Cellular immunity to cartilage link protein in patients with inflammatory arthritis and non-arthritic controls

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Abstract

Objectives—To determine if increased T cell responses to articular cartilage link protein have any correlation with rheumatoid arthritis (RA), and if RA patients with increased responses to link protein also respond to a 17 amino acid peptide covering the 'arthritogenic' epitope in mycobacterial hsp65 which is homologous with link protein.

Methods—The reactivity of T cells from both peripheral blood and synovial fluid, to highly purified human cartilage link protein, hsp65, the 17 amino acid peptide, and bovine type II collagen was determined in patients with RA and non-arthritic controls, by measuring the rate of mononuclear cell proliferation in the presence and absence of antigen.

Results—Using peripheral blood mononuclear cells (PBMC), significant reactivity (stimulation index (SI) > 1.5) to link protein was found in 12 of 46 RA patients (26%), but in only four of 44 controls (9%). A greater proportion of RA patients (eight of 17: 47%) were reactive to link protein when mononuclear cells from synovial fluid were tested. SI values, however, were generally low (0.5–3.1) and only one patient showed a PBMC response above a reference range of values calculated from the logarithmic values of the normal control population. No reactivity was observed against a 17 amino acid synthetic peptide including the arthritogenic epitope from the mycobacterial hsp65 to which T cell clones isolated from rats in the adjuvant arthritis model react. However, eight of nine RA patients and all of seven controls reacted to the intact hsp65.

Conclusion—It remains unclear if T cell responses to link protein are involved in the pathogenesis of RA, but it is unlikely that T cells specific for the sequence homologous with the arthritogenic epitope in hsp65 are present in RA patients.

Despite much investigation, the extent to which cellular immune responses within the joints of patients with rheumatoid arthritis (RA) contribute to the pathogenesis of the disease remains unclear. It has been suggested that the most critical mechanism of joint destruction is cytokine secretion by monocytes. However, a review has presented evidence for the importance of T cells in the immunopathogenesis of RA, and concluded that they were central to the initiation and maintenance of synovitis in RA.

Potential autoantigens under investigation include components of articular cartilage matrix which consists of proteoglycan aggregates within a network of collagen fibrils. In the proteoglycan aggregate, the binding of aggrecan monomers to hyaluronan is stabilised by link protein. The structure, function, and distribution of link protein and structurally related proteins have been reviewed by Neame et al. The structure of proteoglycan aggregates changes with aging, and there is a gradual increase in the heterogeneity of proteoglycan monomers. In addition, proteolytic cleavage results in release of glycosaminoglycan (GAG)-rich fragments of aggrecan monomers into the synovial fluid. Proteolysis may result in the release into the synovial fluid of some link protein and the hyaluronan binding region of aggrecan; these tend to accumulate in the cartilage matrix in association with hyaluronan.

In certain animal models, arthritis can be induced by type II collagen and the less abundant type IX and XI collagens. Arthritis has been demonstrated after intra-articular injection of heterologous proteoglycans in rabbits and chondroitin sulphate depleted human fetal cartilage proteoglycan (HFPG) was shown to give rise to polyarthritids and ankylosing spondylitis in female Balb/c mice. In both collagen induced arthritis and proteoglycan induced arthritis, it is possible to induce secondary arthritis by adoptive transfer of cells from immunised donors to naive syngeneic recipients. More recently, two T cell lines and two T cell hybridomas reactive with HFPG were raised from arthritic Balb/c mice after immunisation with chondroitin sulphate depleted HFPG. One of the T cell lines and the two T cell hybridomas recognised homologous sequences in the G1 domain of adult human aggrecan and link protein, which suggests that link protein is a potential autoantigen. Furthermore, in the rat adjuvant arthritis model, arthritogenic T cell clones have been developed which recognise an epitope on a mycobacterial heat shock protein (hsp65), having some homology with link protein. These clones are also capable...
of eliciting delayed type hypersensitivity responses to preparations of articular cartilage extracts rich in link protein.

Increased T cell responses to type II collagen and cartilage proteoglycan have been described in some patients with RA, although other groups failed to detect antiproteoglycan responses. Recently, we have shown that RA patients have increased T cell proliferative responses to deglycosylated human articular cartilage aggrecan. Here, we report the results of a survey of cellular immunity to human articular cartilage link protein, to bovine type II collagen, and to a synthetic peptide corresponding to the arthritogenic epitope in hsp65 in patients with inflammatory arthritis and normal age matched controls.

Patients and methods

Patients and Controls

Two consultant rheumatologists collected samples from patients attending the hospital as outpatients or receiving inpatient treatment. All the patients had active definite or classical RA. The mean age of the 47 patients tested with link protein was 59.5 years (SD 13.2, range 29–79). Control subjects without arthritic symptoms were recruited from local voluntary associations and had a mean age of 50.2 years (SD 13.5, range 21–82). For tests involving subpopulations of each group, the subgroups were selected randomly without age bias. Each patient or control provided samples of heparinised and clotted blood and, in addition, synovial fluid was aspirated from the knee joints of 17 patients as part of their medical treatment.

Antigens

Link protein was purified as previously described. Briefly, macroscopically normal human articular cartilage obtained from patients who had undergone total hip or knee replacement was minced and extracted with 4 mol/l guanidium hydrochloride (GuHCl). The extract was then fractionated by associative, followed by dissociative, caesium chloride density gradient centrifugation. The link protein rich A1D6 fraction was then mixed with AH-Sepharose 4B coupled to hyaluronic acid and the link protein eluted using 4 mol/l GuHCl. Link protein was dialysed extensively against 1 mol/l sodium chloride, 10 mmol/l 3-(N-morpholino)propanesulphonic acid (MOPS), pH 7.0. In this buffer, link protein forms stable oligomers in solution. The link protein was further purified by fractionation on a Sephadex G-150 Superfine column in MOPS buffer and fractions were assessed for purity on sodium dodecyl sulphate-polyacrylamide gel electrophoresis. The 4–20% gradient gels stained with Coomassie Brilliant Blue R-250 followed by silver staining. The Novex Xcell electrophoresis system (Novex, San Diego, USA) was used to perform Western blotting of protein bands in unstained gels onto nitrocellulose. The blots were probed with the link protein specific monoclonal antibody 9/30/8/AA and the aggrecan G1 and G2 domain specific antibody 12/21/1C6 (Developmental Hybridoma Bank). Purified link protein was used in mononuclear cell proliferation assays after sterile filtration through a 0.2 μm pore size membrane. Link protein solutions were stored at 4°C and the protein concentration was determined using the bicinchoninic assay (BCA) for protein (Pierce).

Because the link protein preparation contained MOPS, the effect of MOPS upon T cell proliferation was investigated. MOPS was not found to be generally inhibitory to mitogen (concanavalin A (con A)) or antigen (purified protein derivative (PPD) or tetanus toxin) driven proliferation at concentrations less than 300 μmol/l (data not shown). Nevertheless, those individuals (four of 96) with a high sensitivity to MOPS mediated inhibition, as demonstrated by a significantly reduced stimulation index in the presence of MOPS, were removed from the study.

Bovine nasal type II collagen was a kind gift of Dr S Ayad and purified Mycobacterium bovis derived hsp65 was a kind gift of Dr W Van Eden. A synthetic 17 amino acid peptide corresponding to hsp65 residues 180–196 was the kind gift of Dr D B Young. PPD or tetanus toxin were used as control antigens and con A was used as a control mitogen.

Mononuclear cell proliferation assay

Serum was prepared from clotted blood and complement inactivated by heating at 56°C for 30 minutes. Mononuclear cells were purified from heparinised peripheral blood (PBMC) or synovial fluid (SFMC) by centrifugation on Lymphoprep (Nycodent). The mononuclear cells were washed twice with RPMI 1640 (Dutch modification), supplemented with 1% v/v penicillin/streptomycin (Sigma) and 2 mmol/l l-glutamine (Sigma). The cells were then plated at a density of 2 × 10^5 cells per well (200 μl) in supplemented RPMI containing 10% v/v autologous complement inactivated serum in sextuplicate in 96 well flat bottomed plates, with or without the following antigens or mitogens: concanavalin A 1 μg/ml, link protein 20 μg/ml, hsp65 20 μg/ml, bovine nasal type II collagen 20 μg/ml, 17 amino acid hsp65 peptide 1 μg/ml, PPD 50 μ/ml, tetanus toxin 0.01%. The concentration of link protein and hsp65 were selected after a pilot scale dose-response study (unpublished results). The cells were incubated for five days at 37°C in 5% carbon dioxide and then pulsed with 3H-thymidine (1μCi/well; specific activity 2Ci/mmol). After six hours the cells were harvested onto glass fibre filter mats and the level of incorporation of thymidine determined by liquid scintillation counting. Lymphocyte responses were expressed as stimulation indices (SI), defined as the ratio of the level of incorporation of 3H-thymidine in mononuclear cells exposed to antigen to the level of incorporation in unstimulated cells (negative control).
STATISTICAL METHODS

Two different criteria were used to compare T cell responses to link protein in patients and controls. First, positive T cell responses were defined arbitrarily as a significantly increased thymidine incorporation compared with zero addition controls (p < 0.05 by the Mann-Whitney U test), where SI was at least 1.5. The proportion of individuals in each group showing positive responses were compared using Fischer's exact test. Second, by taking the natural logarithm of the SIs in the control population, it was possible to define a range of values which did not significantly differ from a normal distribution, and to calculate an upper limit for this range (defined as 2 SD above the mean of the transformed values). The number of patients and controls with SIs greater than this value were then compared. The SI of PBMC and SFMC (in those from whom synovial fluid could be obtained) were compared using the paired t test.

Results

Enhanced T cell proliferative responses to link protein, defined as an SI value > 1.50, were given by the PBMC of 26% of the RA patients tested—a significantly greater proportion (p < 0.05 by Fischer's exact test) than in the non-arthritic control group, of whom fewer than 10% showed increased responses (table 1). Table 2 shows the range of stimulation indices in patients and controls. The distribution of the natural logarithm of SI values in normal controls did not vary substantially from normal (x2, p = 0.409), with a mean of 0.08 (SD 0.42). An upper limit on the control reference range was set 2 SD above the mean, which corresponded to an SI value of 2.5 (95% confidence interval 2.1 to 3.2). Only one of the SI values of the patient group and two of those from the controls fell outside the reference range.

In those 17 patients from whom synovial fluid was obtained, eight (47%) had SI values >1.5 when the SFMC were challenged with link protein. In 16 of the 17 patients, the proliferative responses of the SFMC and PBMC to link protein could be compared directly; no significant difference between the SI for PBMC and that for SFMC was found by the paired t test.

PBMC and SFMC cells from responders and non-responders to link protein were tested for proliferative responses to type II collagen, but only one of the patients tested demonstrated a significant SI: this was with PBMC but not the SFMC of a patient unresponsive to link protein. Cells from both responders and non-responders were also challenged with the synthetic 17 amino acid peptide corresponding to hsp65 residues 180-196. None of the patients tested (table 1) showed increased SI values, whereas a high proportion of RA patients (eight of nine) and non-arthritic controls (seven of seven) showed increased responses to intact hsp65.

Neither the frequency nor the magnitude of increased responses to link protein appeared to be correlated with the duration or severity of the disease, or to the age or gender of the patient. There was no significant difference in the response of patients and controls to mitogen or control antigen.

Discussion

We have previously found that antibodies to link protein were present in the serum of patients with RA, but not at significantly greater frequency or titre than in control individuals. In the present study we have shown that a greater proportion of RA patients have peripheral blood derived T cells reactive to link protein (defined as an SI of 1.5 or greater) than non-arthritic individuals. The frequency of such responses was greater still in synovial fluid derived T cells. However, as the SI values for PBMC for all but one of the patients fell within the control range, it was not possible to show a relationship between the magnitude of these responses and the presence or stage of RA. It seems likely that sub-groups of both RA patients and controls show weak T-cell proliferative responses to link protein.

Evidence from adjuvant induced arthritis had led to the hypothesis that molecular mimicry between mycobacterial heat shock proteins and cartilage proteoglycan could lead to autoimmune responses and arthritis. In the adjuvant arthritis model, T cell cloning gave rise to both arthritogenic (A2b) and protective (A2c) T cell clones which recognised the same epitope (AA 180-188) on M.tuberculosis hsp65. This peptide has 44% homology with rat chondrosarcoma link protein and 33% with human cartilage link protein. Transfer of the arthritogenic clone A2b to naive animals gave rise to enhanced delayed-type hypersensitivity responses to preparations of cartilage proteoglycans rich in aggrecan and link protein. In this study, although we found a high percentage of patients and controls responsive to hsp65 (which is not surprising in light of widespread
immunisation with *M.tuberculosis*, no reactivity to a 17 amino acid synthetic peptide incorporating the sequence AA 180-188 was observed, regardless of whether patients showed reactivity to link protein or to intact *M.bovis* hsp65. We must conclude that in the patients tested this epitope is not immunodominant, and the responses to link protein are not attributable to cross reactivity with it. In a study analysing the pattern of responses of T cells reactive with *M.tuberculosis* in the synovial fluid of RA patients, 12 different antigenic specificities were seen in a panel of 15 T cell clones purified from four patients. It seems unlikely that T cell responses in the joint are directed principally against those sequences in auto-antigens homologous with mycobacterial proteins.

Link protein shares significant homology with the G1 and G2 domains of aggrecan, and T cell responses to aggrecan in the murine HFPG induced model of arthritis have been investigated. From the results of one study it has been suggested that the predominant T cell epitopes on aggrecan may be in the chondroitin sulphate attachment region; however, in another study T cell lines and hybridomas have been shown to cross react with link protein. The responses to link protein reported here are somewhat different from RA patients and normal control T cell responses to deglycosylated aggrecan. First, RA patients showed a greater incidence of T cell proliferative responses to aggrecan than to link protein. Second, compared with those of RA patients, the stimulation indices given by responsive normal controls were similar against link protein, but generally lower against aggrecan.

Cellular immunity to link protein may play a role in the initiation or maintenance of the chronic inflammatory response in at least some RA patients, though it is difficult to distinguish if such immune responses are causal or a result of the destructive process. Articular cartilage is believed to be an immunologically privileged site and the continual, often aggressive erosion of cartilage in patients with RA may release novel matrix epitopes able to elicit secondary immune responses to a wide range of matrix components. It has been shown that the level of release of hyaluronan binding region into synovial fluid is much greater in RA patients with advanced cartilage degeneration than in those with little or no cartilage damage. It is also possible that such epitopes are released during cartilage damage unrelated to the presence of inflammatory arthritis, including that caused by degenerative arthritis or by trauma, which may explain the presence of T cells responsive to link protein in the control population. Fragments from proteoglycans and associated matrix components may be cross reactive with pathological organisms, or altered antigen processing may result in the presentation of cryptic epitopes to T cells, as has been observed in the Balb/c mouse model of proteoglycan induced arthritis. Further work on the processing of proteoglycans by cells from RA patients and normal individuals and mapping T cell epitopes on these antigens is needed to determine the relevance of our findings.

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