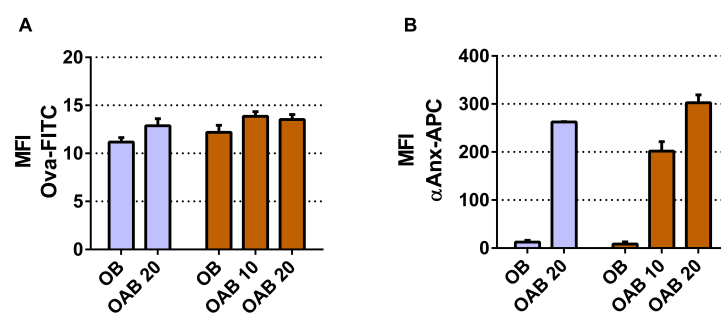
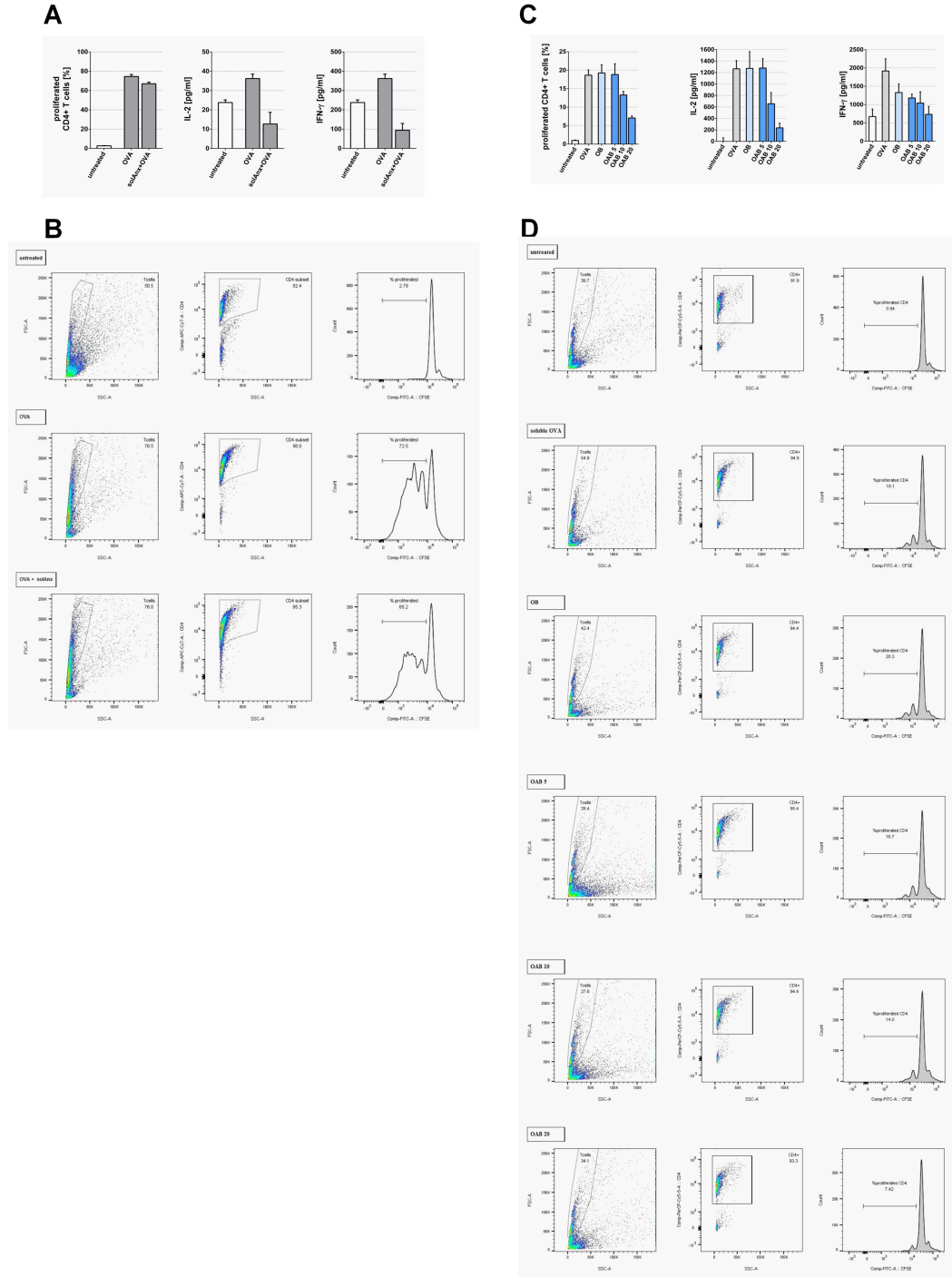


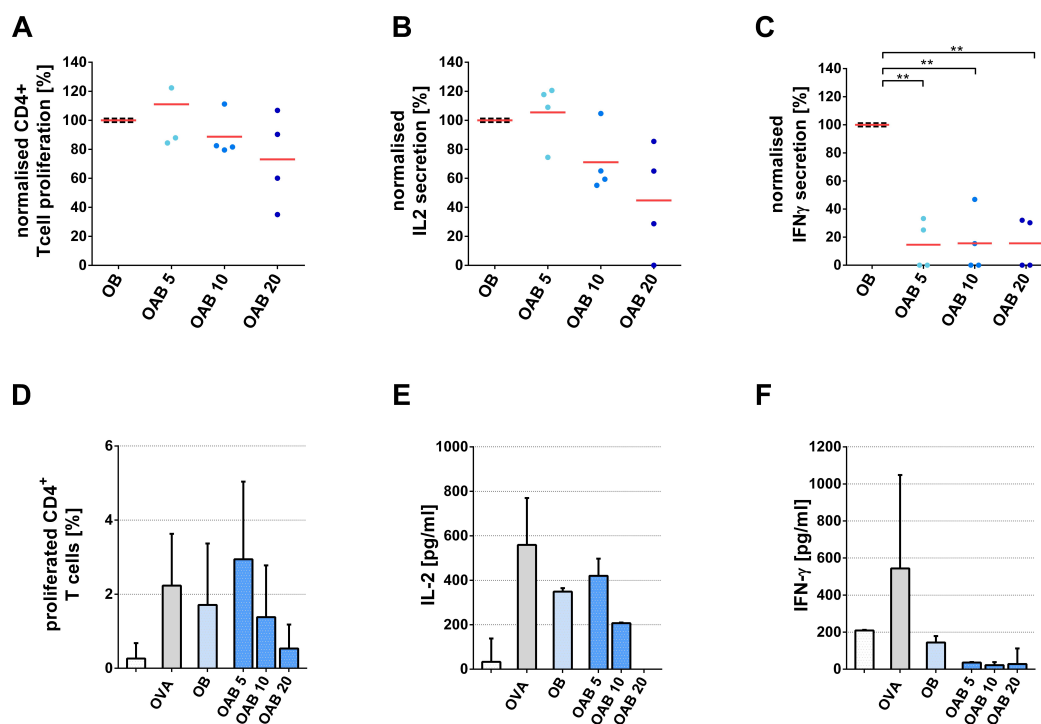
1. Supplementary material



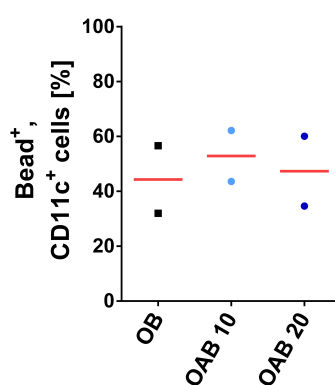
SFig. 1. Bead coating controls. Bead coating was analyzed by FC. Proteins were detected directly (Ova-FITC) or by staining with a fluorescent antibody (α Anx-APC). Different bar colors indicate different bead charges



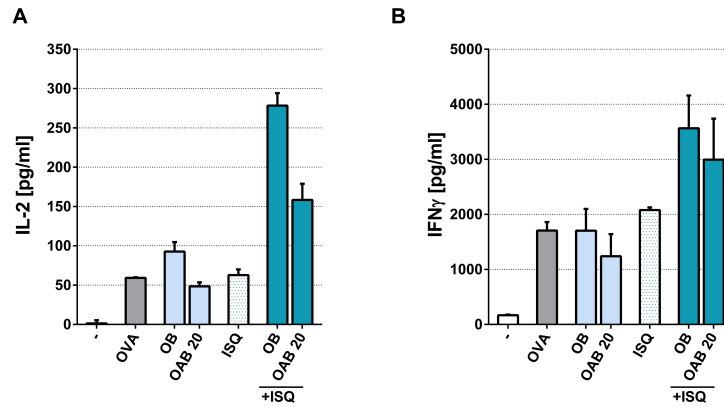
SFig. 2. Exemplary flow cytometry plots of attenuated T cell responses. BMDC were incubated with beads coated with OVA (OB) or OVA and Anx (OAB) and additionally stimulated with CpG. CFSE-labeled OT-II T cells were added and co-cultured for 5 days before the T cells were harvested to analyze proliferation by FC. Additionally, cytokine secretion was analysed by ELISA. (A-B) exemplifies one individual experiment (measured in triplicates) from the summarized data shown in Fig1A-C. (C-D) exemplifies one individual experiment (measured in triplicates) from the summarized data shown in Fig1G-H.



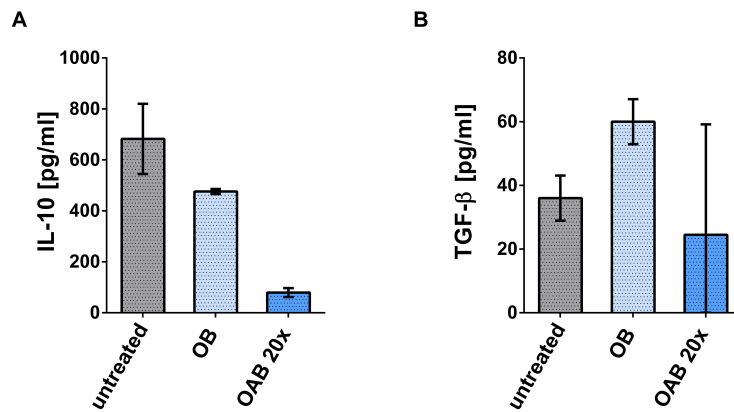
SFig. 3. Treatment of unstimulated BMDC with particulate Anx attenuates T cell responses. BMDC were incubated with beads coated with OVA (OB) or OVA and Anx (OAB). Only if indicated, the cells were additionally stimulated with CpG. CFSE-labeled OT-II T cells were added and co-cultured for 5 days before the T cells were harvested to analyze proliferation by FC. Additionally cytokine secretion was analyzed by ELISA. (A-C) T cell responses to ascending bead concentrations of 4 independent experiments. Dots represent individual experiments. (D-F) exemplifies the raw data of one individual experiment (measured in triplicates) from the summarized data shown in A-C and Fig1G-I.



SFig. 4. Bead uptake is not influenced by different amounts of Anx. BMDC were pre-treated with fluorescent beads [14×10^6 /ml] and stimulated with CpG. 2 days later, BMDC were harvested, stained for CD11c and their bead content was analyzed by FC.



SFig. 5. Exogenous ISQ increased cytokine secretion without affecting OAB-mediated inhibition. BMDC were pre-treated with beads and CpG before they were co-cultured with OT-II T cells. Cytokine secretion was measured after 2 or 5 days of co-culture for IL-2 and IFN γ , respectively. For the indicated conditions, ISQ [0,005 μ M] was added at least 30 min prior to the T cells. (A-B) exemplifies cytokine secretion without normalization of one individual experiment from the normalized and summarized data shown in Fig2E-F.



SFig. 6. OT-II T cells produce limited amounts of IL-10 and TGF- β . BMDC were pre-treated with beads and CpG before they were co-cultured with OT-II T cells. Cytokine secretion was measured after 5 days of co-culture. (A) exemplifies the IL-10 secretion of one individual experiment from the summarized data shown in Fig4B. (B) exemplifies the TGF- β secretion of one individual experiment from the summarized data shown in Fig4C.