

Figure S1

The Pearson correlations calculated for the technical replicated (six samples analyzed in duplicates on separate chips 16_2, 18_3, 27_2, 28_3, 47_3 and 21_2). For each technical replicate, the biopsy location and sex is also listed. Overall positive correlations ($r^2 \geq 0.9$) were observed between all technical replicates.

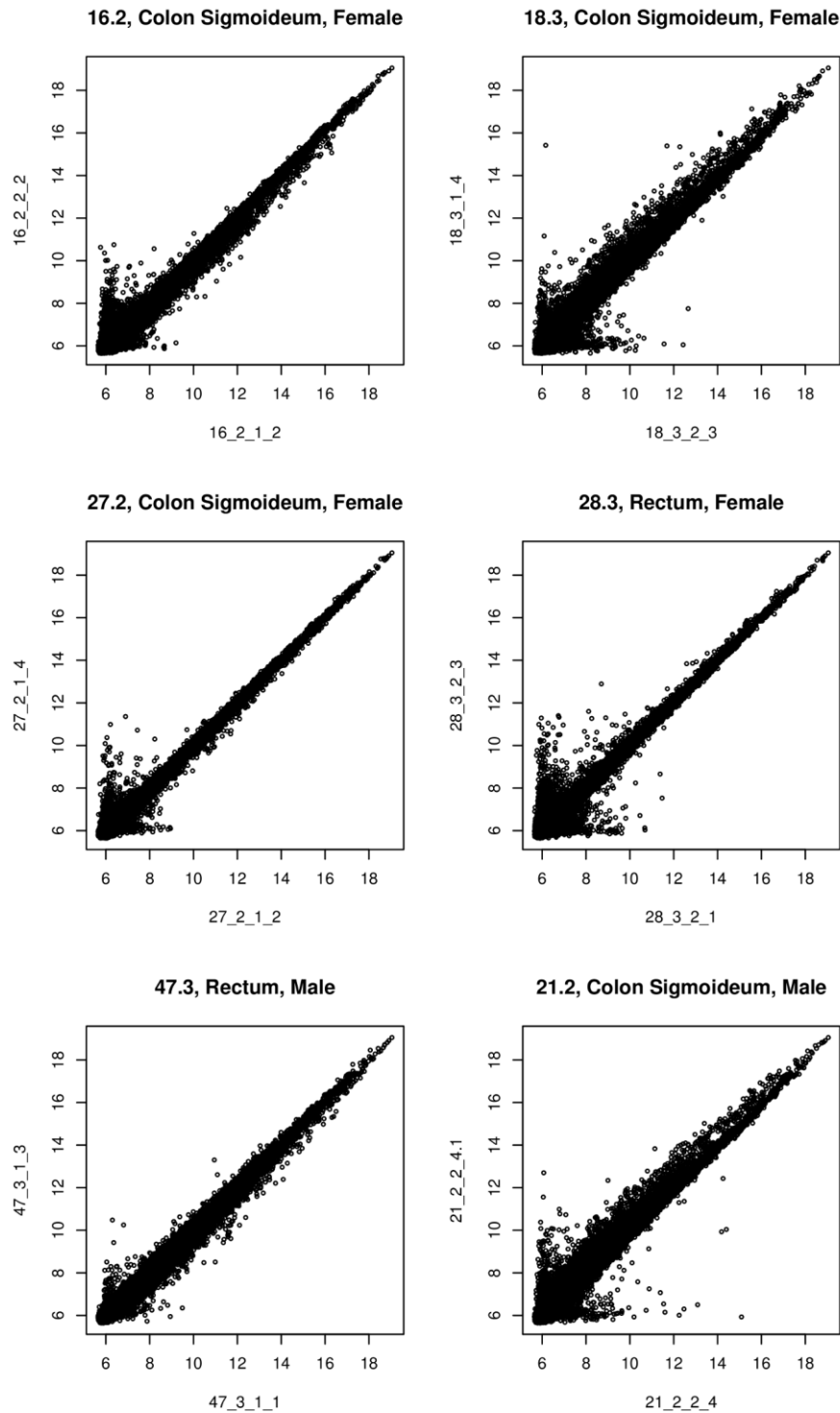


Figure S2

The scatterplot matrices describing the variation explained by the first four principal components for 90 biopsy samples. The different clinical subgroups iCD, iUC, niCD, niUC and controls are depicted in red, green, blue, cyan and black respectively.

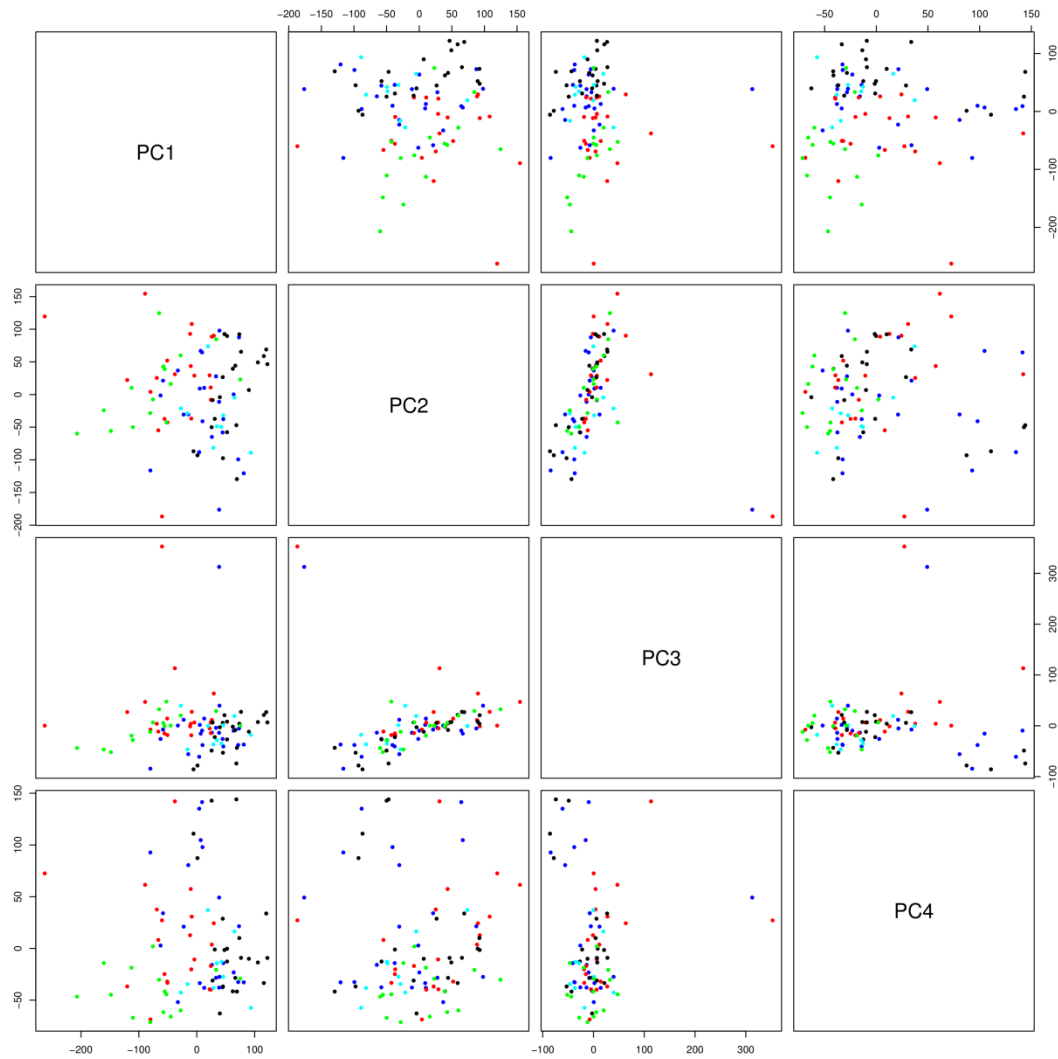


Figure S3

Unsupervised hierarchical clustering of the most dynamic probes (coefficient of variance >0.05) targeting lncRNAs (S3A) and protein-coding genes (S3B) across the samples in different clinical subgroups.

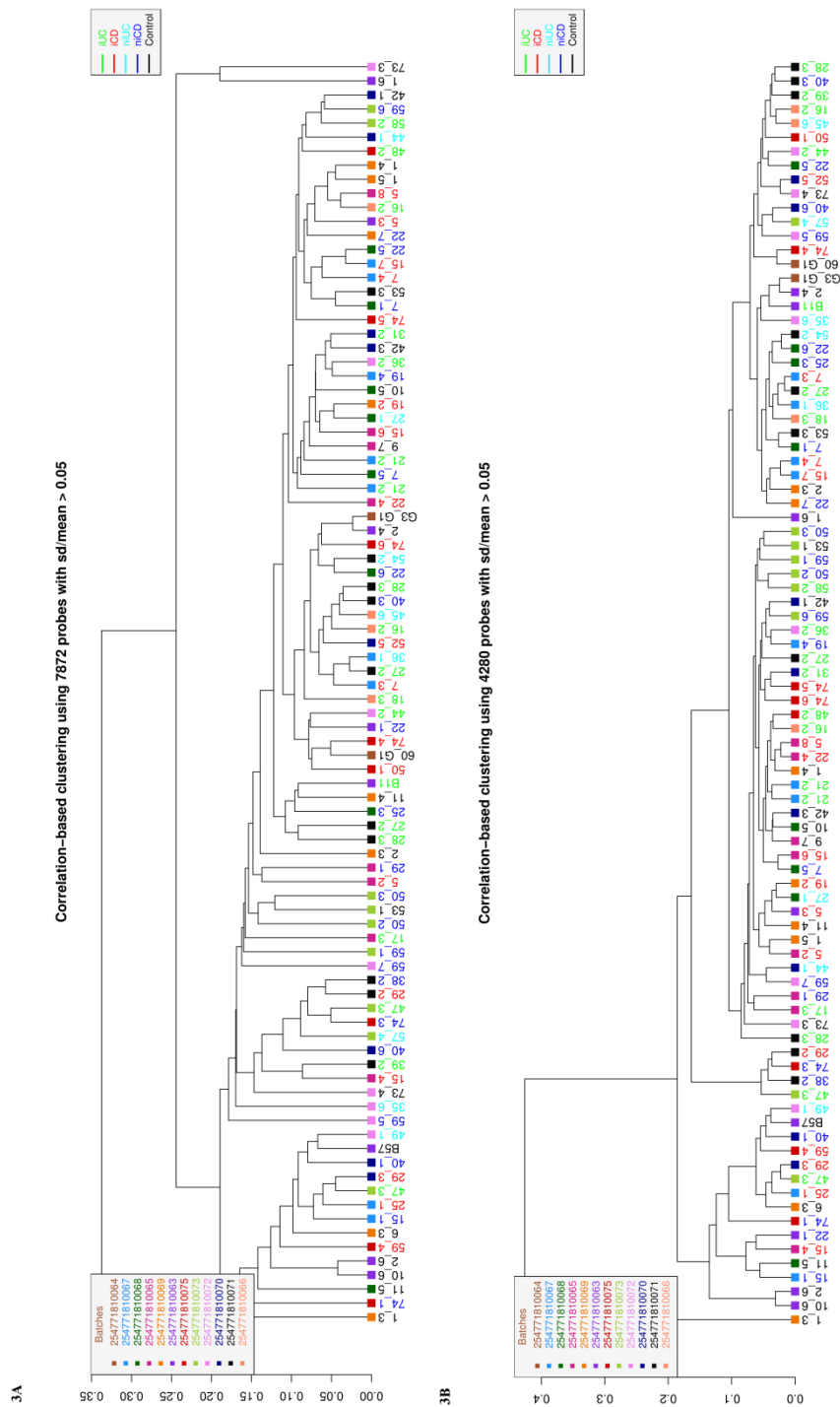
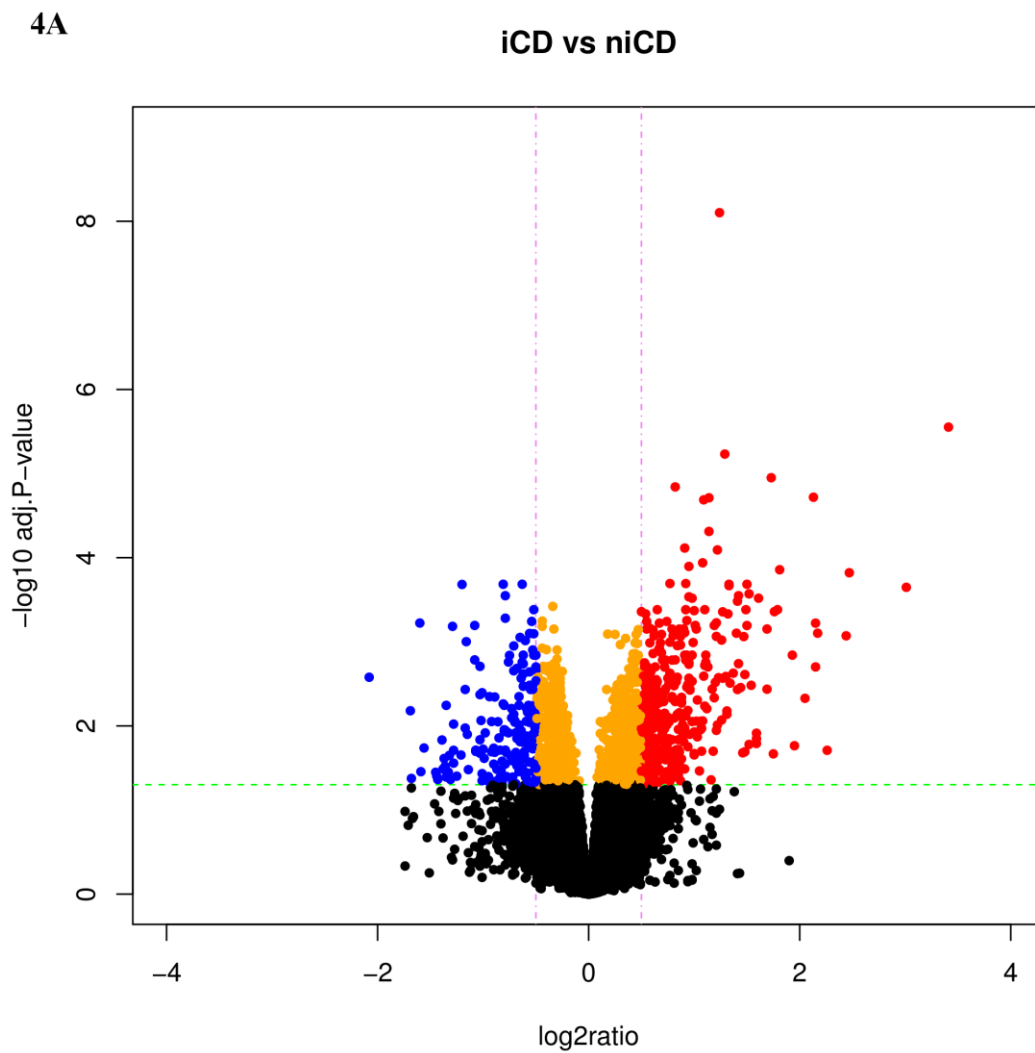


Figure S4

The log2 ratio and $-\log_{10}$ adj.P-values plotted and represented as volcano plots for the non-inflamed tissues contrasts iCD vs niCD (S4A) and iUC vs niUC (S4B). The probes in red, blue and orange colors represents up-regulated ($FC > 1.5$ and adj.P-value < 0.05), down-regulated ($FC < -1.5$ and adj.P-value < 0.05) and significant with small fold change ($FC > -1.5$ and < 1.5), respectively. The non-significant probes are represented in black color.



4B

iUC vs niUC

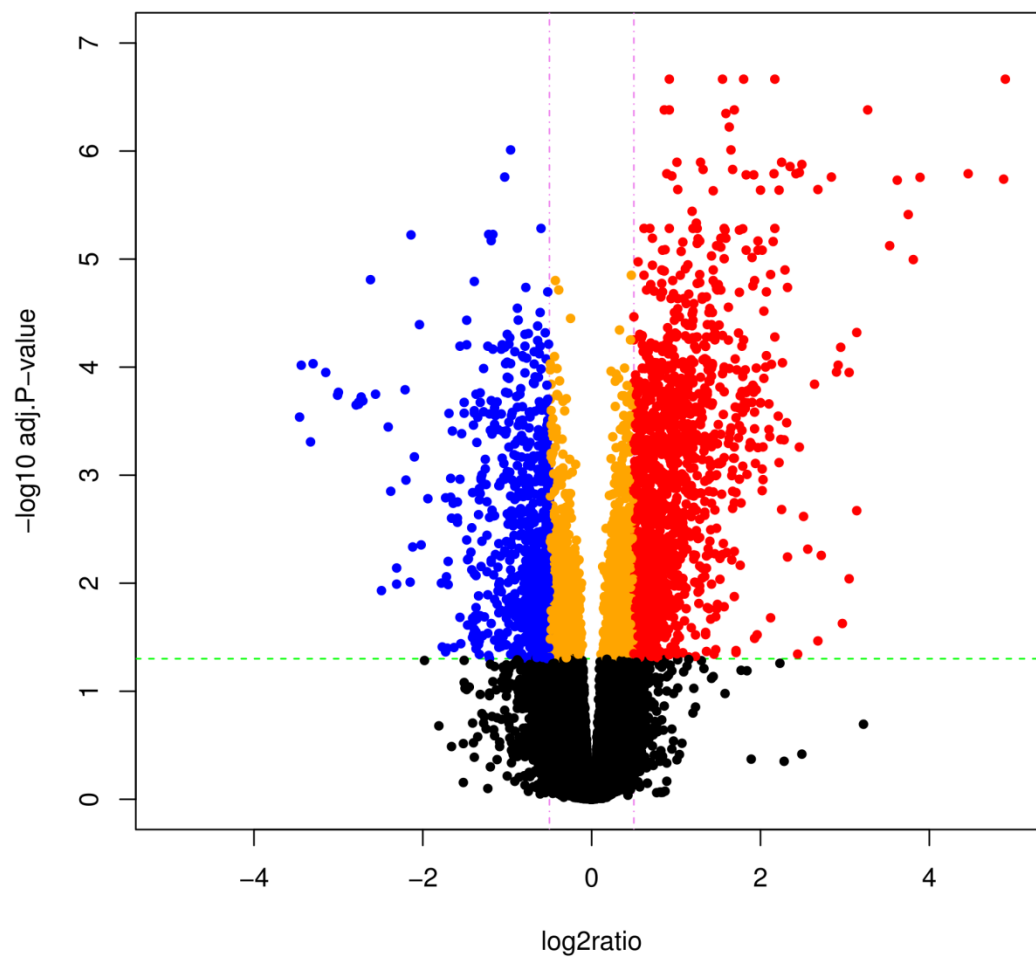


Figure S5

The expression map of the top 40 differentially expressed lncRNAs and protein-coding genes in iCD vs iUC (patients in red, controls in blue) based on unsupervised hierarchical clustering. The clustering algorithm was unable to distinguish between the iCD and iUC samples. The log2 normalized expression values are shown in the color key.

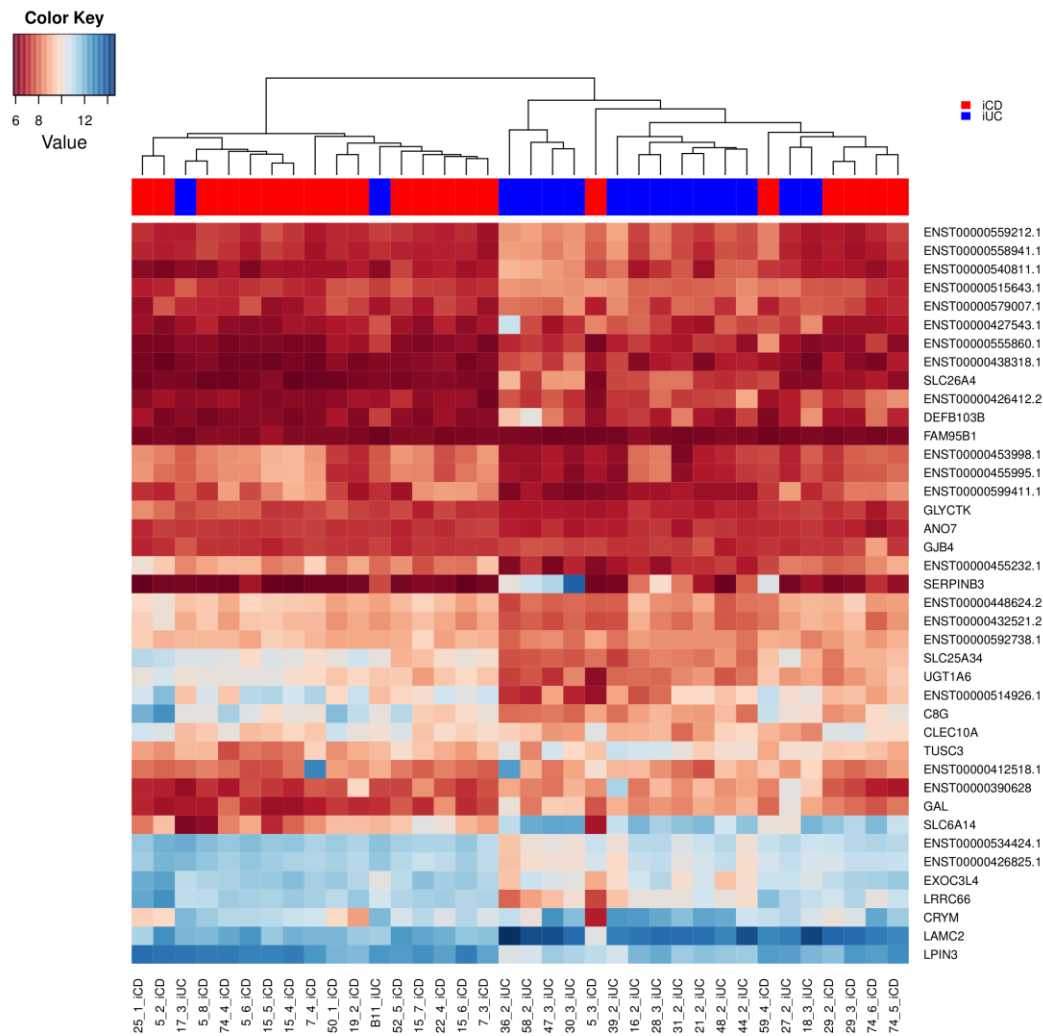
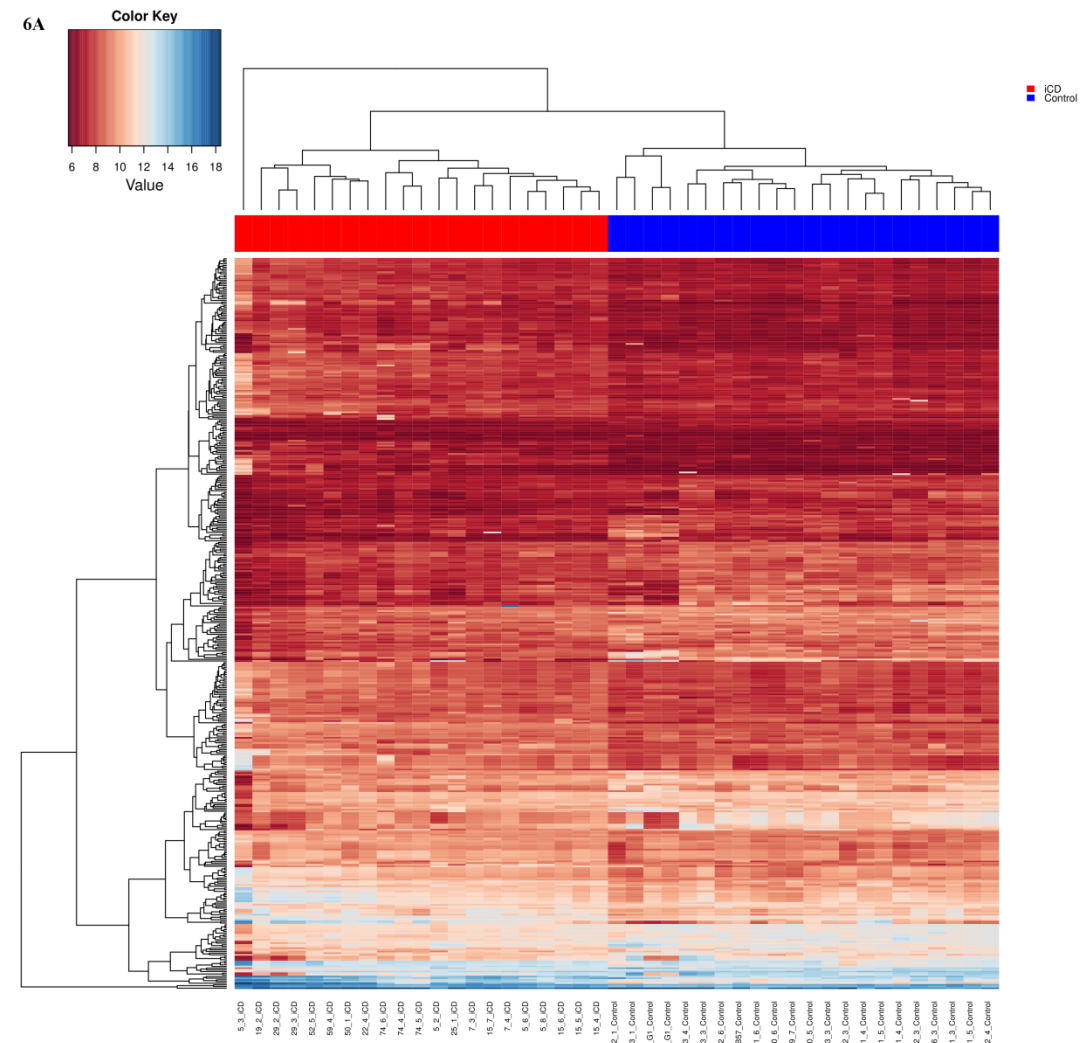


Figure S6

The expression map of the total differentially expressed lncRNAs and protein-coding genes in iCD vs controls (S6A) and iUC vs controls (S6B) (patients in red, controls in blue) based on unsupervised hierarchical clustering. The log₂ normalized expression values are shown in the color key.



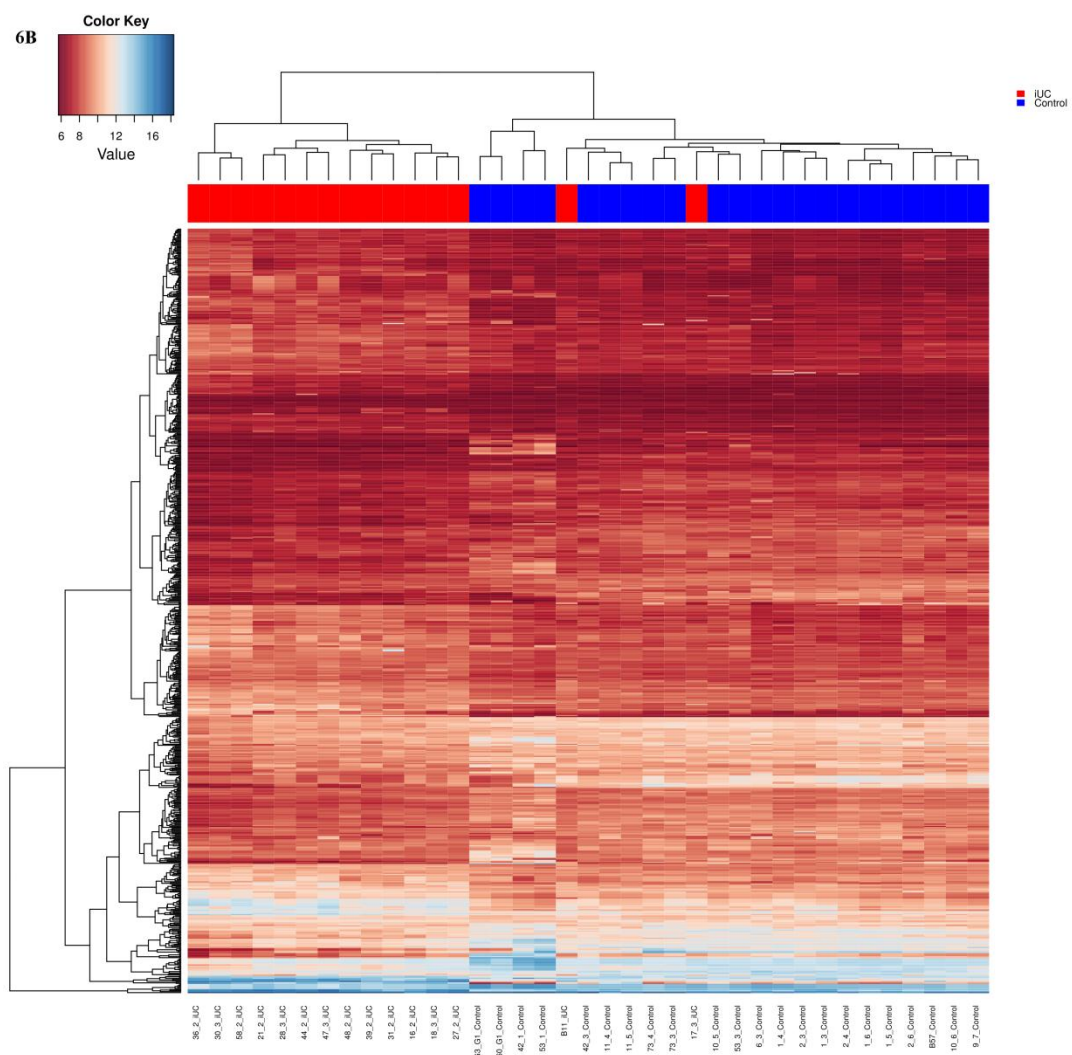


Figure S7

Dendrogram of samples and heatmap of clinical parameters. Linear regression model and weighted correlation network analysis (WGCNA) were used to investigate the impact of clinical parameters of samples on disease diagnosis.

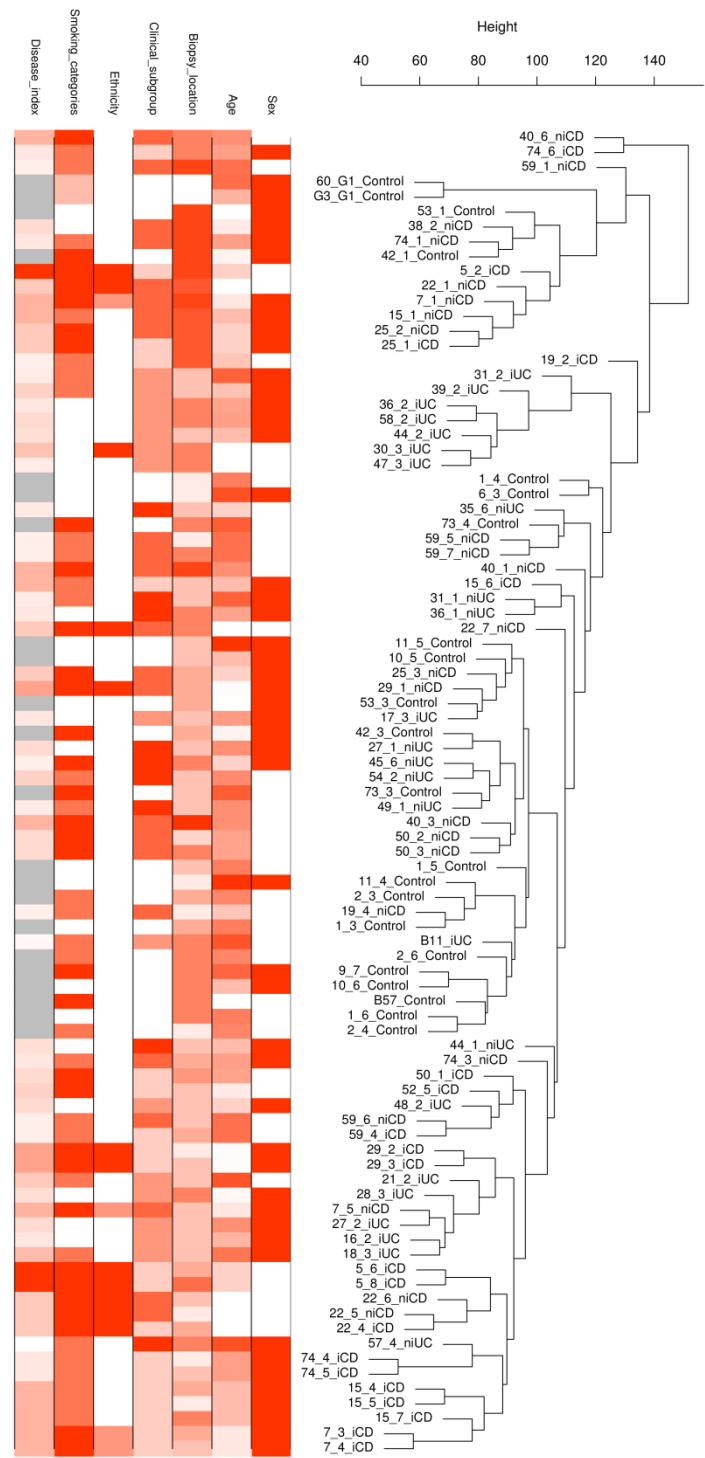


Figure S8

Receiver operating characteristic (ROC) curve analysis for age (a), sex (b), disease index (c), smoking (d) classification using differentially expressed lncRNAs in all five contrasts. ROC analysis for randomly selected same number of lncRNAs (e) was also performed to rule out any selection bias in our analysis.

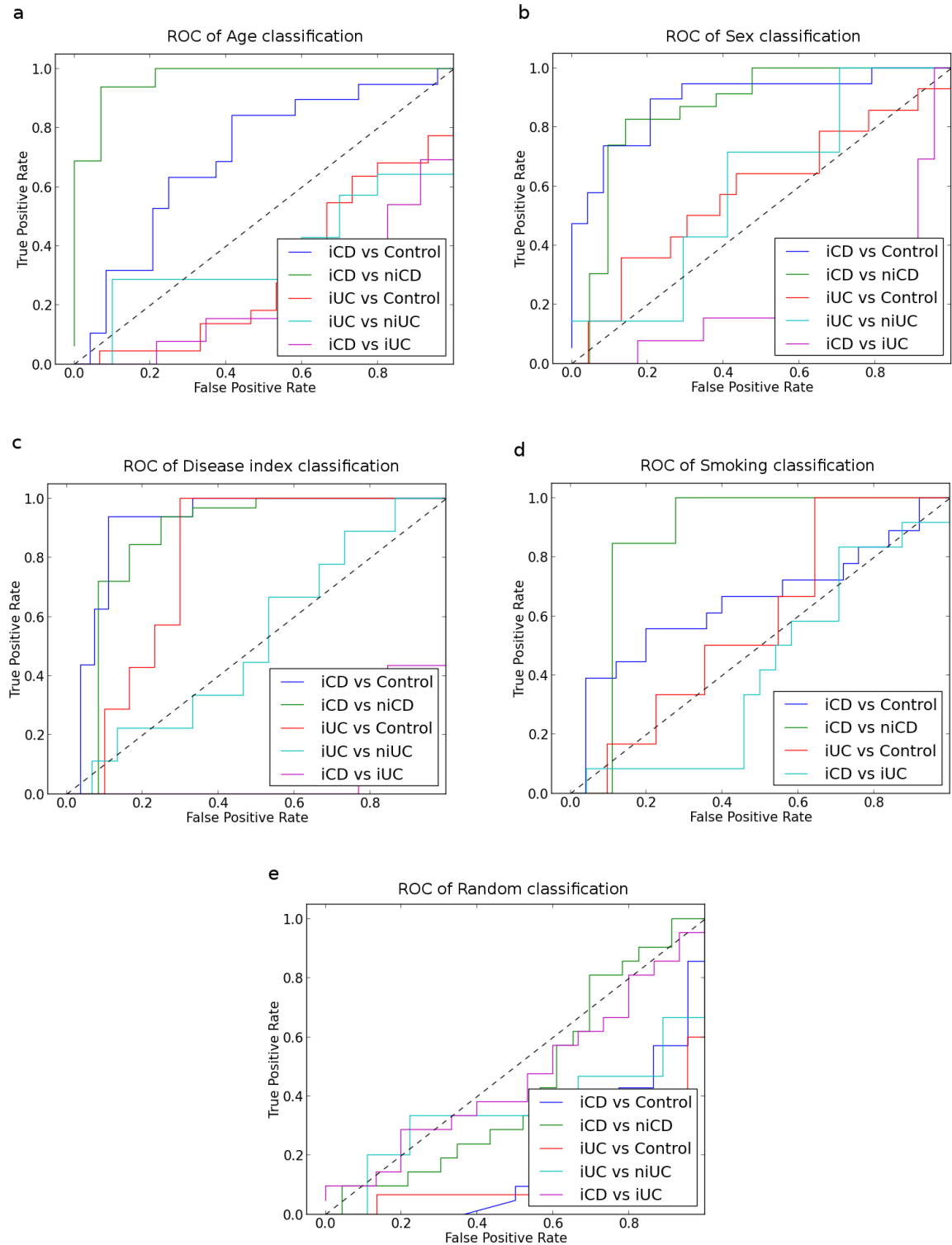


Figure S9

Overlap of differentially expressed genes identified by LIMMA and SVM. SVM was used to cross-validate the differentially expressed genes identified by LIMMA.

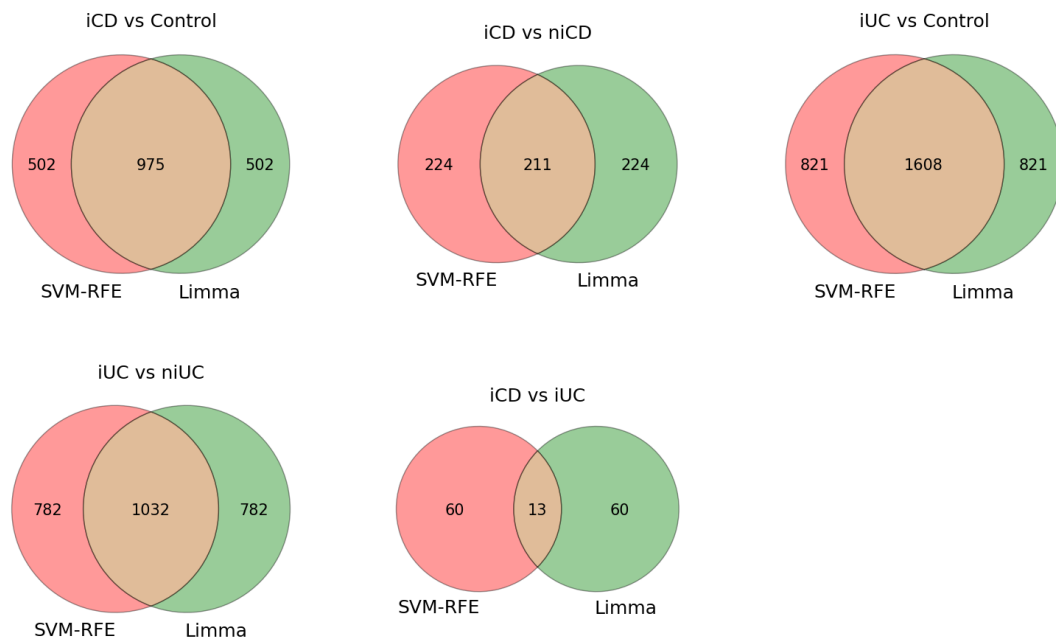


Figure S10

Co-expression network is built by hierarchical clustering and Dynamic Tree Cut. Modules are clusters of highly interconnected genes. The “brown”, “green” and “red” modules are enriched for differentially expressed genes between iUC / iCD and Control.

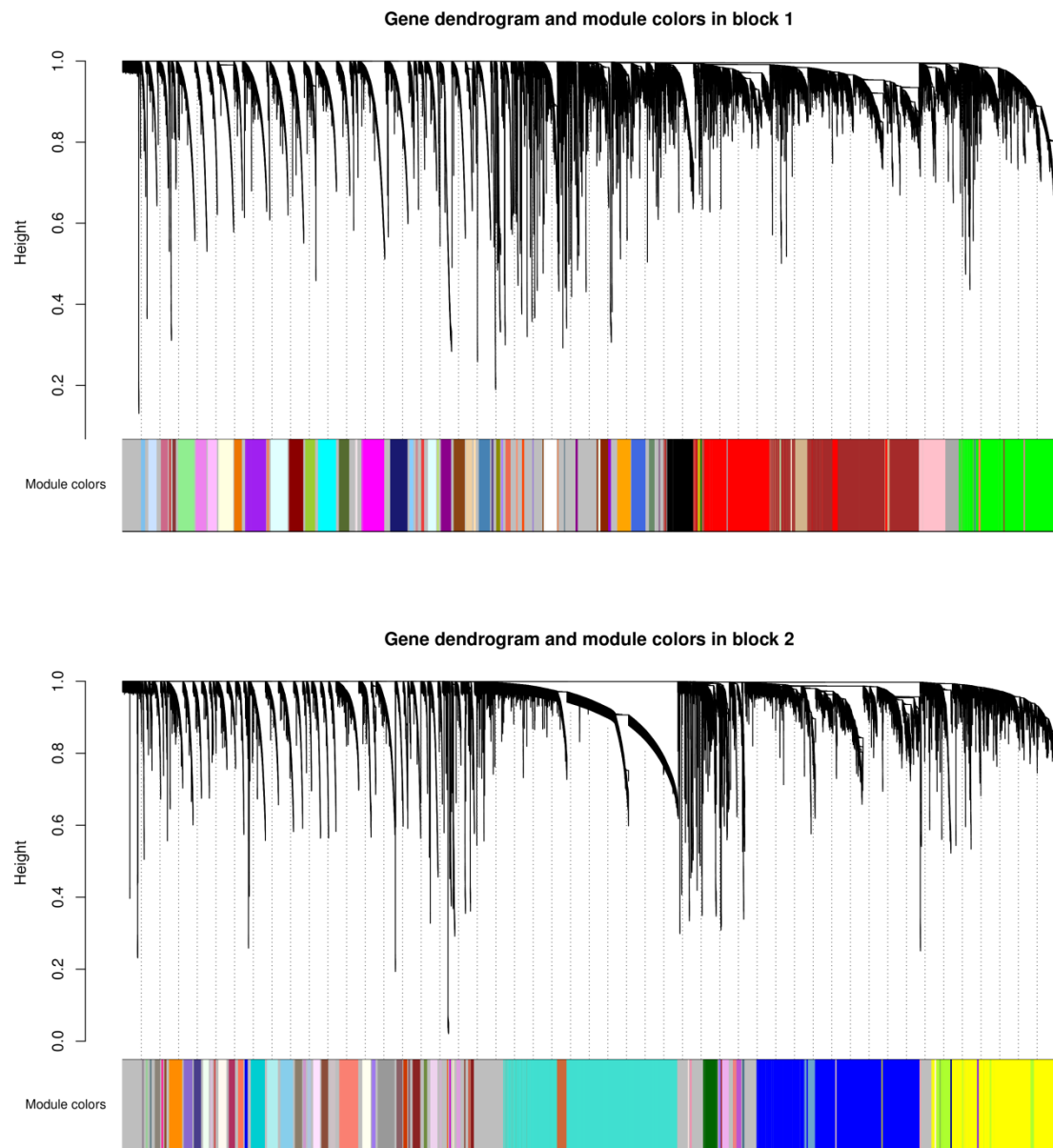


Figure S11

Module significance is determined as the average absolute gene significance measure for all genes in a given module

