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# Elucidating aromatic acid tolerance at low pH in *Saccharomyces cerevisiae* using adaptive laboratory evolution

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Toxicity from the external presence or internal production of compounds can reduce the growth and viability of microbial cell factories and compromise productivity. Aromatic compounds are generally toxic for microorganisms, which makes their production in microbial hosts challenging. Here we use adaptive laboratory evolution to generate *Saccharomyces cerevisiae* mutants tolerant to two aromatic acids, coumaric acid and ferulic acid. The evolution experiments were performed at low pH (3.5) to reproduce conditions typical of industrial processes. Mutant strains tolerant to levels of aromatic acids near the solubility limit were then analyzed by whole genome sequencing, which revealed prevalent point mutations in a transcriptional activator (Aro80) that is responsible for regulating the use of aromatic amino acids as the nitrogen source. Among the genes regulated by Aro80, *ESBP6* was found to be responsible for increasing tolerance to aromatic acids by exporting them out of the cell. Further examination of the native function of Esbp6 revealed that this transporter can excrete fusel acids (byproducts of aromatic amino acid catabolism) and this role is shared with at least one additional transporter native to *S. cerevisiae* (Pdr12). Besides conferring tolerance to aromatic acids, *ESBP6* overexpression was also shown to significantly improve the secretion in coumaric acid production strains. Overall, we showed that regulating the activity of transporters is a major mechanism to improve tolerance to aromatic acids. These findings can be used to modulate the intracellular concentration of aromatic compounds to optimize the excretion of such products while keeping precursor molecules inside the cell.

adaptive laboratory evolution | aromatic acid | tolerance | transporter

*Saccharomyces cerevisiae* has proven to be a great microbial host for producing compounds of plant origin (1, 2). A frequent hurdle when constructing new microbial cell factories is the toxicity of the final product or some of the intermediary metabolites that can result from introducing heterologous enzyme activities. Compounds with aromatic groups have shown particular toxicity (3–5) and mitigating their inhibition on growth can help accelerate the design and improve the performance of new cell factories.

Aromatic acids, such as coumaric acid, can be used as platform chemicals but are also important precursors for producing plant natural products such as flavonoids (6) and phenols (7). Even at low concentration, coumaric acid and related compounds (i.e., phenylpropanoic acids) are known inhibitors of *S. cerevisiae* growth (4, 8), making their presence inside the cell a potential issue when engineering the production of natural products that require them as precursors. Given the limited information of how phenylpropanoic acids inhibit growth, using techniques such as adaptive laboratory evolution (ALE) can help generate tolerant strains and further elucidate the genetic basis underlying tolerance phenotypes and how these compounds stress their host (9).

Acid tolerance in *S. cerevisiae* has been studied in detail for many organic acids and usually requires the cell to up-regulate transporters (e.g., Pdr12) to reduce the concentration of the acid anions inside the cell and proton pumps (Pma1) to restore a neutral pH in the cytosol (reviewed in refs. 10–12). Organic acids are known to exhibit a pH-dependent toxicity because only the protonated form of the acid can cross the membrane, making them more toxic at lower pH (10). Regarding tolerance to aromatic acids in *S. cerevisiae*, the deletion of *PDR12* was shown to induce sensitivity to fusel acids (phenylacetic acid and indoleacetic acid) (13). Furthermore, tolerance to benzoic acid has also been shown to be mediated by Pdr12 (14) or other transporters like Tpo2 (15).

In this work, we used ALE to study tolerance mechanisms of *S. cerevisiae* to two aromatic acids: coumaric acid and ferulic acid. Tolerance to either aromatic acid was traced back to mutations in a transcription activator involved in aromatic amino acid (AAA) catabolism (*ARO80*). Further analysis of possible targets of Aro80 showed that increased expression of the transporter *ESPB6* (previously not known to transport aromatic acids) could reproduce the tolerance phenotypes observed. Since Esbp6 is active when AAAs are used as the nitrogen source, we explored its role in these conditions and determined that this transporter can export fusel acids from the cytosol. The role of Esbp6 in fusel acid tolerance partly overlaps with the function of another

## Significance

Understanding the tolerance mechanism to inhibiting chemicals can help in the design of more robust microbial strains for use in industrial bioprocesses. Aromatic acids are important bioactive molecules and precursors for many plant natural products. This study focused on analyzing *Saccharomyces cerevisiae* mutants tolerant to aromatic acids and revealed that regulating the transport of these compounds from the cytosol is the main mechanism to improve tolerance. Using this knowledge, the transport of aromatic acid can be fine tuned to improve the secretion of aromatic compounds and improve the robustness of yeast strains for industrial use.

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transporter, Pdr12, but the mechanism of activation between the two differs. While *ESBP6* is activated by Aro80 in response to the presence of AAAs, *PDR12* is up-regulated by War1 possibly as a response to the presence of fusel acids.

## Results

**Adaptive Laboratory Evolution and Mutant Screening.** ALE was utilized to generate multiple independent lineages of evolved populations with increased tolerance to the coumaric and ferulic acids. First, the native tolerance of *S. cerevisiae* to each aromatic acid was determined by measuring growth in medium with different concentrations of each compound. Since organic acids are known to exhibit a pH-dependent toxicity (higher toxicity at low pH), we performed the growth tests and evolution at a pH of 3.5 using citrate-phosphate buffer as described by Kildegaard et al. (16). Both aromatic acids have a maximum solubility close to 1 g/L in the conditions used, and a starting concentration of 0.2 g/L for either acid was enough to apply an adequate level of selection pressure on the growth of *S. cerevisiae* for the ALE experiments.

To maximize the genetic diversity that could be obtained during the evolutionary paths, five single clones were used as a seed culture to start five independent replicates for each acid. The growth of each replicate was monitored periodically during evolution and the concentration of the acids was slowly increased to encourage incremental increases in fitness (maximum growth rate) by using a strategy similar to a previous publication (17). The experiment was allowed to continue as long as the fitness increased significantly for a certain time period (18). When the fitness appeared to reach a plateau, the ALE experiment was terminated, and an aliquot of each evolved population was plated to isolate individual clones. Overall, all evolution replicates reached the solubility limit for each aromatic acid (1 g/L), which shows the ALE experiments succeeded in producing mutants with high tolerance. It is worth noting that the solubility and relative toxicity of both aromatic acids change with pH. At low pH (3.5), the maximum solubility is low, but their inhibiting effect is quite high. The ALE mutants obtained here can tolerate 1 g/L of either aromatic acid at pH 3.5 but have the potential to tolerate much higher acid concentrations at higher pHs. On average, ALE experiments for coumaric acid tolerance were terminated after 90 transfers and ferulic acid tolerance was evolved for 69 transfers.

Individual clones were isolated and screened from the independently evolved populations to identify those with increased tolerance and to use for further studying. Within each evolution experiment, we picked 10 individual clones from culture plates and grew them in medium with inhibiting concentrations of each aromatic acid (0.8 g/L). For each set of clones, we chose the 3 with the highest growth rate for further characterization. Fig. 1A and B shows the average growth rates and maximum optical densities for the 3 best performing clones within each evolution replicate (EVO1-EVO5) in comparison to a reference strain (wild type [WT]). Generally, all selected clones showed a superior performance in terms of increased tolerance and fitness compared to the reference strain. Most of the coumaric acid evolved strains were able to achieve a maximum growth rate above  $0.30\text{ h}^{-1}$ , which is considerably higher than the reference strain (WT) but not as high as the reference strain in medium without aromatic acids ( $0.41\text{ h}^{-1}$ ). The only exception was found in the clones isolated from the C-EVO4 population, whose average growth rate was only  $0.23\text{ h}^{-1}$ . Consistent with the performance of cells during the preliminary acid inhibition test, the selected ferulic acid evolved clones showed a lower fitness than coumaric acid evolved clones for the same acid concentration.

In order to understand if the evolved phenotypes relied on degradation of the aromatic acids, the cultures of two evolved clones for each acid were analyzed by high performance liquid chromatography (HPLC). For all four cultivations the concentration

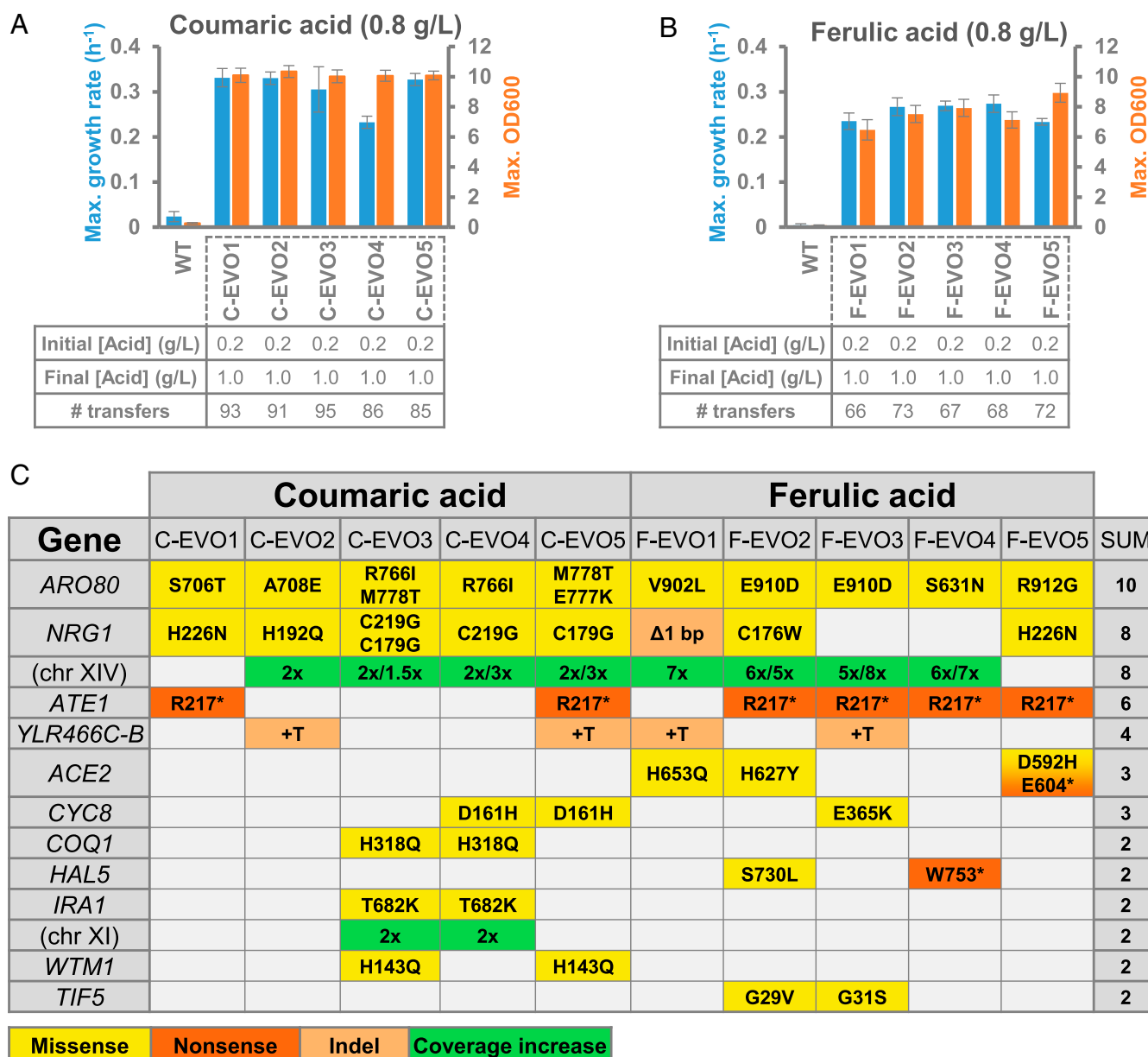
of coumaric acid and ferulic acid remained stable at 800 mg/L, indicating that the evolved phenotypes relied on physiological adaptation to tolerate the presence of aromatic acids. Since it has previously been shown that another strain of *S. cerevisiae* can degrade aromatic acids (19), it was surprising to realize that none of the tested ALE mutants seemed to degrade these compounds.

**Whole Genome Sequencing of Evolved Mutants.** To investigate the genetic basis behind the tolerance phenotypes obtained, we performed whole genome sequencing on three mutant clones exhibiting the highest growth rates within each independently evolved population (for a total of 30 clones). Fig. 1C shows an overview of the genes mutated in at least two independently evolved populations (full list provided in *SI Appendix, Tables S3 and S4*). Among the most common genetic changes, we found mutations in *ARO80*, *NRG1*, and an increase (up to eightfold) in read depth for certain sections of chromosome XIV. Mutations in *ARO80* seem of crucial importance given their presence in all of the clones sequenced for both aromatic acids. The diversity of mutations found in *ARO80* is also a good indicator for the importance of this gene in tolerance to aromatic acids. From a total of 10 independent evolution experiments, we found 9 distinct amino acid substitutions (Fig. 1C) but no cases of nonsense mutations. This suggests that selective pressure for *ARO80* mutations is high, but loss of function does not appear to be responsible for the tolerance phenotype. Aro80 is a transcription factor previously shown to activate genes needed for using AAAs as the nitrogen source (20, 21). In the presence of AAAs, Aro80 induces the transcription of genes from the Ehrlich pathway (22), leading to the assimilation of nitrogen from these compounds and excretion of fusel acids and fusel alcohols. Given the similarity of coumaric and ferulic acids to some of the metabolites in the Ehrlich pathway, mutations in *ARO80* indicated that genes involved in detoxifying such compounds were probably being activated.

Trying to connect mutations in *NRG1* to aromatic acid tolerance is not as straightforward as for *ARO80*. *NRG1* codes for a transcriptional repressor involved in controlling the expression of many glucose-repressed genes (23) and also regulating tolerance to  $\text{Na}^+$ ,  $\text{Li}^+$ , and growth in alkaline conditions (24). Although *NRG1* was shown to be required for tolerance to other stressing conditions, the reason for it being frequently mutated in aromatic acid tolerant strains is not known.

Increases in coverage of chromosome XIV were used to pinpoint a gene related to the tolerance phenotype by looking at the intersection of coverage increases. Computing the intersection between the DNA regions affected in coumaric and ferulic acid tolerance mutants revealed a 35-kb segment that is common to all mutants (spanning open reading frames [ORFs] from YNL123W to YNL141W). We decided to focus our interest in this mutated region by comparing the documented targets of Aro80 (retrieved from Yeasttract) (25) with regions of chromosome XIV that are affected by a coverage increase. From this approach, it was concluded that there is one gene that met the criteria (*SI Appendix, Table S5*). The gene in question is *ESBP6* (*YNL025C*) and codes for a protein that has similarity to monocarboxylate permeases. This gene was shown to be induced by Aro80 in the presence of AAAs (21) but its role has never been uncovered. Its cellular localization is not clear according to YeastRGB (26), but one high throughput dataset suggested that Esbp6 is located in the cell periphery (27). Furthermore, an overexpression mutant of *ESBP6* was shown to confer tolerance to 6% lactic acid (28).

*ATE1*, *ACE2*, and *CYC8* are also mutated in multiple evolution lineages. Mutations in *ACE2* were shown to prevent yeast cells from separating from each other after cell division, which leads to the formation of agglomerates (29, 30). Even though the inner cells inside the agglomerates may benefit from some



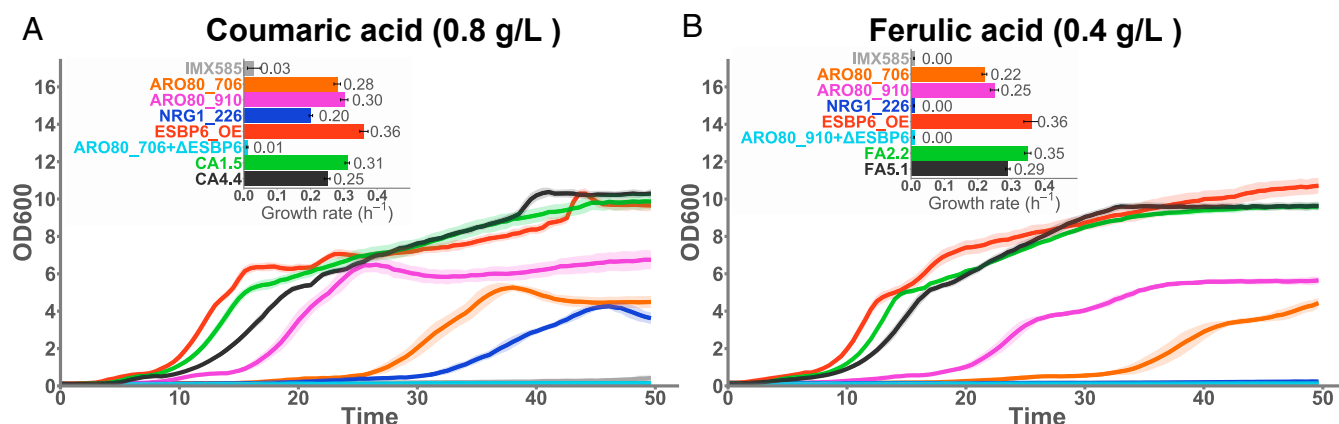
**Fig. 1.** Summary of the results from the adaptive laboratory evolution experiments for tolerance to coumaric acid and ferulic acid. Average growth rates ( $\text{h}^{-1}$ ) and maximum optical densities (OD600) achieved by the three best performing mutants isolated from each of the five populations evolved for coumaric acid tolerance (A) and ferulic acid tolerance (B). Error bars represent the SD from the three best mutants (each with three replicates). (C) Table of mutations for genes mutated in more than one population. Information regarding the individual growth rates, maximum optical densities, and lists of all mutations detected can be found in [SI Appendix, Tables S1–S4](#).

protection against harsh conditions, we have previously observed that mutations in *ACE2* contribute very little to improving the growth rate in the presence of organic acids (31). Regarding the mutations in *CYC8*, we previously showed that mutations in this gene can confer tolerance to pimelic acid (31). However, the exact mechanism leading to the tolerance was never uncovered. Finally, mutations in *ATE1* were found to be the same (R217\*) in all mutants where they were present. This is an indication that this mutation probably arose in the seed culture used to start the ALE experiments and is not relevant for the tolerance phenotypes.

**Reverse Engineering of Mutations to Determine Causality.** In order to find the genetic causes for the tolerance phenotypes, we implemented a selection of the most frequent mutations in a

strain with a clean background. To test the contribution of mutations in *ARO80*, we selected two different point mutations found in this gene: one mutation taken from a lineage evolved in coumaric acid (*ARO80*<sup>S706T</sup> found in C-EVO1) and another present in two lineages evolved in ferulic acid (*ARO80*<sup>E910D</sup> found in F-EVO2 and F-EVO3). Fig. 2 shows the growth curves for *ARO80*<sup>S706T</sup> (*ARO80*<sub>706</sub>) and *ARO80*<sup>E910D</sup> (*ARO80*<sub>910</sub>) mutants in the presence of inhibiting concentrations of coumaric acid or ferulic acid. Both point mutations tested for *ARO80* confer significant tolerance to coumaric and ferulic acid in comparison to a reference strain (IMX585). However, it is clear from Fig. 2 that the mutant *ARO80*<sup>E910D</sup> reaches the stationary phase faster than *ARO80*<sup>S706T</sup>. Although the maximum growth rates do not differ much between the two *ARO80* mutants tested,





**Fig. 2.** Phenotypic characterization of the reverse engineered mutants in the presence of the aromatic acids. (A) Growth curves and maximum growth rates of a reference strain (IMX585) in comparison to five reverse engineered strains (*ARO80*<sup>S706T</sup>, *ARO80*<sup>E910D</sup>, *NRG1*<sup>H226N</sup>, *ESBP6*\_OE, and *ARO80*<sup>S706T</sup>+ $\Delta$ *ESBP6*) and two evolved mutants tolerant to coumaric acid (CA1.5 and CA4.4). Cells were grown in 96-well plates with a volume of 250  $\mu$ L in medium with 0.8 g/L of coumaric acid. (B) Growth curves and maximum growth rates of a reference strain (IMX585) in comparison to five reverse engineered strains (*ARO80*<sup>S706T</sup>, *ARO80*<sup>E910D</sup>, *NRG1*<sup>H226N</sup>, *ESBP6*\_OE, and *ARO80*<sup>E910D</sup>+ $\Delta$ *ESBP6*) and two evolved mutants tolerant to ferulic acid (FA2.2 and FA5.1). Cells were grown in 96-well plates with a volume of 250  $\mu$ L in medium with 0.4 g/L of ferulic acid. All cultivations were performed for three biological replicates (each with three technical replicates) and the average OD600s are shown as solid lines with the interval encompassing the SD represented as a shaded area. The maximum growth rates represent the average of individually calculated growth rates for all replicates and the errors bars represent the SD.

the length of the lag phase is shorter for *ARO80*<sup>E910D</sup>. When we compare the *ARO80* mutants to the ALE mutants where each mutation was taken from (CA1.5 and FA2.2), we see that mutations in *ARO80* are not enough to replicate the full tolerance phenotype, especially in terms of the lag phase length and capacity to grow after the diauxic shift.

Frequent mutations in *NRG1* in both coumaric and ferulic acid tolerant mutants provided justification to test if point mutations in this gene were contributing to the tolerance phenotypes. Since the mutation *NRG1*<sup>H226N</sup> appeared in mutants for both aromatic acids, we constructed a mutant displaying this mutation (*NRG1*\_226) and tested its growth in the presence of coumaric acid or ferulic acid (Fig. 2). In coumaric acid, the strain *NRG1*\_226 performs better than the control strain, but it shows a long lag phase and lower maximum growth rate (0.20 h<sup>-1</sup>) than the evolved clones and *ARO80* mutants. In ferulic acid, no growth was detected, even at 0.4 g/L. These findings suggested that the effect of *NRG1* is minor on its own and could be dependent on the presence of additional mutations to confer a significant fitness improvement.

The increase in read coverage for chromosome XIV would be difficult to reverse engineer given the length of DNA involved. Nevertheless, the presence of a target of Aro80 in the affected areas suggested that a single gene could be the reason behind this genetic alteration. To test this hypothesis, we introduced an extra copy of *ESBP6* under the control of a strong promoter (*P*<sub>TEF1</sub>) in a clean background strain, resulting in strain *ESBP6*\_OE. As shown in Fig. 2, *ESBP6*\_OE grows robustly in medium with inhibiting amounts of coumaric or ferulic acid. In comparison to all ALE mutants tested, the strain *ESBP6*\_OE matches or outperforms them in terms of maximum growth rate and length of the lag phase. To evaluate if *ESBP6* was responsible for the tolerance exhibited by *ARO80* mutants, we deleted this gene in each of the *ARO80* single mutant background (strains *ARO80*<sup>S706T</sup>+ $\Delta$ *ESBP6* and *ARO80*<sup>E910D</sup>+ $\Delta$ *ESBP6*). The growth profiles of both strains showed that knocking out *ESBP6* completely abolishes the tolerance phenotype observed for *ARO80*<sup>S706T</sup> and *ARO80*<sup>E910D</sup>, revealing that *Esbp6* has an essential role in tolerance to aromatic acids.

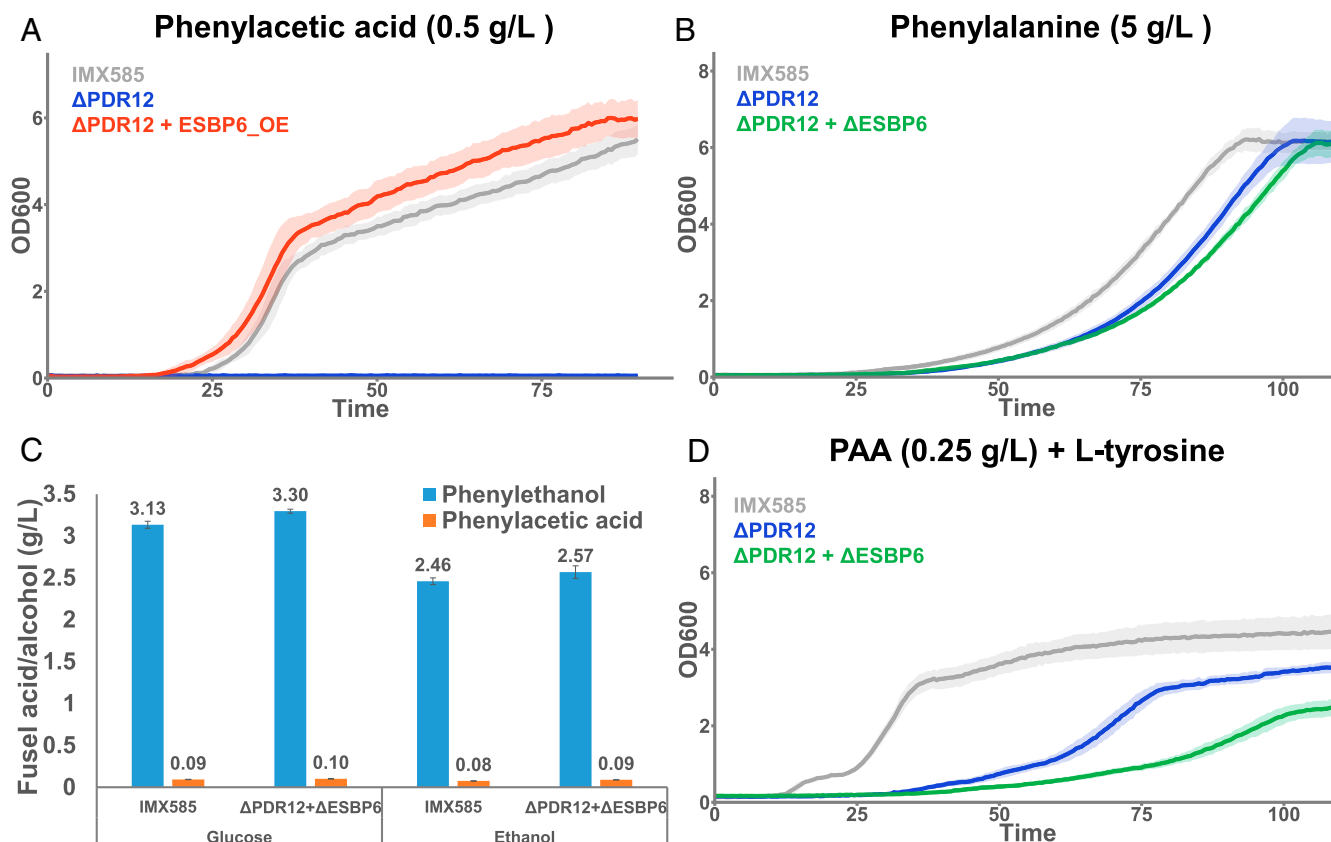
Overall, testing the impact of each mutation in a clean background strain revealed that the tolerance phenotypes evolved

here were likely caused by the up-regulation of *ESBP6*. This was achieved via evolution by point mutations in the transcription factor *ARO80* that controls *ESBP6* and by an increase in the number of copies of the *ESBP6* locus in the genome.

***ESBP6*'s Role in *S. cerevisiae*.** Testing individual mutations from tolerant mutants (Fig. 2) revealed that *ESBP6* overexpression resulted in the highest tolerance to aromatic acids. Although the native function of *ESBP6* was unclear, the expression of this gene was shown to be induced in the presence of AAAs by the transcription factor Aro80 (21). When *S. cerevisiae* uses AAAs as the nitrogen source, Aro80 induces the expression of genes from the Ehrlich pathway. Among the compounds involved in the Ehrlich pathway, fusel acids (phenylacetic acid, 4-hydroxyphenylacetic acid, and indoleacetic acid) are the most similar to coumaric and ferulic acids. The exact way fusel acids are excreted to the medium is not known, but the presence of the ABC transporter Pdr12 was shown to be required for tolerance to phenylacetic acid (PAA) and indoleacetic acid (IAA) (13).

In order to understand if *Esbp6* has a role in fusel acid tolerance, we compared the growth of a *PDR12* deletion mutant with and without the overexpression of *ESBP6* ( $\Delta$ *PDR12*+*ESBP6*\_OE and  $\Delta$ *PDR12*, respectively) in medium with PAA or IAA. Fig. 3A shows that in the presence of 0.5 g/L of PAA, the *PDR12* deletion mutant ( $\Delta$ *PDR12*) exhibits a growth disability when compared to the reference strain (IMX585), confirming the essentiality of Pdr12 in these conditions. When *ESBP6* is overexpressed on a  $\Delta$ *PDR12* background ( $\Delta$ *PDR12* + *ESBP6*\_OE), growth is restored to wild-type levels (Fig. 3A). The observation that *Esbp6* can compensate for Pdr12 absence shows that they have overlapping roles in tolerance to aromatic acids.

Similar tests performed in the presence of IAA showed that at a concentration of 0.1 g/L, the deletion of *PDR12* does not affect growth (SI Appendix, Fig. S14). This observation contradicts a previous report (13) about the role of *PDR12* in tolerance to IAA, but it should be noted that a different media formulation and method for growth evaluation were used. Even though the inactivation of *PDR12* showed little effect on tolerance to IAA, overexpressing *ESBP6* was still enough to improve growth in comparison to the reference strain (SI Appendix, Fig. S14), suggesting a certain role for this gene in tolerance to this fusel acid. Besides PAA and IAA, we



**Fig. 3.** Influence of *PDR12/ESBP6* on fusel acid tolerance and AAA metabolism. (A) Growth curves in medium with 0.5 g/L of phenylacetic acid for a reference strain (IMX585) in comparison to a *PDR12* deletion mutant ( $\Delta$ PDR12) and a double mutant lacking *PDR12* and displaying the overexpression of *ESBP6* ( $\Delta$ PDR12+*ESBP6*<sub>OE</sub>). Cells were grown in 96-well plates with a volume of 250  $\mu$ L in medium with 0.5 g/L of phenylacetic acid. (B) Growth curves in medium with 5 g/L of L-phenylalanine as nitrogen source for a reference strain (IMX585) in comparison to a *PDR12* deletion mutant ( $\Delta$ PDR12) and a double deletion mutant lacking *PDR12* and *ESBP6* ( $\Delta$ PDR12+ $\Delta$ ESBP6). Cells were grown in 96-well plates with a volume of 250  $\mu$ L in medium with 5 g/L of L-phenylalanine as nitrogen source and 20 g/L of ethanol as carbon source. (C) Extracellular concentration of phenylacetic acid and phenylethanol in cultures of a reference strain (IMX585) and a double deletion mutant ( $\Delta$ PDR12+ $\Delta$ ESBP6) grown in medium with 20 g/L of glucose or 10 g/L of ethanol as carbon source. Cells were grown in shake flasks with a volume of 25 mL in medium with the mentioned nitrogen and carbon sources. Error bars represent the SD from three biological replicates. (D) Growth curves in medium with 0.25 g/L of phenylacetic acid and 0.5 g/L of L-tyrosine as the nitrogen source for a reference strain (IMX585) in comparison to a *PDR12* deletion mutant ( $\Delta$ PDR12) and a double deletion mutant lacking *PDR12* and *ESBP6* ( $\Delta$ PDR12+ $\Delta$ ESBP6). Cells were grown in 96-well plates with a volume of 250  $\mu$ L in medium with 0.5 g/L of L-tyrosine as the nitrogen source and 0.25 g/L of phenylacetic acid. All growth curves were performed for three biological replicates (each with two technical replicates) and the average OD600s are shown as solid lines with the interval encompassing the SD represented as a shaded area.

also tested the effect of 4-hydroxyphenylacetic acid on the growth of *S. cerevisiae*, but no toxicity was detected up to 2 g/L, which is in accordance with the lower toxicity of this fusel acid to *S. cerevisiae* (13).

Since *PDR12* and *ESBP6* were shown to be important for fusel acid tolerance, we then tested if single and double disruption of these genes affects growth of *S. cerevisiae* when growing on AAAs. Fig. 3B shows that when growing on L-phenylalanine (precursor of PAA), even the simultaneous inactivation of *ESBP6* and *PDR12* (strain  $\Delta$ PDR12+ $\Delta$ ESBP6) seems to have little effect. This could be a consequence of two different aspects: the amount of fusel acids produced is below the toxicity level needed to observe a phenotype; and there are additional transporters capable of excreting fusel acids to the medium. To test the first alternative, the amount of PAA and phenylethanol (PE) excreted by cells growing on L-phenylalanine as the nitrogen source was quantified. As shown in Fig. 3C, independently of the carbon source or strain used, the major Ehrlich pathway byproduct excreted is PE. Although a small quantity of PAA is still produced, the amount is much lower than what is needed to induce toxicity. In fermentative conditions the production of PE was expected because the high NADH/NAD<sup>+</sup> promoted the

production of reduced byproducts. By using ethanol as a carbon source we expected that the growth in full oxidative conditions would promote fusel acid production. However, the results in Fig. 3C showed that the effect of the carbon source on the amount of fusel acid produced was quite low and that factors other than the cell's oxidative state may be necessary for forcing the production of fusel acids. It is worth noting that all our tests were performed at low pH (3.5), which favors the reentry of any excreted carboxylic acids into the cell.

Given the low levels of PAA produced by cultures grown with AAAs as the nitrogen source, the role of *ESBP6* in these conditions could not be directly confirmed. In order to force a response, we added PAA to cultures grown with L-tyrosine or L-tryptophan as nitrogen sources. Adding 0.5 g/L of PAA was enough to completely abolish growth of a *PDR12* deletion strain when L-tyrosine or L-tryptophan were provided as the nitrogen source (SI Appendix, Fig. S1B). However, by reducing the concentration of PAA to 0.25 g/L, we could see that a *PDR12* deletion mutant ( $\Delta$ PDR12) could grow in the presence of L-tyrosine (Fig. 3D) but slower than the reference strain (IMX585). Furthermore, when *ESBP6* was deleted in addition to *PDR12* ( $\Delta$ PDR12+ $\Delta$ ESBP6) we observed

that the resulting mutant was still viable but its growth was reduced in comparison to the *PDR12* deletion mutant (Fig. 3D). These results indicate that when L-tyrosine is used as the nitrogen source Pdr12 is the main source of tolerance to PAA, but Esbp6 also contributes to it. It is worth noting that the basal growth displayed by the double deletion mutant suggests that other genes may also be contributing to tolerance to PAA.

Besides the growth tests on PAA, we also tried to confirm the roles of *PDR12* and *ESBP6* on IAA tolerance in the presence of AAA. As shown in *SI Appendix, Fig. S1C*, growth on 0.1 g/L of IAA with L-tryptophan as the nitrogen source confirmed that the role of Pdr12 in tolerance to this fusel acid seems minor (*SI Appendix, Fig. S1C*). However, when *ESBP6* is also inactivated there is a clear reduction in growth. This observation confirms that for the fusel acid IAA, Esbp6 plays a more important role than Pdr12.

***ESBP6* Influence on Coumaric Acid Production Strains.** Since Esbp6 confers tolerance to several aromatic acids, we tested if overexpressing its coding gene would also improve the secretion of this class of compounds. Using strains engineered to overproduce coumaric acid, we quantified the amount of secreted product with and without the overexpression of *ESBP6*. The production tests were carried in the same conditions as the evolution experiments (low pH) to maximize the potential effect of improving product secretion. Fig. 4A shows the impact of overexpressing *ESBP6* on two coumaric acid production strains, QL01 and QL38, constructed and characterized elsewhere (32). We selected two strains with varying levels of coumaric acid production to observe if this would affect our results. As shown in Fig. 4A, the coumaric acid production levels increased almost by the same amount in the lower and higher producing strains (47% increase vs. 38% increase). The consistent improvement indicates that increased levels of Esbp6 can promote the export of coumaric acid from the cell and can be used as a strategy to potentially reduce the level of aromatic acids in the cytosol. Although low pH conditions were optimal to observe the effect of *ESBP6* on coumaric acid production, the titers of coumaric acid were considerably lower than previously reported for experiments run at pH 6.0 (32). The medium pH affects the relative abundance of protonated organic acids and consequently

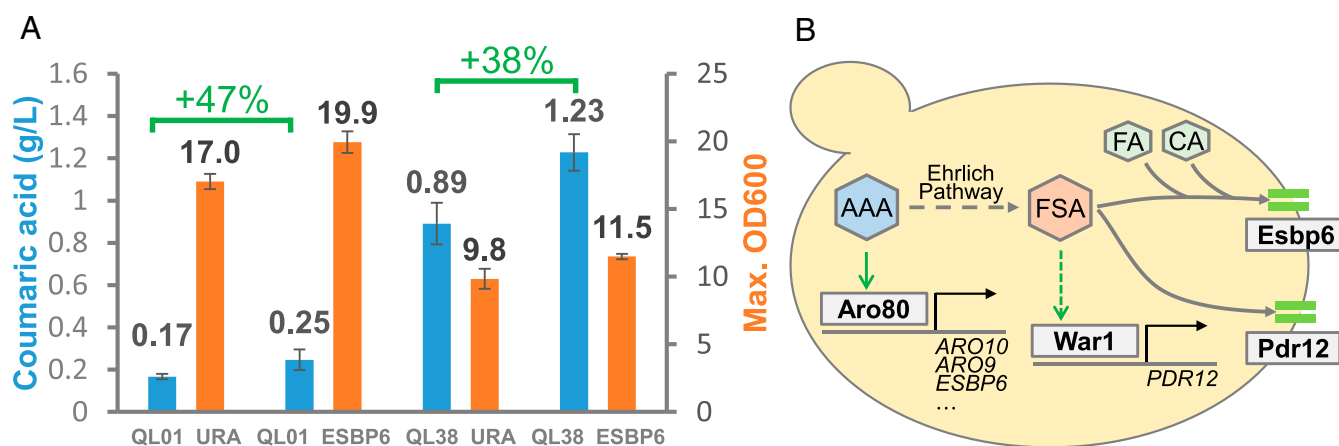
their ability to enter the cell by diffusion. At low pH, besides secreting the coumaric acid produced biosynthetically, the cell must also secrete the increased flow of coumaric acid reentering the cell by diffusion. Besides boosting the export of coumaric acid, the overexpression of *ESBP6* also improved cell growth by 17% in both strains. These results show that besides increasing the secretion of coumaric acid, *ESBP6* can reduce the stress burden on the cell and improve the biomass yield.

## Discussion

ALE and subsequent analysis of mutations has been an indispensable tool to study metabolic pathways and general physiology of eukaryotic cells. Here, using ALE we have generated mutant strains tolerant to two aromatic acids, coumaric and ferulic acids, at levels near their solubility limit. By reverse engineering selected mutations, we showed that increasing the expression of the transporter gene *ESBP6* was a main mechanism to tolerate aromatic acids that arose during the ALE experiments. In most of the tolerant mutants, the likely up-regulation of Esbp6 was associated with nine different mutations in *ARO80* and duplication of segments from chromosome XIV that included the *ESBP6* locus.

Since Esbp6 is activated in the presence of AAAs, we investigated its potential role in medium containing AAAs as nitrogen source and in the presence of fusel acids (aromatic acids derived from AAA). This analysis showed that Esbp6 had a role in tolerance to fusel acids, but its importance is difficult to pinpoint because of redundant transporters present in *S. cerevisiae* and hurdles in designing experiments where the cell is forced to produce fusel acids instead of fusel alcohols. The overlap in function between Esbp6 and Pdr12 was also examined and it revealed that both transporters can confer tolerance to some fusel acids but have different modes of activation (Fig. 4B). While *ESBP6* is activated by Aro80 when AAAs are used as nitrogen sources, Pdr12 is activated through War1 possibly by direct activation by fusel acids.

Besides providing insight into the metabolism of aromatic compounds in *S. cerevisiae*, the work described here can also have implications in the design of cell factories for natural product synthesis. By up-regulating the transport of aromatic acids, cell factories that produce this type of compounds can be



**Fig. 4.** Influence of *ESBP6* overexpression on coumaric acid production. (A) Coumaric acid titer and maximum OD600 in strains with (QL01\_ESBP6 and QL38\_ESBP6) and without (QL01\_URA and QL38\_URA) the overexpression of *ESBP6*. Error bars represent the SD from three biological replicates. (B) Schematic illustration of the interaction between aromatic acids, the transcription factors Aro80 and War1, and the transporters Esbp6 and Pdr12. In the presence of AAAs, Aro80 activates the transcription of genes necessary for degrading these compounds, which leads to the production of fusel acids and fusel alcohols. Fusel acid presence putatively induces the transcription of *PDR12* (through War1) and both Pdr12 and Esbp6 are responsible for excreting these compounds to the medium. Abbreviations: AAA, aromatic amino acids (L-phenylalanine, L-tyrosine, and L-tryptophan); FSA, fusel acids (phenylacetic acid, 4-hydroxyphenylacetic acid, and indoleacetic acid); CA, coumaric acid; and FA, ferulic acid.

improved by increasing the flux of product from the cytosol to the medium and by allowing cells to tolerate higher levels of product in the medium (33). Fig. 4A shows that increased levels of *ESBP6* can lead to a significant improvement in the production titer and higher cell density. Since diffusion of extracellular organic acids into the cells is increased at lower pH (10), by performing these production tests at pH 3.5 we replicated the harsh conditions of industrial fermentations where the product is present in severely inhibiting amounts. The improvement in product titer and OD600 showed that *ESBP6* overexpression can promote the export of acid anions from the cytosol and improve the biomass yield probably by reducing the stress burden imposed by excess aromatic acids inside the cell. We also would like to point out that previous studies achieved much higher coumaric acid titers in fed-batch fermentations but using a considerably higher pH. Although using a higher pH can reduce the toxicity of organic acids (10), in industrial conditions it is desirable to keep pH low to reduce the costs related to product purification (34).

Furthermore, cell factories that use aromatic acid as intermediates can potentially benefit from the down-regulation of the transporters for these classes of compounds if these intermediates are not toxic. By forcing the aromatic acid precursors to accumulate in the cytosol, the flux to natural products (such as flavonoids) can possibly be increased.

## Materials and Methods

A more detailed description of the materials and methods used in this work is provided in [SI Appendix, Supplementary Information Text, Materials and Methods](#).

**Yeast Strains and Medium.** The strain GL01 (31) was used for all ALE experiments and the strain IMX585 (35) was used for the reverse engineering work. Both strains were derived from the *S. cerevisiae* CEN.PK113-7D background. The full list of strains used and constructed in this work is provided in [SI Appendix, Table S6](#).

ALE experiments and growth tests were performed in the minimal medium described in ref. 36 and buffered at pH 3.5 by adding 140 mL/L of 0.5 M citrate solution and 60 mL/L of 1 M Na<sub>2</sub>HPO<sub>4</sub>. For testing strains in alternative nitrogen sources, (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> was replaced with 6.6 g/L of K<sub>2</sub>SO<sub>4</sub> and the required amino acid was added in the appropriate concentration (5 g/L of L-phenylalanine, 5 g/L of L-tryptophan, or 0.5 g/L of L-tyrosine).

**Strain Construction.** The reverse engineered strains containing deletions, point mutations, and insertions were constructed using the single gRNA

method described elsewhere (35). The list of primers used to engineer each mutation is available in [SI Appendix, Table S7](#) and the list of plasmids used and constructed in this study is provided in [SI Appendix, Table S8](#).

**Adaptive Laboratory Evolution.** Adaptive laboratory evolution experiments were conducted on an automated platform using a liquid handling robot as described previously (17, 37).

**Growth Screening.** The growth characterization of the postevolution mutants and reverse engineered strains was performed in a Growth Profiler 960 (EnzyScreen) using 96-half-deep-well microplates (with transparent bottom) with a total culture volume of 250  $\mu$ L, agitation at 250 rpm, temperature controlled at 30 °C, and initial OD600 of 0.05.

**DNA Extraction, Sequencing, and Data Analysis.** The Blood & Cell Culture DNA Mini Kit (Qiagen) was used to extract genomic DNA from 3 mL of the overnight yeast culture ( $\sim 5 \times 10^8$  cells) using the protocol recommended by the manufacturer. Sequencing was performed with the NextSeq (High Output Kit) with 2  $\times$  150 paired-end reads targeting a genome coverage of 100 $\times$  per sample.

Mutations in evolved clones were identified using bresEq. 0.30.2 (38) with Bowtie 2.2.8 as aligner (39) and using the reference genome of CEN.PK113-7D (40) as described before (31).

**Phenylacetic Acid and Phenylethanol Quantification.** Extracellular concentration of phenylacetic acid and phenylethanol in cultures of a reference strain (IMX585) and a double deletion mutant ( $\Delta$ PDR12+ $\Delta$ ESBP6) were quantified by HPLC using a method previously described by Li et al. (32).

**Coumaric Acid Production Tests.** Coumaric acid production tests were performed in 100-mL shake flasks containing 20 mL of the minimal medium without glucose and pH adjusted to 3.5. Six tablets of FeedBeads (SMFB08001, Kuhner Shaker) were added to the culture as the source of glucose. Cultures were inoculated with an initial OD600 of 0.05 and shaken at 200 rpm in an orbital incubator, at 30 °C during 96 h. For the strains QL38\_URA and QL38\_ESBP6, 1% of galactose was also added to induce genes controlled by the GAL promoters. *p*-coumaric acid was quantified using HPLC as described by Li et al. (32).

**Data Availability.** All study data are included in the article and [SI Appendix](#).

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