Continuous Administration of Anti–Interleukin 10 Antibodies Delays Onset of Autoimmunity in NZB/W F1 Mice

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Summary

We have previously shown that continuous administration of anti–interleukin 10 (anti-IL-10) antibodies (Abs) to BALB/c mice modifies endogenous levels of autoantibodies, tumor necrosis factor α (TNF-α), and interferon γ, three immune mediators known to affect the development of autoimmunity in “lupus-prone” New Zealand black/white (NZB/W)F1 mice. To explore the consequences of IL-10 neutralization in NZB/W F1 mice, animals were injected two to three times per week from birth until 8-10 mo of age with anti-IL-10 Abs or with isotype control Abs. Anti-IL-10 treatment substantially delayed onset of autoimmunity in NZB/W F1 mice as monitored either by overall survival, or by development of proteinuria, glomerulonephritis, or autoantibodies. Survival at 9 mo was increased from 10 to 80% in anti-IL-10–treated mice relative to Ig isotype–treated controls. This protection against autoimmunity appeared to be due to an anti-IL-10–induced upregulation of endogenous TNF-α, since anti-IL-10–protected NZB/W F1 mice rapidly developed autoimmunity when neutralizing anti-TNF-α Abs were introduced at 30 wk along with the anti-IL-10 treatment. Consistent with the protective role of anti-IL-10 treatment in these experiments, continuous administration of IL-10 from 4 until 38 wk of age accelerated the onset of autoimmunity in NZB/W F1 mice. The same period of continuous IL-10 administration did not appear to be toxic to, or cause development of lupus-like autoimmunity in normal BALB/c mice. These data suggest that IL-10 antagonists may be beneficial in the treatment of human systemic lupus erythematosus.

The (New Zealand black × New Zealand white [NZB × NZW]) F1 hybrid mouse develops a severe autoimmune disease that closely resembles SLE in humans (1-3). NZB/W F1 mice spontaneously develop a fatal immune complex–mediated glomerulonephritis around 6-9 mo in female animals, and 12-18 mo in male animals. Previous attempts to define the underlying cause of autoimmunity in NZB/W F1 mice have focused on the potential role of MHC genes. Indeed, IFN-γ, a cytokine which upregulates expression of MHC class II antigens in a wide variety of cell types (4-6), accelerates the development of autoimmunity in NZB/W F1 mice (7-10). More recently, several studies (11-14) have suggested that the TNF-α gene, which is located within the MHC complex (15-17), may be involved in the pathogenesis of lupus nephritis in NZB/W F1 mice. These studies reveal that NZB/W F1 mice produce exceptionally low levels of TNF-α, and that this correlates with a restriction fragment length polymorphism in the TNF-α gene (11), and a polymorphism in simple dinucleotide tandem repeats in the 5' regulatory region of the TNF-α gene (13, 14). A causative role for these observations is implied by the fact that replacement therapy with recombinant TNF-α significantly delays development of nephritis in NZB/W F1 mice (11, 12).

IL-10 is a cytokine produced by subsets of activated T cells, B cells, and macrophages which mediates a variety of both immunostimulatory and immunosuppressive properties in mouse and human in vitro assays (for a review see references 18 and 19). In a recent effort to evaluate the physiological role of IL-10, we have treated BALB/c mice continuously from birth until 8 wk of age with neutralizing anti-IL-10 Abs (20, 21). Consistent with the known in vitro properties of IL-10 (18, 19), anti-IL-10–treated mice are characterized by elevated levels of endogenous IFN-γ and TNF-α (20, 21). The elevated IFN-γ in turn leads to the depletion of a numerically small subset of B lymphocytes, termed Ly1, CD5, or B-1 B cells (20), a population from which most murine autoantibodies are derived (22). These studies in normal mice suggested that neutralization of IL-10 may produce some de-

� Abbreviation used in this paper: NZB/W, New Zealand black/white.
sirable consequences in NZB/W F₁ mice, i.e., elevation of endogenous TNF-α and reduction of autoantibody production, and some undesirable consequences, i.e., elevation of endogenous IFN-γ. In this report, we directly address the overall effects of continuous anti-IL-10 treatment on development of autoimmunity in NZB/W F₁ female mice. Our data indicate that neutralization of IL-10 significantly delays onset of autoimmunity in these mice because of an upregulation of endogenous TNF-α.

Materials and Methods

Mice. NZB/W F₁ mice were bred in our colony at DNAX Research Institute using NZB females purchased from The Jackson Laboratory (Bar Harbor, ME) and NZW males purchased from Simonsen Laboratories (Gilroy, CA). Only female F₁ mice were used in this study because of their more rapid onset of autoimmunity.

Anti-IL-10 Treatment. Groups of 17-23 B/W F₁ female mice were treated with either a rat IgM (designated SXC.1) or a rat IgG1 (designated JES.2A5) mAb specific for murine IL-10. Details about the production and characteristics of these Abs are given elsewhere (23, 24). Similarly, groups of 21-23 age-matched B/W F₁ female mice were treated with either of two isotype-matched control mAbs, designated J5/D (IgM) or GL113 (IgG1). Anti-IL-10 and isotype control mAbs were harvested as ascites from athymic nude (nu/nu) mice, purified by two sequential ammonium sulfate precipitation steps, dialyzed against PBS, and quantitated by protein electrophoresis and measurement of optical density. Treatment consisted of 3 parts: during the first week after birth, 0.2 mg of mAb per mouse was injected intraperitoneally either four or two times per week for the IgM and IgG1 Abs, respectively. During the second week, 0.5 mg of mAb per mouse was injected intraperitoneally either three or two times per week for the IgM and IgG1 Abs, respectively. From 3 wk of age and onwards, 1 mg of mAb per mouse was injected intraperitoneally either three or two times per week for IgM and IgG1 Abs, respectively.

Assessment of Renal Disease. Renal disease was assessed either via the development of proteinuria or by histological evaluation of glomerulonephritis. Proteinuria was measured colorimetrically using dip sticks (Albustix; Miles Laboratories, Inc., Elkhart, IN), and graded according to the following code: trace, 10 mg/dl; 1+, 30 mg/dl; 2+, 100 mg/dl; 3+, 300 mg/dl; and 4+, 1,000 mg/dl. Severity of glomerulonephritis was evaluated histologically by tissue sectioning and staining, and was graded on a 0-4+ scale based on the intensity and extent of histopathologic changes, as described previously by Berden et al. (25). Briefly, the score system for evaluating severity of glomerulonephritis was as follows: a grade of 0 was given to kidneys without glomerular lesions; 1+ lesions corresponded to minimal thickening of the mesangium; 2+ lesions contained noticeable increases in both mesangial and glomerular cellularularity; 3+ lesions were characterized by the preceding conditions with superimposed inflammatory exudates and capsular adhesions; and in 4+ lesions, the glomerular architecture was obliterated in >70% of glomeruli, and tubular cast formation was extensive.

Autoantibody ELISA. Serum antibodies specific for double- or single-stranded DNA (ds or ssDNA) were quantitated by ELISA as described elsewhere (26). Briefly, 5 μg/ml ds- or ss-DNA was used to coat ELISA plates (Flow Laboratories, McLean, VA) in an overnight incubation at 4°C. Antigen-coated plates were subsequently blocked for 1 h with PBS containing 0.05% Tween 20, 0.02% NaN₃, and 0.1% BSA, then incubated for 1 h at room temperature with test or standard sera diluted from 1:100 in two-fold serial dilutions. Plates were then washed with PBS-0.05% Tween 20, and incubated for 1 h with 1 μg/ml horseradish peroxidase (HRP)-conjugated anti-mouse IgG or IgM (Zymed Laboratories, Inc., S. San Francisco, CA). Absorbance was measured using a microplate reader (Vmax; Molecular Devices, Menlo Park, CA) with optical density read at 490 nm. A 1:100 dilution of this standard serum was arbitrarily assigned a value of 100 U/ml.

Serum Concentration of TNF-α. Serum levels of TNF-α were evaluated using a cytokine-specific ELISA, commercially available from Endogen, Inc. (Boston, MA).

Results

Continuous Administration of Anti-IL-10 Antibodies Delays Onset of Autoimmunity in NZB/W F₁ Mice. In normal mice, continuous administration of anti-IL-10 Abs from birth until adulthood alters the animals' levels of IFN-γ, TNF-α, and autoantibodies (20, 21), three mediators known to affect the development of autoimmunity in lupus-prone NZB/W F₁ mice (3, 7-14). To determine whether IL-10 neutralization in NZB/W F₁ mice may indeed influence their development of autoimmunity, groups of 20-23 female NZB/W

![Figure 1](https://example.com/figure1.png)
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Figure 2. Development of proteinuria in NZB/W F1 female mice treated with anti-IL-10 Abs. Groups of 22 female mice were injected three times per week from birth and throughout the entire study with either SXC.1 anti-IL-10 Ab (O) or an isotype control Ab (Q). Development of renal disease was assessed by measuring proteinuria using AlbuStix dip sticks from weeks 12–42. Data show proportion of mice with more than 300 mg protein/dl in their urine. Similar results were obtained in two additional experiments.

Figure 3. Autoantibody production in anti-IL-10–treated NZB/W F1 mice. NZB/W F1 female mice were injected with SXC.1 anti-IL-10 Ab (O) or an isotype control (Q) three times per week from birth and throughout the entire study. Sera collected at 5 or 6 mo were evaluated for content of IgG Ab binding ds-DNA using an ELISA described in Materials and Methods. Each circle represents an individual mouse. Serum IgM anti-ds DNA Abs were also decreased in anti-IL-10–treated NZB/W F1 mice (data not shown).

Figure 4. Histological evaluation of severity of renal disease in control and treated NZB/W F1 mice. Kidneys were collected, sectioned, and stained by standard procedures. Severity of glomerulonephritis was graded according to the following score system: a grade of 0 was given to kidneys without glomerular lesions; 1+ lesions correspond to minimal thickening of the mesangium; 2+ lesions contained noticeable increases in both mesangial and glomerular cellularity; 3+ lesions were characterized by the preceding conditions with superimposed inflammatory exudates and capsular adhesions; and in 4+ lesions, the glomerular architecture was obliterated in >70% of glomeruli, and tubular cast formation was extensive. Results show number of animals in each group with a particular grading of renal disease, with each circle representing an individual mouse.

IL-10 Neutralization Delays Onset of Autoimmunity in NZB/W F1 Mice because of Elevation of TNF-α. Several studies have implicated atypically low endogenous TNF-α levels as a contributing factor in the pathogenesis of autoimmunity in NZB/W F1 mice (11–14). Since IL-10 neutralization...
Figure 5. Histological examination of kidneys from anti-IL-10-treated NZB/W F1 mice (PAS staining; H250). (A) A typical grade 4 lesion observed in the isotype control-treated mice shown in Fig. 4. Note accumulation of amorphous PAS-positive materials in the mesangial matrix, and protein casts in renal tubules. (B) A grade 1 lesion in the anti-IL-10 Ab-treated group of mice. Note slight thickening of the mesangium.

IL-10 Administration Accelerates the Onset of Autoimmunity in NZB/W F1 Mice. The experiments described above suggest that IL-10 neutralization in NZB/W F1 mice delays onset of autoimmunity in these animals by elevating their endogenous levels of TNF-α. This conclusion was supported by separate experiments which showed that continuous administration of IL-10 to NZB/W F1 mice caused a small but significant acceleration of onset of autoimmunity. Groups of 20 female NZB/W F1 mice were injected intraperitoneally three times per week from 4 to 35 wk of age with either 1 μg recombinant murine IL-10 or an equivalent volume of PBS. As shown in Fig. 8, expt. 1, PBS injected animals showed a typical survival curve, generally dying between 25 and 40 weeks. In IL-10 treated animals, the survival curve showed a more rapid onset of autoimmunity, with death occurring on average 5 to 6 weeks earlier (Fig. 8, expt. 1). Similar results were obtained in a second independent experiment involving identical numbers of animals monitored for a slightly shorter period of time (Fig. 8, expt. 2). The accelerated death of IL-10-treated NZB/W F1 mice was accompanied by accelerated development of proteinuria and glomerulonephritis (data not shown), indicating that the survival curves reflected death.

Figure 6. Serum TNF-α levels in anti-IL-10-treated NZB/W F1 mice. NZB/W female mice were injected with SXC.1 anti-IL-10 Ab (●) or an isotype control (○) three times per week from birth and throughout the entire study. Sera collected at 5, 7, or 8 mo were evaluated for TNF-α content using a cytokine-specific ELISA. Each circle represents an individual mouse.
due to autoimmunity. Administration of equivalent amounts of IL-10 to normal BALB/c mice from 4 to 38–42 wk of age did not induce autoimmune disease in these animals, as evaluated by animal survival (Fig. 8) or development of proteinuria (data not shown).

Discussion

We show here that continuous treatment of lupus-prone NZB/W F1 mice from birth until 38–42 wk of age with anti-IL-10 Abs substantially delays onset of autoimmunity in these animals. Conversely, continuous administration of IL-10 to NZB/W F1 mice from 4 to 38 wk of age causes a significant acceleration in the development of autoimmunity in these animals, while having no overt consequence in normal BALB/c mice. The protective effect of anti-IL-10 Ab administration to NZB/W F1 mice can be explained by the elevation of endogenous TNF-α levels that results from this treatment (21 and Fig. 6). This conclusion can be drawn from the fact that neutralizing anti-TNF-α Abs reversed the protective effect of anti-IL-10 treatment when introduced into the anti-IL-10–treated animals some 30 wk after experiment initiation (Fig. 7). These data predict a potential clinical use for IL-10 antagonists in treatment of human SLE. Such treatment may be particularly relevant to the subset of SLE patients expressing DR2+ DQw1+ histocompatibility antigens, since PBL from these patients produce atypically low levels of TNF-α (27). Conversely, the use of chronic long-term IL-10 administration may be contraindicated in SLE patients. On this point, it should be noted however that single-dose or short-term IL-10 administration had no significant effect on NZB/W F1 mice (our unpublished observations), and long-term IL-10 administration had no significant effect on normal BALB/c mice (Fig. 8).

The mechanism of anti-IL-10–mediated protection of NZB/W F1 mice from onset of autoimmunity suggests a protective role for TNF-α in this disease. Indeed this has already been demonstrated experimentally by direct cytokine administration (11, 12). While the basis of TNF-α–mediated protection has not yet been elucidated, several known properties of this cytokine suggest potential mechanisms of action in this disease model. One possible explanation relates to the fact that TNF-α significantly decreases IFN-γ–induced upregulation of MHC class II expression on different mature cells collected from NZB/W F1 mice (14, 28). Aberrant up-regulation of class II antigen expression is thought to be a causative mechanism in a variety of endocrine autoimmune diseases (29). Thus, suppression of this effect may be advantageous in NZB/W F1 mice. Alternatively, whereas TNF-α does not suppress humoral immunity in normal or lupus mice (14, 30), there have been some reports that long-term administration of this cytokine causes suppression of cell-mediated immunity in these animals (30). The level of protection afforded by IL-10 neutralization in NZB/W F1 mice is comparable to that previously obtained with direct administration of TNF-α (11, 12). The possibility of using either approach for treatment of SLE patients depends of course on translation of the current findings to the human disease situation, and the long-term toxicity of these reagents.

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