Differential Effects of Anti-Fas Ligand and Anti-Tumor Necrosis Factor α Antibodies on Acute Graft-Versus-Host Disease Pathologies

By Koichi Hattori, Takao Hirano, Hiroaki Miyajima, Norifumi Yamakawa, Masatoshi Tateno, Kazuo Oshimi, Nobuhiko Kayagaki, Hideo Yagita, and Ko Okumura

Both tumor necrosis factor α (TNFα) and Fas ligand (FasL) have been implicated in the pathogenesis of graft-versus-host disease (GVHD). In this study, we examined the ameliorating effects of neutralizing anti-FasL and/or anti-TNFα monoclonal antibody (MoAb) in a lethal acute GVHD model in mice. Whereas the treatment with either anti-FasL or anti-TNFα MoAb alone significantly delayed the mortality and improved the body weight, a complete protection was achieved by the administration of both MoAbs. Pathological examination indicated differential effects of anti-FasL or anti-TNFα MoAb on GVHD-associated pathologies. Hepatic lesion was improved by anti-FasL but not anti-TNFα MoAb. In contrast, intestinal lesion was improved by anti-TNFα but not anti-FasL MoAb. Cutaneous and splenic lesions were improved by either MoAb. The combination of both MoAbs improved all these lesions. These results indicate that FasL and TNFα differentially contribute to the GVHD pathologies and a complete protection from mortality can be achieved by neutralization of both FasL and TNFα.

© 1998 by The American Society of Hematology.
spleens, and bone marrow were subjected to histopathological examination.

**Histopathology.** Tissues were fixed in 10% buffered formalin and paraffin-embedded. Sections were stained with hematoxylin and eosin and examined under microscopy.

**Flow cytometric analysis.** Splenocytes were prepared from normal CBF1, GVHD, or MoAb-treated mice on day 21 and stained with fluorescein isothiocyanate (FITC)-conjugated anti-H-2K\(^d\) (AF6-88.5; PharMingen), and phycoerythrin (PE)-conjugated anti-CD4 (RM4-5; PharMingen), anti-CD8 (53-6.7; PharMingen), or anti-B220 (RA3-6B2; PharMingen) MoAbs followed by APC-conjugated avidin (PharMingen). Cells (1 \(\times\) 10\(^4\)) were analyzed on FACS Vantage and analyzed by Cell Quest program (Becton Dickinson, San Jose, CA).Recipient and donor lymphocytes were identified as H-2K\(^d\)\(^+\)K\(^b\)\(^+\) and H-2K\(^d\)\(^-\)K\(^b\)\(^+\) cells, respectively. Cell numbers of CD4\(^+\) T, CD8\(^+\) T, and B220\(^+\) B cells of recipient or donor origin were calculated from the total numbers of splenocytes recovered, and the percentages of each subpopulation were determined by the three-color analysis.

**Statistical analysis.** Significant differences between experimental groups were determined using the Mann-Whitney U test for the survival rate or using the Student’s t-test for the body weight. P values less than .05 were considered statistically significant.

**RESULTS**

**Effect of anti-FasL and/or anti-TNF\(\alpha\) MoAb on GVHD-induced mortality and weight loss.** A lethal acute type of GVHD was induced by IV injection of B6 splenocytes into CBF1 mice. As represented in Fig 1A, all the recipients administrated with control IgG died within 21 days. In these mice, clinical symptoms of acute GVHD, such as hair ruffling, lesser mobility, and weight loss (Fig 1B), became apparent within 2 weeks. Administration of either anti-FasL MoAb or anti-TNF\(\alpha\) MoAb alone significantly delayed but not completely reduced the mortality, and 1 of 10 or 4 of 10 mice survived at day 60, respectively (Fig 1A). In these surviving mice, no significant weight loss was observed as compared with the age-matched normal mice until day 14, but their growth was retarded after the discontinuation of the treatment at day 14 (Fig 1B) with clinical symptoms of GVHD.

In contrast, all the recipient mice treated with both anti-FasL and anti-TNF\(\alpha\) MoAbs survived over 60 days (Fig 1A). Even after the discontinuation of the treatment at day 14, no apparent clinical symptoms of acute GVHD were observed in these mice and their growth was comparable to that of the age-matched normal mice (Fig 1B).

**Effect on GVHD-associated histopathologies.** In the liver from the control mice undergoing GVHD, a massive infiltration of mononuclear cells and fibrosis were observed mainly in the periportal areas (Fig 2A). A similar hepatic pathology was observed in the liver from the anti-TNF\(\alpha\)-treated mice (Fig 2B). In contrast, such inflammatory changes were minimal in the liver from the anti-FasL–treated mice (Fig 2C). The gut from the control mice undergoing GVHD exhibited a dilatation, flattening of the villi, and elevation and atrophy of the crypts, which are characteristics of intestinal GVHD (Fig 2F). Similar changes were observed in the gut of anti-FasL–treated mice, although structural integrity of the villi was partially improved as compared with the GVHD control (Fig 2H). In contrast, all these lesions were almost absent in the anti-TNF\(\alpha\)–treated mice (Fig 2G).
The skin from the control mice undergoing GVHD exhibited severe inflammatory infiltrates with intraepidermal lymphocytes and dyskeratotic cells, ulceration, loss of hair follicles, and destruction of rete ridges (Fig 2K). Such changes were not observed in either anti-FasL–treated (Fig 2M) or anti-TNFα–treated (Fig 2L) mice.

The spleen from the control GVHD mice showed a marked lymphoid atrophy, structural disorganization, and focal necrosis (Fig 2P). Such changes were minimal in either anti-FasL–treated (Fig 2R) or anti-TNFα–treated (Fig 2Q) mice. In the recipients treated with both anti-FasL and anti-TNFα MoAbs, no apparent lesion was observed in the liver (Fig 2D), intestine (Fig 2I), skin (Fig 2N), or spleen (Fig 2S) as compared with normal mice (Fig 2E, J, O, and T).

Effect on GVHD-associated lymphoid hypoplasia. Cell numbers of CD4<sup>T</sup>, CD8<sup>T</sup>, and B220<sup>B</sup> cells of recipient (H-2K<sup>d</sup><sup>−</sup>K<sup>b</sup><sup>−</sup>) or donor (H-2K<sup>d</sup><sup>+</sup>K<sup>b</sup><sup>+</sup>) origin in the splenocytes from normal CBF1, GVHD, or MoAb-treated mice on day 21 were calculated from the total numbers of recovered, and the percentages of each subpopulation were determined using three-color flow cytometric analysis (Table 1). In the splenocytes from GVHD mice, both CD4<sup>+</sup> and CD8<sup>+</sup> T cells and B220<sup>+</sup> B cells of host origin were severely decreased as compared with normal CBF1 mice. The treatment with either anti-FasL or anti-TNFα MoAb alone partially but substantially prevented the loss of all these lymphocyte subpopulations, and almost complete protection was achieved by the treatment with both MoAbs. It was also noted that donor-derived CD4<sup>+</sup> and CD8<sup>+</sup> T cells were increased in the anti-FasL– and/or anti-TNFα–treated mice as compared with the GVHD mice, representing a chimeric state of these recipients. As represented in Table 1, 60% to 67% of T cells and 47% to 58% of B cells were donor origin. This

**Table 1. Effect of Anti-FasL and/or Anti-TNFα Antibodies on GVHD-Associated Lymphoid Hypoplasia**

<table>
<thead>
<tr>
<th>Mice</th>
<th>Cell No. (&lt;10&lt;sup&gt;7&lt;/sup&gt;)</th>
<th>CD4&lt;sup&gt;+&lt;/sup&gt;</th>
<th>CD8&lt;sup&gt;+&lt;/sup&gt;</th>
<th>B220&lt;sup&gt;+&lt;/sup&gt;</th>
<th>CD4&lt;sup&gt;+&lt;/sup&gt;</th>
<th>CD8&lt;sup&gt;+&lt;/sup&gt;</th>
<th>B220&lt;sup&gt;+&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td>2.1 ± 0.11</td>
<td>1.5 ± 0.57</td>
<td>2.8 ± 0.80</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>GVHD</td>
<td>0.2 ± 0.02</td>
<td>0.2 ± 0.02</td>
<td>0.7 ± 0.06</td>
<td>0.4 ± 0.06</td>
<td>0.3 ± 0.07</td>
<td>1.3 ± 0.60</td>
<td></td>
</tr>
<tr>
<td>Anti-FasL</td>
<td>1.2 ± 0.05</td>
<td>1.7 ± 0.06</td>
<td>1.8 ± 0.33</td>
<td>2.2 ± 0.70</td>
<td>2.5 ± 0.84</td>
<td>1.6 ± 0.52</td>
<td></td>
</tr>
<tr>
<td>Anti-TNFα</td>
<td>1.2 ± 0.46</td>
<td>1.1 ± 0.29</td>
<td>1.4 ± 0.02</td>
<td>2.4 ± 0.30</td>
<td>2.0 ± 0.30</td>
<td>1.9 ± 0.10</td>
<td></td>
</tr>
<tr>
<td>Anti-FasL + anti-TNFα</td>
<td>1.6 ± 0.20</td>
<td>1.9 ± 0.45</td>
<td>2.0 ± 0.65</td>
<td>3.3 ± 0.27</td>
<td>3.3 ± 0.39</td>
<td>2.8 ± 0.90</td>
<td></td>
</tr>
</tbody>
</table>

Cell numbers of CD4<sup>+</sup> T, CD8<sup>+</sup> T, and B220<sup>+</sup> B cells of host (H-2K<sup>d</sup><sup>−</sup>K<sup>b</sup><sup>−</sup>) or donor (H-2K<sup>d</sup><sup>+</sup>K<sup>b</sup><sup>+</sup>) origin in the splenocytes from normal CBF1, GVHD, or anti-FasL antibody-treated and/or anti-TNFα antibody-treated mice on day 21 were calculated from the total numbers of splenocytes recovered, and the percentages of each subpopulation were determined using three-color flow cytometric analysis. Data represent the mean ± SD of 5 mice in each group.
chimeric state appeared to be stable, because no further change in the numbers of host and donor lymphocytes was observed on day 60 in the recipients treated with both anti-FasL and anti-TNFα MoAbs (not shown).

**DISCUSSION**

In this study, we explored the ameliorating effects of neutralizing MoAbs against FasL and TNFα, both of which have been implicated in the pathogenesis of GVHD, in a murine model of lethal acute GVHD. Whereas the treatment with either anti-FasL or anti-TNFα MoAb alone significantly delayed the mortality and improved the weight loss, a complete protection was achieved by the combination of both MoAbs. Histological examination indicated differential effects of these MoAbs on GVHD-associated pathologies.

Recent studies have implied that FasL plays a critical role in the development of hepatic and cutaneous lesions and lymphoid atrophy. Baker et al. showed that, when the FasL-deficient gld mice were used as the T-cell donor in a major histocompatibility complex (MHC)-matched but minor-mismatched allogenic BMT model of acute GVHD, only minimal signs of hepatic and cutaneous GVHD pathology were observed and the lymphoid atrophy in the spleen was improved. However, intestinal GVHD was not abrogated and neither weight loss nor mortality was improved. In contrast, Braun et al. reported a significantly delayed mortality in the recipients of Fasl-defective T cells in a MHC-mismatched spleen cell transfer model. We used the parent (B6) to F1 (CBF1) spleen cell transfer model and found that the treatment with anti-FasL MoAb delayed the mortality and improved the weight loss, consistent with the observation by Braun et al. The apparent discrepancy from the observation by Baker et al. in the FasL contribution to mortality remains to be resolved by testing the effect of anti-FasL MoAb in the BMT model. Our histological observations are consistent with those by Baker et al., indicating a critical contribution of FasL to the development of hepatic and cutaneous, but not intestinal, lesions and splenic atrophy.

The ameliorating effect of anti-TNFα treatment observed in this study was essentially consistent with that reported by Piguet et al. They described that the administration of an anti-TNFα polyclonal antibody reduced the mortality at day 40 by 50% and abolished the weight loss on day 18. In our present study, the administration of an anti-TNFα MoAb similarly reduced the mortality and abolished the weight loss. They also showed that the GVHD-associated pathologies in the skin and gut, but not those in the liver, were prevented by the anti-TNFα treatment. These observations are also consistent with ours, indicating a critical contribution of TNFα to the development of intestinal and cutaneous, but not hepatic, lesions.

When both anti-FasL and anti-TNFα MoAbs were administered in combination, all of these histological lesions in the liver, intestine, skin, and spleen were minimal. It was notable that all the recipients survived more than 60 days and grew well as normal mice without growth retardation observed in the recipients treated with either anti-TNFα or anti-FasL MoAb alone, which may result from hepatic or intestinal damage, respectively. These results verified that FasL and TNFα differentially contribute to the GVHD pathologies as follows: (1) hepatic GVHD is predominantly mediated by FasL; (2) intestinal GVHD is predominantly mediated by TNFα; and (3) cutaneous GVHD, splenic atrophy, weight loss, and mortality are mediated by both FasL and TNFα. Importantly, FasL and TNFα in combination appear to mostly account for all these GVHD-associated pathologies observed in the present study.

In the histological examination, the treatment with either anti-FasL or anti-TNFα MoAb improved the splenic atrophy. Flow cytometric analysis for the lymphocyte subpopulations indicated that the GVHD-associated elimination of host lymphocytes (both T and B cells) was prevented partially by either MoAb alone and almost completely by the combination of both MoAbs. This suggests that both TNFα and FasL contribute to cytotoxic elimination of host lymphocytes by host-reactive donor T cells. Alternatively, this may be also due to blocking of suppressive effects of TNFα and FasL on hematopoiesis (our unpublished data). It was also noted that donor-derived T cells were increased in the anti-FasL- and/or anti-TNFα-treated mice as compared with the GVHD mice. This may result from inhibition of activation-induced apoptosis, in which both FasL and TNFα have been implicated. The preservation of both host and donor lymphocytes represents a chimeric state of the recipients, which appears to be stable over 60 days in the anti-FasL/TNFα-treated mice. It remains to be determined whether a tolerance to the host alloantigen has been established in the donor T cells.

In conclusion, a complete protection was achieved by administration of both anti-FasL and anti-TNFα MoAbs in a murine model of lethal acute GVHD. Although our observations were made in a parent into F1 model in which no cytotoxic conditioning was used before the transplant and thus cannot necessarily be directly extrapolated to allogeneic transplantation as performed clinically today, our present findings may provide insights that would be useful for the treatment of GVHD. The phase I-II clinical trials with a humanized anti-TNFα MoAb for the treatment of refractory acute GVHD have resulted in limited success. We recently succeeded to generate a humanized version of anti-human FasL MoAb (manuscript in preparation), which may be useful for the clinical treatment of severe acute GVHD patients in combination with the anti-TNFα MoAb.

**ACKNOWLEDGMENT**

The authors thank C. Ushiyama for technical assistance and helpful suggestions.

**REFERENCES**

21. Sytwu HK, Liblau RS, McDevitt HO: The roles of Fas/APO-1 (CD95) and TNF in antigen-induced programmed cell death in T cell receptor transgenic mice. Immunity 5:17, 1996
Differential Effects of Anti-Fas Ligand and Anti-Tumor Necrosis Factor α Antibodies on Acute Graft-Versus-Host Disease Pathologies

Koichi Hattori, Takao Hirano, Hiroaki Miyajima, Norifumi Yamakawa, Masatoshi Tateno, Kazuo Oshimi, Nobuhiko Kayagaki, Hideo Yagita and Ko Okumura