Calcineurin antagonists inhibit interferon-gamma production by downregulation of interleukin-18 in human mixed lymphocyte reactions.

Masahiko Kuinose, Okayama University
Hiromi Iwagaki, Okayama University
Yoshinori Morimoto, Okayama University
Hideo Kohka, Okayama University
Kenta Kobashi, Okayama University
Hiroshi Sadamori, Okayama University
Masaru Inagaki, Okayama University
Naoto Urushihara, Okayama University
Takahito Yagi, Okayama University
Noriaki Tanaka, Okayama University
Calcineurin antagonists inhibit interferon-gamma production by downregulation of interleukin-18 in human mixed lymphocyte reactions.*

Masahiko Kuinose, Hiromi Iwagaki, Yoshinori Morimoto, Hideo Kohka, Kenta Kobashi, Hiroshi Sadamori, Masaru Inagaki, Naoto Urushihara, Takahito Yagi, and Noriaki Tanaka

Abstract

Tacrolimus (FK-506) and cyclosporin A (CsA) are calcineurin antagonists used widely as T-cell immunosuppressants; however, their relative efficacy on the production of interleukin-18 (IL-18) remains undefined. We have examined the effects of FK-506 and CsA on the cytokine generation of human peripheral blood mononuclear cells (PBMCs) in mixed lymphocyte reaction (MLR) with lipopolysaccharide (LPS). We studied the levels of interleukin-18 (IL-18), IL-12, IL-10, IL-6, IL-2 and interferon-gamma (IFN-gamma) in the supernatant in allo-MLR by ELISA assay. Supernatant levels of IFN-gamma, IL-2, IL-6, IL-10 and IL-12 were detected 12 h after MLR and markedly increased thereafter. In contrast, production of IL-18 was detected at 12 h, reached a near maximum level at 24 h and decreased at 72 h. These results suggested that IFN-gamma production depended on IL-18, IL-12 and IL-2 in the early phase of MLR and depended mainly on IL-12 and IL-2 in the late phase. Both calcineurin antagonists inhibit the generation of IL-18, which plays a large role in allogeneic cell interactions, in macrophages and they also promote an equivalent down-regulation of T helper 1 (Th1) and Th2 responses in a concentration-dependent manner. About 90% of IFN-gamma production induced by MLR was inhibited by an anti-IL-18 antibody, showing that IL-18 can trigger IFN-gamma production in MLR. These results suggest that dual signaling consisting of antigen-driven nuclear factor of activated T cells (NFAT) activation and LPS-mediated NF-kappaB activation is crucial for IL-18 production in macrophages, and that IL-18 can trigger IFN-gamma production in T-cells by MLR.

KEYWORDS: tacrolimus, cyclosporin, calcineurin antagonist

*PMID: 11061569 [PubMed - indexed for MEDLINE]
Copyright (C) OKAYAMA UNIVERSITY MEDICAL SCHOOL
Calcineurin Antagonists Inhibit Interferon-gamma Production by Downregulation of Interleukin-18 in Human Mixed Lymphocyte Reactions

Masahiko KUNIYONE, Hiromi IWAGAKI*, Yoshinori MORIMOTO, Hideo KOHKA, Kenta KOBASHI, Hiroshi SADAMORI, Masaru INAGAKI, Naoto URUSHIHARA, Takahito YAGI and Noriaki TANAKA

Department of Surgery I, Okayama University Medical School, Okayama 700-8558, Japan

Tacrolimus (FK-506) and cyclosporin A (CsA) are calcineurin antagonists used widely as T-cell immunosuppressants; however, their relative efficacy on the production of interleukin-18 (IL-18) remains undefined. We have examined the effects of FK-506 and CsA on the cytokine generation of human peripheral blood mononuclear cells (PBMCs) in mixed lymphocyte reaction (MLR) with lipopolysaccharide (LPS). We studied the levels of interleukin-18 (IL-18), IL-12, IL-10, IL-6, IL-2 and interferon-γ (IFN-γ) in the supernatant in allo-MLR by ELISA assay. Supernatant levels of IFN-γ, IL-2, IL-6, IL-10 and IL-12 were detected 12 h after MLR and markedly increased thereafter. In contrast, production of IL-18 was detected at 12 h, reached a near maximum level at 24 h and decreased at 72 h. These results suggested that IFN-γ production depended on IL-18, IL-12 and IL-2 in the early phase of MLR and depended mainly on IL-12 and IL-2 in the late phase. Both calcineurin antagonists inhibit the generation of IL-18, which plays a large role in allogeneic cell interactions, in macrophages and they also promote an equivalent down-regulation of T helper 1 (Th1) and Th2 responses in a concentration-dependent manner. About 90% of IFN-γ production induced by MLR was inhibited by an anti-IL-18 antibody, showing that IL-18 can trigger IFN-γ production in MLR. These results suggest that dual signaling consisting of antigen-driven nuclear factor of activated T cells (NFAT) activation and LPS-mediated NF-κB activation is crucial for IL-18 production in macrophages, and that IL-18 can trigger IFN-γ production in T-cells by MLR.

Key words: tacrolimus, cyclosporin, calcineurin antagonist

Interleukin-18 (IL-18) was identified as a cytokine that induced interferon-γ (IFN-γ) in the sera of mice with septic shock (1). IL-18 is secreted from lipopolysaccharide (LPS)-activated macrophages but also from a wide variety of cells. Subsequent studies have shown that IL-18 performs a wide variety of activities in various cells, and recent reviews summarize these activities (2, 3). The cytokine was found to enhance production of T helper 1 (Th1)-type cytokines (IFN-γ and granulocyte/macrophage-colony stimulating factor (GM-CSF)) but not Th2 cytokines (IL-4 and IL-10) in nonadherent murine spleen cells and human peripheral blood mononuclear cells (PBMCs) with T cell receptor (TCR)/CD3 stimulation (1, 4, 5). It was also reported that IL-18 induced the production of IL-2 and IFN-γ and proliferation of TCR/CD3-stimulated murine Th1 clones; however, the cytokine did not affect the production of IL-4 nor the proliferation of Th2 clones in vitro (6, 7). In a previous study, we demonstrated that IL-18 upregulated ICAM-1 expression in a KG-1 monocytic cell line through an IFN-γ-independent pathway (8) and that IL-18 is involved in allo-mixed lymphocyte reactions (MLR) (9).

Tacrolimus (FK-506) and cyclosporin (CsA) are calcineurin antagonists used widely as T-cell immunosuppressants to prevent allograft rejection after solid organ transplantation. Investigations have indicated that uses for these agents also include prevention of graft-versus-host disease after allogeneic bone marrow transplantation and treatment of severe, steroid-dependent asthma. FK-506 and CsA act through ligation of distinct intracellular binding protein targets (FK binding protein for FK-506,
cyclophilin A for CsA). These drug-binding protein complexes inhibit the function of calcineurin, which normally activates various transcription factors, including nuclear factor of activated T cells (NFAT) and nuclear factor-κB (NF-κB), that regulate the expression of key cytokines and signaling proteins (9, 10). The potential differential regulation of human antigen-stimulated T-cell subsets by calcineurin antagonists has not been fully described. We herein present a comprehensive study of the effects of calcineurin antagonists on cytokine generation, including that of IL-18, in an allo-MLR system.

Materials and Methods

Reagents. One mM each of FK-506 and CsA were dissolved in ethanol and further diluted in medium and added to the cultures. FK-506 and CsA were generous gifts from Fujisawa Pharmaceutical Co., Ltd. (Ibaraki, Japan) and Sandoz Ltd (Basel, Switzerland), respectively. Anti-recombinant human (rHu) IL-18 antibody (Ab) and anti-HuIFN-γ Ab were prepared as described elsewhere (4). Anti-HuIL-12 Ab and anti-HuIL-2 Ab were purchased from PharMingen (San Diego, CA, USA).

Culture conditions in mixed lymphocyte reaction (MLR). Human peripheral blood mononuclear cells (PBMC) were isolated from buffy coats of 5 healthy volunteers by centrifugation on a density gradient of Ficoll-Paque (Pharmacia Uppsala, Sweden). The PBMC were then washed 3 times in an RPMI 1640 medium (Nissui Co., Tokyo, Japan) supplemented with 10% (v/v) heat-inactivated fetal bovine serum (FBS), 20 μg/ml kanamycin, and 100 μg/ml streptomycin and penicillin (Sigma Chemical Co., St. Louis, MO, USA). The PBMC were suspended at a final concentration of 1.25 × 10⁶ cells/ml in the RPMI 1640 medium supplemented with 10% (v/v) heat-inactivated FBS. In the MLR, cells from an individual were mixed with cells from an unrelated person. The final concentration of cells was adjusted to 2.5 × 10⁵ cells/ml. Triplicate wells were incubated with 5 μg/ml LPS (Sigma). The plates were incubated at 37°C in 5% CO₂/95% air for 72 h, after which the supernatants were aspirated and stored at −20°C until being assayed for cytokines. When the effect of calcineurin antagonists or other reagents was to be examined, all reagents were added to the media at the start of incubation and cultured for 48 h under the same condition. All experiments were performed in at least triplicate samples.

Cytokine assays. The cytokines were measured using ELISAs employing the multiple Abs sandwich principle (for IL-18, MBL; for other cytokines, Quantikine, R & D systems, Minneapolis, MN, USA). The detection limits of the ELISAs for IL-18, IL-12, IL-10, IL-6 and IFN-γ were 10 pg/ml, and for IL-2 it was 5 pg/ml.

Statistical analysis. Data are expressed as mean ± SEM. For mean comparisons, one-way analysis of variance (ANOVA) was used. Differences were considered statistically valid at P < 0.05.

Results

1) Kinetics of IL-18 production in MLR

IL-18 production was observed 12 h after MLR, reached a near maximum level at 24 h, and decreased at 72 h (Fig. 1).

2) Kinetics of Th1 cytokine productions in MLR

Production of IL-12, IL-2 and IFN-γ was upregulated at 12 h and increased during the culture period. (Fig. 2).

3) Kinetics of Th2 cytokine productions in MLR

Production of IL-6 and IL-10 was upregulated at 12 h and reached a near maximum level at 24 h (Fig. 3).

4) Effect of FK-506 and CsA on IL-18 production in MLR

Both calcineurin antagonists induced significant concentration-dependent inhibition of IL-18 production in MLR. The concentration of FK-506 required for complete inhibition was approximately 10 times lower than that of CsA (Fig. 4).

5) Effect of FK-506 and CsA on Th1 cytokine productions in MLR

Both calcineurin antagonists induced significant concentration-dependent inhibition of Th1 cytokines produced in MLR. The concentration of FK-506 required for complete inhibition was approximately 10 times lower than that of CsA (Fig. 5).

6) Effect of FK-506 and CsA on Th2 cytokine productions in MLR

Both calcineurin antagonists induced significant concentration-dependent inhibition of Th2 cytokines produced in MLR. The concentration of FK-506 required for complete inhibition was approximately 10 times lower
than that of CsA (Fig. 6).

(7) Blocking test in MLR

To investigate cytokine interactions in MLR, we examined the effect of anti-IL-18 Ab, anti-IL-12 Ab, anti-IL-2 Ab and anti-IFN-γ Ab on production of IFN-γ and IL-10 at 48 h. The addition of these Abs significantly reduced IFN-γ production in MLR. In contrast, amounts of IL-10 were significantly downregulated after the addition of anti-IL-18 Ab and anti-IFN-γ Ab, but they were not downregulated by the addition of anti-IL-12 Ab and anti-IL-2 Ab (Fig. 7).

---

**Fig. 1**
Kinetics of IL-18 production in MLR. Closed circles show the levels of IL-18 in pooled supernatants from MLR. Values of IL-18 are expressed as pg/ml. Assays were performed in triplicate, and values are means ± SEM.

**Fig. 2**
Kinetics of Th1 cytokine productions in MLR. Closed circles show the levels of cytokines in pooled supernatants from MLR. Values of IL-12, IL-2 and IFN-γ are expressed as pg/ml. Assays were performed in triplicate, and values are means ± SEM.
**Fig. 3** Kinetics of Th2 cytokine productions in MLR.
Closed circles show the levels of cytokines in pooled supernatants from MLR. Values of IL-10 and IL-6 are expressed as pg/ml. Assays were performed in triplicate, and values are means ± SEM.

**Fig. 4** Effect of FK506/CsA on IL-18 production in MLR. FK506 and CsA were added to the media at the start of incubation and cultured for 48 h. Assays were performed in triplicate, and values are mean ± SEM.
Fig. 5  Effect of FK506/CsA on Th1 cytokine productions in MLR.
FK506 and CsA were added to the media at the start of incubation and cultured for 48 h. Assays were performed in triplicate, and values are mean ± SEM.

Fig. 6  Effect of FK506/CsA on Th2 cytokine productions in MLR.
FK506 and CsA were added to the media at the start of incubation and cultured for 48 h. Assays were performed in triplicate, and values are mean ± SEM.
Discussion

FK-506 and CsA are calcineurin antagonists. These agents act, in part, through the inhibition of nuclear transcription factors such as NFAT and NF-κB. Functionally, FK-506 and CsA inhibit the transcription of numerous cytokines and immunomodulators involved in T-cell activation and proliferation (8, 9). Previous studies have demonstrated the ability of calcineurin antagonists to downregulate IFN-γ (12–15). However, cellular activation in these studies was achieved with mitogens; therefore, the relevance of these systems to antigen-presenting cell (APC) and antigen-driven Th1 and Th2 phenotypic responses is unclear.

The MLR is an important in vitro model for studying allosresponsiveness. Using the MLR, we can measure the disparity in major histocompatibility complex (MHC) antigens between individuals (16, 17). Because cytokines play a crucial role in the posttransplantation response, cytokine release and interactions in MLR are important in the field of transplantation medicine. In human MLR, responder cells recognize alloantigens on the surface of stimulator cells and undergo proliferation. It has been shown that IFN-γ is secreted predominantly into the supernatant and that the IFN-γ response depends on differences in HLA-DR between the 2 individuals (18, 19).

The current study showed that production of IL-18 was upregulated at 12 h, reached a near maximum level at 24 h, and decreased at 72 h. The decrease of IL-18 at 72 h might be attributable to its consumption and degradation or to the negative effect on the IL-18-initiating cytokine cascade caused by the accumulation of Th2 cytokines. Production of both IL-12 and IL-2 was observed at 12 h and then increased markedly. Production of IFN-γ was detected at 12 h and increased there after (Fig. 1, Fig. 2). These results suggested that production of IFN-γ depends on IL-18, IL-12 and IL-2 in the early phase (12–48 h) and predominantly on both IL-12 and IL-2 in the late phase (48–72 h) of the human MLR system.

The synergistic actions of IL-18 and IL-12 have been observed in the production of IFN-γ (20–23). Yoshimoto et al. demonstrated that IL-12 upregulated the expression of IL-18 receptors in T cells, Th1 cells and B cells, which enabled the synergistic production of IFN-γ (21). In contrast, IL-18 can stimulate IFN-γ production in an IL-12-independent manner in KG-1, a monocytic cell line (8). It is quite likely that the cooperative coexistence of IL-18 and IL-12 can induce a strong Th1 response in the human MLR system.

The potential differential regulation of human antigen-stimulated T cell subsets by calcineurin antagonists has not been fully described. The present research has demonstrated concentration-dependent inhibition by calcineurin antagonists of IL-18, Th1 and Th2 cytokine productions in the MLR system (Fig. 4, Fig. 5, Fig. 6). These results suggested that calcineurin antagonists...
promoted an equivalent down-regulation of Th1 and Th2 responses in the human MLR system. However, a number of reports have suggested a relative resistance of Th2 cytokines to the effects of calcineurin antagonists (24–35). Differences in cell populations and methods of cellular activation used in these experiments complicate their interpretation and are likely to account for the contradictory results.

It has been reported that mitogenic stimulation of phenotypically specific CD4+CD4+ T cell clones results in nonphenotypically selective cytokine generation; however, subsequent restimulation with antigen and antigen-presenting cells (APCs) will restore phenotypic specificity (36, 37). Thus, mitogen stimulation appears to obviate phenotypic specificity. Moreover, mitogen stimulation is clearly less physiologic than antigen stimulation. Finally, it has been reported that sensitivity to calcineurin antagonists may vary with the quantity of the activation signal applied to the responder cells (38).

We also found differences in cytokine production profiles of IFN-γ and IL-10 between IL-18 and other cytokines in the MLR system. In the blocking test, the addition of anti-IL-18 Ab and anti-IFN-γ Ab significantly (P < 0.01) reduced the production of IFN-γ and IL-10. In contrast, treatment with anti-IL-12 Ab and anti-IL-2 Ab significantly (P < 0.01) inhibited IFN-γ production but not IL-10 production, as shown in Fig. 5. These results suggested that IL-18 and IFN-γ play a role in the induction of IL-10, which may function as a negative feedback for excessive Th1 response in the MLR system.

In contrast, IL-12 and IL-2 do not induce IL-10 directly, at least in the MLR system. It has also been reported that treatment with IL-18 and anti-CD3 Ab produced IL-2 and IFN-γ in Th1 cells (39, 40). IL-18 strongly induced NF-κB activation, which was detected by electrophoretic mobility shift assay using an NF-κB binding site of the IL-2 promoter and reporter gene analysis in Th1 cells (41). Taking these results into consideration, it is likely that IL-18 is an important triggering cytokine in allogeneic cell interactions and is present in the most upstream of cytokine cascades in immune responses, at least in the MLR system.

IL-12 has also reportedly induced IFN-γ production by Th1 cells, and the immune responses induced by IL-12 are similar to those induced by IL-18 in vivo (41). However, the signals for IFN-γ production by IL-18 have been revealed to be different from those induced by IL-12, since IL-18 with IL-12 synergistically induced IFN-γ production (42), which was consistent with the results of the present study. As a unique signal activated by IL-12, it was reported that STAT4 was essential for IFN-γ production. Furthermore, it has been reported that IL-18 could not induce STAT4 activation in Th1 cells (7). A recent report suggested that dual signaling consisting of IL-18-induced NF-κB activation and TCR/CD3-mediated NFAT activation is crucial for IFN-γ production in Th1 cells (43).

As described above, IL-18 is secreted from LPS-activated macrophages. Although LPS alone did not induce IL-18 production in PBMCs (9), the combination of the LPS/MLR system dramatically induced IL-18, as shown in the present study. Furthermore, calcineurin antagonists inhibited IL-18 production dose-dependently in human antigen-stimulated PBMCs. These results suggested that dual signaling consisting of LPS-induced NF-κB activation and antigen-driven NFAT activation is crucial for IL-18 production in macrophages.

In conclusion, the present study clearly demonstrated the importance of IL-18 in allogeneic cell interactions and that its generation in the human MLR system was strongly inhibited by either FK-506 or CsA. Although there is evidence that calcineurin antagonists can effect T cell responses through interference with antigen presentation from APCs (44), the similarity of PBMCs in the MLR system of this study mitigate against significant confounding of these data. The current study also suggested that dual signaling consisting of antigen-driven NFAT activation and LPS-mediated NF-κB activation is crucial for IL-18 production in monocytes/macrophages.

Recent research has clarified the mechanism of Caspase-1-independent, Fas/Fas ligand-mediated IL-18 secretion from macrophages. Administration of Propionibacterium acnes upregulated functional Fas expression of macrophages in an IFN-γ-dependent manner, and these macrophages became able to secrete mature IL-18 upon stimulation with FasL (45). However, which molecules induce IL-18 has not yet been fully elucidated. Further research is warranted for elucidating the production of IL-18.

References


34. Mori A, Suko M, Nishizaki Y, Kaminuma O, Kobayashi S, Matsuzaki G, Yamamoto K, Ito K, Tsuuroka N and Okudaira H: IL-5 production by CD4+ T cells of asthmatic patients is suppressed by glucocorticoids and the immunosuppressants FK506 and cyclosporin A. Int
October 2000

41. Ahn HJ, Maruo S, Tomura M, Mu J, Hamaoka T, Nakanishi K, Clark

Calcineurin Antagonists Regulate IL-18 Responses

44. Little RG II nd, Ebertowski LA and David CS: Inhibition of alloantigen presentation by cyclosporine. Transplantation (1990) 49, 937-944.

Received April 27, 2000; accepted June 9, 2000.