

# Anti-CTLA-4 Immunotherapy Does Not Deplete FOXP3<sup>+</sup> Regulatory T Cells (Tregs) in Human Cancers



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## Abstract

**Purpose:** CTLA-4 was the first inhibitory immune checkpoint to be identified. Two mAbs, ipilimumab (IgG1) and tremelimumab (IgG2), which block the function of CTLA-4, have demonstrated durable clinical activity in a subset of patients with advanced solid malignancies by augmenting effector T-cell-mediated immune responses. Studies in mice suggest that anti-CTLA-4 mAbs may also selectively deplete intratumoral FOXP3<sup>+</sup> regulatory T cells via an Fc-dependent mechanism. However, it is unclear whether the depletion of FOXP3<sup>+</sup> cells occurs in patients with cancer treated with anti-CTLA-4 therapies.

**Experimental Design:** Quantitative IHC was used to evaluate the densities of intratumoral CD4<sup>+</sup>, CD8<sup>+</sup>, and FOXP3<sup>+</sup> cells in stage-matched melanoma ( $n = 19$ ), prostate cancer ( $n = 17$ ), and bladder cancer ( $n = 9$ ) samples treated with

ipilimumab and in paired melanoma tumors ( $n = 18$ ) treated with tremelimumab. These findings were corroborated with multiparametric mass cytometry analysis of tumor-infiltrating cells from paired fresh melanoma tumors ( $n = 5$ ) treated with ipilimumab.

**Results:** Both ipilimumab and tremelimumab increase infiltration of intratumoral CD4<sup>+</sup> and CD8<sup>+</sup> cells without significantly changing or depleting FOXP3<sup>+</sup> cells within the tumor microenvironment.

**Conclusions:** Anti-CTLA-4 immunotherapy does not deplete FOXP3<sup>+</sup> cells in human tumors, which suggests that their efficacy could be enhanced by modifying the Fc portions of the mAbs to enhance Fc-mediated depletion of intratumoral regulatory T cells.

See related commentary by Quezada and Peggs, p. 1130

## Introduction

CTL-associated protein 4 (CTLA-4) is an inhibitory immune checkpoint receptor that plays a crucial role in modulating T-cell responses (1, 2). In preclinical studies, transient blockade of CTLA-4 resulted in tumor eradication and durable antitumor immunity (3). These findings translated into the clinical development of fully human mAbs against CTLA-4, including ipilimumab and tremelimumab; an approach termed immune checkpoint blockade that has proven to be a paradigm shift in cancer therapy (4).

Initial preclinical studies primarily implicated effector T (Teff) cells in mediating the therapeutic activity of CTLA-4 blockade (1, 3); however, later studies demonstrated the contribution of both the effector and regulatory T-cell compartments in promoting

antitumor responses (5–10). Regulatory T cells (Tregs) maintain immune cell homeostasis and suppress antitumor immune responses. They express CD4, CD25 and forkhead box protein P3 (FOXP3), a transcription factor required for their function. In addition, Tregs constitutively express high levels of CTLA-4 on their surface (11–13), and FOXP3 is known to direct CTLA-4 transcription (14, 15), which led to the hypotheses that CTLA-4 blockade may impair the suppressive activity of Tregs or directly deplete Tregs via antibody-dependent cell-mediated cytotoxicity (ADCC; refs. 16, 17).

More recent preclinical murine studies demonstrated that the therapeutic activity of anti-CTLA-4 mAbs (clones 9D9, 9H10, and 4F10) was dependent upon selective depletion of intratumoral Tregs via ADCC. In these studies, tumor regression was associated with an increase in the intratumoral Teff to Treg cell ratio, due to the preferential expansion of Teff cells and the selective depletion of Tregs (5, 7). Treg depletion was attributed to tumor-infiltrating macrophages expressing the Fc receptor, FcγRIV (5). A murine anti-CTLA-4 mAb containing an IgG2a (clone 9D9) constant region was shown to exhibit enhanced ADCC-dependent intratumoral Treg depletion when compared with IgG2b and IgG1 murine anti-CTLA-4 mAbs (7).

Ipilimumab, the first fully human IgG1 mAb against CTLA-4, demonstrated durable clinical activity and prolonged survival in patients with advanced melanoma (18, 19). It has also been tested in other cancer types including non-small cell lung cancer, renal cell carcinoma, urothelial carcinoma, and prostate cancer (4). Similarly tremelimumab, an anti-CTLA-4 IgG2 mAb, has also been clinically studied across several tumor types (20). The humanized IgG1 mAb isotype binds to most human Fc receptors,

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### Translational Relevance

In preclinical murine models, anti-CTLA-4 mAbs promote cancer regression by increasing the frequency of effector T cells within the tumor microenvironment and selectively depleting intratumoral regulatory T cells (Treg) via an Fc-dependent mechanism. However, it is unclear whether this occurs in patients with cancer treated with anti-CTLA-4 therapies. Herein, we show that treatment with IgG1 ipilimumab and IgG2 tremelimumab mAbs, which have distinct FcγR-mediated mechanisms, increases the density of intratumoral CD4 and CD8 T cells, and does not deplete FOXP3<sup>+</sup> Tregs across different human tumors. These data suggest that there is an opportunity to improve the clinical efficacy of anti-CTLA-4 immunotherapies by modifying the mAbs to promote the depletion of intratumoral Tregs.

and it has the ability to induce ADCC; whereas, the IgG2 isotype is known to have less ADCC activity (21). Clinical studies have been unclear in defining the role of IgG1 ipilimumab and IgG2 tremelimumab in depleting intratumoral Tregs in human cancers (9, 22).

The current study herein was conducted to gain insight on the effect of anti-CTLA-4 immunotherapy on the density of intratumoral FOXP3<sup>+</sup> Tregs across several human cancer types, including melanoma, bladder cancer, and prostate cancer. Our data provide evidence that CTLA-4 blockade does not decrease the density of FOXP3<sup>+</sup> Tregs in human tumors.

## Materials and Methods

### Patient material

Formalin-fixed, paraffin-embedded (FFPE) tumor samples were collected from a cohort of stage-matched untreated and ipilimumab-treated primary prostate cancer patients ( $n = 17$ ), primary bladder cancer patients ( $n = 9$ ), and metastatic melanoma patients ( $n = 19$ ). The median time of tumor biopsy collection following last treatment with ipilimumab was 8 weeks for prostate and bladder tumors (post-dose 2 treatments) and 18 weeks for melanoma (post-dose 2–3 treatments). For the mass cytometry or CyTOF assay, tumor-infiltrating cells were obtained from paired fresh melanoma tumors pre- and post-ipilimumab treatment ( $n = 5$ ). The tumor samples were retrieved from The University of Texas MD Anderson Cancer Center Department of Pathology and Institutional Tumor Bank. All patients provided informed consent on The University of Texas MD Anderson Cancer Center institutional-approved laboratory protocol, PA13-0291. FFPE tumor samples from a cohort of paired metastatic melanoma patients ( $n = 18$ ) treated with tremelimumab were obtained from the Department of Hematology and Oncology, University of California, Los Angeles institutional-approved protocol, 11-003254-AM-00019. The median time of posttreatment biopsy collection was 5 weeks following treatment with tremelimumab (post-dose 1 treatment). The institutional review boards of The University of Texas MD Anderson Cancer Center and University of California, Los Angeles specifically approved these studies. All patients signed the informed consent forms and were given a unique code, thus protecting their identity.

### IHC

IHC was performed on FFPE tumor tissue sections. The tumor tissues were fixed in 10% formalin, embedded in paraffin, and transversely sectioned. Four-micron sections were used for the IHC analyses. The sections were stained with mouse anti-human mAbs against CD4 (Novocastra, CD4-368-L-A), CD8 (Thermo Scientific, MS-457-S), FOXP3 (Abcam, ab96048-206D), and CD68 (Dako, M0876). All sections were counterstained with hematoxylin, dehydrated, mounted, and processed with peroxidase-conjugated avidin/biotin and 3'-3'-diaminobenzidine (DAB) substrate (Leica Microsystems). The IHC slides were scanned and digitalized using the scanscope system from Scanscope XT, Aperio/Leica Technologies. The IHC analysis was solely performed in the center of the tumor (CT) region across all tumor types.

Quantitative analysis of IHC staining was conducted using the image analysis software ImageScope-Aperio/Leica. Five random areas (1 mm<sup>2</sup> each) were selected using a customized algorithm for each marker to determine the number of positive cells per high power field (20× magnification), and the average of the five fields was taken. The pathologists in our team carefully reviewed the data generated by the automated image analysis Aperio software, and the data were expressed as a density score (total number of positive cells per 1 mm<sup>2</sup> area). IHC staining was interpreted in conjunction with H&E-stained sections, which facilitated the identification of malignant cells and thus directed IHC quantification.

### CyTOF

Tumor-infiltrating cells from paired fresh melanoma tumors pre- and post-ipilimumab treatment ( $n = 5$ ) were used for the CyTOF analysis. The tumor sample was dissociated with GentleMACS system (Miltenyi Biotec) as per the manufacturer's instructions and cultured overnight in a 96-well plate with RPMI1640 medium; supplemented with 10% human AB Serum, 10 mmol/L HEPES, 50 μmol/L β-mercaptoethanol, penicillin/streptomycin/l-glutamine, and 50 U/ml human IL2. The cells were then collected and analyzed for seven analytes including CD45 (Fluidigm, HI30), CD3 (Fluidigm, UCTH1), CD4 (Fluidigm, RPA-T4), CD8 (Fluidigm, RPA-T8), CD56 (Fluidigm, NCAM16.2), FOXP3 (Fluidigm, PCH101), and CTLA-4 (Fluidigm, 14D3). The data was analyzed by gating on CD45<sup>+</sup> live singlets using ViSNE (23) and FlowJo.

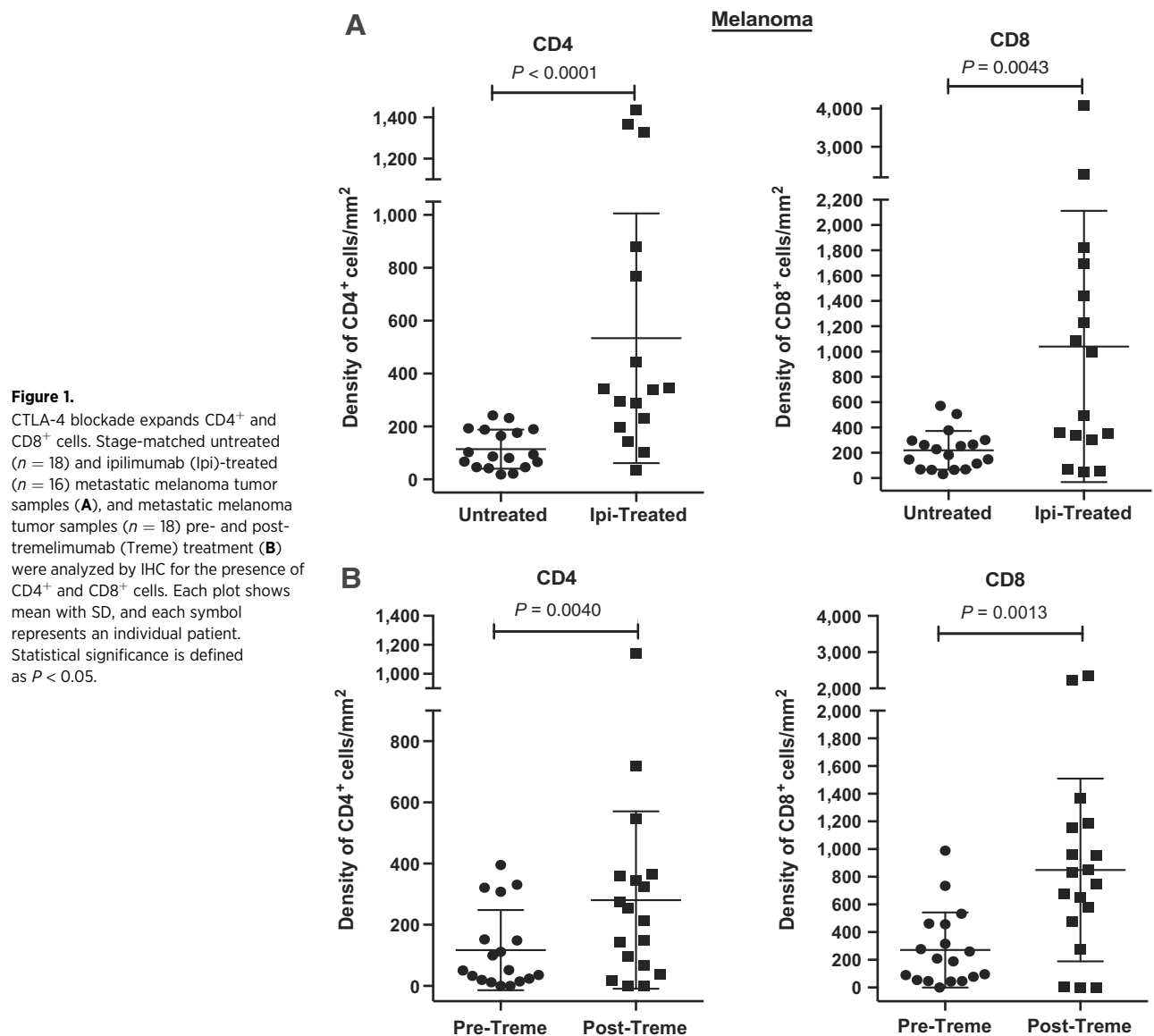
### Statistical analysis

The statistical analysis for each immune subset between the untreated versus treated cohorts was performed using Mann-Whitney unpaired *t* test for the unpaired tumor sections and Wilcoxon paired *t* test for the paired tumor sections (*P* value; Graphpad Prism). Statistical significance was defined as *P* < 0.05.

## Results

### Anti-CTLA-4 immunotherapy increases the density of intratumoral CD4<sup>+</sup> and CD8<sup>+</sup> cells

Preclinical and early clinical studies suggest that CTLA-4 blockade promotes antitumor activity by increasing the frequency of intratumoral Teff cells (5, 7, 8, 10, 24, 25). To confirm these findings using tissues obtained from patients treated with two different anti-CTLA-4 mAbs, we examined the immune profile of melanoma tumors by IHC. Our data demonstrate that both



ipilimumab (Fig. 1A; Supplementary Fig. S1A) and tremelimumab (Fig. 1B) significantly increased the intratumoral densities of both CD4<sup>+</sup> and CD8<sup>+</sup> cells. Similar results have been reported in human melanoma upon treatment with tremelimumab (26). Consistent with these findings, we observed increased densities of intratumoral CD4<sup>+</sup> and CD8<sup>+</sup> cells in bladder and prostate tumors obtained from patients treated with ipilimumab (Supplementary Fig. S2A and S2B).

#### Anti-CTLA-4 immunotherapy does not deplete intratumoral FOXP3<sup>+</sup> cells in human cancers

As it is postulated that human IgG1 mAbs promote ADCC to a greater extent than IgG2 (21), we evaluated the potential differential effects of ipilimumab (IgG1) and tremelimumab (IgG2) on the densities of FOXP3<sup>+</sup> cells in tumor tissues from patients with advanced melanoma. Surprisingly, both ipilimumab (Fig. 2A; Supplementary Fig. S3) and tremelimumab (Fig. 2B) did not deplete, but instead significantly increased the density of FOXP3<sup>+</sup>

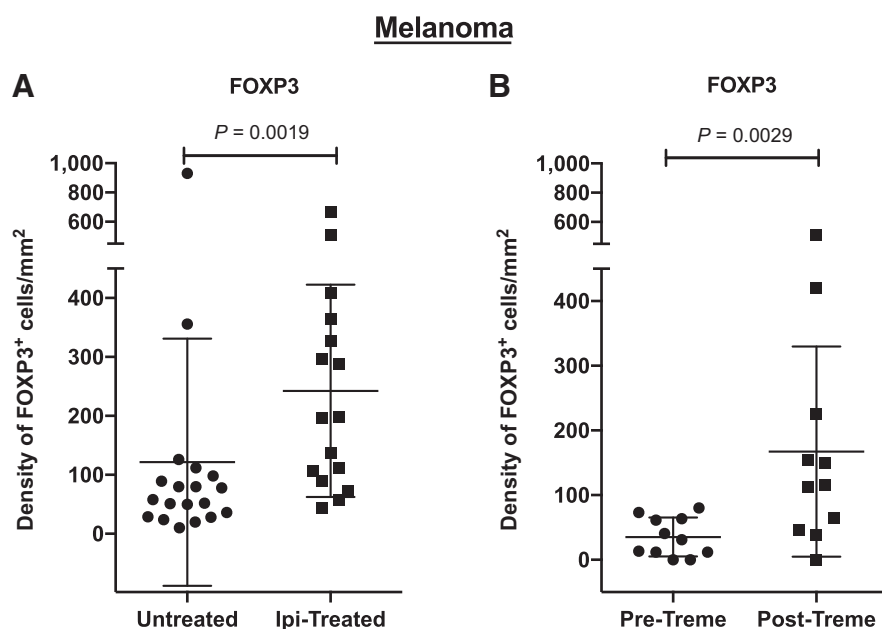
cells within the tumor microenvironment. Of note, pretreatment biopsies from two patients with melanoma, which were negative for FOXP3<sup>+</sup> cells, showed FOXP3<sup>+</sup> cellular infiltration post-tremelimumab treatment (Fig. 2B).

Furthermore, in bladder and prostate tumors obtained from patients treated with ipilimumab, there was no reduction in intratumoral FOXP3<sup>+</sup> cells (Fig. 3; Supplementary Fig. S3). Taken together, the data demonstrate that both ipilimumab and tremelimumab, despite having two different IgG isotypes with potentially different mechanisms of action, similarly augment the densities of intratumoral CD4<sup>+</sup> and CD8<sup>+</sup> cells, but do not deplete FOXP3<sup>+</sup> cells.

#### Presence of tumor-associated macrophages in baseline and ipilimumab-treated biopsies

In humans, ADCC is primarily mediated by FcγRIIIA (CD16a; ref. 27), which is expressed on macrophages and natural killer (NK) cells. The lack of intratumoral Treg depletion following treatment

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**Figure 2.** Effect of CTLA-4 blockade on FOXP3<sup>+</sup> cells in human melanoma. Stage-matched untreated ( $n = 19$ ) and ipilimumab (Ipi)-treated ( $n = 16$ ) metastatic melanoma tumor samples (**A**), and metastatic melanoma tumor samples pre- and post-tremelimumab (Treme) treatment ( $n = 11$ ; **B**) were analyzed by IHC for the presence of FOXP3<sup>+</sup> cells. Each plot shows mean with SD, and each symbol represents an individual patient. Statistical significance is defined as  $P < 0.05$ .

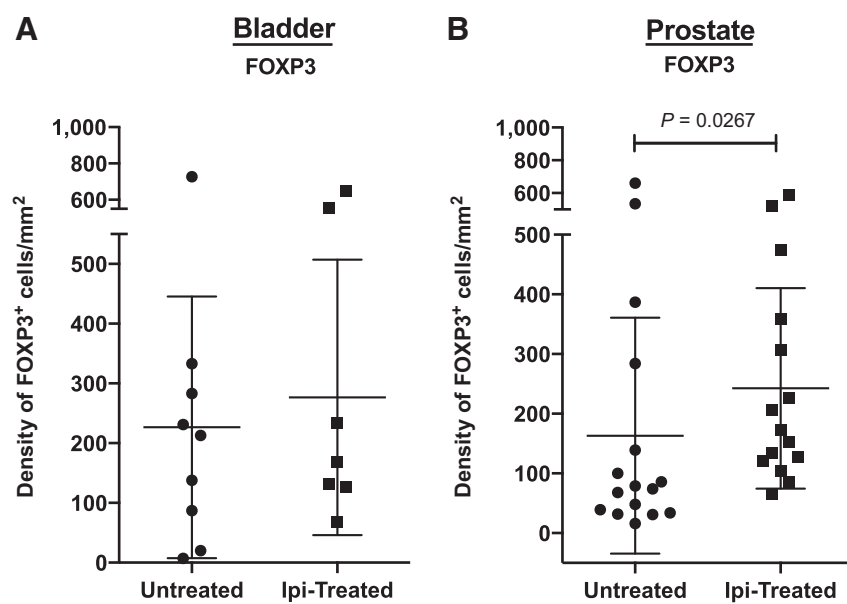
with ipilimumab could be attributable to a paucity of macrophages or NK cells and/or FcγRIIIA expression within the tumor microenvironment. Therefore, we sought to evaluate the effect of ipilimumab on intratumoral CD68<sup>+</sup> macrophages in patients with advanced melanoma. Compared with intratumoral CD4<sup>+</sup> and CD8<sup>+</sup> cell densities, there was considerable infiltration of CD68<sup>+</sup> macrophage in the untreated tumor samples, which did not significantly change following treatment with ipilimumab (Fig. 4).

## Discussion

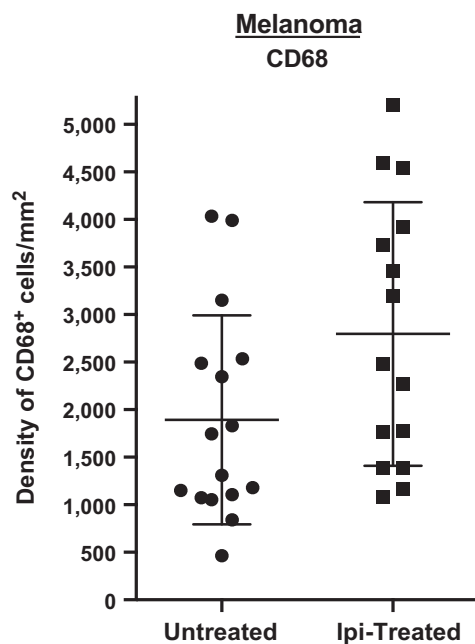
Anti-CTLA-4 immunotherapy has demonstrated significant antitumor activity in preclinical and clinical studies (3, 6, 18, 19, 28), and it has emerged as a paradigm shift in cancer therapy. Although the effect of CTLA-4 blockade on T<sub>H</sub>17 cells has been well

established, its effect on human FOXP3<sup>+</sup> Tregs within the tumor microenvironment remains unclear. While some studies report a decrease in the frequency of circulating or intratumoral Tregs (5, 7, 9, 29, 30), other studies did not result in Treg depletion following anti-CTLA-4 treatment (11, 25, 26, 31, 32). This study has been undertaken to gain a better understanding of the effect of anti-CTLA-4 therapy on intratumoral FOXP3<sup>+</sup> Tregs in human cancers.

Preclinical studies have reported that CTLA-4 blockade results in an increase in the T<sub>H</sub>17 to Treg ratio (5–7). Multiple studies have been conducted to evaluate Tregs in human tumors following anti-CTLA-4 therapy (24–26, 28, 32). However, most of these studies relied on the evaluation of intratumoral Tregs by flow cytometry with the calculation of FOXP3<sup>+</sup> Tregs as a percentage of total cells. In previous studies, FOXP3<sup>+</sup> Tregs were evaluated in bladder tumor tissues pre- and post-ipilimumab treatment. We



**Figure 3.** Effect of CTLA-4 blockade on FOXP3<sup>+</sup> cells in bladder and prostate tumors. Stage-matched untreated and ipilimumab (Ipi)-treated bladder tumors ( $n = 9$ ; **A**), and prostate tumors ( $n = 16$ ; **B**) were analyzed by IHC for the presence of FOXP3<sup>+</sup> cells. Each plot shows mean with SD, and each symbol represents an individual patient. Statistical significance is defined as  $P < 0.05$ .



**Figure 4.**

Effect of CTLA-4 blockade on CD68<sup>+</sup> cells. Stage-matched untreated and ipilimumab (Ipi)-treated ( $n = 16$ ) metastatic melanoma samples were analyzed by IHC for the presence of CD68<sup>+</sup> cells. Each plot shows mean with SD, and each symbol represents an individual patient. Statistical significance is defined as  $P < 0.05$ .

observed an increase in the number of total T cells, but a decrease in frequency of FOXP3<sup>+</sup> Tregs, which was measured as a percent of total cells (24, 25). Because the total number of T cells increased in posttreatment tumor tissues, our calculation of FOXP3<sup>+</sup> Tregs as a percent of the total T cells led to data indicating a decrease in frequency of FOXP3<sup>+</sup> Tregs, even though the absolute number of FOXP3<sup>+</sup> cells may not have changed.

In this study, we evaluated immune subsets including CD4<sup>+</sup>, CD8<sup>+</sup>, and FOXP3<sup>+</sup> cells in tumor tissues obtained from patients treated with anti-CTLA-4 mAbs by quantitative IHC. We measured the density of T-cell subsets in stage-matched human tumor tissues and found that anti-CTLA-4 therapy increases the density of intratumoral CD4<sup>+</sup> and CD8<sup>+</sup> cells, but without significant change or depletion of FOXP3<sup>+</sup> cells. In addition, evaluation of paired pretreatment and post-ipilimumab treatment melanoma tissues revealed higher densities of intratumoral CD8<sup>+</sup> cells (Supplementary Fig. S1B) and FOXP3<sup>+</sup> cells (Supplementary Fig. S1C).

Importantly, the timing of biopsy collection post-anti-CTLA-4 therapy (ipilimumab and tremelimumab) did not impact the infiltration of immune subsets (Supplementary Fig. S4). Consistent with this, an IHC study evaluating melanoma tumor samples ( $n = 19$ ) demonstrated higher infiltrates of FOXP3<sup>+</sup> cells in the responding lesions following treatment with tremelimumab (26). Taken together, these results indicate that IHC may be a more reliable tool to examine treatment-related changes in immune subsets within the tumor, as it provides a more direct measure of the absolute numbers of cells in a given surface area.

The limitation of single-stain IHC is that an individual marker may be expressed on more than one cellular population. For example, single staining for CD8 can represent both T cells and NK

cells; likewise, staining for FOXP3 alone can represent T<sub>H</sub>1 and Treg cells. To address this, we collaborated our IHC results with multiparametric CyTOF analysis of tumor-infiltrating cells from paired pre- and post-ipilimumab-treated melanoma tissues. We found that greater than 95% of the CD8<sup>+</sup> population were T cells (CD3<sup>+</sup>CD8<sup>+</sup>); whereas less than 5% of these were NK cells (CD3<sup>+</sup>CD8<sup>+</sup>CD56<sup>+</sup>). More importantly, ipilimumab significantly increased the frequency of CD8<sup>+</sup> T cells but not CD8<sup>+</sup>CD56<sup>+</sup> NK cells (Supplementary Fig. S5). Similarly, to differentiate between T<sub>H</sub>1 and Treg cells, we evaluated coexpression of FOXP3, CD3, CD4, and CTLA-4. The data demonstrated ipilimumab increases the frequency of CD3<sup>+</sup>CD4<sup>+</sup>FOXP3<sup>+</sup>CTLA-4<sup>+</sup> T cells (Tregs), but not CD3<sup>+</sup>CD4<sup>+</sup>FOXP3<sup>-</sup>CTLA-4<sup>+</sup>, and CD3<sup>+</sup>CD4<sup>+</sup>FOXP3<sup>-</sup>CTLA-4<sup>-</sup> T cells (Supplementary Fig. S6).

It has been reported that antibody-mediated depletion can occur via complement-mediated lysis (33) or ADCC (5). In mouse models, the anti-CTLA-4 therapy-mediated depletion of FOXP3<sup>+</sup> Treg cells was dependent on the presence of intratumoral FcγRIV-expressing macrophages (5). In patients, ADCC is essentially mediated through an equivalent FcγR, namely FcγRIIIA (CD16) that is expressed on macrophages and NK cells (27), which may account for the discrepancy between the preclinical and clinical data. Our findings did not demonstrate significant changes in the intratumoral density of CD68<sup>+</sup> macrophages after anti-CTLA-4 therapy; however, we did not evaluate FcγRIIIA expression in these human tumor samples.

In conclusion, our data illustrate that the IgG1 ipilimumab and IgG2 tremelimumab mAbs with distinct FcγR-mediated mechanisms do not deplete intratumoral FOXP3<sup>+</sup> cells in human cancers.

#### Disclosure of Potential Conflicts of Interest

J. Wargo reports receiving speakers bureau honoraria from Dava, and is a consultant/advisory board member for Bristol-Myers Squibb, AstraZeneca, Novartis, Illumina, and Merck. J.P. Allison is an inventor and recipient of royalty from intellectual property related to activation of ICOS signaling for cancer immunotherapy issued to The University of Texas System and Memorial Sloan Kettering Cancer Center (MSKCC) and licensed to Jounce Therapeutics; is an inventor and recipient of royalty from intellectual property related to the use of Newcastle Disease Virus as a biotherapeutic issued to Icahn School of Medicine at Mount Sinai and MSKCC and licensed to Merck; and is an inventor and recipient of royalty on a now expired patent for use of antagonism of the CTLA-4 signaling pathway for cancer immunotherapy issued to the University of California and licensed to Bristol-Myers Squibb. P. Sharma is listed as a co-inventor of a patent regarding the targeting of ICOS for immunotherapy that is owned by Jounce Therapeutics, and is a consultant/advisory member for Jounce, Neon Therapeutics, Constellation, Oncolytics Biotech, BioAtla, Forty-Seven, Apricity, Polaris, Marker Therapeutics, Codiak, Pieris Pharmaceuticals, Kite Pharma, Merck, and BioMx. No potential conflicts of interest were disclosed by the other authors.

#### Authors' Contributions

**Conception and design:** J.P. Allison, A. Ribas, P. Sharma

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**Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.):** J. Scutti, L. Vence, A. Ribas

**Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis):** A. Sharma, S.K. Subudhi, J. Blando, J. Scutti, L. Vence, J. Wargo, J.P. Allison, A. Ribas, P. Sharma

**Writing, review, and/or revision of the manuscript:** A. Sharma, S.K. Subudhi, J. Blando, J. Wargo, J.P. Allison, A. Ribas, P. Sharma

**Administrative, technical, or material support (i.e., reporting or organizing data, constructing databases):** A. Sharma

**Study supervision:** P. Sharma

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