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DeepBiome: A Phylogenetic Tree Informed Deep Neural Network for Microbiome Data Analysis

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Abstract

Evidence linking the microbiome to human health is rapidly growing. The microbiome profile has the potential as a novel predictive biomarker for many diseases. However, tables of bacterial counts are typically sparse, and bacteria are classified within a hierarchy of taxonomic levels, ranging from species to phylum. Existing tools focus on identifying microbiome associations at either the community level or a specific, pre-defined taxonomic level. Incorporating the evolutionary relationship between bacteria can enhance data interpretation. This approach allows for aggregating microbiome contributions, leading to more accurate and interpretable results. We

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Jing Zhai and Youngwon Choi contribute equally to this study.

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Conflict of interest The authors declare that they have no competing interests.

Ethical Approval All utilized microbiome datasets are de-identified secondary data analysis. No ethics approval was required for this study.

Informed Consent No consent to participate was required for this study.

present DeepBiome, a phylogeny-informed neural network architecture, to predict phenotypes from microbiome counts and uncover the microbiome–phenotype association network. It utilizes microbiome abundance as input and employs phylogenetic taxonomy to guide the neural network’s architecture. Leveraging phylogenetic information, DeepBiome is applicable to both regression and reduces the need for extensive tuning of the deep learning architecture, minimizes overfitting, and, crucially, enables the visualization of the path from microbiome counts to disease. It classification problems. Simulation studies and real-life data analysis have shown that DeepBiome is both highly accurate and efficient. It offers deep insights into complex microbiome–phenotype associations, even with small to moderate training sample sizes. In practice, the specific taxonomic level at which microbiome clusters tag the association remains unknown. Therefore, the main advantage of the presented method over other analytical methods is that it offers an ecological and evolutionary understanding of host–microbe interactions, which is important for microbiome-based medicine. DeepBiome is implemented using Python packages Keras and TensorFlow. It is an open-source tool available at <https://github.com/Young-won/DeepBiome>.

Keywords

Metagenomics; Phylogenetic tree; Neural networks; Prediction; Mixed taxonomic levels

1 Introduction

Emerging high-throughput sequencing technologies have vastly improved our understanding of the human microbiome’s role in many diseases [10, 12, 21, 26]. Despite these achievements, our knowledge of the mechanisms behind microbes’ involvement in disease aggravation remains limited. A notable challenge is inferring and visualizing the path from bacteria to disease across various taxonomic levels.

In 16S ribosomal RNA (RNA) sequencing, sequences are clustered into operational taxonomic units (OTUs) at a threshold of 97% sequence similarity [4]. Sequence representatives, those with the fewest mismatches compared to others in a cluster, are used for taxonomic assignment. These assignments are based on databases of known 16S rRNA gene sequences, such as GreenGenes (des), the Ribosomal Database Project [8], and Silva [24]. Consequently, the processed microbiome data comprise an OTU abundance matrix with rows as samples and columns as OTUs, accompanied by a phylogenetic tree. This tree is crucial for understanding the relationships among OTUs and has been utilized in statistical models to better identify microbiome elements associated with host phenotypes [33, 35, 36]. Closely related microbial taxa often form clusters with similar biological functions, but determining the specific taxonomic level at which association signals are clustered remains a challenge. Are these signals concentrated at a shallow phylogenetic depth (e.g., genus), at a deeper level (e.g., phylum), or across various taxonomic ranks?

Several methods have been proposed to integrate the phylogenetic structure into analyses. Chen et al. [5, 6] introduced a Laplacian penalty, constructed from the sum of branch lengths linking any two OTUs on the evolutionary tree, but their methods are not designed

to detect signals at different evolutionary depths. Garcia Tanya et al. [11] proposed a sparse regression model using ℓ_1 and ℓ_2 regularizations to achieve sparsity at multiple taxonomic levels, yet this method requires three tuning parameters and an exhaustive grid search. Furthermore, it can only select taxa at up to three levels, failing to cover the entire range of phylogenetic depths. Wang and Zhao [31] utilized a tree-guided variable fusion method to build predictive models using bacteria at different taxonomic levels based on the assumption that closely related bacteria have similar biological functions. However, Zhou et al. [38] provided a counterexample; in their study, *Corynebacterium* and *Rothia*, belonging to the same order *Actinomycetales*, had opposite effects on lung function changes in an HIV positive population. Xiao et al. [34] developed a generalized linear mixed model using the evolutionary rate as a tuning parameter, capable of identifying clustered signals without prior knowledge of the phylogenetic depth. However, this method is less effective when microbiome effects are clustered at mixed taxonomic levels.

Deep Neural Networks (DNN), an area of growing interest in biomedical research, show promise for analyzing complex microbiome data. Lu et al. [18]'s study applied a DNN model to the gut microbiome, identifying important bacteria associated with body mass index (BMI). Another study by Reiman et al. [25] incorporated phylogenetic data into a Convolutional Neural Network (CNN) architecture to predict various outcomes. They translated taxa abundances at each phylogenetic level into an abundance matrix, capturing the spatial information of taxa in the phylogenetic tree. However, their model, limited to a fixed number of layers (three), resulted in the loss of taxa lineage information and produced neurons lacking biological meaning, thus making the model difficult to interpret. Other deep learning methods, like MDeep [32], comprising multiple convolutional layers followed by fully connected layers, and DeepMicro [22], utilizing various autoencoders, also struggle with interpretability.

We present DeepBiome, a DNN-based predictive model designed to capture microbiome signals at different phylogenetic depths, applicable to both regression and classification problems. It processes microbiome taxonomic abundance data as input and regularizes the neural network architecture according to the phylogenetic structure. This significantly reduces the number of parameters and the tuning burden compared to conventional neural networks, enabling the identification of important taxa associated with outcomes at all taxonomic levels. Phylogeny regularization in DeepBiome is achieved through weight decay, a widely used technique to prevent overfitting and enhance neural network performances [13, 16, 20, 37]. Unlike existing weight decay schemes that assume a global rate of decay, DeepBiome incorporates the bacteria's evolutionary relationship into a differential weight decay regularization matrix, thereby generating an interpretable effect transfer network for modeling and analyzing microbiome data. Simulation studies and analyses of datasets from a shotgun metagenomic study and a lung microbiome study demonstrate DeepBiome's superior performance over commonly used tools such as support vector machines (SVM), regression with ℓ_1 (Lasso) or $\ell_1 + \ell_2$ (Elastic-Net) penalties, DNN without tree regularization, DNN with ℓ_1 penalty, and Random Forests.

2 Methods and Materials

2.1 Microbiome Data Structure

Suppose we have p OTUs from a total of n microbiome samples, along with a phylogenetic tree that outlines the evolutionary relationships among these microbes. In this tree, each OTU is represented as a tip node, while every internal node corresponds to a taxonomic unit, signifying a common ancestor of its descendant taxa. In our study, we focus on consolidating these p OTUs into m genus-level taxa, treating them as our primary units for analysis. Our analysis could also begin at finer taxonomic levels. An illustrative phylogenetic tree, which we refer to throughout our methods and data simulation discussions, is depicted in Fig. 1. Here, let $\mathbf{x} = (x_1, \dots, x_n)$ denote the input data, with $\mathbf{y} = (y_1, \dots, y_n)$ representing the targeted outcomes. For each subject i , $\mathbf{x}_i = (x_{i1}, \dots, x_{im})^T$ indicates the abundance of m genera. The outcome variable y can be continuous, binary, or categorical.

2.2 Architecture of DeepBiome

DeepBiome is a neural network architecture that associates input vectors \mathbf{x} (representing microbiome abundance) with a clinical outcome \mathbf{y} . A major challenge in neural network construction is deciding on the optimal number of layers and neurons. The conventional wisdom advocates for going deep (many layers) and wide (many neurons per layer), a strategy that has achieved great success in various artificial intelligence tasks. These include image pattern recognition and natural language processing. However, this approach requires a significant amount of training data [3, 28], which is often impractical in biomedical studies due to resource constraints. DeepBiome pre-specifies the network architecture according to the phylogenetic tree. The number of hidden layers is determined by the number of taxonomic levels, while the number of neurons in each layer corresponds to the number of taxa at that particular level. Figure 1b illustrates an example DeepBiome architecture. The input layer accepts microbiome abundances. The information is then propagated through multiple layers of the DeepBiome network to the outcome of interest \mathbf{y} . For example, the input vector \mathbf{x} , representing the abundances of $m^{(0)}$ genera, is propagated to the first hidden layer vector $\mathbf{z}^{(1)}$. This layer contains a total of $m^{(1)}$ neurons, corresponding to the number of family-level taxa. Using an $m^{(1)} \times m^{(0)}$ weight matrix $\mathbf{w}^{(1)}$ and an $m^{(1)}$ bias vector $\mathbf{b}^{(1)}$, we have

$$\mathbf{z}^{(1)} = v(\mathbf{w}^{(1)}\mathbf{x} + \mathbf{b}^{(1)}). \quad (1)$$

Each weight parameter $w_{jk}^{(1)}$ represents the effect of the k th input unit on the j th hidden neuron, and $v(\cdot)$ is an activation function. Throughout this paper, we use the rectified linear unit (ReLU) activations [14, 19]

$$\text{ReLU: } v(a) = a^+ = \max(0, a),$$

(2)

but it can be easily changed to other activation functions in our software. Similarly, $\mathbf{z}^{(\ell)} = v(\mathbf{w}^{(\ell)}\mathbf{z}^{(\ell-1)} + \mathbf{b}^{(\ell)})$, $\ell = 2, \dots, L$, where L is the total number of hidden layers in the neural network. The last hidden layer $\mathbf{z}^{(L)}$ is linked to the outcome using either an identity link or a softmax link. Specifically, we use identity link to predict a continuous outcome, $\mathbf{y} = \mathbf{w}^{(L+1)}\mathbf{z}^{(L)} + \mathbf{b}$. For categorical outcomes with K categories, softmax function is adopted to predict the probability of i th subject belonging to c -th category,

$$\Pr(y_i = c) = \frac{e^{(\mathbf{w}\mathbf{z}^{(L)} + \mathbf{b})_c}}{\sum_{q=1}^K e^{(\mathbf{w}\mathbf{z}^{(L)} + \mathbf{b})_q}}. \quad (3)$$

Finally, we use $f_{\theta}(\mathbf{x})$ with parameters $\theta = \{\mathbf{w}, \mathbf{b}\}$ to represent the whole neural network that maps an input \mathbf{x} to an output \mathbf{y} , where $\mathbf{w} = (\mathbf{w}^{(1)}, \dots, \mathbf{w}^{(L)}, \mathbf{w}^{(L+1)})$ and $\mathbf{b} = (\mathbf{b}^{(1)}, \dots, \mathbf{b}^{(L)}, \mathbf{b}^{(L+1)})$.

2.3 Phylogeny Regularization via Weight Decay

We introduce phylogeny regularization through weight decay. We assume that if taxa j and k have an ancestor-descendent relationship, the associations between the corresponding neurons are stronger, i.e., larger weight value $w_{j,k}$. When taxa j and k do not have this ancestral relationship, we assume $w_{j,k}$ to be a small value, i.e., weight decay. Thus, we construct a weight decay matrix ω to regularize weights in the neural network using the evolutionary relationship carried by the phylogenetic tree. If nodes j and k are ancestor-descendent related, $\omega_{j,k} = 1$; if not, $\omega_{j,k}$ is a small value, e.g., 0.01. See Fig. 1c as an illustration.

2.4 Model Training

Given a training set consisting of training pairs $\{\mathbf{x}, \mathbf{y}\}$ and a neural network $f_{\theta}(\mathbf{x})$ with parameters θ , a supervised training procedure is implemented to learn neural network function by minimizing the empirical loss. We define loss to be Mean Squared Error (MSE) for continuous outcomes and standard Cross-Entropy (CE) for categorical outcomes:

$$\begin{aligned} \text{MSE} &= \frac{1}{n} \sum_{i=1}^n (y_i - \hat{y}_i)^2, \\ \text{CE} &= -\frac{1}{n} \sum_{i=1}^n \sum_{q=1}^K y_{i,q} \log \hat{y}_{i,q}, \end{aligned} \quad (4)$$

where $y_{i,k}$ is a binary indicator (0 or 1) indicating whether observation i belongs to class k (i.e., one hot encoding). $y_{i,c} = \Pr(y_i = c) (c \in \{1, \dots, K\})$ is the probability that observation i belongs to class c defined by Eq. (3). We use the holdout validation method to determine

the stopping criterion during training. The training process stops when the holdout validation error achieves the minimum.

Algorithm 1: Phylogeny regularized weight decay in Adam. β_1, β_2 refer to the exponential decay rates for the moment estimates in Adam. $\epsilon = 10^{-8}$ is used to prevent division from zero error [14].

Data:	$\mathbf{y}, \mathbf{x}_i = (x_1, \dots, x_m)$, phylogenetic tree designed matrix ω , learning rate $\alpha \in \mathcal{R}$
Result:	$\hat{\theta}$ such that the lost function $\text{Loss}(\theta)$ is minimized.
1	Initialize: parameter $(\mathbf{w}, \mathbf{b})_{t=0} \in \mathcal{R}$, 1 st and 2 nd moment vector $m_{t=0}^1 \leftarrow \mathbf{0}$ and $m_{t=0}^2 \leftarrow \mathbf{0}$, and step index $t \leftarrow 0$,
2	repeat
3	$t \leftarrow t + 1$
4	$l'_t \leftarrow \nabla_{\theta} \text{Loss}(\omega \circ \mathbf{w}_{t-1}, \mathbf{b}_{t-1})$ (Get gradients w.r.t. stochastic objective at t)
5	$m_t^1 \leftarrow \beta_1 m_{t-1}^1 + (1 - \beta_1) l'_t$ (Update biased 1 st moment estimate)
6	$m_t^2 \leftarrow \beta_2 m_{t-1}^2 + (1 - \beta_2) l''_t$ (Update biased 2 nd moment estimate)
7	$\hat{m}_t^1 \leftarrow m_t^1 / (1 - \beta_1^t)$ (Update bias-corrected 1 st moment estimate)
8	$\hat{m}_t^2 \leftarrow m_t^2 / (1 - \beta_2^t)$ (Update bias-corrected 2 nd moment estimate)
9	$(\mathbf{w}, \mathbf{b})_t \leftarrow (\mathbf{w}, \mathbf{b})_{t-1} - \alpha \cdot \hat{m}_t^1 / (\sqrt{\hat{m}_t^2} + \epsilon)$ (Update parameters)
10	until stopping criterion is met and return $\hat{\theta} = (\omega \circ \hat{\mathbf{w}}, \hat{\mathbf{b}})$;

Adam optimizer, an adaptive gradient algorithm [15], is used to train DeepBiome. Algorithm 1 describes the parameter estimation in Adam with the proposed phylogeny regularization. Specifically, at each update t , the estimated weight w_i is elementwisely multiplied by the corresponding regularization factor ω and $\omega \circ w_i$ is used to predict \hat{y} in the loss function [see Eq. (4)]. Once the stopping criteria is met, Algorithm 1 outputs the estimated parameters of each layer $(\hat{\mathbf{w}}, \hat{\mathbf{b}}) = (\{\omega^{(1)} \circ \hat{\mathbf{w}}^{(1)}, \dots, \omega^{(L)} \circ \hat{\mathbf{w}}^{(L)}\}, \{\hat{\mathbf{b}}^{(1)}, \dots, \hat{\mathbf{b}}^{(L)}\})$. This phylogeny regularization effectively uses biologically meaningful prior knowledge to limit the number of free parameters in the model. Therefore, it avoids overfitting.

2.5 Performance Metrics

We employ several statistical metrics to evaluate the performance of DeepBiome for its prediction, classification and taxa selection performances. For a quantitative outcome, the primary metric is the mean square error (MSE). Additionally, we report the Pearson correlation coefficient ρ between predicted \hat{y}_i and true y_i . For categorical outcomes, i.e., classification problems, we measure their performance using sensitivity [true positive rate (TPR)], specificity, g -measure, accuracy (ACC), precision (PPV), and the F1 score:

$$\text{Sensitivity} = \frac{\text{TP}}{\text{TP} + \text{FN}},$$

$$\text{Specificity} = \frac{\text{TN}}{\text{TN} + \text{FP}},$$

$$g\text{-Measure} = (\text{Sensitivity} \times \text{Specificity})^{\frac{1}{2}},$$

$$\text{ACC} = \frac{\text{TP} + \text{TN}}{\text{TP} + \text{TN} + \text{FP} + \text{FN}},$$

$$\text{PPV} = \frac{\text{TP}}{\text{TP} + \text{FP}},$$

$$\text{F1score} = 2 \times \frac{\text{PPV} \times \text{TPR}}{\text{PPV} + \text{TPR}} = \frac{2\text{TP}}{2\text{TP} + \text{FP} + \text{FN}},$$

where TP is “true positive” (or recall), TN is “true negative”, FP is “false positive”, and FN is “false negative”. The F1 score is the harmonic mean of precision and sensitivity. An F1 score reaches its best value at one when the prediction has perfect precision and recall and the worst at zero. Note that the F1 score does not take the true negative into account. We use the *g*-measure, the geometric mean of sensitivity and specificity, to assess the performance of a binary classifier. Same as an F1 score, a *g*-Measure reaches its best value at one when the sensitivity and specificity are both perfect (one) while the worst at zero if any of the sensitivity and specificity is zero. We also report AUC (area under the receiver operating characteristics), which reports the capability of a model to distinguish between classes. Sensitivity, specificity, *g*-Measure, and ACC across all hidden layers (see Table 1) are used to report the selection accuracies.

3 Results

3.1 Simulation Studies

We conduct extensive simulation studies to evaluate the performance of DeepBiome and compare it with conventional methods in three different schemes, i.e., linear regression, binary, and multiclass ($K \geq 3$) classification design. Throughout the simulation experiments, we use sample size $n = 1000$ and split them into a training set (75%, $n_{\text{training}} = 750$) and a test set (25%, $n_{\text{test}} = 250$). Different proportions of the split give qualitatively similar results (not shown). All results are obtained based on 1000 replicates. Simulation scenario 1 covers continuous outcome models; simulation scenario 2 is for binary outcome cases, and simulation scenario 3 considers the situation when the outcome variable is categorical. Model robustness is evaluated in simulation scenario 4 when tree structure is misspecified and microbiome abundances contain measurement errors.

Detailed procedures to generate microbiome abundance data have been described before by our group [35, 36]. We use a Dirichlet Multinomial (DM) distribution with the mean proportion vector and the dispersion parameter estimated from a real pulmonary microbiome

dataset to generate OTU counts. We then aggregate 2964 OTUs to 48 genus according to the phylogenetic tree [30]. A forward propagation approach is described below to generate \mathbf{y} .

1. Read the phylogenetic tree to obtain the number of phylogenetic levels and nodes at each level. The microbiome data are then summarized at genus, family, order, class, and phylum level as shown in Fig. 1. Based on a real lung microbiome dataset, the number of nodes are $m^{(0)} = 48$, $m^{(1)} = 40$, $m^{(2)} = 23$, $m^{(3)} = 17$, and $m^{(4)} = 9$, respectively.
2. Construct the weight matrix $\mathbf{w}^{(1)} \in \mathbb{R}^{m^{(1)} \times m^{(0)}}$ to propagate the input layer $\mathbf{x}_{\text{genus}}$ to the 1st hidden layer by

$$\mathbf{h}^{(1)} = \mathbf{w}^{(1)}\mathbf{x}_{\text{genus}} + \mathbf{b}^{(1)}.$$

The bias vector $\mathbf{b}^{(1)} \in \mathbb{R}^{m^{(1)} \times 1}$ follows a standard normal distribution $\mathcal{N}(0, \sigma_c^2)$ with $\sigma_c^2 = 4$. Suppose we have node j at the genus level and k at the family level, then

$$w_{j,k}^{(1)} \sim \begin{cases} \text{Uniform}(-0.5, 1) & \text{associated with output,} \\ \mathcal{N}(0, 0.01) & \text{not associated with output.} \end{cases} \quad (5)$$

3. Multiply the $w_{j,k}^{(1)}$ by a small value $\omega_{j,k} = 0.01$, if family taxa k is not a direct ancestor of genus taxa j ; otherwise, $w_{j,k}^{(1)}$ stays the same.
4. Activate the neurons using ReLU in Eq. 2,

$$\mathbf{x}_{\text{family}} = \mathbf{z}^{(1)} = v(\mathbf{w}^{(1)}\mathbf{x}_{\text{genus}} + \mathbf{b}^{(1)}).$$

5. Repeating steps 2–4 to compute the $\mathbf{x}_{\text{order}}$, $\mathbf{x}_{\text{class}}$, and $\mathbf{x}_{\text{phylum}}$.
6. Simulate the continuous or categorical output layer \mathbf{y} as follow

$$\hat{\mathbf{y}} = \mathbf{w}^{(4)}\mathbf{x}_{\text{phylum}} + \mathbf{b}^{(4)}$$

$$\Pr(y_i = c) = \frac{(e^{\mathbf{w}^{(4)}\mathbf{x}_{\text{phylum}} + \mathbf{b}^{(4)}})_c}{\sum_{q=1}^K (e^{\mathbf{w}^{(4)}\mathbf{x}_{\text{phylum}} + \mathbf{b}^{(4)}})_q}, \quad (6)$$

where $K = 2$ for binary classification and $K \geq 3$ for multi-categorical classification.

The following simulation schemes are considered to examine the robustness of DeepBiome method

1. Abundance contains measurement errors at genus levels. We assume that 10% of the associated genus reads are misclassified to one randomly selected genus from

the same phylum. The microbiome abundance data with measurement errors are then used for training models.

2. The phylogenetic tree for training models is misspecified (see Fig. 2).
 - At class level, the genera that belong to *Clostridia* and *Flavobacteria* are mis-classified to *Bacilli* and *Bacteroidia*.
 - At order level, the genera that belong to *Coriobacteriales* and *Flavobacteriales* are mis-classified to *Actinomycetales* and *Bacteroidales*.

We use fivefold cross-validation to choose the tuning parameters for regularized linear regression models. R package `randomForest` is adopted for Random Forest analysis [17]. Fivefold cross-validation is used to obtain hyper-parameters in `randomForest` [i.e., the number of features selected at each split (`mtry`) and the number of trees generated in a forest (`ntree`)] so that out of bag (OOB) error is minimized. The values of `mtry` and `ntree` vary across datasets, with a mean (standard deviation) to be 800 (250) and 36 (30), respectively. The predictors that result in a mean decrease in accuracy or MSE are selected. On average, 55% of the predictors are selected. The variance importance metric is defined as the decrease in MSE or accuracy when the variable is randomly permuted. For the deep learning models (i.e., DNN, DNN + ℓ_1 , and DeepBiome), we use a holdout validation set including 20% of the training data and train the models until either validation error increases or 5000 epochs are reached, whichever comes the first. Using Adam [15] optimizer, we set $\beta_1 = 0.9$, $\beta_2 = 0.999$, learning rate $lr = 0.01$ and the mini-batch gradient descent with a mini-batch size of 50. The learning rate decays for each epoch with $lr_{\text{epoch}+1} = lr_{\text{epoch}} \frac{1}{1 + 0.0001}$.

3.1.1 Scenario 1: Regression Design: Table 2 displays the prediction performance for continuous outcomes when the associated taxa are only clustered at one phylogenetic level (i.e., phylum). DeepBiome has a higher Pearson correlation and lower MSE on the test set than the regression methods. It also performs the best among all deep learning models. Table 3 shows the prediction performances under a more complex case, where the associated taxa are clustered at different phylogenetic levels (i.e., phylum and order). All regression schemes perform poorly in this case with low correlation, e.g., 0.6. DeepBiome has over 80% reduction in MSE compared to regression-based methods. The deep learning models, DNN and DNN + ℓ_1 , improve correlation to 0.91 and 0.9, respectively. However, both show a hint of overfitting with lower testing performance. Compared with Random Forest, DeepBiome has slightly higher MSE (0.07 vs. 0.04 shown in Table 2, 0.42 vs. 0.30 shown in Table 3) and slightly lower correlation (0.88 vs. 0.94 shown in Table 2 and 0.92 vs. 0.95 shown in Table 2).

Identifying associated taxa at precise levels is critical. We evaluate the selection performance shown in Fig. 3. Regular regression methods do not discriminate associated taxa; therefore, only the results of penalized regressions are shown. Lasso and Elastic-Net can only select the taxa at one phylogenetic level in the penalized regression schemes. We compute the

performance metrics for higher-level taxa selected using their phylogeny relationship. For example, if genus *Prevotella* is selected, we assume that its ancestor, phylum *Bacteroidetes*, is also selected. In contrast, the selection performance of regularized neural network models is based on the weights estimated at each hidden layer. *g*-Measure for DeepBiome ranges from 0.8 to 0.9 across different taxa levels (Fig. 3, the first and second row). DeepBiome also has excellent sensitivity, specificity, and ACC. Although the Random Forest method performs better in predicting continuous outcomes, its *g*-measures range from 0.56 to 0.6, indicating poor selection ability. DNN + ℓ_1 fails to identify the true microbiome taxa across all phylogenetic levels with much lower *g*-measure, e.g., 0.18 for genus level taxa selection (Fig. 3, the first and second row). Overall, for continuous outcome predictions, Random Forest performs slightly better than DeepBiome. Yet DeepBiome offers better selection ability.

3.1.2 Scenario 2: Binary Classification: For simplicity and for demonstrating DeepBiome's ability to discriminate different levels' taxa, we only consider the case that outcome-associated taxa are clustered at mixed phylogenetic levels for the following simulation scenarios (as shown in Fig. 2b). We assume

- [(1)] the higher the abundance of blue node taxa, the higher the probability of y belonging to the disease group;
- [(2)] the higher the abundance of red node taxa, the higher the probability of y belonging to the healthy control group.

We compare DeepBiome to logistic regression, three penalized logistic regression models, two conventional deep learning networks, and the Random Forest method. In Table 4, we present the metrics for evaluating the classification performance of binary outcomes, including sensitivity, specificity, *g*-Measure, ACC, and AUC. Logistic regression has satisfying sensitivity, but other metrics are not competitive compared to DeepBiome. Logistic regressions tend to have more false positives. In contrast, DeepBiome achieves the best classification performance with the highest specificity, *g*-Measure, ACC, and AUC as, 0.84, 0.87, 0.89, and 0.94, respectively. Interestingly, for binary outcome prediction, DeepBiome outperforms the Random Forest method with higher *g*-measure (0.87 vs. 0.84) and AUC (0.94 vs. 0.85) (see Table 4). Figure 3 (the second row) displays the performance of identifying associated taxa. However, Lasso and DNN + ℓ_1 show good sensitivity at some phylogenetic levels, and *g*-Measure and ACC are much worse than Elastic-Net and DeepBiome. This suggests that Lasso and DNN + ℓ_1 selected many null taxa (false positive). Using the order level as an example, the *g*-Measure value of DeepBiome is 0.91, while the DNN + ℓ_1 is 0.15. The selection performance comparison between DeepBiome and Random Forest indicates that DeepBiome also has superior performance, i.e., *g*-measure of Random Forest vs. DeepBiome was 0.61 vs. 0.83 for genus level taxa selection. It is worth noting that logistic regression with Elastic-Net penalization also offers acceptable selection accuracy, e.g., 0.65 (genus level) vs. 0.81 (class level) (Fig. 3, the third row) (Table 5).

3.1.3 Scenario 3: Multiclass Classification: We evaluate the performance of DeepBiome for multi-category outcomes. We assume, as shown in Fig. 2b, that the blue

node taxa lead to “severe” disease, red ones lead to “mild” disease, and gray node taxa are neutral.

We compare DeepBiome to DNN, $DNN + \ell_1$, the support vector machine (SVM) with different kernels, and the Random Forest method. For SVM, linear and non-linear kernels, such as the radial or polynomial kernels, are evaluated. The default parameter setting is adopted for SVM. Among the SVMs, the linear SVM has the highest accuracy and AUC, while the SVM with radial kernel yields better recall and F1 score. However, all SVMs are inferior to deep learning models. DeepBiome exhibits the highest AUC, i.e., 0.9; AUCs of DNN and $DNN + \ell_1$ are around 0.86. The F1 score of DeepBiome is 0.711, which is 14% higher than the second best, $DNN + \ell_1$. We find that DeepBiome offers the best and most balanced performance with precision and recall, which are 0.72 and 0.71, respectively. Consistent with binary outcome comparisons, DeepBiome also outperforms the Random Forest method for multi-categorical outcome prediction. Since SVM models cannot select the microbiome taxa, we only compare DeepBiome to $DNN + \ell_1$ and Random Forest shown in Fig. 3 (the 4th row). DeepBiome surpasses $DNN + \ell_1$ in all of the evaluation metrics at all phylogenetic levels. For instance, the g -measure of $DNN + \ell_1$ in selecting genus level taxa is only 0.19 while that of DeepBiome is 0.82. Note that across three different outcome types, the selection ability of the Random Forest method remains similar with g -measure ranging from 0.5 to 0.7.

3.1.4 Scenario 4: Robustness Under Tree Misspecification and Measurement Errors of Microbiome Abundance:

Table 6 and Fig. 3 (the 5th row) present the results when microbiome abundances contain sequencing errors. Table 7 and Fig. 3 (the 6th row) show the results when using a misspecified phylogenetic tree to train and test the model. Like scenario 1, we simulate continuous outcomes and compare DeepBiome with linear regression, penalized regressions, conventional DNN, ℓ_1 -regularized DNN, and Random Forest. When the model is trained using data with measurement errors (case 1), the performance of all methods decreases compared with scenario 1 using data without errors (Table 6; see also Table 3). For example, for the Random Forest method, the MSE is 0.04 (Table 2) without sequencing error vs. 0.42 with sequencing error (Table 6). For DeepBiome, MSE rises from 0.07 (without sequencing error) to 0.24 (with sequencing error). These results indicate that although Random Forest offers better predictive ability when the outcome is continuous, it is more sensitive to sample contamination than DeepBiome. The average Pearson’s ρ of DeepBiome is 0.95, while those of DNN, $DNN + \ell_1$, and Random Forest are 0.87, 0.91, and 0.92 respectively.

Table 7 displays the prediction performance under case 2 (i.e., using a misspecified phylogenetic tree to train models). DeepBiome outperforms other methods in both MSE and Pearson’s ρ except Random Forest. MSE of Random Forest is 0.04 (without misspecified tree) vs. 0.19 (with misspecified tree) while DeepBiome is 0.07 (without misspecified tree) vs. 0.32 (with misspecified tree). Since DeepBiome relies on a polygenetic tree for regularization, the impact of using a misspecified tree to DeepBiome is larger than a Random Forest.

Figure 3 (the 5th and the 6th row) show the ability to identify associated microbiome taxa. When the abundance data contain measurement errors, the sensitivity decreases in penalized regression and deep learning methods. For example, the specificity of DeepBiome at the genus level is 0.67 (compared to 0.95 using correct tree information), leading to a decreased g -measure, i.e., from 0.84 to 0.80. DeepBiome tends to select less associated taxa when input abundance data contain measurement errors. The Lasso, Elastic-Net, and DNN + ℓ_1 perform similarly to scenario 1. Even if DeepBiome uses a wrong tree structure to guide the model, it still has decent performance with a g -Measure of 0.80 at the finest level (i.e., genus).

3.2 Disease Prediction Using Shotgun Metagenomics

The resolution of the shotgun metagenomics data can reach species and strain levels, providing in-depth information to quantify the association between microbiota and human health. As the cost of shotgun metagenomics sequencing keeps decreasing, the number of available human metagenomics datasets keeps increasing. Using eight large-scale publicly available metagenomic datasets, Reference [23] benchmarked statistical learning tools for disease classification. Publicly available software and uniformly processed microbiome profiles (<http://segatalab.cibio.unitn.it/tools/metaml>) were also provided. For all eight datasets, species-level taxonomic profiling and relative abundances data were processed using MetaPhlan2 [29], and the detailed sequence processing procedures were reported [23].

We apply DeepBiome to a type 2 diabetes (T2D) cohort among one of the eight studies with the largest sample size. It includes the species-level relative abundances from 170 Chinese T2D patients and 174 controls. We use relative abundances of 572 species-level taxa (i.e., 210 genus-level taxa) and the corresponding phylogenetic tree to predict T2D status and select associated microbiome clusters. Table 8 shows the performance of T2D prediction based on five-fold cross-validation. Although Ridge and Lasso regressions have the highest specificity, their low sensitivity suggests that these methods tend to predict all subjects as healthy. DeepBiome performs the best among all methods with the highest g -Measure, accuracy, and AUC, which are 0.620, 0.643, and 0.694, respectively. Consistent with our simulation results, DeepBiome shows better predictive power than the Random Forest method. For example, AUC is 0.61 for the Random Forest method and 0.69 for DeepBiome. Figure 4 demonstrates the taxa selected by DeepBiome. We have selected 86 species, 32 genera, 14 families, 7 orders, and 4 classes. Among these taxa, 32 species, 20 genera, 8 families, 3 orders, and 3 classes are positively associated with T2D, indicating that the higher the abundance of those taxa, the higher the probability of subjects having T2D.

3.3 Computational Efficiency

DeepBiome is implemented in Python 3.6 based the TensorFlow [1, 2] and Keras [7] framework. It can be built on Python 3.4, 3.5, and 3.6. All simulations are performed using a workstation equipped with Intel(R) Xeon(R) CPU E5-2650 v4 processor with 24 cores @ 2.20 GHz and one NVIDIA GeForce GTX TITAN X GPU with 3072 CUDA cores @ 1 GHz and 12 GB memory. DeepBiome requires 290 ± 69 s to fully train the network for one replicate with 1000 samples, 50 minibatches, and 5000 epochs. For the same data,

DNN takes 282 ± 67 s, and $\text{DNN} + \ell_1$ takes 282 ± 67 s. DeepBiome and all other deep learning approaches take less than 0.004 s for prediction. All real data analysis is performed on a MacBook Pro with 2.8 GHz Intel Core i7 processor and 16 GB 2133 MHz LPDDR3 memory.

4 Discussion and conclusions

The proposed DeepBiome, a phylogenetic tree-regularized deep learning model, can be used for prediction and classification tasks. We provide comprehensive simulation experiments and real data applications to demonstrate the superiority of DeepBiome. For regression tasks, our results suggest that, compared to sparse regression and other deep learning models, DeepBiome has a competitive performance, particularly when microbiome taxa associated with the outcome are clustered at different phylogenetic levels. DeepBiome also excels in complex classification tasks with higher accuracy and AUC. More importantly, DeepBiome enables an intuitive visualization of the microbiome–phenotype association network.

Deep learning models gain lots of popularity due to their supremacy in imaging and natural language analysis. However, typical biomedical studies can rarely afford the huge amount of training data required for hyper-parameter tuning [9, 27]. DeepBiome regularizes the neural network structure towards the phylogenetic structure inherent in the microbiome data through weight decay. This way, it greatly reduces the number of parameters, including the architecture itself, to be tuned and trained, avoids overfitting, and allows visualization of the pathway from microbiome counts to phenotypes. The limitations of DeepBiome include the possibility of violation of the assumptions: (1) microbiome abundances classified in the same cluster have similar effects to outcomes of interests, and (2) phylogenetic tree structure translates to effects aggregation structure.

In real-world applications, the number of features (e.g., microbial species or genes) may differ between training and testing datasets, posing a significant challenge for most machine learning models, including DeepBiome. We have the following considerations:

1. Prior to training, align the features of both datasets by selecting a common set of features or using techniques like canonical correlation analysis to find a harmonized feature space.
2. DeepBiome's architecture can handle inputs of varying dimensions.
3. Train DeepBiome on the larger feature set and then apply transfer learning techniques to adapt the model to the smaller feature set in the testing phase.

We defer the details of the investigation to future research.

5 Availability and Requirements

Project name: DeepBiome

Project home page: <https://github.com/Young-won/DeepBiome>

Operating system(s): Platform independent

Programming language: Python

Other requirements: None

License: Open source

Any restrictions to use by non-academics: None

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Data Availability

AGP metadata and the operational taxonomic unit (OTU) table were downloaded from <https://github.com/biocore/American-Gut/tree/master/data>. The phylogenetic tree was extracted from the *.biom* file that contains OTU tables and the taxonomic information. Lung microbiome data is available upon request. Consent for Publication All utilized microbiome datasets are de-identified secondary data analysis. No consent for publication was required for this study.

Abbreviations

DeepBiome	A phylogenetic tree informed deep neural network for microbiome data analysis
RNA	Ribosomal ribonucleic acid
OTUs	Operational Taxonomic Units
HIV	Human immunodeficiency virus
DNN	Deep neural network
BMI	Body mass index
CNN	Convolutional neural network AGP: American gut project
SVM	Support vector machine
Adam	An adaptive gradient algorithm
MSE	Mean square error
AUC	Area under the receiver operating characteristics
TPR	True positive rate
ACC	Accuracy

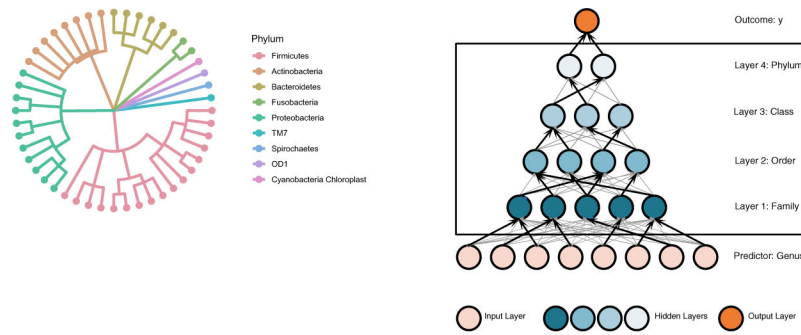
PPV	Precision
TP	True positive (recall)
TN	True negative
FP	False positive
FN	False negative
T2D	Type 2 diabetes
COPD	Chronic obstructive pulmonary disease
FEV1	Forced expiratory volume in 1 s
SPT	Supraglottic predominant taxa
BPT	Background predominant taxa

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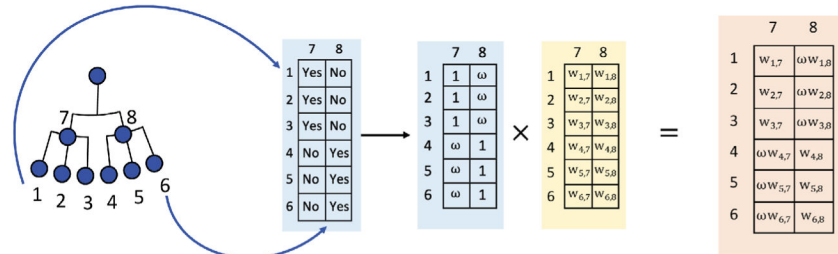
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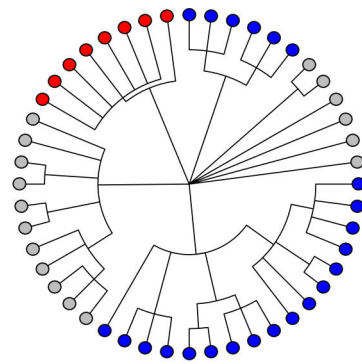


(a) An example phylogenetic tree with 48 genera (b) Network layout of DeepBiome architecture

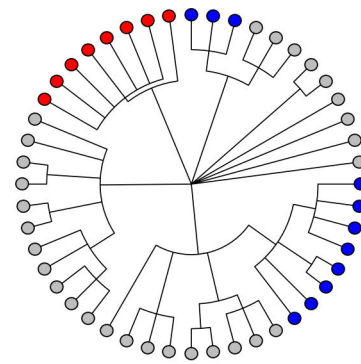


(c) Phylogenetic tree regularized weight decay procedure

Fig. 1. DeepBiome architecture. **a** A phylogenetic tree with 48 genera as tip nodes. Color represents phylum types. **b** Network layout of DeepBiome architecture. The input layer is genus level microbiome abundance. Each hidden layer represents one phylogenetic level, e.g., family, order, class, and phylum. The dark lines represent relationship defined by a phylogenetics tree. The gray lines represent association between layers. **c** Phylogenetic tree regularized weight decay. Suppose we have a simple tree as shown in the left panel, which has 6 genera (taxa 1–6) and 2 classes (7–8). Genera 1–3 belong to class 7 and genera 4–6 belong to class 8. The ancestor–descendent information is embedded into a 6×2 matrix. Without loss of generality, we use ω to indicate a regularization factor with small value (e.g., 0.01). For tree regularized weight decay, the weight estimation matrix $w_{6 \times 2}$ is multiplied with this phylogenetic embedded matrix $\Omega_{6 \times 2}$ elementwisely, denoted by $\Omega_{6 \times 2} \circ w_{6 \times 2}$ (Color figure online)



(a) Phylum Level



(b) Phylum & Order Levels

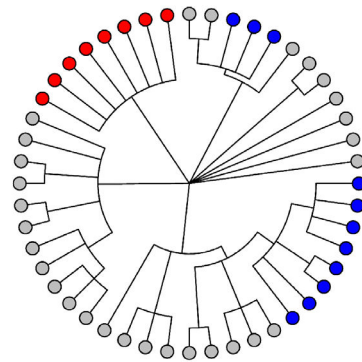
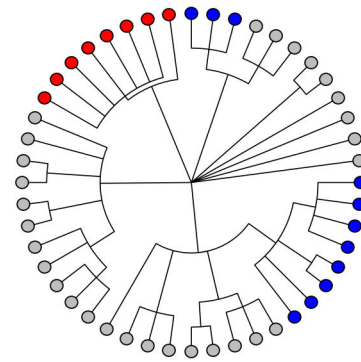
(c) Simulation scenario 4:
Tree in network simulation(d) Simulation scenario 4:
Tree in model learning

Fig. 2. Simulation specifications. Outcome associated taxa (blue and red) are specified at **a** the phylum level and **b** a mixture of phylum and order levels. The blue nodes represent “bad” taxa which result in disease status or are negatively associated with continuous phenotype, e.g., FEV1. The red nodes represent “good” taxa which result in a healthy status. In simulation scenario 4, we evaluate the impact of the misspecified phylogenetic tree: **c** indicates the true phylogenetic tree used in simulation scenario 4 and **d** indicates the phylogenetic tree used in model learning (same as the tree shown in **b**) (Color figure online)

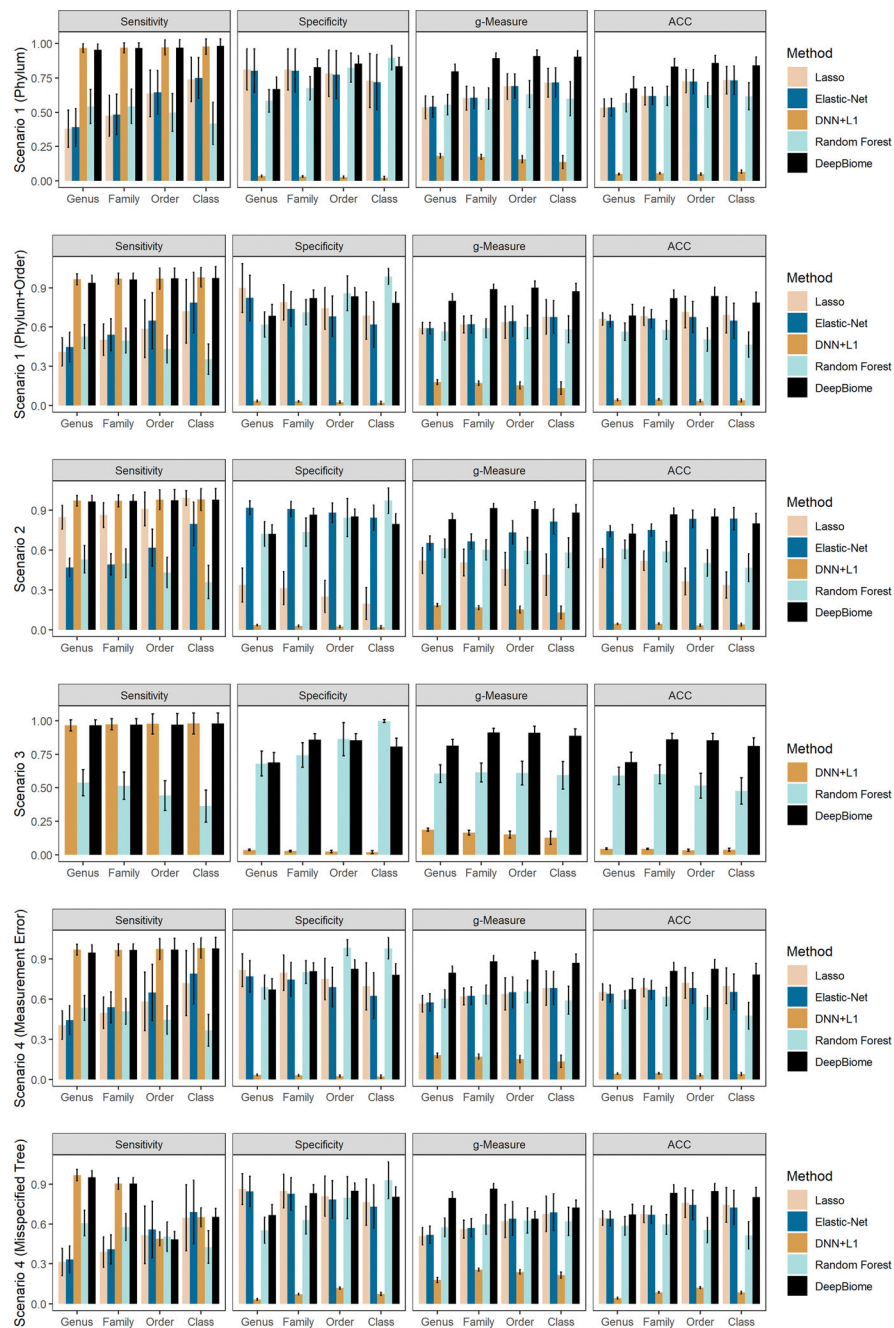


Fig. 3. Taxa selection performance under four simulation schemes at each phylogenetic level. Sensitivity, specificity, g-Measure, and accuracy (ACC) were used to evaluate taxa selection performance. The vertical bar represents the standard deviation over 1000 simulation replicates

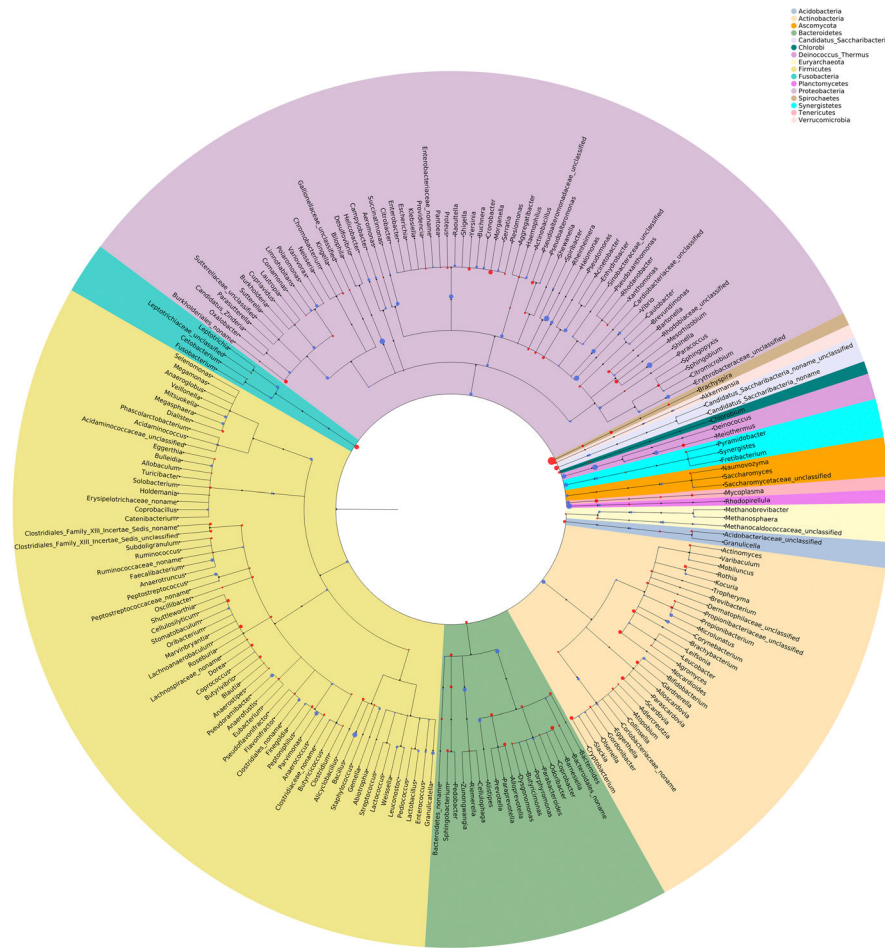


Fig. 4. DeepBiome selected T2D-associated taxa using data from a real metagenomic sequencing study. Estimated weights were overlaid on the phylogenetic tree. The red and blue nodes indicate taxa have positive and negative weights, respectively. The size of colored nodes represents the magnitudes of the weights. Black nodes represent non-selected taxa (Color figure online)

Table 1

Metrics used to assess the performance of outcome prediction and microbiome taxa selection

		Prediction		Selection
		Regression	Binary	Multiclass
Metrics		MSE	Sensitivity	Sensitivity
		Pearson's ρ	Specificity	PPV
			g-Measure	F1 score
			ACC	ACC
			AUC	AUC

MSE mean squared error, *PPV* positive predictive value, *ACC* accuracy, *AUC* area under the receiver operating characteristic (ROC) curve

Table 2

Scenario 1: mean squared error (MSE) and Pearson correlation coefficient between predicted and true outcomes for continuous outcome

Method	Testing				Training			
	MSE		Correlation		MSE		Correlation	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Linear Regression	0.104	0.024	0.824	0.049	0.087	0.011	0.851	0.023
Ridge	0.104	0.022	0.824	0.049	0.09	0.012	0.851	0.023
Lasso	0.100	0.023	0.833	0.048	0.092	0.013	0.843	0.025
Elastic-Net	0.100	0.023	0.833	0.048	0.092	0.012	0.844	0.025
DNN	0.076	0.040	0.874	0.077	0.032	0.034	0.947	0.067
DNN+ ℓ_1	0.075	0.040	0.875	0.073	0.034	0.039	0.945	0.068
Random Forest	0.044	0.017	0.936	0.032	0.044	0.008	0.943	0.016
DeepBiome	0.071	0.036	0.882	0.069	0.043	0.034	0.929	0.061

The associated taxa are clustered at the phylum level

DNN deep neural network, $DNN + \ell_1$ Lasso (least absolute shrinkage and selection operator) penalized deep neural network

Table 3

Scenario 1: mean squared error (MSE) and Pearson correlation coefficient between predicted and true outcomes for continuous outcome

Method	Testing				Training			
	MSE		Correlation		MSE		Correlation	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Linear Regression	1.561	0.146	0.639	0.035	1.337	0.068	0.694	0.018
Lasso	1.479	0.115	0.662	0.034	1.411	0.075	0.678	0.020
Ridge	1.546	0.121	0.639	0.034	1.361	0.075	0.694	0.018
Elastic-Net	1.481	0.117	0.662	0.034	1.405	0.076	0.680	0.020
DNN	0.457	0.522	0.905	0.118	0.164	0.337	0.964	0.091
DNN+ ℓ_1	0.456	0.516	0.904	0.122	0.176	0.362	0.963	0.085
Random Forest	0.296	0.050	0.949	0.011	0.300	0.028	0.948	0.006
DeepBiome	0.423	1.474	0.916	0.139	0.256	0.463	0.944	0.110

The associated taxa are clustered at the phylum and order levels

DNN deep neural network, *DNN* + ℓ_1 Lasso (least absolute shrinkage and selection operator) penalized deep neural network

Table 4

Scenario 2: classification performance for binary outcomes

Method	Testing						Training													
	Sensitivity		Specificity		g-Measure		ACC		AUC		Sensitivity		Specificity		g-Measure		ACC		AUC	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Logistic	0.920	0.030	0.725	0.066	0.815	0.038	0.860	0.026	0.822	0.033	0.950	0.021	0.785	0.049	0.863	0.030	0.900	0.020	0.867	0.026
Lasso	0.965	0.016	0.583	0.086	0.747	0.056	0.848	0.027	0.774	0.041	0.973	0.006	0.619	0.081	0.774	0.053	0.866	0.023	0.796	0.040
Ridge	0.957	0.018	0.461	0.077	0.661	0.055	0.805	0.027	0.709	0.037	0.970	0.007	0.522	0.062	0.711	0.043	0.835	0.016	0.746	0.030
Elastic-Net	0.998	0.004	0.022	0.020	0.121	0.082	0.699	0.030	0.510	0.010	0.998	0.002	0.022	0.013	0.140	0.046	0.702	0.015	0.510	0.006
DNN	0.887	0.049	0.725	0.148	0.788	0.133	0.837	0.038	0.897	0.039	0.932	0.032	0.869	0.158	0.887	0.146	0.913	0.043	0.958	0.033
DNN+ ℓ_1	0.887	0.048	0.727	0.146	0.790	0.129	0.838	0.038	0.898	0.036	0.931	0.031	0.868	0.154	0.887	0.141	0.912	0.042	0.958	0.030
Random Forest	0.731	0.063	0.977	0.013	0.844	0.036	0.902	0.021	0.854	0.031	0.730	0.044	0.976	0.008	0.844	0.024	0.902	0.011	0.853	0.021
DeepBiome	0.918	0.042	0.835	0.111	0.870	0.093	0.892	0.044	0.941	0.051	0.952	0.033	0.922	0.106	0.932	0.094	0.943	0.041	0.974	0.048

ACC accuracy, AUC area under the receiver operating characteristic (ROC) curve, DNN deep neural network, DNN + ℓ_1 Lasso (least absolute shrinkage and selection operator) penalized deep neural network

Table 5

Scenario 3: classification performance for multi-categorical outcomes

Method	Testing						Training													
	ACC		Precision		Recall		F1		AUC		ACC		Precision		Recall		F1		AUC	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
SVM-Linear	0.667	0.032	0.525	0.060	0.493	0.027	0.508	0.039	0.667	0.027	0.694	0.017	0.571	0.034	0.520	0.019	0.544	0.024	0.679	0.018
SVM-Radial	0.659	0.033	0.527	0.092	0.506	0.024	0.514	0.050	0.665	0.027	0.821	0.012	0.807	0.035	0.663	0.017	0.728	0.020	0.809	0.015
SVM-Polynomial	0.576	0.034	0.443	0.090	0.394	0.026	0.414	0.048	0.566	0.030	0.767	0.014	0.837	0.012	0.622	0.025	0.714	0.019	0.742	0.026
DNN	0.752	0.045	0.620	0.136	0.599	0.105	0.599	0.115	0.856	0.041	0.854	0.063	0.741	0.188	0.718	0.149	0.718	0.166	0.941	0.044
DNN+ ℓ_1	0.760	0.042	0.668	0.139	0.612	0.090	0.618	0.104	0.863	0.043	0.864	0.058	0.794	0.160	0.731	0.130	0.739	0.146	0.948	0.037
Random Forest	0.871	0.022	0.861	0.031	0.802	0.033	0.820	0.033	0.807	0.037	0.870	0.012	0.859	0.017	0.800	0.022	0.821	0.021	0.806	0.024
DeepBiome	0.815	0.066	0.720	0.151	0.714	0.115	0.711	0.135	0.900	0.080	0.880	0.072	0.804	0.170	0.793	0.135	0.791	0.157	0.942	0.084

ACC accuracy, AUC area under the receiver operating characteristic (ROC) curve, DNN deep neural network, $DNN + \ell_1$ Lasso (least absolute shrinkage and selection operator) penalized deep neural network, SVM support vector machine

Table 6

Scenario 4: mean squared error (MSE) and Pearson correlation coefficient between predicted and true outcome (continuous), when the input microbiome abundance data contain measurement errors

Method	Testing				Training			
	MSE		Correlation		MSE		Correlation	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Linear Regression	1.569	0.154	0.639	0.036	1.336	0.066	0.694	0.018
Ridge	1.551	0.128	0.639	0.036	1.358	0.073	0.694	0.018
Lasso	1.488	0.119	0.661	0.034	1.408	0.073	0.679	0.020
Elastic-Net	1.490	0.121	0.660	0.034	1.402	0.075	0.681	0.019
DNN	0.619	0.682	0.873	0.137	0.188	0.317	0.961	0.068
DNN+ ℓ_1	0.445	0.351	0.909	0.081	0.129	0.234	0.974	0.050
Random Forest	0.419	0.072	0.922	0.016	0.425	0.037	0.920	0.008
DeepBiome	0.243	0.400	0.950	0.087	0.117	0.244	0.976	0.052

The associated taxa are clustered at the phylum and order levels

DNN deep neural network, *DNN* + ℓ_1 Lasso (least absolute shrinkage and selection operator) penalized deep neural network

Table 7

Scenario 4: mean squared error (MSE) and Pearson correlation coefficient between predicted and true outcome (continuous), when using an mis-specified phylogenetic tree

Method	Testing				Training			
	MSE		Correlation		MSE		Correlation	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Linear Regression	0.872	0.163	0.683	0.046	0.737	0.074	0.726	0.027
Lasso	0.826	0.144	0.706	0.047	0.780	0.080	0.710	0.030
Ridge	0.866	0.138	0.683	0.046	0.752	0.081	0.726	0.027
Elastic-Net	0.826	0.144	0.706	0.047	0.779	0.079	0.711	0.028
DNN	0.437	0.208	0.849	0.077	0.167	0.166	0.944	0.058
DNN+ ℓ_1	0.434	0.214	0.850	0.080	0.166	0.171	0.944	0.065
Random Forest	0.194	0.050	0.939	0.019	0.198	0.025	0.937	0.009
DeepBiome	0.316	0.261	0.892	0.094	0.195	0.207	0.933	0.075

The associated taxa are clustered at the phylum and order levels

DNN deep neural network, $DNN + \ell_1$ Lasso (least absolute shrinkage and selection operator) penalized deep neural network

Table 8 Predict Type 2 Diabetes (T2D) status using microbiome counts in the metagenomics sequencing study

Method	Testing						Training															
	Sensitivity		Specificity		g-Measure		ACC		AUC		Sensitivity		Specificity		g-Measure		ACC		AUC			
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD		
Logistic	0.533	0.067	0.509	0.086	0.517	0.034	0.526	0.021	0.521	0.029	1.000	0.000	1.000	0.000	1.000	0.000	1.000	0.000	1.000	0.000	1.000	0.000
Lasso	0.331	0.332	0.819	0.183	0.364	0.335	0.558	0.107	0.575	0.076	0.330	0.320	0.884	0.118	0.395	0.364	0.615	0.093	0.607	0.103	0.607	0.103
Ridge	0.167	0.373	0.892	0.241	0.124	0.277	0.517	0.082	0.529	0.066	0.187	0.418	0.935	0.146	0.159	0.355	0.569	0.134	0.561	0.136	0.561	0.136
Elastic-Net	0.409	0.284	0.785	0.156	0.476	0.272	0.578	0.108	0.597	0.067	0.409	0.292	0.785	0.090	0.528	0.310	0.66	0.089	0.653	0.102	0.653	0.102
DNN	0.585	0.174	0.647	0.184	0.596	0.075	0.616	0.070	0.681	0.054	0.767	0.118	0.792	0.106	0.773	0.023	0.78	0.020	0.853	0.026	0.853	0.026
DNN+ ℓ_1	0.582	0.199	0.680	0.195	0.606	0.100	0.625	0.086	0.678	0.079	0.742	0.106	0.808	0.114	0.768	0.029	0.776	0.026	0.856	0.034	0.856	0.034
Random Forest	0.520	0.020	0.710	0.020	0.610	0.010	0.590	0.010	0.610	0.010	0.640	0.050	0.700	0.050	0.670	0.041	0.670	0.040	0.670	0.039	0.670	0.039
DeepBiome	0.640	0.180	0.645	0.200	0.620	0.066	0.643	0.054	0.694	0.037	0.836	0.142	0.888	0.086	0.856	0.052	0.863	0.046	0.948	0.015	0.948	0.015