Elevated expression of the chemokine receptor CCR4 in tumors is associated with poor prognosis in several cancers. Here, we have determined that CCR4 was highly expressed in human renal cell carcinoma (RCC) biopsies and observed abnormal levels of CCR4 ligands in RCC patient plasma. An antagonistic anti-CCR4 antibody had antitumor activity in the RENCA mouse model of RCC. CCR4 inhibition did not reduce the proportion of infiltrating leukocytes in the tumor microenvironment but altered the phenotype of myeloid cells, increased NK cell and Th1 cytokine levels, and reduced immature myeloid cell infiltrate and blood chemokine levels. In spite of prominent changes in the myeloid compartment, the anti-CCR4 antibody did not affect RENCA tumors in T cell–deficient mice, and treatment with an anti–class II MHC antibody abrogated its antitumor activity. We concluded that the effects of the anti-CCR4 antibody required the adaptive immune system and CD4+ T cells. Moreover, CCL17-induced IFN-γ production was reduced when Th1-polarized normal CD4+ T cells were exposed to the CCR4 ligand, evidencing the involvement of CCR4 in Th1/Th2 regulation. The anti-CCR4 antibody, alone or in combination with other immune modulators, is a potential treatment approach to human solid cancers with high levels of CCR4-expressing tumor-infiltrating leukocytes and abnormal plasma CCR4 ligand levels.
A CCR4 antagonist reverses the tumor-promoting microenvironment of renal cancer

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Elevated expression of the chemokine receptor CCR4 in tumors is associated with poor prognosis in several cancers. Here, we have determined that CCR4 was highly expressed in human renal cell carcinoma (RCC) biopsies and observed abnormal levels of CCR4 ligands in RCC patient plasma. An antagonistic anti-CCR4 antibody had antitumor activity in the RENCA mouse model of RCC. CCR4 inhibition did not reduce the proportion of infiltrating leukocytes in the tumor microenvironment but altered the phenotype of myeloid cells, increased NK cell and Th1 cytokine levels, and reduced immature myeloid cell infiltrate and blood chemokine levels. In spite of prominent changes in the myeloid compartment, the anti-CCR4 antibody did not affect RENCA tumors in T cell–deficient mice, and treatment with an anti–class II MHC antibody abrogated its antitumor activity. We concluded that the effects of the anti-CCR4 antibody required the adaptive immune system and CD4+ T cells. Moreover, CCL17-induced IFN-γ production was reduced when Th1-polarized normal CD4+ T cells were exposed to the CCR4 ligand, evidencing the involvement of CCR4 in Th1/Th2 regulation. The anti-CCR4 antibody, alone or in combination with other immune modulators, is a potential treatment approach to human solid cancers with high levels of CCR4-expressing tumor-infiltrating leukocytes and abnormal plasma CCR4 ligand levels.

Introduction

Tumor microenvironments possess complex chemokine networks that contribute to the extent and phenotype of the host infiltrate (1–3). In addition, malignant cells may gain functional chemokine receptors, often as a consequence of oncogenic mutations, allowing them to respond to distant chemokine gradients during metastatic spread (4, 5).

The chemokine receptor CCR4 is expressed on circulating and tissue-resident T cells, being predominantly associated with a Th2 phenotype (6–8), as well as on other T helper cells (9). CCR4 is also highly expressed on circulating Tregs and on Tregs recruited at tumor sites in ovarian cancer (10) and in glioblastoma (11). In ovarian cancer, the CCR4 ligand CCL22 is found both in the tumor tissue and in macrophages isolated from ascitic fluid (9). In hepatocellular carcinoma, malignant cell–produced CCL22 recruited CCR4+ Tregs that facilitated immune escape of malignant cells (12). Similarly, in breast cancer, CCR4+ Tregs, recruited by CCL22 in the tumor microenvironment, are predictive of a worse prognosis (13). A second breast cancer study found reduced overall survival and high CCR4 expression in tumor biopsies (14). Finally, in a cohort of 753 patients with gastric adenocarcinoma, positive staining for CCR4 was also associated with a poorer prognosis (15).

CCR4 also plays a role in hematological malignancies, and there are now clinical trials of an anti-CCR4 antibody, mogamulizumab, that has enhanced antibody-dependent cell-mediated cytotoxicity (ADCC) activity. Mogamulizumab is approved in Japan for the treatment of relapsed adult T cell leukemia (ATL) (16) and has also been tested in patients with relapsed peripheral T cell lymphoma (PTLC) and cutaneous T cell lymphoma (CTLC) (17). The treatment is indicated for patients with CCR4-positive leukemia cells, but might also act by reducing the number of Tregs in cancer patients (18).

In this article, we have investigated CCR4 as a target in renal cell carcinoma (RCC) using patient samples and an orthotopic mouse RCC model. We have found abnormal levels of CCR4 and its ligands in human RCC biopsies and plasma samples. In preclinical experiments we found that Affi-S, a fully human anti-CCR4 antibody with antagonistic activity (described in ref. 19), has antitumor activity in a renal cancer model. Inhibition of CCR4 did not reduce the proportion of CCR4-positive infiltrating leukocytes in the tumor microenvironment but altered the phenotype of the immune infiltrate, affecting in particular the phenotype of myeloid cells and increasing the number of infiltrating NK cells. These effects were dependent on the adaptive immune system and required functioning CD4+ T cells. The antibody also altered the phenotype of tumor-associated macrophages (TAMs) in the B16 melanoma model. Inhibition of CCR4, alone or in combina-
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**B–D), suggesting that CCR4 may be important in the trafficking of tumor-associated leukocytes.**

**Results**

*CCR4 and its ligands in human renal cell carcinoma.* This study was prompted by the finding of abundant *CCR4* mRNA in biopsies from renal cancers as compared with normal kidney (Figure 1A). *CCR4* protein was also detected by IHC on malignant cells and leukocytes in a tissue microarray (TMA) constructed from 57 advanced RCC patient biopsies (Figure 1B and Supplemental Figure 1A; supplemental material available online with this article; https://doi.org/10.1172/JCI82976DS1). Of the 173 cores in the TMA, 157 showed positive *CCR4* staining. 75% of the biopsies were classified as clear cell, with others classified as papillary RCC. There was a significant positive correlation between *CCR4* positivity and the extensive T cell (CD3+) or macrophage (CD68+) infiltrates in the tumor cores (Supplemental Figure 1, B–D), suggesting that *CCR4* may be important in the trafficking of tumor-associated leukocytes.

The *CCR4* ligands CCL22 (also known as MDC) and CCL17 (also known as TARC) were also expressed in the RCC tumors (Figure 1B and Supplemental Figure 1A). *CCR4* was weakly expressed in normal kidney, but the ligands could be detected in normal kidney tubules (Supplemental Figure 1A).

We next compared plasma concentrations of CCL17 and CCL22 from patients with advanced RCC with age-matched controls. The *CCR4* and its ligands CCL17 and CCL22 were analyzed by IHC in a TMA of renal cancer biopsies from human patients. Each biopsy was scored 0, no staining; 1, weak staining; 2, strong staining for *CCR4*, CCL17, and CCL22. A total of 173 biopsy cores from 57 patients were stained for *CCR4* and CCL22, and 145 cores from 48 patients for CCL17. (C–E) Plasma levels of CCL17 and CCL22 and the CCL17/CCL22 ratio in RCC patient plasma were compared with those from normal individuals of matched age using Meso Scale Discovery System Ultra-Sensitive plates. *n* = 47 for RCC patients, *n* = 26 for normal individuals; 2-tailed Student’s t test, ***P = 0.0001 for CCL17 (C), CCL22 (D), and CCL17/CCL22 (E). (F and G) Kaplan-Meier survival curves for progression-free survival (PFS; F) and overall survival (OS; G) for RCC patients with CCL17/CCL22 high (above the median) or low (*n* = 57). For progression-free survival, hazard ratio 0.436, 95% CI 0.239–0.797; for overall survival, hazard ratio 0.552, 95% CI 0.306–0.995.
determined by flow cytometry. The human cell lines had detectable intracellular CCL17 and CCL22 (Supplemental Figure 2), and these chemokines were also present in the tissue culture medium during 3 days of incubation (CCL17 300 pg/10⁶ cells, CCL22 2 ng/10⁶ cells). Renca cells also secreted CCL17 (200 pg/10⁶ cells) and CCL22 (10 pg/10⁶ cells) in the medium during 3 days of incubation.

Both CCR4 ligands stimulated migration of the human cell line 786-O (Supplemental Figure 3, A and B). Similar data were examined (Supplemental Figure 1, E and F), suggesting that activity of both chemokines is important in RCC biology. In our cohort, CCR4 expression, as determined by IHC on the TMAs, was not predictive of clinical outcome (data not shown).

Renal cancer cell lines have functional CCR4 receptors. As we had detected CCR4 and its ligands in malignant cells in tumor biopsies, we next studied RCC cancer cell lines. RCC cell lines 786-O and A498 (human) and RENCA (murine) expressed cell surface CCR4 as determined by flow cytometry. The human cell lines had detectable intracellular CCL17 and CCL22 (Supplemental Figure 2), and these chemokines were also present in the tissue culture medium during 3 days of incubation (CCL17 300 pg/10⁶ cells, CCL22 2 ng/10⁶ cells). RENCA cells also secreted CCL17 (200 pg/10⁶ cells) and CCL22 (10 pg/10⁶ cells) in the medium during 3 days of incubation.

Both CCR4 ligands stimulated migration of the human cell line 786-O (Supplemental Figure 3, A and B).
Affi-5 did not influence the growth or viability of either human or murine RCC cells in normal or low serum or their release of CCR4 ligands (data not shown).

RENCA cells labeled with luciferase were grown orthotopically in their syngeneic hosts, WT BALB/c mice, by injection into the renal capsule of the left kidney. Mice reached humane end point during the course of tumor growth between 17 and 21 days after implantation. Anti-CCR4 antibody treatment significantly inhibited tumor burden as measured by tumor weight and bioluminescence as compared with treatment with an isotype control antibody (Figure 2, B and C). Figure 2B shows the mean tumor weights at end point from 6 independent experiments with 20 mg/kg Affi-5 versus an isotype control, while Figure 2C shows a typical experiment using bioluminescence as a measure of tumor growth. 10 mg/kg Affi-5 also had significant antitumor activity (Figure 2D). There was a significant reduction in the serum concentration of the CCR4 ligand CCL17 in treated mice (Figure 2E). In contrast, the serum concentration of CCL22 was low and did not change following treatment (Figure 2F). Analysis of CCL17 and CCL22 expression in tumor lysates showed that, adjusted for tumor size, CCL17 levels were stable while CCL22 levels were higher in anti-CCR4–treated tumors (data not shown), suggesting that the decrease in CCL17 circulating levels might reflect a reduction in tumor size.

**Actions of the anti-CCR4 antibody on TAMs.** We considered based on the data presented previously, as well as the published literature, that the mechanism of anti-CCR4 inhibition on tumor growth could involve direct effects on malignant cells and/or on leukocytes. To investigate mechanisms of action of the anti-CCR4 antibody, we studied single-cell suspensions from the treated tumors. Several cell types were positive for CCR4 staining in control tumors: macrophages (CD45+CD11b+F4/80+) and different T cell subtypes, such as CD4+ (CD45+CD3+CD4+FoxP3–), CD8+ lymphocytes (CD45+CD3+CD8+), and Tregs (CD45+CD3+CD4+FoxP3+) (Supplemental Figure 4). NK cells were also weakly positive for CCR4 staining (Supplemental Figure 4). Compared

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**Figure 3. Effects of anti-CCR4 on the RENCA TAMs.** BALB/c mice were injected with RENCA-luc cells and treated with Affi-5 (T) or isotype control (C). Mice were sacrificed 17 days after surgery, and tumors were dissociated and characterized by flow cytometry. (A) Tumor-infiltrating macrophages (gated as CD45+CD11b+F4/80+) per milligram of tumor for 5 independent experiments are shown. Tumors were dissociated and characterized by flow cytometry for 2 representative experiments. Two-tailed Student’s t test, **P = 0.0008 and ***P = 0.0085. n = 19 for C, n = 17 for T. (B) Geometric mean of fluorescence intensity (MFI) for MHCI and MR staining on macrophages for 5 independent experiments, and staining for isotype-treated and Affi-5–treated dissociated tumors for 1 representative experiment. Two-tailed Student’s t test, ***P = 0.0008 and **P = 0.0085. n = 19 for C, n = 17 for T. (C) RNA was extracted from macrophages (CD45+CD11b+F4/80+) sorted by flow cytometry from dissociated tumors. The ratio between arginase and Nos2 expression was determined by real-time PCR in 2 independent experiments pooled together (Mann-Whitney U test, *P = 0.035), with n = 6 for C, n = 7 for T. (D) Cells were dissociated at end point from dissected tumors from BALB/c mice treated with Affi-5 (T) or isotype control (C) and plated overnight in the presence of brefeldin A. The fold change in the number of macrophages (CD45+CD11b+F4/80+) positive for intracellular TNF-α is shown from 2 pooled experiments (Mann-Whitney U test, **P = 0.002, n = 11 for C and n = 15 for T).
The arginase/Nos2 expression ratio was significantly lowered by anti-CCR4 treatment (Figure 3D). Purified Affi-5–treated macrophages also showed increased intracellular TNF, further evidence for an M1 cytotoxic phenotype (Figure 3E).

Taken together, these data imply that the phenotype of macrophages from anti-CCR4–treated tumors was altered compared with isotype control–treated tumors and displayed several characteristics associated with an antitumor response.

We next asked whether CCR4 inhibition could have similar activities in another mouse cancer model, testing Affi-5 in the B16 melanoma model. As shown in Supplemental Figure 5, treatment altered the TAM phenotype, with a significant increase in MHCII in myeloid cells in 2 separate experiments (Supplemental Figure 5A). There was, however, no effect on MR expression (Supplemental Figure 5B) or tumor weight (Supplemental Figure 5C) in this rapidly growing model.

**Figure 4. Involvement of T cells in the actions of the anti-CCR4 antibody.** BALB/c mice were injected with RENCA-luc cells and treated with Affi-5 (T) or isotype control (C). Mice were sacrificed 17 days after surgery, and tumors were dissociated and characterized by flow cytometry. (A and B) Number of CD45+CD3+ (A) and CD4+FoxP3+ (Tregs), CD4+FoxP3− (CD4eff), and CD8+ (B) cells/mg of tumor in tumors from isotype-treated (C) or Affi-5–treated (T) mice for 4 experiments pooled together. Two-tailed Student’s t test, (A) **P = 0.003 and (B) *P = 0.01 (Tregs and CD8), *P = 0.02 (CD4eff), with n = 15 for C, and n = 14 for T. (C) Ratio of CD3+CD4+FoxP3+ or CD3+CD8+ to CD3+CD4+FoxP3+ (Treg) lymphocytes in isotype-treated and Affi-5–treated mice for 4 experiments. (D) Two control–treated (control) and 2 Affi-5–treated (treated) tumors from BALB/c mice were lysed, and an amount equivalent to 200 μg tumor lysate was incubated on Proteome Profiler Mouse Cytokine Array Panel A membranes. Average signal for each cytokine was normalized to signal from control-treated tumors. Fold change compared with control, grouped for Th1 cytokines and Th2 cytokines, is shown (2-tailed Student’s t test, P = 0.0173). (E and F) CD3 cells were isolated at end point from dissected tumors from BALB/c mice treated with Affi-5 (T) or isotype control (C). Lymphocytes were stimulated with PMA and ionomycin for 4 hours in the presence of brefeldin A and stained for intracellular IFN-γ. The percentage of IFN-γ−positive CD4+ cells (E) or CD8+ cells (F) is represented. Two independent experiments pooled together are shown (2-tailed Student’s t test, *P = 0.028 for IFN-γ−positive CD4+ cells [E], n = 5 for C, n = 6 for T). (G) Percentage of CD8+ cells positive for granzyme B (GZMB) is represented, from 4 independent experiments.
microenvironment. The number of CD3+ cells per milligram of tumor was significantly higher in Affi-5–treated tumors compared with controls (Figure 4A). As CCR4 has been implicated in the recruitment of Tregs at tumor sites (12, 13), we hypothesized that treatment with anti-CCR4 would reduce the number of CCR4+ Tregs, as observed in adult T cell leukemia patients treated with mogamulizumab (18). However, the number of Tregs per milligram of tumor was higher in anti-CCR4–treated tumors (Figure 4B). Also the number of CD4+ effector cells and CD8+ cells per milligram of tumor was increased with treatment (Figure 4B). As a result of these changes, the ratios of CD4+ T effector or CD8+ T cells to Tregs in tumors were unaffected by Affi-5 treatment (Figure 4C).

However, there was an increase in the amount of Th1 cytokines compared with Th2 cytokines in the Affi-5–treated tumor lysates (Figure 4D), which could help explain the M2/M1 switch observed in the TAMs.

We next explored in more detail the phenotype of CD4+ effector and CD8+ cells. The number of CD4+ cells positive for IFN-γ expression was increased in treated tumors (Figure 4E), while it was unaltered for the CD8+ cells (Figure 4F). Moreover, CD8 staining for granzyme B was not significantly altered by treatment (Figure 4G). Collectively, these data suggested a role for CD4+ cells in the antitumor activity of the anti-CCR4 antibody.

Other treatment-induced changes to tumor-infiltrating leukocyte populations. We observed repeatedly a significant increase in tumor NK cells as a proportion of total CD45+ cells (Figure 5A). Moreover, CD8 staining for granzyme B was not significantly altered by treatment (Figure 4A). Collectively, these data suggested a role for CD4+ cells in the antitumor activity of the anti-CCR4 antibody.

**Other treatment-induced changes to tumor-infiltrating leukocyte populations.** We observed repeatedly a significant increase in tumor NK cells as a proportion of total CD45+ cells (Figure 5A), while the proportion of CD8+ cells was not altered (data not shown). The effects on NK cells was due to changes in cell number, as a significant increase in terms of NK cells/mg tumor could be observed (P = 0.03) (Figure 5B). Also, myeloid-derived suppressor cells (MDSCs), characterized as CD45+CD11b+Gr1+, constituted a reduced percentage of the CD45+ infiltrate in tumors from treated compared with control mice (Figure 5C). As for the MDSCs, there was some variability between experiments in terms of numbers of cells, so we expressed the results as fold change in cells/mg tumor, which could help explain the M2/M1 switch observed in the TAMs.
mg tumor. Pooling results from 4 experiments, we see a significant reduction in treated tumors (P = 0.017) (Figure 5D). A similar reduction of granulocytic (Gr1hi) and monocytic (Ly6Chi) MDSCs was seen in the spleen of treated mice compared with control mice (Figure 5E). To determine whether the accumulated CD11b+Gr1+ cells have a suppressive phenotype and could really be identified as MDSCs, we performed immunosuppression assays using T cells as effectors. Increasing amounts of MDSCs from the tumors of untreated mice effectively suppressed proliferation of activated CD4+ and CD8+ cells isolated from naïve splenocytes (Figure 5F). Similarly, splenic MDSCs isolated from tumor-bearing mice suppressed proliferation of CD4+ and CD8+ cells (data not shown).

Figure 6. Effects of anti-CCR4 on the RENCA tumors require CD4+ cells. (A–D) BALB/c nu/nu mice were injected with 1 × 10^5 RENCA-luc cells and treated with Affi-5 (T) or isotype control (C) (10 mg/kg) twice weekly starting 48 hours after surgery. Mice were sacrificed at 17 days after surgery, and tumor weight was determined (n = 9 C, n = 9 T, not significant). Geometric mean of fluorescence intensity for MHCII (B) and MR (C) staining on macrophages (CD45^+CD11b^+F4/80^+), for isotype-treated and Affi-5-treated dissociated tumors; n = 4. (D) Percentage of NK cells (CD45^+CD3^-DX5^+) among the CD45^+ population; n = 4. (E–H) BALB/c mice were injected with 1 × 10^5 RENCA-luc cells and treated with Affi-5 (T) or isotype control (C) (10 mg/kg) twice weekly starting 48 hours after surgery. Treatment with anti-MHCII or the relevant isotype control (10 mg/kg) was started 1 day prior to surgery and continued with 3 doses per week. Mice were sacrificed 17 days after surgery, tumor weight was determined (n = 6 for each group), and tumors were dissociated and characterized by flow cytometry. (E) Blocking of MHCII has a significant effect on tumor weight (2-way ANOVA, *P = 0.049). Bonferroni post-test showed significant difference (P < 0.05) in weight of Affi-5-treated tumors in the presence versus absence of anti-MHCII. (F and G) Geometric mean of fluorescence intensity (MFI) for MHCII (F) and MR (G) staining on macrophages (CD45^+CD11b^+F4/80^+). There is a significant difference between MHCII and MR expression of macrophages from Affi-5-treated tumors in the presence or absence of anti-MHCII (Kruskal-Wallis test with Dunn post-test, *P < 0.05 and 1-way ANOVA with Bonferroni post-test, **P < 0.001, with n = 3–4 for each group). (H) Percentage of NK cells (CD45^+CD3^-DX5^+) among the CD45^+ population. There is a significant difference (1-way ANOVA with Bonferroni post-test, ***P < 0.001) in the percentage of NK cells from Affi-5-treated tumors in the presence versus absence of anti-MHCII.
Further investigation of CCR4 receptor on tumor CD4+ T cells. As in our model we observed changes to cells of both the adaptive and innate immune response, and some of our evidence pointed to involvement of CD4+ T cells, we wanted to better understand the interplay between these different components. Affi-5 did not inhibit RENCA tumor growth in T cell–deficient nude mice (Figure 6A), and there was also no effect on macrophage phenotype or extent of NK cell infiltrate in tumors in nude mice (Figure 6, B–D). This suggested that adaptive immunity, especially via CD4+ T cells, was upstream of the actions on the cells of the innate immune system.

To further investigate a role for CD4+ T cells in the antitumor actions of Affi-5, we combined this agent with a neutralizing antibody to MHCII. This completely abrogated the effects of the anti-CCR4 antibody on RENCA tumor weight (Figure 6E), macrophage MHCII (Figure 6F), and MR expression (Figure 6G), and percentage of NK cells in the tumor microenvironment (Figure 6H).

We concluded that CD4+ T cells are essential mediators of the actions of the anti-CCR4 receptor antibody Affi-5, and were required for the observed changes in macrophage phenotype and proportion of NK cells.

CCR4 function in normal CD4+ T cells. Our results led us to question whether CCR4 might be involved in direct regulation of Th1 and Th2 responses in normal CD4+ T cells. To explore this, we developed an in vitro assay in which CD4+ T cells were purified from splenocytes of healthy mice and polarized to a Th1 response with IL-12 and IL-2 in the presence of anti-CD3 and anti-CD28.

Figure 7. CCL17 can inhibit Th1 responses in vitro. (A and B) CD4+ cells were isolated from spleens of healthy mice and stimulated with IL-2 and IL-12 in the presence of anti-CD3– and anti-CD28–coated beads. CCL17, CCL22, Affi-5 (10 μg/ml), or isotype control was added after an overnight incubation; after 3 days cells were stimulated with cell stimulation cocktail, harvested, stained for intracellular IFN-γ, and analyzed by flow cytometry. Results of 7 and 3 independent experiments are shown for CCL17 and CCL22, respectively (A), together with representative plots of 1 experiment. Results of 4 independent experiments are shown in B, with representative plots of 1 experiment. *P < 0.05, **P < 0.01, 1-way ANOVA with Bonferroni post-test.
beads. This treatment stimulated production of IFN-γ over the course of 3 days. When CCL17 was added to the CD4+ cells 1 day after the initial stimulation, a significant reduction in the production of IFN-γ was observed (Figure 7A). Although CCL22 produced a similar trend, it was markedly weaker than CCL17 in inhibiting Th1 polarization (Figure 7A). The action of CCL17 on CD4+ cells in vitro was abolished by addition of the anti-CCR4 antibody (Figure 7B). These results indicate that CCL17 might play a role in directly inhibiting the Th1 response, and provide more mechanistic insight into the action of Affi-5. CCL17 in the tumor microenvironment might be secreted by many cell types, including M2-polarized macrophages (21). To support this hypothesis, we measured mRNA levels for Ccl17 and Ccl22 in the different cell populations of the tumor microenvironment. Macrophages showed the highest expression of the two chemokines (Supplemental Figure 6), although a contribution from other cell types cannot be excluded.

Is ADCC involved in the antitumor action of the anti-CCR4 antibody? Finally, as ADCC is implicated in the mechanisms of action of the anti-human CCR4 antibody currently used clinically in treatment of hematological malignancies (16), we investigated the role of ADCC in the actions of Affi-5. While it is possible that murine Fc receptors would interact with a human antibody (22), both a defucosylated (which was used in all the experiments presented up to this point) and a fucosylated version of Affi-5 antibody had similar and significant antitumor effects (Figure 8A). Moreover, the CCR4 antagonist Affi-5 retained antitumor activity on RENCA cells in which CCR4 was silenced by shRNA (Figure 8, B and C). This result indicates that the antitumor effect of Affi-5 occurs primarily through modulation of non-malignant cells in the tumor microenvironment. This is not entirely unexpected, since in vivo CD45+ cells, which include RENCA cells, expressed low levels of CCR4 (Supplemental Figure 4). However, as this antibody has a reported ADCC activity against human lymphoma cells (19), it may act by ADCC on other tumor cells where CCR4 expression is higher.

Discussion

In this study, we provide a comprehensive analysis of the expression of the chemokine receptor CCR4 and its ligands CCL17 and CCL22 in a solid tumor. We present evidence that CCR4 is expressed at significant levels in renal cancer biopsies, where it is associated with the extent of immune infiltrate. Also, expression of CCL17 and CCL22 is altered in renal cancer tissue and in the plasma of patients. In fact, a high CCL17/CCL22 ratio in plasma is associated with a worse prognosis. This is reminiscent of what has been observed in other solid tumors, where there is high CCR4 expression that is generally associated with a poor prognosis (14, 15). CCL22 is also detected in the tumor microenvironment of ovarian, hepatocellular, and breast cancer (10, 12, 13). Our work is further supported by a recent multivariate analysis of CCR4 expression in 53 RCC patient...
biopsies, where CCR4 expression was an independent risk factor for poor prognosis and overall survival (23). Taken together, these observations suggest that CCR4 is an attractive therapeutic target in solid cancers.

In the current work, we have not considered a role for the chemokines CCL2 and CCL5, which may also bind to CCR4, as CCL17 and CCL22 have the highest affinity for the receptor, but in future studies it would be interesting to assess the effect of CCR4 inhibitions on their local and systemic levels.

In this study we report for the first time to our knowledge that an anti-CCR4 antibody has activity in a solid cancer model. As CCR4 is expressed on a number of different immune cells, we had expected that the CCR4 antibody would reduce the number of tumor-infiltrating leukocytes as part of its mechanism of action, but this did not occur. In particular, we did not observe an effect on the number of infiltrating Tregs, which are thought to be recruited though CCR4 in the tumor microenvironment of different tumor types (11–13). In fact, the antibody caused unexpected changes in the phenotype of myeloid cells in the RENCA tumor microenvironment from potentially pro- to antitumor. TAMs mainly consist of a population with little cytotoxicity for tumor cells because of their limited production of NO and proinflammatory cytokines. At the same time, TAMs also possess poor antigen-presenting capability (20). This has led to the notion of depleting TAMs from the tumor microenvironment is an interesting target in cancer therapy. However, it was shown recently that inhibition of the macrophage cell surface receptor CSF1R with a small molecule inhibitor in a model of glioblastoma was able to reduce tumor progression by reducing the M2 polarization of TAMs (24). This work proved that modification of TAM tumor-promoting functions may have a significant impact on tumor growth and that depletion is not strictly necessary for an effective TAM-targeted therapy. In our work, the use of an anti-CCR4 antibody achieved, through a different mechanism of action, a similar change of TAM phenotype, which resulted in a reduction of tumor growth. Findings in human renal cancer provide an interesting correlate to our experimental data, suggesting that stimulating an M2/Th2 to M1/Th1 response may be of therapeutic value. In a recent analysis of the intratumoral immunologic profile of RCC biopsies, the expression of M2 macrophage markers correlated with a poor prognosis and high tumor NOS2 mRNA levels with a good prognosis (25). To confirm the importance of macrophages in the mechanism of action of Affi-5, a depletion experiment with clodronate liposomes was set up, but the partial depletion of the macrophages was accompanied by an M1 polarization of the remaining macrophages (data not shown), thus invalidating this model for testing our hypothesis. The potential of the anti-CCR4 antibody was also suggested in the B16 melanoma experiments, where we found an increase in MHCII expression. However, this “partial switch” was not enough to generate an antitumor effect.

As part of the multiple effects of the anti-CCR4 antibody, we also observed a significant increase in NK cells, as well as reduction in MDSCs and circulating CCL17, which can additionally contribute to decreased tumor growth. To determine the contribution of these different populations infiltrating the tumors, we attempted to deplete NK cells and MDSCs in this model. However, while systemic depletion was successful, we never obtained a satisfactory depletion in the tumor microenvironment. As proven by the experiment in nude mice, these multiple effects are not directly mediated by leukocytes of the innate immune system, but are dependent on the adaptive immune system. Moreover, disrupting the MHCI-TCR interaction abolished the therapeutic effect of the anti-CCR4 antibody and also impacted innate immunity, thus proving an essential role for CD4+ cells, linked to their ability to secrete IFN-γ. Our in vitro assay further supported the hypothesis that the CCR4/CCL17 axis may be involved in maintaining Th2 responses. To our knowledge, the effects described here of CCR4 inhibition in this tumor microenvironment are novel.

We had also predicted that the anti-CCR4 antibody Affi-5 may have ADCC activity in the RENCA model, especially on the malignant cells, but the RENCA cells in vivo had low expression of CCR4. CCR4 is strongly expressed in several human T cell malignancies. Mogamulizumab (KW-0761; Poteligeo), a humanized fucosylated anti-CCR4 antibody that markedly enhances ADCC, is used in the treatment of patients with relapsed or refractory CCR4-positive ATL in Japan (26). Recently it also received approval in Japan for relapsed or refractory CCR4-positive peripheral T cell lymphoma (PTCL) and cutaneous T cell lymphoma (CTCL). We could find no evidence that Affi-5 was working via ADCC in our model system, but this does not preclude such an action in other models or patients where malignant cells have higher levels of CCR4. Neither, we would suggest, does it preclude mogamulizumab having other actions in the tumor microenvironment. Nor could we find evidence that the anti-CCR4 antibody had direct effects on the CCR4-expressing malignant cells. This may be because the CD45– population, which contained the malignant cells, expressed low levels of CCR4 in the tumor microenvironment. In addition, RENCA tumors growing in nude mice did not respond to the anti-CCR4 antibody. As anti-CCR4 inhibited RENCA cell migration in vitro, it is possible that the antibody had an antimetastatic effect, but it was not possible to measure this in our model system.

In summary, we have described here a therapeutic strategy to target solid tumors with significant CCR4 expression in the tumor microenvironment. As targeting CCR4 in hematologic malignancies has shown manageable side effects, this approach could readily be translated into the clinic. Moreover, this opens the possibility for evaluation of combinations of CCR4 inhibition with other immune-modulatory agents. Inhibition of CCR4 had multiple actions in the RCC experimental tumor microenvironment — but predominantly there was evidence of a Th2/M2 to Th1/M1 switch. CCR4 inhibition also increased MHCII expression on TAMs in the B16 model. As neither Treg infiltration nor CD8 activation was affected and the antibody did not alter levels of CTLA4 and PD-L1 in the renal tumor microenvironment, there is a strong rationale for a combination with immune checkpoint blockade. Also combinations with anti-CD40 agonistic antibodies would be an attractive option, as stimulating macrophage activation in a Th1-skewed environment may increase a host antitumor response.

Methods
Reagents. Recombinant human chemokines CCL17 (catalog 300-30) and CCL22 (catalog 300-36) were purchased from Peprotech. Recombinant mouse CCL17 (529-TR) and CCL22 (439-MD) were purchased from R&D Systems.
RNA isolation and real-time PCR. RNA from sorted macrophages was extracted with the RNeasy Micro Kit (QIAGEN) and amplified with the Ovation PicoSL WTA System V2 (NuGEN). Real-time RT-PCR analysis was performed using TaqMan assays (Applied Biosystems) — CCR4 (Hs99999919_m1), 18S (4310893E), arginase 1 (Mm00475988_m1), Nos2 (Mm00440502_m1) — with the ABI StepOnePlus instrument (Applied Biosystems).

Renal tissue and patient plasma samples. Patients' samples were collected from patients who had locally advanced or metastatic RCC, who had progressed after first-line cytokine-based therapy (for locally advanced disease), or who were intolerant to first-line cytokine-based therapy (for locally advanced or metastatic disease). The TMA mainly comprised clear cell renal carcinomas (75%), with some biopsies classified as papillary renal cancer. Clear cell renal carcinomas were identified using standard IHC and carbonic anhydrase IX (CAIX) staining. CCR4 was detected on the malignant cells from 153 of 173 malignant tumors, clear cell and non-clear cell, in our TMA. Controls were age-matched individuals with no malignancies.

IHC. Paraffin-embedded sections (4 μm) were dewaxed and dehydrated, and antigen retrieval was performed by microwaving sections in Antigen Unmasking Solution (Vector Laboratories, H-3300) for 9 minutes. After blocking with the appropriate serum, samples were incubated overnight at 4°C using primary antibodies: CCR4 (ab16669 1:300; Abcam), CCL17 (ab182793 1:100; Abcam), CCL22 (500-P107 1:20; Peprotech), CD3 (A0452, 1:100; Dako), CD68 (M0876 1:50; Dako). Following incubation with a biotinylated secondary antibody (anti-goat, anti-rabbit, or anti-mouse IgG, 1:200; Vector Laboratories) for 30 minutes at room temperature, antigens were revealed with 3,3′-diaminobenzidine (Sigma-Aldrich). Omission of the primary antibody and isotype control antibody were used as negative controls. The scoring for intensity of staining on positive cells was as follows: 0 (no expression), 1 (low expression), and 2 (high expression).

ELISA and Meso Scale Discovery System. Human CCL17 and CCL22 were determined from plasma with Meso Scale Discovery System plates (Human TARC Ultra-Sensitive Kit, K151BGC-1; and Human MDC Ultra-Sensitive Kit, K151BAC; Meso Scale Diagnostic). Mouse CCL17 and CCL22 were determined from plasma or serum with Mouse CCL17/TARC Quantikine ELISA Kit (MCC170) or Mouse CCL22/MDC Quantikine ELISA Kit (MCC220) from R&D Systems.

Cell culture. 786-O and B16F0 cells were obtained from ATCC, while RENCA cells were cultured in DMEM containing 10% FBS and 1% p/s. Streptomycin (p/s), and 1% glutamine. RENCA culture medium was further supplemented with 20 mM PBS, 145 mM NaCl, and 1% heat-inactivated FBS after blocking the right kidney weight from the weight of the tumor-bearing left kidney.

Tumor growth was monitored after administration of luciferin (3 mg/mouse, Sigma-Aldrich) with the IVIS Imaging System 100 (Xenogen Biosciences). Mice were sacrificed between days 17 and 12 after surgery.

The anti-MHCII blocking antibody (clone M5/114) and the isotype control (LTF-2) were obtained from BioXCell and administered i.p. at 10 mg/kg 1 day prior to surgery and 3 times/week thereafter.

For the melanoma model, 8- to 12-week-old C57BL/6 mice from Charles River Laboratories were injected subcutaneously with 1 × 10⁴ B16F0 cells resuspended in PBS. Affi-5 and the appropriate isotype control were injected i.p. twice weekly at 20 mg/kg, starting on day 2 after surgery. Tumor weight was determined at the end of the experiment by subtracting the right kidney weight from the weight of the tumor-bearing left kidney.

RNA isolation and real-time PCR. RNA from sorted macrophages was extracted with the RNeasy Micro Kit (QIAGEN) and amplified with the Ovation PicoSL WTA System V2 (NuGEN). Real-time RT-PCR analysis was performed using TaqMan assays (Applied Biosystems) — CCR4 (Hs99999919_m1), 18S (4310893E), arginase 1 (Mm00475988_m1), Nos2 (Mm00440502_m1) — with the ABI Ste-
CD3) and plated (5 × 10^4/well) in round-bottom 96-well plates. Mouse T-Activator CD3/CD28, Invitrogen) at a ratio of 1:2 (beads/cell). Some of the cells were left without stimulus, while the remaining cells were pulsed with PMA (50 ng/ml, Sigma-Aldrich) and ionomycin (1 μg/ml) for 3 hours. At the end of the incubation, cells were washed; blocked as above; stained with α-CD45 (catalog 48-0451), αLy6-G (Gr1) (catalog 35-5931), αF4/80 (catalog 47-4801, 1:150), αCD11b (catalog 11-0112), for 30 minutes at 4°C. DAPI (2.5 μg/ml) was added prior to sorting, which was performed with a BD FACS Aria II cell sorter.

**Suppression assay.** Tumors were dissociated as for flow cytometry staining and pooled. MDSCs were purified according to the manufacturer’s instructions using the Myeloid-Derived Suppressor Cell Isolation kit (Miltenyi Biotec). Naïve CD3+ cells were isolated from spleens of healthy mice using the Dynabeads FlowComp Pan T kit (Invitrogen). Before stimulation, CD3+ cells were pre-labeled with 5 μM CFSE (eBioscience) for 5 minutes at 37°C in medium and washed. Some of the cells were left without stimulus, and the remaining cells were stimulated with anti-CD3/anti-CD28–coated beads (Dynabeads Mouse T-Activator CD3/CD28, Invitrogen) at a ratio of 1:2 (beads/cell) and plated (5 × 10^4/well) in round-bottom 96-well plates. MDSCs were added to the wells at 1:1, 1:2, and 1:8 ratios (CD3/MDSCs) and incubated for 3 days. At the end of the incubation period, cells were collected and stained for viability, CD11b, CD3, CD4, and CD8 and analyzed by flow cytometry. Staining was performed for 30 minutes at 4°C. Cells were washed, fixed in 2% formaldehyde, and analyzed using a BD LSR Fortessa cytometer. Analysis was performed with FlowJo software.

**Intracellular flow cytometry.** Tumors were dissociated and dissociated as described above. For macrophage staining, cells were plated (0.5 × 10^5/500 μl in a 24-well plate) and incubated overnight with Brefeldin A (Sigma-Aldrich), 20 μg/ml. The following day, cells were stained in PBS + 2% heat-inactivated FBS + 2 mM EDTA after blocking with αCD16/CD32, 1:200 (14-0161, eBioscience) for 15 minutes. Staining antibodies were from eBioscience and were diluted 1:200 unless otherwise specified: αCD45 (catalog 48-0451), αLy6-G (Gr1) (catalog 35-5931), αF4/80 (catalog 47-4801, 1:150), αCD11b (catalog 11-0112), and Fixable Viability Dye eFluor 780 or 506 (eBioscience) diluted 1:1,000. Staining was performed with Fixable Viability Dye eFluor 780 or 506 (eBioscience) diluted 1:1,000 for 30 minutes at 4°C. DAPI (2.5 μg/ml) was added prior to sorting, which was performed with a BD FACS Aria II cell sorter.

**Immunofluorescence.** Renal carcinoma 786-O and RENCA cells were cultivated on a chamber slide (Nalge Nunc International) for 1-2 days. Cells were then fixed for 30 minutes with 4% formaldehyde and permeabilized with 0.5% Triton X-100 in PBS for 10 minutes. Samples were blocked with 1% BSA for 2 hours at room temperature, then incubated overnight at 4°C with 25 μg/ml αCCR4 (IMG-322, IMGENEX, Novus Biologicals). After washing, samples were incubated with Alexa Fluor 594–conjugated secondary antibody, 1:2,000, for 2 hours at room temperature. Finally, samples were washed and mounted with ProLong Gold DAPI (Invitrogen, P36931). Cells were then visualized using a Zeiss LSM 510 confocal microscope.

**RNA interference for CCR4 in renal cancer cell lines.** Commercially available shRNAs, based on the PRS vector, were purchased from Origene Technologies. Four non-overlapping sequences were provided for each available shRNA sequence, based on the pRS vector, were purchased from Origene Technologies. Four non-overlapping sequences were provided for each target human (TR314127) and mouse (TR500386) CCR4. A non-specific shRNA sequence (shGFP, TR30003) and empty vector (TR20003) served as controls. Phoenix packaging cell line was transfected overnight with LipofectAMINE 2000 (Invitrogen) and 5 μg of the shRNA plasmid DNA. After an incubation with complete medium at 33°C and 5% CO2 for 16 hours, the supernatant was collected, filtered, and diluted 8:10 in RCC medium with the addition of Polybrene.

**Cytokine expression.** Frozen tumors were lysed with a gentleMACS M tube (Miltenyi Biotec) in PBS with Complete Protease Inhibitors (Roche). Triton X-100 was added (%), and lysates were cleared by centrifugation and applied to Proteome Profiler Mouse Cytokine Array Panel A membranes (R&D Systems). An equivalent amount of 200-μg tumor lysate was incubated according to the manufacturer’s instructions. Film exposures were quantified using NIH ImageJ software, subtracting a lane background.

**In vitro Th1 polarization assay.** Splenocytes from healthy BALB/c mice were obtained by mashing spleens through a 70-μm strainer and lysis of red blood cells. CD4+ cells were purified with the Miltenyi Biotec CD4+ T cell purification kit according to the manufacturer’s instructions. CD4+ cells were resuspended in Iscove’s modified DMEM (10%FBS, 50 μM β-mercaptoethanol, 8 mM glucose) and stimulated with 1:1 Dynabeads Mouse T-Activator CD3/CD28 (Invitrogen), mouse IL-2 20 ng/ml, and mouse IL-12 5 ng/ml (from R&D Systems). The day after, the indicated concentrations of CCL17 or CCL22 in combination with Affi-5 (30 μg/ml) were added to the plate. Experiments with Affi-5 were performed in Corning Ultra-Low attachment plasticware (Corning Life Sciences). Cells were stimulated with restimulation cocktail plus transport inhibitor from eBioscience for 5 hours and stained for 30 minutes at 4°C with αCD16/CD32 (14-0161, eBioscience, 1:200), Fixable Viability Dye eFluro 450 1:1000, αCD4 (560783, BD Horizon CD4, BD Biosciences, 1:300) in 50 μl FACS buffer. After washing, cells were permeabilized and stained with αIFN-γ (17-7311, eBioscience, 1:100) or isotype control. After washing, cells were analyzed using a BD LSR Fortessa cytometer.

**RNA interference for CCR4 in renal cancer cell lines.** Commercially available shRNAs, based on the PRS vector, were purchased from Origene Technologies. Four non-overlapping sequences were provided to target human (TR314127) and mouse (TR500386) CCR4. A non-specific shRNA sequence (shGFP, TR30003) and empty PRS vector (TR20003) served as controls. Phoenix packaging cell line was transfected overnight with LipofectAMINE 2000 (Invitrogen) and 5 μg of the shRNA plasmid DNA. After an incubation with complete medium at 33°C and 5% CO2 for 16 hours, the supernatant was collected, filtered, and diluted 8:10 in RCC medium with the addition of Polybrene.
RENCA or 786-O cells at 30% confluence were infected for 10 hours twice within 48 hours. Virus-infected cells were selected with 1.5 μg/ml puromycin (InvivoGen). For long-term silencing in vivo, RENCA-luc cells were infected with 3 lentiviral GIPZ vectors targeting mouse CCR4 from Thermo Fisher Scientific. Viral particles were obtained by transfecting HEK293T cells with the calcium phosphate method with 21 μg shRNA construct, 7 μg VSVG construct, 14 μg HIV construct (from Addgene), in a 14-cm dish, and collecting the supernatant for 24 hours. RENCA-luc cells were infected for 24 hours in a 6-well plate with 1 ml supernatant. Virus-infected cells were selected with 0.3 μg/ml puromycin (Sigma-Aldrich). After verifying the silencing, cells infected with vector V3LM_M_439088 were chosen for the in vivo experiment.

**Statistics.** All data are expressed as mean ± SEM. Differences were considered significant at P < 0.05, using a Student’s t test (2-tailed), ANOVA test, or nonparametric test as appropriate, performed with the statistical analysis software Prism (GraphPad Software). P values are specified.

**Study approval.** Patient samples were collected under the Multi-centre Research Ethics Committee (MREC); Ethical Number MREC 02/8/78. All patients provided written informed consent. All experimental procedures were performed according to a protocol approved under Home Office licence 70/4711 and according to the Animal Welfare and Ethical Review Body (AWERB) of Queen Mary University of London.

**Author contributions**

CB designed, performed, and interpreted most of the experiments, and wrote the manuscript. MNK designed, performed, and interpreted the experiments shown in Figure 1 and Supplemental Figures 1-3. TS, AM, and EM contributed to designing, performing, and interpreting some experiments and editing the manuscript. RT performed most of the surgery experiments. MJ conducted and analyzed the B16 melanoma studies. MC measured and analyzed the chemokine levels in patients. HK contributed to the design of the research studies. UBH and ARD provided the Affi-5 antibody and contributed to the design of the experiments. LF and RWW contributed to the design of the research studies and editing the manuscript. TP provided patient samples, patient information, and intellectual input. SAQ contributed to designing and interpreting experiments analyzing adaptive immunity. FRB conceived of, designed, and supervised the project and experimental plan, interpreted experiments, and wrote the manuscript.

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