

The landscape of toxic intermediates in the metabolic networks of pathogenic fungi reveals targets for antifungal drugs

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The burden of fungal infections for humans, animals and plants is widely underestimated and comprises deadly infections as well as great economic costs. Despite that, antifungal drugs are scarce and emergence of resistance in fungal strains contributes to a high mortality. To overcome this shortage, we propose toxic intermediates and their controlling enzymes in metabolic pathways as a resource for new targets and provide a web-service, *FunTox-Networks* to explore the landscape of toxic intermediates in the metabolic networks of fungal pathogens. The toxicity of metabolites is predicted by a new random forest regression model and is available for over one hundred fungal species. Further, for major fungal pathogens, metabolic networks from the KEGG database were enriched with data of toxicity and regulatory effort for each enzyme to support identification of targets. We determined several toxic intermediates in fungal-specific pathways like amino acid synthesis, nitrogen and sulfur assimilation, and the glyoxylate bypass. For the latter, we show experimentally that growth of the pathogen *Candida albicans* is inhibited when the detoxifying enzymes Mls1 and Hbr2 are deleted and toxic glyoxylate accumulates in the cell. Thus, toxic pathway intermediates and their controlling enzymes represent an untapped resource of antifungal targets.

1 | INTRODUCTION

Fungi represent important human pathogens which cause a wide range of diseases from harmless superficial infections of skin and nails, affecting roughly a quarter of the worldwide population [1], to life-threatening infections with millions of patients per year. Despite lower incidence rates globally of invasive fungal infections than e.g. tuberculosis, Brown *et al.* [2] estimates a comparable number of deaths (around 1.5 millions) due to the very high mortality rates of invasive fungal infections. For example, in France the total burden of serious fungal infections is estimated to affect more than 1% of the population each year [3] and the current COVID-19 pandemic is accompanied by life-threatening secondary fungal infections, predominately by SARS-CoV-2-associated pulmonary aspergillosis (CAPA) [4] or mucormycosis [5].

The increasing incidence of invasive fungal infections is linked to advances in medical care that can save lives of immunocompromised patients, which however are at high risk of fungal infections [6, 7]. While most immunocompetent hosts are able to clear or control fungal infections, immunocompromised patients can suffer from blood-borne disseminating infections with often very high mortality rates of more than 50 % [2]. However, there is only a very limited number of antifungal agents. This is mostly due to the much closer phylogenetic relationship between fungi and humans compared to bacterial pathogens, which constrains the options for agents that do not harm the human host [8]. Furthermore, resistance to the widely used azole antifungals is increasingly reported [9] and polyenes antifungals are only cautiously used due to their toxicity to mammalian cells [9]. Due to these difficulties, novel approaches for speeding up the discovery of antifungal drugs and their targets are urgently needed [10, 11].

The metabolic flexibility of pathogenic fungi is a cornerstone of their virulence [12] and, additionally, enzymes of fungal-specific pathways in central metabolism such as ergosterol biosynthesis are key targets for antifungal drugs [10, 11, 13]. Hence, the metabolism of pathogenic fungi has come under scrutiny to establish novel antifungal targets and develop highly efficient new antimycotics [14, 15]. Since the metabolism of fungi is highly dynamic during host interactions [12, 16], modeling of metabolic regulation by dynamic optimization is of high relevance to unveil optimal regulatory strategies in fungal pathways as well as to elucidate key enzymes regulating pathway flux [17]. In recent years, we and others have uncovered optimality principles behind fast pathway activation strategies and efficient pathway regulation across a wide range of bacteria [18–24], which are in principle transferable to eukaryotes such as fungi, too. To combat pathogens, our previous observation that enzymes upstream of toxic intermediates are tightly regulated is of special interest, since we hypothesized that an upregulation (downregulation) of key enzymes regulating flux before (after) a toxic intermediate can lead to self-poisoning [23]. Further, as shown across prokaryotes [23], the inference of metabolic hubs and key regulated enzymes by estimators like the promoter length can be used in a reverse approach to identify highly regulated enzymes and to support target identification for potential antimicrobials. This reasoning provides a new avenue for the identification of drug targets using endogenously produced cytotoxins. While metabolic networks have recently been investigated extensively to find key enzymes for virulence using the concept of elementary modes [25], there has been, to our knowledge, no systematic screen for toxic intermediates in metabolic networks of fungal pathogens so far. However, the importance of toxic intermediates has been recognized in the fields of pathway evolution [26] and metabolic engineering [27]. Similarly, an anti-cancer therapy using endogenous toxic metabolites to stop cell growth was recently proposed [28].

For a large-scale prediction of toxicity across multiple organisms, quantitative structure - activity relationship (QSAR) models have proven to be valuable in *in silico* screening of inhibitors [29, 30], in improving yield by identification of toxic intermediates [31, 32] and in risk assessment [33, 34]. These machine learning approaches can help to fill the gap between empirical measured toxicity data in databases like ChEMBL [35] and pathway intermediates which have unknown toxicity. For fungi, Prado-Prado *et al.* [36] developed a multi-target spectral moment QSAR to classify drugs with activity against several species. However, to take advantage of novel machine learning approaches as well as

much larger toxicity assays stored in the ChEMBL-database, we here used a random forest regression model to predict the toxicity of intermediates in fungal metabolism based on chemical features. Using a multi-output model [37, 38], we were able to include data from all fungi with toxicity measurements as well as different types of measurement of toxicity to train a single regression model for different fungal species and additional models for host organisms. Moreover, we integrated our predictions with data on the degree of regulation of individual enzymes in metabolic networks derived from KEGG [39]. To this end intra-species protein-protein interactions networks were inferred using the newly developed FungiWeb database (<https://fungiweb.bioapps.biozentrum.uni-wuerzburg.de>) and used here to overcome the lack of regulatory data like large-scale transcription factor networks in fungal species. The enriched networks together with a cytotoxicity prediction can be displayed in a web-service *FunTox-Networks* (<http://funtox.bioinf.uni-jena.de>).

As proof-of-principle for the use of our tool to identify targets for the enrichment of antifungal intermediates, we used *FunTox-Networks* to identify several toxic intermediates in fungal-specific pathways. As promising targets, we identified toxic intermediates and their controlling enzymes in pathways of nitrogen and sulfur assimilation, in amino acid synthesis, and in the catabolism of fatty acids as alternative carbon sources. Especially for the latter, where glyoxylate is a toxic intermediate, we show that detoxifying enzymes like malate synthase (Mls1) and an aminotransferase (Hbr2) are potential targets for novel antifungal drugs.

2 | RESULTS

2.1 | Prediction of metabolite toxicity with a random forest regression model

Based on the toxicity assay data in the ChEMBL database, we trained separate QSAR regression models for the prediction of metabolite toxicity in fungal species and their human as well as murine host (see section 4.1 for details). After carefully filtering wrong annotations and removal of outliers the data set for fungal species included 122,474 data points for 112 fungal species, 653,035 data points for human cells and 10,703 data points for mice (see section 4.1 for details). Due to the best performance among other regression models on a subset of the data, the random forest regression approach was chosen to build the QSAR models (see Supp. 6). The final random forest regression model for fungal species obtained a reasonable coefficient of determination of $R^2 = 0.64$, which is slightly lower for the models of human ($R^2 = 0.57$) and murine ($R^2 = 0.59$) cells. The model quality is underlined by low root mean squared error (RMSE) values between 0.30 (fungi), 0.30 (human), and 0.32 (mice). For our fungal model we fulfill the recommended criteria [40] of a high $R^2 > 0.6$ and a low RMSE, which should be lower than 10 % of the range of predicted toxicity values (between -1.73 and 4.59 in training data). Other QSAR toxicity regression models achieve higher predictive power but lack universality by considering only a single species like *Escherichia coli* [32] or by focusing on a certain group of substances with even more precise predictions [41].

While our random forest regression models show good performances on test and training data, we further used annotated compounds from the KEGG database to validate the predictions of our models. To this end, we predicted the toxicity of compounds with biological roles like antibiotics or carbohydrates in important fungal species and host organisms (see Fig. 1A). As expected, antifungal compounds were predicted to be significantly more toxic than non-antibiotic compounds in fungal species (Wilcoxon rank-sum test $P < 1^{-16}$). The regression model clearly discriminates antifungals also from other antibiotics like antibacterials (Wilcoxon $P < 1^{-16}$). One should note that toxicity of antibiotics is relatively high even in hosts, which is reasonable since many antibiotics cause side effects harming human cells and therefore are often not applied in high concentrations [42, 43]. Our model predicts carbohydrates as the least toxic compound group in fungi and other organism which is in line with their common biological functions and their high

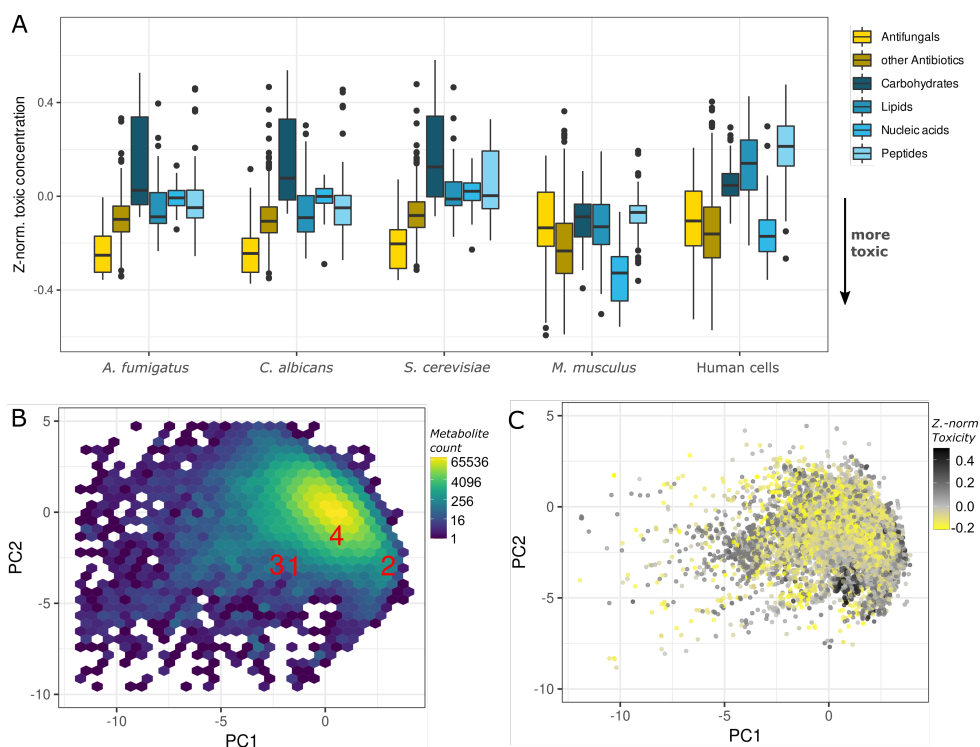


FIGURE 1 **A** Toxicity prediction of KEGG compounds and their categories in fungi and host organisms. Distribution of toxic concentration are depicted in yellow for antibiotics and in blue for common metabolites. **B** Applicability Domain of the QSAR model based on a PCA of molecular descriptors (descriptor space) with the exemplary metabolites (red): ATP (1), glyoxylate (2), amphotericin B (3), fluconazole (4). The density of learning data in descriptor space is depicted from dark blue to yellow. **C** Distribution of metabolites from KEGG based in the same descriptor space as in A and toxicity in *C. albicans* indicated in gradient from yellow (toxic) to black (non-toxic).

abundance in standard growth media and diet. From these results we conclude that our QSAR model is able to estimate the toxicity of metabolites across a wide range of compound classes and organisms.

A main goal behind our QSAR model is that it is accessible for all researchers and we therefore provide the web-service *FunTox-Networks*. In addition to the toxicity prediction of KEGG metabolites, the web-service can be queried to predict toxicity for other or novel compounds via their Simplified Molecular Input Line Entry Specification (SMILES) or International Chemical Identifier (InChI) representation [44, 45]. Furthermore, we provide, as recommended by the Organisation for Economic Co-operation and Development (OECD) [46], information about the confidence of predictions as an applicability domain of our random forest regression model. Firstly, we provide the standard deviation across predictions of all trees as additional output, which has been shown to reflect the accuracy of predictions [47]. Secondly, a principal component analysis of trained metabolites and their molecular descriptor values was performed to visualize the descriptor space of training data (see Fig. 1B). Using this presentation we see that training data covers not only the known toxic antifungals like fluconazole or amphotericin B well, but is also nearly identical to the descriptor space obtained from all metabolites of pathways in KEGG (see Fig. 1C).

Importantly, although the toxicity prediction is mainly based on molecular descriptors, we did not observe distinct

regions or clusters in the principal component analysis of molecular descriptors for toxic and non-toxic metabolites. This observation indicates that toxic metabolites are uniformly distributed across the entire chemical space (see Fig. 1C). Moreover, this demonstrates that our random forest regression model is able to recognize small structural differences that are associated with toxicity. An example is the well known toxic byproduct methylglyoxal ($C_3H_4O_2$) formed during glycolysis [48]. Our model correctly predicts a high toxicity of methylglyoxal e.g. in the common fungal pathogen *Candida albicans* ($Z_{tox} = -0.177$) and no toxicity of the structurally highly similar metabolite pyruvate ($C_3H_4O_3$, $Z_{tox} = 0.007$).

2.2 | Integration of intermediate toxicity and enzyme regulation in pathway maps

Since we demonstrated a close relationship between toxicity of pathway intermediates and optimal points of pathway regulation in our previous work [23], we integrated metabolite toxicity and regulatory effort on KEGG pathway maps to identify suitable targets of deregulation to accumulate self-poisoning intermediates in pathogenic fungi (see section 4.2 and Fig. 5B for details). Thus, we provide enriched KEGG pathway maps of seven major fungal pathogens or model organisms and their hosts (human cells and mice, see Tab. 2). To infer key regulated enzymes with toxic intermediates, we used the number of transcription factors controlling an enzyme and, for pathogenic fungi, a score representing the connectivity of an enzyme within its intraspecies protein-protein interaction (PPI) network. In those networks each interaction is further characterized by its confidence, e.g. experimental validation (see section 4.2). Interestingly, yeast data show that this measure correlates with the number of interactions as well as with the number of post-translational modification (PTM) sites of a gene ($\rho = 0.52$ and $\rho = 0.24$, respectively; Spearman correlation with each $P < 10^{-16}$, see also Supp. 8). Since PPI networks have not been reconstructed for all fungi, we used the promoter length as easily calculable (intergenic distance) estimator to infer transcriptional regulation in the case of *Arthroderma benhamiae* and *Aspergillus flavus* (see section 4.2).

In addition to enriched KEGG pathway maps, *FunTox-Networks* provides tables containing all pathways and compounds that can be searched and sorted by key features. These overview tables also show that very diverse compound groups within the KEGG pathway maps are predicted to be highly toxic. Examples in *C. albicans* include heme-like compounds (siroheme, precorrin), activated fatty acids (acyl-CoA) and highly reactive acids like acetic acid, which all are known to be cytotoxic [49–51]. These data further emphasize the predictive quality of our model. Examples of KEGG pathways which contain many toxic intermediates are fatty acid metabolism (map01040, map00062, map00071), propanoate metabolism (map00640), glyoxylate metabolism (map00630), and porphyrin metabolism (map00860).

Because we can predict toxicity in fungal species as well as for their potential hosts, our web-service additionally provides an interactive plot to compare the toxicity of KEGG metabolites in two species. This enables the identification of metabolites which are more toxic to the pathogen than to host cells or *vice versa* (see Fig. 2). While the former metabolites are interesting antifungals that are tolerable by the host, the latter can comprise potential small-molecule virulence factors of pathogens. Due to the advent of large-scale transcriptomics able to generate measurements during host-pathogen interaction (e.g. dual-species RNA-Seq of host and pathogen in parallel), this plot of species-specific toxicity can be enriched with previously published expression data for genes coding for the adjacent enzymes [16, 52, 53].

Fig. 2 depicts how this view of data can be used to get insights into molecular host-pathogen interactions. Firstly, a distinct group of acyl-CoA compounds is predicted to be toxic in the pathogens *C. albicans* (Fig. 2A) and *C. glabrata* (Fig. 2B) as well as in the human host. However, the adjacent enzyme coding genes, which are mainly related to fatty-acid oxidation, are upregulated in *C. glabrata*, but not in *C. albicans* after 1h of co-incubation with human blood. In contrast, genes encoding the enzymes of the glyoxylate bypass are strongly upregulated in *C. albicans* and not in *C. glabrata*. Further, toxicity of glyoxylate is primarily predicted in fungal cells and not in host cells. Interestingly, despite the

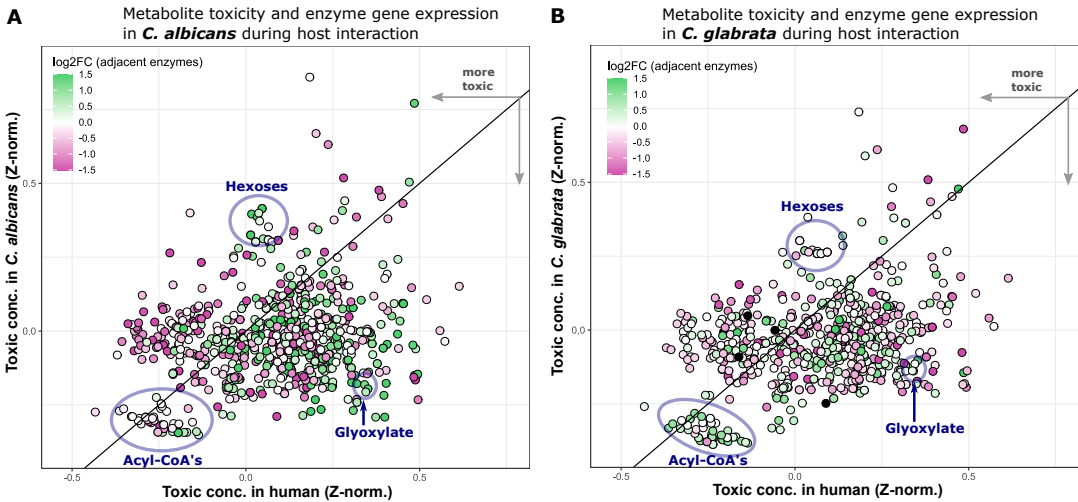


FIGURE 2 Integration of enzyme gene expression and metabolite toxicity in pathogenic fungi upon interaction with human blood cells [16]. Each dot represents the predicted toxicity of KEGG metabolites in the human host (x-axis) and in the fungal pathogen *C. albicans* (subpanel A) or *C. glabrata* (subpanel B). The color shows the up- or downregulation (green to pink) of adjacent enzyme coding genes (producing or consuming the metabolite) after 1h incubation with human blood for each *Candida* species as measured by Kämmer et al. [16].

upregulation of genes encoding the enzymes of the glyoxylate bypass in *C. albicans*, which indicates glucose starvation, a group of non-toxic hexoses are connected to upregulated enzyme coding genes or transporter genes in sugar metabolism (see Fig. 2A).

To summarize, our toxicity prediction and web-service *FunTox-Networks* offers not only the prediction of toxicity for pathway intermediates, but also a detailed integration with the pathway topology in KEGG maps, regulation of enzymes, respectively the expression of their genes during host-pathogen interactions. This allows to identify new potential targets for antifungals by searching for toxic pathway intermediates and the identification of key regulated enzymes controlling the intermediate's accumulation.

2.3 | Toxic intermediates in fungal specific metabolic pathways

To prove its usefulness, we employed *FunTox-Networks* to search for enzymes which are suitable targets for antifungal interventions. We limited our search to metabolic pathways which are specific to fungi and enzymes which have no homolog in humans. Since fungal species are able to grow in a great variety of conditions, many fungal specific metabolic pathways with toxic intermediates are related to resource acquisition, such as carbon, nitrogen, and sulfur assimilation (see Fig. 3). We found known and new targets in the synthesis of amino acids and in fatty acid metabolism, as well as in ergosterol synthesis, which is a main target for the antifungal class of azoles.

Ergosterol synthesis is especially suitable for antifungal intervention in our sense, as azoles are not only inhibiting the synthesis by blocking the Lanosterol-14 α -demethylase, but also lead to the accumulation of toxic 14-methyl sterol intermediates like 14 α -methyl-3,6-diol [54].

To harbor suitable antifungal targets a pathway needs to be active during host invasion and is ideally essential for virulence. Because pathogenic fungi have different strategies to invade the host and conquer different host niches like

inhibition of Mls1 since the toxic intermediate glyoxylate is accumulated.

2.4 | The glyoxylate bypass as primary example for toxic intermediates and drug target search

To verify the potential of toxic intermediates as antifungal drug targets, we experimentally investigated the glyoxylate bypass in *C. albicans*. In addition to the two key enzymes Icl1 and Mls1, the genome annotation as well as metabolic databases like KEGG suggest two additional enzymes which can detoxify glyoxylate (see Fig. 4A). We therefore analyzed the transcriptional response of *C. albicans* to various organic acids [67]. In addition to *ICL1* or *MLS1*, we created a deletion strain lacking the putative alanine glyoxylate aminotransferase gene (*HBR2*), because it shows a strong coregulation with the glyoxylate bypass enzymes (see Fig. 4B) and its homolog in baker's yeast codes for an enzyme that is able to detoxify glyoxylate to pyruvate [68]. In contrast to this, the glyoxylate reductase gene (*GOR1*) is coregulated with the isocitrate dehydrogenase gene *IDP2* of the TCA cycle (see Fig. 4A and B). Hence, we excluded Gor1 as candidate for the control of glyoxylate accumulation when glyoxylate bypass enzymes are active.

To study the importance of glyoxylate and their detoxifying enzymes, we performed growth experiments in glucose-rich medium at a low pH to ensure membrane-permeability of externally supplemented glyoxylate (cf. section 4.3). While externally added glyoxylate leads only to minor growth inhibition of the wild-type *C. albicans* strain, single deletion strains of *MLS1* or *HBR2* show intermediate, and a double deletion mutant (*mls1Δ/Δ hbr2Δ/Δ*) a significant growth inhibition (see Fig. 4C). This supports the view that glyoxylate detoxification is ensured by both, Mls1 and Hbr2.

C. albicans faces a glucose-poor environment during phagocytosis and relies on the glyoxylate bypass for survival. Having shown that *C. albicans* depends on Mls1 and Hbr2 for glyoxylate detoxification, we performed additional experiments where glyoxylate is produced intracellularly. To this end, we tested growth with ethanol as the only carbon source. As expected, the wild-type and *hbr2Δ/Δ* strains grew slowly on ethanol, and the glyoxylate bypass mutants were unable to grow. In addition to growth inhibition, we observed that strains lacking the glyoxylate detoxification enzymes (*mls1Δ/Δ* and *mls1Δ/Δ hbr2Δ/Δ*) die off earlier when ethanol is the sole carbon source than the non-glyoxylate producing strain *ic1Δ/Δ*, which otherwise is similarly unable to grow (see Fig. 4D).

Lastly, we confirmed glyoxylate accumulation in our mutant strains by gas chromatography-mass spectrometry (GC-MS) after growth in glucose-rich medium. We found that glyoxylate accumulated in *mls1Δ/Δ* and reached even higher concentrations in the double mutant *mls1Δ/Δ hbr2Δ/Δ*. This is accompanied by the enrichment of the TCA cycle intermediates citrate and malate. Interestingly, in the single mutant *mls1Δ/Δ* glycine accumulated which supports our hypothesis that glyoxylate is detoxified by Hbr2 via the conversion of alanine to glycine (see Fig. 4A). From the above experiments we can conclude that glyoxylate accumulation cannot be achieved in *C. albicans* by inhibition of only the malate synthase, as it has been shown for *M. tuberculosis* [66]. Moreover, we discovered that the so-far uncharacterized aminotransferase Hbr2 is part of the more complex glyoxylate detoxification in *C. albicans*, which provides new avenues for drug target search. Overall, we show that toxic intermediates are an untapped resource for the development of antifungal drugs by the characterization of enzymes as well as the underlying transcriptional regulation, which control accumulation of pathway intermediates during host-pathogen interactions.

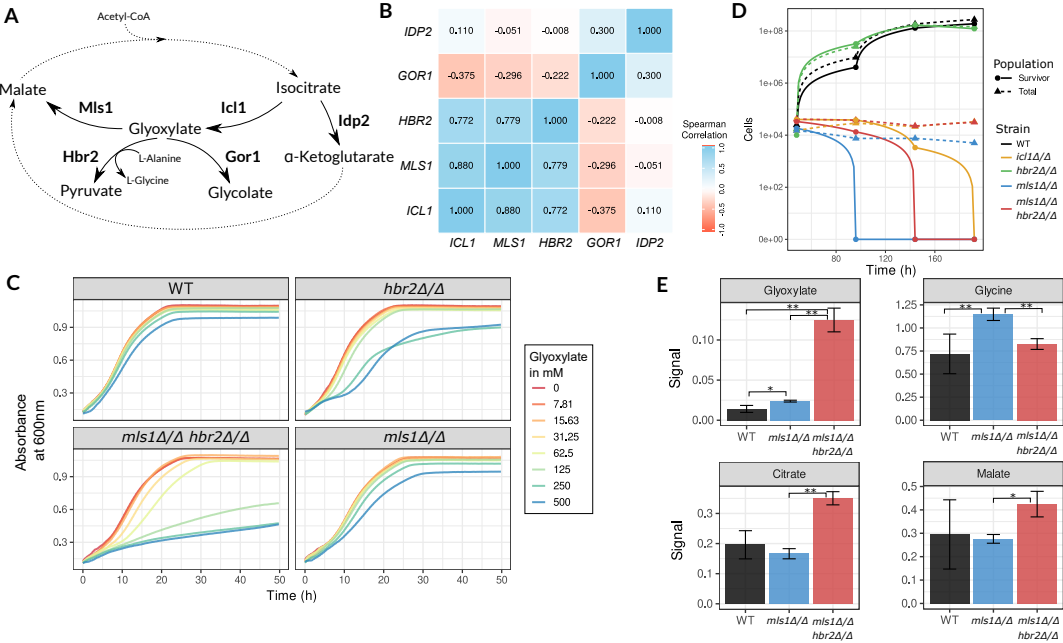


FIGURE 4 Investigation of glyoxylate detoxifying enzymes in *C. albicans*. **A** Overview of glyoxylate bypass as well as related metabolites and enzymes. **B** Correlation matrix of transcription for glyoxylate bypass genes (*ICL1*, *MLS1*), genes for potential detoxifying enzymes (*HBR2*, *GOR1*) and for the decarboxylation of isocitrate (*IDP2*) in response to various organic acids [67]. **C** Growth curve of *C. albicans* and deletion strains in glucose (2%) media with addition of varying amounts of glyoxylate. The mean of the absorbance at 600nm of three biological replicates is shown for each glyoxylate concentration. **D** Long term survival assays of *C. albicans* strains in SD medium and ethanol as sole carbon source. Surviving yeast cells were determined on YPD agar plates after the indicated incubation times. **E** Measurement of metabolites by gas chromatography–mass spectrometry (GC-MS) of *C. albicans* strains grown in YNB broth without amino acids and 2% glucose. Error bars indicate the standard errors of mean (SEM) for three replicates and significance (T-tests) is shown for selected comparisons between strains with values * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$. In all figures WT refers to the wild-type strain BWP17+Clp30.

3 | CONCLUSION AND OUTLOOK

In summary, we were able to build a new QSAR model for the prediction of metabolite toxicity for fungal species and also for their hosts during infection. Based on a specialized normalization scheme, our random forest regression models show a good performance in predicting the toxicity of different compound categories such as toxic antifungals or non-toxic carbohydrates. Since it was our goal to support antifungal drug target identification, we interleaved metabolite toxicity prediction with metabolic pathway maps for main fungal pathogens and their hosts and provide it as the web-service *FunTox-Networks*. To test our approach with a real-life application, we used this new resource and identified multiple potential toxic intermediates in fungal-specific metabolic pathways. We found a very promising target in the malate synthase and the aminotransferase Hbr2 of *C. albicans* which synergistically control the detoxification of glyoxylate. This can become important when fatty acids are used as a carbon source by *C. albicans* e.g. during the confrontation with phagocytic immune cells [69]. In our experiments we observed a growth defect of deletion strains related to the accumulation of toxic glyoxylate. Therefore we conclude that antifungal drugs inhibiting malate synthase and the

aminotransferase Hbr2 would be more effective than inhibitors of the isocitrate lyase which are currently being tested [70]. Moreover, it reveals that a comprehensive study of the regulatory circuits controlling pathway intermediate accumulation provides an untapped resource for the discovery of novel types of antifungals.

While we focused here on fungal pathogens of humans, fungi that infect plants and animals are a significant concern in agriculture and biodiversity, leading not only to high economic costs but also affecting human health, since they can cause crop failures and famine [71]. The principle of toxic intermediates as guides to valuable antifungal drug targets can also be applied to fungal plant pathogens. Toxicity prediction via *FunTox-Networks* can therefore become a valuable resource for research and drug development in this field, too. Beyond the use for antifungal target selection, the prediction of intermediate toxicity can also be of advantage in industrial processes involving yeasts, which try to optimize yield and efficiency by reducing the accumulation of toxic intermediates [72]. Especially in metabolic engineering or synthetic biology, where foreign or redesigned metabolic pathways are incorporated into yeast, the knowledge of potential toxic intermediates and their control can be of great importance [73]. Interestingly, a rewiring of the glyoxylate bypass regulation was necessary to improve the titer reached in production of glycolic acid which is an important chemical compound and can be produced in yeasts by glyoxylate reduction [74, 75]. However, production is hampered by the tight repression of the isocitrate lyase when glucose is present and glyoxylate accumulation is avoided rigorously which underlines the importance of our findings.

Taken together, knowledge about toxicity of pathway intermediates as well as their regulation enables various applications in antifungal drug target search or industrial use of fungal species.

4 | MODEL AND METHODS

4.1 | Toxicity prediction

To train the regression model for predicting the toxicity of metabolites in fungi as well as their hosts (humans, mice), we retrieved the corresponding taxonomy-based activity data listed in the ChEMBL database [35] (Release 27, January 2021). The three data sets for fungi, human and mice were then filtered and checked for the correct taxonomy, target type (only 'organism' or 'cell' and not 'protein inhibition' etc.), non-valid data, only precise toxicity measurements ('=' relation) and uniform direction of toxicity standard types (lower value indicating higher toxicity). To learn the regression model, we used a cut-off of at least 50 activity data measurements for each standard type (type of toxicity measure) and at least 100 measurements for each organism. Using these filtering steps, we obtained 122,474 activity measurements for fungi and 653,035 for humans and as well as 10,703 for mice, respectively, which were used as input for the machine learning approach (see Fig. 5). This machine learning pipeline and data is also documented and stored in a open repository (<http://doi.org/10.5281/zenodo.3529162>).

To extract structural features to train a QSAR, we retrieved the Simplified Molecular Input Line Entry Specification (SMILES) [44] for all compounds from the above described data set. Using the python package Mordred [76], we calculated molecular descriptors of several properties such as size, polarity and topology (see Tab. 1). To rule out that descriptor choice weakens prediction performance, multiple sets of descriptors with minimal correlation to each other were tested. However, prediction performance was similar based on this set and we therefore conclude, that our selection of descriptors is sufficient to represent the chemical space.

In contrast to other QSAR models, we built a multi-output model to train the model on all fungal species as well as on different toxicity measurements (standard types) using a normalization scheme as described previously [37, 38]. Hence, the activity data (tox) is standardized by Z-normalization with the mean (avg) and standard deviation (sd) of all

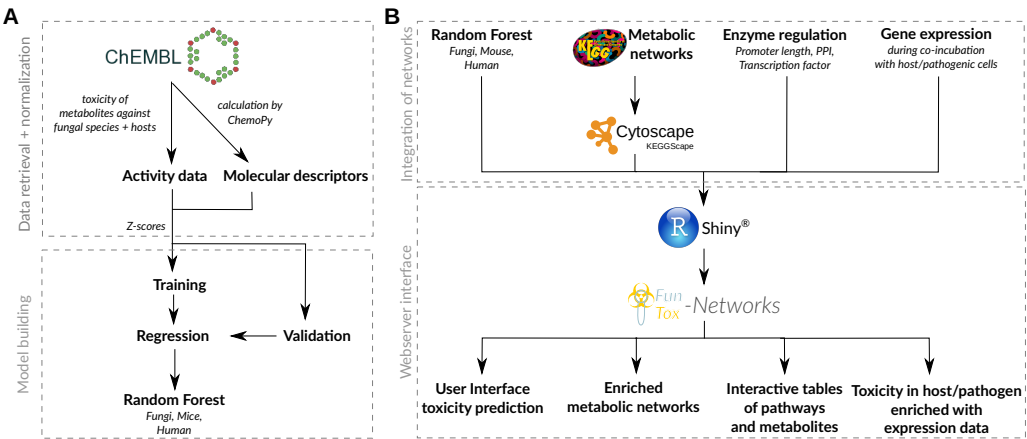


FIGURE 5 **A** Schematic workflow of data preparation and model building to predict metabolite toxicity in fungi, mice and human. **B** Overview of used databases, data and tools to integrate toxicity prediction and metabolic networks and the provided interfaces in the web-service *FunTox-Networks*.

TABLE 1 Overview of used molecular descriptors.

Molecular Descriptor (<i>m</i>)	Category
Number of rings (nRing)	Constitutional descriptor
Balaban index (BalabanJ)	Topological descriptor
Number of multiple bonds (nBondsM)	Molecular property
Number of hydrogen bond acceptors (nHBAcc)	Molecular property
Partition coefficient (SLogP)	Molecular property
Wiener Index (WPath)	Molecular property
Topological polarity surface area (TopoPSA)	Molecular property
Molecular weight (MW)	Constitutional descriptor

activity data under the same standard type (s):

$$Z_{tox} = \frac{tox_s - avg_s}{sd_s} \quad (1)$$

The normalized activity data (Z_{tox}) is estimated through a regression by structural features (molecular descriptors), which are Z-normalized for fields (f) describing the influence of assay characteristics (organism f_o , standard type f_s , cell type f_c and data curation level f_d). Together with the average Z-normalized toxicity for the combination of organism as well as standard type (Z_{exp}) and the eight molecular descriptors (see Tab. 1) this results in 33 features describing the relationship of toxicity, assay characteristics and molecular structure:

$$Z_{tox} = Z_{exp} + \underbrace{Z_{md}(f_o, m_R) + \dots + Z_{md}(f_d, m_R)}_{\text{Rings}} + \dots + \underbrace{Z_{md}(f_o, m_{MW}) + \dots + Z_{md}(f_d, m_{MW})}_{\text{Molecular weight}} \quad (2)$$

After removal of outliers (e.g. extreme values in ChEMBL for (non-)toxic compounds) based on a simple linear model (upper and lower 5% residuals), a 10 % randomly chosen subset of the resulting data was used to identify the best method for regression by applying the rRegr package [77] comparing several machine learning approaches (see Supp. 6). The decision tree-based random forest regression method [78] performed best in terms of deviation and correlation and therefore random forests were trained on the full data set using the R package 'ranger' [79]. As optimal parameters for the machine learning approach, we used five randomly selected features at each split ($mtry$) and one hundred trees ($ntree$), which has proven to be a good trade-off between performance and computational costs in previous machine learning approaches [80] and in for our model (cf. our analysis documented in <http://doi.org/10.5281/zenodo.3529162>).

Furthermore, cross-validation was performed by splitting the data set in 4 splits with each run using 75% of the full data set as training and 25% as test data set to calculate the cross-validated R^2_{CV} (or Q^2) [81]. Since values of R^2_{CV} were close to the coefficient of determination R^2 , we did not observe overfitting of our model for all splits (cf. our analysis documented in <http://doi.org/10.5281/zenodo.3529162>) and using all features described in equation 2. Therefore all features were used to train the final random forest models for fungal species, human and mice on the full data set.

For QSAR models it is advised to provide information about the certainty of a prediction, called applicability domain [46]. The robustness of toxicity prediction can be inferred by analyzing the chemical space of compounds used for learning and by the confidence of predictions made by the random forest regression. To this end, the standard deviation across all trees of the random forest, which is an indicator of prediction accuracy [47], is given as additional output. Furthermore, we performed a principle component analysis (PCA) of the molecular descriptor space of all compounds used to train the random forest regression to visualize similarity of queried to trained compounds [82]. Both is provided in the web-service of our toxicity prediction.

4.2 | Integration of intermediate toxicity and metabolic networks

For the identification of possible drug targets in metabolic networks, pathways of the major fungal pathogens and their hosts were enriched with the predicted toxicity of metabolites and regulatory data of enzymes (see Tab. 2). Pathway maps were retrieved using the R package KEGGREST in the KEGG Markup Language (KGML) format [39, 83] for the organisms listed in Tab. 2. Toxicity of KEGG compounds was predicted with our random forest regression model with MIC (minimal inhibitory concentration) in fungal species as standard type and IC50 (half maximal inhibitory concentration) for human cells respectively LD50 (median lethal dose) for mice, since these were the preferred standard

types in the data set for training.

TABLE 2 Organisms with toxicity prediction, metabolic networks from KEGG and estimated regulatory effort of enzymes.

Organism	KEGG ID	Description	Regulatory data
<i>Arthroderma benhamiae</i>	abe	Dermatophyte	Promoter length
<i>Aspergillus fumigatus</i>	afm	Saprophyte, (invasive) aspergillosis	PPI score
<i>Aspergillus flavus</i>	afv	Saprophyte and crop pathogen (invasive) aspergillosis	Promoter length
<i>Candida albicans</i>	cal	Commensal, (invasive) candidiasis	PPI score
<i>Candida glabrata</i>	cgr	Commensal, (invasive) candidiasis	PPI score
<i>Cryptococcus neoformans</i>	cnb	Saprophyte, (invasive) cryptococcosis	PPI score
<i>Homo sapiens</i>	hsa	Host	Transcription factor
<i>Mus musculus</i>	mmu	Host	Transcription factor
<i>Saccharomyces cerevisiae</i>	sce	Important fungal model organism rare opportunistic pathogen	PPI score

We used three different approaches to infer the strength of regulation of individual enzymes referred to as regulatory effort controlling an enzyme. Intuitively, the number of transcription factors controlling an enzyme is a direct measurement of regulatory effort and is used for human and murine enzymes, where genome-wide data is available from the RegNetwork database [84].

For fungi with only scarce knowledge of gene regulatory interactions, we used information that were obtained from the FungiWeb-Database (<https://fungiweb.bioapps.biozentrum.uni-wuerzburg.de>). Since no experimentally validated large-scale protein-protein networks on a genome-scale for these species exists we used an interolog-based method to infer these networks from the established and validated human and yeast protein-protein networks (see Remmele et al. 2015 [85]). For this approach we obtained the intraspecies networks from *Homo sapiens* and *Saccharomyces cerevisiae* from the 14 active partners of the International Molecular Exchange (IMEx) consortium [86] and the protein orthology information from Inparanoid8 [87], *Candida* Genome Database (CGD) [88] and *Aspergillus* Genome Database (AspGD) [89] via an automated pipeline. Based on the experimental evidence, size of the experiment and number of publication that support a source interaction we calculated for each edge of these networks a modified version of the MINT-Score (Molecular INteraction database [90]) as a measurement of the interaction reliability. From the PPI networks of *A. fumigatus* (176,584 interactions, 4,086 interactors), *C. albicans* (97,614 interactions, 1,984 interactors), *C. glabrata* (297,419 interactions, 4,604 interactors) and *C. neoformans* (191,274 interactions, 3,203 interactors) we estimated the regulatory effort of an enzyme by the connectivity (vertex degree) in the network of the species and weighted each edge by their evidence according to the reliability score. The resulting PPI score provides a good estimate of regulation and interaction on post-translational level, which we could confirm for yeast (see supplement 8).

For organisms where neither transcription factors nor a PPI score are available (see Tab. 2), the promoter length determined as the intergenic distance was used as in previous studies for prokaryotes [21, 23]. Promoter lengths have proven to be good proxies of regulatory effort due to the reduced genome sizes of fungi compared to higher eukaryotes

which makes a loss of non-functional sequences more likely [91]. In consequence, the length of promoter regions should reflect the number of functional elements such as transcription factor binding sites contained in them. In agreement with this hypothesis, we found for the well-studied regulatory network of yeast that promoter length correlates with the number of transcription factors (see Supp. 8).

We enriched pathways maps with color-coded information on regulatory effort and toxicity information in KGML files. We used Cytoscape [92] with the extension KEGGScape [93] to convert KGML formats to JavaScript Object Notation (JSON), which can be used in a javascript-based network viewer provided by Cytoscape. The collection of pathway maps is embedded in the R Shiny [94] application *FunTox-Networks* and accessible as a web-service (<http://funtox.bioinf.uni-jena.de>). Moreover, pathway maps and compounds are summarized in tables that can be searched, filtered and downloaded to facilitate the search of toxic intermediates and the controlling enzymes to identify drug targets.

4.3 | Investigation of glyoxylate detoxifying enzymes in *C. albicans*

Identification and co-regulation of enzymes

We used metabolic information of *Saccharomyces cerevisiae* from MetaCyc [95] to identify additional enzymes which degrade glyoxylate. In addition to the malate synthase (EC: 2.3.3.9) this includes the glyoxylate reductase (EC: 1.1.1.79) and the alanine-glyoxylate transaminase (EC: 2.6.1.44). Based on sequence homology to yeast counterparts we determined GOR1 (orf19.2989), respectively HBR2 (orf19.1078) and calculated the Spearman's rank correlation of their gene expression in a published data set of the response in *C. albicans*' transcription to organic acids [67].

Strains and mutant generation

All *C. albicans* strains were stored as glycerol stocks at -80°C and streaked on YPD plates for growth at 30°C before use. Mutants were generated by standard heat shock transformation procedures (PCR-amplified pFA plasmids with $\approx 100\text{bp}$ homology regions [96]) using the BWP17 strain (lacking *URA3*, *HIS1* and *ARG4* [97]). Uridine prototrophy was restored with Clp10, and BWP17 + Clp30 (restored for all three markers) was used as an isogenic control. For additional deletion of HBR2, the dominant SAT flipper method was employed as described previously [98], using pSFS5 and the In-Fusion cloning system (Takara Biotech) for creation of a deletion cassette with *SacI* and *KpnI* restriction sites. The final mutant was cured of the SAT1 cassette and contained only the FRT site in place of HBR2. All primers are listed in supplementary table 3, and all mutants are listed in supplementary table 4.

Growth assays and inhibition by glyoxylate

The *C. albicans* deletion mutants (*mls1Δ/Δ*, *hbr2Δ/Δ*, *mls1Δ/Δ hbr2Δ/Δ*) or wild type (SC5314) were grown over night in yeast extract peptone dextrose (YPD) medium at 30°C , 180 rpm and then washed three times with distilled water. For the assay, $10\mu\text{L}$ of the cell suspension in phosphate-buffered saline (PBS) was mixed (in 96 well plates) with $190\mu\text{L}$ buffered Synthetic Defined (SD) medium (1×YNB, w/o amino acids/ammonium sulfate; 5 g/L ammonium sulfate; 2 g/L glucose; 100mM phosphate buffer, $\text{pH} = 3$). The medium also contained glyoxylate, in a 1:2 dilution series from 500mM down to 7.8mM , plus a control without any glyoxylate. The strains were grown in triplicates in each glyoxylate concentration for 50 hours at 30°C in a Tecan Infinite 200 microplate reader. The absorption at 600nm was measured every 15 minutes after 10s of orbital shaking.

Long term survival assays

The *C. albicans* deletion mutants (*icl1Δ/Δ*, *mls1Δ/Δ*, *hbr2Δ/Δ*, *mls1Δ/Δ hbr2Δ/Δ*) or wild type (SC5314) were grown for long periods in SD medium without carbon source or with 2g/L of a C2 compound (ethanol). For this, overnight YPD cultures of *C. albicans* (30°C, 180 rpm) were washed three times with PBS. The cells were counted in a Neubauer chamber, and 20mL of each SD medium was inoculated at a concentration of 10⁵ cells/mL. The strains were grown at 30°C for 8 days, and samples (1mL) taken every 48 hours for cell number quantification with a Neubauer chamber. A serial 1:10 dilution (down to 10⁶ cells/mL) in PBS was used to assess the number of living cells in the sample: Three 10μL spots of each dilutions were pipetted on YPD agar plates and incubated for 48 hours at 30°C for counting of living cells. The plates were then incubated for further 48 hours at 30°C to allow growth of smaller colonies from lagging cells, which were added to the count.

Measurement of glyoxylate in *Candida albicans* strains

Overnight pre-cultures of *C. albicans* wild type (SC5314) and deletion mutants *mls1Δ/Δ* and *mls1Δ/Δ hbr2Δ/Δ* were used to inoculate the corresponding main cultures with an OD600 of 0.1. After a cultivation over a period of 12h in 50mL YNB broth without amino acids and 2% glucose as carbon source at 30°C, samples were harvested in triplicates and intracellular metabolites extracted. For this purpose, the cells were centrifuged at 4°C and 4000g for 5min, followed by three washing steps with 20mL cold 0.9% NaCl solution. The pelleted cells were resuspended in ethanol supplemented with 2.5μg/mL [U-¹³C₅]-ribitol or [2,2,3,3,4,4-¹³C₆]-glutaric acid as internal standard. Resuspended cells were set out by ultrasonic treatment at 70°C for 15min for cell lysis. The suspensions were incubated for 2min on ice followed by the addition of 0.75mL H₂O. Metabolites were extracted by addition of 1mL chloroform followed by harshly mixing for 1min and centrifugation for 5min at 4000g and 4°C. 0.8mL of the resulting upper polar phase was sampled and triplicates were dried in a vacuum concentrator over night at 4°C. Polar dried metabolites were automatically derivatized in a two step procedure; first, 15μL of 2% methoxyamine hydrochloride in pyridine was added and samples were incubated at 55°C for 90min under shaking. Second, MTBSTFA (N-methyl-N[tert]-butyldimethylsilyl trifluoroacetamide w/1% tert-butyldimethylchlorosilane) was added in an equal amount and samples were incubated for 60min at 55°C. Gas-chromatography coupled to mass-spectrometry (GC-MS) measurements were performed using an Agilent 7890B GC / Agilent MSD 5977B instrument. Derivatized metabolites were separated by a capillary column (Phenomenex ZB 35, 30m length, 0.25mm in diameter, 0.25μm film thickness) and a SSL Liner Agilent 5190-3171: 900 μL (splitless, single taper, wool, Ultra). Carrier gas was Helium with a flow rate of 1.0mL/min. The temperature in the GC oven was 2min at 100°C with increased steps of 10°C per minute up to 300°C. Total run time of 26min per sample. Electron impact ionization at 70eV were operated by an Agilent MSD 5977B with an extractor source and the transferline temperature at 280°C. The mass spectrometry was held at 230°C and the quadrupole at 150°C. The detection of metabolites were done in scan and sim mode. To acquire the full mass spectra scanning from 70 $\frac{m}{z}$ to 800 $\frac{m}{z}$ at a scan rate of 4 $\frac{scans}{s}$. SIM data for glyoxylate was performed with the *mz* of 160, 161, 162, 202, 203, 204 each a dwell time of 15ms. Data was processed with the MetaboliteDetector software. Statistical analyses were performed. Standard errors of mean (SEM) were shown. Students T-tests were performed and p values are calculated with two way analysis. Significance values were considered as significant with values *p < 0.05, **p < 0.01, ***p < 0.001.

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SUPPORTING INFORMATION

Data and scripts of machine learning pipeline are documented and stored here: <http://doi.org/10.5281/zenodo.3529162>.

Details of machine learning procedure

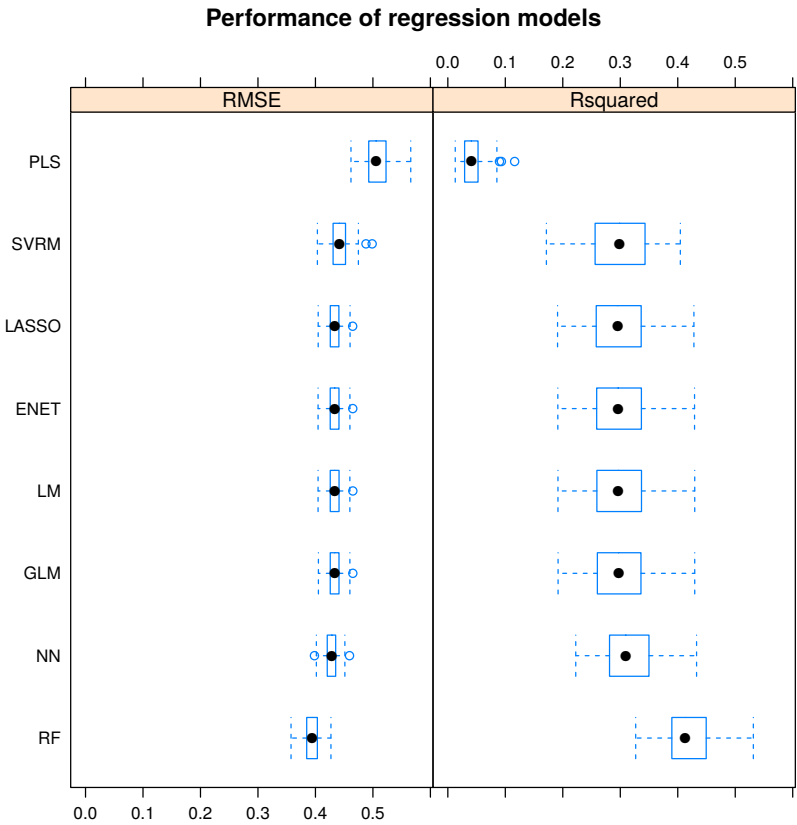


FIGURE 6 Training results of different regression models measured by the root mean squared error (RMSE) and the coefficient of determination (R^2). Models: Multiple Linear regression (LM), Generalized Linear Model with Stepwise Feature Selection (GLM), Partial Least Squares Regression (PLS), Lasso regression (LASSO), Elastic Net regression (ENET), Support vector machine using radial functions (SVRM), Neural Networks regression (NN), Random Forest (RF).

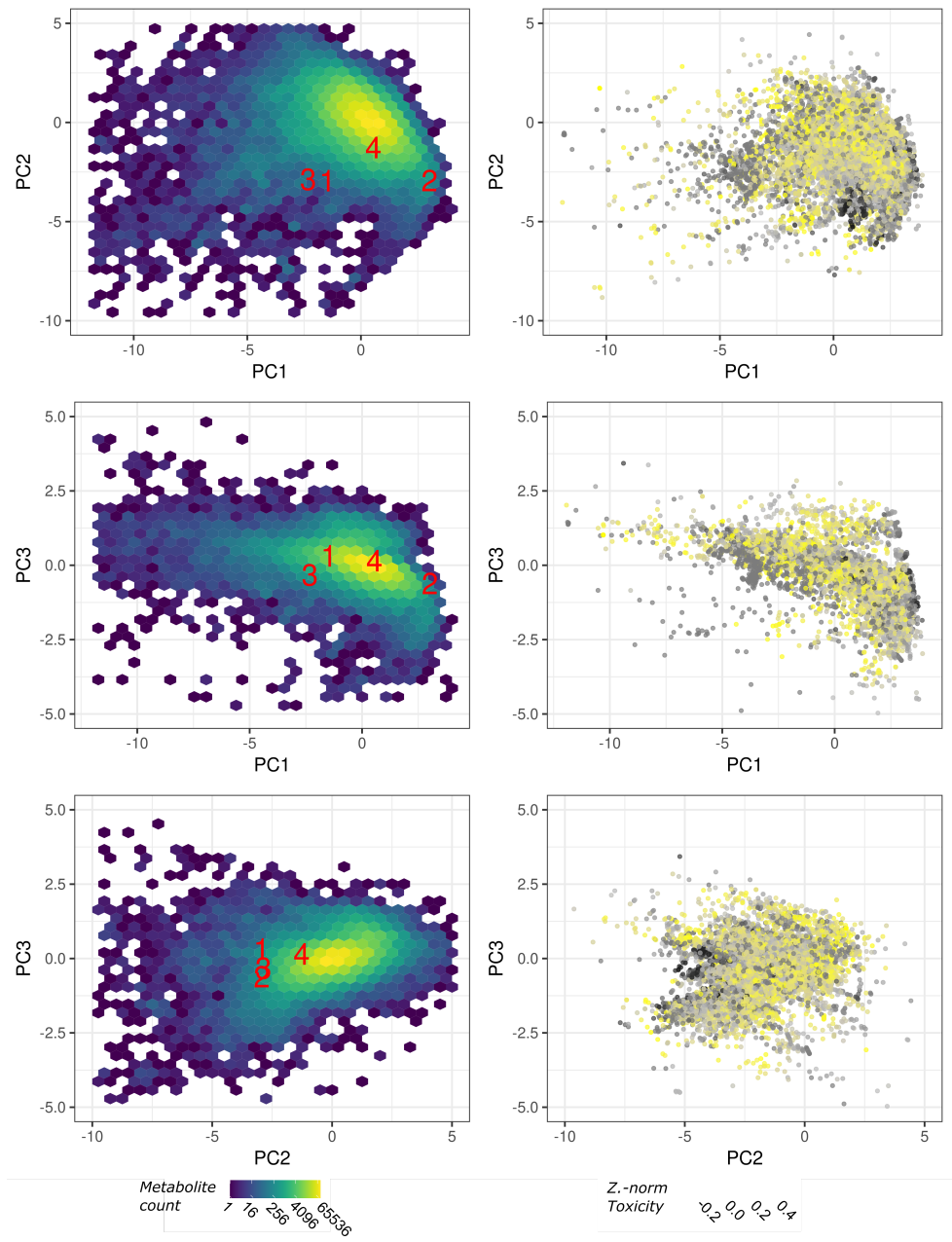


FIGURE 7 Additional plots including third principle component for applicability domain (left) and toxicity distribution (right) of metabolites as shown in

Regulatory effort estimates in yeast

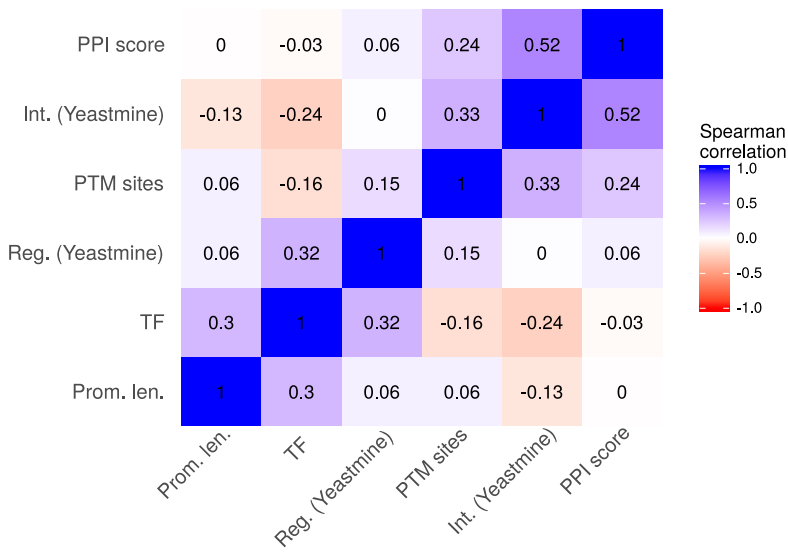


FIGURE 8 Correlation of promoter length, number of transcription factors (TF) from Yeastract [99], number of regulators as well as interactions listed in YeastMine [100], number of PTM sites from dbPTM [101] and the PPI score for each gene in yeast.

Primer and mutant list

TABLE 3 List of primers

Primer name	Sequence
ICL1-fwd	TTTTTAATACCCTTTTTCTTTTTCTTTTTCTTCCTCTATACTTATACCTTT TATTCTAATATAAATTAAAGAATAAACATTAATAATATCTACCGAAGCTTCGTA CGCTGCAGGTC
ICL1-rev	TTACTAGACAGATCATCTCAAATGAATACCGTCTTTTGTTTTTGTATATTAT AAGTTCTGTTTCTTTACTAATTTACACTTCTATCCCTCAAATTATCTGATATCA TCGATGAATTCGAG
MLS1-fwd	TAAGTTTGAATTCTCTTTTCCTTTTTCTTATTATTTTACTTTTACATTTATATATA TAAATATTACACACAGCTTTGTATATATATTAACCAAGTTACATAGAAGCTTCGT ACGCTGCAGGTC
MLS1-rev	ATATTTTCATGAATAAACATAAACAAACTAATAAAAAAAGCTACTTTCATACT ATTTAAATTACAAATTGAAAACGTTTCCCGAACATTCTTTTTTTATCTGATATC ATCGATGAATTCGAG
HBR2-upstream-fwd	TATAGGGCGAATTGGAGCTCCAAGTGGTAGTGGTGGTGGT
HBR2-upstream-rev	CCGCCACCGCGGTGGTCAAAAGAATAAAAAATAAAAAGTAA
HBR2-downstream-fwd	TCGAGGGGGGGCCCGCAATTTAGTAATTGAATTTAGGTC
HBR2-downstream-rev	GGGAACAAAAGCTGGGTACCGTGTGGCATCCTATTCGGTC

TABLE 4 List of strains and mutants

Identifier	Name	Parent	Genotype	
C55	SC5314	-	(Wild type)	
M2251	BWP17+Clp30	BWP17	<i>URA3::imm434/URA3::imm434</i>	<i>HIS1::hisG/HIS1::hisG</i> <i>ARG4::hisG/ARG4::hisG</i> <i>RPS1::(URA3 HIS1 ARG4)/RPS1</i>
M2577	<i>icl1Δ/Δ</i>	BWP17	<i>URA3::imm434/URA3::imm434</i>	<i>HIS1::hisG/HIS1::hisG</i> <i>ARG4::hisG/ARG4::hisG</i> <i>ICL1::HIS1/ICL1::ARG4</i> <i>RPS1::URA3/RPS1</i>
M2582	<i>mls1Δ/Δ</i>	BWP17	<i>URA3::imm434/URA3::imm434</i>	<i>HIS1::hisG/HIS1::hisG</i> <i>ARG4::hisG/ARG4::hisG</i> <i>MLS1::HIS1/MLS1::ARG4</i> <i>RPS1::URA3/RPS1</i>
M2692	<i>hbr2Δ/Δ</i>	BWP17+Clp30	<i>URA3::imm434/URA3::imm434</i>	<i>HIS1::hisG/HIS1::hisG</i> <i>ARG4::hisG/ARG4::hisG</i> <i>HBR2::FRT/HBR2::FRT</i> <i>RPS1::(URA3 HIS1 ARG4)/RPS1</i>
M2696	<i>mls1Δ/Δ/hbr2Δ/Δ</i>	M2582	<i>URA3::imm434/URA3::imm434</i>	<i>HIS1::hisG/HIS1::hisG</i> <i>ARG4::hisG/ARG4::hisG</i> <i>MLS1::HIS1/MLS1::ARG4</i> <i>HBR2::FRT/HBR2::FRT</i>