Rapamycin and abundant TCR stimulation are required for the generation of stable human induced regulatory T cells

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Received 8 October 2020; Revised 7 July and 8 October 2020; Accepted 12 November 2020
doi: 10.1002/cti2.1223

Clinical & Translational Immunology 2020; 9: e1223

Abstract

Objectives. Regulatory T cells (Tregs) are a vital sub-population of CD4+ T cells with major roles in immune tolerance and homeostasis. Given such properties, the use of regulatory T cells for immunotherapies has been extensively investigated, with a focus on adoptive transfer of ex vivo expanded natural Tregs (nTregs). For immunotherapies, induced Tregs (iTregs), generated in vitro from naïve CD4+ T cells, provide an attractive alternative, given the ease of generating cell numbers required for clinical dosage. While the combination of TGF-β, ATRA and rapamycin has been shown to generate highly suppressive iTregs, the challenge for therapeutic iTreg generation has been their instability. Here, we investigate the impact of rapamycin concentrations and α-CD3/CD28 bead ratios on human iTreg stability. Methods. We assess iTregs generated with various concentrations of rapamycin and differing ratios of α-CD3/CD28 beads for their differentiation, stability, expression of Treg signature molecules and T helper effector cytokines, and Treg-specific demethylation region (TSDR) status. Results. iTregs generated in the presence of TGF-β, ATRA, rapamycin and a higher ratio of α-CD3/CD28 beads were highly suppressive and stable upon in vitro re-stimulation. These iTregs exhibited a similar expression profile of Treg signature molecules and T helper effector cytokines to nTregs, in the absence of TSDR demethylation. Conclusion. This work establishes a method to generate human iTregs which maintain stable phenotype and function upon in vitro re-stimulation. Further validation in pre-clinical models will be needed to ensure its suitability for applications in adoptive transfer.
INTRODUCTION

Regulatory T cells (Tregs) are a sup-population of CD4+ T cells with immune-suppressive and immune-modulatory properties. With such properties, Tregs form a vital part of immune homeostasis, providing tolerance to self and non-pathogenic foreign antigens, and down-regulating immune responses once pathogens are cleared in order to minimise tissue-damage.1 Currently, human natural Tregs (nTregs) are defined by the surface phenotype CD4+CD25hiCD127lo, and expression of the master regulator of Treg-lineage, FOXP3.2-7 Within the CD4+CD25hiCD127FOXP3+ population, there are two main subgroups of nTregs: thymic-derived thymic Tregs (tTregs) and peripherally induced peripheral Tregs (pTregs). tTregs are generated within the thymus from self-reactive CD4+ T cells and account for 5–10% of circulating CD4+ T cells.8 Most self-reactive CD4+ T cells are negatively selected and deleted by apoptosis to establish central tolerance; however, some self-reactive CD4+ T cells with relatively high affinity for self-antigens receive signals to differentiate into nTregs by the induction of FOXP3 expression.9 pTregs express a T-cell receptor (TCR) repertoire skewed towards foreign antigens allowing them to curtail autoimmune responses by self-reactive conventional T cells that have escaped central tolerance and thus prevent autoimmunity.10,11 In contrast, peripheral naive CD4+ T cells, which normally account for 30–50% of circulating CD4+ T cells in adults aged between 18 to 70,12,13 can acquire FOXP3 expression upon activation, becoming pTregs. As pTregs are differentiated from conventional T cells, they express a TCR repertoire skewed towards foreign antigens and help establish tolerance to commensal microbiota, environmental and food allergens, and foetal alloantigens during pregnancy.11,14 The induction of pTregs can be mimicked in vitro to generate induced Tregs (iTregs).14,15

iTregs provide an attractive alternative to Treg-based immunotherapies in allogeneic transplantation. Currently, most Treg-based immunotherapies employ adoptive transfer of ex vivo expanded nTregs.16 To generate the required number of cells for clinical dosage, which requires up to 5 billion Tregs per patient, nTregs need to be ex vivo expanded for a prolonged period of time due their low frequency in peripheral blood.17,18 Thus, it would be beneficial to generate large number of Tregs in a shorter time frame, by differentiating iTregs from naive CD4+ T cells, which are at a significantly higher frequency in peripheral blood. Furthermore, the TCR repertoire of iTregs is potentially more relevant for allogeneic transplantation, as rejection of the donor tissue occurs in response to foreign antigens against which the nTreg pool has not been educated.15 In addition, the broader TCR repertoire of iTregs compared with nTregs has advantages in the generation of antigen-specific Tregs, potentially providing a more targeted therapy.14,19

The pivotal point in the generation of iTregs has been the discovery of differentiation induction molecules converting naive CD4+ T cells into pTregs. One particular environment in which pTregs are present in significant numbers is the gut.20,22 The gut mucosal environment contains TGF-β, all-trans retinoic acid (ATRA), and short chain fatty acids, such as butyrate, which have been shown to promote pTreg differentiation.22,23 Additionally, manipulation of ex vivo iTreg generation has validated various molecules such as IL-2, rapamycin and progesterone as enhancer of pTreg differentiation.15 To date, different approaches using combinations of these molecules have been explored to generate human iTregs.24-32 In particular, the combination of TGF-β, ATRA and rapamycin has been shown to generate highly suppressive iTregs that are stable upon resting26; however, phenotypic instability of iTregs upon re-stimulation remains a major challenge significantly limiting the use of iTregs for therapeutic applications.

Here, we aimed to optimise an iTreg differentiation method for the robust and reproducible production of iTregs with phenotypic and functional stability in vitro. Various concentrations of rapamycin have been utilised previously for the induction of human iTregs, ranging from 0.45 to 100 ng mL−1.26,27,29,33-36 In addition, the effect of rapamycin concentration (1, 10 and 100 ng mL−1) on the proportion of cells...
that differentiate into iTregs has been studied; however, the effect on iTreg stability remains unknown. Thus, we investigated which concentration of rapamycin is optimal for the generation of stable iTregs, comparing low-dose rapamycin of 1 ng mL\(^{-1}\), medium-dose rapamycin of 10 ng mL\(^{-1}\) and high-dose rapamycin of 100 ng mL\(^{-1}\). Various TCR stimulation methods appear across the literature for the generation of human iTregs, with no clear consensus. In particular, different ratios of \(\alpha\)-CD3/CD28 beads (3:1 and 1:10 bead to cell) have been utilised. Hence, we explored the impact of TCR stimulation bead ratio by using 1:1 and 1:10 \(\alpha\)-CD3/CD28 beads to represent ‘high’ and ‘low’ ratios of \(\alpha\)-CD3/CD28 beads, respectively. We assessed phenotype and suppressive function of iTregs generated under various conditions after initial stimulation and upon re-stimulation without iTreg-differentiating factors. We further characterised the functionality of iTregs by measuring expression of Treg signature molecules and T helper signature cytokines. In addition, phenotypic stability and T helper signature cytokine production of iTregs upon challenge with Th17-polarising cytokines were evaluated.

Lastly, Treg-specific demethylation region (TSDR) methylation status of iTregs was analysed.

**RESULTS**

**Phenotype and function of iTregs generated with varied rapamycin concentration**

To investigate the impact of rapamycin concentration on human iTreg induction, naïve CD4\(^+\) T cells and naïve nTregs were isolated from human peripheral blood (Supplementary figure 1). Naïve CD4\(^+\) T cells were differentiated into iTregs over a 7-day stimulation using TGF-\(\beta\), ATRA, IL-2, \(\alpha\)-CD3/CD28 beads (1:1 bead to cell) and various concentrations of rapamycin (0, 1, 10 and 100 ng mL\(^{-1}\); these iTregs were termed iTreg-0, iTreg-1, iTreg-10 and iTreg-100, respectively). Naïve nTregs were stimulated using IL-2 and \(\alpha\)-CD3/CD28 beads (1:1 bead to cell) as a positive control for Treg phenotype and function (nTreg), and naïve CD4\(^+\) T cells were stimulated using IL-2 and \(\alpha\)-CD3/CD28 beads (1:1 bead to cell) without iTreg differentiation components as a mock stimulation control (Tconv). Following the 7-day stimulation, all cell types were rested for 7 days in the presence of IL-2 and were assessed after 3 days and 7 days of rest. After a 3-day rest, cells were assessed for expression of Treg markers FOXP3 and CD25. FOXP3 and CD25 expression levels were measured via flow cytometry and expressed as a percentage (%FOXP3/CD25\(^+\)) of the viable CD4\(^+\) population, and as protein expression levels by MFI (Supplementary figure 2). In each experiment, CD25 and FOXP3 MFI were normalised to the highest raw MFI and represented as nMFI (%).

Tconv cells were 77% FOXP3/CD25\(^+\), iTreg-0 cells were 87% FOXP3/CD25\(^+\), nTreg and iTreg-1 cells were 91% FOXP3/CD25\(^+\), and iTreg-10 and iTreg-100 cells were 95% FOXP3/CD25\(^+\). %FOXP3/CD25\(^+\) of Tconv was significantly lower than nTreg, iTreg-1, iTreg-10 and iTreg-100 (\(P = 0.0398, 0.0343, 0.0105\) and 0.0088). FOXP3 MFI of Tconv was significantly lower than nTreg, iTreg-10 and iTreg-100 (\(P = 0.0121, 0.0017\) and 0.0011) while FOXP3 MFI of iTreg-0 was significantly lower compared with iTreg-10 and iTreg-100 (\(P = 0.0080\) and 0.0049). CD25 MFI of iTreg-10 and iTreg-100 were significantly higher than other cell types (Figure 1a; raw MFI for CD25 and FOXP3 shown in Supplementary figure 3; nTreg: \(P = 0.0228\) and 0.0180, Tconv: \(P = 0.0017\) and 0.0013, iTreg-0: \(P = 0.0217\) and 0.0172, and iTreg-1: \(P = 0.0330\) and 0.0260). Generated iTregs were then assessed for their suppressive function. Cell Trace Violet (CTV) was used to track cell proliferation of responder cells (naïve CD4\(^+\) T cells) in the presence of \(\alpha\)-CD3/CD28 beads (1:5 bead to cell) and different ratios of Tregs (1:1 to 1:8 Treg to Tresponder). Based on the positive proliferation control (CTV-stained and stimulated with no Tregs), suppression was calculated (Supplementary figure 4). Suppressive activities of iTreg generated in the presence of rapamycin showed no significant differences compared with nTreg, while Tconv and iTreg-0 displayed significantly lower suppressive activities than nTreg, iTreg-1, iTreg-10 and iTreg-100 at most Treg:Tresponder ratios (1:1 – Tconv: \(P = 0.0409, 0.0250, 0.0233\) and 0.0250, 1:2 – Tconv: \(P < 0.0001\) for all and iTreg-0: \(P = 0.0688, 0.0331, 0.0226\) and 0.0250, 1:4 – Tconv: \(P < 0.0001\) for all and iTreg-0: \(P = 0.0020, P < 0.0001, 0.0001\) and 0.0001, 1:8 – Tconv: \(P < 0.0001\) for all and iTreg-0: \(P = 0.0038, P < 0.0001, 0.0001\) and 0.0001). Additionally, suppressive activities of Tconv were significantly lower than iTreg-0 (Figure 1b; 1:2 – \(P = 0.0387\) and 1:4 – \(P = 0.0336\)). After a 7-day rest, FOXP3 and CD25 expression were assessed again.
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iTreg-0, iTreg-1, iTreg-10 and iTreg-100 retained their %FOXP3*CD25* at 87%, 78%, 86%, 91% and 92%, respectively. % FOXP3*CD25* of Tconv was significantly lower than iTreg-0, iTreg-1, iTreg-10 and iTreg-100 at 45% (P < 0.0001, P = 0.0004, P < 0.0001, 0.0001 and 0.0001, respectively). While no significant differences in FOXP3 MFI between Tconv and other cell types were observed, CD25 MFI of iTreg-0 was significantly lower than iTreg-1, iTreg-10 and iTreg-100 (Figure 1a; raw MFI for CD25 and FOXP3 shown in Supplementary figure 5; iTreg-0: P = 0.0096 and 0.0045 and iTreg-100: P = 0.0022 and 0.0011). Furthermore, cell growth rates at the end of the stimulation cycle were assessed, which showed iTreg-10 and iTreg-100 with significantly lower cell growth rate compared with nTreg, Tconv and iTreg-0 (Supplementary figure 6; iTreg-10: P = 0.0135, 0.0578, 0.0267 and iTreg-100: P = 0.0064, 0.0267, 0.0021).

Reductions in FOXP3 expression and suppressive activities have been observed in human iTregs upon re-stimulation without iTreg differentiation components,25,26 which could occur in vivo resulting in generation of non-functional ex-Tregs. Thus, the impact of rapamycin concentration on human iTreg stability upon in vitro re-stimulation was evaluated. Following a 7-day stimulation and 3-day rest, cells were re-stimulated using IL-2 and α-CD3/CD28 beads (1:1 bead to cell) without the iTreg differentiation factors for 7 days, then extensively washed and rested for 3 days in the presence of IL-2. After re-stimulation, FOXP3 and CD25 expression and suppressive function of cells were assessed. nTreg, iTreg-1, iTreg-10 and iTreg-100 retained their % FOXP3*CD25* at 89%, 81%, 89% and 90%. % FOXP3*CD25* of iTconv was significantly lower than nTreg, iTreg-0, iTreg-1, iTreg-10 and iTreg-100 at 38% (P < 0.0001, P = 0.0007, P < 0.0001, 0.0001 and 0.0001). iTreg-0 exhibited significantly lower %FOXP3*CD25* than nTreg, iTreg-1 and iTreg-10 and iTreg-100 at 66% (P = 0.0032, 0.0127, 0.0038 and 0.0028). FOXP3 MFI of Tconv and iTreg-0 were significantly lower than nTreg (P = 0.0349 and 0.0496). Tconv showed significantly lower CD25 MFI compared with nTreg, iTreg-1, iTreg-10 and iTreg-100 (P = 0.0002, 0.0011, 0.0003 and 0.0002), and iTreg-0 displayed significantly lower CD25 MFI than nTreg, iTreg-10 and iTreg-100 (Figure 2a; raw MFI for CD25 and FOXP3 shown in Supplementary figure 7; P = 0.0053, 0.0085 and 0.0051). Furthermore, iTreg-1, iTreg-10 and iTreg-100 retained their suppressive function with no significant differences compared with nTreg. Tconv and iTreg-0 exhibited significantly lower suppressive activities than nTreg, iTreg-1, iTreg-10 and iTreg-100 (Figure 2b; 1:1 – Tconv: P = 0.0001, 0.0001, P < 0.0001 and 0.0001, and iTreg-10: P = 0.0134, 0.0125, 0.0036 and 0.0055, 1:2 – Tconv: P < 0.0001, P = 0.0006, 0.0001 and 0.0003, iTreg-10: P = 0.0012, 0.0375, 0.0098 and 0.0202, 1:4 – Tconv: P < 0.0001, P < 0.0099, 0.0046 and 0.0037, iTreg-10: P = 0.0019, 0.1149, 0.0630 and 0.0525 and 1:8 – Tconv: P = 0.0356, 0.1588, 0.0427 and 0.0315). Notably, assessment of cell growth rates upon re-stimulation revealed significantly higher fold expansion in iTreg-10 and iTreg-100 than in nTreg, Tconv and iTreg-0 (iTreg-10: P = 0.0303, 0.0005 and 0.0010, and iTreg-100: P = 0.0042, 0.0001 and 0.0002). Moreover, cell growth rate of iTreg-1 was significantly higher than Tconv and iTreg-0 (Supplementary figure 8; P = 0.0051 and 0.0114).

Phenotype and function of iTregs generated with differing ratios of α-CD3/CD28 beads

To investigate the impact of TCR stimulation bead ratios on human iTreg induction, iTregs were generated from naïve CD4+ T cells using TGF-β, ATRA, 100 ng mL⁻¹ rapamycin, IL-2 and differing ratios of α-CD3/CD28 beads (1:1 and 1:10 bead to cell; termed iTreg-10 as previous and iTreg-1:10). As above, stimulated naïve nTreg and naïve CD4+ T cells were used as controls (nTreg and Tconv), and cells were stimulated and rested for 7 days each. After 3 days of rest, cells were assessed for their FOXP3 and CD25 expression and suppressive function. iTreg-1:10 showed significantly lower % FOXP3*CD25* than nTreg (91%), Tconv (85%) and iTreg-100 (95%) at 54% (P = 0.0015, 0.0036 and 0.0009). This was accompanied with a significant difference in FOXP3 MFI between iTreg-1:10 and nTreg (P = 0.0452). Additionally, Tconv showed significantly lower FOXP3 MFI than nTreg (P = 0.0422). CD25 MFI of iTreg-100 was significantly higher than nTreg, Tconv and iTreg-1:10 (Figure 3a; raw MFI for CD25 and FOXP3 shown in Supplementary figure 9; P = 0.0002, 0.0003 and P < 0.0001). The suppressive activities...
Figure 1. The impact of rapamycin concentrations on iTreg differentiation. Naïve nTregs were stimulated with a 1:1 α-CD3/CD28 beads and IL-2 (nTreg). Naïve CD4\(^+\) T cells were mock-stimulated with a 1:1 α-CD3/CD28 beads and IL-2 (Tconv). iTregs were differentiated from naïve CD4\(^+\) T cells using a 1:1 α-CD3/CD28 beads, IL-2 TGF-β, ATRA and various concentrations of rapamycin (0, 1, 10 and 100 ng mL\(^{-1}\); iTreg-0, iTreg-1, iTreg-10 and iTreg-100). Following a 7-day stimulation, cells were rested up to 7 days with IL-2. After 3 days of rest, (a) expression of FOXP3 and CD25 (b) suppressive activities were evaluated. After 7 days of rest, (c) expression of FOXP3 and CD25 was evaluated. MFI of FOXP3 and CD25 were normalised to highest raw MFI value in each experiment and represented as nMFI (%). Raw MFI for (a and c) Supplementary figures 3 and 5, respectively. For (b), grey-shaded histogram represents positive control (no Treg control). Data are represented as mean \pm SE, N = 3 in three independent experiments. For each donor (N), technical triplicates were utilised, and the average of technical replicates was used for each datapoint. Statistical significance identified by RM one-way (a and c) and RM two-way (b) ANOVA with Dunnett’s multiple comparisons test: *P < 0.05, **P<0.01, ***P<0.001, ****P<0.0001.
Figure 2. The impact of rapamycin concentrations on iTreg stability upon re-stimulation. Following a 7-day stimulation and a 3-day rest, cells were re-stimulated using a 1:1 α-CD3/CD28 beads and IL-2 without iTreg differentiation factors for 7 days then rested with IL-2 for 3 days. After 3 days of rest, (a) expression of FOXP3 and CD25 (b) suppressive activities were evaluated. MFI of FOXP3 and CD25 were normalised to highest raw MFI value in each experiment and represented as nMFI (%). Raw MFI for (a) is shown in Supplementary figure 7. For (b), grey-shaded histogram represents positive control (no Treg control). Data are represented as mean ± SEM, N = 3 in three independent experiments. For each donor (N), technical triplicates were utilised, and the average of technical replicates was used for each datapoint. Statistical significance identified by RM one-way (a) and RM two-way (b) ANOVA with Dunnett’s multiple comparisons test: *P < 0.05, **P<0.01, ***P<0.001, ****P<0.0001.
of Tconv were significantly lower than nTreg, iTreg-100 and iTreg-1:10 (Figure 3b; 1:2 – $P = 0.0348, 0.0009$ and $0.0493$, and 1:4 – $P = 0.0109, 0.0002$ and $0.0479$, and 1:8 – $P = 0.04151, 0.0316$ and $0.6079$). After a 7-day rest, expression of FOXP3 and CD25 were further assessed. %FOXP3$^+$CD25$^+$ of Tconv and iTreg-1:10 were significantly lower than nTreg (86%) and iTreg-100 (93%) at 38% and 48%, respectively (Tconv: $P = 0.0027$ and 0.0012, and iTreg-1:10: $P = 0.0090$ and 0.0037). FOXP3 MFI of iTreg-100 were significantly higher than nTreg and Tconv ($P = 0.0436$ and 0.0244). CD25 MFI of iTreg-100 remained significantly higher than nTreg, Tconv and iTreg-1:10 ($P = 0.0174, 0.0002$, and 0.0002), and nTreg exhibited significantly higher CD25 MFI than Tconv and iTreg-1:10 (Figure 3c; raw MFI for CD25 and FOXP3 shown in Supplementary figure 10 $P = 0.0039$ for both). Additionally, evaluation of cell growth rates showed that fold expansion of iTreg-1:10 is significantly lower than nTreg, Tconv and iTreg-100 ($P = 0.0064, 0.0010$ and 0.0391). Cell growth rate of iTreg-100 was significantly lower than in Tconv (Supplementary figure 11; $P = 0.0179$).

Following this up, the impact of TCR stimulation bead ratios on human iTreg stability upon in vitro re-stimulation was assessed. As above, after initial stimulation and rest, cells were re-stimulated using IL-2 and α-CD3/CD28 beads (1:1 bead to cell) for 7 days, then rested for 3 days in the presence of IL-2. Upon re-stimulation, FOXP3 and CD25 expression and suppressive activities of cells were evaluated. Significantly lower %FOXP3$^+$CD25$^+$ were observed in Tconv and iTreg-1:10 than in nTreg (91%) and iTreg-100 (91%) at 52% and 63%, respectively (Tconv: $P = 0.0042$ and 0.0039, and iTreg-1:10: $P = 0.0185$ and 0.0168). While Tconv showed significantly lower FOXP3 MFI than nTreg and iTreg-100 ($P = 0.0078$ and 0.0244), iTreg-1:10 displayed no significant differences in FOXP3 MFI compared with nTreg and iTreg-100. CD25 MFI showed similar trends to %FOXP3$^+$CD25$^+$, with Tconv and iTreg-1:10 exhibiting significantly lower CD25 MFI than nTreg and iTreg-100 (Figure 4a; raw MFI for CD25 and FOXP3 shown in Supplementary figure 12; Tconv: $P = 0.0011$ and 0.0003, and iTreg-1:10: $P = 0.0016$ and 0.0004). Moreover, no significant differences were shown with suppressive activities of nTreg, iTreg-100 and iTreg-1:10, while Tconv displayed significantly lower suppressive activities compared with nTreg, iTreg-100 and iTreg-1:10 (Figure 4b, 1:1 – $P = 0.0582, 0.0062$ and 0.0167, 1:2 – $P = 0.0009, 0.0002$ and 0.0009, and 1:4 – $P = 0.0010, 0.0030$ and 0.0206). In addition, cell growth rates upon re-stimulation were measured. Tconv exhibited significantly lower fold expansion compared with iTreg-100 and iTreg-1:10 ($P = 0.0179$ and 0.0117), and cell growth rate of nTreg was significantly lower than iTreg-1:10 (Supplementary figure 13; $P = 0.494$).

**Functional characterisation of iTregs generated with varied concentrations of rapamycin and differing ratios of α-CD3/CD28 beads**

To provide a broader understanding of iTregs generated under different conditions, their functionality was further characterised. After an initial 7-day stimulation and 3-day rest, expression levels of Treg signature molecules and CD4$^+$ T cell effector cytokines were assessed by measuring CTLA4 and IL2 gene expression and production of active TGF-β, IL-4, IL-6, IL-10, TNF-α, IFN-γ and IL-17A. While no significant differences were observed in CTLA4 expression amongst cell types (Figure 5a), Tconv exhibited significantly higher IL2 expression compared with nTreg, iTreg-1, iTreg-10, iTreg-100 and iTreg-1:10 (Figure 5b; $P = 0.0415, 0.0423, 0.0423, 0.0430$ and 0.0445). No significant differences were shown in production of active TGF-β (Figure 5c), IL-4, IL-6 and IL-17A. Furthermore, IL-10 production in nTreg was significantly higher than Tconv, iTreg-0, iTreg-1, iTreg-10, iTreg-100 and iTreg-1:10 ($P = 0.0401, 0.0489, 0.0493, 0.0400, 0.0391$ and 0.0387). Tconv produced significantly more TNF-α compared with nTreg, iTreg-0, iTreg-1, iTreg-10, iTreg-100 and iTreg-1:10 ($P < 0.0001, P = 0.0130, 0.0012, 0.0002, 0.0002$ and 0.0192), and iTreg-0 and iTreg-1:10 produced significantly more TNF-α compared with nTreg ($P = 0.0102$ and 0.0070). Lastly, IFN-γ production in Tconv was significantly greater than nTreg, iTreg-0, iTreg-1, iTreg-10, iTreg-100 and iTreg-1:10 (Figure 5d; $P = 0.0017, 0.0056, 0.0021, 0.0017$ and 0.0016).

**Th17-polarising challenge of iTregs generated with varied concentrations of rapamycin and differing ratios of TCR stimulation bead**

It has previously been shown that iTregs can convert to pathogenic Th17 cells in a pro-inflammatory...
Figure 3. The impact of α-CD3/CD28 bead ratios on iTreg differentiation. Naive nTregs were stimulated with a 1:1 α-CD3/CD28 beads and IL-2 (nTreg). Naive CD4⁺ T cells were mock-stimulated with a 1:1 α-CD3/CD28 beads and IL-2 (Tconv). iTregs were differentiated from naive CD4⁺ T cells using IL-2 TGF-β, ATRA and 100 ng mL⁻¹ rapamycin and various ratios of α-CD3/CD28 beads (1:1 and 1:10 bead to cell; iTreg-100 as before and iTreg-1:10). Following a 7-day stimulation, cells were rested up to 7 days with IL-2. After 3 days of rest, (a) expression of FOXP3 and CD25 (b) suppressive activities were evaluated. After 7 days of rest, (c) expression of FOXP3 and CD25 was evaluated. MFI of FOXP3 and CD25 were normalised to highest raw MFI value in each experiment and represented as nMFI (%). Raw MFI for (a and c) is shown in Supplementary figures 9 and 10, respectively. For (b), grey-shaded histogram represents positive control (no Treg control). Data are represented as mean ± sem, N = 3 in three independent experiments. For each donor (N), technical triplicates were utilised, and the average of technical replicates was used for each datapoint. Statistical significance identified by RM one-way (a and c) and RM two-way (b) ANOVA with Dunnett’s multiple comparisons test: *P < 0.05, **0.01, ***0.001, ****0.0001.
environment, and that challenging iTregs with Th17-polarising cytokines can compromise their FOXP3 expression and protective abilities in vivo. Therefore, the stability of iTregs upon Th17-polarising challenge was investigated by culturing them for 3 days with Th17-polarising cytokines IL-1β, IL-6, IL-21, and IL-23, α-CD3/CD28 beads (1:10 bead to cell) and low-dose IL-2, after initial 7-day stimulation and 3-day rest. Low-dose IL-2 was utilised to maintain cell viability with minimal
stabilisation of FOXP3 expression and inhibition of Th17 polarisation during the challenge. After the challenge, FOXP3 and CD25 expression, production of CD4+ T cell cytokines and gene expression of Th17 master regulator, RORC, were assessed. While no significant differences were observed in %FOXP3+CD25+ and FOXP3 MFI for all cell types after the challenge, challenged nTreg, Tconv, iTreg-0, iTreg-1, iTreg-10, iTreg-100 and iTreg-1:10 displayed significant increases in their CD25 MFI compared with ‘unchallenged’ cells (Figure 6a; raw MFI for CD25 and FOXP3...
shown in Supplementary figure 14; \( P = 0.0439, 0.0239, 0.0318, 0.0040, 0.0370, 0.0228 \) and 0.0097). All cell types exhibited no significant differences in production of IL-17A (Figure 6b). Additionally, no significant differences were shown with IL-2, IL-4, IL-6, IL-10, TNF-\( \alpha \) and IFN-\( \gamma \) after the challenge (Supplementary figure 15). Notably, both unchallenged and challenged cells lacked RORC expression (Supplementary figure 16).

**TSDDR methylation status of iTreg generated with varied concentrations of rapamycin and differing ratios of a-CD3/CD28 beads**

The demethylation status of Treg-specific demethylation region (TSDDR) is an important indicator of stability of FOXP3 expression.\(^{47,48} \) Methylation levels of 11 CpG motifs in the TSDDR were measured via targeted bisulphite pyrosequencing. DNA was collected after an initial 7-day stimulation and 3-day rest. TSDDR methylation levels of Tconv, iTreg-0, iTreg-1, iTreg-10, iTreg-100 and iTreg-1:10 were above 90% in average with each CpG motifs showing various levels of methylation between 78% and 100%, while nTreg showed TSDDR methylation level of 45% in average with each CpG motifs showing various levels of methylation between 40% and 67% (Figure 7).

**DISCUSSION**

In this study, we optimised an iTreg differentiation method utilising TGF-\( \beta \), ATRA, rapamycin, IL-2 and a-CD3/CD28 beads to differentiate and expand human iTregs from naive CD4\(^+\) T cells, which are 7- and 20-fold more frequent in human peripheral blood compared with nTregs and naive nTregs. (Supplementary figure 17). These molecules have previously been shown to induce or enhance iTreg generation. TGF-\( \beta \) secreted by CD103\(^+\) intestinal dendritic cells (DCs) plays an essential role in the generation of pTregs in the gut by inducing binding of the transcription factors Smad2 and Smad3 to conserved non-coding DNA sequence 1 (CNS1) region of FOXP3 locus,\(^ {14,15} \) which is crucial for FOXP3 induction in pTregs but not in nTregs.\(^ {49} \) Additionally, ATRA secreted by CD103\(^+\) intestinal DCs reinforces binding of Smad3 to CNS1 region through histone acetylation of Smad3-binding region, preventing potential Th17 polarisation.\(^ {14,15,50} \) Furthermore, the mTOR (mammalian target of rapamycin)-targeting drug, rapamycin, has been shown to induce FOXP3 expression, promote expansion and stabilise FOXP3 expression via inhibition of mTORC1 complex.\(^ {51,52} \) Rapamycin also enhances purity of generated iTregs by selectively inhibiting the activation of conventional T cells,\(^ {8} \) as Tregs utilise an IL-2R-dependent STAT5 pathway for activation while conventional T cells require the mTOR pathway for activation.\(^ {53,54} \) Lastly, IL-2 promotes FOXP3 expression and Treg expansion, and inhibits Th17 polarisation.\(^ {15,55} \) The combination of TGF-\( \beta \), ATRA, rapamycin and IL-2 has previously been used to successfully generate iTregs with superior in vitro suppressive activities.\(^ {26,27} \) However, these iTregs lost FOXP3 expression upon re-stimulation and were then not able to prevent the onset of xenogenic graft-versus-host disease in a humanised mouse model.\(^ {26} \)

We demonstrated that any concentration of rapamycin in conjunction with 1:1 beads robustly generate highly suppressive iTregs that are phenotypically and functionally stable upon in vitro re-stimulation, without compromising cell viability (Supplementary figure 18). While all three concentrations of rapamycin-induced high expression of FOXP3 and CD25, medium- and high-dose rapamycin resulted in elevated FOXP3 and CD25 expression after initial stimulation, compared with low-dose rapamycin. Conversely, iTregs generated without rapamycin were not as suppressive as nTregs or rapamycin-induced iTregs despite having comparable FOXP3 and CD25 expression to nTregs. These iTregs lost FOXP3 and CD25 expression upon re-stimulation. This demonstrated the importance of rapamycin for suppressor function and stability upon in vitro re-stimulation. Interestingly, with 1:10 beads, FOXP3 and CD25 expression of rapamycin-induced iTregs were not comparable to nTregs and yet were highly suppressive after initial stimulation and upon re-stimulation. This was in line with a previously published protocol utilising the combination of TGF-\( \beta \), ATRA, high-dose rapamycin and 1:10 beads, even though the authors reported CD25-FOX3\(^+\) iTregs prior to their functional analysis.\(^ {27} \) Moreover, with 1:10 beads, extremely limited cell growth was observed, which together indicated cells have limited accessibility to beads when 1:10 beads are utilised. iTregs generated with 1:10 beads showed stabilisation of FOXP3 expression similar to nTregs after a 7-day rest and upon re-stimulation, while their CD25 expression remained low compared
Figure 6. The impact of rapamycin concentrations and α-CD3/CD28 bead ratios on iTreg stability upon challenge with Th17-polarising cytokines. Following a 7-day stimulation and a 3-day rest, cells were challenged using a 1:10 α-CD3/CD28 beads, low-dose IL-2, IL-1β, IL-6, IL-21 and IL-23 for 3 days. Unchallenged controls were stimulated using a 1:10 α-CD3/CD28 beads and low-dose IL-2 for 3 days. After 3 days of challenge, (a) expression of FOXP3 and CD25 and (b) production of IL-17A were assessed. MFI of FOXP3 and CD25 were normalised to highest raw MFI value in each experiment and represented as nMFI (%). Raw MFI for (a) is shown in Supplementary figure 14. For (b), the cells were stimulated with PMA and ionomycin for 6 hours prior to collection of cell supernatants. Cell-free medium was used as a control to measure background cytokine levels in the medium which was supplemented with heat inactivated human serum. Background cytokine levels in the medium were negligible (below the lowest standards). Data are represented as mean ± sem, N = 3 in three independent experiments (a) or in one independent experiment (b). For each donor (N), technical triplicates (a) or technical duplicates (b) were utilised, and the average of technical replicates was used for each datapoint. Statistical significance identified by a paired t-test: *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001.
with nTregs and rapamycin-induced iTregs with 1:1 beads. Together, these suggested that using a high ratio of α-CD3/CD28 beads is beneficial for generation of iTregs. In addition, while we compared different ratios of α-CD3/CD28 beads, direct comparison to other forms of TCR stimulation, such as plate-bound α-CD3 and soluble α-CD28,24,26 would provide valuable insights, as different forms would present the cells with varying TCR stimulation intensity and density. Indeed, the importance of TCR signal intensity on iTreg generation has been extensively investigated. While some have demonstrated that sub-optimal TCR stimulation, which results in low AKT activation, favors FOXP3 induction and iTreg generation,56–58 it has also been shown that low doses of strong TCR stimulation induce stable and persistent iTregs.59 Given this controversy, it has been proposed that the ‘Goldilocks principle’, which has been associated with development of tTregs,60 could apply to iTregs, requiring a fine balance in TCR stimulation with the stimulation being ‘just right’.61,62 Furthermore, while CD28 co-stimulation is necessary for the development of tTregs, it has been shown to promote mTOR activation and inhibit differentiation of naive T cells into iTregs.62–64 Thus, it is possible that rapamycin counteracts activation of AKT/mTOR pathways driven by α-CD3 and α-CD28 on the beads,26 ultimately modulating TCR stimulation intensity to favor iTreg generation, which could explain why iTregs generated with medium- and high-dose rapamycin displayed reduced cell growth.

Notably, with 1:1 α-CD3/CD28 beads, even mock-stimulated Tconvs highly expressed FOXP3 and CD25, which were only lost upon in vitro re-stimulation. In contrast, previously published protocols utilising 1:10 α-CD3/CD28 beads and plate-bound α-CD3 and soluble α-CD28 showed moderate expression of FOXP3 in mock-stimulated Tconvs. Evidently, it has been shown that stimulation-induced FOXP3 expression in Tconvs is dependent on the amount of stimulation used. In particular, almost 100% of Tconvs exhibited FOXP3 expression similar to nTregs under 1:1 α-CD3/CD28 beads, while less than 40% of Tconvs expressed FOXP3 under 1:16 α-CD3/CD28 beads or plate-bound α-CD3 and soluble α-CD28.65 In addition, mock-stimulated Tconvs exhibited moderate suppressive activities, which was in line with murine Tconvs acquiring potent suppressor function upon activation.66 Dose-dependent suppression shown with mock-stimulated Tconvs was comparable to iTregs generated without rapamycin at some ratios, which was shown with
a previously published protocol. Of note, in this protocol, mock-stimulated Tconv and iTregs generated without rapamycin displayed substantial differences in their FOXP3 expression, which suggests that suppressive capacity in these cells may be unspecific suppression instead of being induced by FOXP3 expression. This highlights the need for assessment of phenotype and function upon in vitro re-stimulation for therapeutic applications of Treg products, given the used stimulation can induce non-Treg-like and non-stable FOXP3. Furthermore, mock-stimulated Tconv and ‘no rapamycin’ iTregs displayed limited cell growth upon re-stimulation. This suggests that perhaps 1:1 α-CD3/CD28 beads over-stimulated these cells, inducing T-cell exhaustion, while rapamycin-induced iTregs were able to expand upon re-stimulation, potentially as a result of rapamycin dampening TCR-driven exhaustion.

Tregs have versatile modes of suppression via expression of various suppressive molecules and also are able to produce T helper effector cytokines. Interestingly, expression levels of canonical Treg suppressive molecules in cell types – no differences observed in active TGF-β production, higher IL-10 production in nTregs, and differences shown in CD25 expression – did not correlate with their suppressive activities. Of note, while no differences were shown in CTLA4 gene expression, CTLA4 functionality is largely controlled by externalisation of CTLA4 proteins to the cell surface, thus this did not indicate much. In addition, while it has been previously shown that expression levels of FOXP3 can be directly correlated with suppressive activity, FOXP3 expression levels after initial stimulation were not indicative of their suppressive activities. Furthermore, rapamycin-induced iTregs with 1:1 α-CD3/CD28 beads were most comparable to nTregs in terms of their T helper effector cytokine production profile. Expectedly, mock-stimulated Tconv displayed most discrepancies, showing high expression of IL2 and producing more pro-inflammatory cytokines such as TNF-α and IFN-γ. While nTregs and rapamycin-induced iTregs exhibited repressed IL2 expression in line with nTregs not being able to produce IL-2 for autocrine signalling, iTregs generated without rapamycin displayed comparable IL2 expression to mock-stimulated Tconv. In addition, TNF-α production by iTregs generated without rapamycin and rapamycin-induced iTregs with 1:10 α-CD3/CD28 beads were comparable to mock-stimulated Tconv. These reinforced the importance of rapamycin and high ratio of α-CD3/CD28 beads for iTreg differentiation. Notably, while TNF-α production in rapamycin-induced iTregs with 1:1 α-CD3/CD28 beads was substantially lower than mock-stimulated Tconv, they still produced a considerable amount of TNF-α. Considering the pro-inflammatory characteristics of TNF-α, this could affect in vivo functionality of these cells, despite showing no differences in suppression of responder T-cell proliferation compared with nTregs in vitro. Indeed, suppression of cytokine production in responder T cell is another aspect of Treg-mediated suppression, which may be altered by TNF-α. Overall, in-depth analysis of transcriptomes and proteomes may be required to fully understand which factors and genes are attributing to discrepancies in the suppressor function and the phenotype of these cells. Indeed, recent transcriptomic and proteomic profiling of iTregs generated by multiple differentiation protocols revealed iTreg-specific molecular pathways and molecules, even though these protocols were not able to generate stable iTregs.

The tendency of iTregs to switch to a pro-inflammatory phenotype, such as Th17, in a pro-inflammatory microenvironment, needed to be assessed, as conversion of iTregs into pathogenic pro-inflammatory T cells in vivo is the major challenge to the use of iTregs for therapeutic purposes. In addition, iTregs can be polarised to Th17-like Tregs, which express Th17 signature markers while retaining regulatory phenotype and function, under pro-inflammatory conditions. Given that IL-1β, IL-6, IL-21 and IL-23 are known promoters of Th17 polarisation, mock-stimulated Tconv were expected to display Th17 phenotypes upon challenge. Unexpectedly, all cell types including mock-stimulated Tconv exhibited no differences in their FOXP3 expression, T helper effector cytokine production and RORC gene expression upon challenge with Th17-polarising cytokines, while showing elevated CD25 expression. In particular, negligible amounts of IL-17A were produced, and RORC gene expression was undetectable in all cell types with and without challenge, possibly as a result of high expression of FOXP3 in these cells. While, this is in line with TGF-β/ATRA-induced human iTregs showing increased expression of Treg suppressive molecules, PD-1 and GITR, upon treatment with...
IL-1β and IL-6,_generation without a positive control which expresses Th17 markers upon challenge. Thus, a more physiological environment may be required to validate iTreg stability under pro-inflammatory conditions. Indeed, humanised graft-versus-host disease (GVHD), skin transplantation and islet transplantation models which present in vivo reactivation and pro-inflammatory milieu have been utilised to assess Tregs.\textsuperscript{26–28,29,32,82,83} In conjunction, the safety and effectiveness of iTregs can be corroborated through these humanised models as well, to ensure clinical relevance of this iTreg differentiation protocol.

Currently, demethylation of Treg-specific demethylation region (TSDR), the CpG-rich CNS2 region of FOXP3 locus, is thought to be the hallmark of stable FOXP3 expression, thus stable Treg phenotype.\textsuperscript{14,84,85} In humans, it has been shown that nTregs exhibit full demethylation of TSDR, while Tconvs present fully methylated TSDR.\textsuperscript{48} Even though rapamycin-induced iTregs with 1:1 α-CD3/CD28 beads demonstrated phenotypic and functional stability upon in vitro re-stimulation, demethylation of TSDR in these cells was not observed. Mock-stimulated Tconvs, ‘no rapamycin’ iTregs and rapamycin-induced iTregs with 1:10 α-CD3/CD28 beads also showed no demethylation of TSDR. Thus, this iTreg differentiation protocol could be modified to induce TSDR demethylation. Indeed, hypoxia and vitamin C were shown to enhance expression and activity of TET (ten eleven translocation) enzymes and facilitate demethylation of the TSDR in mice and human.\textsuperscript{86–89} Interestingly, nTregs analysed for methylation status only displayed partial demethylation. As a result of the anonymity of donated buffy coats, we were not able to confirm the gender of the donors, which is crucial for evaluation of TSDR data, as FOXP3 is located on the X chromosome resulting in partial demethylation of TSDR in female caused by X chromosome inactivation.\textsuperscript{90,91} Indeed, given the low inter-donor variation in TSDR demethylation rates of nTregs and similarity of these rates to published data,\textsuperscript{90} it is possible that all donors were females.

In summary, we demonstrated the importance of rapamycin and high ratio of α-CD3/CD28 beads for iTreg differentiation and stability. The combination of TGF-β, ATRA, any concentration of rapamycin, IL-2 and 1:1 α-CD3/CD28 beads generated iTregs which are superior to iTregs generated without rapamycin and iTregs generated with 1:10 α-CD3/CD28 beads. Notably, medium-dose and high-dose rapamycin-induced superior expression of CD25 compared with low-dose rapamycin, without differences in stability and function. These iTregs were highly suppressive, and stable upon in vitro re-stimulation. Expression of Treg signature molecules and T helper effector cytokines in these iTregs was largely comparable to nTregs. Furthermore, these iTregs were stable in the presence of Th17-polarising cytokines with no differences in FOXP3 expression, T helper effector cytokine production and RORC expression; however, further validation in humanised mice models will be required as these results were also observed in mock-stimulated Tconvs. Despite their stability upon in vitro re-stimulation, demethylation of TSDR was not shown for these iTregs. While TSDR demethylation has been thought to be a surrogate for Treg stability, there might be further factors involved in stabilisation of Treg phenotype. Indeed, it has been shown that demethylation of other Treg signature genes such as TNFRSF18, CTLA-4, IKZF4 and IL2RA, dubbed as Treg-specific demethylation patterns (TSDP), is crucial for Treg development and stable FOXP3 expression, as well as TSDR demethylation.\textsuperscript{92}

**METHODS**

**Cell isolation and stimulation**

Human buffy coat (Australian Red Cross) was treated with a RosetteSep Human CD4+ T-cell enrichment cocktail (STEMCELL Technologies, Vancouver, Canada) for 20 min on a platform mixer at 80 rpm. Treated buffy coat was diluted with PBS (+2% foetal calf serum (FCS), Bovogen, Keilor East, Victoria, Australia) prior to isolation of CD4+ T cells by density-gradient centrifugation over Lymphoprep (STEMCELL Technologies). Enriched CD4+ T cells were surface-stained for CD4-APC-H7 (SK3), CD25-PE-Cy7 (M-A251), CD127-FITC (HIL-7R-M21) and CD45RA-PE (HI100; all BD Biosciences, San Jose, CA, USA). CD4+ T cells (naive natural regulatory T cells or naive nTregs) and CD4+ CD25+CD45RA+ T cells (naive CD4+ T cells) were sorted by fluorescence-activated cell sorting (FACS; BD FACSAria Fusion, BD Biosciences). Sorted naive nTregs and naive CD4+ T cells were rested overnight in a complete X-vivo medium at 1 x 10^6 cells mL\(^{-1}\) (X-vivo: serum-free with gentamycin and phenol red, Lonza, Basel, Switzerland). Complete X-vivo medium (cx-vivo) was always supplemented with 2% HEPES (Gibco, Carlsbad, CA, USA), 1% L-glutamine (Hyclone, Logan, UT, USA) and 5% human serum (Heat inactivated; Sigma-Aldrich, St. Louis, MO, USA). 500 U mL\(^{-1}\) of IL-2 (Novartis Vaccines and Diagnostics, Cambridge, MA, USA) was added to complete X-vivo
medium unless otherwise stated. After overnight resting, nTregs were stimulated with 500 U mL⁻¹ of IL-2 and a 1:1 ratio of Human T-expander CD3/CD28 Dynabeads™ (Gibco) in cX-Vivo at 1 × 10⁶ cells mL⁻¹ (nTreg). Naïve CD4⁺ T cells were stimulated with 500 U mL⁻¹ of IL-2 and a 1:1 ratio of expander beads in cX-Vivo containing (1) no other factors (mock stimulation; Tconv) or (2) 5 ng mL⁻¹ of human TGF-β (eBioscience, San Diego, CA, USA), 10 nM of all-trans retinoic acid (ATRA; Sigma-Aldrich), and varying concentrations of rapamycin (0, 1, 10 and 100 ng mL⁻¹). iTreg-0, iTreg-1, iTreg-10 and iTreg-100, respectively; LC Laboratories, Woburn, MA, USA) at 1 × 10⁶ cells mL⁻¹. Naïve CD4⁺ T cells were also stimulated with 500 U mL⁻¹ of IL-2 and a 1:10 ratio of expander beads in cX-Vivo containing 5 ng mL⁻¹ of human TGF-β, 10 nM of ATRA and 100 ng mL⁻¹ of rapamycin (iTreg-1:10). Stimulation was carried out for 7 days. Media was replenished on day 3 and 5 to keep the cell density at 1 × 10⁶ cells mL⁻¹. On day 7 of stimulation, cells were washed three times with PBS (+2% FCS). After washing, expander beads were magnetically removed. Cells were re-suspended in cX-Vivo with 500 U mL⁻¹ of IL-2 at 2 × 10⁶ cells mL⁻¹ and rested up to 7 days. On day 3 of rest, cells were used for flow cytometric analysis, in vitro suppression assay and TSDR methylation assay. On day 7 of rest, cells were used for flow cytometric analysis. Cell cultures were carried out using flat bottom well plates (6, 12, 24 and 48 wells depending on total cell numbers) and culture flasks (T25, T75 and T175 depending on total cell numbers). Within an experiment, culturing formats were usually equivalent.

**iTreg stability evaluation**

For evaluation of iTreg stability, cells were (1) re-stimulated without iTreg differentiation components and (2) challenged with Th17-polarising cytokines. For (1), cells were re-stimulated using 500 U mL⁻¹ of IL-2 and a 1:1 expander beads without addition of TGF-β, ATRA and rapamycin, for 7 days at 1 × 10⁶ cells mL⁻¹ then rested with 500 U mL⁻¹ of IL-2 for 3 days at 2 × 10⁶ cells mL⁻¹, after an initial 7-day stimulation and 3-day rest. On day 3 of rest, cells were used for flow cytometric analysis and in vitro suppression assay. For (2), cells were challenged for 3 days with Th17-polarising cytokines, IL-1β, IL-6, IL-21 and IL-23 (10 ng mL⁻¹ for all, Biolegend, San Diego, CA, USA), in the presence of 25 U mL⁻¹ of IL-2 and a 1:10 expander beads at 1 × 10⁶ cells mL⁻¹, after an initial 7-day stimulation and 3-day rest. Cells were stimulated with 25 U mL⁻¹ of IL-2 and a 1:10 expander beads at 1 × 10⁶ cells mL⁻¹ as unchallenged controls. On day 3 of challenge, cells were washed three times with PBS (+2% FCS). After washing, expander beads were magnetically removed. Cells were used for flow cytometric analysis, cytokine production assays and reverse transcription quantitative polymerase chain reaction.

**Flow cytometric analysis for CD25, FOXP3 and viability**

1 × 10⁶ cells were stained for viability (Fixable Viability Stain 780, FVS780; BD Biosciences) and CD4-FITC (OKT4, BD Biosciences) and CD25-BV421 (M-A251, BD Biosciences). Cells were fixed and permeabilised (Foxp3/Transcription factor staining buffer set, eBioscience) then intracellularly stained for FOXP3-BB700 (236A/E7, BD Biosciences). Compensation controls were utilised and freshly isolated naïve CD4⁺ T cells were used for as a control. All samples were analysed on a BD FACS Canto II flow cytometer (BD Biosciences), and the data analysed with FCS Express 6 (De Novo Software, Pasadena, CA, USA).

**In vitro suppression assay**

Human peripheral blood mononuclear cells (PBMCs) cells were isolated from a fresh buffy coat by density-gradient centrifugation as described above. Naïve CD4⁺ T cells were isolated using EasySep™ Human Naïve CD4⁺ T Cell Isolation kit (STEMCELL Technologies). Isolated naïve CD4⁺ T cells were allogeneic to the Tregs. Naïve CD4⁺ T cells were labelled with Cell Trace Violet (CTV; Thermo Fisher Scientific, Waltham, MA, USA) as per manufacturer's protocol. 96-well round bottom plates were seeded with CTV-labelled naïve CD4⁺ T cells (5 × 10⁶ cells per well; ‘Tresponder’), and Tregs were added to the wells at various ratios of Treg:Tresponder (1:1, 1:2, 1:4 and 1:8). Human T-expander CD3/CD28 Dynabeads™ (bead:Tresponder ratio of 1:5) was added to each well. CTV-labelled naïve CD4⁺ T cells with and without beads were used as positive and negative controls, respectively. Naïve CD4⁺ T cells without CTV labelling were used as unstimanted control. Cell density control was prepared by mixing positive and negative controls at 1:1 ratio. The plate was incubated at 37 °C in 5% CO₂ for 5 days. IL-2 free cX-vivo medium was used for suppression assay. After a 5-day incubation, cells were stained with FVS780 then analysed by flow cytometry (FACS Canto II). Percentage proliferation was measured by CTV dilutions. Percentage suppression was calculated as follows: 100 × (1 − (percentage proliferation in the experiment sample divided by percentage proliferation in the positive control)).

**Cytokine production assays**

Cells were washed three times then re-suspended in cX-vivo supplemented with cell stimulation cocktail (2 μL mL⁻¹ final concentration; PMA and ionomycin; eBioscience). Cells were incubated at 37 °C in 5% CO₂ for 6 h. After 6-h incubation, cells were spun down and supernatants were collected. Supernatants were stored at −80 °C until assays. Cytokine production was analysed using Human Th1/Th2/Th17 cytokine bead array (CBA) kit (BD Biosciences) and LEGENDplex™ Human Free Active/Total TGF-β1 Assay kit (Biolegend, San Diego, California, USA), as per manufacture’s protocols. Cell-free cX-vivo medium was used as a control to measure background cytokine levels as a result of human serum used to supplement the medium. Samples were analysed by flow cytometry (FACS Canto II), and the data were analysed by FCP Array software (BD Biosciences) and LEGENDplex™ Data Analysis software (Biolegend). Human Th1/Th2/Th17 CBA kit measured IL-2; however, these data were excluded as a result of carry-over from recombinant IL-2 added to the medium.

**Flow cytometric analysis for CD25, FOXP3 and viability**

1 × 10⁶ cells were stained for viability (Fixable Viability Stain 780, FVS780; BD Biosciences) and CD4-FITC (OKT4, BD Biosciences) and CD25-BV421 (M-A251, BD Biosciences). Cells were fixed and permeabilised (Foxp3/Transcription factor staining buffer set, eBioscience) then intracellularly stained for FOXP3-BB700 (236A/E7, BD Biosciences). Compensation controls were utilised and freshly isolated naïve CD4⁺ T cells were used for as a control. All samples were analysed on a BD FACS Canto II flow cytometer (BD Biosciences), and the data analysed with FCS Express 6 (De Novo Software, Pasadena, CA, USA).
Reverse transcription quantitative polymerase chain reaction

$5 \times 10^5$ cells were pelleted, and supernatants removed. Cell pellets were lysed, and RNA was isolated using RNAqueous™ Total RNA Isolation Kit (Ambion, Austin, TX, USA) as per manufacturer’s protocol. Purified RNA was converted to cDNA using iScript Reverse Transcription Supermix (Bio-Rad, Hercules, CA, USA) as per manufacturer’s protocol. cDNA was then used for measurement of gene expression via quantitative polymerase chain reaction on a CFX Connect Real-Time PCR Detection System (Bio-Rad) using TaqMan™ Gene Expression Master Mix (Applied Biosystems, Foster City, CA, USA) and TaqMan primers for RORC (Hs01076112_m1), CTLA4 (Hs00175480_m1), IL2 (Hs00174114_m1) and RPL13A (Hs04194366_g1; Housekeeping gene; Applied Biosystems for all), as per manufacturer’s protocol.

TSDR methylation assay

$1 \times 10^6$ cells were spun down, and all supernatants were discarded, leaving cell pellets. Cell pellets were stored at $-80\, ^\circ \text{C}$ until assay. Assay was conducted by EpigenDx (Hopkinton, MA, USA) using the assay ADS783-FS1 and ADS783-FS2 (Ensembl Transcript ID: ENST00000376207) which assessed the methylation status of 11 CpG sites in the CNS2 by targeted bisulphite pyrosequencing of genomic DNA isolated from the cell pellets. This region covered CpG sites $-2376$ (CpG#44), $-2371$ (CpG#43), $-2330$ (CpG#42), $-2322$ (CpG#41), $-2312$ (CpG#40), $-2309$ (CpG#39), $-2303$ (CpG#38), $-2299$ (CpG#37), $-2291$ (CpG#36), $-2282$ (CpG#35) and $-2263$ (CpG#34) relative to the FOXP3 ATG start codon. Internal low (3.2%), medium (63.3%) and high (91.7%) methylation controls were utilised.

Statistics

Statistical significance ($P < 0.05$) was analysed using the GraphPad Prism 8 (San Diego, CA, USA). RM one-way ANOVA with Dunnett’s multiple comparisons test, RM two-way ANOVA with Dunnett’s multiple comparisons test and the paired two-tailed t-test were used to identify statistical significance. N numbers indicate biological replicates (separate buffy coats). Within each biological replicate, technical triplicates were utilised, except for CBA and Legendplex which used duplicates and TSDR methylation assay which used no technical replicates. For each data point, average value of technical replicates was used.

ACKNOWLEDGMENTS

The authors acknowledge funding from The Hospital Research Foundation, Adelaide Australia, support of Australian Red Cross Blood Services for providing human buffy coat for cell isolation, and Dr Randall Gros of South Australian Health and Medical Research Institute for operating fluorescence-activated cell sorter.

AUTHOR CONTRIBUTIONS

Juewan Kim: Conceptualization; Data curation; Formal analysis; Investigation; Methodology; Project administration; Visualization; Writing—original draft. Christopher M Hope: Conceptualization; Methodology; Supervision; Writing—review & editing. Griffith Perkins: Investigation; Writing—review & editing. Sebastian Stead: Investigation; Visualization; Writing—review & editing. Jacqueline Scaffidi: Investigation. Francis Kette: Investigation. Robert Carroll: Supervision; Writing—review & editing. Simon C Barry: Conceptualization; Supervision; Writing—review & editing. Patrick Toby Coates: Conceptualization; Funding acquisition; Supervision; Writing—review & editing.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

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Supporting Information
Additional supporting information may be found online in the Supporting Information section at the end of the article.