

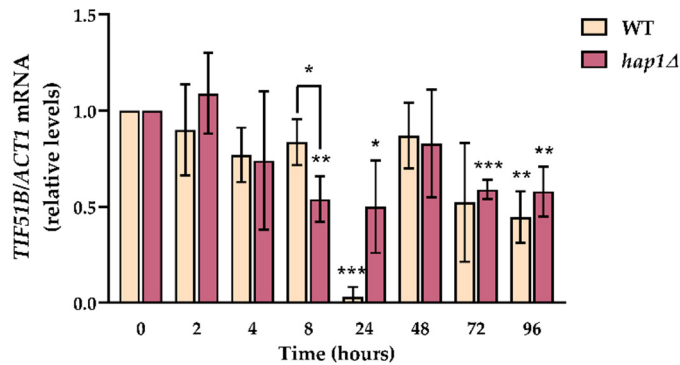
Table S1. Yeast strains used in this study

Name	Genotype	Source
BY4741	MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0</i>	Euroscarf
<i>hap1Δ</i>	BY4741 MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0 YLR256w::kanMX6</i>	This study
<i>hap4Δ</i>	BY4741 MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0 YKL109w::kanMX4</i>	Euroscarf
<i>lia1Δ</i>	BY4741 MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0 YJR070c::kanMX4</i>	Euroscarf
<i>mpc1Δ</i>	BY4741 MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0 YGL080w::kanMX4</i>	Euroscarf
<i>msn2Δ</i>	BY4741 MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0 YMR037c::kanMX4</i>	Euroscarf
<i>pda1Δ</i>	BY4741 MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0 YER178w::kanMX4</i>	Euroscarf
<i>snf1Δ</i>	BY4741 MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0 YDR073w::kanMX4</i>	Euroscarf
<i>tif51A-1</i>	BY4741 MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0 tif51A-1::kanR</i>	(Li <i>et al.</i> 2011)
<i>tif51A-3</i>	BY4741 MATa <i>ura3Δ0 leu2Δ0 his3Δ1 met15Δ0 tif51A-3::kanR</i>	(Li <i>et al.</i> 2011)

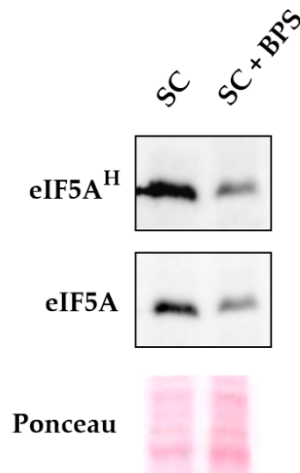
Table S2. Oligonucleotides used in this study

Primer	Sequence (5'-3')
Gene expression detection by RT-qPCR	
ACT-F	TCGTTCCAATTTACGCTGGTT
ACT-R	CGGCCAAATCGATTCTCAA
CYC1-F	AGATGTCTACAATGCCACACC
CYC1-R	CCCTTCAGCTTGACCAGAGT
eIF2A-F	ATTCTACTCCGGCCCCATCT
eIF2A-R	TCTAGTTTGTCACCGACGGC
HAP1-F	TTGGACCTTCCTCACGAATC
HAP1-R	CAGGATCTTCACTGCCCATT
SDH1-F	CTCCAAGTTGACTTTGCTCAGAA
SDH1-R	ACGCGGAACCGTTTACAGA
TIF51-1	TCGACAATCTTACATGGTCT
TIF51A-1	CGATTCTACTTCTGTAGCCA
TIF51B-1	CTACACTTTAGTTCCCTTAC
Gene disruption by PCR	
HAP1-F1	GAAATAGAAGAAAAAGAAAAAAAAAAAAAGGGAACAATAGGTTAGCGGATCCCCGGGTTAATTAA
HAP1-R1	TTACATTATCAATCCTTGCGTTTCAGCTTCCACTAATTTAGATGAGAATTCGAGCTCGTTTAAAC

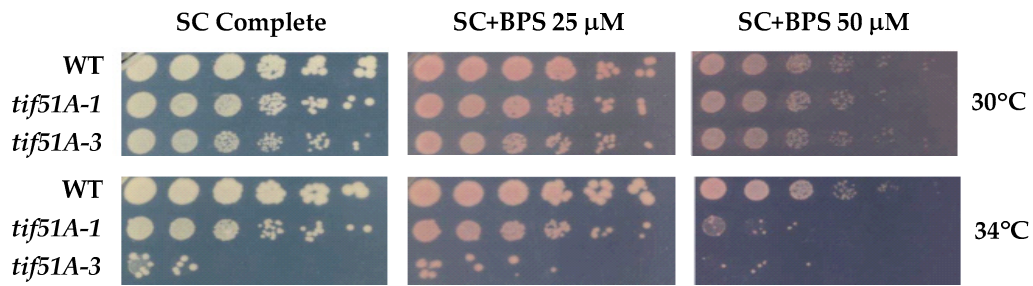
Supplementary Figures



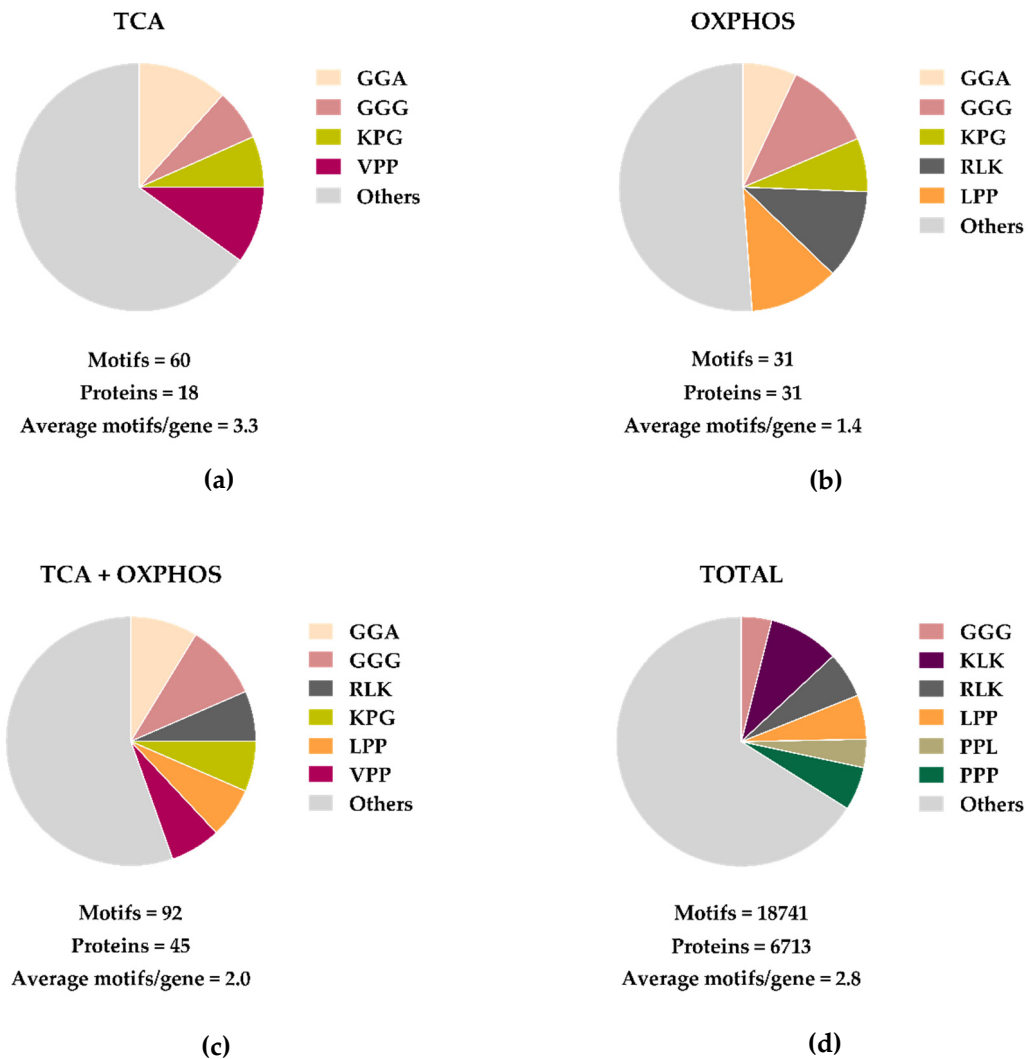
Supplementary Figure S1. Metabolic switch from fermentation to mitochondrial respiration yields a rapid drop of eIF5A but later recovery. WT and *hap1Δ* cells were grown in YPD medium during 96 hours and samples were collected at indicated time points. Relative *TIF51B* mRNA levels were determined. Results are shown as means±S.D. from three independent experiments and expressed relative to the value at time 0. Statistical significance was measured by Student's t-test relative to time 0. *p < 0.05, **p < 0.01, ***p < 0.001.



Supplementary Figure S2. Western blotting of eIF5A and hypusinated eIF5A in WT cells cultured in SC Complete medium with or without the addition of BPS (100 μM) for six hours. Red ponceau was used as loading control.



Supplementary Figure S3. Growth of the WT, *tif51A-1* and *tif51A-3* strains was tested in SC Complete medium with or without the addition of BPS at 25 μM or 50 μM at the indicated temperatures.



Supplementary Figure S4. Distribution of eIF5A-dependent motifs in the proteins of TCA and OXPHOS. Distribution of the 43-highest score eIF5A-dependent ribosome pausing motifs [2] in the proteins of TCA (a), OXPHOS (b), TCA + OXPHOS (c) and total proteins of yeast genome (d).