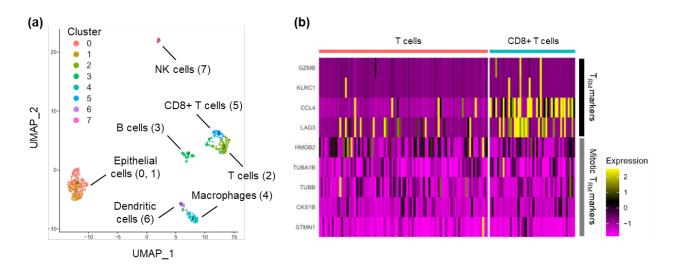
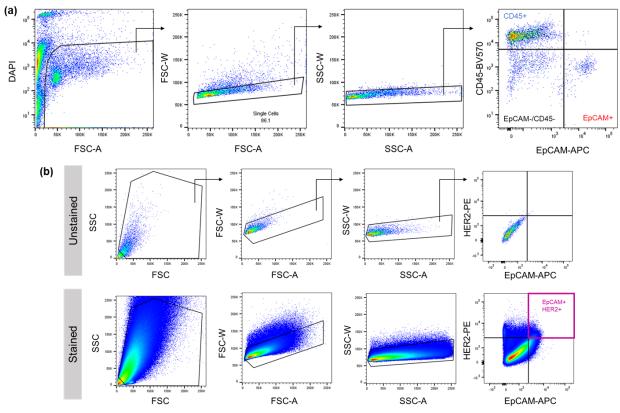
## SUPPLEMENTARY FIGURES



## Supplementary Figure 1. Single-cell transcriptomic analysis shows cellular diversity in a research core biopsy specimen.

(a) High-dimensional uniform manifold approximation and projection (UMAP) clustering of single-cell transcriptomic data from a triple-negative breast core biopsy specimen shows a diverse composition of epithelial, lymphoid and myeloid cells.

(b) Heatmap of single-cell transcriptome of all T cells identified in the triple-negative breast core biopsy specimen. T cells positive for non-mitotic  $T_{RM}$  markers (top) were enriched in the CD8+ T cell cluster, while few or none of them were positive for mitotic  $T_{RM}$  markers (bottom).  $T_{RM}$  markers were derived from a previous publication (Savas P et al., 2018). See also Figure 3C.



Supplementary Figure 2. Gating strategies were used to define specific cell populations in research core biopsy specimens.

(a) Gating strategy used to identify viable (DAPI-negative) EpCAM-positive, CD45-positive or double-negative cells. See also Figure 2c.

(b) Gating strategy used to identify formalin-fixed EpCAM/HER2-positive cells. See also Figure 3b.

## SUPPLEMENTARY TABLE

Specimen -Clinical specimen -Research specimen	Primary pathologic diagnosis	Grade	Cellularity
1	IDC	1	< 5%
	IDC	1	< 5%
2	IDC	1	10%
	IDC	1	< 1%
3	Fibroadenoma	N/A	N/A
	Fibroadenoma	N/A	N/A
4	IDC	2-3	20%
	IDC	2	20%
5	ILC	2-3	30%
	ILC	2	30%
6	IDC	3	20%
	IDC	3	30%
7	IDC	3	60%
	IDC	3	10%
8	ILC	2	30%
	ILC	2	30%
9	IDC	2	50%
	Non-diagnostic, debris	N/A	N/A
10	IDC	2	40%
	IDC	2	40%

Legend: IDC, invasive ductal carcinoma; ILC, invasive lobular carcinoma.

Supplementary Table 1. Concordance between clinical and research biopsy specimens.