NPDR Supplementary Material

Trang T. Le 1 , Bryan A. Dawkins 2 and Brett A. McKinney 2,3*

 ¹Department of Biostatistics, Epidemiology and Informatics, University of Pennsylvania, Philadelphia, PA 19104
²Department of Mathematics, University of Tulsa, Tulsa, OK 74104
³Tandy School of Computer Science, University of Tulsa, Tulsa, OK 74104

June 17, 2019

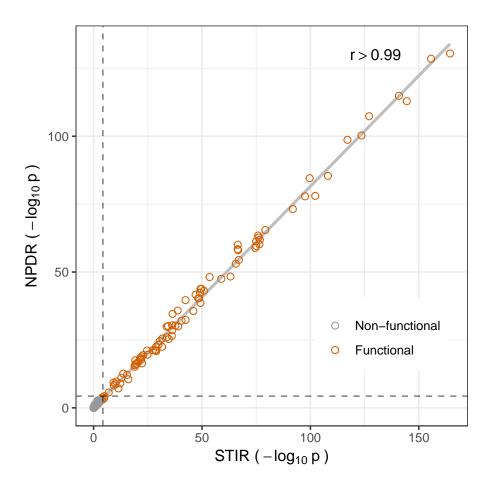
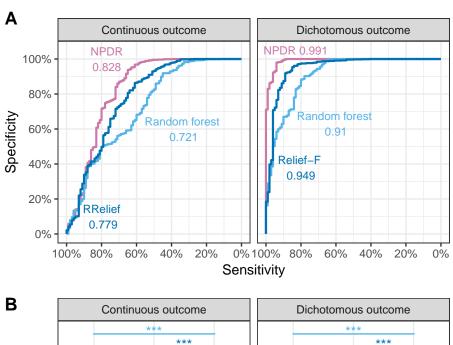


Figure S1: Similarity between NPDR and STIR (dichotomous outcomes). Comparison of $-\log_{10}$ P values for one interaction simulation of m=200 (100 cases and 100 controls) and p=1000 attributes with 100 functional. In 100 replicate simulations, correlation, r, between the two methods ranges from 0.9827 to 0.9994. STIR is based on a t-test of projected distances and NPDR is based on a logistic regression of projected distances. NPDR has the added benefit of handling continuous outcomes and covariate correction.



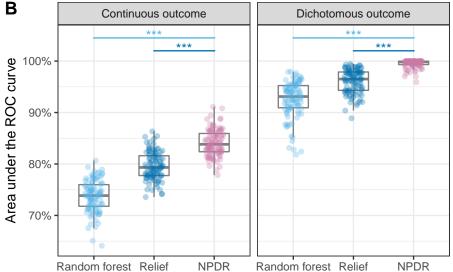


Figure S2: Receiver Operating Characteristic (ROC) curves for Relief, NPDR and random forest feature selection. For one replicate simulation (A), ROCs for continuous outcome data with main effects (left) and dichotomous outcome data with interaction effects (right). The auROC value is given for each method. For 100 replicate simulations of both simulation types (B), NPDR yields statistically significant higher auROC than Relief or random forest (*** indicate P < .0001). All simulations use m = 200 samples and p = 1,000 attributes with 100 functional.