

Modulation of Chemokine Production and Inflammatory Responses in Interferon- γ - and Tumor Necrosis Factor-R1-Deficient Mice during *Trypanosoma cruzi* Infection

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Infection with *Trypanosoma cruzi* causes a strong inflammatory reaction at the inoculation site and, later, in the myocardium. The present study investigates the role of cytokines as modulators of *T. cruzi*-induced chemokine expression *in vivo* and *in vitro*. In macrophage cultures, although the stimulation with interferon (IFN)- γ increases the expression of IP-10, it blocks KC expression. Tumor necrosis factor (TNF)- α , on the other hand, potentiates KC, IP-10, macrophage inflammatory protein-1 α , and JE/monocyte chemoattractant protein-1 expression. Interleukin-10 and transforming growth factor- β inhibited almost all chemokines tested. The role of IFN- γ and TNF- α in chemokine modulation during infection was investigated in *T. cruzi*-infected IFN- γ -deficient (GKO) or TNF-R1/p55-deficient (p55-/-) mice. The expression of chemokines detected in the inoculation site correlated with the infiltrating cell type observed. Although GKO mice had a delayed and intense neutrophilic infiltrate correlating with the expression of KC and macrophage inflammatory protein-2, none of the above was observed in p55-/- mice. The detection of infiltrating T cells, Mig, and IP-10 in the myocardium was observed in wild-type and p55-/-, but not in GKO mice. Together, these results suggest that the regulatory roles of IFN- γ and TNF- α on chemokine expression may play a crucial role in the modulation of the inflammatory response during *T. cruzi* infection and mediate resistance to infection. (*Am J Pathol* 2001, 158:1433-1440)

Trypanosoma cruzi is a hemoflagellate protozoan parasite that is the etiological agent of Chagas' disease, a major public health problem in South and Central America. The infection is primarily characterized by an acute phase with detection of circulating parasites and an impressive inflammatory reaction at the site of parasite penetration. A few days later, significant myocarditis associated with tissue parasitism are noted and these are the most relevant pathological findings during infection with *T. cruzi*.

The resistance of inbred mice to acute infection with *T. cruzi* has been shown to be dependent on interferon (IFN)- γ that activates macrophages to produce nitric oxide (NO) and kill the obligate intracellular amastigote form of the parasite.¹⁻⁶ In addition, tumor necrosis factor (TNF)- α provides a second signal stimulating NO production and anti-*T. cruzi* activity in IFN- γ -activated macrophages. Because *T. cruzi*-infected macrophages produce TNF- α , this cytokine seems to mediate its trypanocidal function via an autocrine pathway.⁷

On the other hand, the down-regulatory cytokines interleukin (IL)-10 and transforming growth factor (TGF)- β are associated with susceptibility to infection⁸⁻⁹ by inhibiting IFN- γ -mediated macrophage activation. Thus, neutralization of endogenous IL-10 leads to an increased *T. cruzi*-induced IFN- γ production and parasite killing.^{9,10} These results suggest that IL-10 may be a potent inhibitor of IFN- γ production during *T. cruzi* infection in mice and that the early resistance to infection is a result of the balance between IFN- γ and IL-10 produced.⁹

The mechanisms by which cytokines control parasite replication and myocarditis in *T. cruzi*-infected mice are still a matter of controversy. One distinct possibility is that cytokines modulate the expression of chemokines that in turn drive the inflammatory infiltrates observed during the acute and chronic phases of infection. In fact, *T. cruzi* infection induces the production of inflammatory and regulatory cytokines,^{7,8,11-13} and the β -chemokines macro-

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phage inflammatory protein (MIP)-1 α , MIP-1 β , RANTES, and JE in macrophages¹⁴ or in cardiomyocytes.¹⁵ The importance of the production of these chemokines to disease outcome and host immunopathology during infection is not known.

In the present study, we evaluated the role of IL-10, TGF- β , IFN- γ , and TNF- α in regulating chemokine production and expression as well as leukocyte influx. We found that IFN- γ , TNF- α , IL-10, and TGF- α are indeed able to modulate the expression of various chemokines in *T. cruzi*-infected macrophages. Moreover, we found that the decreased inflammatory infiltrate in the peritoneum and heart of IFN- γ -deficient (GKO) and in TNF-R1-deficient (p55-/-) mice correlated with the altered expression of chemokines. These results suggest an important role for regulatory cytokines, especially IFN- γ and TNF- α , in the control of chemokine expression and production in experimental *T. cruzi* infection.

Materials and Methods

Experimental Animals

Five- to 6-week-old female C3H/HeJ, C57BL/6 (WT), or IFN- γ -deficient (GKO) or TNF- α receptor p55 (p55-/-) or p75 (p75-/-)-deficient mice were bred and maintained under standard conditions in the animal house of the Department of Immunology, University of São Paulo, Ribeirão Preto, Brazil. Deficient mice were obtained from The Jackson Laboratories (Bar Harbor, ME).

Parasites

The Y strain of *T. cruzi* was used in all experiments. For *in vitro* experiments, trypomastigotes were grown and purified from a monkey fibroblast cell line (LLC-MK₂). For *in vivo* experiments, bloodstream trypomastigotes were obtained from infected animals. The levels of parasitemia were evaluated in 5 μ l of blood drawn from the tail vein.

Differential Counting of Leukocytes and Detection of Chemokine mRNA Expression in Vivo

Mice were infected intraperitoneally with 1×10^4 bloodstream trypomastigotes. At different times after infection peritoneal cavities were washed with cold phosphate-buffered saline (PBS). Total peritoneal host and parasite cells were counted in a hemocytometer; 1×10^4 peritoneal cells were cytopun to slides and stained with May-Grünwald-Giemsa; leukocyte populations were determined after differential counting using standard morphological criteria.

FACS Analysis

For flow cytometry, 1×10^8 cells were incubated in the presence of 5 μ l of the fluorescein isothiocyanate-conjugated antibodies anti-CD3, anti-CD4, anti-CD8, and anti-

CD22 (Pharmingen, San Diego, CA) for 30 minutes at 4°C in the dark according to the supplier's specifications. The cells were then washed and resuspended in PBS-1% formaldehyde. Data acquisition was performed using a FACSsorter, (Becton-Dickinson Immunocytometry System Inc., San Jose, CA). The percentage of each subset was used together with the total cell counting described above to calculate total numbers of each lymphocyte subset in the peritoneal exudate.

Macrophage Cultures for RNA Extraction

BALB/c mouse inflammatory macrophages were harvested from peritoneal cavities 3 days after injection of 1 ml of 3% sodium thioglycollate. The cells were washed and suspended to 1×10^6 cells/ml in RPMI 1640 (Sigma), supplemented with 5% fetal bovine serum, 5×10^{-5} mol/L 2-mercaptoethanol, 2 mmol/L L-glutamine, and antibiotics. The adherent cells were obtained after 2 to 4 hours incubation of single-cell suspensions in 24-well tissue-culture plates at 37°C. Nonadherent cells were removed and trypomastigote forms were added at 1:1 parasite:cell ratio to macrophages and incubated for 6 hours at 37°C in a humidified chamber containing 5% CO₂. The cells were then washed three times and 0.5 ml of Trizol LS reagent was added to each well, incubated at room temperature for 5 minutes, and stored at -70°C until RNA extraction.

Total RNA Extraction and cDNA Preparation by Reverse Transcription

For reverse transcriptase-polymerase chain reaction (RT-PCR) analysis, cells were collected by centrifugation of the peritoneal lavage fluid and RNA was prepared from the cell pellet. The extraction of total RNA was performed using 1 ml of the Trizol LS reagent according to the manufacturer's recommendations (Life Technologies, Inc., Gaithersburg, MD). Each RNA sample was resuspended in water at 0.5 μ g/ μ l. cDNA was synthesized from 2 μ l of sample using Superscript II reverse transcriptase according to the supplier's specifications (Life Technologies, Inc.).

Chemokine mRNA Detection

mRNAs for the C-X-C chemokines (Mig, IP-10, KC, MIP-2), the CC chemokines (MIP-1 α , RANTES, JE/MCP-1), hydroxyphosphoribosyltransferase (HPRT), and β -actin were analyzed by RT-PCR. PCRs were performed using Taq polymerase (Life Technologies, Inc.) in a PTC-100 thermal cycler (MJ Research, Watertown, MA). The primer sequences and PCR product sizes for the genes above have been previously published^{14,16} and validate in the laboratory using plasmid containing the genes for each chemokine. Reaction conditions were 30 cycles of 1 minute at 94°C, 1 minute at 54°C, and 2 minutes at 72°C, with a final extension step of 7 minutes at 72°C. For each set of primers a negative sample (water) was run in

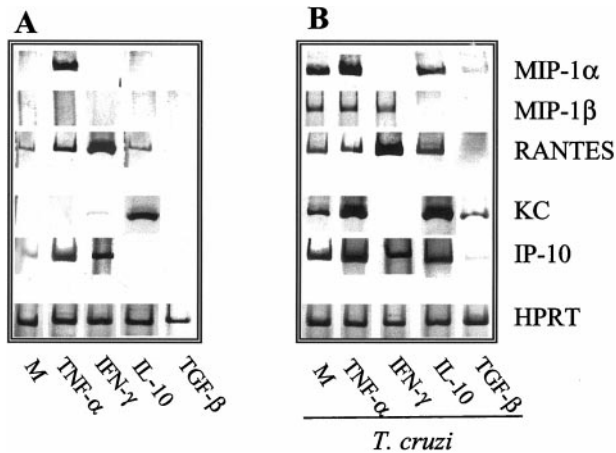


Figure 1. The cytokines IFN- γ , TNF- α , IL-10, and TGF- β regulate *T. cruzi*-induced chemokine mRNA expression by macrophages. Control (A) or *T. cruzi*-infected (B) C3H/HeJ peritoneal macrophages (1:1 parasite:host cell ratio) were pretreated with buffer (M) or IFN- γ , TNF- α , IL-10 (all at 100 U/ml), or TGF- β (50 ng/ml). At 6 hours, total RNA was extracted and mRNA expression was evaluated by RT-PCR. The amplification products were separated by electrophoresis in an acrylamide gel and silver stained. These data are representative of three independent experiments.

parallel. PCR products were separated by acrylamide gel electrophoresis and stained with silver nitrate.

Immunofluorescence

Cryostat sections of heart tissue harvested during the acute phase of the infection were air-dried for 1 hour and fixed in acetone before immunostaining with rabbit anti-mouse IP-10, anti-mouse Mig, or rabbit Ig (control). Sections were then incubated with a biotin goat anti-rabbit IgG (Santa Cruz Biotechnology, Santa Cruz, CA) followed by a 30-minute incubation with streptavidin-APC (Pharmingen, San Diego, CA). For three-color analysis, infiltrating leukocytes were immunostained using the primary antibodies rat anti-mouse CD4-fluorescein isothiocyanate and anti-mouse CD8-PE (Pharmingen) or the respective isotype controls. Nonspecific binding was blocked with PBS-bovine serum albumin 1%. The slides were analyzed using a fluorescent microscope (Leica, Wetzlar, Germany) and the images processed using SlideBook software (Intelligent Imaging Innovations, Inc., Denver, CO).

Results

*IFN- γ , TNF- α , IL-10, and TGF- β Regulate the Expression of Chemokine mRNAs by *T. cruzi*-Infected Macrophages*

Noninfected control macrophages only express little message for the chemokine RANTES (Figure 1A). Addition of TNF- α or IFN- γ increased message for RANTES and IP-10 and TNF- α also induced MIP-1 α message (Figure 1A). IL-10 induced significant message for KC but none for the other chemokines, whereas TGF- β only blocked RANTES expression (Figure 1A). The infection of macrophages with *T. cruzi* induced significant message

for MIP-1 α , MIP-1 β , KC, and IP-10, and enhanced the message for RANTES (Figure 1B). To test the effects of modulatory cytokines on the regulation of chemokines *in vitro*, macrophages were infected with *T. cruzi* together with modulatory cytokines. The addition of TNF- α enhanced the expression of MIP-1 α , KC, and IP-10 and had no apparent effect on RANTES or MIP-1 β expression (Figure 1B). The addition of IFN- γ to infected macrophages down-modulated KC and MIP-1 α and enhanced RANTES and IP-10 expression (Figure 1B). Similarly to its effects on control macrophages, IL-10 increased KC expression in infected cells (Figure 1B). Interestingly, TGF- β decreased the expression of all chemokines tested in these experiments (Figure 1B).

*Disregulated Inflammatory Cell Infiltrates at the Inoculation Site of *T. cruzi* in IFN- γ - and TNF-R1-Deficient Mice*

To evaluate the contribution of IFN- γ and TNF-R p55 and p75 in the response against *T. cruzi* infections, GKO, p55-/- and p75-/- mice were infected intraperitoneally with trypomastigote forms of the Y strain and parasitemia and mortality evaluated throughout the acute phase. As shown in Figure 2A, GKO presented an uncontrolled parasitemia that kept growing until mortality began. Similarly, p55-/- mice had a significantly higher parasitemia than either to wild-type (WT) or p75-/- mice. The survival rates were found to be consistent with the levels of parasitemia observed (Figure 2B), ie, both GKO and p55-/- mice had an early mortality, which was not observed in WT and p75-/- mice. The above data suggest the critical role of IFN- γ and p55 in controlling parasite replication *in vivo*. Once these genes seemed to play an important role for resistance against *T. cruzi*, we questioned whether IFN- γ and TNF-R1 could also play a role in the modulation of the inflammatory cell infiltrates *in vivo*. To achieve this goal the composition of inflammatory infiltrates in the site of inoculation was evaluated in WT, GKO, and p55-/- mice infected with trypomastigote forms of *T. cruzi*.

Kinetic experiments showed that neutrophils peaked early (24 hours) whereas lymphocytes peaked much later (day 8) after *T. cruzi* infection in WT mice (Figure 3, top, middle, and bottom). Two peaks of macrophage infiltration were noted on days 1 and 8. The neutrophil, macrophage, and lymphocyte infiltrate observed in infected GKO mice was late, peaking 9 days after infection, and it was more intense than that observed in WT animals (Figure 3, top, middle, and bottom). No neutrophil or macrophage infiltration was noted in p55-/- mice throughout the observation period. On the other hand, in p55-/- mice lymphocytes were raised as early as 2 days after infection and were persistently elevated throughout the observation period (Figure 3, bottom). The parasite counting at the inoculation site (peritoneal cavity, Table 1) showed a consistent correlation with the parasitemia levels (Figure 2A) and both GKO and p55-/- mice exhibited a significantly higher parasite replication than WT mice.

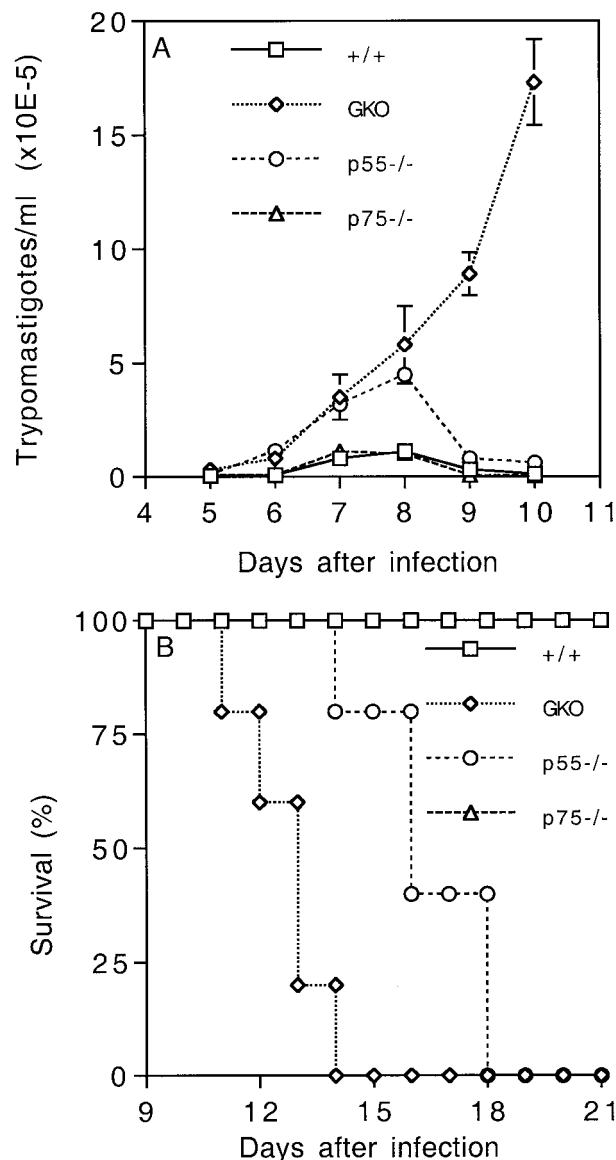


Figure 2. Resistance to *T. cruzi* infection is abolished in GKO and p55^{-/-}, but not in p75^{-/-} mice. WT (open square), GKO (open diamond), p55^{-/-} (open circle), and p75^{-/-} (open triangle) mice were infected intraperitoneally with 1×10^4 trypomastigote forms of *T. cruzi* (Y strain) and parasitemia and mortality evaluated at different days after infection. Results are shown as the mean \pm SEM and the data are representative of two independent experiments ($n = 10$ mice per group).

Immunophenotyping of Infiltrating Lymphocytes at the Site of *T. cruzi* Infection in IFN- γ - and TNF-R1-Deficient Mice

The changes in lymphocyte subtypes at the site of inoculation during the course of *T. cruzi* infection in WT animals were mild with a tendency toward a decrease of CD22⁺ and increase of CD3⁺, CD4⁺, and CD8⁺ lymphocyte numbers in days 5 to 8 after infection. Overall, the lymphocyte subtypes in infected GKO mice were similar to those found in WT animals with a small but significant increase in CD22⁺ cells at day 8 after infection. The major variations of lymphocyte subtype infiltration were observed in p55^{-/-} mice. In these animals, there was an

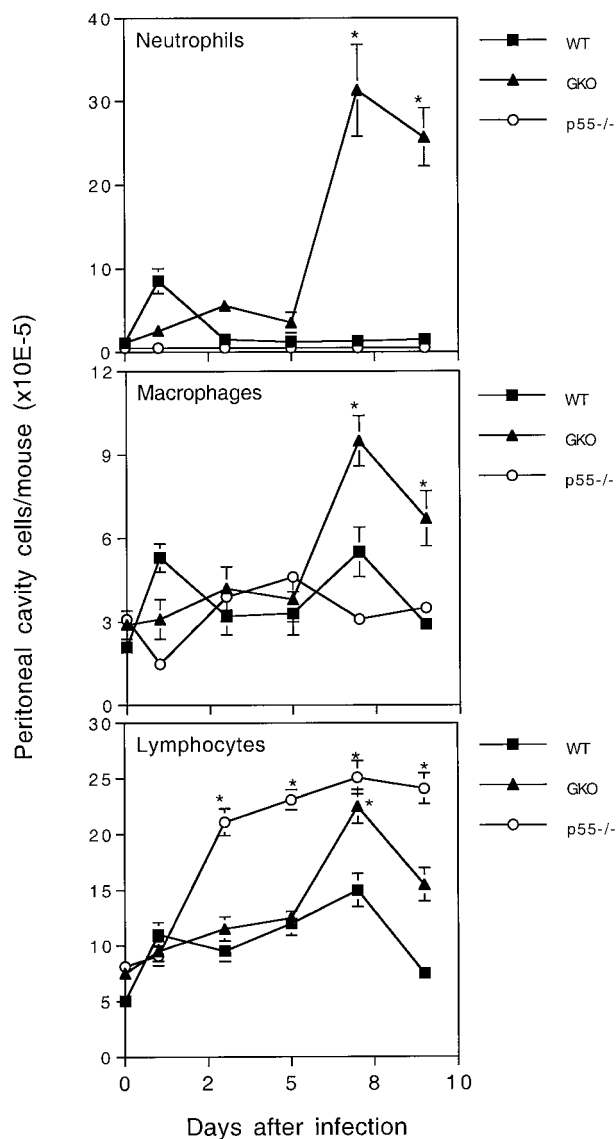


Figure 3. Disregulated cellular infiltration at the site of inoculation with *T. cruzi* in GKO or p55^{-/-} mice. WT, GKO, and p55^{-/-} mice were infected intraperitoneally with 1×10^4 trypomastigote forms of *T. cruzi* (Y strain). After different days of infection, mice were sacrificed and the number of peritoneal leukocytes evaluated. Results are means \pm SD of triplicate counts from three mice per group and are representative of six independent experiments.

increase in the numbers of lymphocyte subtypes from day 1 to 8 of the infection, with CD3⁺, CD4⁺, and CD8⁺ lymphocytes peaking at day 8 and CD22⁺ lymphocytes peaking earlier after infection (Figure 4).

Chemokine mRNA Expression at the Site of Inoculation during *T. cruzi* Infection in IFN- γ and TNF-R1-Deficient Mice

The kinetics of *in vivo* mRNA expression of chemokines was evaluated in peritoneal exudate cells and is shown in Figure 5. In WT mice, there was significant expression of KC mRNA that was limited to the early days after infection and was not accompanied by the expression of MIP-2. In

Table 1. Number of Parasites in the Peritoneal Cavity of *T. cruzi*-Infected Mice

Strain*	Number of parasites per peritoneal cavity			
	Day 3 [†]	Day 5	Day 7	Day 9
WT	7.5 ± 0.5 [‡]	15 ± 2	27.5 ± 3.5	8.3 ± 1.2
GKO	85 ± 27 [§]	220 ± 12 [§]	288 ± 22 [§]	369 ± 11.2 [§]
p55 ^{-/-}	38 ± 6	100 ± 32 [§]	74 ± 10 [§]	55 ± 5.5 [§]

*n = 3 mice per group per time point in triplicate experiments.

[†]Days after infection.

[‡]Mean ± SD (1 × 10⁴) trypomastigote forms per peritoneal cavity lavage.

[§]P < 0.05 when compared to WT mice.

GKO mice, both KC and MIP-2 mRNAs were expressed persistently in higher levels. No expression of KC or MIP-2 was observed in p55^{-/-} mice. Significant levels of IP-10, Mig, and RANTES were detected in WT animals from day 3 after infection. In GKO animals no message for these chemokines was detected, whereas in p55^{-/-} mice the chemokine message was detected in higher levels after 5 days of infection. Interestingly, only low levels of MIP1- α were detected in WT or p55^{-/-} mice whereas mRNA for this chemokine was persistently elevated in infected GKO mice.

Detection of the C-X-C Chemokines Mig and IP-10 in Myocardium of *T. cruzi*-Infected Mice

The modulation of subpopulations of infiltrating lymphocytes and the expression of chemokines in the myocar-

dium of GKO and p55^{-/-} mice were evaluated by immunofluorescence. WT animals showed increased numbers of CD4⁺ and CD8⁺ cells in the myocardium on the ninth day after infection (Figure 6, a and b). In contrast, GKO mice did not show significant increases in the numbers of CD4⁺ or CD8⁺ cells (data not shown). On the other hand and in contrast to the increased number of lymphocytes at the site of inoculation, p55^{-/-} mice showed a trend toward decreasing numbers of CD4⁺ or CD8⁺ cells when compared to WT animals, at least until the ninth day of the infection (Figure 6, c and d). Interestingly, GKO mice showed an increased number of F4/80⁺ cells on the ninth day after infection, which was not observed in WT mice (data not shown). To establish a correlation between the modulation of lymphocyte migration and chemokine expression, serial sections of heart tissue from infected mice were immunostained for the chemokines IP-10 and Mig, well-known lymphocyte chemoattractants. Heart sections of WT animals (Figure 6, a and b) were found to be strongly positive for IP-10 and Mig on the ninth day after infection,

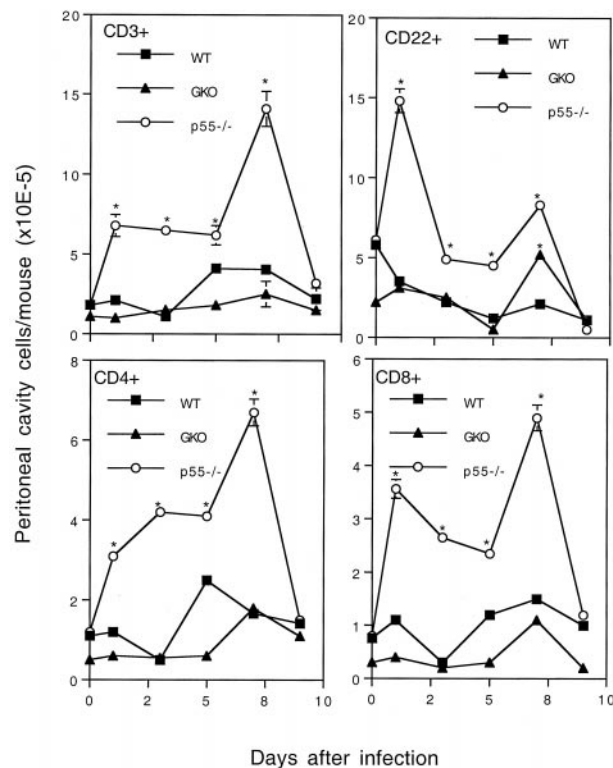


Figure 4. Phenotyping of lymphocytes in peritoneal exudates. Leukocytes obtained from the peritoneum of infected WT, GKO, or p55^{-/-} mice were immunostained with fluorescein isothiocyanate-conjugated antibodies to mouse CD3, CD4, CD8, and CD22 and analyzed by flow cytometry. Results are the means ± SD of the number of each leukocyte subset from three mice per group and are representative of three independent experiments.

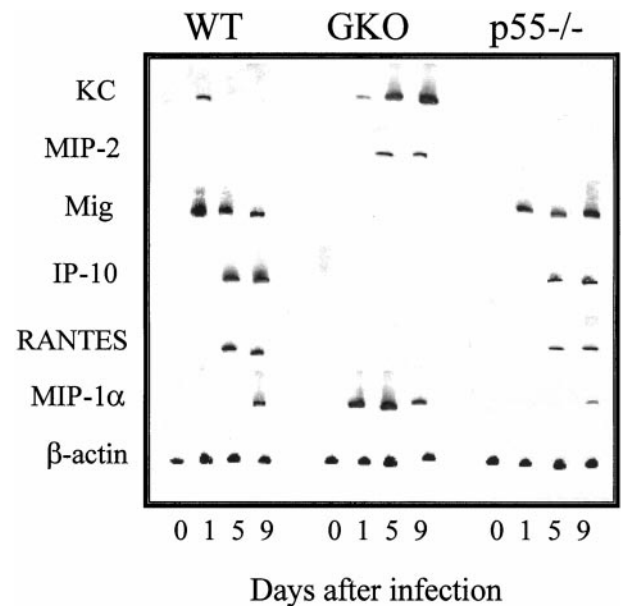


Figure 5. Chemokine mRNA expression at the site of inoculation of *T. cruzi* in IFN- γ or p55-deficient mice. WT, GKO, and p55^{-/-} mice were infected intraperitoneally with 1 × 10⁴ trypomastigote forms of *T. cruzi* (Y strain). Total RNA was extracted from leukocytes infiltrating the peritoneum at different days after infection and the expression of chemokine mRNAs was evaluated using RT-PCR. The amplification products were separated by electrophoresis in a polyacrylamide gel and silver stained. These data are representative of five independent experiments.

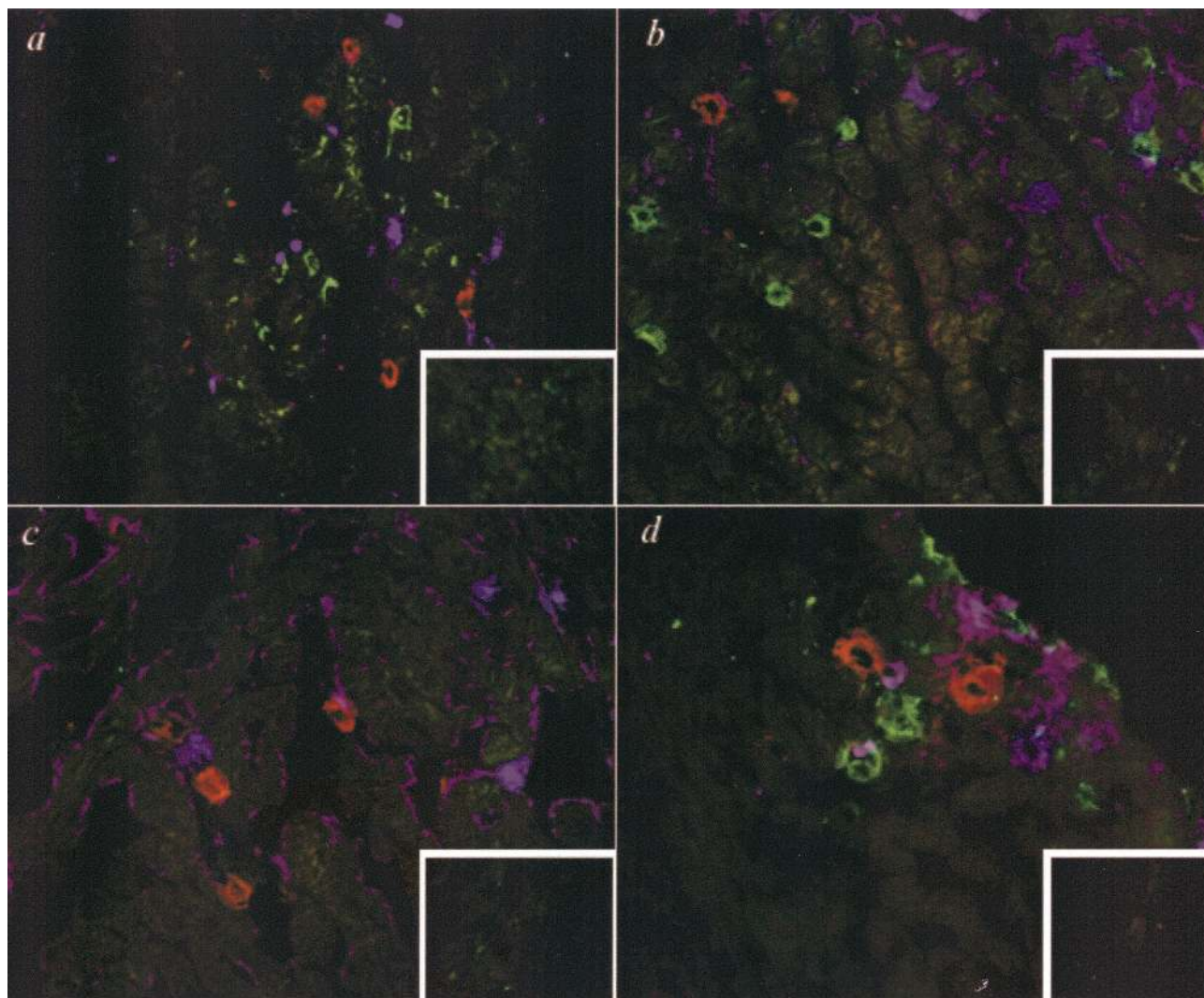


Figure 6. Lymphocyte infiltration in the myocardium of *T. cruzi*-infected p55-deficient mice. WT (a and b) or p55^{-/-} (c and d) mice were infected with 1×10^4 trypomastigote forms of *T. cruzi* (Y strain) and hearts were harvested at the ninth day after infection. Frozen sections of heart tissue were fixed in acetone and immunostained with the antibodies anti-CD4-fluorescein isothiocyanate (green) and anti-CD8-PE (red) and with anti-IP-10 (a and c), anti-Mig (b and d) (blue) (original magnification, $\times 400$). The **inset** shows the lack of specific staining in tissues incubated with control antibodies. These data are representative of four independent experiments.

in agreement with the presence of CD4- and CD8-positive cells in the tissue. On the other hand, we did not detect production of Mig and IP-10 in GKO mice (data not shown), p55^{-/-} mice showed a restricted pattern of Mig staining associated with modest perivascular infiltrates (Figure 6, c and d). To confirm these data we evaluated chemokine mRNA expression in the myocardium of these animals before and in the ninth day after infection. We found a strong infection-induced expression of the C-X-C chemokines Mig and IP-10 in the heart of infected WT mice (Figure 7). Confirming the immunohistochemical observations, GKO mice did not show detectable levels of mRNA for Mig and IP-10, whereas infected p55^{-/-} mice did. Despite the complete absence of Mig and IP-10 expression and CD4 and CD8 cells in GKO mice, a significantly increased number of nests were found in the myocardium (25.0 ± 3.3 parasite nests per section in GKO mice versus 3.5 ± 1.1 and 7.2 ± 1.9 in WT and p55^{-/-} mice, respectively).

Discussion

Although many reports have focused on the cellular compositions of inflammatory infiltrates during infection with *T. cruzi*, either at the inoculation site or in other organs including the heart,^{17,18} the mechanisms that control the differential leukocyte accumulation are not understood. In the present study, we provide data demonstrating the importance of cytokines, especially IFN- γ , for the modulation of chemokine expression *in vivo* and suggesting important roles for chemokines during *T. cruzi* infection.

Our results show that trypomastigote-induced expression of KC could be inhibited by the addition of IFN- γ (Figure 1) in a similar manner to the inhibitory effects of this cytokine on LPS-treated macrophages.¹⁹ The inhibitory effects of IFN- γ on KC expression could also be observed indirectly during infection in GKO mice. These animals had a progressive increase in KC expression which, together with an increased expression of MIP-2, was

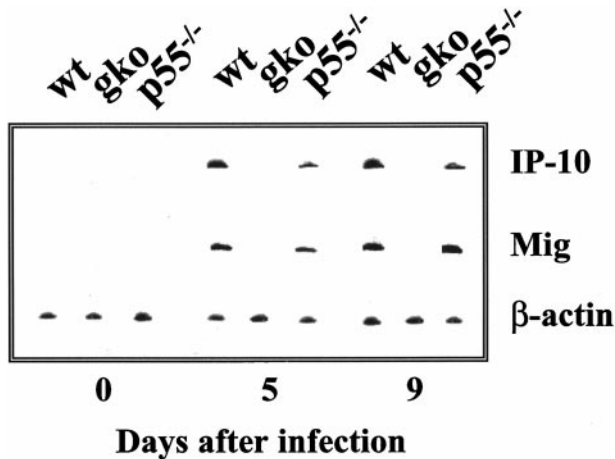


Figure 7. Myocardial chemokine production is reduced in infected GKO or p55^{-/-} mice. Total myocardial RNA was extracted from normal and 9-day-infected WT, GKO, and p55^{-/-} mice. Mig and IP-10 mRNA expression was assessed by RT-PCR. PCR products were electrophoresed in polyacrylamide gels and silver stained. These data are representative of two independent experiments.

likely responsible for the dramatic accumulation of polymorphonuclear neutrophils in the peritoneal cavity (Figure 3). These results suggest that IFN- γ is a potent inhibitor of KC-mediated neutrophil migration *in vivo*.

On the other hand, the enhancing effects of TNF- α on trypanostigote-mediated KC expression (Figure 1) were consistent with the observations made during infection of p55^{-/-} mice. These animals lack TNF- α -driven KC production and did not show expression of KC at any point during infection. Moreover, no neutrophilic infiltrate was found at the inoculation site or in the myocardium. It is possible that, in the absence of TNF- α signaling via the p55 receptor, reduced induction of KC occurred and this led to a failure to recruit neutrophils. Together, these data suggest that KC is a main factor that mediates neutrophil migration *in vivo* during *T. cruzi* infection and that, at least in this infection, TNF-R1 is essential for the production of KC. In contrast, animals deficient in the p75 TNF- α receptor did not differ from WT mice in respect to leukocyte recruitment or infection levels, suggesting that signaling via the latter receptor does not play a role in acute *T. cruzi* infection.

Lymphocyte recruitment *in vivo* was found to be significantly enhanced in p55^{-/-} animals. Interestingly, these animals had lower expression of almost all chemokines so far tested (Figure 5), with the exception of Mig. The accumulation of lymphocytes in the peritoneal cavity of p55^{-/-} animals was predominantly composed of CD3⁺, CD8⁺, and CD22⁺ cells on day 1 and CD3⁺, CD4⁺, and CD8⁺ cells on day 7 after infection (Figure 4). These results suggest that TNF- α down-regulates the recruitment of lymphocytes to the site of infection via activation of the TNF-R1 receptor. A possible beneficial consequence of a limited recruitment of lymphocytes might be to decrease the number of autoreactive cells within the infiltrates. This is in agreement with the increased number of inflammatory cells found in the late phase of *T. cruzi* infection in the heart of p55^{-/-} mice.²⁰ On the other hand, a possible negative consequence would be the

decrease of the number of parasite-specific cells, which might contribute to the less efficient response to infection. Our findings also suggest that, in the absence of an adequate TNF-R1 signaling, Mig seems to be the main mediator of lymphocyte migration *in vivo* during *T. cruzi* infection. However, although GKO mice did not express mRNA for the chemokines Mig, IP-10, and RANTES, these animals showed intense lymphocyte accumulation intraperitoneally, suggesting that other chemokines, such as MIP-1 α and JE/MCP-1 might be involved in lymphocyte recruitment to the latter site (Figures 3 and 4).

The direct correlation found between the presence of CD4- and CD8-positive cells (Figure 6) in the myocardium and the production of the chemokines Mig and IP-10 in this tissue suggests that these chemokines may be involved in the attraction of lymphocyte populations to the hearts of infected mice. Moreover, the limited lymphocytic infiltrates in GKO and p55^{-/-} mice, concomitantly with the complete absence of Mig and IP-10 expression and production in the heart of GKO and low levels in p55^{-/-} mice, strengthen the correlation. This could also be observed during chronic chagasic myocarditis, in which the expression of Mig and IP-10 remained high concomitantly to an intense expression of IFN- γ mRNA.²¹ Considering that Mig and IP-10 are able to induce the preferential migration of IFN- γ -producing cells by signaling via CXCR3,^{22,23} the above results raise the possibility that the cytokines IFN- γ and TNF- α may contribute to the genesis of myocarditis through induction of Mig and IP-10, which in turn, lead to the accumulation of IFN- γ -producing T cells as a part of a positive feedback loop. To provide further support for this observation, recently published data showed that IP-10 is essential for the accumulation of effector T cells that are required for resistance to infection with *Toxoplasma gondii*.²⁴

In GKO mice, not only did we observe a greater number of parasite nests but also a large increase in the size of each nest (data not shown). This is in marked agreement with our recent observation showing that cardiomyocytes controlled the replication of parasites in the presence of IFN- γ or TNF- α in a NO-dependent manner.¹⁵ Of interest, we also showed that *T. cruzi*-infected cardiomyocytes could produce a significant amount of lymphocyte-active chemokines. These results raise the interesting possibility that the production of chemokines by cardiomyocytes could, in turn, increase the infiltration of inflammatory cells in the heart and, then, control the infection via the production of IFN- γ -driven NO production. On the other hand, this cardiomyocyte/lymphocyte cross talk could also be a relevant interaction explaining the major pathological alterations observed in chronic chagasic heart disease.

The role of chemokines and their receptors in modulating innate as well as acquired immunity may not remain only in driving cell trafficking to target organs and tissues. Recent observations also point to direct effects of chemokines in the activation of key cell populations for innate and acquired immunity.²⁵ The results presented herein contribute to our understanding of the genesis and modulation of inflammatory reactions in *T. cruzi*-infected mice both at the inoculation site and in the myocardium. Inas-

much as inflammation seems to play a major role in the pathophysiology of Chagas heart disease, our results suggest that antagonizing the action of chemokines, especially IP-10 and Mig, might be of benefit in the treatment of chagasic myocarditis.

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