Rapamycin Resistant Murine Th9 Cells Have a Stable In Vivo Phenotype and Inhibit Graft-Versus-Host Reactivity

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Abstract

The cytokine micro-environment can direct murine CD4+ T cells towards various differentiation lineages such as Th1, Th2 and Tregs even in the presence of rapamycin, which results in T cells that mediate increased in vivo effects. Recently, a new lineage of T cells known as Th9 cells that secrete increased IL-9 have been described. However, it is not known whether Th9 differentiation occurs in the presence of rapamycin or whether adoptively transferred donor Th9 cells would augment or restrict alloreactivity after experimental bone marrow transplantation. We found that CD4+ T cells that were co-stimulated and polarized with TGF-β and IL-4 in the presence or absence of rapamycin each yielded effector cells of Th9 phenotype that secreted increased IL-9 and expressed a transcription factor profile characteristic of both Th9 and Th2 cells (high GATA-3/low T-bet). Augmentation of T cell replete allografts with manufactured rapamycin resistant Th9 cells markedly reduced both CD4+ and CD8+ T cell engraftment and strongly inhibited allo-specific T cell secretion of IFN-γ. The potency of Th9 cell inhibition of alloreactivity was similar to that of rapamycin resistant Th2 cells. Importantly, rapamycin resistant Th9 cells persisted and maintained their cytokine phenotype, thereby indicating limited differentiation plasticity of the Th9 subset. As such, Th9 differentiation proceeds in the presence of rapamycin to generate a cell therapy product that maintains high IL-9 expression in vivo while inhibiting IFN-γ driven alloreactivity.

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specific pathogen-free facility at the National Institutes of Health. Experiments were performed in accordance with the guidelines of the NIH Animal Care and Use Committee (ACUC). The protocol was approved by the NCI ACUC (protocol number MB-075).

In vitro Generation of Rapamycin-resistant Th9 Cells

Spleen cells from C57BL/6 mice were harvested and lysed using ACK lysing buffer (Quality Biological, Inc., Gaithersburg, MD). B cells were depleted using goat-anti-mouse conjugated magnetic bioparticles (Polysciences Inc., Warrington, PA). To generate CD3, CD28 mAb coated beads; M450 Dynabeads (Dynal ASA, Oslo, Norway) were incubated with anti-murine CD3 and CD28 mAbs (BD Pharmingen, San Diego, CA) in 0.1M borate solution at 37°C overnight. CD3, CD28 mAb coated beads were then washed with PBS containing 3% FBS (Gemini Bio-Products, Woodland, CA), 5mM EDTA (Quality Biological, Inc.) and 0.1% sodium azide (Sigma, St. Louis, MO), and brought up in the same solution at a concentration of 40 x 10^6 beads/ml. Control rapamycin resistant Th1 cells (Th1.R) were generated using αCD3/αCD28-coated beads in RPMI complete media with 20 IU/mL rhIL-2, 20 ng/mL rhIL-7, 20 ng/ml rmIL-12 (Peprotech), 10 mM of Rapamycin (LC Laboratories, Woburn, MA) and 3.3 mM NAG. Control rapamycin resistant Th2 cells (Th2.R) were cultured exactly the same except for the addition of 1000 IU/ml of rmIL-4 instead of rmIL-12. Culture conditions for the manufacture of Th9 cells were exactly the same as for Th1 cells except for the further addition of 5 ng/ml rhTGF-β1. Specific cytokine-containing complete media was added to all cultures daily from days 2 to 6 to maintain a cell concentration between 0.2–1.0 x 10^6 cells/mL. After 6 days in culture, the various T helper cell subsets were used for bone marrow transplant experiments, multiplex bead array, and flow cytometry analysis.

Cytokine Secretion Assays

Rapamycin-generated Th2 (Th2.R) and Th9 (Th9.R) cells were harvested (day 6) and adjusted to 1 x 10^6 cells/ml in 24-well plates with or without αCD3/αCD28 beads. Twenty-four-hour culture supernatants were collected; cytokine content was evaluated by Bio-Plex multiplex sandwich immunoassay (Bio-Rad).

Bone Marrow Transplantation

Host BALB/c (H-2K^b) mice were lethally irradiated (950cGy; 137Cs γ-radiation source; Gamma Cell 40; Atomic Energy of Canada, Ottawa, Canada) and reconstituted with parental B6 (H-2K^b) bone marrow cells (5 x 10^6 cells) and pan T cells (1 x 10^6 cells) intravenously (i.v.). Some cohorts received in vitro cultured Th2.R (10 x 10^6 cells) or Th9.R (10 x 10^6 cells) cells. In vivo alloreactivity was allowed to occur and immune biology was studied at day 14 post-transplant.

Flow Cytometry Analysis of T Cells

Splenocytes from BALB/c mice were obtained at day 14 post-BMT. Day 14 splenocytes were stained with CD4 PE-cy5 (H129.19), H2K^b PE (AF6.80.5), CD3 FITC (145-2C11), CD45.1 APC (A20) and CD45.2 PE-cy7 (104). For intracellular cytokine measurement, T cells were stimulated for 4h with phorbol 12-myristate 13-acetate and ionomycin (Sigma) followed by the addition of Brefeldin A (GolgiPlug) and Monensin (Golgi stop) for the last 2hrs. Cells were then fixed, permeabilized and stained to detect IFN-γ FITC (XMG1.2), IL-9 PE (RM9A4) and IL-4 APC (11B11). All antibody staining was performed in the presence of αCD16/32 (9,3; eBioscience). Cells were then analyzed using LSRII and FlowJo software. All antibodies were obtained from BD Bioscience unless otherwise stated.

In Vivo Tracking of Donor Th2.R and Th9.R Populations and Unmanipulated CD4 and CD8 T Cells

To allow cell tracking, Th2.R and Th9.R cells were generated using splenic T cells from CD45.1 C57BL/6 congenic mice. Anti-CD45.2 PE,cy7(BD Pharmingen) was used to track unmanipulated CD4 and CD8 populations.

Determination of in vivo Allosensitization

Spleen cells collected post-BMT were subjected to: no stimulation, αCD3/αCD28 stimulation, B6 syngeneic DC stimulation or BALB/c allogeneic DC stimulation. For αCD3/αCD28 stimulation, spleen cells were adjusted to a concentration of 0.5 x 10^6 cells/ml and incubated with CD3/CD28 beads. DCs were obtained by culturing marrow cells for 4 days in rmGM-CSF and rmIL-4 (each at 1000 IU/ml; Peprotech); bacterial LPS (1 µg/ml; Calbiochem) was added to final 24 h of DC culture. Expanded DC were washed and used at a ratio of 10:1 (spleen cell:DC). For cytokine analysis, twenty-four-hour culture supernatants were collected; cytokine content (IL-4, IL-9, IFN-γ and TNF-α) was evaluated by Bio-Plex multiplex sandwich immunoassay (Bio-Rad).

Statistics

Statistical analysis was determined using GraphPad Prism 4 software. P values from data presented as histograms, unless stated otherwise, were determined using a two-tailed unpaired t test. Unless stated otherwise, histogram columns represent the mean values for each experiment and error bars indicate the standard error of mean. Single asterisk (*) denotes P<0.05, double asterisk (**) denotes P<0.01, and triple asterisk (***) denotes P<0.001.

Results

Th9 Polarization Occurs in the Presence of Rapamycin

Using intra-cellular cytokine flow cytometry, we found that bulk CD4^+ T cells were capable of being polarized into Th1, Th2 or Th9 cells in the presence or absence of rapamycin (Representative data; Figure 1a). The frequency of IFN-γ secreting cells was reduced in each subset by rapamycin, with the Th2 and Th9 subsets being lower than the Th1 subset (Figure 1b). Both Th9 and Th9.R cultures had an increased frequency of IL-9 secreting cells relative to the other subsets (Figure 1c). Using transcription factor ILC, flow cytometry, we found that T-bet was highly expressed by Th1 and Th1.R cells (Figure 1d and 1e) whereas both Th2/Th2.R and Th9/Th9.R cells preferentially expressed GATA3 (Figure 1d and 1f).

Murine Th9 Cells Expended in Rapamycin Secrete IL-9 In vitro

Using multiplex bead array system, the cytokine secretion profile of Th2.R and Th9.R was tested. Th2.R and Th9.R secreted low levels of IL-2 and IFN-γ (Figure 2a) and similar magnitudes of IL-4 and IL-9 (Figure 2a). The ratio of IL-4 to IL-9 tended to be higher in the Th2.R subset but this difference did not reach statistical significance (Figure 2b); similarly, the ratio of IL-9 to IL-4 tended to be higher in the Th9.R subset but did not reach statistical significance (Figure 2c).
Murine Th9.R Cells Inhibit IFN-γ Mediated GVHR

Murine recipients were reconstituted with BM and allogeneic T cells following TBI. Certain cohorts received additional Th2.R or Th9.R cells that were expanded from CD45.1 C57BL6 donors. At day 14, splenocytes from cohorts that were treated with Th2.R or Th9.R cells secreted reduced IFN-γ and TNF-α in response to allogeneic DC stimulation (Figure 3a, 3b). Co-transfer of Th9.R cells, but not Th2.R cells, resulted in decreased engraftment of donor CD8+ T cells (Figure 3c). However, both Th9.R and Th2.R cells reduced the frequency of IFN-γ producing donor CD8+ T cells (Figure 3d). By comparison, Th2.R cells but not Th9.R cells increased the number of post-transplant donor CD4+ T cells that were derived from the unmanipulated T cell inoculum (CD45.2+)(Figure 3e). Both Th9.R and Th2.R cells reduced the frequency of IFN-γ producing donor CD45.2+CD4+ T cells (Figure 3f). At day 14 post-BMT, the adoptively transferred Th2.R and Th9.R populations were present in vivo at similar levels (absolute number of Th2.R vs. Th9.R cells, ×10^6 per spleen; Mean ± SEM: 9.3±1.9 vs. 7.1±1.7). Similar to their phenotype at the time of adoptive cell transfer, Th9.R and Th2.R cells harvested at day 14 post- BMT preferentially produced either IL-9 or IL-4, respectively (Figure 3g).

Figure 2. Cytokine Profile of CD4+ Th9.R cells. Splenocytes were harvested from C57B6 mice and bulk CD4+ T cells were isolated. CD4+ T cells were polarized to either Th1 or Th2 or Th9 differentiation for 6 days. At day 6, cells were washed and characterized for cytokine production by Multiplex Luminex assay. (A) Shows Th1/Th2 cytokine secretion profile of Th2 and Th9 cells expanded in the presence of rapamycin. (B) Ratio of IL-4 versus IL-9 cytokine production in Th2.R and Th9.R cells. (C) Ratio of IL-9 versus IL-4 cytokine production in Th2.R and Th9.R cells.
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Figure 3. Rapamycin resistant Th9 cells inhibit alloreactivity in vivo. Host BALB/c mice were subjected to TBI (950cGY) and then reconstituted with allogeneic CD45.2 B6 BM (10M) and T cells (1M). In certain cohorts, mice received either Th2.R or Th9.R cells (10M) that were generated from congeneric CD45.1 mice. At day 14, splenocytes were harvested and restimulated with either syngeneic or allogeneic DC. (A) Alloreactive IFN-γ secretion in the different cohorts. (B) Alloreactive TNF-α secretion. (C) Absolute numbers of donor CD45.2+ CD8+ T cell engraftment in the various cohorts at day 14 post-BMT. (D) Absolute numbers of donor CD45.2+ CD8+ T cells that produce IFN-γ after PMA/ionomycin stimulation was detected by I.C. flow cytometry. (E) Absolute numbers of donor CD45.2+ CD4+ T cell engraftment in the various cohorts at day 14 post-BMT. (F) Absolute numbers of post-BMT donor CD45.2+ CD4+ T cells that produce IFN-γ after PMA/ionomycin stimulation. (G) Ex vivo cytokine production of
Th9.R and Th9.R cells at day 14 post BMT. Each experiment had n = 5 animals per cohort. Single asterisk (*) denotes P<0.05, double asterisk (**) denotes P<0.01, and triple asterisk (***) denotes P<0.001.

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Discussion

In this study, we have initiated an evaluation of the in vitro phenotype and in vivo stability of rapamycin resistant Th9 cells during graft-versus-host-reactivity. The data presented here shows that Th9 cells: (1) can be polarized in the presence of rapamycin, (2) possess cytokine phenotype stability in vivo during allogeneic BMT; and (3) inhibit IFN-γ mediated graft-versus-host-reactivity.

The data presented here are the first to demonstrate that Th9 cell differentiation can proceed in the presence of rapamycin. The current finding extends previous observations on the relatively unbiased nature of CD4+ T cell differentiation in rapamycin [21]. The results presented here further confirm that the effect of rapamycin in blocking CD4+ T cell differentiation [12] can be overcome by using an APC free system along with polyclonal stimulation and adequate cytokine availability. Such expanded Th9.R cells were similar in cytokine phenotype and transcription factor profile to their non-rapa counterparts (Th9 cells). Also, in keeping with previous reports on Th9 cells [8] [2], we found that Th9,R cells were similar to Th2.R cells in terms of transcription factor expression and cytokine profile.

For the first time, we have demonstrated the phenotype stability of Th9,R cells in vivo during allogeneic BMT. Differentiation plasticity of T helper subsets during allogeneic BMT [22] [23] raises concerns with respect to the use of T helper cell therapy. Also, previous reports have suggested that Th9 cells are only transient IL-9 producers [24] and are prone to plasticity [25] in vivo during inflammation. In this context, it is noteworthy that we found that Th9.R cells inhibited type I cytokine production of allo-reactive T cells while maintaining their IL-9 producing capacity. Moreover, lineage stability was not exclusive to Th9,R cells but was also observed with respect to Th2.R cells. Taken together, these results indicate that expansion of T helper subsets in rapamycin may render lineage stability in vivo during inflammation, thereby enhancing the fidelity of defined T helper cell therapy.

This report is also the first to demonstrate an immune modulation function of Th9.R cell function during experimental allogeneic BMT. Specifically, we found that Th9.R cells inhibited IFN-γ driven alloreactivity with a potency similar to Th2,R cells. This finding indicates that it is possible to deliver cells with increased IL-9 secretion capacity without exacerbating alloreactivity. Further studies will be required to determine whether transfer of T cells with increased IL-9 secretion capacity do not increase GVHD or whether such cells might improve graft-versus-tumor-effects.

One might argue as others have previously asserted [2], that because the Th2.R and Th9.R cells had a similar in vitro phenotype and mediated similar in vivo inhibition of alloreactivity, these subsets are essentially the same T cell product and without substantial differential effect. However, contrary to this line of reasoning, we found that the Th2,R and Th9,R products had highly differential effects on donor T cell engraftment. Namely, donor Th9,R cells greatly inhibited donor CD8 engraftment during allogeneic BMT; Th2,R cells, on the other hand, did not inhibit donor CD80 T cell engraftment. It is possible that the potent ability of Th9,R cells to limit CD80 T cell engraftment may offer an improved balance to GVHD and GVT effects [9]. Similarly marked differential effects were also observed in terms of donor CD4+ T cell engraftment; that is, whereas Th2,R cell infusion increased donor CD4+ T cell engraftment, Th9,R cell infusion did not increase donor CD4+ T cell numbers post-BMT. Thus relative to the Th2,R population, Th9,R cells were equally efficient in terms of inhibiting IFN-γ driven alloreactivity and more potent in terms of limiting donor CD4+ and CD80 T cell numbers post-BMT. Further studies are therefore warranted to determine the differential effects of Th2,R and Th9,R cell therapy in the modulation of transplant responses after allogeneic bone marrow transplantation.

Author Contributions

Conceived and designed the experiments: CWM PRM. Analyzed the data: CWM PRM. Wrote the paper: CWM DHF SA.

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