Effects of cyclosporin on collagen induced arthritis in mice

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SUMMARY We have studied the effect of the immunosuppressive agent cyclosporin on collagen induced arthritis in mice. Cyclosporin, when given prophylactically, was capable of suppressing the development of collagen induced arthritis and the immunological response to native type II collagen in a dose dependent manner. Furthermore, treatment with cyclosporin, started on the same day as the booster injection with type II collagen, also resulted in inhibition of development of arthritis and of immunity to collagen. These findings suggest that the time of a booster injection, three weeks after the initial immunisation, might be still within the induction phase of arthritis since reinoculation is required to produce a high incidence of arthritis in mice. In addition, therapeutic treatment with cyclosporin did not affect the clinical course of the disease or the immune response to collagen.

Immunisation with native type II collagen induces inflammatory polyarthritis and immunity to collagen in rats and susceptible strains of mice.\textsuperscript{1-6} The disease, which has been shown to have similarities to human rheumatoid arthritis in many respects,\textsuperscript{7} can be transferred by immune sera from arthritic rats to naive mice\textsuperscript{8} and rats.\textsuperscript{9} Furthermore, we reported previously that a serum concentrate from arthritic rats is capable of inducing arthritis in cyclosporin treated, type II collagen tolerant rats\textsuperscript{10} and congenitally athymic nude rats.\textsuperscript{11} These findings suggest that arthritis in rats and mice might be caused by similar mechanisms,\textsuperscript{8} and that arthritis could be induced by humoral immunity in the absence of cell mediated immunity.\textsuperscript{11} Although pathogenetic mechanisms and the histopathology of collagen induced arthritis in mice are similar to those seen in rats,\textsuperscript{3,5} there are definite discrepancies in the immune response to collagen between these two species.\textsuperscript{3,5} In addition, a booster injection and use of complete Freund's adjuvant combined with type II collagen at the time of the primary sensitisation are required to produce a high incidence of arthritis in mice.\textsuperscript{2} The development of arthritis and immune response to collagen are suppressed by pretreating the rat with immunosuppressive agents.\textsuperscript{12-16} The effect of any immunosuppressants on collagen induced arthritis in mice has not been previously reported.

Cyclosporin is a fungal metabolite with potent immunosuppressive properties both in man and animals. Treatment of animals with cyclosporin results in the inhibition of many thymus dependent functions, such as allograft rejection, graft v host rejection, and the development of experimental autoimmune diseases.\textsuperscript{15-20} Recently, we reported that treatment with cyclosporin, begun on the same day as type II collagen immunisation, suppressed the development of collagen induced arthritis in the rat, and that a significant enhancement of arthritis was observed when cyclosporin treatment was started during the immediately preclinical phase of arthritis or after the disease onset.\textsuperscript{13}

In this paper we are investigating the effect of cyclosporin on collagen induced arthritis in mice with three different regimens: (a) started on the same day as the primary immunisation, (b) only after a booster injection, and (c) on the established disease. We have found that treatment with cyclosporin, starting concurrently with either the initial or the booster sensitisation, can inhibit the development of arthritis and the immunological response to collagen, but that cyclosporin does not affect the established disease.
Materials and methods

MICE
DBA/1 mice of both sexes, originating from the Jackson Laboratory, Bar Harbor, Maine, were obtained from Seiwa Experimental Animals Ltd, Fukuoka, Japan. They were allowed at least one week to adapt to their environment and were used at 8-10 weeks old, weighing 16-24 g at the start of the present experiment. The same number of mice of each sex were used for each experiment. Animals were fed ad libitum with standard laboratory food and water.

PREPARATION OF TYPE II COLLAGEN AND PRODUCTION OF COLLAGEN INDUCED ARTHRITIS
Type II collagen was isolated and purified from bovine articular cartilage as previously described.1 The purity was assessed as described elsewhere.15 Lyophilised type II collagen was dissolved in 0.1 M acetic acid at a concentration of 2 mg/ml (g/l). Complete Freund's adjuvant (CFA) was prepared by adding dried, heat killed Mycobacterium butyricum (Difco Laboratories, Inc., Detroit, MI), which was finely powdered with a mortar and pestle, to incomplete Freund's adjuvant (Difco Laboratories, Inc.) at a concentration of 2 mg/ml. Equal volumes of collagen solution and CFA were emulsified by a homogeniser (Polytron PT 10-35; Kinematica, Lucerne, Switzerland) and kept cold in an ice bath. Collagen induced arthritis was produced by intradermal injection of 0.1 ml of the cold emulsion at one site into the base of the tail. Mice were boosted intraperitoneally with 100 μg of type II collagen in 0.017 M acetic acid on day 21.

TREATMENT WITH CYCLOSPORIN
Cyclosporin (kindly provided by Sandoz Ltd, Basel, Switzerland) was dissolved in pure olive oil at a concentration of 20 mg/ml by heating in a water bath to 65°C. It was freshly prepared every three days. Cyclosporin was given subcutaneously under light ether anaesthesia on the days and at the doses indicated in the text. The dose was adjusted according to the daily body weight. Control mice, while immunised with the same amount of type II collagen, received intradermal injection of olive oil alone. Cyclosporin treated mice and control mice were handled identically for administration of the agent and the solvent.

ASSESSMENT OF ARTHRITIS
Mice were examined daily for seven weeks after the injection of type II collagen to record the day of onset and the severity of arthritis. The lesions of the four paws were each graded from 0 to 4 according to the increasing extent of periarticular erythema, swelling, and joint deformity, as described previously.8 21 The maximum possible score was 16.

IMMUNOASSAY OF IgG ANTIBODY LEVELS TO TYPE II COLLAGEN
Blood was routinely collected by retro-orbital bleeding with a capillary tube on days 21 and 47. Serum samples were removed, heat inactivated at 56°C for 30 min, and stored at -80°C until used. Serum antibodies to type II collagen were measured by the enzyme linked immunosorbent assay technique. The methods used were adapted from those of Vollero et al.22 Briefly, microtitre plates were coated with 100 μl of type II collagen at a concentration of 25 μg/ml (mg/l) in coating buffer pH 9.6 and incubated overnight at 4°C. Before assay plates were washed three times with phosphate buffered saline (PBS) containing 0.05% Tween 20. Coating buffer containing 1% bovine serum albumin (BSA) was used to avoid non-specific adsorption. The cuvettes were washed three times before the addition of a 100 μl aliquot of mouse test serum diluted with PBS-Tween containing 1% BSA. The preparation was incubated for two hours at room temperature. After washing three times with PBS-Tween 100 μl of 1:1000 dilution of alkaline phosphatase conjugated anti-mouse IgG (Miles Laboratories, Inc.) was added. After incubation for three hours at room temperature the plates were again washed three times with PBS-Tween, and 100 μl of p-nitrophenyl phosphate substrate (Sigma Chemical Co., St Louis, Mo) dissolved in 10% diethanolamine was added. After incubation at room temperature for 30 min the reaction was stopped by adding 20 μl of 3 M NaOH to each well, and the reaction product was measured by absorbance at 410 nm by an automated device (Dynatech Laboratories, Inc., Alexandria, VA). The quantity of IgG anti-type II collagen antibody was expressed as mg/100 ml of serum by comparison with standard curves obtained from purified mouse anti-type II collagen antibody control. Mouse anti-type II collagen antibody was purified by affinity chromatography on cyanogen bromide Sepharose 4B using procedures provided by the supplier (Pharmacia Fine Chemicals, Piscataway, NJ). The procedures used have been described in detail elsewhere.23

RADIOMETRIC EAR IMMUNOASSAY
Intradermal, delayed type hypersensitivity (DTH) reactions to type II collagen were quantified on day 48 by a radiometric ear immunoassay. Our assay was a modification of that previously described by Vadasz et al24 and Rogers et al.25 Native type II collagen was...
dissolved at a concentration of 1 mg/ml in 0.1 M acetic acid at 4°C for 24 h and then dialysed against cold 0.05 M calcium acetate. 10 µl of the test solution was injected intradermally into the right pinna and the same volume of 0.05 M calcium acetate alone into the left. A 2 µCi pulse of $[\text{\textsuperscript{5}}\text{I}]$Iodo-2'-deoxyuridine ($[\text{\textsuperscript{125}}\text{I}]$UdR, Amersham International plc, Buckinghamshire, England) was given intravenously 10 h later. The mice were killed 16 h after the injection of $[\text{\textsuperscript{125}}\text{I}]$UdR, and the ears were cut off at the hair line. The radioactivity was determined in an automatic gamma counting system (Gamma Trac 1191, Tracor Analytic Inc. Ill.). The radiometric ear index (REI) was expressed as the ratio of radioactivity in the right and left ear samples of an individual mouse. REIs exceeding the mean±2 SD of values derived by testing 24 naive DBA/1 mice with type II collagen were considered to be significant responses. By these criteria REI > 1.4 indicated DTH to bovine type II collagen.

**STATISTICS**
Continuous variables were analysed by their group means (Student's t test) and dichotomous variables by their proportionate group frequencies ($\chi^2$ test). P Values less than 0.05 were considered to be statistically significant.

**Results**

**INCIDENCE OF COLLAGEN INDUCED ARTHRITIS IN DBA/1 MICE**
To find the incidence and day of onset of arthritis in DBA/1 mice 35 mice of each sex were immunised on day 0 with 100 µg of type II collagen in CFA and challenged on day 21 with intraperitoneal injection of 100 µg of type II collagen without adjuvant. The results are shown in Fig. 1. An inflammatory polyarthritis developed in 97.1% of mice. The incidence of arthritis did not depend on the sex of the mice (100% in male mice and 94.2% in female, not significant). The day of onset of the disease was variable: between 23 and 46 days after the initial sensitisation, with a mean of 30-3 days. Sixty of 70 animals developed arthritis by the end of five weeks after primary immunisation. Therefore, therapeutic treatment with cyclosporin was started on day 35. The mean maximum severity of arthritis in male mice was similar to that seen in female mice (7.2±3.3 and 6.9±2.8 respectively).

**Cyclosporin Treatment**

**Effect from the day of the primary immunisation with type II collagen**
Mice received 14 daily subcutaneous injections of various doses of cyclosporin or olive oil starting on the day of the primary immunisation with type II collagen. Treatment with cyclosporin prevented the development of collagen induced arthritis in a dose dependent manner despite a booster injection (Table 1). A complete prevention of the disease was

![Fig. 1](http://ard.bmj.com)  
*Fig. 1  The incidence and day of onset of arthritis in DBA/1 mice. Male (closed circle) and female (open circle) mice were immunised on day 0 with type II collagen in complete Freund's adjuvant and challenged on day 21 with an intraperitoneal injection of type II collagen alone.*

**Table 1  Effect of cyclosporin treatment (days 0–13) on the development of collagen induced arthritis**

<table>
<thead>
<tr>
<th>Cyclosporin dose (mg/kg)</th>
<th>Olive oil</th>
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<tbody>
<tr>
<td></td>
<td>100</td>
</tr>
<tr>
<td>Incidence of arthritis</td>
<td>0/18 (0%)***</td>
</tr>
<tr>
<td>Arthritic index†</td>
<td>—</td>
</tr>
<tr>
<td>Day of onset‡</td>
<td>—</td>
</tr>
<tr>
<td>Antibody level (mg/dl)$§$</td>
<td>21 days</td>
</tr>
<tr>
<td></td>
<td>47 days</td>
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<tr>
<td>DTH skin reaction§</td>
<td>1.2±0.1***</td>
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</tbody>
</table>

*p<0.05 v the oil group; **p<0.01 v the oil group; ***p<0.001 v the oil group.
†Expressed as the mean of maximum arthritic indices±SEM.
‡Based on arthritic mice only (mean±SEM).
§Expressed as the mean±SEM.
achieved by treating the animals daily with 100 mg/kg/day of cyclosporin, whereas no clear effects were produced by 25 mg/kg/day of the drug. The effects were accompanied by the inhibition of REI to collagen measured on day 48 and of anti-type II collagen antibody production measured on days 21 and 47. These findings are in accord with our previous report, which suggested that helper T cell function is mandatory for the production of anti-type II collagen antibody. 15

Effect from the day of a booster injection of type II collagen
Mice were injected daily with cyclosporin at a dose of 100 mg/kg for two weeks from the day of a booster injection. It is apparent from a number of in vitro experiments that cyclosporin exerts its effect at a very early stage of T cell differentiation. 26-29 Cyclosporin, however, again produced statistically significant suppression of the development of arthritis (Table 2). Moreover, serum antibodies to collagen measured on day 47 and REI to collagen measured on day 48 in cyclosporin treated mice were significantly reduced, though no difference was observed in serum antibodies to collagen measured at the time of a booster injection. A small number of cyclosporin treated animals developed disease, and in those that did the disease was less severe. The onset of the disease in cyclosporin treated mice, however, was not delayed.

Effect on the established disease
The response of the established disease to cyclosporin treatment was studied. Only arthritic mice received daily cyclosporin therapy for 10 days beginning 35 days after primary immunisation. Cyclosporin did not affect the course of the disease (Fig. 2), nor did its administration alter the immunological response to collagen in this regimen (Table 3).

We did not observe a significant decrease in weight or any other toxic signs in animals receiving this relatively high dosage of cyclosporin in any of the three experiments.

Discussion
The results presented in this paper show that cyclosporin inhibited the development of collagen-induced arthritis in mice in a dose dependent manner when the agent was given prophylactically. This effect was accompanied by a marked suppression of antibody production and DTH skin response to type II collagen. These findings confirm our previous study, which has shown that a 14 day course of cyclosporin treatment, begun on the same day as type II collagen immunisation, suppressed the development of arthritis and the immunological response to type II collagen in rats. 15 In addition, Brahn and Trentham have shown that the administration of antithymocyte serum, started one day before sensitisation, also suppressed the development of arthritis in rats. 30 Moreover, we and other investigators have found that immunisation with
type II collagen did not induce arthritis or immunity to collagen in congenitally athymic nude rats. On the basis of these findings Brahn and Trentham postulated that collagen induced arthritis is a T cell dependent disease. We have recently shown, however, that arthritis can be passively transferred with a serum concentrate to congenitally athymic nude rats and to cyclosporin treated, type II collagen tolerant rats. These findings indicate that arthritis may be inducible by humoral immunity alone. Taken together, the present results can be explained by the suppressive effect of cyclosporin on helper T cells, whose defined function is manifesting the induction of anticolonagen antibody formation.

On the basis of the data reported previously cyclosporin acts early in the immune response after exposure to an antigen and affects primary immune events, and treatment is found to be most effective if started simultaneously with an antigen presentation. Unique to this study, however, is the demonstration that cyclosporin treatment began concurrently with a booster injection, also significantly suppressed the development of arthritis and the immune response to collagen, though the antibody levels at the time of a booster challenge were very low but detectable. Cyclosporin treated mice that did develop arthritis had less severe disease. In contrast with these findings, we have reported that the treatment initiated during the immediate preclinical phase of arthritis in rats was associated with a significant enhancement of the severity of the disease. The difference in results of the effect of cyclosporin on collagen induced arthritis between rats and mice is not clear; however, histological changes in rats are already evident in the synovium five days after immunisation and treatment of rats with cyclosporin started during the immediate preclinical phase might be too late to improve the course of the disease. This explanation is supported by the ineffectiveness of antithymocyte serum on collagen induced arthritis in rats when the administration of antithymocyte serum is started five days after immunisation. On the other hand, a booster injection is required for the development of arthritis in mice, and treatment with cyclosporin initiated on the same day as a booster injection is sufficiently early to suppress the development of arthritis. This preventive effect of cyclosporin on the development of arthritis in mice suggests that the time of a booster injection, three weeks after immunisation, might be within the induction phase of arthritis.

In the last experiment we showed that cyclosporin did not affect the severity of the disease, nor did it alter antibody response and DTH skin reaction, when given therapeutically. It has been previously reported that cyclosporin had a beneficial effect in a patient with relapsing polychondritis and reversed dermatological plaques in several patients with psoriasis. In addition, Henderson et al implicated cyclosporin as an antiarthritic agent by investigating only its prophylactic effect on the development of collagen induced arthritis in rats. We are concerned, however, by the therapeutic use of cyclosporin based on the data in the present work and in our previous experiments.

In conclusion, cyclosporin was only capable of suppressing the development of collagen induced arthritis in mice when the treatment was started on the same day as either the initial or the booster sensitisation. The results suggest that the time of a booster injection might be within the induction phase of arthritis. Its therapeutic use in mice, i.e. after the onset of clinical features, did not affect the clinical course of arthritis or the immune response to collagen.

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References


