Looking Back to the Future: Predicting in Vivo Efficacy of Small Molecules versus Mycobacterium tuberculosis

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ABSTRACT: Selecting and translating in vitro leads for a disease into molecules with in vivo activity in an animal model of the disease is a challenge that takes considerable time and money. As an example, recent years have seen whole-cell phenotypic screens of millions of compounds yielding over 1500 inhibitors of Mycobacterium tuberculosis (Mtb). These must be prioritized for testing in the mouse in vivo assay for Mtb infection, a validated model utilized to select compounds for further testing. We demonstrate learning from in vivo active and inactive compounds using machine learning classification models (Bayesian, support vector machines, and recursive partitioning) consisting of 773 compounds. The Bayesian model predicted 8 out of 11 additional in vivo actives not included in the model as an external test set. Curation of 70 years of Mtb data can therefore provide statistically robust computational models to focus resources on in vivo active small molecule antituberculars. This highlights a cost-effective predictor for in vivo testing elsewhere in other diseases.

INTRODUCTION

Drug discovery involves a considerable effort in the selection and translation of in vitro leads into molecules with in vivo efficacy in an animal model of the disease. Our collective memories are often short when it comes to decades of research in a single therapeutic area, let alone research of over more than a half-century, and yet the past may hold many insights to aid us in drug discovery efforts in both the present and future. As an example, from the 1940s to the 1960s, significant efforts led to first- and second-line drugs for Mycobacterium tuberculosis (Mtb), which is the causative agent of tuberculosis (TB). This disease has infected approximately 2 billion people and kills 1.3 million people annually. We critically need next-generation active small molecules as tools to query essential infection biology to drive novel therapies. Chemical probes can enable interrogation of Mtb pathways essential to in vivo infection. Next generation drugs must lack cross resistance to current therapeutics, shorten treatment, and address drug-drug interactions with co-administered treatments. Many molecules have been assessed as to their ability to modulate Mtb infection in mice. These data reside in numerous journals and reports that are not readily accessible despite today’s electronic media and databases. Unfortunately, much of the early (pre-1970s) structure-activity relationship (SAR) data from both in vitro and in vivo models appears to have been neglected. Typically, these data are only unearthed on a compound-by-compound basis when we rediscover an agent that was already known from decades ago.

The field’s interest in this historical data may have been unfortunately diminished due to the recent upsurge in whole-cell phenotypic high-throughput screens (HTS) for novel antituberculars that have seen several million compounds tested. Commercial vendor and in-house libraries have been assayed, leading to the clinical candidate SQ-109 and a diarylquinoline hit that was optimized to the drug bedaquiline.

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line.\textsuperscript{19} However, these successful outcomes from screening represent anomalies as the road from hit to drug invokes words such as “valley of death”.\textsuperscript{20} Mtb HTS hit rates are usually below 1%,\textsuperscript{11,21} and most hits require a significant amount of chemical evolution in an effort to identify a probe let alone a drug discovery lead. To date we estimate about 1500 in vitro Mtb hits of interest have been derived from one laboratory alone,\textsuperscript{13–15,22} while GSK has recently published another 177 promising in vitro actives.\textsuperscript{12} Many laboratories have also described Mtb hits; so in total, there are likely close to 2000 compounds from recent efforts that may require triage before advancement of the most promising through the discovery pipeline.

To significantly impact the TB field, these hits, or their evolved analogs, ideally must demonstrate in vivo activity in an animal model. Rarely is the specific Mtb target for these compounds known before in vivo testing. The mouse model of TB infection is considered important for comparative assessment of different treatments and optimization of TB drug dosing schedules.\textsuperscript{23} There has also been considerable development of the acute\textsuperscript{24} and chronic mouse models\textsuperscript{25} of Mtb infection. Given current resource limitations that are magnified for a neglected disease such as TB, we are faced with a dilemma: How do we efficiently select among the thousands of hits to decide which to carry forward for in vivo efficacy assessment?

This report details a novel approach to address this critical issue through the curation of 773 molecules that have been tested in the last 70 years in the mouse TB infection model. This in vivo data has never before been curated. We present detailed analyses of the physiochemical and structural properties of both active and inactive molecules as well their chemical property space coverage in an effort to guide the future design of novel antitubercular chemical probes and drugs. Furthermore, we leverage our experience with machine learning models for in vitro activity\textsuperscript{8,22,26–29} to construct and statistically validate these computational models to predict in vivo efficacy in the Mtb-infected mouse model. The computational models are further validated through the correct prediction of 8 of 11 known in vivo actives absent from the training set. The models are also applied to score a set of 177 in vitro drug leads recently reported by GlaxoSmithKline (GSK)\textsuperscript{12} to aid in their prioritization for in vivo assessment.

\section*{EXPERIMENTAL SECTION}

Molecule: Curation, Drawing, Quality Assessment, and Storage. Various search terms were used in PubMed to retrieve papers with compounds tested in murine acute and chronic Mtb infection models. For example, “tuberculosis and in vivo mouse”, “Tuberculosis and efficacy and mouse”, and “comparison and antituberculosis and mouse”. The same search terms were also used in SciFinder (CAS, Columbus, OH) and Web of Knowledge (Thomson Reuters). Individual journals were also searched online (e.g., Tuberculosis, Journal of Medicinal Chemistry, and PLOS journals). The In vivo data was manually curated, and structures were sketched using the Mobile Molecular DataSheet (MMDS) iOS app,\textsuperscript{30} ChemDraw (Perkin-Elmer, Waltham MA), or downloaded from ChemSpider (www.chemspider.com) and combined with pertinent data fields. The data has been made publically available in the CDD TB database (Collaborative Drug Discovery, Inc., Burlingame, CA).\textsuperscript{31} Molecules were classed as active/inactive, and this was generally based on the data in the publications. For example, a reduction of log CFU in lung greater or equal to 1 log was considered active. The initial assembled data set was shown to contain duplicates by utilizing the ACD/ChemFolder version 12 software program (Advanced Chemistry Development, http://www.acdlabs.com/products/km/ackm/chemfolder/). Utilizing the ability to check for duplicates and incorrect structures (valence errors, pentavalent carbons, missing stereochemistry), identified structure issues were manually curated. A total of 18 compounds were either removed from the originally assembled data set or edited to deal with the identified errors.

Molecular Property Distribution. AlogP, molecular weight, number of rotatable bonds, number of rings, number of aromatic rings, number of hydrogen bond acceptors, number of hydrogen bond donors, and molecular fractional polar surface area were calculated from input SD files using Discovery Studio 3.5 (San Diego, CA).

Principal Components Analysis with in Vitro Hits and TB Mobile Data. We compared the 773 compounds with the previously described 745 compounds with known Mtb targets collated from the literature\textsuperscript{27} and available in TB Mobile (version 1)\textsuperscript{32} that were utilized to generate a principal components analysis (PCA) plot with the interpretable descriptors selected previously (AlogP, molecular weight, number of rotatable bonds, number of rings, number of aromatic rings, number of hydrogen bond acceptors, number of hydrogen bond donors, and molecular fractional polar surface area) for machine learning. This PCA model represents essentially the published target-chemistry property space for Mtb. We also compared 1429 Mtb hits (active and nontoxic only, from the NIH screens where IC\textsubscript{90} < 10 μg/mL or 10 μM and a selectivity index (SI) greater than 10 where the SI is calculated from SI = CC\textsubscript{50}/IC\textsubscript{90}) to show how they covered the target-chemistry property space. These analyses can be compared with those previously published which focused on in vitro Mtb data.\textsuperscript{29}

Building and Validating Machine Learning Models with Mouse Mtb in Vivo Data. We have previously described the generation and validation of the Laplacian-corrected Bayesian classifier models developed from Mtb growth inhibition screens of small molecule libraries\textsuperscript{8,28} using Discovery Studio 3.5\textsuperscript{33–37} This approach was utilized with the literature data curated in the course of this study. The following molecular descriptors were used and were calculated from input SD files: molecular function class fingerprints of maximum diameter 6 (FCFP\_6),\textsuperscript{38} AlogP, molecular weight, number of rotatable bonds, number of rings, number of aromatic rings, number of hydrogen bond acceptors, number of hydrogen bond donors, and molecular fractional polar surface area. Models were validated using leave-one-out cross validation in which each sample was left out one at a time, a model was built using the remaining samples, and that model was utilized to predict the left-out sample. Each model was internally validated, ROC plots were generated, and the XV ROC AUC calculated. The Bayesian model was additionally evaluated by leaving out 50% of the data and rebuilding the model 100 times using a custom protocol for validation to generate the ROC AUC, concordance, specificity, and selectivity as described previously.\textsuperscript{8,28} The internal ROC value represents the training set value while the external ROC represents the test set molecules left out. We also compared the resulting Bayesian model with SVM and RP Forest and single tree models built with the same molecular descriptors in Discovery Studio. For SVM models, we calculated interpretable descriptors in Discovery Studio and then used Pipeline Pilot to generate the FCFP\_6 descriptors followed by integration with R.\textsuperscript{39} RP Forest and RP Single Tree models used the standard protocol in Discovery Studio. In the case of RP Forest models, 10
trees were created with bagging. Bagging is short for “Bootstrap AGgregation”. For each tree, a bootstrap sample of the original data is taken, and this sample is used to grow the tree. A bootstrap sample is a data set of the same size as the original one but in which the same data record can be included multiple times. RP Single Trees had a minimum of 10 samples per node and a maximum tree depth of 20. In all cases, five-fold cross validation (leave out 20% of the database five times) was used to calculate the ROC for the models generated.

Model Predictions for Additional Compounds Identified after Model Building. Eleven compounds active in the mouse in vivo model were identified from a 1950s compilation. Of these, only seven were not in the 773 compounds training set. A further four active compounds resulted in 11 compounds that were predicted with the computational models developed (Table 4). For each molecule, the closest distance to the training set was also calculated using the Bayesian model in the calculated properties protocol method in Discovery Studio (a value of zero represents a molecule in the training set, while larger values are further from the training set).

Predictions for GSK Compounds. 177 Mtb leads were recently disclosed by GSK and represent a promising set of small molecules for further exploration as potential antitubercular drug candidates. The GSK set was scored with all of the in vivo models generated in this study. The mean closest distance to the training set was also calculated for the 177 compounds to provide an idea of similarity to the training set. These data were calculated from the outputs of each of the Bayesian models. For each test set molecule a score for closest distance to training set was calculated using Discovery Studio (described earlier). We averaged this number across the 177 molecules, where the “closeness” of a compound to the training set scales inversely with the value. The maximum Tanimoto similarity for each molecule versus the training set was also calculated using MDL fingerprints. Consensus predicted active compounds were identified across all four machine learning models. These compounds were then evaluated using TB Mobile and clustering in Discovery Studio with the 745 compounds in this data set to infer potential targets. ADME/Tox properties for these compounds were generated using Discovery Studio ADMET predictors and custom Bayesian models for PXR, hERG, etc.

Scaffold Analysis Using SAR Table. Scaffold analysis was performed using the SAR Table app (for iOS-based devices such as iPhones and iPads), which provides a user interface for drawing scaffolds and substituents and specifying activity data. It also provides access to analysis functionality such as scaffold-substructure matching, structure—activity model generation, data visualization, and manuscript figure creation.

Statistical Analysis. Means for descriptor values for active and inactive compounds were compared by two-tailed t-tests with JMP v. 8.0.1 (SAS Institute, Cary, NC).

### RESULTS

Data Set Curation, Molecular Property Analysis, and Data Visualization. A total of 773 molecules were collated from the literature for the first time to our knowledge using various search terms in PubMed, SciFinder, and Web of Knowledge (see Experimental Section), for which there was data in acute or chronic mouse models of Mtb infection (Table S1, Supporting Information). Following convention, an “active” compound minimally exhibited a 1 log_{10} reduction in Mtb colony-forming units (CFUs) in the lungs as compared to no-drug control. Occasionally, other types of analyses required our best scientific judgment on the active/inactive boundary; for example, the work of Denny and co-workers used an activity ratio versus PA-824. We considered a value >0.4 as active given the demonstrated 2.5–3 log_{10} reduction in CFUs in the lungs by PA-824 (27–32). Older publications relied on extension of survival compared to negative and positive controls. Foruitously for machine learning model construction, the data set was divided almost equally between actives (N = 362) and inactives (N = 411). Through analysis of simple molecular descriptors (see Experimental Section) we sought to gain insight as to why active and inactive compounds behaved differently (Table S2 and Figure S1, Supporting Information). Among the 773 compounds, statistically significant differences were found between the number of rings, their AlogP, and fractional polar surface area (Table 1). It should be noted that the standard deviations are large for the in vivo data and may stem from the heterogeneity of structures (e.g., between very small molecules, large macrolide antibiotics, and calixarenes) and likely published experimental methods (e.g., different mice strains, dosing concentrations, dosing period, etc.). We would not advise using individual properties alone (like calculated logP) to differentiate in vivo active compounds as there were temporospatial effects for the descriptors, suggesting the addition of further compounds over time increases or decreases differences observed (Figure S1, Supporting Information).

PCA can be used to understand multi-dimensional data represented by the multiple molecular descriptors representing the molecular properties of the training set and shows overlap in this property space between active and inactive compounds (Figure 1A). Our analyses can also be compared to those previously performed with Mtb in vitro data. This overlap would also suggest some complexity in using these individual molecular descriptors alone to distinguish in vivo active compounds. Approved drugs for TB are distributed in this same chemical property space with much larger intravenous TB drugs (representing generally larger molecules with different molecular properties (less hydrophobic)) separated out of the main cluster (Figure 1B).

PCA using molecules known to inhibit specific Mtb targets suggests that the in vivo compounds possess good coverage of known target chemistry property space and extend well outside of it (Figure 2A). However, it should be noted these targets do
not broadly cover the chemistry property space of the \textit{Mtb} metabolome (using molecules extracted from BioCyc)\textsuperscript{51,52} (Figure 2B), and the active \textit{in vivo} compounds also only cover a fraction of the \textit{Mtb} metabolome PCA plot (Figure 2C).

When the same \textit{in vivo} compounds are compared to known \textit{Mtb} \textit{in vitro} actives\textsuperscript{13–15,22} they are also well distributed over the PCA plot (Figure 3A). The majority of the \textit{in vivo} actives overlap with the \textit{in vitro} actives\textsuperscript{13–15,22} suggesting a good coverage of the chemistry property space (and likely requirement for similar molecular features or descriptors), with a small number of the \textit{in vivo} actives exploring property space distinct from that of the \textit{Mtb} \textit{in vitro} actives from recently described screening efforts. This graphical tool can demonstrate differential coverage of chemical property space and may be utilized to explore how candidate molecules compare to molecules previously assayed \textit{in vivo}.

A significant component of the data set (51 of 773 molecules; Table S1, Supporting Information) contains the triazine central scaffold. These compounds were initially brought to our
attention when we identified TCMDC-125802 ((E)-6-(2-((5-nitrofuran-2-yl)methylene)hydrazinyl)-N,N'-diphenyl-1,3,5-triazine-2,4-diamine, Figure 4A; MIC of 62.5 ng/mL against Mtb) through our machine learning models for in vitro antitubercular activity. A literature search highlighted one report of the antitubercular activity of TCMDC-125802 and related
triazines from 1969.53 Figure 4 shows a matrix correlation plot of triazine substituents $R_1$ versus $R_2$ for this published data set based on Mtb in vitro and in vivo data (Figure 4A). The data visualization quickly hones in on the features important at $R_1$ and $R_2$ (i-propylamino, and n-propylamino groups) in cells that are green (Figure 4B). This type of approach could be useful for interrogating other structure–activity relationships within the complete in vivo data set.

**Machine Learning Models for Mtb in vivo Data.** Machine learning models are educated by example and provide an excellent opportunity to discern actives from inactives by both their physiochemical and structural properties.54 They have an additional benefit of enabling prediction of additional sets of small molecules with a significant degree of accuracy.8,22,28,29 Previously, we have reported the validation of support vector machine (SVM), recursive partitioning (RP) Single Tree, and RP Forest models to compare with Bayesian models of in vitro antitubercular efficacy with acceptable (selectivity index >10) Vero cell cytotoxicity.29 These types of models (Bayesian, SVM, and RP) are commonly used for drug discovery applications in virtual screening and balance fitting the training set data with external predictive capability outside of the training set’s chemical property space. Such approaches have been described by us in some detail previously,29,55 and the machine learning with the Mtb in vivo data paralleled our practices with previous Mtb in vitro data sets.8,22,28,29 We utilized FCFP_6 fingerprints55,56 and the following set of readily interpretable molecular descriptors: ALogP,57 molecular weight, number of H-bond
donors, number of H-bond acceptors, number of rings, number of aromatic rings, number of rotatable bonds, and molecular fractional polar surface area (FPSA). The Bayesian model statistics for 773 molecules, generated by leaving out 50% of the data and rebuilding the model 100 times using a custom protocol for validation to produce the cross-validated receiver operator curve area under the curve (XV ROC AUC), concordance, specificity, and sensitivity as described previously, are shown in Table 2. The leave-out 50% × 100 external ROC score was 0.72, while the concordance (66.91), specificity (74.23), and sensitivity (58.46) suggested a bias toward predicting inactive compounds. The 773 molecule Bayesian model provides almost

Table 2. Mean (± sd) Leave-One-Out and Leave-Out 50% × 100 Cross Validation of Bayesian Models

| leave-one-out ROC | leave-out 50% × 100 external ROC score | leave-out 50% × 100 internal ROC score | leave-out 50% × 100 concordance | leave-out 50% × 100 specificity | leave-out 50% × 100 sensitivity |
|-------------------|---------------------------------------|----------------------------------------|-------------------------------|--------------------------------|--------------------------------|
| 0.77              | 0.72 ± 0.02                           | 0.74 ± 0.02                            | 66.91 ± 2.24                  | 74.23 ± 8.96                  | 58.46 ± 9.19                  |

*ROC = receiver operator characteristic. Best split −2.195.
Table 3. Individual Machine Learning Model Cross Validation Receiver Operator Curve Statistics for 773 Molecules Tested in the Mouse in Vivo Model for Mtbr

| Model                  | FP Forest (out of bag ROC) | RP Single Tree (five-fold cross validation ROC) | SVM (with five-fold cross validation ROC) | Bayesian (with five-fold cross validation ROC) |
|------------------------|----------------------------|-----------------------------------------------|-----------------------------------------|------------------------------------------------|
| ROC                    | 0.75                        | 0.71                                          | 0.77                                    | 0.73                                           |

Bayesian five-fold cross validation has sensitivity = 66.3%, specificity = 90.3%, and concordance = 79.0%.

DISCUSSION

A recent review has described a timeline for the introduction of the antitubercular drugs. Several of these date back to the 1940s and 1950s and span up to the late 1960s. Only recently (2012) has a new drug, namely, bedaquiline, been approved. Although there are several drugs in clinical trials, the pipeline is relatively thin for a disease where drug resistance has a significant impact and extensively drug resistant Mtbc is present in nearly 60 countries. The last 10–15 years have witnessed an upswing in high-throughput screening in an attempt to identify molecules that modulate perceived essential targets or from phenotypic screening in whole cells. The result has been the discovery of about 2000 in vitro Mtbc hits and perhaps 100s of promising leads. The next hurdle is likely the selection of appropriate compounds to test in the mouse in vivo model of infection. The history of this model itself dates back to the 1940s, and even though it has limitations in extrapolating to humans, it is the only animal model that has been validated with human subjects in guiding TB drug development. The mouse represents an expensive medium-throughput model and a bottleneck in screening when used for rank-ordering compounds. On the basis of a recent literature analysis of publications over a 12-year period, there was a five-fold increase in use of the TB mouse model from 1997 to 2009. Our own analysis looks at a much longer time period, collecting data from over 70 years and illustrates that between 1970 to 2000 there was a gap in the publication of mouse in vivo data (Figure S1, Supporting Information), with just 55 compounds retrieved in this period out of the total 773. We see no change in this previously reported increased utilization of the mouse model based on the large number of papers describing in vivo data for approximately 200 compounds from the last four years (Table S1, Supporting Information), and it is therefore imperative that we question the current workflow and ask how greater cost- and time-efficiencies can be achieved. These data may also suggest some degree of publication bias toward actives.
While we need to better understand how in vitro efficacy, absorption, metabolism, distribution and excretion (ADME), and in vivo pharmacokinetic profiles are linked to in vivo antitubercular activity, we also need to computationally learn from the collective experiences of the considerable number of generally small molecules (773 in this study alone) that have been tested in the mouse model. These data serve as the training set for an educated computational model to prioritize hits for in vivo testing, minimizing the assessment of compounds likely to fail. Several computational models are available to predict in vitro ADME properties in addition to other physicochemical properties. While we and others have utilized machine learning models to predict Mtb activity in vitro, the studies to date have not analyzed Mtb in vivo data in mice. However, mathematical modeling of pathogen and host interactions pertinent to latent infection have been reported. One reason for this lack of modeling Mtb in vivo is perhaps because the achievement of in vivo efficacy is multi-factorial due to complex interactions between Mtb, host, and drug. Additionally, the data to conduct this in vivo modeling exercise had not been previously curated until now.

The curation of this in vivo data, as with all literature data capture, must be conducted with attention paid to structure deposition errors. For example, after testing in mice, it was suggested that 27753-RP was a highly promising antituberculosis drug. Subsequently, in the absence of supportive data, it was stated that griselimycin(e) while e was not de as the process, we noted that the griselimycin structure does not capture, must be conducted with attention paid to structure.

### Table 4. Test Set of in Vivo Active Compounds Not in the TB in Vivo Models

| name (number or abbreviation relates to original nomenclature) | Forest | Single Tree | Bayesian score | Bayesian class | closest distance | SVM | ref |
|---------------------------------------------------------------|--------|------------|----------------|----------------|-----------------|-----|-----|
| 1070 = anisaldehyde, thiosemicarbazone | 0 | 0 | −3.02 | 0 | 0.35 | 0 | 40 |
| 1493 = 1-(p-methoxybenzyl)-3-thiosemicarbazide | 0 | 0 | −1.19 | 1 | 0.46 | 0 | 40 |
| 2403 = p-nitrobenzaldehyde, thiosemicarbazone | 0 | 0 | −3.70 | 0 | 0.44 | 0 | 40 |
| 2406 = N-threo-[α,α-dichloro-N-[β-hydroxy-(hydroxymethyl)-p-nitrophenethyl]acetamide(chloromycetin) | 0 | 0 | 1.54 | 1 | 0.53 | 1 | 40 |
| 2875 = nicotinamide | 0 | 1 | −1.06 | 1 | 0.40 | 1 | 40 |
| viomycin | 0 | 1 | 10.95 | 1 | 0.27 | 1 | 40 |
| neomycin | 1 | 0 | 10.11 | 1 | 0.01 | 1 | 40 |
| PClH = 2-pyrdylcarboxaldehyde p-nitrobenzoyl hydratine | 1 | 1 | −1.53 | 1 | 0.39 | 1 | 41 |
| Cpd 3 = N-(2-fluoroethyl)-1-((6-methoxy-5-methylpyrimidin-4-yl)-methyl)-1H-pyrrole[3,2-b]pyrdine-3-carboxamide | 1 | 1 | −0.44 | 1 | 0.56 | 1 | 42 |
| Cpd 4 = N-cyclopropylmethyl)-1-((6-methoxy-5-methylpyrimidin-4-yl)-methyl)-1H-pyrrole[3,2-b]pyrdine-3-carboxamide | 0 | 0 | −2.06 | 1 | 0.53 | 0 | 42 |
| indoleamide 3 | 0 | 0 | −6.20 | 0 | 0.31 | 1 | 58 |

*Prediction scores 1 = active, 0 = inactive.*

Our retrospective analysis of the data for small molecules (monotherapy only) in Mtb-infected mice has been critical to the realization of what physicochemical properties and chemical features best describe the actives (Figure S2, Supporting Information) as well as inactives (Figure S3, Supporting Information). This analysis is important because many promising in vitro active compounds do not show in vivo activity.8 The ring count was significantly higher in in vivo active compounds, and the AlogP and fractional polar surface area were lower than in inactives. Intriguingly, our analysis of Mtb in vitro HTS results has shown actives to have a higher calculated logP than inactives. We cannot also discount possible differences between methods for predicting logP. The reliance on a single molecular descriptor may be suboptimal. The curated in vivo data was also analyzed using multiple interpretable descriptors and machine learning models in this study. All four of the methods we have described resulted in similar receiver operator curve (ROC) values (0.71–0.77), indicative of potentially useful models. These models could score small molecules (absent in the training set) to prioritize antitubercular hits and leads for crucial in vivo studies. We evaluated the models with 11 known in vivo actives and found the Bayesian model outperformed the SVM and tree methods (Table 4). The Bayesian model correctly identified 8 out of 11 small molecules, absent from the training set, as in vivo actives. Compounds like the recently published indoleamide 3 (Table 4, Figure S6, Supporting Information) may have been poorly predicted due to the absence of features such as the cyclooctyl in the training set molecules, and the most similar compound (missing this feature) was classed as inactive. This points to the need for a structurally diverse training set and understanding the threshold similarity distance for making a prediction.29 Scoring of a set of 177 in vitro active compounds from GSK12 with the different machine learning models showed some variation in the number predicted as likely active in vivo. Twenty-four compounds were predicted as consensus actives (Table S5, Supporting Information). This result represents a potential data set for testing the in vivo model predictions. Their further prioritization for in vivo study could also rely on predicted ADME/Tox profiles and Mtb targets (Table S6, Supporting Information). Additionally, with the attainment of more...
extensive SAR for these chemical series, one could apply the scaffold analysis approach illustrated herein for the triazines.

We have described a data curation and machine learning process that could be further expanded to capture the remaining public Mtb in vivo data that we have been unable to find to date. As older articles prior to 1970 may not have an abstract in PubMed, we also used SciFinder and Web of Knowledge to expand our possible range of journals covered to extract as many molecules as possible. Ultimately, training and test set molecules were extracted from 119 references (Table S1, Supporting Information). Another consideration when curating data from old papers is that there are few clearly drawn structures, and it can be a very complex and tedious process even for experienced medicinal chemists to definitively identify structures. For this reason, some structures and data were excluded because of this uncertainty or structural ambiguity.

While prospectively testing the models would be preferred, the high cost of in vivo experiments (currently ~$5000 per compound) renders this practically difficult, and much of this testing is coordinated by the NIH at academic laboratories. Traditionally, the use of computational machine learning models in other areas for which testing is relatively expensive (e.g., ADME/Tox models like hERG requiring patch clamping data\textsuperscript{31–38}) has involved prospective testing on a long time scale (10 years) as more data is generated by different groups and as more higher throughput techniques are developed. This also enables iterative model building and updating to expand the chemistry property space covered, while also expanding the scope of algorithms tested and compared with each other.

In conclusion, we suggest that the machine learning methods described here utilized for modeling Mtb in vivo data could also be applied to other diseases for which similar information from a pertinent animal model is available. Currently, to our knowledge, there are no databases that cover or curate in vivo animal model data, and other sources like PubChem focus almost exclusively on in vitro HTS data. Several limitations of the current study include poor extrapolation from mouse (or other animal models) to human due to likely ADME differences.\textsuperscript{25,64} In addition, we have not yet pursued modeling combination therapies and potential synergies and antigens of experimental antituberculars with known drug treatments. We have also not removed prodrugs from our data set, and we have combined data from several dosing routes, both of which may add to the noise in the models. It is important to note that the four computational models do not per se replace in vivo studies but instead they may enable prioritization of molecules that are likely to perform well in vivo. This approach has obvious benefits for decreasing costs and reducing animal testing\textsuperscript{36} as well as continuing to enrich and accelerate the tuberculosis drug discovery process by learning from prior data. It may also be a useful approach for design of antituberculars or at the very least prioritizing compounds for host-derived therapeutics for TB.\textsuperscript{84} This work also calls for more sharing of data so that we can avoid repeating the discoveries and failures of the past in the future.

ASSOCIATED CONTENT

Supporting Information

Seven figures, six tables, and supplemental references. All computational models are available from the authors upon request. All molecules used in the models are available in CDD, ChemSpider, and FigShare. This material is available free of charge via the Internet at http://pubs.acs.org.

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Notes

The authors declare the following competing financial interest(s): S.E. is a consultant for Collaborative Drug Discovery, Inc.

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ABBREVIATIONS USED

ADME, absorption, metabolism, distribution, and excretion; CPUs, colony-forming units; DHFR, dihydrofolate reductase; FCFP_6, molecular function class fingerprints of maximum diameter 6; GSK, GlaxoSmithKline; HTS, high-throughput screening; hERG, human ether-a-go-go related-gene; MMDS, Mobile Molecular DataSheet; Mtb, Mycobacterium tuberculosis; NIH, National Institutes of Health; PCA, principal components analysis; PXR, pregnane X-receptor; SAR, structure–activity relationship; RP, recursive partitioning; SI, selectivity index; SVM, support vector machine; TB, tuberculosis; XV ROC AUC, cross-validated receiver operator curve area under the curve

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