

Figure ia: BRCA Top Gain

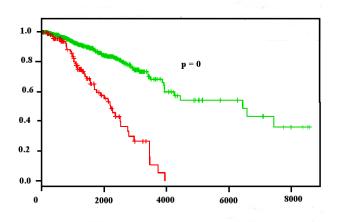


Figure ic: BRCA Top Lost

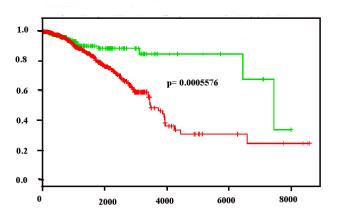


Figure ib: BRCA Specific Top Gain

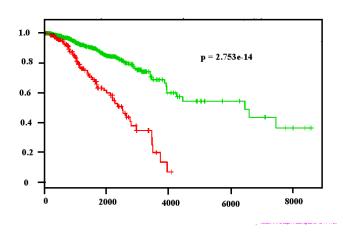


Figure id: BRCA Specific Top Lost

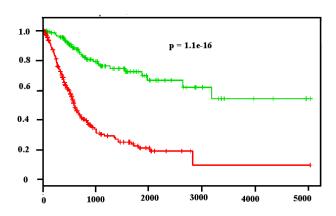


Figure iia: BLCA Top Gain

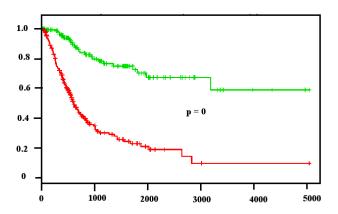


Figure iic: BLCA Top Lost

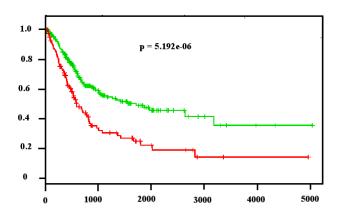


Figure iib: BLCA Specific Top Gain

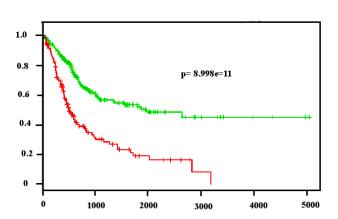


Figure iid: BLCA Specific Top Lost

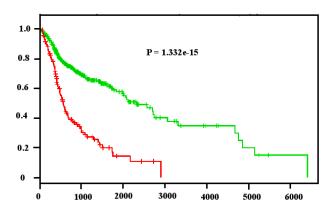


Figure iiia: HNSC Top Gain

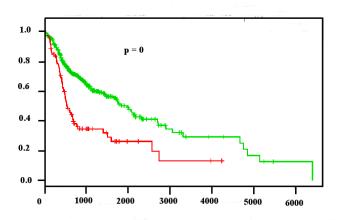


Figure iiic: HNSC Top Lost

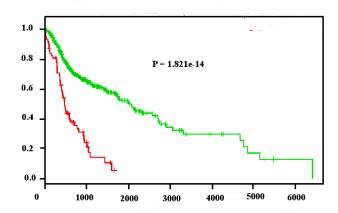


Figure iiib: HNSC Specific Top Gain

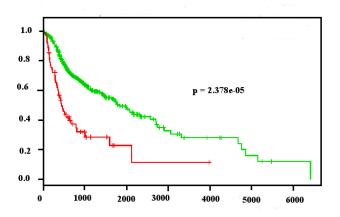


Figure iiid: HNSC Specific Top Lost

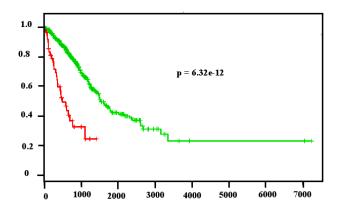


Figure iva: LUAD Top Gain

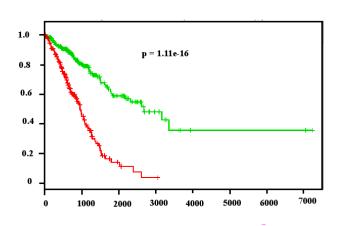


Figure ivc: LUAD Top Lost

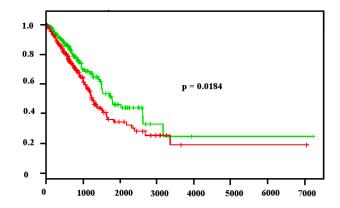


Figure ivb: LUAD Specific Top Gain

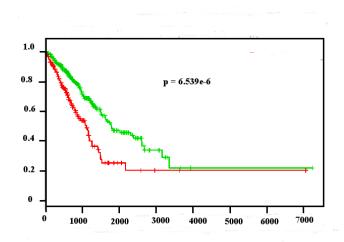


Figure ivd: LUAD Specific Top Lost

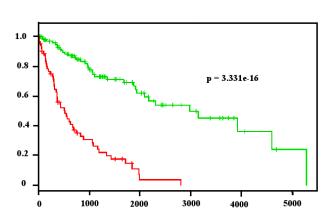


Figure va: LUSC Top Gain

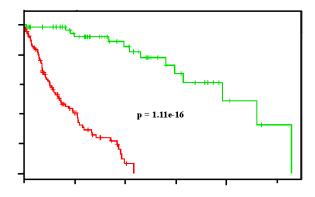


Figure vc: LUSC Top Lost

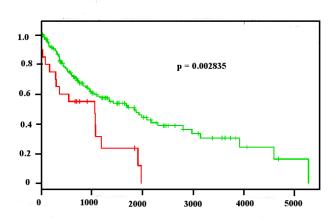


Figure vb: LUSC Specific Top Gain

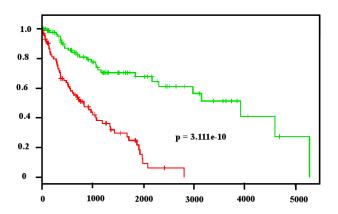


Figure vd: LUSC Specific Top Lost

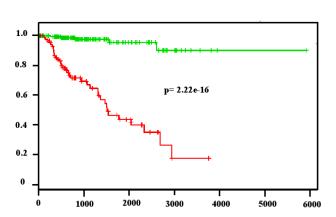


Figure via: KIRP Top Gain

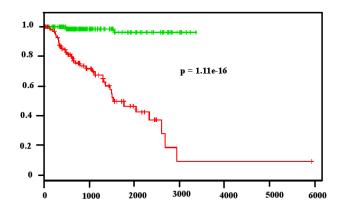


Figure vic: KIRP Top Lost

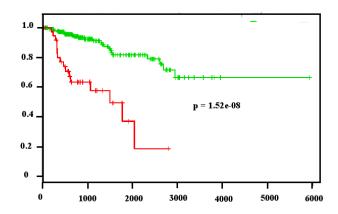


Figure vib: KIRP Specific Top Gain

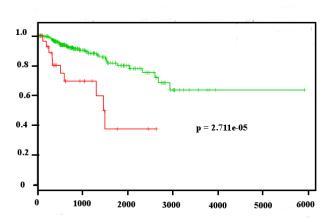


Figure vid: KIRP Specific Top Lost

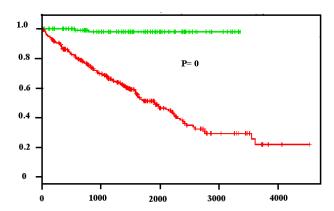


Figure viia: KIRC Top Gain

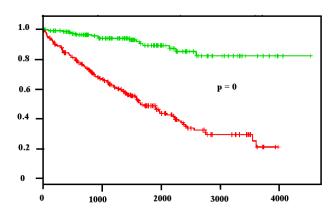


Figure viic: KIRC Top Lost

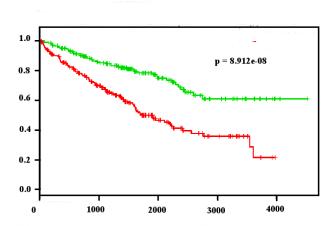


Figure viib: KIRC Specific Top Gain

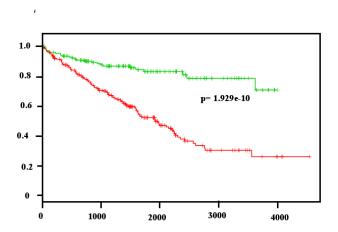


Figure viid: KIRC Specific Top Lost

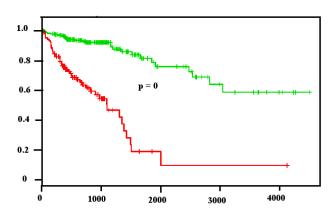


Figure viiia: COAD Top Gain

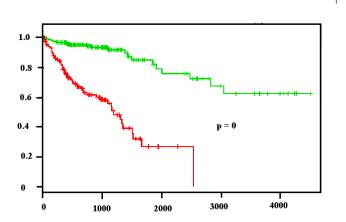


Figure viiic: COAD Top Lost

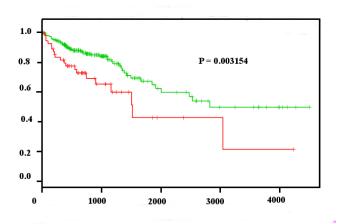


Figure viiib: COAD Specific Top Gain

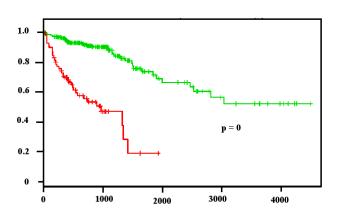


Figure viiid: COAD Specific Top Lost

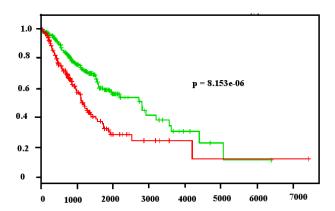


Figure ixa: STES Top Gain

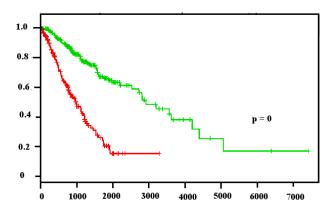


Figure ixc: STES Top Lost

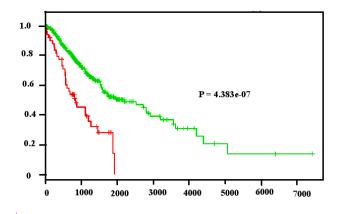


Figure ixb: STES Specific Top Gain

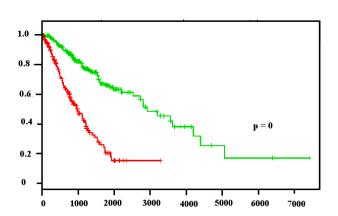


Figure ixd: STES Specific Top Lost

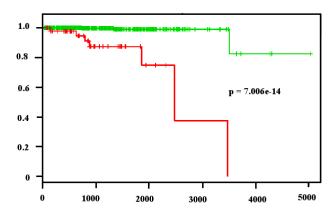


Figure xa: PRAD Top Gain

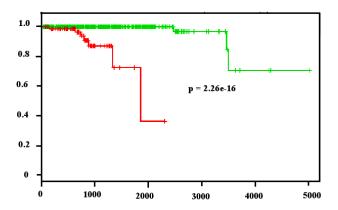


Figure xc: PRAD Top Lost

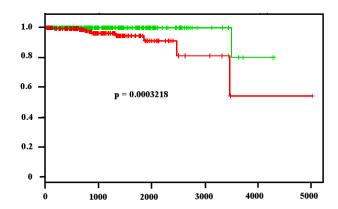


Figure xb: PRAD Specific Top Gain

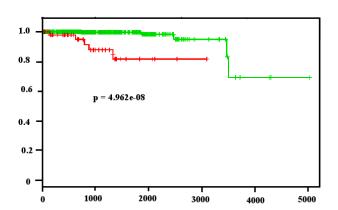


Figure xd: PRAD Specific Top Lost

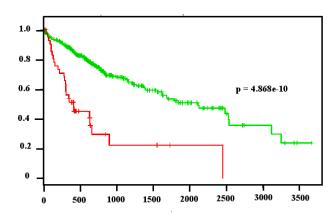


Figure xiia: LIHC Specific Top Gain

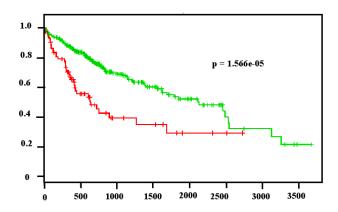


Figure xiib: LIHC Top Lost

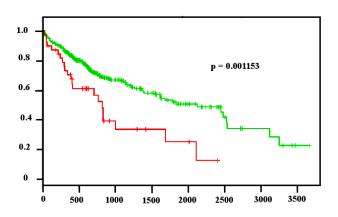


Figure xiic: LIHC Specific Top Lost

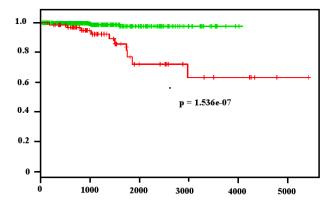
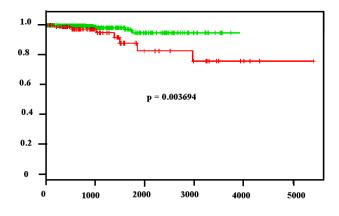


Figure xiiia: THCA Top Gain

Figure xiiib: THCA Specific Top Gain



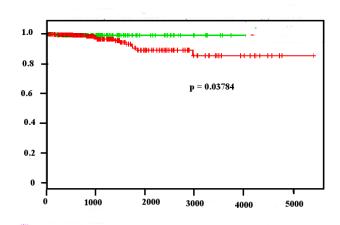


Figure xiiic: THCA Top Lost

Figure xiiid: THCA Specific Top Lost

Supplementary Figure 3 (i - xiiic): Kaplan-Meier survival analysis plots of multigene cancer biomarkers involved in edgetic perturbations. The x axes indicate the number of days until patient death whereas the y axes indicate the probability of patient survival. In all the Figures, the green lines indicate better survival (longer life-span) after cancer diagnosis while the red lines indicate poor survival (shorter life-span) after cancer diagnosis as a result of the proteins involved in edgetic gains or losses. In all the cases, the proteins involved in edgetic perturbations predicted poor survival of the patients (Logrank test p-value < 0.05), indicating their importance in cancer monitoring and prognosis. (a) Overall survival predicted from gene signatures involved in edgetic gains across most patients of a cancer type (except for LIHC), (b) Overall survival predicted from gene signatures involved in edgetic gains across patients showing cancer-specific perturbations, (d) Overall survival predicted from gene signatures involved in edgetic from gene signatures involved in edgetic losses across patients showing cancer-specific perturbations. The names of the prominent proteins with multiple perturbations responsible for the above observations can be found in Supplementary File 7.xls (Tables 7a and 7b).