

Certificate of Analysis

| | |
|-------------------|---|
| Cell count | >2 million per vial |
| Viability | 95% |
| CD4+ | 100.0% |
| CD4+CD25+ | 82.2% |
| CD25+CD127- | 78.5% |
| Sterility Testing | Negative for bacteria, fungi and mycoplasma |

Donor Information

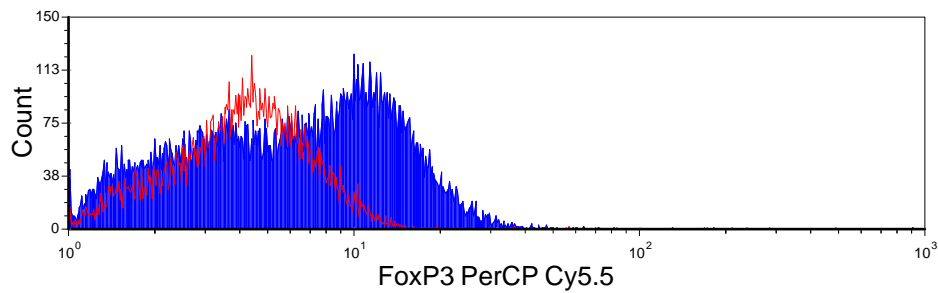
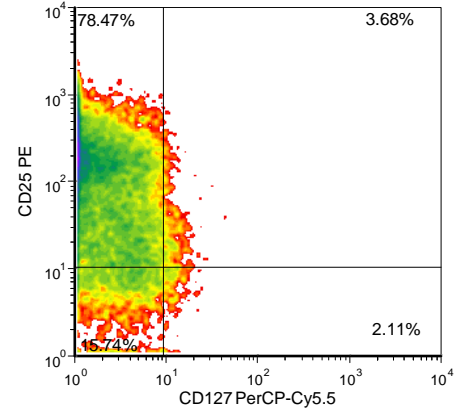
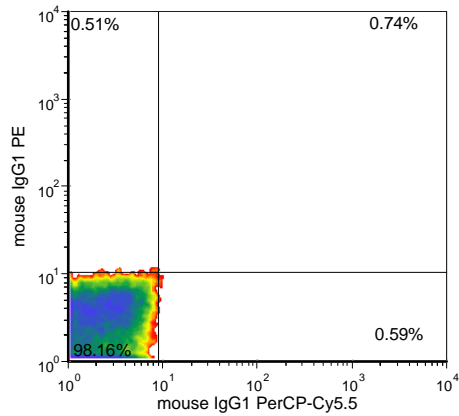
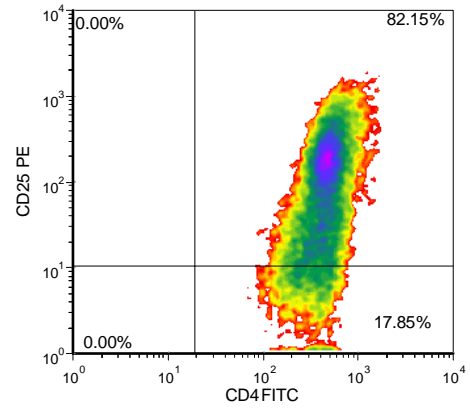
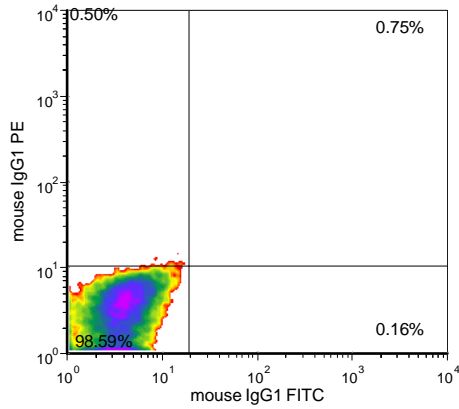
| | |
|----------|------------|
| Donor ID | 345 |
| Sex | Male |
| Age | 40 |
| Race | Caucasian |
| Height | 6'1" |
| Weight | 250 lbs |
| ABO Type | O Positive |

HLA typing

| | Allele 1 | Allele 2 |
|----------|----------|----------|
| HLA-A | 31 | 32 |
| HLA-B | 07 | 40 |
| HLA-C | 03 | 07 |
| HLA-DRB1 | 04 | 15 |

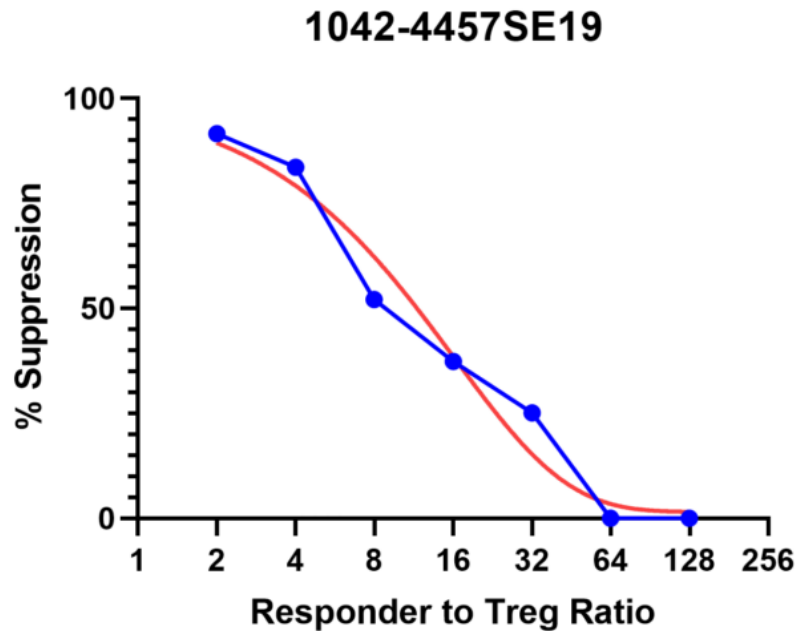
Donors are tested for the blood borne pathogens HIV-1 and 2, Hepatitis B, Hepatitis C and HTLV-1 and are negative. Cells should still be handled as if potentially infectious following biosafety level 2 procedures

FOR RESEARCH USE ONLY. NOT FOR DIAGNOSTIC OR THERAPEUTIC USE.



Red histogram designates intracellular staining with isotype control (mouse IgG PerCP Cy5.5). Filled blue histogram designates intracellular staining with anti-FoxP3 PerCP Cy5.5 antibody.

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Co-Culture of Tregs Resulted in Suppression of Anti- CD3-mediated CD8+ T Cell Proliferation of Allogeneic Peripheral Blood Mononuclear Cells (PBMCs). In short, in a 96 well plate, 250,000 CFSE-labelled allogeneic responder PBMC were plated/well in the presence of 0.2 µg/mL anti-CD3 and varying number of Tregs as shown in the chart above. The plate was incubated in a 37 deg C, 6% CO₂ incubator. After 5 days of culture, individual wells were harvested and labelled with PE-labelled anti-CD8 mAb. Data was acquired on a flow cytometer and the extent of proliferation (Division Index, DI) was analyzed using FCS Express 4.0. % Suppression = 100- (DI in the presence of Treg/DI in the absence of Treg) x 100.