

SUPPLEMENTAL FIGURE 1. Representative *in vitro* saturation binding curves of ¹¹¹In-DTPA-pertuzumab to SKBr-3 (A) or MDA-MB-361 (B) cells treated for 72 h with trastuzumab (14 μ g/mL) and untreated controls. The B_{max} values were calculated from the specific binding by fitting a plot of total cell-bound ¹¹¹In-DTPA-pertuzumab (nmols) *vs*. the concentration of total added radioligand (nmols/L) to a 1-site saturation binding model using Prism® Ver. 4.0 software and then mathematically deriving the non-specific binding for subtraction from the total binding.

SUPPLEMENTAL TABLE 1

3-Day Treatment 3-Week Treatment PBS Trastuzumab PBS Mouse Trastuzumab 39.4 1 39.5 16.2 7.2 2 32.5 13.3 41.1 7.2 3 20.2 10.9 23.2 8.1 4 21.8 n/a n/a 7.8

Subcutaneously with MDA-MB-361 Human Breast Cancer Xenografts at 72 Hours Post-Injection

Tumor Uptake (%ID/g) of ¹¹¹In-DTPA-Pertuzumab of Individual Athymic Mice Implanted

n/a: not available

SUPPLEMENTAL TABLE 2

Immunohistochemical Analysis of Excised MDA-MB-361 Human

Breast Cancer Xenografts Stained for HER2

PBS*	Trastuzumab 3 days*	Trastuzumab 3 weeks
25 ± 17	30 ± 13	n/a^{\dagger}

* Tumors were scored as the percentage of strong, complete membrane

staining. Results are presented as the mean \pm SD of 3 tumors.

[†]Not applicable. Too few viable tumor cells to assign a score.