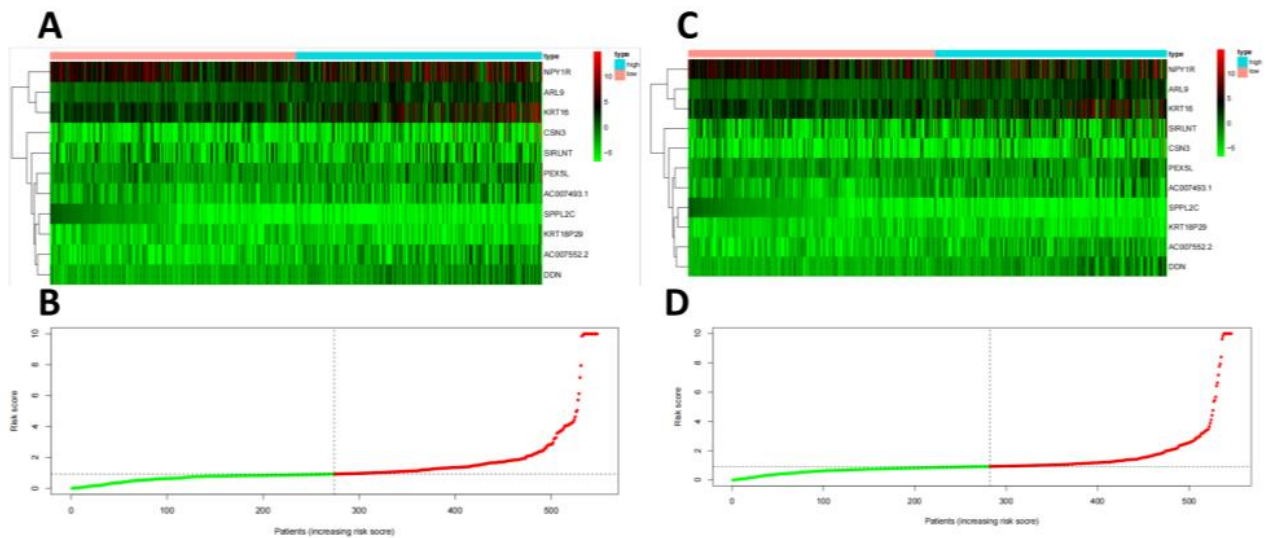


**Figure S1. Differential expression of TNBC-positive and -negative samples at the transcriptome level.**

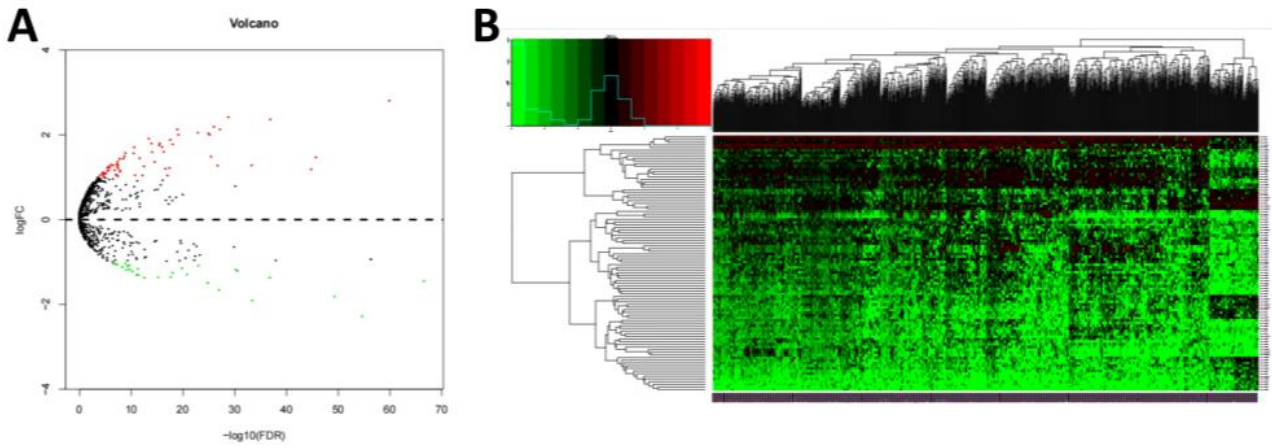
Note: (A) Volcano plot showing the differential RNA expression between TNBC-positive and -negative samples. Red represents significantly downregulated TNBC-positive samples, blue represents significantly upregulated TNBC-negative samples, and gray represents samples with no significant difference. (B) GO enrichment analysis of DEGs in TNBC-positive samples, with the size

of the points indicating the number of genes in the pathway and the color indicating the adjusted P-value. (C) KEGG enrichment analysis of DEGs in TNBC-positive samples, with the size of the points indicating the number of genes in the pathway and the color indicating the adjusted P-value.



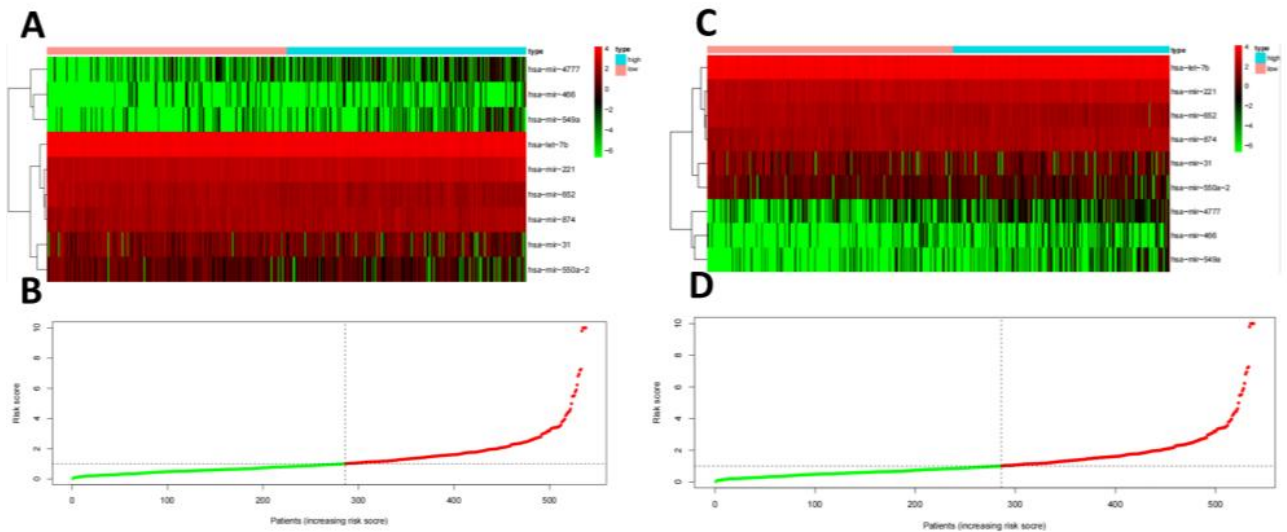
**Figure S2. Gene heatmap and risk curve of miRNA-seq data in the training and validation sets.**

Note: (A) Heatmap of DEGs between high-risk and low-risk groups in the training set. (B) The risk curve of the training set, where the X-axis represents patients sorted by increasing risk score and the Y-axis represents the computed risk score. The green region represents the low-risk group, and the red region represents the high-risk group. (C) Heatmap of DEGs between high-risk and low-risk groups in the validation set. (D) Risk curve of the validation set, with the X-axis representing patients sorted by increasing risk score and the Y-axis representing the computed risk score. The green region represents the low-risk group, and the red region represents the high-risk group.



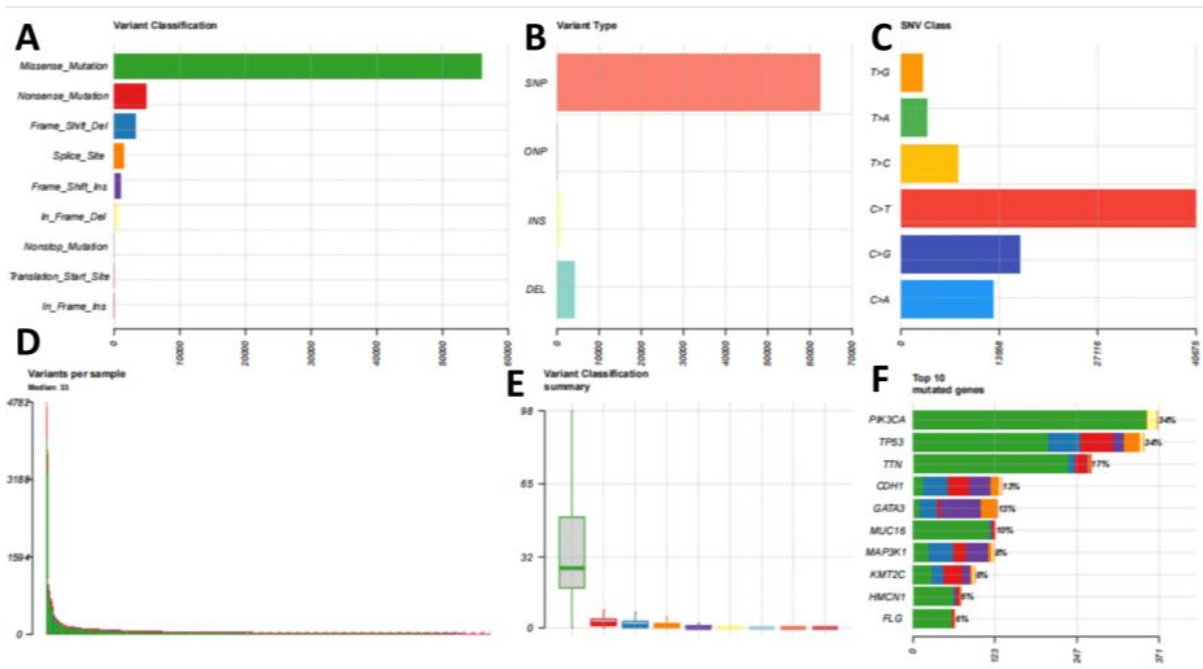
**Figure S3. Differences in transcriptomes between TNBC-positive and -negative samples.**

Note: (A) Volcano plot showing differential miRNA expression between TNBC-positive and -negative samples. Red indicates significantly downregulated expression in TNBC-positive samples, blue indicates significantly upregulated expression, and black indicates no significant difference. (B) Heatmap of miRNA expression differences between TNBC-positive and -negative samples.



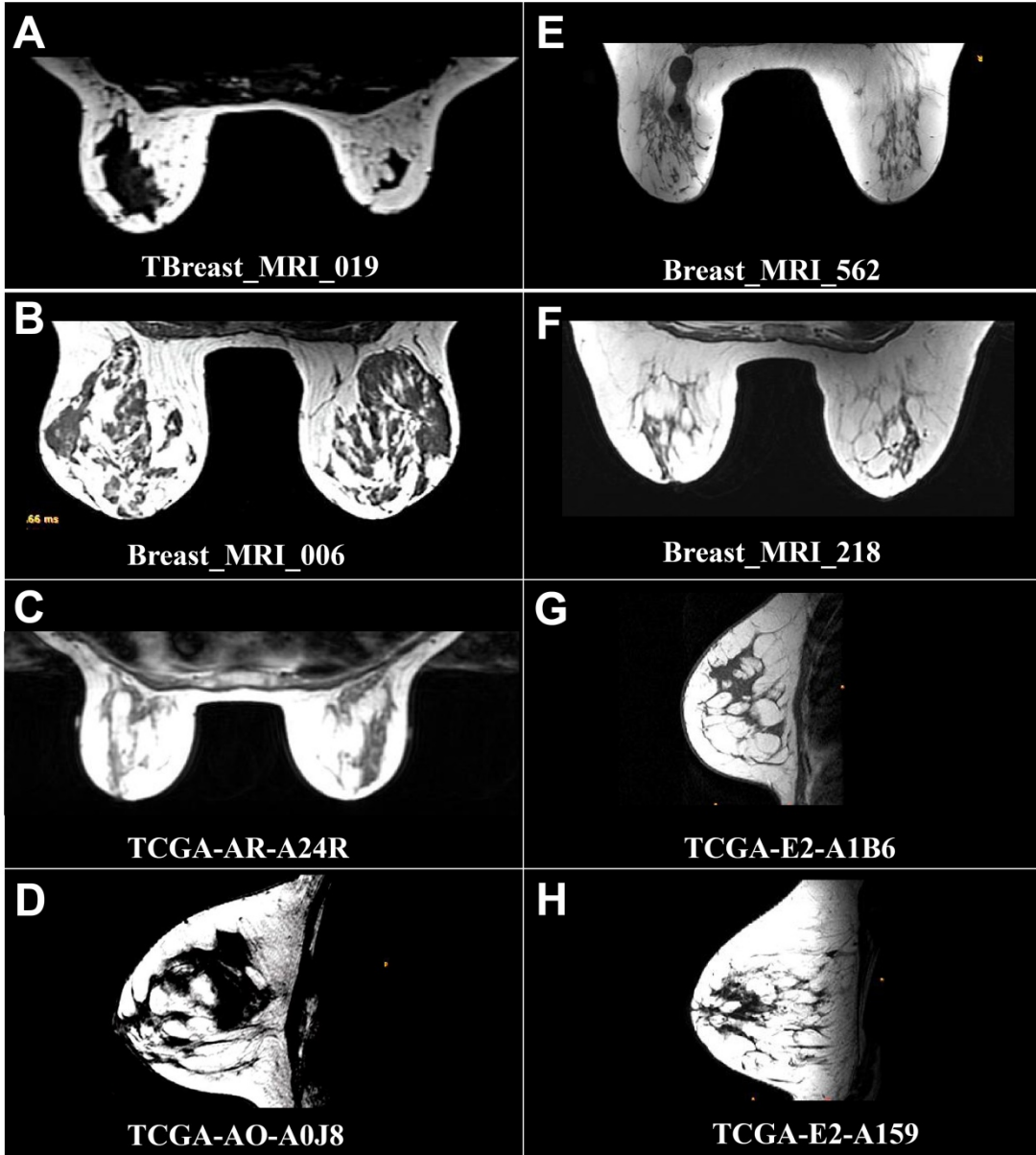
**Figure S4. Heatmap and risk curve of miRNA-seq data training and validation sets.**

Note: (A) Heatmap showing differential gene expression between high-risk and low-risk groups in the training set of the model. (B) Risk curve of the training set, with patients sorted by risk score on the x-axis and computed risk scores on the y-axis. The green region represents the low-risk group, and the red region represents the high-risk group. (C) Heatmap showing differential gene expression between high-risk and low-risk groups in the validation set of the model. (D) Risk curve of the validation set, with patients sorted by risk score on the x-axis and computed risk scores on the y-axis. The green region represents the low-risk group, and the red region represents the high-risk group.



**Figure S5. Gene mutation features in TNBC-positive and -negative samples.**

Note: (A) Variant Classification: types of mutations. (B) Variant Type: Primarily single nucleotide polymorphisms (SNP), which are significantly outnumbering other types. (C) SNV Class: classification of single nucleotide variations (SNVs). (D) Variants per Sample: distribution of the number of variants per sample. (E) Variant Classification Summary: distribution and median of different variant classifications in the form of a boxplot. (F) Top 10 Mutated Genes: top 10 genes with the largest number of mutations.



**Figure S6. Representative misclassified images under the DL model.**

Note: (A-D) False positive samples: Non-TNBC breast samples misclassified as TNBC. (E-H) False negative samples: TNBC samples misclassified as non-TNBC. The numbers below the images indicate the sample identification in the original dataset.