

# Identifying potential cancer driver genes by genomic data integration

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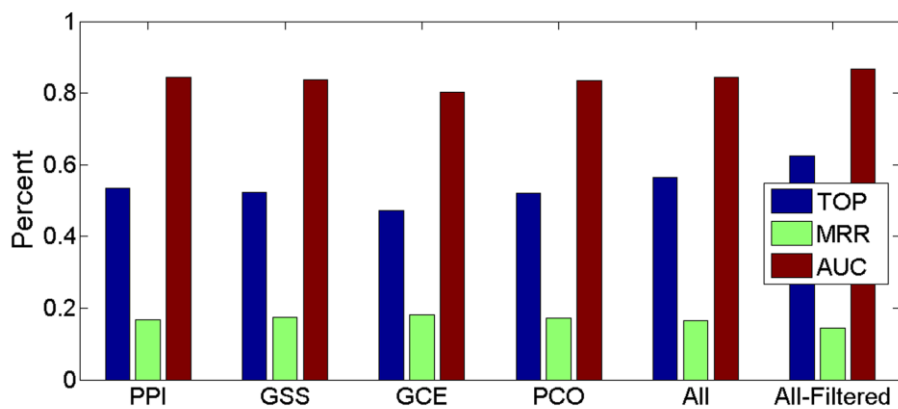
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## Supplementary figures and legends

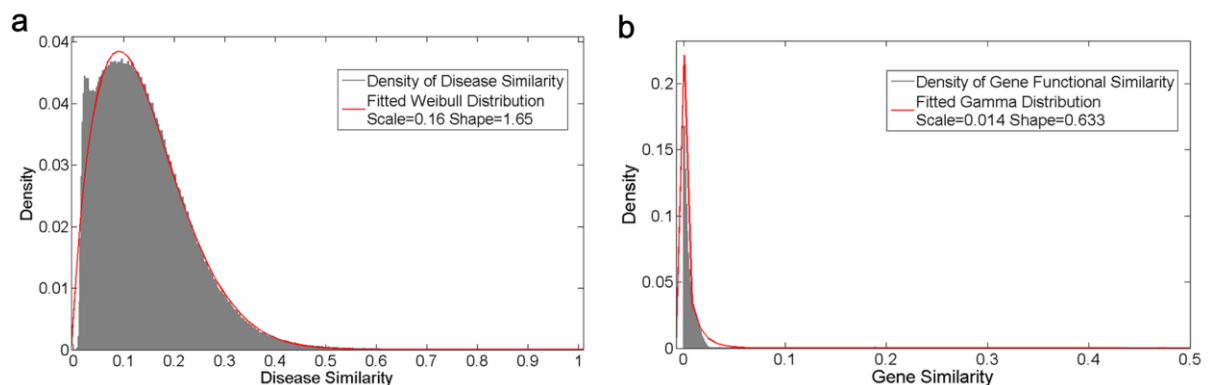
### Supplementary Figure S1. The performances of each data source.

The results were obtained from the validations against random controls based on the heterogeneous network. PPI: Protein-protein interactions. GCE: Gene co-expression patterns. GSS: Gene sequence similarities. PCO: Pathway co-occurrence relationships. All: All the four types omics data integrated, but without noise filtering. All-Filtered: All the four types omics data integrated and noise filtered ( $\beta = 0.25$  and  $\gamma = 0.19$ ). TOP: Top one ranked ratio of positive cases. MRR: The averaged relative ranks of all positive cases. AUC: The area under the receiver operating characteristic curves (ROC).



### Supplementary Figure S2. Statistical analysis of and disease phenotypic similarities and fused gene functional similarities.

Both distributions were fitted by using MATLAB Fitting Toolbox. (a) The density and fitted Weibull distributions of disease phenotypic similarities. (b) The density and fitted Gamma distributions of fused gene functional similarities.





## Supplementary Figure S5. Relationship analysis of 164 genes.

The data were analyzed and outputted by using STRING database.

