Chaperonin Abundance Boosts Bacterial Fitness

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Abstract

The ability of chaperonins to buffer mutations that affect the protein folding pathways of suggests that their abundance should be evolutionarily advantageous. Here, we investigate the effect of chaperonin overproduction on cellular fitness in *Escherichia coli*. We demonstrate that chaperonin abundance confers (a) an ability to tolerate higher temperatures, (b) improved cellular fitness and (c) enhanced folding of metabolic enzymes, which is expected to lead to enhanced energy harvesting potential.

Introduction

Chaperonins are found in nearly every organism across all domains of life, and are essential in all cases tested to date, although in some cases non-essential paralogues are found (Kumar, 2017; Lund, 2009). The GroE chaperonin system of *E. coli*, consisting of the 60 kDa GroEL and the 10 kDa GroES proteins assembled into ring complexes of 14 and 7 sub-units, respectively, is encoded by the *groE* operon (Balchin et al., 2016; Bukau and Horwich, 1998). This operon is expressed principally from two promoters, one utilized in the presence of housekeeping sigma factor σ^{70}, and the other, which is strongly induced due to the accumulation of unfolded proteins, in the presence of the alternative sigma factor, σ^{32} (RpoH) (Kumar et al., 2015; Lund, 2001; Schumann, 2017). As σ^{32} levels respond to unfolded protein, this provides a feedback loop to maintain proteostasis (Kim et al., 2013). When cells are shifted to heat shock temperatures between 42 – 46 °C, GroE levels increase by 5 - 10 fold, reaching up to 12% of the entire cellular proteome (Martin et al., 1992). These increased levels interact more extensively with the proteome and are assumed to prevent misfolding or assist refolding of heat-stressed proteins (Houry et al., 1999; Llorca et al., 1998; Martin et al., 1992). Cells that cannot mount an unfolded protein response due to *rpoH* deletion are extremely temperature sensitive, and selection for pseudo-revertants of these strains at elevated temperatures yields up-promoter mutations in the *groE* promoter (Kusukawa and Yura, 1988). GroE is thus important even under normal growth conditions, and indeed GroEL and GroES are respectively the 20th and 21st most abundant proteins in *E. coli* (excluding ribosomal proteins), with sufficient protein being made under non-stressed conditions to produce approximately 2800 complexes of GroEL and 5700 complexes of GroES (Li et al., 2014). Other chaperones that are also abundant include the ribosome bound trigger factor (TF), which is the 19th, and the Hsp70 homologue, DnaK, which is the 27th most abundant. The

high levels of all these chaperones indicates their key roles in cell growth. Although combined loss of TF and DnaK is deleterious to cells, GroEL and GroES are the only chaperones in *E. coli* that are essential under all conditions (Fayet et al., 1989).

GroE assists the folding of 10-15% cellular proteins (Houry et al., 1999), many of which are essential (Kerner et al., 2005). GroE's ability to fold 'folding-compromised' proteins (Fares et al., 2002; Houry et al., 1999; Kerner et al., 2005; Tokuriki and Tawfik, 2009) is consistent with a 'genetic capacitance' function. Many studies with different heterologous proteins have shown that GroE can enhance their folding (Durao et al., 2015; Ishimoto et al., 2014; Tokuriki et al., 2008; Tokuriki and Tawfik, 2009; Wyganowski et al., 2013). In addition, some deleterious mutations are retained in the genome upon overexpression of *groE*, probably due to chaperonin-buffered folding of polypeptides whose folding pathway has been perturbed (Fares et al., 2002; Sabater-Munoz et al., 2015; Van Dyk et al., 1989; Williams and Fares, 2010). However, since GroE is an active ATPase, its overproduction could be deleterious. In this study, we have assessed the effect of GroE overproduction on the growth characteristics and thermal tolerance of *E. coli*, and used proteomics and *in silico* flux balance analysis to determine the likely impact of chaperonin overproduction on the metabolic advantage and consequent fitness of the organism.

Results

Construction of GroE Overproducing Strains.

To investigate the effect of chaperonin overproduction on *E. coli*, we constructed two chaperonin producing strains, GL-H_t and GL-L_t, which produce high and low levels of GroE (Fig. 1A). These strains were derived from strain *E. coli* LG6 (Horwich et al., 1993), in which the P_{groE} promoter is replaced by the P_{lac} promoter, by transforming with pTrc-GSL, which

overexpress *groE* operon upon induction with lactose, or its parental plasmid pTrc99A. SDS-PAGE confirmed significant overproduction of GroEL (Fig. 1B) and GroES (Fig. 1C) in GL-H_t compared to GL-L_t. From Western blotting of the lysates we estimate that GroEL levels are twenty-fold greater in GL-H_t than in GL-L_t (Fig. 1D). The expression levels of GroEL in GL-L_t were lower than the MG1655, where wildtype P_{groE} promoter drives the expression (Fig. 1B-D) (Chapman et al., 2006).

GroEL-GroES Overproducing Strains showed Enhanced Temperature Tolerance.

As GroE is involved in protection against thermal stress, we analyzed the impact of different GroE levels in GL-H_t and GL-L_t on growth at temperatures ranging from 17°C to 48°C (Fig. 2) (Chilukoti et al., 2015). *E. coli* MG1655 and MG1655 hosting pTrc-GSL were included for comparison. As expected, GL-L_t cells exhibited heat and cold sensitive phenotypes and consequently showed poor growth at many temperatures, consistent with previous observations that sufficient levels of GroES-GroEL are required for growth over a wide temperature range (Ferrer et al., 2003). Further, MG1655 showed much better temperature tolerance than GL-L_t, showing the importance of the heat-shock regulation of the P_{groE} promoter. The strains harboring pTrc-GSL tolerated higher temperatures, up to 48 °C, than the vector-only MG1655, where *groE* expression is temperature regulated, suggesting that higher levels of GroEL-GroES enable higher temperature tolerance.

GroEL-GroES Overproducing Strain Exhibited Fitness Advantage in Competition Culture.

Since higher levels of chaperonins led to a growth advantage, we examined whether this translated to a fitness advantage even under low stress conditions, by competing two strains with different GroE levels. Since two strains showed similar growth profiles on the plates and in independent liquid cultures at 30 °C (Fig. S2), we chose this temperature for the

competition culture. To do these experiments, we needed to be able to control the plasmid borne and chromosomal copies of the *groESL* operon independently. Therefore, we constructed two new strains with a P_{BAD} based plasmid expression system, called GL-H_b (high expression) and GL-L_b (low expression) strains. Similar to GL-H_t, GL-H_b showed several fold higher GroE induction levels (Fig. S1A) and temperature resistance (Fig. S1B). The cultures of GL-H_b and GL-L_b were competed for 20 passages (~700 generations) and their relative fitness(s) were estimated (Fig. 3A) as described in Materials and Methods (Macho et al., 2010; Monk et al., 2008; van Opijnen and Camilli, 2013). The high *groE* expressing GL-H_b outcompeted GL-L_b (Fig. 3B), indicating that chaperonin level is an important fitness determinant.

Proteomic Analysis Reveals Preferential Enrichment of Metabolic Enzymes in GroEL-GroES Overproducing Strains.

Overproduction of a chaperonin is likely to enrich the levels of folded proteins in the cells, while unfolded or misfolded proteins tend to remain insoluble and thereby targeted to either the inclusion bodies or marked for degradation (Samuelson, 2011). Given this context, we investigated the proteomes of GL-H_b and GL-L_b cells, to identify what might account for the differences in fitness. Both strains were grown under identical conditions (Fig S2) and their soluble proteome profiles (on 2D PAGE) were compared for relative abundance (Fig. S3 and Table S1). Although many of the identified proteins were known chaperonin clients of classes I – III (Kerner et al., 2005), class IV (Arora et al., 2016) and class V (Chapman et al., 2006), several proteins identified as being differentially expressed were not known clients (Table S1), suggesting that either chaperonin overexpression can indirectly affect the folding of these non-client proteins or that the chaperonin client base is larger than currently understood.

Interestingly, none of the obligate GroEL clients – class III (Kerner et al., 2005) were enriched in GL-L_b strain, suggesting that the lowered chaperonin level is insufficient to fold these proteins. Notably, the outer membrane proteins, OmpC and OmpF, which are involved in metabolite import and are known GroE clients (Kerner et al., 2005), were enriched in the soluble proteome of GL-L_b. Consequently, these proteins were less abundant in the membrane fractions of GroEL-deficient strains (Fig. S5). Further, the enrichment of TF in GroEL-deficient GL-L_b (Fig. 1B and Table S1), suggested compensatory functions for low GroE. Furthermore, enrichment of several metabolic enzymes in GL-H_b strain, suggested a higher rate of metabolism in this strain. To evaluate this hypothesis, we collated publicly available *E. coli* proteomic data from the paxdb database (Wang et al., 2012), screened for proteins that were co-enriched with GroE across different experiments and identified 404 proteins that showed significant correlation, in expression levels, with GroE (Pearson correlation co-efficient > = 0.7, $P < 0.05$). Interestingly, a GO enrichment analysis of this set of proteins revealed that majority of these proteins were involved in metabolism and energy production, including multiple GO terms related to carbohydrate metabolism (Table S3).

Flux Balance Analysis of oxidative phosphorylation in high- and low-GroEL strains.

The metabolic FBA simulations analysis was carried out using 'TransFlux', an in-house tool with a module to incorporate gene expression/ proteomic profiles in the FBA framework. The proteomic profiles (summarized in supplemental materials and Table S1) and observations from *E. coli* gene expression microarray studies, derived from the Many Microbe Microarrays Database (M3D, www.m3d.mssm.edu) (Faith et al., 2008) were utilized to constrain fluxes though respective reactions, while performing two independent FBA simulations, each of which corresponded to the expression/enrichment profiles of the enzymes enriched in GL-L_b

and GL-H_b strains. As expected, higher flux was observed through several pathways of carbon metabolism including glycolysis, gluconeogenesis, citric acid cycle (TCA cycle) and its anaplerotic reactions, and alternate carbon metabolism, in the simulated GL-H_b strain (Table S2). Apparently, these pathways are supported by enhanced import of glucose and glycerol (Supplementary File 2). Further, pathways corresponding to several glucogenic amino acids metabolism and energy generating oxidative phosphorylation were enriched in this strain. However, the pathways leading to the toxic Methylglyoxal synthesis have been enriched in the GL-H_b strain (Table S2). However, pathways leading to the metabolism of membrane lipids, pyruvic acid, pentose sugars, ubiquinone and salvage of nucleotides are enriched in the GL-L_b strain. Overall, FBA simulations indicated that the metabolic enzymes that were enriched in GL-H_b potentially lead to higher metabolic flux in this strain (Table S2 and Supplementary material). Further inadequate respiration was observed in GL-L_b (Table S2). Consistent with this analysis, the GroEL-deficient strain showed reduced growth relative to the high GroEL strain under low oxygen conditions (Fig. S4).

Discussion

Over- or under-production of chaperonins in several organisms has been demonstrated to perturb rates of proteolysis (Martinez-Alonso et al., 2010), influence growth rates, and alter the expression levels of compensatory chaperones like DnaK (Lemos et al., 2007). Here we present a simple model system to study the effects of GroE overproduction (Fig. 1). We demonstrate that the overexpression of GroE chaperonin results in enhanced thermal tolerance (Fig. 2) and competitive advantage (Fig. 3). GroEL is known to be required for growth at low (Ferrer et al., 2003) and high (Guisbert et al., 2004) temperatures. Consistent with this, the GroEL-deficient strains exhibited both cold and heat sensitive

phenotypes (Fig. 2) Proteomic studies (Table S1) followed by flux balance analysis (Supplementary material) suggest that the acquired fitness advantage could be attributed to an enriched set of metabolic enzymes. Chaperonin depletion was observed to induce the enrichment of the compensatory chaperone, TF (Fig. 1B and Table S1), which may act as a *holdase* for the GroE client proteins (Hartl and Hayer-Hartl, 2002). Interestingly, while GroE is more abundant than TF in *E. coli* (Zou et al., 2014), TF is observed to be abundant in mycoplasma which lack *groE* operon (Bang et al., 2000; Lund, 2009; Musatovova et al., 2006; Weiner et al., 2003), suggesting that higher levels of TF might be needed in such bacteria to compensate for the chaperonin deficiency. The TF - GroEL interplay, owing to their overlapping functions and client-base (Avellaneda et al., 2017; Bhandari and Houry, 2015), has been demonstrated both *in vitro* (Kandror et al., 1995) and *in vivo* in *E. coli* (Kandror et al., 1997). Therefore, it seems likely that TF enrichment in GL-L_b is compensating for GroE depletion and that TF may be holding the substrates (Singhal et al., 2015). Further, the enrichment of the outer-membrane proteins OmpC and OmpF in the soluble proteome of GL-L_b suggests that these known GroEL client proteins failed to reach their final cellular destinations and may have remained soluble, possibly in a TF-bound state. The reduced levels of these metabolite transporters in the membranes of GroEL deficient strain (Fig. S5) might be responsible, in part, for the lower metabolite transport and metabolic flux in this strain (Table S2). The upregulation of TF was not observed in MG1655, even though GroE expression levels were lower, which may be because in the wild-type strain, levels of GroE respond directly to levels of unfolded proteins. The higher fitness of the GroES and GroEL over-producing strains under the conditions of our experiments is likely to be associated with fitness costs under other conditions (Fig. 2 and 3), otherwise it would be expected that higher

expression would have evolved. In this regard, it is interesting to note that growth in the absence of aeration was reduced in GroEL-GroES over-producing strains (Fig. S2).

This study shows a direct relation between abundance of chaperonins and competitive fitness, which begs the question as to why evolution has not selected for strains with higher levels of chaperonin producing capability. The predictions from FBA simulations provide some clues that may answer this. Although enhanced glycolysis, TCA cycle and oxidative phosphorylation in the GL-H_b cells increase cellular energy currency, FBA simulations for the GL-H_b strain also predicted an enhanced production of a toxic side product, methylglyoxal (Table S2), a very toxic three-carbon aldehyde that can inhibit *E. coli* growth at millimolar concentrations (Kayser et al., 2005; Weber et al., 2005). Therefore, evolution might have selected a balance in metabolic states between energy production and methylglyoxal toxicity, which would have, in turn, selected for an optimal level of chaperonin expression. The fact that chaperonins are active ATPases provides another possible answer to this question. Overabundance of chaperonins might be linked to ATP depletion and consequent reduced growth (Sabater-Munoz et al., 2015). Thus high levels of chaperonin expression would have been selected against during the course of evolution. These explanations are not exhaustive and the final level of chaperonin expression selected for is likely to result from a balance of optimizing fitness due to a multitude of factors.

Our analysis showed that GroE over-production results in several pleiotropic consequences that can enhance cellular fitness under certain conditions. These observations need to be probed further, using tools to enhance our understanding of the precise role of the chaperone-client interaction in influencing fitness and, ultimately, evolution. A similar

system could be advantageous in studying the effect of chaperonin overproduction in different microbes, especially those with multiple chaperonins (Kumar, 2017).

Materials and Methods

Materials, Plasmids, Bacterial Strains and Growth Conditions. All chemicals were from Sigma, Inc. Bacterial growth media and media supplements were from HiMedia Laboratories, Inc., Mumbai, India. Phusion polymerase for colony PCR was purchased from New England Biolabs Inc, USA. GroE expression plasmids, pBAD-GSL and pTrc-GSL were generated by cloning GroE operon into NcoI and HindIII sites on plasmids pBAD24 (Guzman et al., 1995) and pTrc99A (Amann et al., 1988), respectively. The *groE* conditional mutant strain, *E. coli* LG6, was a kind gift from Arthur Horwich, Yale University, USA (Horwich et al., 1993). This strain produces GroE at levels similar to the wildtype at 30 °C (Chapman et al., 2006). Oligonucleotide primers were purchased from Integrated DNA Technologies, Inc., Coralville, IA, USA.

Construction and validation of strains producing high and low GroE levels. To enable control of GroE levels independently from the growth temperature, two strains that differentially express *groE* were generated from the *E. coli* strain LG6, in which the chromosomal *groE* promoter has been replaced with a P_{lac} promoter (Horwich et al., 1993). A high level GroE expression strain, GL-H_t (for GroEL High pTrc), was obtained by transforming LG6 with pTrc-GSL and a lower level GroE expression strain, GL-L_t (for GroEL Low pTrc) was obtained by transforming with the control plasmid pTrc99A (Amann et al., 1988). The scheme for the generation of these phenotypes is illustrated in Fig. 1A. To confirm the expression levels, these strains were cultured in the presence of 0.2% D-lactose to induce chromosome and plasmid borne *groE* operons, for 3 h at 30 °C. The resulting cells were suspended in lysis buffer

containing 50 mM HEPES:KOH pH 7.5 and 150 mM NaCl, 1 mM EDTA and 1 mM PMSF, mixed with Lysing Matrix E and lysed by homogenization in FastPrep (M. P. Biomedicals, Irvine, CA, USA). Lysates were centrifuged at 13,000 rpm for 20 min to obtain soluble lysates. The soluble lysates were resolved on 12.5% and 20% SDS-PAGE followed by Coomassie Brilliant Blue staining to detect the levels of GroEL and GroES, respectively. In parallel, these lysates were probed with an anti-GroEL monoclonal antibody (1.10B) at 1:100 dilution and the blots were developed by BCIP/NBT-Purple Liquid Substrate System (Sigma Aldrich Inc., St. Louis, MO, USA). In addition to these strains, two strains that enable independent regulation of the chromosome and plasmid borne copies of *groE* operon were generated by transforming LG6 with pBAD-GSL and pBAD24 to result in high and low expression strains, GL-H_b and GL-L_b, respectively. These strains were cultured in the presence of 0.2% lactose plus 0.2% arabinose to obtain the high and low expression levels (Fig. S1A).

Temperature Sensitivity Assessment. To assess the extent to which GroE overproduction enables temperature tolerance, actively growing cultures of GL-H_t and GL-L_t were normalized for OD_{600}, serially diluted, and spotted onto eight LB agar plates supplemented with 0.2% D-lactose. The plates were incubated at 17 °C, 20 °C, 22 °C, 25 °C, 30 °C, 37 °C, 40 °C, 42 °C, 45 °C, 46 °C and 48°C. Wildtype MG1655 harboring pTrc-GSL or empty vector (pTrc99A), respectively, were included as controls.

Competition and estimation of relative fitness. GL-H_b and GL-L_b cells were subjected to competitive serial culturing as described previously (Smith, 2011; Vulic and Kolter, 2001; Zambrano et al., 1993). Briefly, equal number of cells from these two cultures were mixed and grown in fresh LB supplemented with 0.2% L-arabinose and 0.2% D-lactose. This mixed culture was grown to stationary phase at 30 °C, recovered, labelled Passage-1 and used to

generate the second passage (Fig. 2A). Serial sub-culturing was repeated for further 20 passages (~700 generations). At each passage, a fraction of the cultures was serially diluted up to 10^{-7} dilution in LB broth and spread on LB agar plates supplemented with 0.2% D-lactose, which supports the growth of the cells derived from either strain. The resulting colonies at each passage, in the range of 23-28 colonies, were screened using colony PCR to identify whether colonies were derived from either GL-H_b or GL-L_b cells. Colony PCR with the PBADF (5'-CTGTTTCTCCATACCCGTT-3') and PBADR (5'-CTCATCCGCCAAAACAG-3') primers, which bind upstream and downstream of the MCS on the parental vector pBAD24, results in the amplification of 2.1 and 0.3 kb fragments from the pBAD-GSL and pBAD24 vectors, harbored by the GL-H_b and GL-L_b cells, respectively. Relative competitive index (CI), a measure of relative fitness, was calculated for each phenotype as the ratio of the proportion of a particular cell type at the final and initial generations (Macho et al., 2010; Monk et al., 2008; van Opijnen and Camilli, 2013).

Proteomic Analysis. Equal number of cells from exponentially growing cultures (OD_{600} = ~ 0.6) of GL-H_b or GL-L_b strains were harvested, suspended in lysis buffer (50 mM HEPES:KOH pH: 7.5 and 150 mM NaCl, 1 mM EDTA and 1 mM PMSF), lysed by sonication, and the soluble protein fractions were recovered by centrifugation at 12,000 rpm for 20 min. 200 µg protein from the soluble fractions of each lysate were resolved through 2D PAGE following the standard protocols. Briefly, the lysates were resolved on the first dimension through a 7 cm Immobilized pH Gradient (IPG) strip of 3 - 10 pH range, followed by 10% SDS-PAGE on the second dimension. The separated proteins were stained with Coomassie brilliant blue and intensities of the stained protein spots were compared between the two gels using densitometry. Differentially enriched spots between the two lysates were picked and identified by tandem mass-spectrometry in an LTQ Orbitrap Mass Spectrometer (Thermo

Fisher Scientific Inc., Waltham, MA, USA). The differentially enriched proteins were identified using MASCOT (Hirosawa et al., 1993) search against UniProtKB/TrEMBL (UniProt, 2019) and RefSeq (O'Leary et al., 2016) databases. The spot identification was done in collaboration with the Centre for Cellular and Molecular Platforms, Bangalore, India.

Flux Balance Analysis of the GL-H_b and GL-L_b Strains. *E. coli* genome-scale metabolic network iJO1366 (Blais et al., 2013) was used for performing the FBA simulations. The iJO1366 model was first simulated using a standard energy source (equivalent of a glucose-supplemented minimal media) to obtain the steady state fluxes through each of the reactions (Orth et al., 2011). The objective function of this FBA simulation was to maximize the biomass production, while using some 'default constraints' (lower- and upper-bounds of fluxes through each reaction) derived from the literature (Blais et al., 2013). Following this preliminary assessment of the *E. coli* cell's metabolic potential, two independent FBA simulations were performed, each of which corresponded to the enzyme expression/enrichment profiles of the GL-H_b and GL-L_b strains. During each of these simulations the reaction flux values were appropriately constrained, based on the results from the preliminary assessment and the corresponding enzyme expression/enrichment profiles (Supplementary File 2). Incorporating enzyme expression profiles into FBA simulations was performed with our software tool 'TransFlux', developed in-house and housed at http://www.nccs.res.in/TrasFlux/index.jsp. Details of the parameters and the principles applied in FBA are presented in the supplementary methods section.

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Figures and Legends

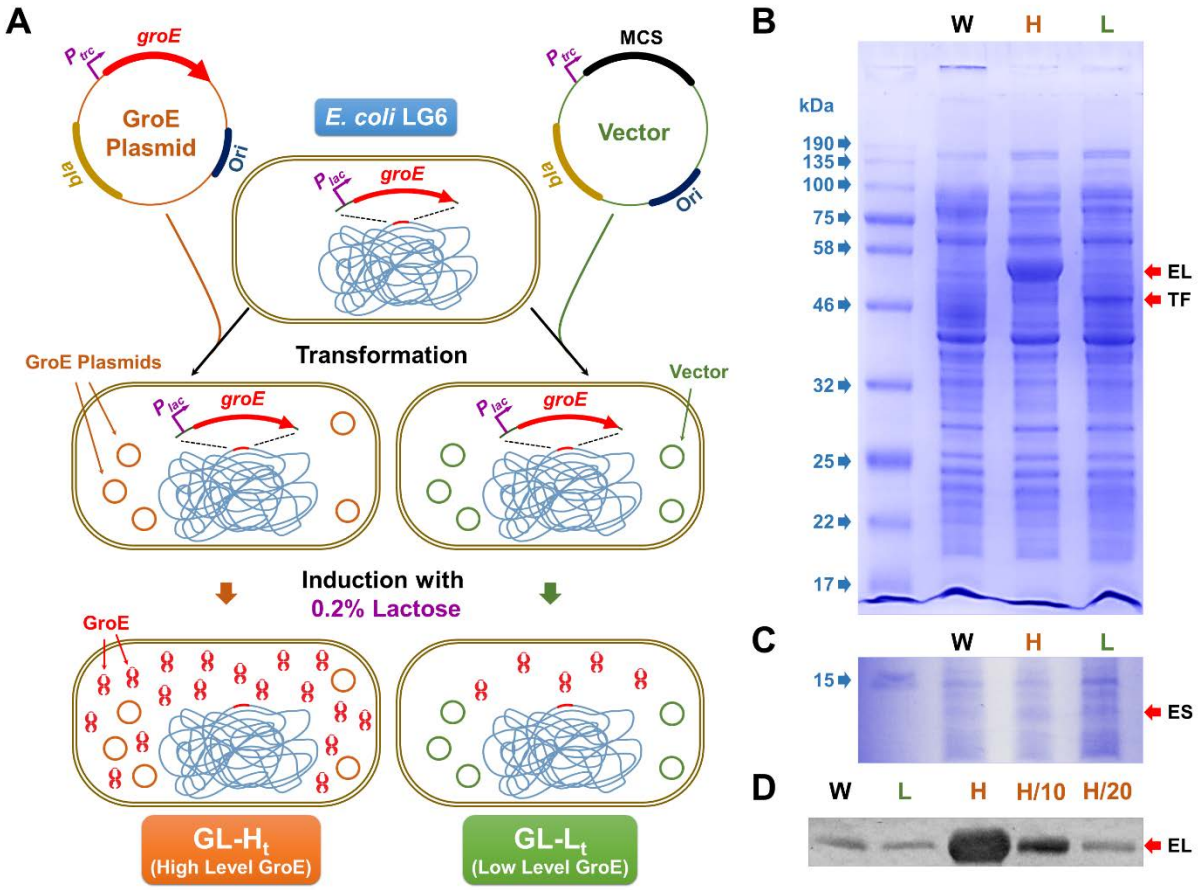

Figure 1. A. Construction of GL-L_t and GL-H_t Strains. In *E. coli* LG6, the *groE* operon is under the control of the inducible P_{trc} promoter. This strain was transformed with either a plasmid expressing *groE* under the control of lactose (GroE Plasmid), pTrc-GSL or its empty vector (Vector), pTrc99a. Upon culturing in the presence of lactose, GL-H_t produces elevated levels of GroEL and GroES, due to the induction of chromosomal and plasmid-borne *groE* operon, while GL-L_t will have lower production of GroES and GroEL due to the induction of only the chromosomal *groE* copy. **B and C. SDS-PAGE Confirms Enhanced Levels of GroE in GL-H_t.** 12.5% (**B**) and 20% (**C**) SDS-PAGE gels comparing the levels of GroEL (EL) and GroES (ES), respectively, in GL-H_t (H), GL-L_t (L) and the wildtype *E. coli* MG1655 (W) strains. Numbers and arrows in blue indicate molecular masses of standards. TF indicates trigger factor. **C. Immunoblotting Estimates Enhancement of GroEL in GL-Ht**. Lysates of MG1655 (W), GL-L_t (L) and GL-L_t (H) were resolved on a 12.5% SDS-PAGE and probed with GroEL specific antibody. H/10 and H/20 indicate the lanes with ten and twenty fold diluted GL-H_t lysates, respectively.

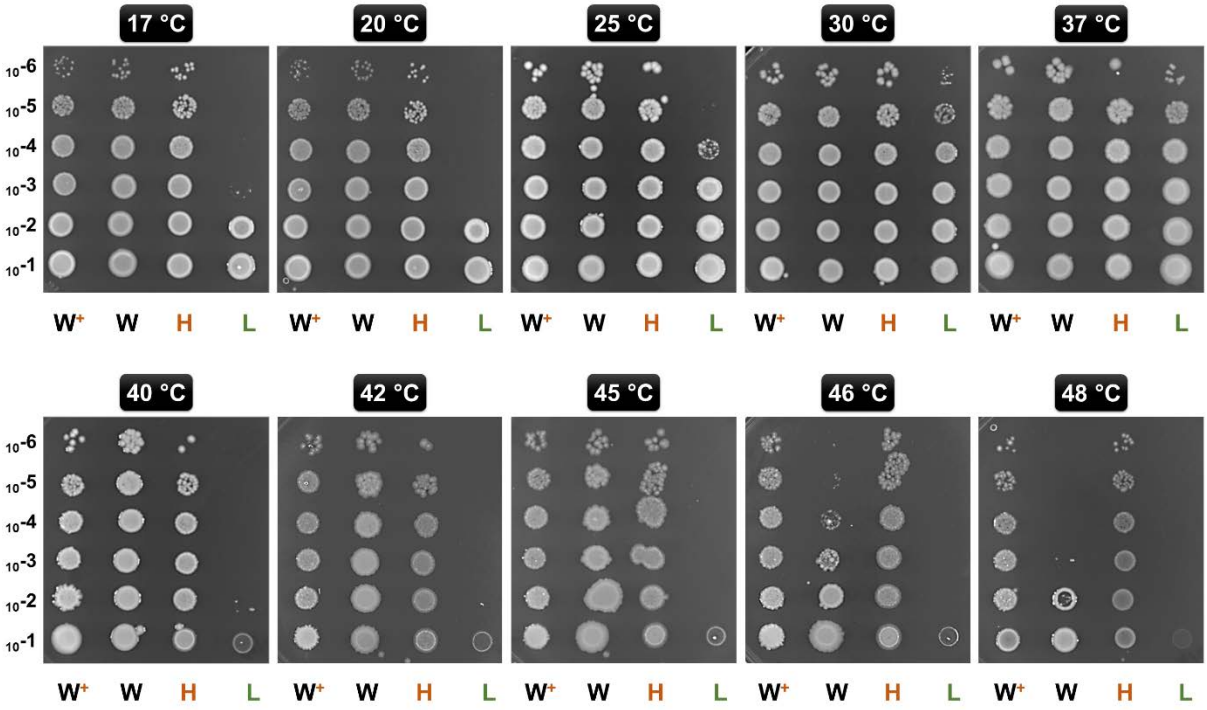

Figure 2. Temperature Tolerance upon Chaperonin Overproduction. Ten-fold serially diluted (10^{-1} through 10^{-6}) exponentially growing cultures of GL-H_t (H), GL-L_t (L), MG16155 (W) and MG1655 with pTrc-GSL (W^+) were spotted onto LB agar plates supplemented with lactose. These plates were incubated at the indicated temperatures.

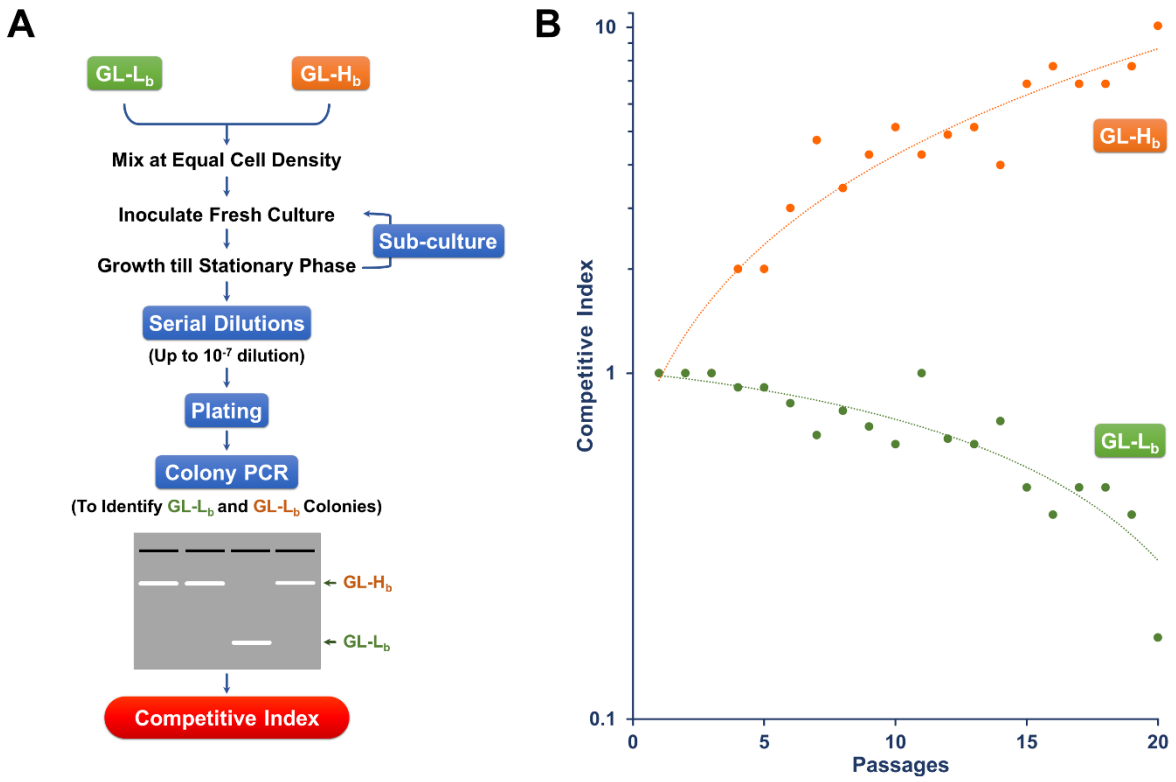

Figure 3. Chaperonin Depletion leads to Lower CI. A. Strategy for Determining the CI for GL-L_b and GL-H_b strains. Stationary phase cultures of GL-L_b and GL-H_b strains were mixed at equal cell density, grown to stationary phase and sub-cultured in fresh media for 20 continuous passages. Cells recovered at each passage were serially diluted as indicated, spread on LB agar plates and the resulting colonies were scored for their phenotype (GL-L_b or GL-H_b), by colony PCR, using vector specific oligonucleotide primers. **B.** CI, as a degree of fitness, was calculated at every passage from a ratio of proportion of the cells with a particular phenotype and plotted as a function of number of passages. CI trend was similar among the three independent experiments.

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