

Additional file 1 for “A sparse Bayesian factor model for the construction of gene co-expression networks from single-cell RNA sequencing count data”

Michael Sekula¹, Jeremy Gaskins¹, Susmita Datta²

¹ Department of Bioinformatics and Biostatistics, University of Louisville, KY 40202, USA

² Department of Biostatistics, University of Florida, FL 32610, USA

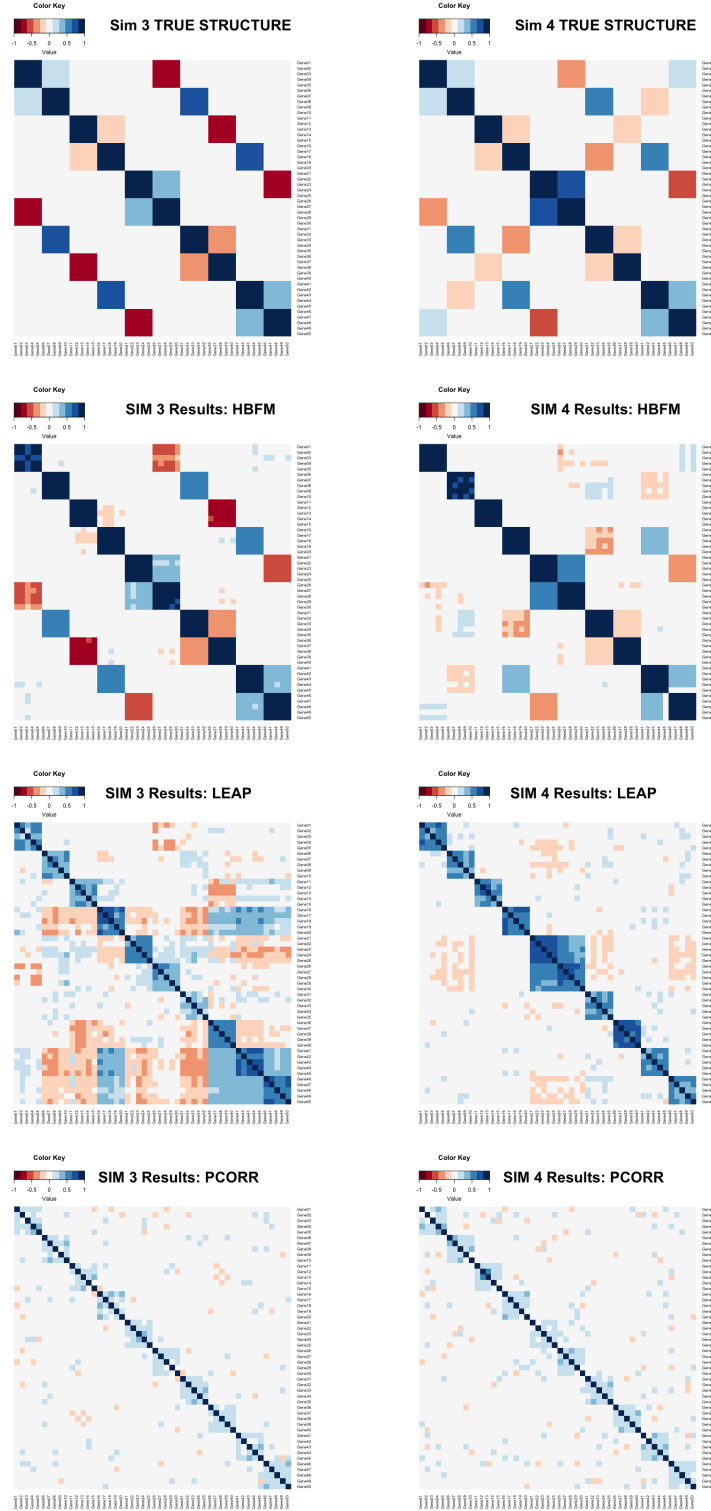


Figure S1: Heatmaps of the “true” correlation structure in Sim 3 ($F = 10, N = 500$) and Sim 4 ($F = 15, N = 500$) and the correlation structures estimated by HBFM, LEAP, and PCORR.

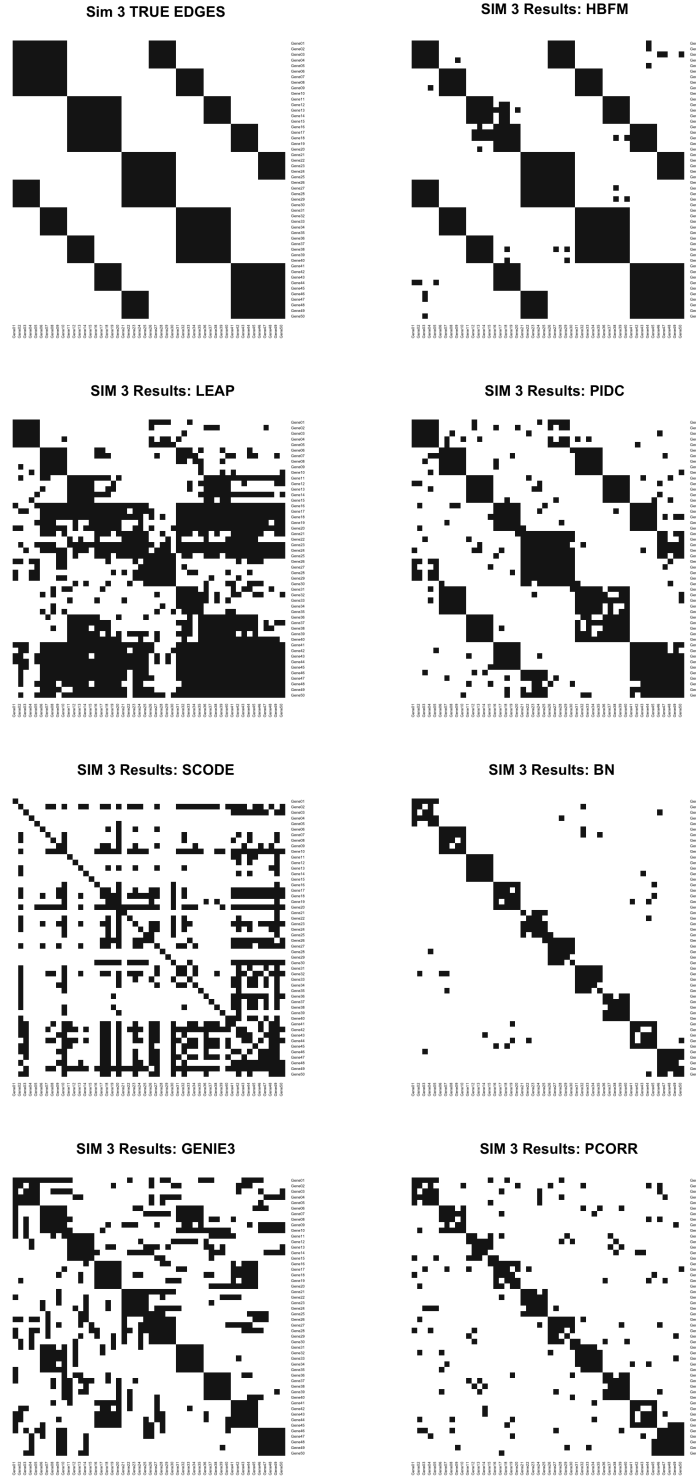


Figure S2: Heatmaps of the “true” network structure in Sim 3 (top left) and the network structures estimated by all network methods. The diagonal has been shaded in each figure to serve as a reference point. The shaded cells (other than the diagonal) indicate significant gene-gene associations. For the networks estimated by PIDC, SCODE, and GENIE3, the number of edges have been fixed to match the number of edges estimated by HBFM.

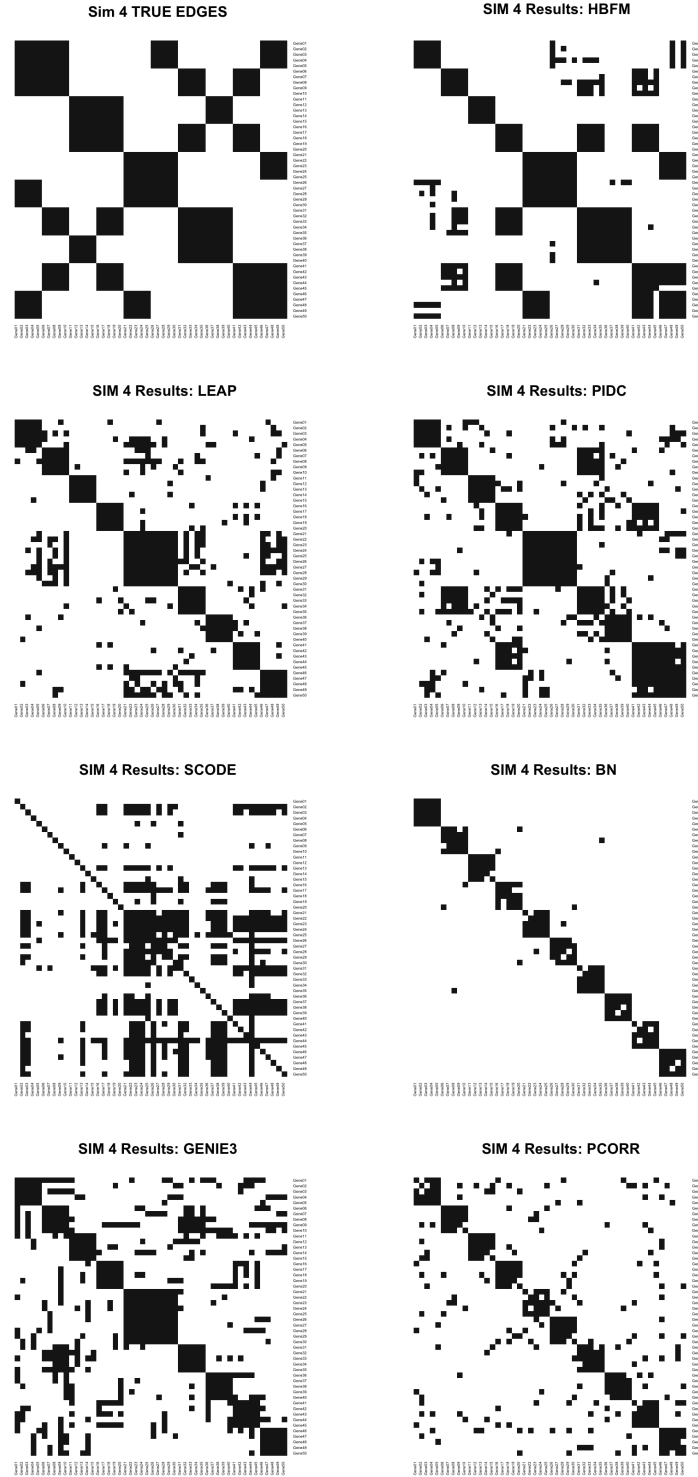


Figure S3: Heatmaps of the “true” network structure in Sim 4 (top left) and the network structures estimated by all network methods. The diagonal has been shaded in each figure to serve as a reference point. The shaded cells (other than the diagonal) indicate significant gene-gene associations. For the networks estimated by PIDC, SCODE, and GENIE3, the number of edges have been fixed to match the number of edges estimated by HBFM.

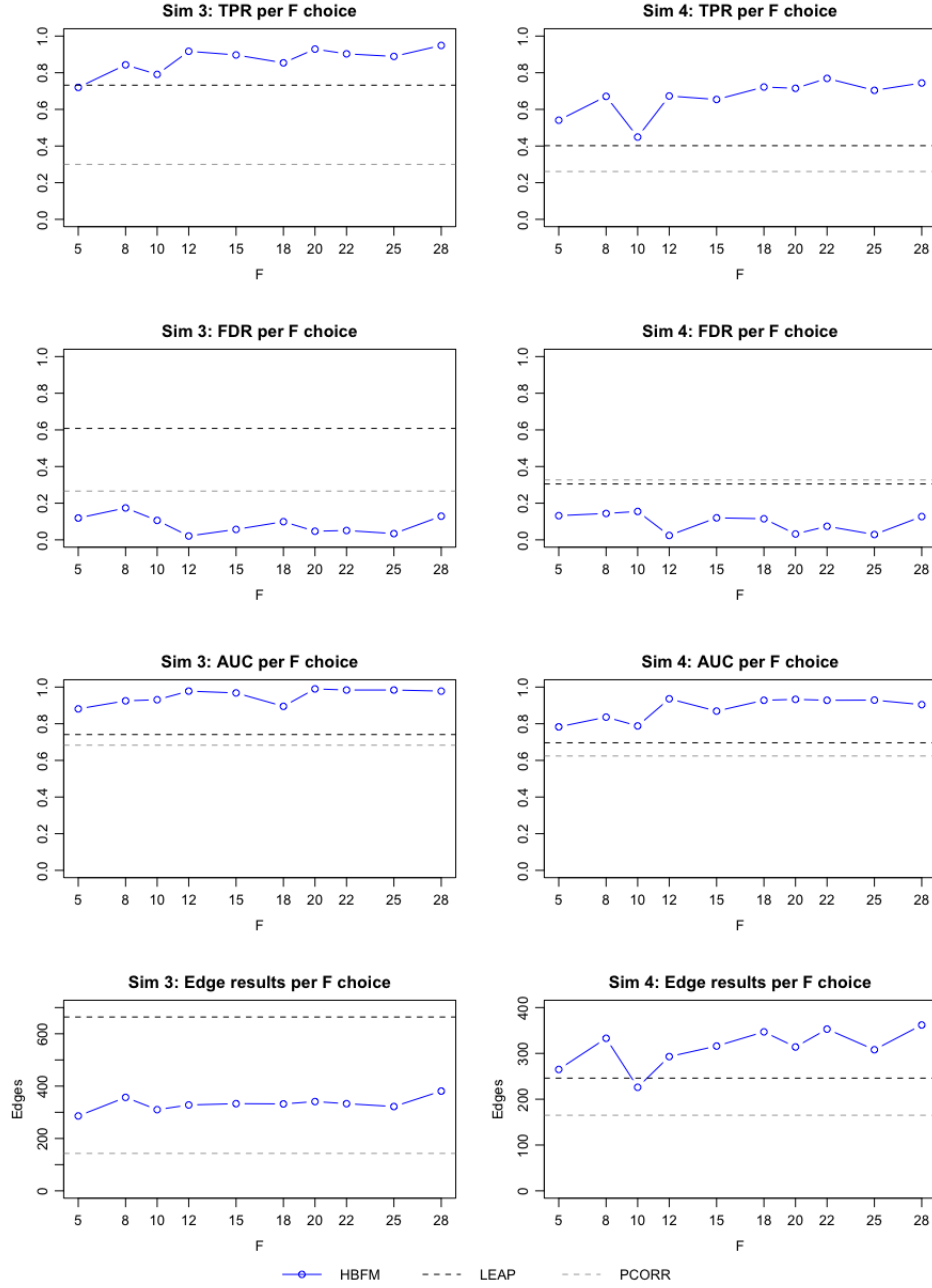


Figure S4: Performance measures for HBFM with different numbers of factors (F) for Sim 3 and Sim 4. The dotted lines represent the performance measures for LEAP and PCORR. Both LEAP and PCORR are correlation-based network methods and are the most similar to our proposed methodology.