

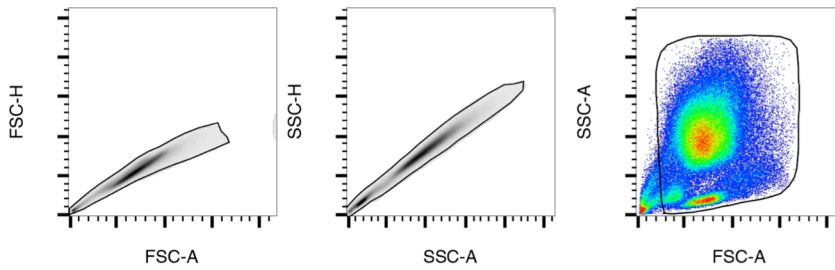
SUPPLEMENTAL MATERIAL

Table S1: Antibody used flow cytometry-based ex vivo BsAb antibody assay

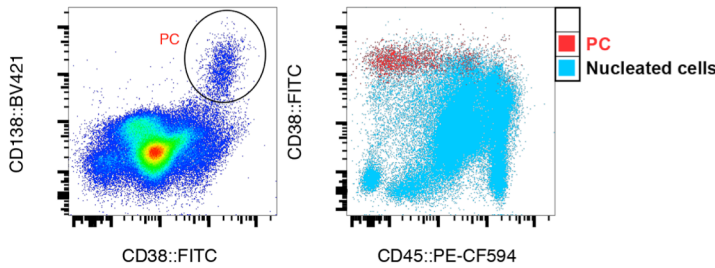
Marker	Channel	Clone	Supplier	Cat #
CD38	FITC	Multi-Epitope	ALPCO	70-38F2
CD269 (BCMA)	PE	19F2	Biolegend	357504
CD45	PE-CF594	HI30	BD Biosciences	562279
7-AAD	PerCP-Cy5.5	Kit	BD Biosciences	559925
CD8	PE-Cy7	RPA-T8	BD Biosciences	557750
Annexin V	APC	Kit	BD Biosciences	550474
CD3	APC-C750	UCTH-1	ALPCO	70-3AC750
CD138	BV421	MI15	BD Biosciences	562935
CD107a	BV510	H4A3	BD Biosciences	563078

Figure S1: Gating strategy for plasma cell and cytotoxic T cells. Following (A) doublet & debris exclusion based on morphological features, (B) plasma cells were defined as CD38^{high}/CD138^{high}/CD45^{-/int} and PC lysis was calculated for each treatment group as follow $[\% \text{ lysis} = \frac{\text{Treated PC count} \times 100}{\text{Untreated PC counts}}]$. (C) Cytotoxic T cells were defined as CD3⁺/CD8⁺/CD45^{high} (C) and CTL distribution was calculated for each treatment group as follow $[\% \text{ CTL} = \frac{\text{Treated CTL count} \times 100}{\text{Untreated CTL counts}}]$.

A Doublet exclusion and debris exclusion gating strategy



B Plasma cell gating strategy



C Cytotoxic T cell gating strategy

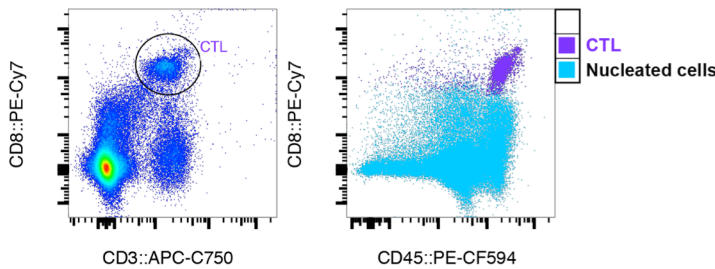


Figure S2: Gating strategy for plasma cell and cytotoxic T cells phenotyping.

(A) Gated from CD38^{high}/CD138^{high}/CD45^{-int} population, live plasma cells were defined for each treatment group as 7-AAD⁻ PC and dead apoptotic cells as 7-AAD⁺/Annexin V⁺ PC. **(B)** Gated from CD38^{high}/CD138^{high}/CD45^{-int} population, BCMA expression was expressed for each treatment group as average frequency of PC expressing BCMA (PC BCMA⁺) and mean fluorescence intensity (BCMA MFI). BCMA expression by T cells (defined as CD3⁺/CD45^{high} population) was used as negative control. **(C)** Gated from CD3⁺/CD8⁺/CD45^{high}, CTL degranulation was determined for each treatment group by calculating mean fluorescence intensity of CD107a.

