



Supplementary Fig. S3: Regions of somatic copy number aberration across cancers and genes dysregulated due to copy number. (A) Plots of the frequency of copy number gain and loss across the genome for four pediatric cancer cohorts in this study. **(B)** IncRNA loci on chromosomes with copy number alterations across the pediatric cancers: NBL, WT, B-ALL, and AML. IncRNAs were evaluated to have differential expression due to copy number using the Wilcoxon rank sum test: highly differential: p -value < 0.05 and $\log |\text{fold change}| > 1.5$ and moderately differential: p -value < 0.1 and $\log |\text{fold change}| > 1.0$. Points are colored based on loci in an amplified or deleted region of the chromosome and if the IncRNA is highly or moderately differentially expressed.