

# Managing Variable Dependence in Eigenvector Selection for Phylogenetic Eigenvector Regression

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## Description

Phylogenetic Eigenvector Regression (PVR) is a powerful tool used in evolutionary biology to account for phylogenetic relationships among species when examining ecological and evolutionary patterns. This method leverages eigenvectors derived from a phylogenetic distance matrix to capture the phylogenetic structure within a dataset, allowing researchers to isolate and analyze the influence of phylogeny on various traits. A critical aspect of PVR is the selection of these eigenvectors, as the choice can significantly impact the model's explanatory power and interpretability. One of the primary challenges in this context is managing variable dependence in eigenvector selection, which can complicate the analysis and interpretation of results. Eigenvectors in PVR are derived from a phylogenetic distance matrix, which encapsulates the evolutionary relationships among the species under study. These eigenvectors represent orthogonal axes in a multidimensional space, each capturing different aspects of phylogenetic variation. Ideally, the selected eigenvectors should account for the major sources of phylogenetic signal in the data. However, eigenvectors are often correlated due to the shared evolutionary history they represent. This interdependence among eigenvectors can lead to redundancy, where multiple eigenvectors may capture overlapping phylogenetic information, potentially skewing the regression results and reducing the model's efficiency. To address variable dependence in eigenvector selection, researchers typically employ criteria that prioritize eigenvectors based on their explanatory power. One common approach is to select eigenvectors corresponding to the largest eigenvalues, as these typically explain the most significant portions of phylogenetic variation. However, this method does not entirely eliminate the issue of dependence, as eigenvectors with large eigenvalues can still be correlated. Advanced selection techniques, such as stepwise selection or penalized regression methods like Lasso, can help mitigate this by introducing regularization and thereby reducing the influence of collinear eigenvectors. These methods aim to balance model complexity with explanatory power, ensuring that the chosen eigenvectors provide distinct and complementary insights into phylogenetic relationships. Another approach to managing variable dependence involves the use of statistical criteria that assess the contribution of each eigenvector to the overall model fit. Measures such as the Akaike Information Criterion (AIC) or Bayesian Information Criterion (BIC) can be applied to select a subset of eigenvectors that optimally balance goodness-of-fit with model simplicity. By penalizing the inclusion of redundant or marginally informative eigenvectors, these criteria help in constructing more parsimonious models. Additionally, cross-validation techniques can be employed to validate the robustness of the selected eigenvectors and ensure that the model generalizes well to new data. Incorporating biological insights into the eigenvector selection process can also be beneficial. For example, researchers can prioritize eigenvectors that correspond to known evolutionary splits or significant phylogenetic events. This biologically informed selection can enhance the interpretability of the results and align the statistical model with biological reality. Moreover, this approach can reduce the likelihood of overfitting by focusing on the most ecologically and evolutionarily relevant sources of phylogenetic variation. The implications of variable dependence in eigenvector selection extend beyond statistical concerns, influencing the biological conclusions drawn from PVR analyses. Overlooking variable dependence can lead to overestimation or underestimation of phylogenetic effects, thereby misinforming ecological and evolutionary inferences. For instance, if redundant eigenvectors are included, the model might overemphasize certain phylogenetic signals, leading to biased estimates of trait evolution or ecological interactions.

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## Conflict of Interest

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