

1 **Figure S1. Analyzation of B cell subsets and IL-10 expression in HCC PBMC with public**
2 **scRNA-seq data.** (A) tSNE plotting showed the B cell clusters in Normal and HCC PBMC. (B)
3 The expression of markers in B cell subsets. (C) ssGSEA analyzation on different B cells in
4 different histology, B cell related hallmark biological functions and GO_BP are obviously
5 repressed in HCC B cells globally. (D) Biological process of target B subclasses
6 (B10/MZB/FoB) were also analyzed by ssGSEA. (E-F) The landscape of interaction of B cell
7 in both health and HCC PB. (G) Expression of IL-10 in B10 cell and other B cell subtypes at
8 single-cell mRNA level. (H) Survival analysis in TCGA LIHC project, total 364 patients with
9 clinical information, and figure KM-curve, dividing the cohort into IL10-high and IL10-low
10 groups.

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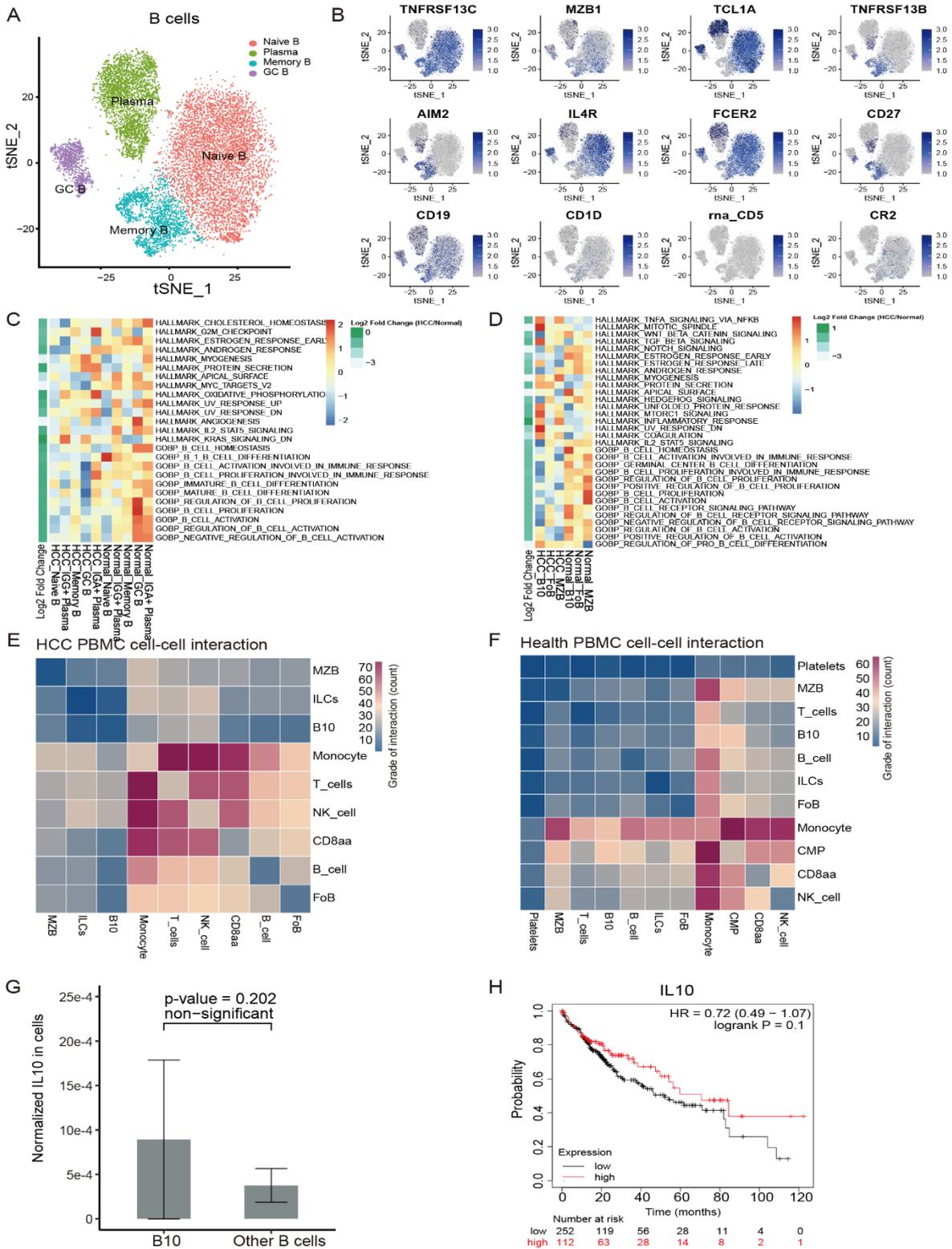


Figure S1

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16 **Table S1. Marker Genes by Cell.**

17 Marker genes of lymphocyte subsets in the three datasets.

18 **Table S2. DEG_10x_vs_zhang's data.**

19 Differentially expressed genes (DEGs) of PBMC between HCC and healthy donors are
20 analyzed by R packages DESeq2 (v1.16.1). DESeq2 provides an acceptable strategy to
21 determine the differences in single-cell datasets. We use $p_{\text{adjust}} < 0.01$ and $\log_2\text{fold-change} >$
22 0.3 (or < -0.3) as a cutoff to interpret significant differentially expressed genes to plot heatmap.

23 **Table S3. Cell interaction.**

24 Cell-cell interactions are operated by CellPhoneDB, and ggplot2 was used for drawing dot
25 diagrams.

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