Table S3	. Description of TFs identified by DREM
TFs	Description
ABF1	DNA binding protein with possible chromatin-reorganizing activity; involved in transcriptional activation, gene silencing, and DNA replication and repair
	Copper-sensing transcription factor; involved in regulation of genes required for high affinity copper transport;
MAC1	required for regulation of yeast copper genes in response to DNA-damaging agents; undergoes changes in
	redox state in response to changing levels of copper or MMS
RPN4	Transcription factor that stimulates expression of proteasome genes; Rpn4p levels are in turn regulated by the 26S proteasome in a negative feedback control mechanism; RPN4 is transcriptionally regulated by various
	stress responses; relative distribution to the nucleus increases upon DNA replication stress
GCN4	bZIP transcriptional activator of amino acid biosynthetic genes; activator responds to amino acid starvation;
UCIN	expression is tightly regulated at both the transcriptional and translational levels
YAP7	Putative basic leucine zipper (bZIP) transcription factor
SKN7	Nuclear response regulator and transcription factor; physically interacts with the Tup1-Cyc8 complex and recruits Tup1p to its targets; part of a branched two-component signaling system; required for optimal induction of heat-shock genes in response to oxidative stress; involved in osmoregulation; relocalizes to the cytosol in response to hypoxia
	Regulator of ribosomal protein (RP) transcription; has forkhead associated domain that binds phosphorylated
FHL1	proteins; recruits coactivator Ifh1p or corepressor Crf1p to RP gene promoters; also has forkhead DNA-binding domain though in vitro DNA binding assays give inconsistent results;
MSN2	Stress-responsive transcriptional activator; activated in stochastic pulses of nuclear localization in response to
	various stress conditions; binds DNA at stress response elements of responsive genes; relative distribution to
	nucleus increases upon DNA replication stress
MSN4	Stress-responsive transcriptional activator; activated in stochastic pulses of nuclear localization in response to
	various stress conditions; blinds DIVA at stress response elements of responsive genes, inducing gene
	expression, involved in diauxic sinit Trimeric heat shock transcription factor: activates multiple genes in response to highly diverse stresses:
HSF1	recognizes variable heat shock elements (HSEs) consisting of inverted NGAAN repeats: monitors translational
	status of cell through an ROC (Ribosomal Quality Control)-mediated translation-stress signal: involved in
	diauxic shift; posttranslationally regulated
NDC1	Transcriptional repressor; recruits the Cyc8p-Tup1p complex to promoters; mediates glucose repression and
NKGI	negatively regulates a variety of processes including filamentous growth and alkaline pH response; activated in
	stochastic pulses of nuclear localization in response to low glucose
EK II 1	Forkhead family transcription factor; minor role in expression of G2/M phase genes; negatively regulates transcription elongation; positive role in chromatin silencing at HML. HMR: facilitates clustering and
	activation of early-firing replication origins; binds to recombination enhancer near HML, regulates donor
	preference during mating-type switching; relocalizes to cytosol in response to hypoxia
ACE2	Transcription factor required for septum destruction after cytokinesis: phosphorylation by Chk1n blocks
	nuclear exit during M/G1 transition, causing localization to daughter cell nuclei, and also increases Ace2p
	activity; phosphorylation by Cdc28p and Pho85p prevents nuclear import during cell cycle phases other than
	cytokinesis; part of RAM network that regulates cellular polarity and morphogenesis
MBP1	Transcription factor; involved in regulation of cell cycle progression from G1 to S phase, forms a complex with
	Swi6p that binds to MluI cell cycle box regulatory element in promoters of DNA synthesis genes
	Transcription cofactor; forms complexes with Swi4p and Mbp1p to regulate transcription at the G1/S
SWI6	transition; involved in meiotic gene expression; also binds Stb1p to regulate transcription at START; cell wall
	stress induces prosphorylation by Mipkip, which regulates Swibp localization; required for the unfolded
	protein response, independently of its known transcriptional coactivators
SID4	co znic cruster transcriptional activator, office regulation of gluconeogenesis: regulated by Softo protein kineses
5124	localized to the nucleus
MCM1	Transcription factor; involved in cell-type-specific transcription and pheromone response; plays a central role
	in the formation of both repressor and activator complexes; relocalizes to the cytosol in response to hypoxia