Immunoglobulin Treatment Reduces Atherosclerosis in apo E Knockout Mice

Antonino Nicoletti,‡ Srini Kaveri,‡ Giuseppina Caligiuri,* Jean Bariety,‡ and Göran K. Hansson*‡
*Center for Molecular Medicine, Karolinska Institute, S-17176 Stockholm, Sweden; and ‡INSERM U430, Hôpital Broussais, Paris, France

Abstract

Atherosclerosis is associated with immune activation. T cells and macrophages infiltrate atherosclerotic plaques and disease progression is associated with formation of autoantibodies to oxidized lipoproteins. In the apo E knockout mouse, a genetic model of cholesterol-induced atherosclerosis, congenital deficiency of macrophages, lymphocytes, or interferon-γ receptors result in reduced lesion formation. We have now evaluated whether immune modulation in the adult animal affects disease development. Injections of 7-wk-old male apo E knockout mice with polyclonal immunoglobulin preparations (ivIg) during a 5-d period reduced fatty streak formation over a 2-mo period on cholesterol diet by 35%. Fibrofatty lesions induced by diet treatment for 4 mo were reduced by 50% in mice receiving ivIg after 2 mo on the diet. IvIg treatment also reduced IgM antibodies to oxidized LDL and led to inactivation of spleen and lymph node T cells. These data indicate that ivIg inhibits atherosclerosis, that it is effective both during the fatty streak and plaque phases, and that it may act by modulating T cell activity and/or antibody production. Therefore, immunomodulation may be an effective way to prevent and/or treat atherosclerosis. (J. Clin. Invest. 1998. 102:910–918.) Key words: antibody • atherosclerosis • immunoglobulin • T cells

Introduction

Atherosclerosis is associated with an activation of the immune system. Atherosclerotic plaques are infiltrated by macrophages and T lymphocytes, some of which immunospecifically recognize oxidized LDL (oxLDL) (1), which also accumulate in plaques. Autoantibodies to oxLDL are also found in atherosclerotic plaques (2, 3) together with activated components of the complement cascade (4, 5). In fact, the titer of auto-antibodies to oxLDL may correlate with progression of atherosclerosis in humans (6). However, it is still unclear to what extent interference with immunopathologic mechanisms may modulate the progression of the disease.

The recently developed apolipoprotein E knockout (apo E KO) mouse offers new opportunities to study pathogenetic mechanisms. These mice develop atherosclerotic lesions similar to those found in humans within weeks after birth (7). Although the atherosclerotic disease of these mice is based solely on a genetic disturbance of lipid metabolism, activated macrophages and CD4+ T cells infiltrate their lesions from the start (8) and autoantibodies to oxLDL similar to those found in humans appear in their peripheral blood (9). This suggests that the immune response is directly linked to the metabolic abnormality and that it is central to the pathogenesis of the disease. Congenital deficiency of macrophages, lymphocytes, and the Th1 effector pathway, by cross-breeding apo E KO mice with op/op mutant mice (10), T and B cell–deficient RAG-1 KO mice (11), and interferon-γ receptor KO mice (12), respectively, have resulted in reduction of lesions by 80, 40, and 60%, respectively. These results underline the importance of immune mechanisms in atherogenesis, but since they are based on genetic defects they do not clarify whether immunomodulation may be used to treat or prevent atherosclerosis.

Intravenous immunoglobulins (ivIg) are preparations of normal polyclonal IgG obtained from large pools of plasma from healthy donors. In addition to its use as substitution therapy for primary and secondary antibody deficiencies, ivIg is used increasingly in patients with autoimmune and systemic inflammatory diseases (13, 14). The mode of action of ivIg is still unclear and may involve both Fc- and V region–dependent mechanisms (for review see reference 15). Fc fragments of Ig may block Fc receptors on phagocytic cells of the reticuloendothelial system, inhibit antibody synthesis by B cells, modulate suppressor and helper functions of T cells, affect the production of cytokines by monocytes/macrophages, and interfere with complement-mediated tissue damage. IvIg have also been shown to contain antibodies that recognize idiotypic determinants on autoantibodies. Modulation of B and T cell function including cytokine production has been observed during ivIg treatment in rodent models of autoimmune diseases (16–19). Several of these mechanisms could conceivably be beneficial in atherosclerosis. For instance, ivIg could regulate the production of autoantibodies against oxLDL, modulate T cell functions in plaques, control complement activation, and affect phagocytosis and cytokine production by macrophages. In the present communication, we show that ivIg treatment of apo E KO mice inhibits the progression of atherosclerosis.

Methods

Biological preparations/immunogens/immunomodulators. IvIg (Sandoglobulin®) was a gift of the Central Laboratory of the Swiss Red Cross (Bern, Switzerland). When reconstituted for therapeutic use, Sandoglobulin® contains 50 mg/ml of IgG, 25–35 mg/ml sucrose, 6–10 mg/ml glucose, and 40–100 mM NaCl (osmolality 515 mosmol). For this study, a stock solution of 100 mg/ml (0.6 mM) of ivIg was prepared in serum-free RPMI containing L-glutamine, penicillin 50 U/ml, and streptomycin 50 mg/ml. The ivlg solution was dialyzed overnight against a DME medium to remove the stabilizing agents of the Sandoglobulin® preparation. The final osmolality of the medium was

1. Abbreviations used in this paper: Con A, Concanavalin A; ivIg, intravenous Ig preparation; KO, knockout; MDA, malondialdehyde; oxLDL, oxidized LDL.
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270 mosmol. As a control preparation, we used HSA (Amersham Pharmacia Biotech, Uppsala, Sweden) made as fraction V of human serum pooled from multiple donors and containing 10 mg/ml HSA dissolved in the above medium.

**Experimental protocol.** apo E KO mice were obtained from The Jackson Laboratory (Bar Harbor, ME). This strain was developed on a 129/J background and had been backcrossed six times to C57BL/6J mice (20). At 5–7 wk of age, male apo E KO mice were put on a “Western” diet (see below) and injected with ivlg or HSA as described in Table I. Two series of experiments were carried out. In the first one, the effect of ivlg on fatty streak development was assessed 1 and 2 mo after five ivlg injections. In the second series of experiments, the effect on the development of fibrofatty atherosclerotic plaques was analyzed after 4 mo of diet. The mice in the experimental groups received an intraperitoneal injection of 10 mg ivlg daily over a 5-d period. In the control groups, the mice received 10 mg HSA instead of ivlg. In the second series of experiments, a control group was added that was left completely untouched except for diet treatment. The animal experiments were approved by the regional ethical board.

**Diet.** The mice were placed on a Western diet (AB AnalyCen, Liköping, Sweden) that contained 16.7% protein and 21.2% fats. It was based on corn starch, casein, glucose, saccharose, cocoa butter, cellulose, minerals, cholesterol, and a vitamin mix. The cholesterol content was 0.15% and the energy content was 16.4 MJ/kg.

**Tissue processing.** Mice were killed by exsanguination under carbon dioxide anesthesia. The blood was collected and allowed to clot. Serum was separated by centrifugation and stored at $-20^\circ$C. The cholesterol concentration was assessed using a Unimate kit (Hoffman-La Roche, Basel, Switzerland) and a Cobas Mira analytical system (Hoffman-La Roche). The vasculature was perfused with sterile PBS. The spleen and the inguinal, mesenteric, axillary, and submandibular lymph nodes were harvested for in vitro assays and FACS® analysis.

In experiment 1, the whole aorta was fixed in PBS with 1% formaldehyde and stained with oil red-O after removal of the adventitia under a macroscope. Specimens containing the root, arch, thoracic, and abdominal aorta were immersed for 3 s in 60% isopropanol and stained for 10 min in a saturated oil red-O solution. They were rinsed in wa-

**Table I. Experimental Design**

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For abbreviations, please see text.

![Figure 1](https://doi.org/10.1172/JCI119892)

**Figure 1.** Effects of section angle on lesion estimates. (A) Computer simulation of an artery with an atherosclerotic lesion shows the effect of the section angle on the lesion estimate. Notice that the fraction area of lesion (lesion surface/vessel surface at cross-section) is relatively insensitive to angles at sectioning, whereas the absolute lesion surface is overestimated by $>15\%$ at a tilting of only $20^\circ$. (B) Regression between the volume fraction of the lesion with both the lesion surface and the fraction area of lesion (surface area of the lesion/surface area of the vessel).
and mounted en face on slides under coverslips within Kaiser’s glycerol gelatin (Merck, Darmstadt, Germany). The cryosections were counterstained with hematoxylin.

In experiment 2, the root of the aorta was dissected under a microscope and frozen in OCT embedding medium for serial cryosectioning covering 0.8 mm of the root. The first section was harvested when the first cusps became visible in the lumen of the aorta. Five sections of 10 μm in thickness were harvested per slide and thus 18 slides per mouse were prepared. Sections at 0, 200, 400, 600, and 800 μm distance from the cusps were stained with oil red-O, counterstained with hematoxylin, and mounted under coverslips.

**Quantitation of atherosclerotic lesions.** The oil red-O–stained en face specimens were analyzed at a magnification of 2.5. The image was captured on a microcomputer and a program developed in Quips language (Leica, Cambridge, UK) permitted a quantitation of the surface area of the lesion and the total surface area of the tissue sample. The ratio of these two surfaces provided the fraction area of the lesion. The oil red-O–stained cryosections were analyzed at a magnification of five using a similar program to quantify the cross-section surface area of the lesion and the cross-section surface area of the vessel. The fraction area of the lesion was calculated by dividing the surface of the lesion by the surface of the vessel. This allowed us to correct for errors caused by oblique sections that could otherwise lead to overestimation of the surface area occupied by a lesion (Fig. 1A). The most important value to estimate is actually the volume fraction of the lesions. This can be calculated as the volume of the lesion divided by the volume of the vessel. Knowing the distance between the sections and the cross-section surface area of both the lesion and the vessel, it is possible to compute the volume [S(surface area × distance)] of both the vessel and of the lesion and finally the volume fraction. As shown in Fig. 1A, the fraction area of the lesion was strongly correlated to the volume fraction of the lesion (r² = 0.88), whereas the surface of the lesion not normalized by the dimension of the vessel was less well correlated (r² = 0.75).

**Immunohistochemistry.** Cryosections were processed for immunohistochemistry as described previously (8) with minor modifications. In brief, CD4 (1:50), CD8 (1:50), and I-Ab (1:25) antibodies (PharMingen, San Diego, CA) were applied to acetone-fixed cryosections. After a washing step, the second antibody (a biotinylated rabbit anti–immunoglobulin antibody) and ABC alkaline phosphatase kit (Vector Laboratories, Burlingame, CA) were used. The enzyme was revealed with 0.2 mg/ml naphthol AS.MX (Sigma Chemical Co., St. Louis, MO), 0.002 mg/ml dimethyl formamide (Merck, Darmstadt, Germany) in 0.1 M Tris, pH 8.2, 1 mg/ml Fast Red TR salt (Sigma), and 0.33 mg/ml levamisole (Sigma). Sections were counterstained with hematoxylin.

**LDL preparations.** Human LDL (d = 1.019–1.063 g/ml) was obtained under sterile conditions by ultracentrifugation of human plasma collected from 10 donors. Malondialdehyde-modified LDL (MDA-LDL) were produced as described previously (21). In brief, MDA was generated by acidic reaction of malondialdehyde bis dimethylacetal tetramethoxypropane (Sigma) with HCl (4 M) for 10 min at 37°C. After neutralization of the MDA solution, 100 μM of MDA (0.5 M) was added to 1 mg of LDL. After 3 h of incubation at 37°C, free MDA was removed by running a PD10 Sephadex column (Pharmacia, Uppsala, Sweden). oxLDL was produced by an overnight oxidation with 5 mM CuSO₄, as described previously (1). The degree of oxidation was evaluated by a spectrofluorometric analysis (excitation at 350 nm and emission at a wavelength of 433 nm) as described by Steinbrecher (22). The fluorescent intensity of the oxLDL preparations was always > 0.5 as compared with the fluorescent intensity of native LDL < 0.2.

**ivIg and anti-ivIg antibodies in recipient mice.** An ELISA was set up to quantitate the titer of circulating levels of human ivIg in mouse sera. 96-well flat-bottomed plates (Corning Costar Corp., Cambridge, MA) were coated with mouse sera and incubated overnight at 4°C. The plates were washed three times with PBS-0.05% Tween, blocked with PBS-5% BSA, and washed with PBS-Tween. An anti-human IgG, Fcγ antibody conjugated to alkaline phosphatase (Southern Biotechnology, Birmingham, AL) was added at 1:1,000 and the plates were incubated for 1 h at 37°C under gentle agitation. After PBS-Tween washes, 25 μl of substrate solution was added (a 5 mg tablet of p-nitrophenyl phosphate [Sigma] in 5 ml 0.1 M NaCl, 5 mM MgCl₂, 0.1 M Tris-HCl). Immunoreactivity was determined by absorbance at 405 nm using an automated plate reader (Labsystems Multiskan, Helsinki, Finland). Levels of remaining ivIg in mouse sera were calculated by subtracting the absorbance of controls without sera.

The levels of antibodies to ivIg in mouse sera were assessed by coating 96-well plates with 50 μl ivIg (10 μg/ml). After an overnight incubation at 4°C and washing and blocking steps, mouse sera were added at 1:10 dilutions and the plates were incubated for 1 h at 37°C. After washing and blocking steps, the titer of anti-ivIg antibodies was revealed by using an anti-mouse antibody conjugated to alkaline phosphatase as described above. Titers of antibodies to ivIg were calculated by subtracting the absorbance of controls without serum.

**Autoantibody titers to nLDL, MDA-LDL, and oxLDL.** The titer of autoantibodies was measured by ELISA as described above. Plates were coated overnight with native LDL, MDA-LDL, and oxLDL at 10 μg/ml. Mouse sera were added at 1:100 for the IgM titers and at 1:10 for IgG1 and IgG2a titers. Finally, alkaline phosphatase–conjugated detecting antibodies were plated at the following dilutions: anti–mouse IgM (Southern Biotechnology): 1:1,000; G1-6.5 anti–mouse IgG1 and R19-15 anti–mouse IgG2a (PharMingen): 1:250. The detection step was as described above. To check whether variations observed in the titer of specific immunoglobulins were due to variations of the overall concentration of immunoglobulin isotype, the titers of total IgG1 and IgG2a were measured using the following capture/detecting antibody pairs: anti–mouse IgG1 (Serotec, Oxford, UK)/HRP-conjugated anti–mouse IgG1 (Serotec); anti–mouse IgG2a (R12-4, PharMingen)/ALP-conjugated anti–mouse IgG2a (R19-15; PharMingen). The HRP-conjugated antibodies were revealed by adding one 5 mg tablet of o-phenylenediamine dihydrochloride (Sigma) to 10 ml of substrate buffer with 4 μl H₂O₂, and absorbance was determined at 450 nm. As controls for autoantibodies, the antigen or the serum was omitted, while the controls for total IgG1 and IgG2a omitted the capture antibody or the serum. In all cases, titers of antibodies were calculated by subtracting the absorbance by values obtained in the controls.

**Cell culture.** Spleens and lymph nodes were processed separately but were pooled in each group of animals. Lymph node cells and spleen cells were prepared by meshing the tissues on 100-μm nylon filters followed by a series of washing steps in DME with 5% FBS.
Figure 2. Effect of ivIg on fatty streak formation. apo E KO mice were injected with HSA or ivIg and then fed a Western diet for 2 mo (experiment 1). En face specimens from the aortic root were stained with oil red-O and photographed at a magnification of 2.5. (A) HSA treatment. Notice large lesion in the left part of the root. (B) ivIg-treated mouse. No large lesions are found.

Figure 3. ivIg reduces fatty streak formation. apo E KO mice were injected with HSA or ivIg and then fed a Western diet for 2 mo (experiment 1). The lesion area (percentage of total surface area) was determined in oil red-O-stained en face specimens of the aorta. Lesion surface and vessel surface were determined using the Leica Quantimet image analysis system. Fraction area of lesion (percentage) is lesion surface/vessel surface. Means±SEM. †Significantly different from HSA treatment. P < 0.01.

(DME-5%). Red blood cells were lysed in the spleen cell preparations with ACK lysing buffer (0.15 M NH₄Cl, 1 mM KHCO₃, 0.1 mM Na₂EDTA, pH 7.3). 300,000 cells/well were plated into 96-well flat-bottomed plates in the presence of the following stimuli: Concanavalin A (Con A, 2.5 μg/ml; Sigma), oxLDL and native LDL (3.125 and 6.25 μg/ml), and ivIg and HSA (20 mg/ml). Each type of stimulation was performed in duplicate. The plates were incubated for 2 d at 37°C. 0.5% CO₂. 50 μl of supernatant was removed from each well to check the production of IL-2 and IL-4 by a CTL-L2 bioassay and were replaced by 50 μl of DME-5% containing 1 μCi of [³H]thymidine. After an additional day of incubation, the cells were harvested and transferred to glass fiber filters (Wallac, Turku, Finland) using an Inotech (Wohlen, Switzerland) cell harvester. The filters were dried at 60°C for 1 h, sealed in a sample bag with melt-on scintillator sheets (Wallac), and placed in a Wallac Microbeta β-counter. Results expressed as counts per minute reflect the incorporation of [³H]thymidine into DNA and thus the proliferation of the cells. For the CTL-L2 assay, 5,000 CTL-L2 cells were added to the 50 μl of supernatant and 1 μCi of [³H]thymidine was added after 1 d of incubation. After 18 h, the cells were harvested as described above.

FACS® analysis. A fraction of the cells prepared for cell culture was used for a FACS® analysis to check the different cells populations and their activation state. Cells were stained for 30 min at 4°C using purified anti-CD3 and anti-CD25, biotin-conjugated anti-CD4 and anti-CD19, and FITC-conjugated anti-CD8 and anti-CD69 antibodies (PharMingen). Cells were washed and analyzed with a FACS-Calibur® (Becton Dickinson, Mountain View, CA) flow cytometer.

Statistical analysis. Results are expressed as means±SEM. Data were analyzed by one-factor ANOVA using Statview 4.1 software (Abacus Concept Inc., Berkeley, CA). Differences between groups were considered significant if P < 0.05. Regression coefficients were obtained by the least squares method.

Results

IvIg treatment. 7-8-wk-old apo E KO mice were injected intraperitoneally with 10 mg human ivIg (Sandoglobulin®) or HSA daily for five consecutive days and then fed an atherogenic diet for 2 or 4 mo. The injected ivIg was detectable in mice sera after 1 mo but the levels were close to background at 2 and 4 mo after treatment, indicating clearance from the plasma compartment with kinetics similar to that of endogenous Ig. A weak anti–human Ig reactivity could be detected plasma bodies (PharMingen). Cells were washed and analyzed with a FACS-550® (Becton Dickinson, Mountain View, CA) flow cytometer.

Figure 3. Effect of ivIg on fatty streak formation. apo E KO mice were injected with HSA or ivIg and then fed a Western diet for 2 mo (experiment 1). En face specimens from the aortic root were stained with oil red-O and photographed at a magnification of 2.5. (A) HSA treatment. Notice large lesion in the left part of the root. (B) ivIg-treated mouse. No large lesions are found.

Table III. Effect of ivIg on the Inflammatory Cell Infiltrate of the Lesions

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</tr>
<tr>
<td>CD8⁺</td>
<td>3.2±0.2</td>
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<tr>
<td>I-A⁺</td>
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Lesions of the aortic root were analyzed. Data were obtained by dividing the number of positively stained cells by all hematoxylin-stained cells inside the internal elastic lamina. Three random microscopic fields were analyzed at ×400.
smaller lesions were detected in the descending thoracic and abdominal parts of the aorta (Figs. 2 and 3). The surface area covered by fatty streak lesions was quantified in oil red-O–stained samples and specimens from ivIg-treated mice were compared with HSA-treated controls. In mice treated with ivIg, the fraction area of lesion was reduced by 35% in the aortic root and 32% in the arch (Figs. 2 and 3).

**Effect of ivIg on atherosclerosis.** In a second series of experiments, mice were kept on the Western diet for 4 mo to permit the formation of fibrofatty atherosclerotic plaques (23). ivIg and HSA injections were given either at the start of feeding, i.e., at 7 wk of age, after 2 mo on the diet, i.e., at 15 wk of age, or both at 7 and 15 wk of age. An additional control group was left untreated and all mice were killed at 22 wk of age (i.e., after 4 mo on the diet).

This treatment resulted in extensive lesion formation in both HSA-treated and untreated mice (Fig. 4). Both these groups exhibited lesion formation covering 12–15% of the cross-sectioned aortic root (Fig. 5). Mice treated with ivIg at 0 and 2 mo of diet (the ivIg[0/2-4] group) or only at 2 mo (ivIg[2-4]) showed a 50% reduction of lesion formation, with only 6–8% of the cross-section area covered by lesions (Figs. 4 and 5). In all groups, lesions were fibrofatty plaques with lipid-rich core regions covered by fibrous cap with smooth muscle cells (Figs. 4 and 6). The cellular composition of the lesions did not differ significantly between the groups. In particular, the inflammatory cell infiltrate as assessed by the number of CD4+, CD8+, or I-A^b^ cells divided by the number of hematoxylin-stained cells was not significantly different between the different groups (Fig. 6, Table III, and data not shown). In contrast, ivIg injections at the beginning of the diet treatment did not confer protection against atherosclerosis (Fig. 5).

**Effect of ivIg on immune responses.** ivIg treatment did not affect the relative proportions of CD3^+^, CD4^+^, or CD8^+^ T cells, CD19^+^ B cells, or the proportion of activated cells expressing CD25 or CD69, as determined by FACS® analysis of spleen and lymph node cell suspensions (data not shown). However, spleen and lymph node cells of ivIg-treated apo E

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**Figure 4.** Effect of ivIg on the formation of fibrofatty plaques. apo E KO mice were fed a Western diet for 2 mo, injected with ivIg or HSA, and then continued on the diet for another 2 mo (experiment 2). The aortic root was cryosectioned and stained for lipids with oil red-O. Magnification, ×5. (a–e) Five consecutive sections from HSA-treated mice covering the root of the aorta, taken every 200 μm from the origin. (f–j) Five consecutive sections from ivIg-treated mice. Lesions of ivIg-treated mice are smaller and cover a smaller fraction of the inner circumference of the aortic root than those of HSA-treated control mice.

**Figure 5.** ivIg treatment inhibits formation of fibrofatty plaques. apo E KO mice were fed a Western diet for 4 mo (experiment 2) and were injected with HSA or ivIg immediately before the start of diet treatment (a), after 2 mo on the diet (b), or both at 0 and 2 mo (c). A fourth group of mice (d) received the diet but no injections. Lesion surface and vessel surface were determined using the Leica Quantimet image analysis system. Fraction area of lesion (lesion surface/vessel surface) was measured on cryo-sections from the aortic root. Means ± SEM. *Significantly different from HSA-treated group, P < 0.01.
KO mice exhibited a lower basal proliferation rate than those of HSA-treated or untreated apo E KO mice (Figs. 7 and 8). T cells of ivIg-treated mice were able to respond to Con A by proliferation, indicating that they were anergized rather than eliminated (Figs. 7 and 8). This was supported by the finding that addition of ivIg to spleen cells from control mice reduced T cell proliferation in response to Con A. The data obtained on the proliferation were corroborated by IL-2/IL-4 production assays (data not shown).

IgM antibodies to oxLDL were significantly reduced 2 mo after ivIg treatment, implying that the B cell response was also hampered by the treatment (Fig. 9). This effect was no longer discernible after 4 mo (data not shown). No effect were observed on IgG (Fig. 9), IgG1, or IgG2a antibodies to oxLDL (data not shown).

Discussion

This study shows that ivIg treatment inhibits atherosclerosis in cholesterol-fed apo E KO mice. Both the development of fatty streaks and the progression to fibrofatty atherosclerotic plaques could be inhibited by this treatment. This obviously suggest that similar ivIg-modulated mechanisms operate in both stages of the disease.

To reduce fatty streaks, it was sufficient to inject ivIg once at the start of treatment, whereas protection against atherosclerotic plaques required ivIg to be administered after 2 mo on the atherogenic diet. Since ivIg was cleared from the circulation after 1–2 mo, these data suggest that protection depends on the presence of the infused immunoglobulins in the animal. The lack of protection against atherosclerosis by the fatty streak–preventing protocol (i.e., ivIg injections at the beginning of treatment) suggests that prevention of fatty streaks may not be sufficient to prevent subsequent formation of atherosclerotic plaques. This may reflect a capacity of ivIg to inactivate but not eliminate atherogenic factors. If this is the case,
such factors would presumably be reactivated when ivIg disappears from the circulation. Alternatively, different factors present in the ivIg preparation act in the fatty streak phase and the phase of plaque growth.

The precise atheroprotective mechanism(s) conferred by ivIg treatment have not been identified but it is interesting that the atheroprotective effect was paralleled by effects on T and B cells. T cells of ivIg-treated mice exhibited a lower basal proliferation rate but could be activated to proliferate by addition of the polyclonal mitogen, Con A. This suggests that ivIg inhibits T cell activation. Such an interpretation was supported by the observation that in vitro treatment of spleen cell cultures with ivIg blocked T cell activation. If T cells infiltrating fatty streaks and atherosclerotic plaques are also anergized, this would reduce immune activation in the artery. Since most plaque T cells are proinflammatory Th1 effector cells, a conceivable consequence is reduced macrophage activation. The recent finding that elimination of Th1 signaling by targeted disruption of the interferon-γ receptor gene inhibits atherosclerosis in apo E KO mice by 60% (12) is in agreement with such an interpretation. Interestingly, ivIg preferentially down-regulates T cell cytokine secretion, whereas monokine production is less affected (24).

In addition to its effect on T cells, ivIg also reduced autoantibody titers to oxLDL species. Since a significant reduction was found for IgM but not IgG antibody classes, this effect was probably exerted directly on the B cell. A drop in the titer of autoantibodies after administration of ivIg has been reported previously in several autoimmune manifestations (14). Recently, the B cell/humoral immunity system was shown to play a major role in the pathogenesis of transplant arteriopathy (25) but its role in atherosclerosis has remained obscure. Few B cells are found in developing atherosclerotic plaques but advanced plaques of fat-fed rabbits were reported to contain significant amounts of Ig-secreting plasma cells (26). Anti-oxLDL antibody titers appear to reflect disease activity but the pathogenic mechanisms of these antibodies remain unclear. While IgM antibodies do not mediate rapid antigen elimination via Fc receptors, their interaction with oxLDL in the arterial intima could initiate complement activation. This may secondarily lead to cell death, proteolysis, release of chemotactic agents, and local inflammation. Therefore, a reduction of the IgM anti-oxLDL titer might dampen vascular inflammation.

In addition to the immunomodulatory effects of ivIg, the possibility that it confers protection against atherosclerosis by transfer of specific antibodies remains an exciting scenario. Microorganisms have been implicated recently in the pathogenesis of atherosclerosis and there is strong epidemiological evidence for an association between *Chlamydia pneumoniae* and cardiovascular disease in humans. The ivIg preparations contain thousands of different antibody specificities, which can neutralize a large series of pathogens. However, it is unlikely that more than a small proportion of these microorganisms would affect vascular disease in both humans and rodents. However, antibodies directed against several cell surface molecules of relevance in immunomodulation have also been identified in ivIg preparations (27–30). In addition, soluble
forms of CD4, CD8, and class I and class II molecules have been identified in ivIg (31, 32). Therefore, a fractionation of ivIg, and/or the use of specific antibodies in experimental models of atherosclerosis, should shed light on the role of specific antibodies in the atheroprotective action of ivIg.

In conclusion, we have shown that treatment of atherosclerosis-prone apo E KO mice with five injections of ivIg protects against fatty streak development over a 2-mo period of cholesterol feeding. ivIg reduces IgM antibodies to modified LDL. Sera were obtained from apo E-deficient mouse for the first time, that treatment by immunomodulation inhibits the formation of fibrofatty plaques in these mice. ivIg treatment was associated with anergization of T cells and reduction of IgM anti-oxLDL antibodies, but the precise mechanism by which ivIg confers protection against atherosclerosis remains unclear. However, the current results show, for the first time, that treatment by immunomodulation inhibits atherogenesis. The ivIg treatment protocol presented here is unlikely to be useful in preventing cardiovascular disease in humans, but the results are encouraging for investigators exploring immune-based therapy as a way to treat atherosclerosis.

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