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Detailed Terms
Rational, combinatorial, and genomic approaches for engineering L-tyrosine production in Escherichia coli

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Although microbial metabolic engineering has traditionally relied on rational and knowledge-driven techniques, significant improvements in strain performance can be further obtained through the use of combinatorial approaches exploiting phenotypic diversification and screening. Here, we demonstrate the combined use of global transcriptional machinery engineering and a high-throughput L-tyrosine screen towards improving L-tyrosine production in Escherichia coli. This methodology succeeded in generating three strains from two separate mutagenesis libraries (rpoA and rpoD) exhibiting up to a 114% increase in L-tyrosine titer over a rationally engineered parental strain with an already high capacity for production. Subsequent strain characterization through transcriptional analysis and whole genome sequencing allowed complete phenotype reconstruction from well-defined mutations and point to important roles for both the acid stress resistance pathway and the stringent response of E. coli in imparting this phenotype. As such, this study presents one of the first examples in which cell-wide measurements have helped to elucidate the genetic and biochemical underpinnings of an engineered cellular property, leading to the total restoration of metabolite overproduction from specific chromosomal mutations.

Genetic modifications identified by the rational analysis of pathway stoichiometry, kinetics, and regulation (such as gene deletions/overexpressions and pathway deregulation) have been routinely used in metabolic engineering applications with remarkable success. Despite this, however, many important multigenic phenotypes still remain inaccessible through these methods. Even when pathways are well elucidated, unexpected disconnects between genotypes and predicted phenotypes often hamper the engineering of highly interconnected biological systems. Combinatorial metabolic engineering has emerged in recent years to address this issue using the concept and tools of phenotypic diversification and screening (1). Unlike traditional rational approaches, these methods require very little a priori knowledge of the metabolic network’s properties and interactions and instead depend on the type of diversity generated and the efficiency of the high-throughput assay used. One such technique, global transcription machinery engineering (gTME), has been shown to be particularly effective in introducing phenotypic diversity by reprogramming the cellular transcriptional network (2). In Escherichia coli, mutating two plasmid-encoded components of the RNA polymerase—rpoD, encoding the principal sigma factor σ70—and rpoA, encoding the α subunit—led to improvements in a number of cellular traits, including ethanol tolerance, butanol tolerance, lysine overproduction, and hyaluronic acid production (3, 4). Both polymerase subunits play natural roles in dictating rates of transcription through their interactions with the −10 and −35 regions of the promoter (rpoD) and upstream-promoter (UP) elements, activators, and repressors (rpoA) (Fig. S1), thus making them important knobs for modulating global transcriptional profiles and, by extension, for generating complex phenotypes.

Interest in developing a robust microbial production process for the amino acid L-tyrosine has remained high due to its use in a myriad of pharmaceutical and industrial applications. Apart from its common use as a dietary supplement (5, 6), L-tyrosine also serves as an important precursor for the Parkinson’s disease drug 3, 4-dihydroxy-L-phenylalanine (L-DOPA) (7), as well as melanin-based UV absorbers, cation exchangers, drug carriers, and amorphous semiconductors (8). In addition, both p-coumaric acid and p-hydroxybenzoyl alcohol can be synthesized directly from L-tyrosine and appear frequently as key components in several polymers, adhesives and coatings, pharmaceuticals, biocosmetics, and flavoronoid product precursors (9–11). Work to-date on L-tyrosine and, more broadly, aromatic amino acid (AAA) production in E. coli has focused primarily on rational metabolic engineering strategies to alter regulation and increase pathway flux (12–15). As such, the use of less targeted combinatorial approaches such as gTME presents an as-yet untapped potential for optimizing this important cellular property.

In a recent publication, we described the development of a high-throughput screen for L-tyrosine production based on the accumulation of the dark pigment melanin (16). Here we present the combined application of this assay with gTME for improving L-tyrosine production over a rationally engineered parental strain. We successfully isolated three strains from two plasmid-based mutagenesis libraries (rpoA, rpoD) which exhibited significant increases in both L-tyrosine yields and titers. Unusually, the observed phenotypes were found to be dependent on both the plasmid-encoded rpoA/rpoD gene and unidentified mutations within the bacterial chromosome. Characterization of these isolates using cell-wide assays such as transcriptional analysis and whole genome sequencing allowed us to identify the exact origins for these improvements and, more significantly, reconstruct the phenotype through the introduction of specific chromosomal mutations. Evaluation of this reconstructed strain in large-scale fermentations yielded L-tyrosine titers of up to 13.8 g/l after 36 h at very high volumetric productivities.

Results

Rational Engineering of a Parental Strain. We first sought to construct a high-performing parental strain through the introduction of rational modifications based on known properties of the AAA biosynthetic pathway (Fig. S2). In particular, prior work in our lab (17) helped us identify several strategies for overcoming key kinetic and regulatory limitations in the pathway: (i) deletion of tyrH to circumvent transcriptional regulation, (ii) deletion of pheA to eliminate the loss of AAA pathway intermediates to competing reactions, and (iii) overexpression of feedback resistant derivatives of 3-deoxy-D-arabinoheptulosonate-7-phosphate (DAHP) synthase (aroG<sup>thr</sup>) and chorismate mutase/prephenate dehydrogenase ( tyrA<sup>thr</sup>) to increase flux through the AAA pathway. To avoid potential issues associated with marker and plasmid origin incompatibilities and to eliminate the need for antibiotics and
inducers, the tyrA\textsuperscript{fr}-aroG\textsuperscript{fr} operon was placed under constitutive control and integrated into the bacterial chromosome. Expression levels of these genes were optimized through the use of varied strength promoters (R, Y, W, B, P\textsubscript{t}) from a synthetic constitutive promoter library (Table S1) (18). However, because the L-tyrosine titers of these single integration strains still fell short of their plasmid-based counterparts (Fig. 1A), we also explored the integration of a second copy of tyrA\textsuperscript{fr}-aroG\textsuperscript{fr} under the promoters W, B, and P\textsubscript{t}. From this set of strains, P2, which contains two P\textsubscript{t}-driven copies of tyrA\textsuperscript{fr}-aroG\textsuperscript{fr} and achieves a titer of 418 mg/L L-tyrosine after 24 h, was selected as the starting strain for the construction of our gTME-derived libraries (Fig. 1B).

Isolation and Sequence Analysis of gTME-derived Mutants. In a recent publication, we described a high throughput screen for L-tyrosine production based on the tyrosinase-mediated conversion of L-tyrosine to the dark and diffusible pigment melanin (16). When implemented in an agar plate format, high L-tyrosine overproducers could be identified by the extent of colony pigmentation. We applied this simple visual assay to screen two separate gTME libraries based on the mutagenesis of the RNA polymerase subunits, rpoA and rpoD. Libraries consisting of 10\textsuperscript{5}–10\textsuperscript{6} viable colonies were created by transforming plasmid-encoded rpoA and rpoD mutants into the rationally engineered parental strain P2.

Our search led to the isolation of three strains (rpoA\textsubscript{14}, rpoA\textsubscript{27}, and rpoD\textsubscript{3}) exhibiting 91–113% increases in L-tyrosine production over parental P2 (Fig. 2A). Of note, final L-tyrosine concentrations, which ranged between 798 and 893 mg/L for the three isolates, are reported after 48 h due to the slower growth rates of these strains; however, growth and L-tyrosine production profiles show that levels reach their peaks after just 28 h (Fig. S3).

Sequence analysis of the mutant rpoA/rpoD plasmids revealed interesting mechanistic insights into transcriptional diversification for two of the three strains. Although the wild-type σ\textsuperscript{70} protein typically contains four conserved domains (19), the altered rpoD gene in strain rpoD\textsubscript{3} encoded a truncated form comprising just the latter part of a D521E-substituted region 3 and region 4 (Fig. 2B). We hypothesize that overexpression of this shortened protein could result in anti-σ (such as Rsd) or RNA polymerase holoenzyme titration events which could affect transcriptional preferences. Sequencing of the mutant rpoA plasmid from strain rpoA\textsubscript{14} revealed the presence of two amino acid substitutions (V257F, L281P) within the C-terminal domain of the α subunit (αCTD), in close proximity to residues involved with regulatory factor and UP element interactions (Fig. 2B) (20). The appearance of mutations in key secondary structural elements of the αCTD (the “nonstandard helix” and an α-helix, respectively) (21) suggests that structural deviations may shift the relative alignment of amino acid promoter contacts and, as a result, alter the affinity of the RNA polymerase enzyme for some of its targets. This theory is supported by other rpoA mutations and strains uncovered in our lab (4). Surprisingly, no changes were found in the promoter sequence or coding region of rpoA from strain rpoA\textsubscript{27} (Fig. 2B), which suggests that plasmid-based overexpression of even the wild-type gene can lead to phenotypically advantageous transcriptional modifications.

Requirement for and Interchangeability of Mutant Plasmids and Backgrounds. The recovery of a wild-type rpoA plasmid from rpoA\textsubscript{27}, which by itself has not been shown to be beneficial for L-tyrosine production, introduced the possibility that unidentified chromosomal mutations in the isolated strains may either partially or fully contribute to the observed phenotypes. To explore these scenarios, we constructed and tested a series of strains containing either the wild-type or mutated rpoA/rpoD plasmids in the backgrounds of the parental strain P2 or the plasmid-cured isolates (henceforth referred to as mutant or modified chromosomal backgrounds). We also assessed the individual contributions of the mutant rpoA/rpoD plasmids and mutant backgrounds on cellular phenotype. Unusually, we found that the L-tyrosine production properties of all three gTME strains were uniquely dependent on the presence of both the rpoA/rpoD plasmid and a modified chromosomal background (Fig. 3A), thus highlighting a possible synergism between the plasmid-encoded components.

**Fig. 1.** L-tyrosine production of rationally engineered strains after 24 h. (A) L-tyrosine production of E. coli K-12 ΔpheA ΔtyrR strains containing plasmid-based overexpression of the tyrA\textsuperscript{fr}-aroG\textsuperscript{fr} operon, a single integrated copy, or two chromosomal copies. Plasmid-based copies of tyrA\textsuperscript{fr}-aroG\textsuperscript{fr} under the control of the synthetic constitutive promoters (R, Y, W, B, P\textsubscript{t}) were provided on pZE-kan\textsuperscript{Rn}::tyrA\textsuperscript{fr}aroG\textsuperscript{fr} (Fig. 2A,B). (B) L-tyrosine production of wild-type E. coli K-12, a double deletion mutant (E. coli K-12 ΔpheA ΔtyrR), and parental strain P2.

**Fig. 2.** Properties of mutants isolated from gTME libraries. (A) L-tyrosine titers of isolated mutants after 48 h. (B) Sequence analysis of plasmid-encoded mutant rpoA and rpoD. Numbers above wild-type (WT) rpoD-encoded σ\textsuperscript{70} indicate conserved regions of the protein. NSH stands for the “nonstandard helix” portion of the rpoA-encoded α subunit’s C-terminal domain.

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and other altered proteins residing on the chromosome. Of note, the exception of strain rpoA27, the presence of a wild-type or mutant rpoA/rpoD plasmid did not lead to an enhanced frequency of mutation when compared to the parental strain P2 (Fig. S4). We therefore assume that the high rate of chromosomal variation in our isolates arose out of a requirement for these specific background mutations to increase L-tyrosine production and, as a result, be identified by our screen.

Even more surprisingly, we discovered that transforming the recovered rpoA1/rpoD plasmids into each of the three mutant chromosomal backgrounds led to several unique strains with high L-tyrosine titers (Fig. 3A). One possible explanation for this unexpected result is that the low-level expression of the plasmid-encoded copy of rpoA/rpoD may be sufficient to allow the detection of low-likelihood of having recovered the same chromosomal mutation(s) in all three isolates.

**Transcriptional Analysis Reveals Roles for the Acid Stress and Stringent Responses in E. coli.** To elucidate the biochemical significance of these plasmid and background mutations, we conducted a full transcriptional analysis of the parental strain P2 and the three gTME isolates, rpoD3, rpoA14, and rpoA27. Although several hundred genes were found to be differentially expressed, specific patterns of pathway up- and down-regulation offered notable insights into the underlying changes in these strains.

An analysis of the upregulated genes in the plasmid-bearing strains returned only a handful of loci for rpoD3 and rpoA27 (and none for rpoA14), with a significant number of these presumed hits possessing functions related to acid stress resistance (Table 1). Indeed, several of these genes were found to reside within the same transcriptional network, a complex regulon with at least three separate layers of control exerted by the proteins encoded by gadE, ydeO, and evgA (in ascending hierarchical order) (22).

Although hundreds of downregulated genes were uncovered by our analysis, overlaying the transcriptional data on top of a preassembled network (using Ecocyc’s Pathway Tools Omics Viewer) revealed interesting global reductions in several pathways, including ribosomal protein and RNA formation, fatty acid elongation, de novo purine/pyrimidine biosynthesis, and DNA replication (Table 2). Taken altogether, these transcriptional patterns implicated a stringent response in *E. coli*, a metabolic state triggered by amino acid starvation and other conditions of nutritional stress (23–25). This cellular shift from proliferation and growth to maintenance and stress survival (including, notably, acid stress resistance) is mediated by the small nucleotide (p)pGpp, a regulator whose levels are jointly controlled by the relA-encoded (p)pGpp synthase and the spoT-encoded (p)pGpp synthase/hydrolase.

Because several of these cellular changes were observed in all three gTME mutants, we suspected that the acid resistance pathway and/or the stringent response may in fact play important roles in L-tyrosine overproduction. If this were the case, then altering the expression of known regulators of these pathways may have interesting implications for engineering this particular phenotype. To test this hypothesis, we constitutively overexpressed four regulators—gadE, ydeO, evgA, and a truncated form of relA (26)—to ascertain their effects on L-tyrosine production.

While the overexpression of all four genes in the parental strain P2 typically yielded only minimal increases in titer, a completely different cellular response was observed upon transfer of the evgA and relA plasmids into the modified backgrounds (Fig. 4). Indeed, strains containing one of the mutant chromosomal backgrounds and either evgA or relA exhibited L-tyrosine titers that closely matched those of the original isolates. Testing both weak and strong constitutive expression also revealed gene- and background-specific optima for each combination. Given

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**Table 1. List of upregulated genes related to acid resistance**

<table>
<thead>
<tr>
<th>Gene name</th>
<th>Gene ID</th>
<th>Function</th>
<th>Log fold change</th>
<th>Q value*, %</th>
<th>Log fold change</th>
<th>Q value, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>b3517</td>
<td>gadA</td>
<td>glutamate decarboxylase</td>
<td>2.236</td>
<td>0</td>
<td>1.974</td>
<td>11.05</td>
</tr>
<tr>
<td>b1493</td>
<td>gadB</td>
<td>glutamate decarboxylase</td>
<td>2.699</td>
<td>0</td>
<td>2.515</td>
<td>0</td>
</tr>
<tr>
<td>b1492</td>
<td>gadC</td>
<td>glutamate-γ-aminoxybutyrate antiporter</td>
<td>2.027</td>
<td>0</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>b3512</td>
<td>gadE</td>
<td>transcriptional activator</td>
<td>2.137</td>
<td>0</td>
<td>2.160</td>
<td>0</td>
</tr>
<tr>
<td>b3510</td>
<td>hdeA</td>
<td>acid stress chaperone</td>
<td>1.950</td>
<td>0</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>b3509</td>
<td>hdeB</td>
<td>acid stress chaperone</td>
<td>2.115</td>
<td>0</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>b3511</td>
<td>hdeD</td>
<td>acid resistance membrane protein</td>
<td>1.707</td>
<td>0</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>b3506</td>
<td>slp</td>
<td>starvation lipoprotein</td>
<td>1.609</td>
<td>10.17</td>
<td>1.897</td>
<td>5.67</td>
</tr>
</tbody>
</table>

*Q values represent the lowest false discovery rate at which that gene is called significant. It is like the well-known P value, but adapted to multiple testing situations.
such remarkable phenotypic sensitivity to promoter strength, it seems plausible that additional fine-tuning of evgA and relA expression may yield even greater titers than those observed here.

Whole Genome Sequencing Identifies Beneficial Substitutions in hisH and purF. Although specific cellular pathways were found to be important for L-tyrosine overproduction, the identities of chromosomal alterations necessary for the desired phenotype remained unknown. We therefore conducted whole genome sequencing on parental P2 and the three gTME strains to parse out the identities of these mutation(s). Auspiciously, our analysis revealed only a single base pair (bp) change in each of the three gTME strains—an L82R substitution in the hisH-encoded imidazole glycerol phosphate (IGP) synthase subunit of strain rpoA14, a V5G shift in the purF-encoded amidophosphoribosyltransferase gene of strain rpoD3, and a T → C nucleotide substitution 17 bp upstream of the purF gene in strain rpoA27 (Fig. S5 and Table S2). Furthermore, reintroduction of these specific mutations into a clean parental P2 background resulted in high L-tyrosine titers when strains were also supplied with their respective ppoA or purD plasmids (reconstructed strains were named rpoA14R, rpoA27R, and rpoD3R) (Fig. S6). These results clearly demonstrate the importance of each hisH and purF mutation in establishing an L-tyrosine overproduction phenotype. More significantly, the identification and validation of these chromosomal mutations have led us to the construction of a completely genetically defined strain, rpoA14R, which possesses a titer of 902 mg/l L-tyrosine and a yield of 0.18 g L-tyrosine/g glucose in 50 ml shake flask cultures. To put these numbers into perspective, this yield on glucose is more than 150% greater than a classically improved L-phenylalanine auxotroph (DPD4195) currently used for the industrial production of L-tyrosine (cultivated under similar conditions) (27) and, when excluding glucose diverted to biomass, represents 85% of the maximum theoretical yield.

Performance of rpoA14R in Large-Scale Bioreactors. To evaluate the performance of rpoA14R in larger-scale fermentations, we ran 24 bioreactors using two different minimal medium formulations—MOPS minimal medium (Teknova) (28) and R medium (29). When cells were cultivated in MOPS minimal medium, strain characteristics remained comparable to those previously observed in 50-ml shake flask cultures, with yields on glucose of 0.204 g L-tyrosine/g glucose and a specific growth rate of 0.275 h⁻¹. Unfortunately, however, cell growth was severely hampered (OD₆₀₀ of 10), leading to a titer of just 5.7 g/L L-tyrosine after 56 h (Fig. 5A). Switching to another defined synthetic medium (R medium) led to much higher maximum productivities (2.1 g L-tyrosine/l/h), titers (13.8 g/l after 36 h), and growth rates (0.405 h⁻¹) but with a lower overall yield (0.120 g L-tyrosine/g glucose) (Fig. 5B). Future fermentation optimization should therefore aim at identifying limiting nutrients in MOPS minimal medium to extend the growth and production phase of the culture or, alternatively, modifying the composition of R medium to boost L-tyrosine yields on glucose.

Discussion

In this study, we have demonstrated the clear potential of utilizing a combinatorial metabolic engineering approach for optimizing cellular phenotype. This methodology led to the identification of rpoA/rpoD mutants capable of enhancing L-tyrosine production over a rationally engineered parental, albeit in a mutated genetic background. An additional unique feature of this work is the successful identification of the specific chromosomal mutations and global regulators that, in combination, allowed the complete recovery of the L-tyrosine overproduction phenotype.

Fig. 4. Overexpression of evgA and relA in parental and mutant backgrounds. The genes evgA and a truncated form of relA were individually overexpressed in four different backgrounds—the parental strain P2 and the cured backgrounds of rpoA14, rpoA27, and rpoD3. Two promoter strengths were tested for each gene in order to ascertain the effects of both low- and high-level constitutive expression on cellular phenotype. All strong promoters were P₅. Weak promoters for evgA and relA were AA and II, respectively (Table S1). L-tyrosine titers were measured after 48 h cultivation in 50 ml MOPS minimal medium.

Table 2. List of downregulated pathways in rpoD3, rpoA14, and rpoA27

<table>
<thead>
<tr>
<th>Pathway</th>
<th>rpoD3</th>
<th>rpoA14</th>
<th>rpoA27</th>
</tr>
</thead>
<tbody>
<tr>
<td>Arginine synthesis</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>Isoleucine synthesis</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>Leucine/valine synthesis</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>Histidine synthesis</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>Tryptophan synthetesis</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>Lysine synthesis</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>Glutamate synthesis</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>De novo purine/pyrimidine biosynthesis</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>DNA replication</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>Ribosomal proteins and RNA</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>Fatty acid elongation</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
</tbody>
</table>

Examination of the transcriptional data revealed that the overexpression of one of two transcriptional regulators—evgA or relA—could completely supplant the phenotypic requirement for either a mutant ppoA or purD gene. This interchangeability provides compelling evidence regarding the specific biochemical mechanisms induced by these mutant transcriptional components and clearly implicates both acid resistance and the stringent response as important pathways for eliciting L-tyrosine overproduction in E. coli. However, because (p)ppGpp levels are known to directly or indirectly govern the onset of both responses (23–25), it is plausible that these similar phenotypic outcomes are in fact linked by a common mode of action. Although we still lack a complete mechanistic understanding of how alterations in these global cellular processes can lead to increases in L-tyrosine titers, these results provide a solid footing for future investigations aimed at deciphering the underlying mechanism.

Whole genome sequencing of our strains was required to identify spontaneous chromosomal alterations found to be essential
for a L-tyrosine overproduction phenotype. Although large-scale sequencing projects oftentimes return more mutations than can be functionally characterized (30), only a single base-pair change was identified in each of our three gTME strains. These individual mutations were isolated to the genes hisH and purF, two loci encoding proteins possessing striking similarities in both function and structure. Each belongs to the class of glutamine amidotransferases, enzymes which utilize glutamine as a source of an amide nitrogen with the concomitant production of glutamate. Remarkably, both of the internal substitutions recovered during our analysis (V5G in rpoD3 and L82R in rpoA14) map to helical structures directly adjacent to two catalytic cysteine residues with important roles in glutamate binding (31–33). It is therefore possible that these chromosomal mutations may exert their effects by altering the substrate affinities, and hence, activities of these enzymes. Such a result would then point to relative glutamine/glutamate levels as a possible control point for L-tyrosine production, a theory with plausible grounds given the requirement for enzymes which utilize glutamine as a source of an amide nitrogen with the concomitant production of glutamate. Following digestion and ligation, preparations of the resulting pHACM-derived plasmid. Following three to four rounds of reinoculation, routine chemical transformation protocols were utilized in the construction of strains containing a combination of wild-type or mutant backgrounds and plasmids.

**Transcriptional Analysis.** Strains P2, rpoA14, rpoA27, and rpoD3 were grown in 50 ml MOPS minimal medium (Teknova) (28) to an OD600 of approximately 0.4. Triplicates samples of RNA (on three separate days) were then extracted using the QIAGEN RNeasy Mini Kit according to the manufacturer's protocol. Microarray services were provided by the David Koch Institute for Integrative Cancer Research Microarray Technologies Core Facility using the GeneChip E. coli Genome 2.0 Arrays (Affymetrix). Arrays were run in triplicate with biological replicates to allow for statistical confidence in differential gene expression. Microarray data was deposited in the Gene Expression Omnibus database under accession number GSE21652. Expression profile deviations between mutants and parental were identified by a Significance Analysis of Microarrays (SAM) analysis using SAM 3.0 (35).

**Whole Genome Sequencing and SNP Detection.** Whole genome sequencing for strains P2, rpoA14, rpoA27, and rpoD3 was performed using the Illumina/Solexa Genome Analyzer System. Briefly, genomic DNA from all four strains was extracted using the Wizard Genomic DNA Purification Kit (Promega). Samples were then fragmented and prepared for paired-end sequencing using the Paired-End DNA Sample Prep Kit (Illumina). Samples were analyzed and processed at the Massachusetts Institute of Technology Biopolymers Laboratory, with duplicate lanes used for each strain. Sequence alignment and analysis, including the detection of insertions, deletions, and SNPs, was performed by the David Koch Institute Bioinformatics Facility. All sequence deviations from P2 were additionally validated by Sanger sequencing.

**Fed-batch Fermentations of rpoA14** 
RpoA14 was cultivated in 5 ml LB medium until the mid-exponential phase, then transferred into 2–50 ml MOPS minimal or R medium flask cultures at a starting OD600 of 0.1. These were cultivated at 37 °C with 225 rpm orbital shaking until the mid-exponential phase and were subsequently used as the inoculum for a 1.5–5 ⃯ culture (5% by volume). Fed-batch fermentations for rpoA14R were performed in 2–1 glass vessels using the BioFlo110 modular fermentor system (New Brunswick Scientific). Two separate experiments were conducted using MOPS minimal medium and R medium (29) supplemented with 0.5 g/l L-phenylalanine and 68 μg/ml chloramphenicol. Fermentations were performed at 37 °C with
the pH automatically adjusted to 7.0 using a 30% solution of ammonium hydroxide. Dissolved oxygen (pO2) was maintained at ≥25% by agitation speed (100–1,000 rpm) and gas mix control (air:oxygen). Foam formation was controlled by the addition of Tego Antifoam K911 (Evonik Goldschmidt). Operations were controlled and recorded with the BioCommand Lite Data Logging software (New Brunswick Scientific).

**Analytical Methods.** The L-tyrosine content of cell-free culture supernatants was analyzed via HPLC with a Waters 2690 Separations module and a Waters 996 Photodiode Array detector set to a wavelength of 278 nm. Samples were analyzed via HPLC with a Waters 2690 Separations module and a Waters 996 Photodiode Array detector set to a wavelength of 278 nm. Samples were separated on a Waters Resolve C18 column with 0.1% (vol/vol) trifluoroacetic acid (TFA) in water (solvent A) and 0.1% (vol/vol) TFA in acetonitrile (solvent B) as the mobile phase. The following gradient was used at a flow rate of 1 ml/min: 0 min, 95% solvent A + 5% solvent B; 8 min, 20% solvent A + 80% solvent B; 10 min, 80% solvent A + 20% solvent B; 11 min, 95% solvent A + 5% solvent B. Cell densities of cultures were determined by measuring their absorbance at 600 nm with an Ultrospec 2100 pro UV/Vis spectrophotometer (Amersham Biosciences).

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