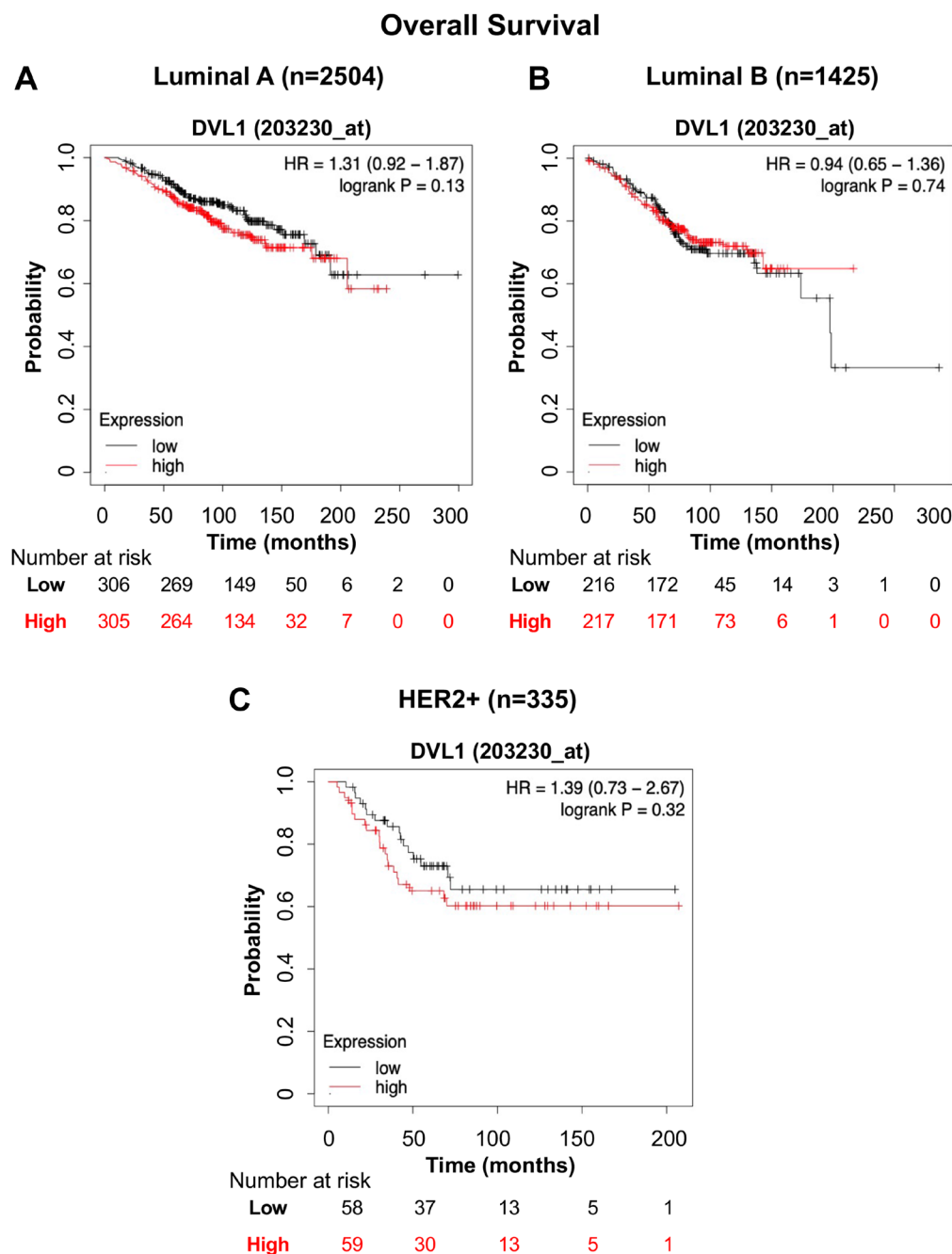
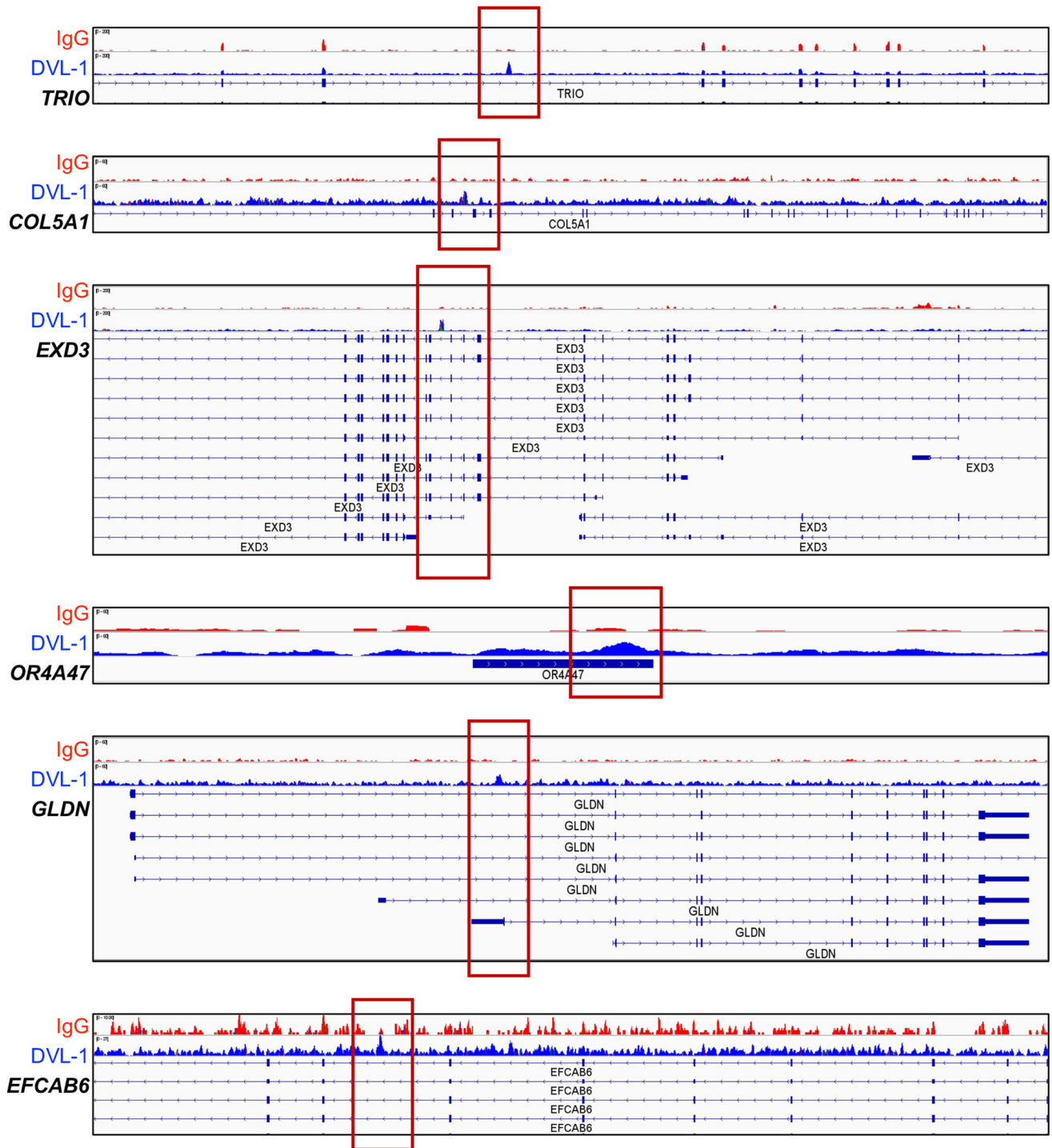


Genomic profiling of DVL-1 and its nuclear role as a transcriptional regulator in triple negative breast cancer

SUPPLEMENTARY MATERIALS



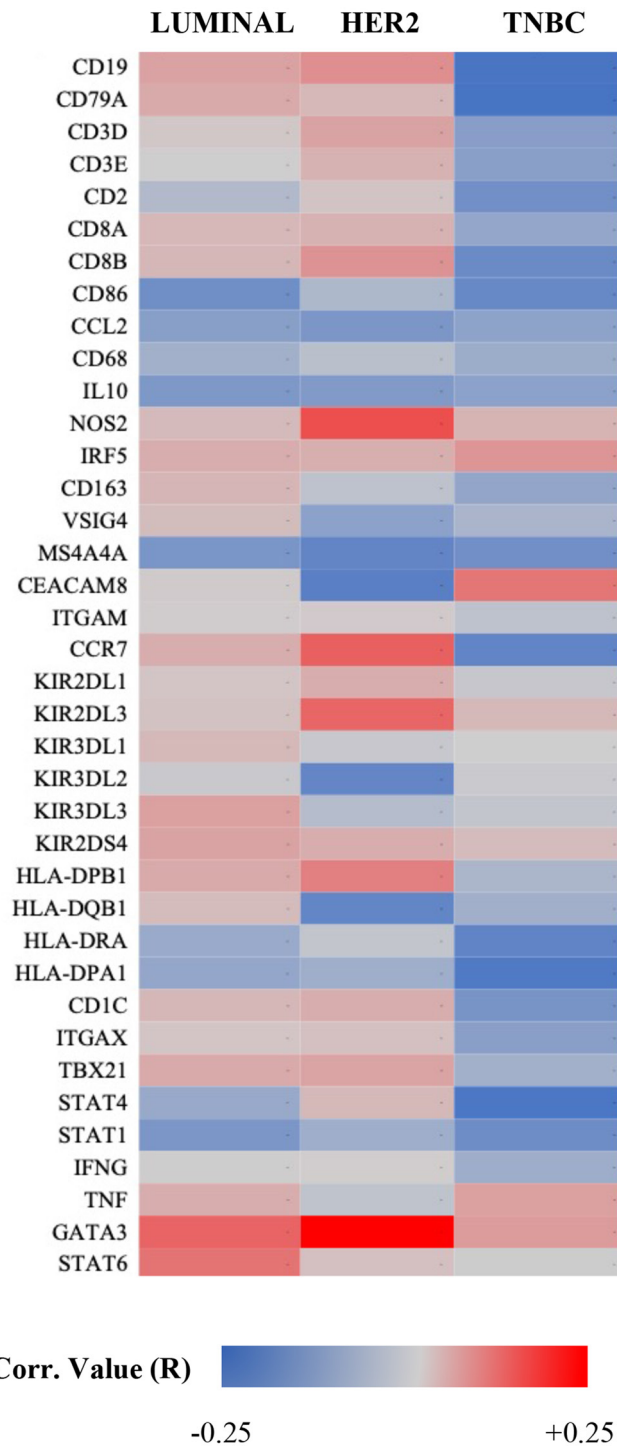
Supplementary Figure 1: Association of DVL-1 expression with overall survival (OS) in different breast cancer subtypes. Kaplan Meier database was used to compare survival rates between high and low expression of DVL1 (203230_at) in different subclasses of breast cancer. Survival curves of OS and DMFS in (A) Luminal A ($n = 2504$), (B) Luminal B ($n = 1425$), and (C) HER2+ ($n = 335$).



Supplementary Figure 2: DVL-1 localizes at various cancer-associated genes in MDA-MB-231 cells. An assembly of IgG (first row) and DVL-1 (second row) ChIP-Seq data in MDA-MB-231 for the *TRIO*, *COL5A1*, *EXD3*, *OR4A47*, *GLDN*, and *EFCAB6* genes, visualized by IGV. The true peaks representing enrichment of DVL-1 at a specific gene location is highlighted within the box.



Supplementary Figure 3: DVL-1 localizes at immune cell genes in MDA-MB-231 cells. An assembly of IgG (first row) and DVL-1 (second row) ChIP-Seq data in MDA-MB-231 for the *CD8B*, *CD1C*, *MS4A4A*, *STAT5B*, and *STAT4* genes, visualized by IGV. The true peaks representing enrichment of DVL-1 at a specific gene location is highlighted within the box.



Supplementary Figure 4: Heatmap representing correlation between DVL-1 expression and markers of immune cells in breast cancer. (Source: Table 2).