

Supplementary Figures to A General Framework for Integrative Analysis of Incomplete Multi-omics Data

Dan-Yu Lin, Donglin Zeng, and David Couper

Department of Biostatistics, University of North Carolina, Chapel Hill, NC 27599-7420

Figure Legends

Figure S1. Quantile-quantile plots for testing $\beta = 0$ under the Joint Model when the detection limit is 1.

Figure S2. Simulation results under the Joint Model when the biomarker has the t -distribution with 5 degrees of freedom. The red, black, green, and blue curves pertain to the proposed method, complete-case analysis, imputation at limit, and imputation at mid-point, respectively.

Figure S3. Simulation results under the Joint Model when the biomarker is the logarithm of the standard exponential random variable. The red, black, green, and blue curves pertain to the proposed method, complete-case analysis, imputation at limit, and imputation at mid-point, respectively.

Figure S4. Simulation results under the Joint Model when the biomarker has the standard normal distribution and the inverse-normal transformation is performed on the observed values. The red, black, green, and blue curves pertain to the proposed method, complete-case analysis, imputation at limit, and imputation at mid-point, respectively.

Figure S5. Simulation results under the Joint Model with $\beta = 0$ when the biomarker has 30% missing values: (A) effect of the SNP genotype on the quantitative omics variable (i.e., α_G); (B) effect of the quantitative omics variable on the phenotype (i.e., γ); and (C) effect of the genotype on the phenotype (i.e., β_G). The bias and standard error of the

parameter estimator and the power or type I error of the association test are plotted against the detection limit of the quantitative omics variable. The red, black, green, and blue curves pertain to the proposed method, complete-case analysis, imputation at limit, and imputation at mid-point, respectively. The silver curve pertains to the mean of the standard error estimator for the proposed method.

Figure S6. Simulation results under the Joint Model with $\beta = 0.2$ when the biomarker has 30% missing values: (A) effect of the SNP genotype on the quantitative omics variable (i.e., α_G); (B) effect of the quantitative omics variable on the phenotype (i.e., γ); and (C) effect of the genotype on the phenotype (i.e., β). The bias and standard error of the parameter estimator and the power of the association test are plotted against the lower detection limit of the quantitative omics variable. The red, black, green, and blue curves pertain to the proposed method, complete-case analysis, imputation at limit, and imputation at mid-point, respectively. The silver curve pertains to the mean of the standard error estimator for the proposed method.

Figure S7. Simulation results under the Joint Model with $\beta = 0.2$ when there are two SNPs in the model: (A) effect of the SNP with MAF of 0.4 on the quantitative omics variable (i.e., α_G); (B) effect of the quantitative omics variable on the phenotype (i.e., γ); and (C) effect of the SNP with MAF of 0.4 on the phenotype (i.e., β). The bias and standard error of the parameter estimator and the power of the association test are plotted against the lower detection limit of the quantitative omics variable. The red, black, green, and blue curves pertain to the proposed method, complete-case analysis, imputation at limit, and imputation at mid-point, respectively. The silver curve pertains to the mean of the standard error estimator for the proposed method.

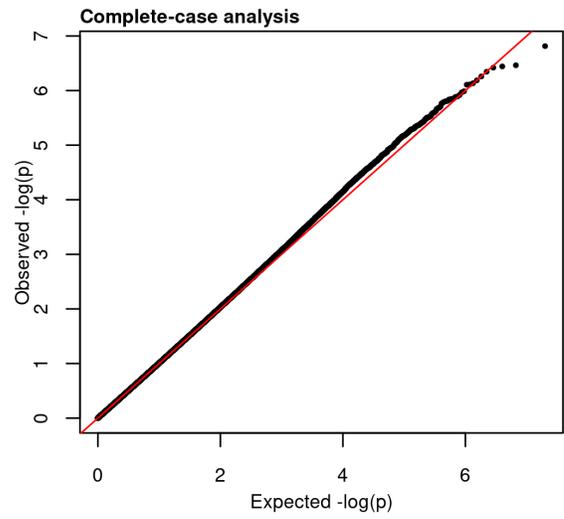
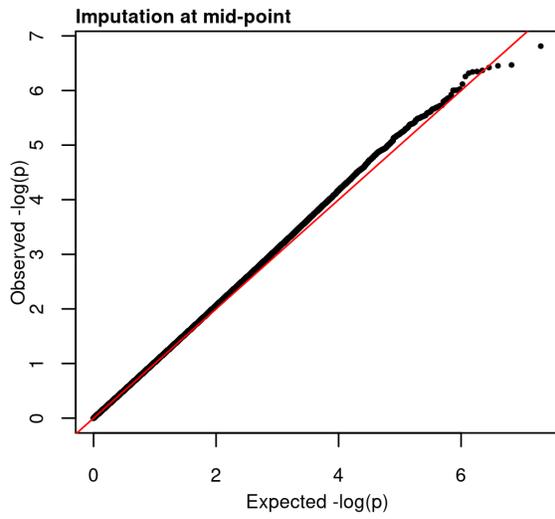
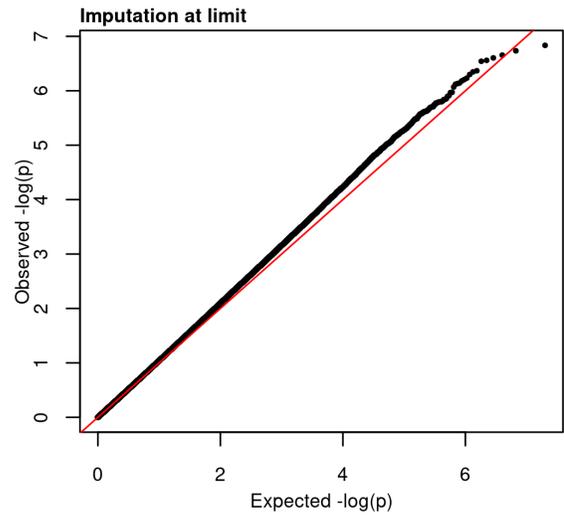
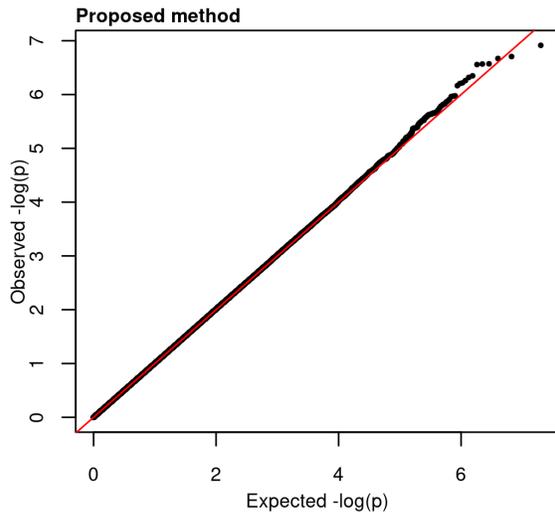


Figure S1

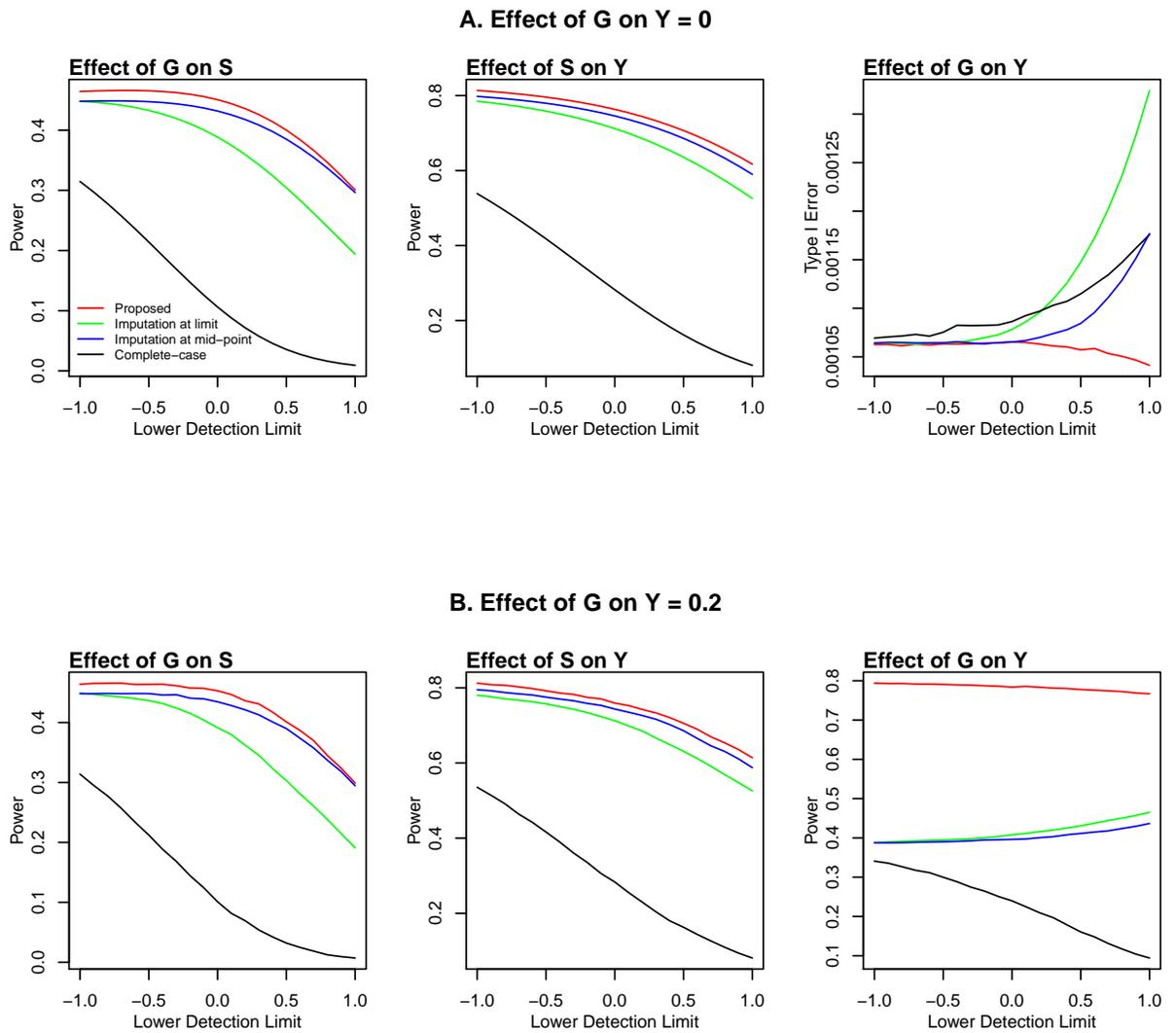


Figure S2

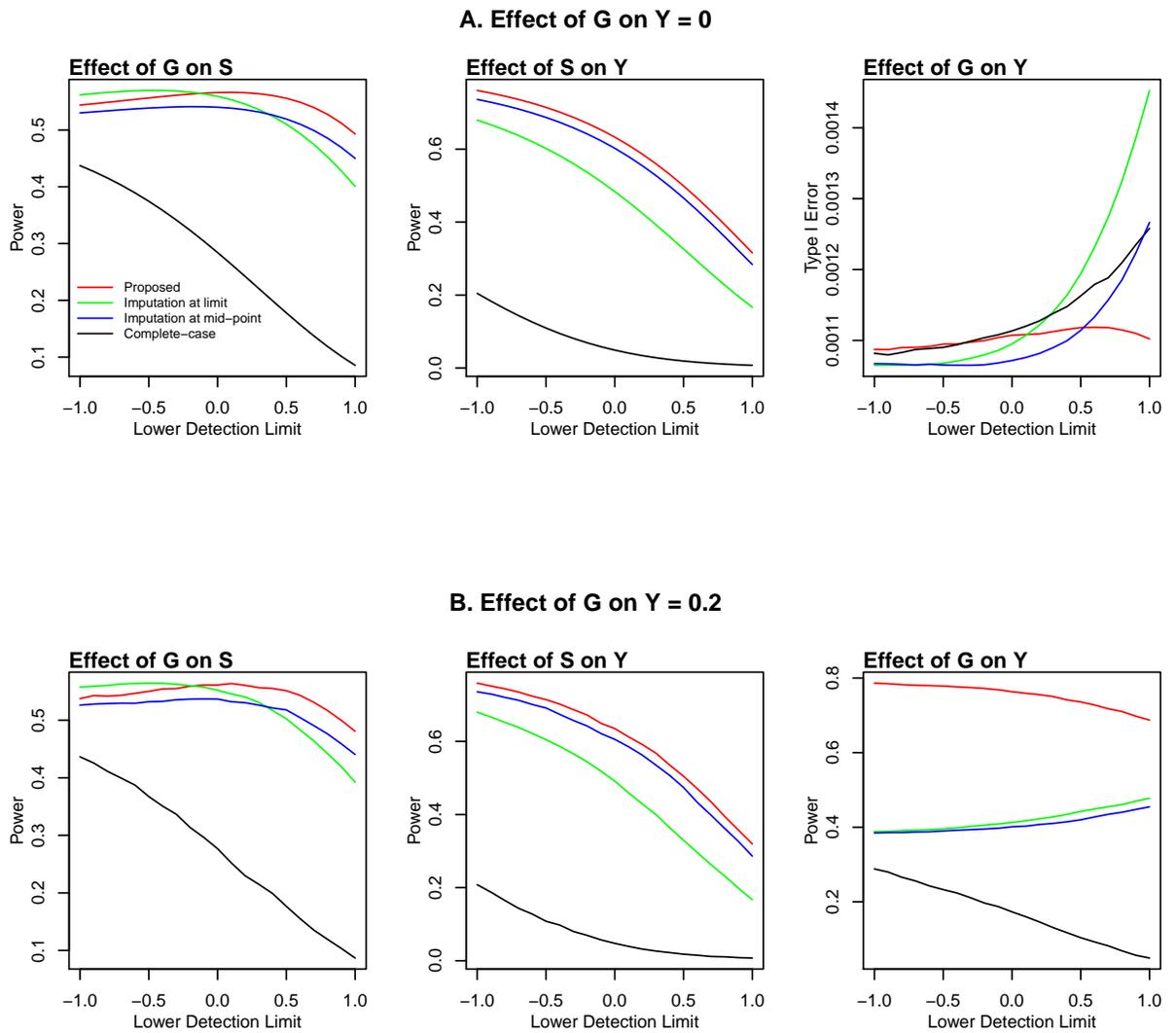


Figure S3

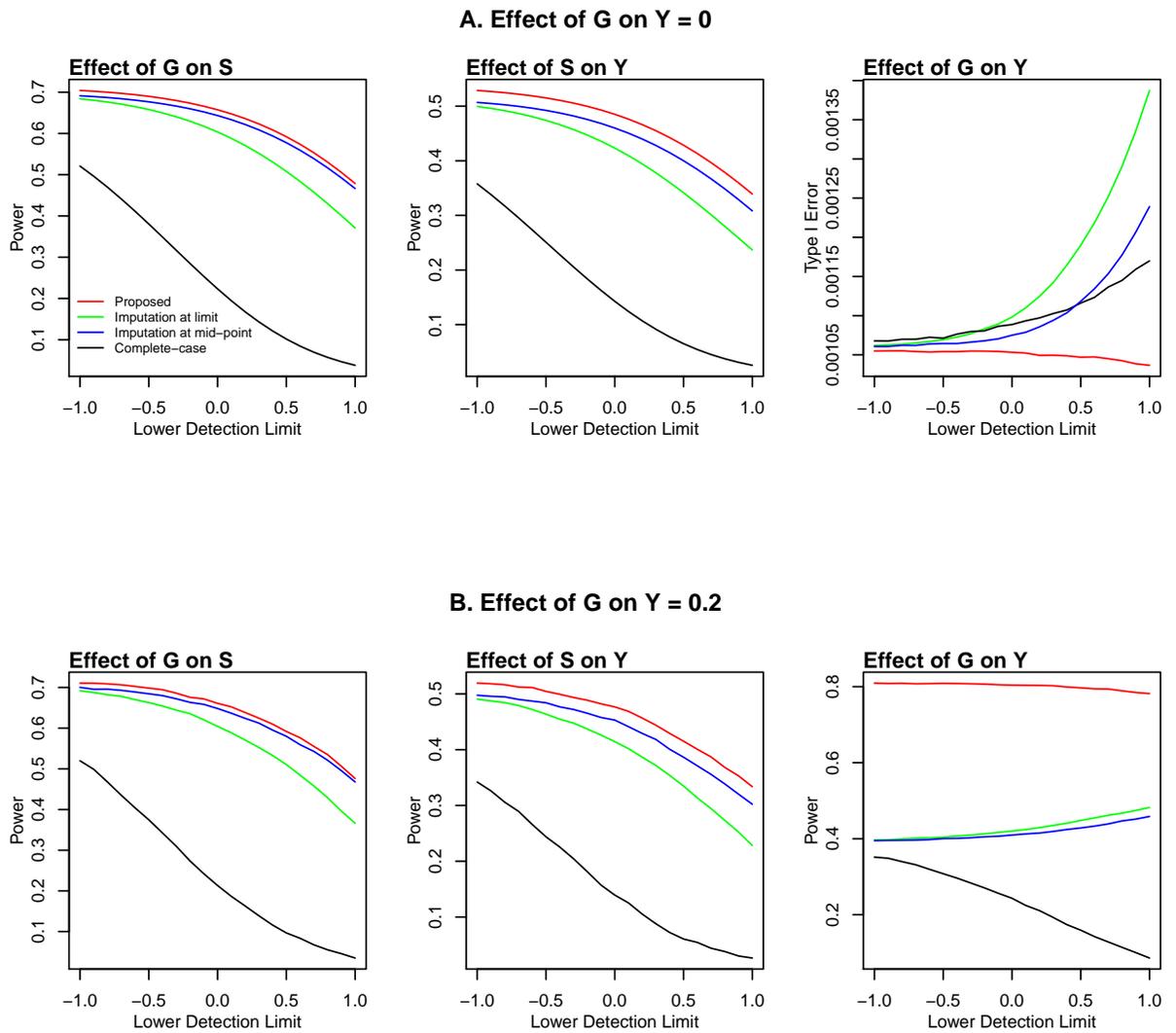


Figure S4

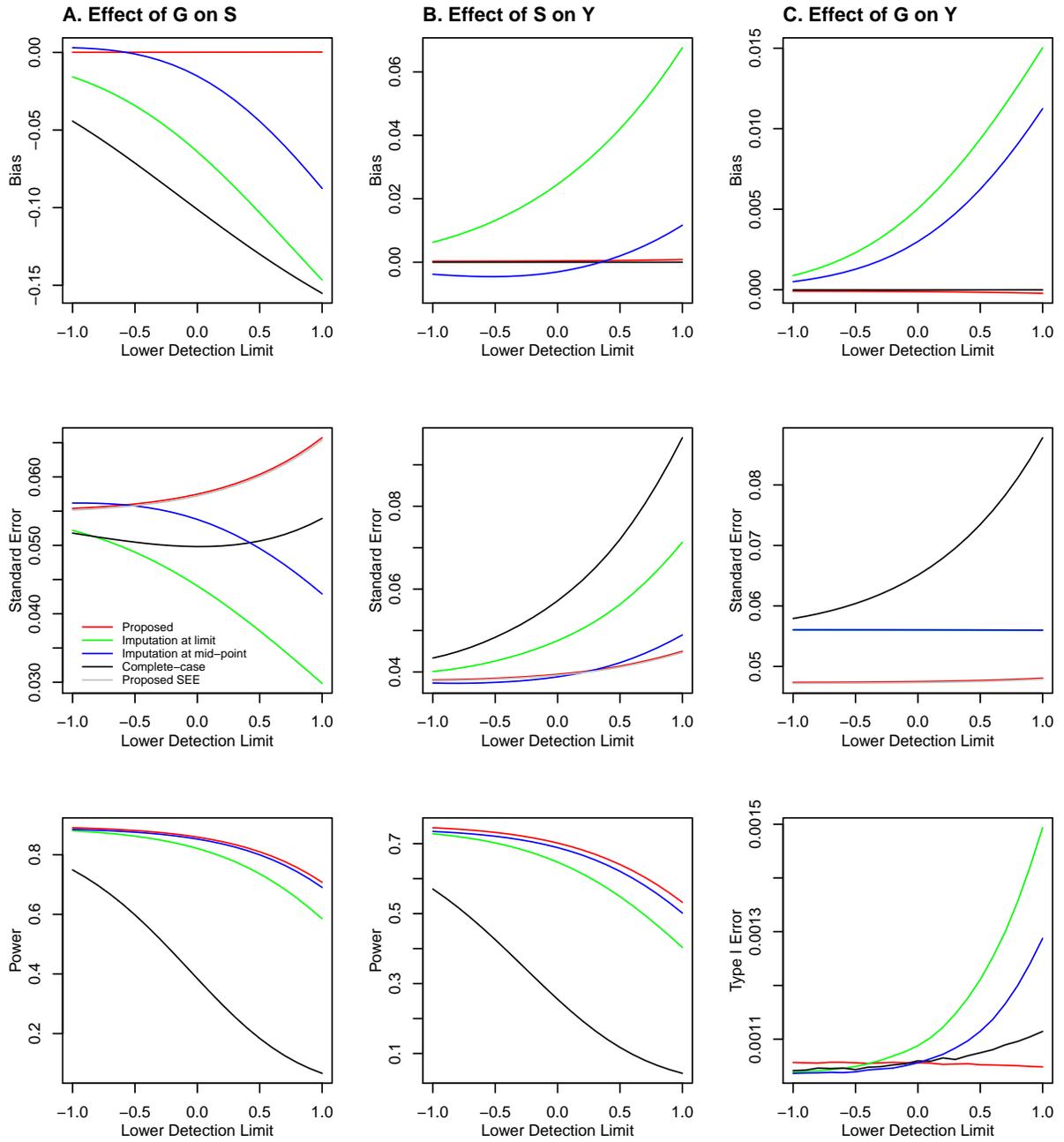


Figure S5

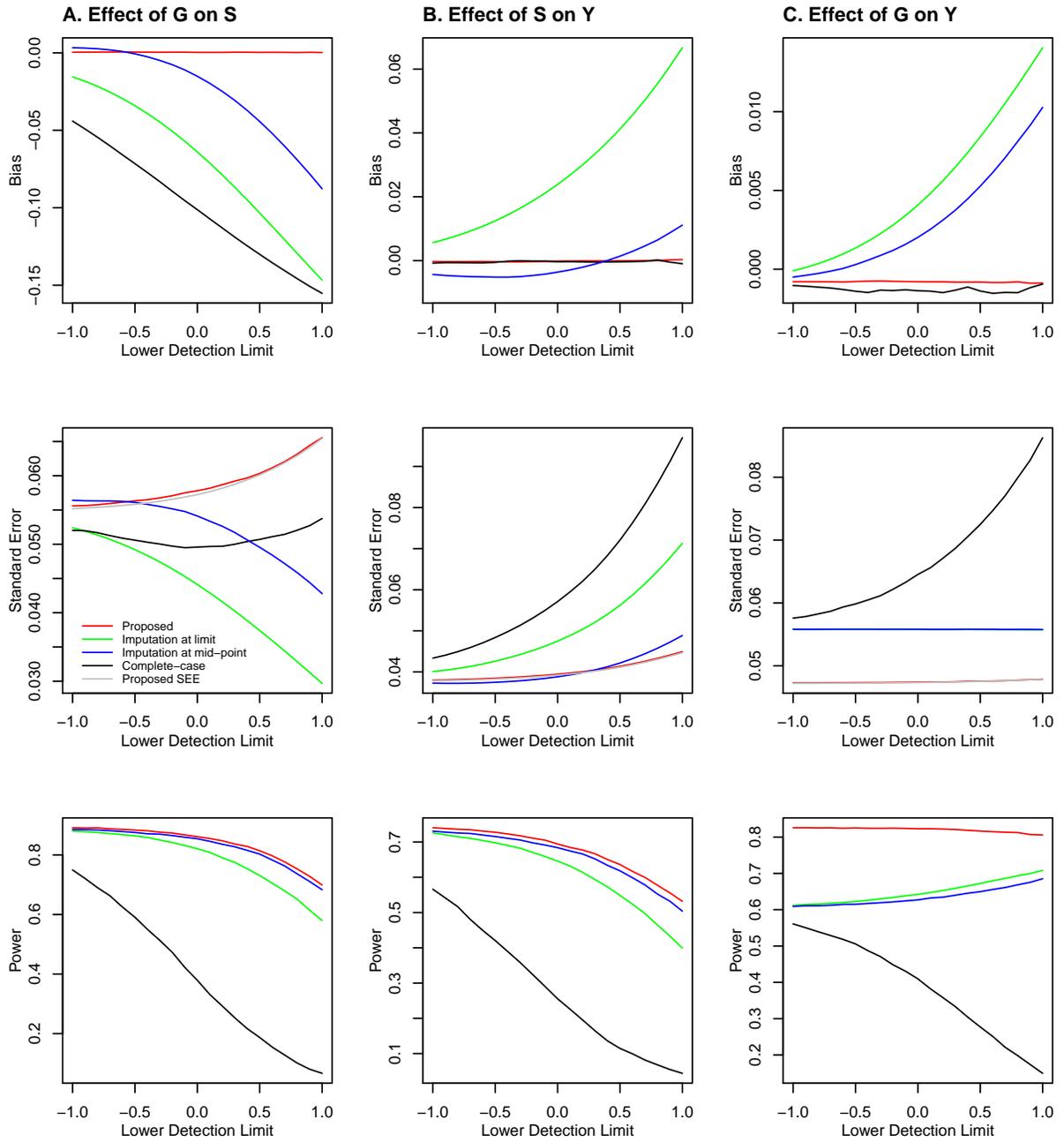


Figure S6

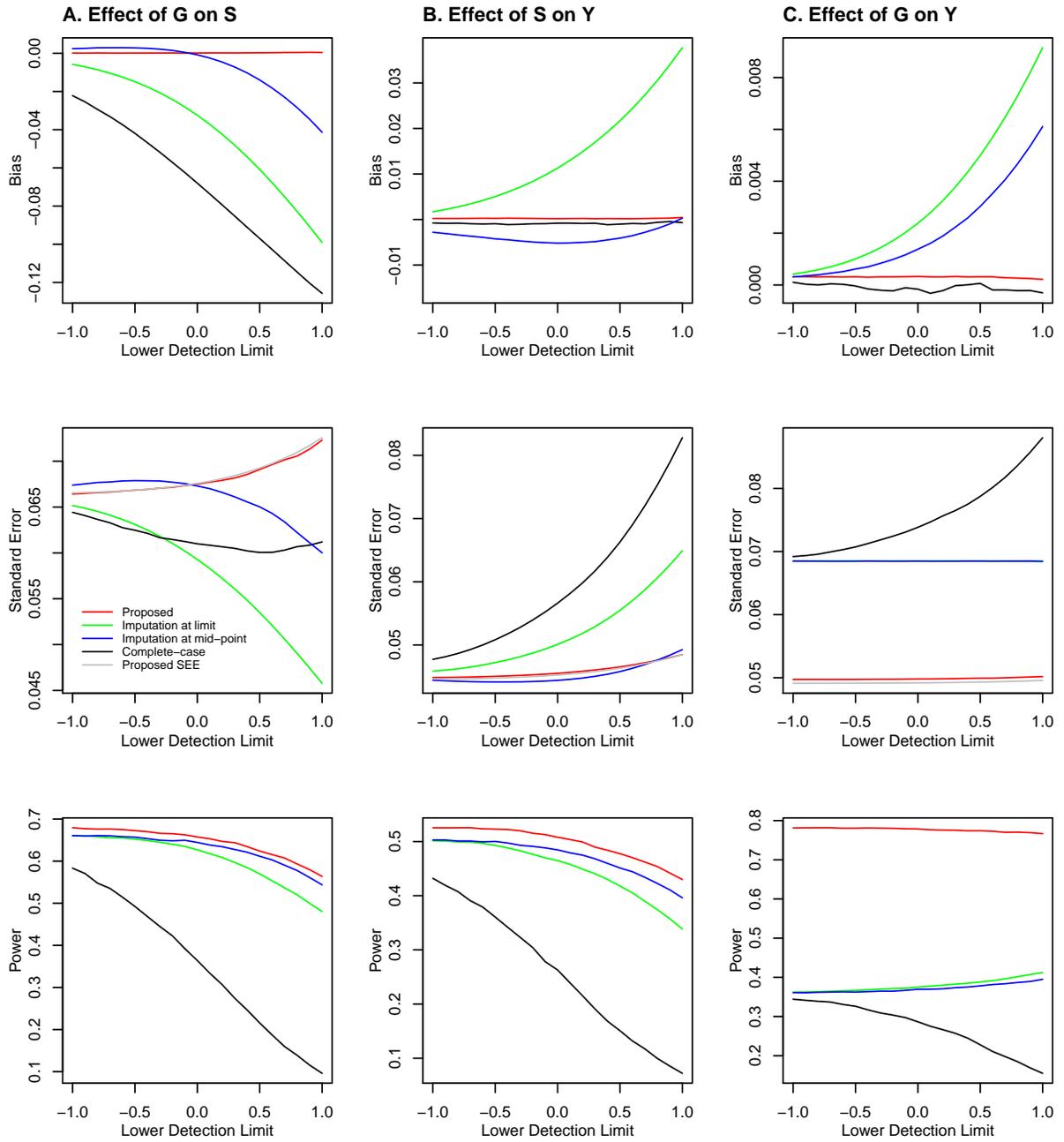


Figure S7