-F249K*. Each allele caused a strong Spt<sup>-</sup> phenotype detected as His<sup>+</sup> and Lys<sup>+</sup> phenotypes with these strains that carry both *his4-912∂* and *lys2-128∂* reporters (Table 1). Neither this nor other phenotypes associated with these alleles were strongly suppressed by H2A-N39K.