Anti-TNF immunotherapy reduces CD8+ T cell–mediated antimicrobial activity against Mycobacterium tuberculosis in humans

Heiko Bruns,1 Christoph Meinken,2 Philipp Schauenberg,3 Georg Härter,4 Peter Kern,4 Robert L. Modlin,5 Christian Antoni,3 and Steffen Stenger1

1Institute for Medical Microbiology and Hygiene, University Hospital of Ulm, Ulm, Germany. 2Institute for Clinical Microbiology, Immunology and Hygiene, and Department for Internal Medicine 3, University Erlangen-Nürnberg, Erlangen, Germany. 3Section of Infectious Diseases and Clinical Immunology, University Hospital of Ulm, Ulm, Germany. 4Department of Medicine, Division of Dermatology, UCLA David Geffen School of Medicine, Los Angeles, California, USA.

The incidence of tuberculosis is increased during treatment of autoimmune diseases with anti-TNF antibodies. This is a significant clinical complication, but also provides a unique model to study immune mechanisms in human tuberculosis. Given the key role for cell-mediated immunity in host defense against Mycobacterium tuberculosis, we hypothesized that anti-TNF treatment impairs T cell–directed antimicrobial activity. Anti-TNF therapy reduced the expression in lymphocytes of perforin and granulysin, 2 components of the T cell–mediated antimicrobial response to intracellular pathogens. Specifically, M. tuberculosis–reactive CD8+CCR7–CD45RA+ effector memory T cells (TEMRA cells) expressed the highest levels of granulysin, lysed M. tuberculosis, and infected macrophages and mediated an antimicrobial activity against intracellular M. tuberculosis. Furthermore, TEMRA cells expressed cell surface TNF and bound the anti-TNF therapeutic infliximab in vitro, making them susceptible to complement-mediated lysis. Immune therapy with anti-TNF was associated with reduced numbers of CD8+ TEMRA cells and decreased antimicrobial activity against M. tuberculosis, which could be rescued by the addition of CD8+ TEMRA cells. These results suggest that anti-TNF therapy triggers a reduction of CD8+ TEMRA cells with antimicrobial activity against M. tuberculosis, providing insight into the mechanism whereby key effector T cell subsets contribute to host defense against tuberculosis.

Introduction
The treatment of patients with monoclonal anti-TNF antibodies (infliximab) has led to a 4-fold increased incidence of tuberculosis (1), representing a major complication of anti-TNF therapy. Despite this potential side effect, infliximab treatment has revolutionized the treatment of severe forms of RA (2, 3), Crohn disease, psoriatic arthritis, and ankylosing spondylitis (AS) (4). The rationale for immune intervention with TNF-neutralizing agents resulted from 2 decades of basic research showing that TNF is a central component in the cascade of cytokines that trigger cell recruitment, inflammation, and ultimately tissue destruction (5). However, TNF is also a key molecule in host immunity to intracellular bacteria, most notably tuberculosis (6). Mice deficient in TNF production are highly susceptible to primary infection with Mycobacterium tuberculosis (7), and depletion of TNF by treatment with anti-TNF antibodies results in reactivation of latent disease (8–11). TNF fails to elicit antimycobacterial activity in human host cells in vitro (12–15), although in vivo, the critical role is illustrated by the reactivation of latent disease in anti-TNF-treated individuals (16, 17).

A key aspect of host defense against tuberculosis is the antigen-specific adaptive T cell response. Traditionally, the T cell–mediated response against M. tuberculosis was believed to be dominated by IFN-γ–releasing CD4+ T cells. However, it has become increasingly clear that cytotoxic CD4+ T cell subsets (18–20) and CD8+ T cells contribute to protective immunity against M. tuberculosis (21). CD8+ T cells contribute to host defense by the release of Th1 cytokines and the coordinate interaction of chemotactic, lytic, and antimicrobial molecules, including the antimicrobial pathway mediated by perforin and granulysin (22, 23).

Granulysin is a member of the saposin-like family of lipid-binding proteins (24) that is expressed in intracellular granules of cytotoxic lymphocyte subsets, including NK cells (25), CD4+ T cells (19, 26), CD8+ T cells (22, 23, 27), NKT cells (28), and TCRδ T cells (29). The major functions are lysis of tumor cells (30), support of inflammation (31, 32), and killing of microbial pathogens (22, 25, 33–36). Current knowledge of the relevance of the perforin/granulysin pathway of antimicrobial activity in vivo is limited because suitable animal models are not available, largely because there is no granulysin homolog in mice.

The use of anti-TNF monoclonal antibodies as immune therapy in patients with autoimmune disease is associated with an increased incidence of tuberculosis and provides what we believe to be a unique clinical model to search for immunological mechanisms that are critical for M. tuberculosis containment and that predispose to the reactivation of latent tuberculosis in humans. We therefore hypothesized that anti-TNF immune therapy alters a key T cell population required for host defense against tuberculosis in humans. The results presented here demonstrate that granulysin–expressing CD8+CCR7–CD45RA+ effector memory T cells (TEMRA cells) decreased during anti-TNF treatment. This subset had anti-
microbial activity and may be critical for maintaining latency in patients infected with *M. tuberculosis*.

**Results**

We hypothesized that anti-TNF immune therapy leads to an alteration of the adaptive T cell response against *M. tuberculosis*, thereby triggering reactivation of latent tuberculosis. Given the key roles of perforin and granulysin in a T cell–mediated antimicrobial pathway, we initially studied expression of these proteins in patients receiving anti-TNF therapy. Patients with active RA (*n* = 9) or AS (*n* = 8; Table 1) that were scheduled to receive therapy with anti-TNF antibodies (infliximab) were recruited, and the expression of perforin and granulysin in lymphocytes was compared before the first infliximab infusion and before the second infusion 2 weeks later by intracellular flow cytometry (Figure 1). During the course of immunotherapy, perforin and granulysin expression dropped significantly in all donors tested (*P* < 0.001), indicating that the perforin/granulysin axis of antimicrobial activity was disturbed by anti-TNF treatment. The levels of perforin and granulysin expression in the peripheral blood of patients with autoimmune diseases was not significantly different compared with healthy donors (perforin, 19% ± 17% in 36 healthy donors; granulysin, 23% ± 21% in 39 healthy donors).

Because TNF had no direct effect on the gene or protein expression of granulysin in PBMCs (data not shown), we reasoned that changes in lymphocyte subsets were responsible for the reduction of perforin+ and granulysin+ lymphocytes. Because granulysin is expressed in numerous lymphocyte subgroups (19, 22, 23, 25–29), we initially sought to identify subsets with prominent granulysin expression in the peripheral blood. We labeled PBMCs from 27 healthy donors with granulysin and markers for T cells (CD4 and CD8), NK cells (CD56), and B cells (CD19). The number of granulysin+ cells ranged from 11% to 46% and was 23% on average (data not shown). The majority of granulysin+ cells were NK cells (51% ± 27%) or CD8+ T lymphocytes (28% ± 13%; Figure 2).

CD8+ T cells can be subdivided into functional subsets, including naïve T cells, memory T cells, and at least 2 types of effector T cells (37–39). These subsets were defined according to the expression of CCR7 and CD45RA: naïve cells are CCR7+CD45RA+, central memory T (TCM) cells are CCR7+CD45RA+, effector memory T (TEM) cells are CCR7−CD45RA+, and TEMRA cells are CCR7−CD45RA+. To identify CD8+ T cell subsets with pronounced expression of granulysin, we quadruple-labeled lymphocytes of healthy donors with antibodies to granulysin, CD8, CD45RA, and CCR7 (Figure 3). The majority of CD8+ ‘granulysin+’ cells were TEM (91% ± 4%), and approximately half were TEMRA cells. CD8+ NK or NK cells expressed no marked amounts of intracellular granulysin (Figure 3A). Similarly, perforin was predominantly expressed in TEM (96% ± 6%), with a clear preference for TEMRA (73% ± 18%; Figure 3B). This expression pattern was very similar to that of granulysin and further supported the functional link between lytic and antimicrobial effector molecules in this subset.

To evaluate whether CD8+ TEM or TEMRA cells mediate anticytotoxic activity, we first measured cytotoxic activity of purified TEM cells, TEMRA cells, TCM cells, or PBMCs against autologous monocytes infected with virulent *M. tuberculosis*. TEM cells, most notably TEMRA cells, were highly efficient in lysing infected macrophages (Figure 4A). Significant lysis was detected at an effector/target ratio of 3:1 (12% ± 2%) and reached 36% ± 3% at an effector/target ratio of 30:1. Total PBMCs and TEM cells induced less than 50% lysis compared with TEMRA cells at effector/target ratios of 10:1 and 30:1. Lytic activity was antigen specific, because lysis of uninfected monocytes was below 7% in all samples (data not shown). Because perforin-mediated pore formation facilitates granulysin-mediated antimicrobial activity, we next cocompatible with *The Journal of Clinical Investigation*. For more information, see <http://www.jci.org>.

Table 1

<table>
<thead>
<tr>
<th>Patient no.</th>
<th>Disease</th>
<th>Disease duration (yr)</th>
<th>Age (yr)</th>
<th>Sex</th>
<th>Erythrocyte sedimentationa</th>
<th>TSTb</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>RA</td>
<td>6</td>
<td>58</td>
<td>M</td>
<td>47/24</td>
<td>Negative</td>
</tr>
<tr>
<td>2</td>
<td>RA</td>
<td>10</td>
<td>48</td>
<td>F</td>
<td>65/40</td>
<td>Negative</td>
</tr>
<tr>
<td>3</td>
<td>RA</td>
<td>7</td>
<td>34</td>
<td>F</td>
<td>6/5</td>
<td>Negative</td>
</tr>
<tr>
<td>4</td>
<td>RA</td>
<td>6</td>
<td>61</td>
<td>M</td>
<td>51/75</td>
<td>Negative</td>
</tr>
<tr>
<td>5</td>
<td>RA</td>
<td>22</td>
<td>60</td>
<td>F</td>
<td>29/15</td>
<td>Negative</td>
</tr>
<tr>
<td>6</td>
<td>RA</td>
<td>5</td>
<td>28</td>
<td>M</td>
<td>19/17</td>
<td>Negative</td>
</tr>
<tr>
<td>7</td>
<td>RA</td>
<td>28</td>
<td>47</td>
<td>M</td>
<td>42/30</td>
<td>Negative</td>
</tr>
<tr>
<td>8</td>
<td>RA</td>
<td>12</td>
<td>64</td>
<td>M</td>
<td>41/12</td>
<td>Negative</td>
</tr>
<tr>
<td>9</td>
<td>AS</td>
<td>6</td>
<td>33</td>
<td>M</td>
<td>18/3</td>
<td>Negative</td>
</tr>
<tr>
<td>10</td>
<td>AS</td>
<td>3</td>
<td>37</td>
<td>F</td>
<td>8/2</td>
<td>Negative</td>
</tr>
<tr>
<td>11</td>
<td>AS</td>
<td>1</td>
<td>30</td>
<td>M</td>
<td>1/3</td>
<td>Negative</td>
</tr>
<tr>
<td>12</td>
<td>AS</td>
<td>2</td>
<td>31</td>
<td>M</td>
<td>67/2</td>
<td>Negative</td>
</tr>
<tr>
<td>13</td>
<td>AS</td>
<td>14</td>
<td>36</td>
<td>M</td>
<td>20/3</td>
<td>Negative</td>
</tr>
<tr>
<td>14</td>
<td>AS</td>
<td>8</td>
<td>43</td>
<td>F</td>
<td>32/10</td>
<td>15 mm</td>
</tr>
<tr>
<td>15</td>
<td>AS</td>
<td>9</td>
<td>29</td>
<td>M</td>
<td>39/7</td>
<td>11 mm</td>
</tr>
<tr>
<td>16</td>
<td>AS</td>
<td>11</td>
<td>52</td>
<td>M</td>
<td>16/5</td>
<td>12 mm</td>
</tr>
<tr>
<td>17</td>
<td>RA</td>
<td>12</td>
<td>22</td>
<td>F</td>
<td>25/17</td>
<td>18 mm</td>
</tr>
</tbody>
</table>

M, male; F, female. aValues (in mm) show sedimentation at 1 hour/2 hours. bPositive values denote diameter of induration.

*Figure 1*

Decreased expression of lytic and antimicrobial effector molecules in patients treated with infliximab. PBMCs from patients with active RA or AS were stained for perforin (*n* = 7) or granulysin (*n* = 17) before and 2 weeks after beginning therapy. The percentage of granulysin+ and perforin+ lymphocytes was determined by flow cytometry. Shown are results from all individual donors tested.
tured purified TEMRA cells, TEM cells, TEM cells, and PBMCs with M. tuberculosis–infected monocytes and determined mycobacterial survival after 36 hours. The pattern of antimicrobial activity was very similar to the lytic activity (Figure 4B). Reduced bacterial growth was most remarkable for TEMRA cells (47% ± 6%) compared with TEM cells (28% ± 3%), with little or no effect for TEM cells or PBMCs. Antimicrobial activity was antigen specific, because T cells purified from donors negative for tuberculin skin test (TST–donors) had no effect on mycobacterial survival (Figure 4B). Taken together, these results show that CD8+ TEMRA cells account for the majority of antimycobacterial effector cells in the peripheral blood.

To determine whether immunotherapy with anti-TNF antibodies modifies the frequency of CD8+ TEMRA cells in vivo, we monitored the numbers of this antimicrobial lymphocyte subset in 7 patients during treatment. While the percentage of CD8+ T cells, TEM cells, and TEM cells did not change during the first 2 weeks of therapy, the frequency of TEMRA cells decreased in all donors during infliximab treatment by 29% on average (range, 15%–55%; Figure 5A and Figure 3). The majority of granulysin+ cells in the peripheral blood are cytotoxic lymphocytes. PBMCs from healthy donors were stained with PE-Cy5–conjugated anti-CD4, PE-conjugated anti-CD56, allophycocyanin-conjugated anti-CD19, PerCP-conjugated anti-CD8 (filled histograms), or appropriate isotype controls (open histograms). All samples were then stained for granulysin using FITC-conjugated donkey anti-rabbit as a secondary antibody. Granulysin+ cells were gated according to the isotype (control rabbit serum, not shown) and analyzed for the expression of cell surface markers. The percentages of CD4+, CD56+, CD19+, and CD8+ cells within the granulysin gate are indicated. Shown is a typical result of 27 donors.

Figure 2

Figure 3

The majority of CD8+ and granulysin+ or perforin+ T cells are effector cells. PBMCs from healthy donors were stained with PerCP- or allophycocyanin-conjugated anti-CD8 and anti-granulysin (detected with donkey anti-rabbit biotin and streptavidin) or PE-conjugated perforin. Additional labeling was performed using allophycocyanin-conjugated anti-CD45RA and FITC-conjugated anti-CCR7 to detect TEM cells, PE-conjugated anti-CD161 and FITC-conjugated anti-Vα24 to detect NKT cells, and PE-conjugated CD56 and FITC-conjugated anti-CD16 to detect NK cells. Samples with appropriate isotypes were included in all experiments. (A) CD8+granulysin+ cells were gated, and the expression of additional markers within this gate was determined. Numbers denote the percentage of positive effector cells, NKT cells, or NK cells within the population of CD8+granulysin+ cells. For each sample, 1 × 10^6 cells were acquired. Shown is a typical result of 8 donors. (B) CD8+perforin+ cells were gated, and the expression of additional markers within this gate was determined. Numbers denote the percentage of positive effector cells, NKT cells, or NK cells within the population of CD8+perforin+ cells. For each sample, 1 × 10^6 cells were acquired. Shown is a typical result of 4 donors.
The total lymphocyte counts, as determined by automated counting (range, 5,212–8,691 μl), and the percentage of CD8+ T cells (Figure 5A) remained stable in individual patients during therapy. Accordingly, the relative decrease in the percentage of TEMRA cells was reflected in a decrease in the absolute number of TEMRA cells (range, 521–894 cells/μl before therapy, 364–622 cells/μl during therapy) in all 7 patients. An increase in the percentage of naive T cells compensated for the reduced frequency of TEMRA cells (data not shown).

The number of TEMRA cells returned to pretherapeutic levels after 3 months of therapy in all 7 donors and was stable at 1 year in the 5 donors that were available for follow-up (Figure 5B). Interest-
ingly, the majority of tuberculosis cases occur during the first 3 months of anti-TNF therapy (40), within the time period of reduced TEMRA cell levels.

To investigate the mechanism by which anti-TNF immunotherapy resulted in a decrease of CD8⁺ TEMRA cells, we explored whether freshly isolated PBMCs from patients with active RA express membrane TNF on the cell surface. The expression of membrane TNF, as measured by labeling with biotinylated infliximab, was higher in lymphocytes of 4 RA patients (mean, 2.1%) than in those of 4 healthy controls (mean, 0.3%; Figure 6, A and B). Further subset analysis demonstrated that 8.3% ± 3.6% of CD8⁺ TEMRA cells expressed infliximab, with the level undetectable in healthy controls (Figure 6, C and D), indicating an enrichment of infliximab⁺ cells within the CD8⁺ TEMRA cell subset in RA patients. A considerable proportion of T cells expressing membrane TNF was CD8⁻ (63% ± 17%; Table 2). Most likely, these cells are CD4⁺ lymphocytes that play a major role in the pathogenesis of RA. Within the CD8⁻ TEMRA cell subset in RA patients. A considerable proportion of T cells expressing membrane TNF was CD8⁻ (63% ± 17%; Table 2). Most likely, these cells are CD4⁺ lymphocytes that play a major role in the pathogenesis of RA. Within the CD8⁻ TEMRA cells, TEMRA cells were by far the most prominent subset expressing membrane TNF (69% ± 10%, Table 2), implicating their central role in anti-TNF–triggered immune modulation.

Based on these observations, we reasoned that exogenously added complement would be activated to attack anti-TNF-coated TEMRA cells. Freshly isolated PBMCs from patients with active RA were cultured with infliximab or a control antibody before addition of rabbit complement. The frequency of TEMRA cells decreased in the PBMCs from all 4 patients treated with infliximab compared with control antibody–treated PBMCs (Figure 7). The percentage of infliximab⁺ TEMRA cells (8.3% ± 3.6%, Figure 6B) was similar to the average decrease in CD8⁺ TEMRA cells by complement treatment in vitro (6.8% ± 2.6%, Figure 7). Taken together, these observations suggest that antimicrobial CD8⁺ TEMRA cells are a preferred target for complement-mediated lysis in infliximab treated patients.

To evaluate whether the reduction in TEMRA cells is reflected in the ability to kill M. tuberculosis, we measured antimicrobial activity of PBMCs from patients with active RA were labeled for membrane TNF with infliximab (detected with PE-conjugated streptavidin), CD8, CD45RA, and CCR7. Shown are the percentage of total CD8⁺ TEMRA cells, based on gating of infliximab⁺ cells, and the percent distribution of subsets within all CD8⁺ infliximab⁺ T cells.

Table 2
Characterization of membrane TNF–expressing CD8⁺ T cells in RA patients

<table>
<thead>
<tr>
<th>Patient no.</th>
<th>Total CD8⁺</th>
<th>Naive</th>
<th>TEM</th>
<th>TEMRA</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>31%</td>
<td>9%</td>
<td>23%</td>
<td>4%</td>
</tr>
<tr>
<td>2</td>
<td>24%</td>
<td>18%</td>
<td>1%</td>
<td>1%</td>
</tr>
<tr>
<td>3</td>
<td>57%</td>
<td>8%</td>
<td>23%</td>
<td>8%</td>
</tr>
</tbody>
</table>

PBMCs from patients with active RA were labeled for membrane TNF with infliximab (detected with PE-conjugated streptavidin), CD8, CD45RA, and CCR7. Shown are the percentage of total CD8⁺ T cells, based on gating of infliximab⁺ cells, and the percent distribution of subsets within all CD8⁺ infliximab⁺ T cells.
PBMCs in patients before and during anti-TNF therapy. Monocytes were isolated from TST+ patients before and after the beginning of anti-TNF therapy and infected with virulent M. tuberculosis. Infected monocytes were incubated with autologous PBMCs, and the number of viable bacilli was determined after 24 and 96 hours of incubation. PBMCs from patients before therapy limited bacterial growth more efficiently than did PBMCs from patients during anti-TNF therapy at both time points (Figure 8A). This was not the result of differences in the uptake of bacilli by macrophages, because the initial load was similar in both groups, as determined by acid fast staining and plating of cell lysates before addition of PBMCs (data not shown). In contrast, there was no significant difference in antigenspecific IFN-γ release between PBMCs obtained before and during therapy (Figure 8B). This result supports a model whereby lytic and antimicrobial CD8+ TEMRA cells comprise a functionally unique subset that complements the cytokine-producing (e.g., IFN-γ) CD4+ T lymphocytes in order to eliminate M. tuberculosis.

To determine whether the relative deficiency of CD8+ TEMRA cells was responsible for the reduced antimicrobial activity, PBMCs from TST+ infliximab-treated patients were replenished with autologous, sorted CD8+ TEMRA cells, and mycobacterial growth was measured as described above. The initial bacterial load was 3.3 ± 0.7 × 10^4 CFU, reflecting the facts that not all mycobacteria added to the culture were taken up and that not all macrophages take up mycobacteria (21% ± 12% of the macrophages were infected). The relative inability of PBMCs from patients during anti-TNF therapy to limit the intracellular proliferation of virulent M. tuberculosis was rescued by CD8+ TEMRA cells (Figure 9).

Although the percentage reduction of CFUs induced by TEMRA cells was within an order of magnitude, M. tuberculosis infection in vivo is slow and protracted, and the time of in vitro assay was only 36 hours, so that a cumulative antimicrobial effect over time could have a profound effect on the number of bacilli during the course of infection. TEM cells were also effective, whereas TEMCM cells and B cells had no effect on mycobacterial viability, reflecting the levels of granulysin expression (Figures 2 and 3).

In summary, these experiments identify CD8+ ‘granulysin’ TEMRA cells as antimicrobial effector cells in humans. Furthermore, these findings indicate that treatment of individuals with anti-TNF antibodies reduces the frequency of granulysin+ TEMRA cells, providing a mechanism for the reactivation of latent tuberculosis infection.

**Discussion**

The reactivation of latent tuberculosis is a clinically significant complication of immunotherapy with anti-TNF antibodies, but it also provides an opportunity to investigate mechanisms of human host defense against mycobacterial infection in humans. Here we found that anti-TNF treatment induced a marked decrease in the IFN-γ production by PBMCs from patients with active RA or AS as compared with before therapy. The reduced antimicrobial activity of PBMCs correlates with the relative deficiency of CD8+ TEMRA cells and the reduced granulysin expression resulting from anti-TNF treatment. Infliximab therapy reduces the antimicrobial activity of PBMCs. PBMCs from TST+ patients (n = 4) with active RA or AS were collected before and 2 weeks after the onset of anti-TNF therapy and frozen in liquid nitrogen to allow for simultaneous measurement of antimycobacterial activity. Matched pairs were thawed, and the number of viable bacilli was determined by plating the cell lysates at 24 and 96 hours after infection. Shown is the mean ± SEM number of CFUs of all 4 donors tested. (A) The number of viable bacilli was determined by plating the cell lysates at 24 and 96 hours after infection. Shown is the mean ± SEM number of CFUs of all 4 donors tested.
of perforin and granulysin in lymphocytes, 2 components of direct T cell-mediated antimicrobial activity against intracellular \textit{M. tuberculosis}. This reduction was caused by decreased frequency of the T$_{EMRA}$ cell subset, which expressed granulysin and mediated a direct antimicrobial activity against \textit{M. tuberculosis} in macrophages. T$_{EMRA}$ cells expressed membrane TNF and bound infliximab, becoming susceptible to complement-mediated lysis. Together, these data identify the T$_{EMRA}$ cell subset as a requirement for optimal protection against tuberculosis in humans.

The experimental evidence implicating CD8$^+$ T cells as pivotal in host defense in tuberculosis began with adoptive transfer and cell depletion studies in vivo in mouse models (41–43). Subsequent work showed that disruption of MHC class I antigen presentation to CD8$^+$ T cells severely impaired host defense (44). Human \textit{M. tuberculosis}-reactive CD8$^+$ T cell lines and clones were found to lyse \textit{M. tuberculosis}-infected macrophages in an antigen-specific manner and to restrict the growth of \textit{M. tuberculosis} in macrophages (45, 46). Our own prior studies indicated that CD8$^+$ T cells contained one or more microbicidal effector molecules in their cytotoxic granules that are able to directly kill intracellular \textit{M. tuberculosis} (47), including the antimicrobial protein granulysin (22). Here, linking T cell phenotypic and functional changes that occur during anti-TNF immunotherapy, we identified that a key granulysin itself is an antimicrobial peptide with a broad range of antimicrobial activities against Gram-positive and Gram-negative organisms, including \textit{M. tuberculosis}. The mechanism by which granulysin kills bacteria depends upon its positive charge and $\alpha$-helical structure facilitating binding to bacterial membranes and subsequent lysis of the pathogen (48). The tissue expression of granulysin correlated with the ability of the host to restrict infection in the mycobacterial disease leprosy (26). Plasma granulysin levels were found to correlate with host defense responses in human tuberculosis (49–51). Although granulysin is likely to contribute to host defense against microbial infection, it may also contribute to tissue injury by inducing necrosis of normal cells. For example, granulysin has previously been shown to induce keratinocyte death in Stevens-Johnson syndrome and toxic epidermal necrolysis (52), leading to skin blister formation. It remains to be determined whether granulysin contributes to tissue damage in human tuberculosis, such as the caseation necrosis that is a hallmark of the disease. Although our present findings as well as previous studies into the role of granulysin involve human donors and by nature are somewhat correliative, it is important to note that granulysin has no murine homolog, such that there are no adequate mouse models to study its role in vivo.

The demonstration that CD8$^+$ T$_{EMRA}$ cells are required for optimal host defense against tuberculosis is consistent with, yet also extends, prior findings. T$_{EMRA}$ cells were previously recognized as the terminally differentiated stage of T$_{EM}$ cells that develop under the driving force of homeostatic mediators such as IL-15 (39, 53, 54). This multistep maturation process involves conversion of CCR7$^+$ CD45RA$^+$ lymphocytes to CCR7$^+$ CD45RA$^-$ cells that express high levels of perforin, granzymes (37), and granulysin. There is a positive correlation between the frequency of T$_{EMRA}$ cells and efficient control of HIV-1 infection (55, 56), and a maturation defect of T$_{EMRA}$ cells may represent an HIV-induced evasion mechanism (53). In Ebstein-Barr virus disease and cytomegalovirus-infected patients, T$_{EMRA}$ cells are a major component of the antigen-specific effector cell pool in the blood and affected tissue and may account for the successful control of these viruses that regularly occurs in immunocompetent individuals (53, 57, 58). A role for CD8$^+$ T$_{EMRA}$ cells in tuberculosis was previously suggested by one elegant study measuring \textit{M. tuberculosis}–specific responses by pentamer staining. In healthy, protected TST$^+$ children, the majority of antigen-specific CD8$^+$ cells were T$_{EMRA}$ cells (59), whereas in patients that failed to control the tubercle bacilli, antigen-specific CD8$^+$ cells were predominantly T$_{CM}$ cells (59). In another study, children with active tuberculosis had an increased frequency of CD8$^+$ CD45RO$^-$ CD28 CD27$^+$ effector cells, which are probably closely related to
T\textsubscript{EMRA} cells (60). Cells within this subset showed clonal expansion in severe disease, making a protective function counterintuitive. One previous report demonstrated that the frequency of T\textsubscript{EMRA} cells remains unchanged during the first 3 days of anti-TNF treatment (61). However, the observation period of the present study spanned the first weeks of therapy, which covers the time frame during which reactivation of tuberculosis occurs. The role of T\textsubscript{EMRA} cells in host defense against tuberculosis warrants further investigation in other clinical settings, including in patients with tuberculous pleuritis, in which cell-mediated immunity is strong (62), and in cases of tuberculosis in the context of HIV infection, in which cell-mediated immunity is impaired (63). Additional studies will clarify whether CD8\textsuperscript{+} T\textsubscript{EMRA} cells, in addition to directly recognizing and eliminating M. tuberculosis–infected macrophages, also participate in the formation and maintenance of granulomas, as would be predicted by their unique and specific chemokine receptor expression pattern (39, 64).

Our present results provide a functional analysis of T\textsubscript{EMRA} cells in bacterial infection and favor a role for T\textsubscript{EMRA} cells in protective immunity in human tuberculosis. This insight also provides a mechanism for the reactivation of tuberculosis that occurs during anti-TNF immunotherapy. Among several infectious diseases that occur during infliximab therapy, the greatest threat comes from pathogens causing granulomatous infections, most notably M. tuberculosis (40). A likely explanation is that granuloma formation depends on coordinated cellular trafficking, which is altered by anti-TNF treatment (16, 65–67). The demonstration that T\textsubscript{EMRA} cells express membrane TNF, bind anti-TNF antibodies, and subsequently become susceptible to complement-mediated lysis provides a cellular mechanism by which anti-TNF therapy influences adaptive immunity.

The expression of membrane TNF on PBMCs reflects the disease activity observed in our patients, as increased levels of TNF are a hallmark of autoimmune diseases and the rationale for the therapeutic intervention with TNF-neutralizing agents (5). The majority of TNF is present as a transmembrane protein, which needs to be cleaved by the metallocproteinase TNF–α–converting enzyme (TACE) to be released into the extracellular space. While this process is very efficient in macrophages and dendritic cells, much of the TNF produced by T cells remains in the membrane form (68, 69). Therefore, the increased expression of membrane TNF on lymphocytes from patients with RA (Figure 6) reflects the inappropriate T cell activation that is responsible for the active disease diagnosed in our patients.

Patients treated with anti-TNF antibodies (e.g., infliximab) have a 5-fold higher risk for developing tuberculosis than do patients treated with soluble TNF receptors (e.g., etanercept; ref. 40). Also, the majority of tuberculosis reactivation occurs during the first 3 months of initiating infliximab therapy, whereas etanercept-mediated disease occurs much later (40). This different impact on host defense against M. tuberculosis is explained by previous findings on the interactions of TNF-neutralizing agents with membrane TNF. Etanercept binds poorly and with low avidity to transmembrane TNF (70, 71). Accordingly, etanercept is much less efficient than infliximab in triggering complement-dependent cytotoxicity (70, 71). The present data provide a rationale for determining whether T\textsubscript{EMRA} cells are preserved during etanercept treatment.

Previous reports on anti-TNF–mediated effects on T cell responses to mycobacterial antigens showed a decreased expression of membrane TNF in stimulated CD4\textsuperscript{+} T cells (72), a variable effect on the number of IFN-γ–releasing cells (72, 73), and no changes in proliferation (72). The decrease of T\textsubscript{EMRA} cells described in the present study clearly favors the establishment of an antiinflammatory environment. While this could be a disadvantage in protection against intracellular pathogens, the decrease of T\textsubscript{EMRA} cells may contribute to the clinical success of anti-TNF therapy. For example, complement-mediated lysis of T\textsubscript{EMRA} cells may be the mechanism by which the number of perforin\textsuperscript{+} cells and elevated granyme level in fluids and tissues of RA patients (74, 75) is normalized. Taken together, our own and others’ studies (61, 76–83) strongly suggest that the correction of inappropriate T cell responses is a key mechanism by which anti-TNF therapy antagonizes the hyperinflammatory state that accounts for autoimmune diseases.

In summary, our findings demonstrated that anti-TNF therapy resulted in a significant reduction of CD8\textsuperscript{+} T\textsubscript{EMRA} cells, thereby disturbing an important component of protective immunity against intracellular bacteria. Therefore, CD8\textsuperscript{+} T\textsubscript{EMRA} cells should be considered as a target for manipulating immune responses: strengthening this compartment would be beneficial for infections, and counterregulation could improve autoimmune disease.

**Methods**

**Cell culture reagents.** Cells were cultured in complete medium, consisting of RPMI 1640 (Biochrom) supplemented with glutamine (2 mM; Sigma-Aldrich), 10 mM HEPES, 13 mM NaHCO\textsubscript{3}, 100 μg/ml streptomycin, 60 μg/ml penicillin (all from Biochrom), and 5% heat-inactivated human AB serum (Cambrex).

**Antibodies and reagents.** The following antibodies were used for flow cytometry: PE-Cy5.5–conjugated anti-CD4, FITC-conjugated anti-CD16, allophycocyanin-conjugated anti-CD19, allophycocyanin-conjugated anti-CD45RA, biotin-conjugated anti-mouse IgG1, and allophycocyanin/PE-conjugated streptavidin (all from CalTag); PerCP/allophycocyanin-conjugated anti-CD8, PE/allophycocyanin-conjugated anti-CD56, and PE-conjugated anti-perforin (all from BD Biosciences); FITC-conjugated anti-CCR7 (R&D Systems); FITC-conjugated anti-Vu24 (Immunotech); PE-conjugated anti-CD161 (Beckman Coulter); polyclonal granulysin rabbit serum (gift from A. Krensky, Stanford University, Stanford, California, USA); infliximab (Essex Pharma); etanercept (Wyeth Pharma); and Cy5-conjugated donkey anti-rabbit and biotin-conjugated goat anti-rabbit (Dianova). The following reagents were used: paraformaldehyde, saponin, sodium azide (Sigma-Aldrich), rabbit complement (BAG Healthcare), and EZ-Link NHS-Biotin Reagent (Pierce).

**Flow cytometry.** PBMCs were stained with cell surface markers for 30 minutes. Cells were then fixed with 4% paraformaldehyde (10 minutes at 4°C) and incubated with permeabilization buffer (0.5% saponin, 5% FCS, and 0.05% Tween 80; Sigma-Aldrich) and 10% Middlebrook 7H9 broth (BD Biosciences) supplemented with 1% glycerol (Roth); 0.05% Tween 80 (Sigma-Aldrich); and 10% Middlebrook oleic acid, albumin, dextrose, and catalase enrichment (BD Biosciences). Aliquots from logarithmically growing cultures were frozen in PBS containing 10% glycerol, and representative vials were thawed and enumerated for viable...
CFU on Middlebrook 7H11 plates. Staining of bacterial suspensions with fluorochromic substrates differentiating between live and dead bacteria (BacLight; Invitrogen) revealed greater than 90% viability of the bacteria. Several precautions were taken to minimize clumping of mycobacteria: (a) culture conditions (rotation, tweezing) were chosen to support the growth of single-cell suspensions; (b) before in vitro infection, M. tuberculosis bacilli were sonicated to disrupt small aggregates of bacteria; and (c) the MOI was selected such that there were only 1–3 bacilli per infected cell.

**Infection of monocytes.** Monocytes were infected with single-cell suspensions of M. tuberculosis at MOI 5 in complete medium without antibiotics and supplemented with 5% non–heat-inactivated human serum. After 3 hours of incubation at 37°C, monocytes were washed 3 times to remove extracellular bacteria. Adherent cells were detached using 1 mM EDTA. The efficacy of infection was determined in each experiment by acid fast stain (auramin rhodamine) and was 25% ± 13%.

**Cytotoxicity assay.** Uninfected and infected monocytes were labeled with 100 μCi 35S (ICN Biomedicals) for 1 hour and plated in a 96-well V-bottom plate at a final concentration of 5,000 targets/100 μl. CD8 effector cells were purified from the mononuclear fraction by negative selection using magnetic beads per the manufacturer’s recommendations (Miltenyi Biotec). Purified CD8 cells were then labeled with allopurinol-conjugated anti-CD45RA and FITC-conjugated anti-CCR7 and sorted using a cell sorter (MoFlow) as CCR7+/CD45RA− (Tcm), CCR7+/CD45RA+ (Tcm), CCR7−/CD45RA− (naive), and CCR7−/CD45RA+ (TEmra). The purity of the cell populations exceeded 97% in all experiments. On the same day, effector cells were lyzed with 0.3% saponin (Sigma-Aldrich) to release intracellular fluorochromic substrates differentiating between live and dead bacteria using magnetic beads per the manufacturer’s recommendations (Miltenyi Biotec). The purity of the cell populations exceeded 97% in all experiments.

**Target cell lysis was calculated by quantifying 51Cr release.** The spontaneous release of 51Cr in the absence of T cells was less than 10%. Results were expressed as percent specific lysis, calculated as follows: (experimental 51Cr release – spontaneous release)/(maximum release – spontaneous release).

**Quantification of mycobacterial growth.** To measure mycobacterial growth, cells were lysed with 0.3% saponin (Sigma-Aldrich) to release intracellular bacteria. At all time points, an aliquot of unlysed infected cells was harvested and counted. This allowed exact quantification of cells as well as determination of cellular viability by trypan blue exclusion. Recovery of cells was greater than 80% in all experiments, with cell viability regularly exceeding 90% of total cells. Lysates of infected cells were resuspended vigorously, transferred into screw-cap tubes, and sonicated in a preheated water bath sonicator for 5 minutes at 37°C.

**Patients.** Patients were recruited from 2002 to 2008 in the outpatient departments of the Institute for Clinical Immunology at the University of Ulm, Germany, and the European Union (TB-VAC). The expert associate of Martin Röllinghoff and Jochen K. Kalden throughout the study.

Received for publication January 5, 2009, and accepted in revised form March 25, 2009.

Address correspondence to: Steffen Stenger, Universitätsklinikum Ulm, Institut für Medizinische Mikrobiologie und Hygiene, Albert Einstein Allee 11, 89081 Ulm, Germany. Phone: 49-500-65300; Fax: 49-500-65302; E-mail: steffen.stenger@uniklinik-ulm.de.

9. Bean, A.G., et al. 1999. Structural deficiencies in granuloma formation in TNF gene-targeted mice underlie the heightened susceptibility to aerosol Mycobacterium tuberculosis infection, which is


