Induction of regulatory T cells by green tea polyphenol EGCG

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\textbf{ABSTRACT}

Regulatory T cells (Treg) are critical in maintaining immune tolerance and suppressing autoimmunity. The transcription factor Foxp3 serves as a master switch that controls the development and function of Treg. Foxp3 expression is epigenetically regulated by DNA methylation, and DNA methyltransferase (DNMT) inhibitors can induce Foxp3 expression in naïve CD4\textsuperscript{+} T cells. We showed that EGCG, a major green tea polyphenol, could act as a dietary DNMT inhibitor, and induced Foxp3 and IL-10 expression in CD4\textsuperscript{+} Jurkat T cells at physiologically relevant concentrations in vitro. We further showed that mice treated with EGCG in vivo had significantly increased Treg frequencies and numbers in spleen and lymph nodes and had inhibited T cell response. Induction of Foxp3 expression correlated with a concomitant reduction in DNMT expression and a decrease in global DNA methylation. Our data suggested that EGCG can induce Foxp3 expression and increase Treg frequency via a novel epigenetic mechanism. While the DNMT inhibitory effects of EGCG was not as potent as pharmacologic agents such as 5-aza-2′-deoxycytidine, the ability of dietary agents to target similar mechanisms offers opportunities for potentially sustained and longer-term exposures with lower toxicity. Our work provides the foundation for future studies to further examine and evaluate dietary strategies to modulate immune function.

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2. Materials and methods

2.1. Cell culture and in vitro treatments

Human Jurkat leukemic CD4+ T cell line was obtained from ATCC (Manassas, VA), and was maintained in RPMI1640 medium supplemented with 10% fetal bovine serum. Jurkat T cells were adjusted to a cell concentration of 1 x 10^6 cells/mL, and were incubated in the presence or absence of Aza (Sigma, St. Louis, MO) (5 μM) or EGCG (Sigma) at 2, 10, or 50 μM for 24 h to 72 h. For EGCG treatments, media were removed each day and cells were replenished with fresh media containing EGCG. For green tea treatments, green tea (2%, w/v) was brewed for 2 min in boiling water with constant stirring, and sterilized by filtration using a 0.22 μm filter. Green tea was purchased from Harney & Sons (Millerton, NY). EGCG content was determined by HPLC [14]. The average EGCG concentration in 2% green tea was 598 ± 31 μg/mL.

2.2. Animal studies and in vivo treatments

Eight-week old Balb/c male mice were purchased from Jackson Laboratory (Bar Harbor, ME). All mice were housed in a temperature- and humidity-controlled environment. Food and water were provided ad libitum. Mice were either left untreated, or injected i.p. daily with 1 mg EGCG per mouse (50 mg/kg) for a total of 7 days. Mice were sacrificed on day 8 by CO2 asphyxiation, and lymphoid organs including thymus, spleen and lymph nodes from individual mice were collected. All procedures involving animals and their care were conducted in accordance with the guidelines as specified in the animal protocol approved by the Oregon State University Institutional Laboratory Animal Care and Use Committee.

2.3. Plasma EGCG

Plasma EGCG was measured by HPLC following enzymatic hydrolysis. Plasma (200 μL) collected from untreated mice or at various time points from mice administered EGCG i.p. was mixed with 20% of 10% ascorbic acid containing 2.5% diethylenetriamine pentaacetic acid (DTPA). The sample was then subjected to enzymatic hydrolysis (45 min, 37°C) using 790 U β-glucuronidase and 38 U sulfatase prepared in 0.4 M sodium phosphate buffer (pH 5.0) containing 2.5 mM DTPA. Samples were rapidly chilled on ice and EGCG was extracted three times using 2 mL of ethyl acetate. Extracts were combined, dried under nitrogen gas, reconstituted in 5.0 L of 10% ascorbic acid containing 2.5 mM diethylenetriamine pentaacetic acid (DTPA). The sample was then subjected to enzymatic hydrolysis. Plasma EGCG was measured by HPLC [14]. The average EGCG concentration in 2% green tea was 598 ± 31 μg/mL.

2.4. Foxp3, IL-10, and DNA methyltransferase expression

Total RNA was isolated using Trizol reagent (Invitrogen, Carlsbad, CA). One microgram of total RNA was reverse transcribed into cDNA using SuperScript III First-Strand Synthesis SuperMix for qRT-PCR (Invitrogen). Real time PCR was performed using gene-specific primers as follows: Foxp3 (forward: 5′-CGGACATCTTGGTAGTACG-3′, reverse: 5′-TTTGCGGTATGTG-3′), IL-10 (forward: 5′-CTGTAGGATTTAAGGTGATCT-3′, reverse: 5′-TTAGATGTCGGTGTGTCGTT-3′), DNMT1 (forward: 5′-GTTGGGACTTGCTTCCTTG-3′, reverse: 5′-TGAAGCTGATGCTGCTTCAC-3′), DNMT3a (forward: 5′-CAGCAAGAAGCCTATCGAGTG-3′, reverse: 5′-AGTGGACTGGAACAAATACCT-3′), DNMT3b (forward: 5′-ACTGGAATCCAGGCAAAGC-3′, reverse: 5′-ACTGGAATCCTACAGGACAT-3′), and GAPDH (forward: 5′-CGAGATCCCCC1AAATCA-3′, reverse: 5′-TTCACCACAGACACAT-3′).

2.5. Genomic DNA isolation and global DNA methylation analyses

Genomic DNA was isolated using DNeasy Blood and Tissue kit (Qiagen, Valencia, CA). Global DNA methylation was measured using SuperSense Methylated DNA Quantification Kit (Epigentek, Brooklyn, NY), and was reported as relative fluorescence units (RFU) per 100 ng genomic DNA.

2.6. Quantification of Treg frequency and numbers

Spleen, lymph nodes, and thymus from individual mouse were made into single cell suspensions. Red blood cells were removed from splenocytes by incubating with red blood cell lysis buffer (150 mM NH4Cl, 1 mM KHCO3, 0.1 mM EDTA, pH 7.2). Total cell number from each organ was determined using the Z1 Coulter Particle Counter (Beckman Coulter). The frequency of Treg was determined by flow cytometry using Mouse Regulatory T Cell Staining kit (eBiosciences, San Diego, CA), where Treg were identified by a combination of cell surface expression of CD4 and CD25, and intracellular expression of Foxp3. Data were acquired using FACSCalibur (BD Biosciences, San Jose, CA). A minimum of 20,000 events in the lymphocyte gate were collected. Data analyses were performed using Summit software (DakoCytomation, Carpinteria, CA). Total Treg was calculated based on Treg frequency and total cell numbers per lymphoid organ.

2.7. Ex vivo T cell functional assays

T cell proliferation and IFNγ production in mice treated with EGCG were assessed ex vivo. Splenocytes from individual mice were seeded at 2 x 10^6 cells per well in 96-well tissue culture plates, and stimulated with 5 μg/mL plate-bound anti-CD3 antibodies (eBiosciences) for 48 h. At the end of incubation, culture supernatants were collected to determine IFNγ production using Mouse IFNγ Ready-Set-Go ELISA kit (eBiosciences). T cell proliferation was determined using standard MTT proliferation assay [16].

Treg suppression assay was performed as described [17] with the following modifications: CD4^+CD25^+ Treg and CD4^+CD25^- T cells were isolated from the spleens of individual untreated or EGCG-treated mice using MACS cell separation columns and mouse CD4^+CD25^+ Regulatory T Cell Isolation Kit (Miltenyi Biotec, Auburn, CA). Graded doses of purified CD4^+CD25^- Treg isolated from untreated or EGCG-treated mice were co-cultured in 96-well plates with 1 x 10^6 cells CD4^+CD25^- responder T cells (isolated from untreated mice only) in the presence of 1 μg/mL plate-bound anti-CD3 (eBioscience) and 1 x 10^5 accessory cells (mitomycin C-treated, T cell depleted splenocytes) for 72 h at 37°C. Responder:Treg ratios ranged from 1:1 to 16:1 per well. T cell proliferation
was determined using standard MTT proliferation assay. Percent suppression was determined by the following equation:

$$1 - \left( \frac{[OD(CD4^{+}CD25^{-} + CD4^{+}CD25^{+})]}{[OD(CD4^{+}CD25^{-} + CD4^{+}CD25^{+})]} \right) \times 100$$

### 2.8. Statistical analysis

Statistical analyses were performed using GraphPad Prism Version 5.02 (GraphPad, La Jolla, CA). All data were reported as mean ± SEM. P values were determined using unpaired t test or one-way ANOVA where appropriate. Statistical significance was defined as $P < 0.05$.

### 3. Results

#### 3.1. EGCG and green tea induced Foxp3 and IL-10 expression in Jurkat T cells

The ability of EGCG to induce Foxp3 expression was tested in Jurkat T cells, a human CD4+ leukemic T cell line. Aza, a potent DNMT inhibitor that has been reported to induce Foxp3 expression in CD4+ T cells was used as a positive control [6]. Both EGCG and Aza induced a significant increase in Foxp3 expression compared to baseline control (Fig. 1). In agreement with previous reports, Aza was a potent inducer of Foxp3 expression, and Jurkat T cells treated with 5 μM Aza had an average of 4.6-fold and 29.3-fold increase in Foxp3 expression at 24 h and 72 h, respectively (Fig. 1B). To determine if EGCG at physiologically relevant concentrations (<10 μM) [18–20] can similarly induce Foxp3 expression, Jurkat T cells were treated with 2 μM and 10 μM EGCG for 72 h and tested for Foxp3 expression (Fig. 2). The expression of IL-10, an immunosuppressive cytokine that is expressed in Treg, was also tested. Significant induction of both Foxp3 (Fig. 2A) and IL-10 (Fig. 2B) expression were detected in Jurkat T cells treated with 2 μM and 10 μM EGCG. Next, to determine if green tea, as a whole food, could similarly induce Foxp3 and IL-10 expression, Jurkat T cells were treated with diluted green tea containing the equivalence of 10 μM and 50 μM EGCG for 72 h (Fig. 3). Both Foxp3 and IL-10 expression were significantly induced (5.7-fold and 7.1-fold, respectively) with green tea at 50 μM EGCG equivalence compared to untreated cells. Cells treated with green tea at 10 μM EGCG did not significantly induce Foxp3 and IL-10 expression.

#### 3.2. EGCG decreased global DNA methylation and DNMT expression in Jurkat T cells

EGCG is a dietary DNMT inhibitor that can reactivate methylation-silenced genes by DNA demethylation [10,11]. We tested whether EGCG treatment would reduce DNA methylation and DNMT expression (DNMT1, DNMT3a and DNMT3b) in Jurkat T cells. Jurkat T cells treated with Aza and EGCG at all three EGCG doses tested had significantly reduced global DNA methylation compared to baseline (Fig. 4A). Administration of 10 μM (Fig. 4B)...
Baseline

10 μM
50μM
0
2
4
6
8
10

Foxp3
IL10

4

Gene expression (Fold-change)

EGCG contents in green tea

Fig. 3. Green tea induced Foxp3 and IL-10 expression in Jurkat T cells in vitro. Jurkat T cells were left untreated (baseline), or treated with diluted green tea containing the equivalence of 10 and 50 μM EGCG for 72 h. At the end of incubation, cells were harvested and analyzed for Foxp3 (open bar) or IL-10 (solid bar) expression by real time PCR. Gene expression was normalized to GAPDH housekeeping gene. Data represent mean normalized fold-change ± SEM. *P<0.05 versus baseline. Results are representative of two independent experiments.

3.3. EGCG increased Treg frequencies and numbers in vivo

To determine the effects of EGCG on Treg numbers in vivo, Balb/c mice were injected i.p. with EGCG daily for 7 days (50 mg/kg). To assess systemic availability of EGCG using this treatment regimen, we measured plasma EGCG concentrations at various time points following i.p. EGCG administration. Plasma EGCG peaked between 5 and 30 min (7.0 ± 0.4 μM and 7.8 ± 0.8 μM, respectively), decreased to 2.0 ± 0.3 μM by 2 h post-injection, and returned to baseline levels below detection limit (<0.1 μM) by 24 h post-injection which was comparable to untreated controls. Plasma EGCG levels in mice that received seven daily EGCG injections had similar EGCG bioavailability compared to mice that received a single injection. Significant increases in both Treg frequencies and numbers were observed in the spleens, pancreatic lymph nodes, and mesenteric lymph nodes of mice treated with EGCG (Fig. 5 and Table 1). For example, in
Table 1
Treg frequencies and absolute numbers in various lymphoid organs in untreated and EGCG-treated mice.

<table>
<thead>
<tr>
<th>Lymphoid organ</th>
<th>Untreated CD4⁺ Treg (%)</th>
<th>EGCG-treated CD4⁺ Treg (%)</th>
<th>Untreated total Treg #</th>
<th>EGCG-treated total Treg #</th>
</tr>
</thead>
<tbody>
<tr>
<td>Spleen</td>
<td>15.5 ± 0.6</td>
<td>22.1 ± 1.7*</td>
<td>2.8 ± 0.2 x 10⁶</td>
<td>5.0 ± 0.6 x 10⁶</td>
</tr>
<tr>
<td>PLN*</td>
<td>12.0 ± 0.3</td>
<td>14.6 ± 0.5</td>
<td>1.3 ± 0.2 x 10⁵</td>
<td>4.7 ± 0.2 x 10⁵</td>
</tr>
<tr>
<td>MLN*</td>
<td>10.1 ± 0.6</td>
<td>12.4 ± 0.1</td>
<td>2.7 ± 0.5 x 10⁵</td>
<td>6.5 ± 1.2 x 10⁵</td>
</tr>
<tr>
<td>ILN*</td>
<td>12.3 ± 0.5</td>
<td>13.9 ± 0.6</td>
<td>2.6 ± 0.3 x 10⁵</td>
<td>2.8 ± 0.4 x 10⁵</td>
</tr>
</tbody>
</table>

*CD4⁺Foxp3⁺ Treg were identified by flow cytometry (n=4 per treatment group).

Table 2
Frequencies and absolute numbers of T and B cells in the spleens of untreated and EGCG-treated mice.

<table>
<thead>
<tr>
<th>Lymphocytes frequency (%)</th>
<th>Absolute number (per spleen)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD4⁺</td>
<td>CD8⁺</td>
</tr>
<tr>
<td>Untreated</td>
<td>30.9 ± 0.4</td>
</tr>
<tr>
<td>EGCG</td>
<td>29.6 ± 1.4</td>
</tr>
</tbody>
</table>

* n = 4 mice per treatment group.

In contrast, the Foxp3 locus in Treg is completely demethylated, allowing for Foxp3 expression. In addition to DNA methylation, histone acetylation/methylation, and Foxp3 acetylation status also contribute to the stability of Foxp3 expression [21,22]. In humans, Treg is a rare cell population and CD4⁺CD25⁺Foxp3⁺ Treg represents only 1–3% of total T cells. Strategies that can induce and/or expand Treg have potentially significant clinical applications for the treatment of autoimmune diseases and prolonging allograft survival. Naturally occurring Treg arise in the thymus, but Treg can also be induced by peripheral conversion of naive Foxp3⁺CD4⁺CD25⁻ T cells into Foxp3⁺ Treg by tolerogenic antigen treatments in vivo, or in vitro exposure of naive CD4⁺CD25⁻ T cells to TGFβ (adaptive/induced Treg) [6]. However, in vitro TGFβ-induced Treg have been shown to have unstable Foxp3 expression, attributed to incomplete demethylation at the Foxp3 locus. In contrast, treatment of CD4⁺CD25⁻ T cells with DNMT inhibitors such as Aza results in complete demethylation of the Foxp3 locus and induces stable Foxp3 expression, and exerts immune suppressive function. The use of pharmacologic agents to alter epigenetically regulated genes for expansion of Treg has been of considerable clinical interest. However, while demethylating agents such as Aza induces stable Foxp3 expression, its use is associated with strong cellular toxicity. As an alternative, the use of natural products including dietary phytochemicals with known epigenetic modulatory activities can potentially be utilized for epigenetic therapy with relative ease of administration and lower toxicity. Several dietary polyphenols have been reported to affect DNA methylation. For example, the consumption of genistein, an isoflavone from soybean altered the methylation patterns in mice [23,24], and tea polyphenols (catechin, epicatechin, and EGCG) and flavonoids (quercetin, fisetin, and myricetin) had DNMT inhibitory activities in vitro [10,13,25]. Among the various dietary constituents, EGCG, the major polyphenol in green tea, has demonstrated to be one of the more potent DNMT inhibitors, and this inhibitory activity has been associated with the reactivation of methylation-silenced genes in cancer cells. EGCG has been associated with a variety of health promoting activities, including anti-inflammatory and anti-carcinogenic effects [9]. Several studies demonstrated that EGCG and green tea polyphenols have immune suppressive effects on T cell in vitro, and limited in vivo studies showed EGCG had inhibitory effects on autoimmune disease progression [26–29]. However, to date the mechanisms by which EGCG suppresses immune (and autoimmune) responses is not well understood. We hypothesize that one of the mechanisms by which EGCG suppresses T cell function and inhibits autoimmune response...
EGCG inhibit DNMT enzymatic activity [25]. In our in vitro culture system, we showed that EGCG also reduced the gene expression of all three DNMTs (DNMT 1, DNMT3a, and DNMT3b), which correlated with reduced DNA methylation in EGCG-treated Jurkat T cells (Fig. 4). As expected of a dietary DNMT inhibitor, EGCG was not as potent as Aza in inducing Foxp3 expression. Nevertheless, Foxp3 expression was induced with EGCG at physiologically relevant concentrations (Figs. 2 and 3).

Our in vivo animal data suggested that EGCG could increase the frequency and number of Treg in various lymphoid organs (Fig. 5 and Table 1). In one human study, oral ingestion of green tea extract resulted in plasma EGCG concentration of ~4 μM at 90 min [19]. Our EGCG bioavailability data indicated that i.p. EGCG injections resulted in peak plasma EGCG concentration of ~7–8 μM between 5 and 30 min post-injection, and reduced to 2 μM by 2 h post-injection. Thus the in vivo effects observed in the current study were achieved using EGCG concentrations that were within physiologically attainable doses of EGCG. While our data suggest that EGCG could increase Treg in vivo, additional tracking experiments will need to be performed to definitively conclude whether Treg is being induced from naive T cell populations, or whether increased Treg is due to expansion of existing Treg. Furthermore, in addition to DNMT inhibitory activities, EGCG is known to exert multiple effects on numerous additional targets, including the induction of cell cycle arrest and apoptosis, and the inhibition of MAP kinase and growth factor-related signaling [31]. We do not propose that Foxp3 is the only target of EGCG, nor do we suggest that the increase in Treg is solely due to epigenetic mechanisms, as EGCG likely acts via a combination of mechanisms in modulating immune response. Nevertheless the DNMT inhibitory effect of EGCG on Foxp3 expression represents a novel and previously uncharacterized mechanism of action that can potentially contribute to the induction and/or enhancement of Treg frequency and function.

The use of dietary approaches to modulate immune function is a novel and potentially promising strategy that can affect immune regulation and disease outcomes with lower toxicity. In autoimmune diseases, environmental factors, including dietary factors, contribute significantly to diseases susceptibility and outcome [32–34]. Indeed, a recent study indicated that dietary intervention early in life was feasible in decreasing the risk of autoimmune type 1 diabetes [35]. In addition, epidemiological studies indicate that populations in countries which traditionally consume large quantities of green tea, such as China and Japan, have one of the lowest disease incidences of autoimmune type 1 diabetes [36]. The use of dietary strategies to epigenetically promote Treg differentiation and Foxp3 expression represents a novel mechanism that has the potential to modulate immune responses and autoimmunity. Although dietary agents may not be as potent as pharmacologic agents, their ability to target similar mechanisms offers opportunities for sustained and longer-term exposures without associated toxicity. Our work provides the foundation for future studies to further evaluate dietary strategies to modulate immune function.

Fig. 6. EGCG reduced T cell proliferation and IFNγ production ex vivo. Balb/c mice (4 mice per group) were left untreated (open bar), or injected i.p. with 50 mg/kg EGCG daily for 7 days (solid bar). At the end of treatments, spleens from individual mice were harvested and splenocytes were stimulated ex vivo with 5 μg/mL plate-bound anti-CD3 for 48 h. T cell proliferation was determined by MTT assay (A), and IFNγ production in the culture supernatant was determined by IFNγ-specific ELISA (B). Splenocytes cultured in media only (unstimulated control) had minimal T cell proliferation and IFNγ production (data not shown). Data represent mean ± SEM. Results are representative of three independent experiments. *P<0.05 versus untreated control. (C) Suppressive function of Treg isolated from untreated (●) or EGCG-treated (▲) mice were determined using standard Treg suppression assay. 1 × 10⁵ CD4⁺CD25⁺ responder T cells were co-cultured with purified Treg isolated from individual untreated or EGCG-treated mice at various responder:Treg ratios in the presence of 1 μg/mL plate-bound anti-CD3 and 1 × 10⁵ mitomycin C-treated accessory cells. Suppression of cell proliferation was determined by MTT assay after 72 h incubation. Data represent mean ± SEM. Results are representative of two independent experiments.

is by specifically inducing and/or enhancing Treg numbers and function via epigenetic mechanisms. Supporting this hypothesis is a recent report that showed EGCG enhanced the functionality of human Treg in vitro [30]. Dietary polyphenols inhibit DNMT activities by various mechanisms, and EGCG has been shown to directly

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