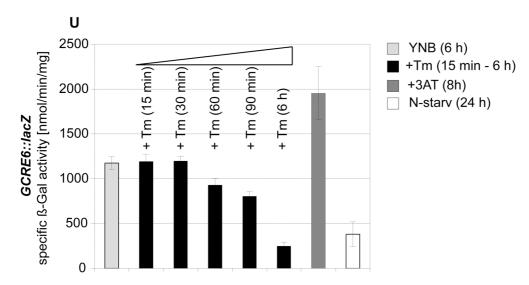
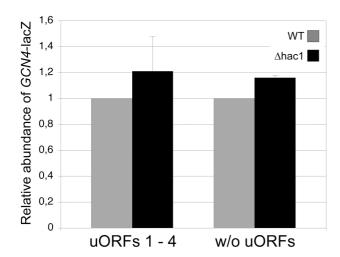
## **Supplementary Figure 1**



## Supplementary Figure 1. Kinetics of *GCRE::lacZ* repression after tunicamycin-mediated ER stress.

Expression of the *GCRE6::lacZ* reporter gene was determined in the haploid  $\Sigma1278b$  wild type strain RH3409 under non-starvation (YNB, light gray bar) in comparison to different starvation or stress conditions. Amino acid (3AT, dark gray bar) and nitrogen starvation (N-starv, white bar) were induced as described in Figure 2B. For ER stress conditions cells were incubated either in the presence of 1  $\mu$ g/ml Tm for 6 h (Tm, black bar) or cells were initially grown for 6 h under non-stress conditions before incubating for indicated time points with 1  $\mu$ g/ml Tm (black bars; Tm induction for 15, 30, 60 and 90 min). Specific  $\beta$ -galactosidase were analyzed as described in Figure 2B.

## **Supplementary Figure 2**



Supplementary Figure 2. *GCN4-lacZ* expression is independent of Hac1. Relative *GCN4-lacZ* levels of haploid wild type strain (RH2816) as well haploid *HAC1* deletion strain (RH3351) expressing the *GCN4-lacZ* fusion gene either with (p180) or without (p227) all four uORFs were determined after growth for 6h in YNB. Bars depict means of at least three independent measurements of  $\beta$ -galactosidase activities.

## **Supplementary Table 1**

 $\beta$ -Galactosidase assays of individual 400 bp *FLO11* promoter elements cloned into the UAS of a *CYC1::lacZ* reporter construct.

	Wild type ( <i>Mata/α</i> ) Units (nmol/min/mg)		∆hac1/∆hac1 Units (nmol/min/mg)		∆gcn4/∆gcn4 Units (nmol/min/mg)	
	YNB	+3AT	YNB	+3AT	YNB	+3AT
FLO11	27	227	27	75	12	28
No insert	19	39	22	50	8	17
2/1	36	61	40	82	15	30
3/2	107	121	123	138	33	69
4/3	61	78	46	106	12	25
5/4	36	60	35	64	8	23
6/5	674	1530	336	768	120	176
7/6	806	920	323	254	108	158
8/7	58	69	34	50	9	15
9/8	70	92	60	121	21	44
10/9	100	333	69	211	27	72
11/10	287	255	822	537	172	463
12/11	92	99	95	94	23	55
13/12	33	64	31	59	11	28
14/13	33	64	29	81	12	30
15/14	28	68	35	101	11	34

The diploid wild type strain (RH2656) as well as diploid  $\Delta hac1/\Delta hac1$  (RH3412) and  $\Delta gcn4/\Delta gcn4$  (RH2658) mutant strains were transformed with indicated *CYC1::lacZ* reporter constructs (53) and expression was assayed under non-starvation conditions (YNB) compared to amino acid starvation induced by adding 10 mM 3AT (3AT). The received  $\beta$ -galactosidase units (nmol/min/mg) are given for each sequence element in each strain and represent mean values of at least three independent measurements.