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Heterogeneity and Clinical Significance of Glomerular-binding Antibodies in Systemic Lupus Erythematosus

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Abstract

We used an ELISA employing extracts of human glomerular basement membrane (GBM) to detect, characterize, and evaluate the clinical significance of glomerular-binding IgG in patients with SLE nephritis. Most patients with SLE nephritis exhibited GBM-binding IgG, although many patients with active nonrenal SLE or symptomatic, drug-induced lupus had similar reactivity, albeit at lower levels. IgG binding to GBM in SLE nephritis patients was decreased by DNase pretreatment of GBM, restored after DNase with nuclear antigens (most notably with nucleosomes), inhibited by exogenous nuclear antigens (particularly nucleosomes), but unaffected by exposure to DNase/high ionic strength. The characteristics of IgG binding to GBM largely paralleled the patients’ underlying autoimmune response, which was dominated either by antibodies to DNA/nucleosomes or to nucleosomes alone. Binding of lupus sera to nonrenal extracellular matrix (even with nucleosomes) was not equivalent to GBM. Collagenase pretreatment of GBM variably decreased IgG binding, depending on the level and type of binding. SLE nephritis patients with high levels of GBM-binding IgG exhibited more severe disease clinically, but the same renal histopathology, as patients with lower levels. The level of GBM-binding IgG at presentation did not predict the therapeutic response, but decreased in responders to therapy. In sum, glomerular-binding IgG in lupus nephritis bind to epitopes on chromatin, which adheres to GBM in part via collagen. These autoantibodies appear necessary, but not sufficient, for the development of nephritis, and correlate with clinical and histopathologic parameters of disease activity. (J. Clin. Invest. 1996. 98:1373–1380.) Key words: lupus nephritis • autoantibodies • glomerular basement membrane • systemic lupus erythematosus • nucleosomes

Introduction

The prevailing wisdom regarding SLE nephritis is that this disorder is mediated by anti-DNA antibodies (reviewed in reference 1). The presence of anti-DNA antibodies correlates with nephritis in both mice and human patients, anti-DNA antibodies are concentrated within human and murine glomeruli in vivo, administration of DNA to autoimmune mice accelerates the progression of nephritis, and anti-DNA mAbs produce nephritis by adoptive transfer (2–9). Substantial data have accumulated which suggest, however, that anti-DNA antibodies are neither necessary nor sufficient for nephritis to occur (reviewed in reference 1). Immunogenetic studies of murine lupus have demonstrated that not all mice with anti-DNA antibodies develop nephritis and that nephritis can occur in their absence (10, 11). Only certain anti-DNA mAbs are pathogenic when administered to nonautoimmune mice (7, 9). Moreover, patients with lupus nephritis do not invariably have anti-DNA antibodies (3, 12, 13).

Apropos to these issues, recent studies have reinforced the concept that SLE is an antichromatin disease; e.g., that the autoimmune response is against chromatin with the development of antibodies directed to various epitopes on chromatin including but not limited to DNA (14–19). The possibility that antinucleosomal antibodies may generally be involved in nephritis has also been suggested by recent experimental and clinical studies (15, 18, 20, 21). In one recent clinical study of SLE, antibodies to epitopes on chromatin were found to be more highly associated with nephritis than antibodies to DNA (15).

The concept that lupus nephritis involves the participation of a family of antinucleosomal antibodies has also been supported by recent work by our group on identifying glomerulotropics antibodies in lupus nephritis. In our studies, we used glomerular and glomerular basement membrane (GBM)-binding ELISAs to define glomerular-binding antibodies in murine lupus serum and to produce glomerular-binding mAbs. The antibodies detected by these assays in MRL lpr serum react specifically with glomeruli in vitro, concentrate in glomeruli in vivo, and correlate with the presence of nephritis (9, 22, 23). Analysis of MRL lpr serum and mAbs derived from these mice has demonstrated that such antibodies comprise a family of IgGs that react with various epitopes on chromatin adherent to GBM type IV collagen (24, 25). These data support the hypothesis that the immunopathogenesis of lupus nephritis involves the contribution of multiple autoantibodies that bind to the glomerulus by virtue of chromatin adherence to GBM collagen.

In other recent work, we have found that glomerular-binding antibodies are associated with nephritis in SLE patients (26). The intent of the current work was to characterize these

1. Abbreviations used in this paper: dsDNA, double-stranded DNA; ECM, extracellular matrix; GBM, glomerular basement membrane; H, histone.
human glomerular-binding antibodies, and thereby test the above hypothesis for human SLE. Towards this end, we developed a human GBM ELISA similar to that which we used in our murine studies (24, 25). We used this assay to characterize glomerulotrophic antibodies in lupus patients, and to assess both the diagnostic and prognostic import of such antibodies. The results we obtained suggest that we should reorient our conceptualization of lupus nephritis to a disease mediated by antichromatin autoantibodies rather than simply anti-DNA antibodies. Moreover, the data suggest that the development of nephritis is not simply a function of the presence of these autoantibodies.

Methods

Patient characteristics and treatment. Sera were obtained from 63 patients with established lupus nephritis who were enrolled in a randomized prospective study at the National Institutes of Health. Entry required (a) a diagnosis of SLE based on the American College of Rheumatology classification criteria (27), and (b) glomerulonephritis. Glomerulonephritis was defined as an active urine sediment (>10 red blood cells/hpf and/or cellular casts without evidence of infection on two or more urinalyses) or >3 g proteinuria/24 h plus histological evidence of proliferative glomerulonephritis on a renal biopsy obtained within 3 mo of study entry. Two patients were not biopsied because of uncontrolled hypertension or anticoagulation therapy, but both patients exhibited an active sediment and >3 g proteinuria/24 h.

Sera from 15 patients with clinically active nonrenal lupus and from 33 patients with clinically inactive lupus were obtained from patients in the Rheumatology Clinic at Washington University. These patients also fulfilled the American College of Rheumatology criteria for SLE (27). The former patients had clinically active disease requiring treatment at the time of sample procurement, but did not fulfill the American College of Rheumatology criterion for nephritis. The latter patients had no evidence of active lupus at the time of sample procurement and for 1 yr thereafter. Patients with symptomatic, drug-induced lupus and with drug-induced autoantibodies without symptoms have been described in detail (28, 29). The patients with symptomatic, drug-induced lupus included eight patients with lupus induced by procainamide, six by hydralazine, seven by quinidine, and one each with lupus induced by acebutalol, penicillamine, methyl-

dopa, and timolol. All asymptomatic patients were treated with GBM or DNA-binding IgG in lupus sera, sera were exposed to conditions of high ionic strength and DNase simultaneously before assay using a variation of a published method (20). Serum was diluted 1:5 (v/v) in 3 M NaCl containing MgCl2 5 mM and 100 μg/ml DNase, and then incubated with one of the following nuclear antigens at 10 μg/ml before assay: histones (purified mixture of H1, H2A, H2B, H3, and H4; Calbiochem-Novabiochem Corp., La Jolla, CA), double-stranded (ds)DNA (λ phage DNA; Sigma), or nuclease (Worthington), as previously described (24). In the inhibition studies, serum was preincubated with histones, dsDNA, or nucleosomes (at a final concentration of 100 μg/ml) before assay, as described (24).

To determine a potential contribution of immune complexes to GBM or DNA-binding IgG in lupus sera, sera were exposed to conditions of high ionic strength and DNase simultaneously before assay using a variation of a published method (20). Serum was diluted 1:5 (v/v) in 3 M NaCl containing MgCl2 5 mM and 100 μg/ml DNase, and incubated 16 h at 37°C. Before assay, serum was further diluted in 1 mM EDTA containing 10 mg/ml BSA to inactivate the DNase, achieve isotonicity, and produce a final dilution of 1:100 for assay. The binding of DNase/high ionic strength–treated serum was compared to that of untreated serum. DNase was active in the high ionic strength buffer (data not shown).

Anti-dsDNA and antinuclear antigen ELISAs. The anti-dsDNA assay was performed using a previously published ELISA protocol that uses λ phage dsDNA coated on poly-L-lysine (Sigma; 25). Binding to intact chromatin, oligonucleosomes (2–8 nucleosomes in length, stripped of histone H1), histone H2A-H2B–DNA complexes, histone (H3-H4)–DNA complexes, histone H2A-H2B complexes without DNA, histone (H3-H4), complexes without DNA, or individual histones (H1, H2A, H2B, H3, and H4) were also performed by ELISA as published (25). Individual histones used as assay substrates were purchased from Calbiochem-Novabiochem, while other nuclear antigens were prepared as described (36).

Statistical analysis. GBM and nuclear antigen ELISAs were run in duplicate and are presented as average values. GBM reconstitution and inhibition studies were performed in triplicate. Data for these experiments were normalized to the GBM binding of unmanipulated serum (=100%) to facilitate comparisons between patients. Regres-
sion analysis of nuclear antigen reactivity and GBM binding was performed using the least squares method. Comparisons of clinical data between the subgroups of patients with lupus nephritis were performed by Mann-Whitney U test, and median values are shown. Comparison of multiple groups used one-way ANOVA combined with Student’s t test, since analysis was restricted to a limited number of comparisons. Comparison of two groups used Student’s t test. Means±SEM are shown for the groups.

**Results**

Quantification of GBM-binding antibodies in patients with SLE. As shown in Fig. 1, the average level of IgG binding to GBM in lupus nephritis patients was substantially and significantly greater than that observed in normal individuals. Virtually all lupus nephritis patients (81%) exhibited GBM-binding IgG greater than the normal range (i.e., >0.05 OD). The average IgG binding to GBM in lupus nephritis patients was also significantly greater than that in patients with inactive lupus, although a number of patients with inactive lupus also exhibited levels greater than normal controls (Fig. 1). As indicated above, none of the patients with inactive lupus (even with elevated levels of GBM-binding IgG) developed nephritis during the next year. Additionally, levels of GBM-binding IgG did not correlate with a previous history of nephritis in these patients (data not shown). The average level of GBM-binding IgG in patients with lupus nephritis was not significantly greater than that in patients with active lupus but without clinical evidence of renal disease (Fig. 1).

We additionally quantified GBM-binding IgG in the sera of patients with drug-induced lupus. As shown in Fig. 1, the average level of GBM-binding IgG in patients with symptomatic, drug-induced lupus was similar to that in patients with active lupus without renal disease and was not significantly different from that in patients with SLE nephritis. In contrast, patients with drug-induced SLE without symptoms (i.e., asymptomatic but serologically abnormal) exhibited an average level of GBM-binding IgG similar to normal controls (Fig. 1).

Qualitative characterization of GBM-binding IgG in lupus nephritis patients. We used the group of inactive lupus patients to separate patients with active lupus nephritis into two groups: those with high GBM-binding IgG (>mean ± 2×SD of the inactive population or >0.285 OD) and those with lower levels (<0.285 OD). 27% of the nephritis patients fell into the former category. Qualitative characterization of the GBM-binding IgG in representative patients with high levels demonstrated substantial similarities among patients. GBM binding was uniformly diminished by DNase pretreatment of the GBM (average decrease of 67±6%, Fig. 2 A). In contrast, DNase did not disrupt the binding of the commercial anti-GBM antibody (data not shown). Binding of these sera was also efficiently reconstituted by incubating the DNase-treated GBM with either dsDNA alone or nucleosomes before assay (Fig. 2 A). In several patients (Nos. 25, 100, and 106), however, histones alone were able to reconstitute binding, although less efficiently than either DNA or nucleosomes (Fig. 2 A). Inhibition studies using exogenous nuclear antigens largely corroborated the reconstitution studies (Fig. 2 B). Nucleosomes and dsDNA consistently inhibited binding, whereas the effect of histones was more modest and variable.

Profiling the nuclear antigen–binding characteristics of the sera from these patients demonstrated an overall parallelism between the characteristics of IgG binding to GBM and the underlying autoimmune response to nuclear antigens (Fig. 2 C). The autoimmune response of this group of patients was characterized by substantial binding to dsDNA, chromatin, and subnucleosomes. Many of these patients, however, exhibited considerable binding to individual histones or histone–histone complexes. Although no common histone-binding pattern was observed among patients, the ability of histones to reconstitute GBM binding after DNase treatment in this group of patients was correlated with reactivity to the H2A–H2B dimer (r = 0.85, P = 0.017 for the correlation between histone reconstitution of GBM binding and H2A–H2B binding; P > 0.05 for other correlations).

The qualitative characteristics of GBM-binding IgG from representative patients with lower levels of such IgG were distinct from this first group. In these patients, GBM binding was comparably disrupted by DNase pretreatment of the GBM (average decrease in binding of 64±9%, Fig. 3 A). Binding was reconstituted most consistently and effectively with nucleosomes, however, and less efficiently (and more variably) with histones or dsDNA alone (Fig. 3 A). Inhibition studies in these patients exhibited two different patterns (Fig. 3 B). One subset of these patients (Nos. 16, 71, 96, and 117) exhibited inhibition with nucleosomes and DNA, and variable inhibition with histones. In contrast, a second subset of patients (Nos. 77, 86, and
exhibited little inhibition (and occasionally enhanced binding) in these experiments. These latter patients were those who exhibited the most exuberant reconstitution of GBM binding with histones.

Similar to the patients with high levels of GBM-binding IgG, an overall parallelism between IgG binding to GBM and the underlying autoimmune response to nuclear antigens was observed in patients with lower levels (Fig. 3 B). The autoimmune response in these latter sera was dominated by uniform reactivity with chromatin and subnucleosomes with modest-to-negligible reactivity to dsDNA. The binding of these sera to individual histones or histone–histone complexes was also modest with the exception of binding to H1 in several of the sera. No correlation between the ability of histones to reconstitute GBM binding and binding to individual histones or histone–histone complexes was observed ($P > 0.05$ for all correlations).

Because DNase treatment of GBM did not completely prevent the binding of IgG from lupus nephritis sera, we conje-

Figure 2. Characterization of GBM binding and nuclear antigen reactivity in sera from patients with GBM-binding IgG > 0.285 OD. The characteristics of IgG binding to GBM in representative patients with GBM-binding IgG > 0.285 OD was determined using both reconstitution and inhibition strategies. GBM binding is normalized to 100%, and individual patients are referred to by number on the x axis. The binding of these same sera to a panel of defined nuclear antigens was concurrently determined. (A) Effect of DNase pretreatment of GBM on binding, and the ability of nuclear antigens to reconstitute binding after DNase. (B) Inhibition of GBM binding by exogenous nuclear antigens. (C) Nuclear antigen binding. Nuc, nucleosomes; Oligo, oligonucleosomes; Chr, chromatin; T-D, H3-H4 tetramers with DNA; D-D, H2A-H2B dimers with DNA; T, H3-H4 tetramers; D, H2A-H2B dimers.

Figure 3. Characterization of GBM binding and nuclear antigen reactivity in sera from patients with GBM-binding IgG < 0.285 OD. The characteristics of IgG binding to GBM in representative patients with GBM-binding IgG < 0.285 OD was determined using both reconstitution and inhibition strategies. GBM binding is normalized to 100%, and individual patients are referred to by number on the x axis. The binding of these same sera to a panel of defined nuclear antigens was concurrently determined. (A) Effect of DNase pretreatment of GBM on binding and the ability of nuclear antigens to reconstitute binding after DNase. (B) Inhibition of GBM binding by exogenous nuclear antigens. (C) Nuclear antigen binding.
tured that this residual binding might be caused by immune complexes binding to GBM (e.g., anti-DNA antibodies complexed to DNA or nucleosomes; 20, 25, 37). To test for this possibility, we exposed serum from patients with high levels of GBM-binding IgG to high ionic strength/DNase, and then examined for an effect on binding to GBM, DNA, or poly-l-lysine (pre-coat for the anti-DNA assay). This manipulation, however, did not significantly alter binding to these substrates (data not shown). We subsequently postulated that the residual binding after DNase might be caused by retained histones that can bind to GBM with high affinity (34, 35). Consequently, we examined the effect of DNase followed by high ionic strength on GBM binding. Using sera with high GBM-binding IgG, we observed that the combination of these two treatments abrogated GBM binding (decrease of 96±2%, n = 7).

To examine the specificity of GBM binding, we next determined whether nonrenal ECM could substitute for GBM in the binding assay. As shown in Fig. 4, lupus nephritis patients, patients with active nonrenal lupus, and patients with inactive lupus were differentiated by their respective levels of GBM-binding IgG, but not by the level of IgG binding to placental ECM (which is similar in composition to the GBM, see Methods). The inability of placental ECM to substitute for GBM could not be rectified by adding nucleosomes to the placental ECM substrate (Fig. 4).

As we have noted with murine lupus, type IV collagen within the GBM is necessary for autoantibody binding because of the adherence of nuclear antigens to this protein (24, 25). Autoantibody binding to type IV collagen per se is negligible (24, 25). We tested these statements in human lupus by examining the effect of collagenase on GBM binding of lupus nephritis sera. As shown in Fig. 5, GBM binding of lupus nephritis sera was generally decreased by preincubation of the GBM with collagenase, though substantial variability was noted among patients (ranging from a decrease of 0 to 100%). The average decrease in a set of 25 lupus nephritis sera (14 with GBM binding > 0.285 and 11 with GBM binding < 0.285 OD) was 65±5%. The effect of collagenase on GBM binding tended to be less substantial in sera with GBM binding > 0.285 OD vs. sera with GBM binding < 0.285 OD: 57±5% vs. 76±9% inhibition, respectively (P = 0.057). Such sera could also be distinguished, based on the characteristics of IgG binding to GBM (see above) and clinical associations (see below). Lupus nephritis sera did not bind to human type IV collagen per se (data not shown).

**Association of GBM-binding IgG in lupus nephritis patients with clinical parameters and renal histopathology.** We next sought to determine whether the level of GBM-binding IgG (which, as noted above, was correlated with the overall characteristics of IgG binding to GBM) could be related to the clinical status, histopathology, or prognosis of the patient. Patients with high levels of GBM-binding IgG overall had evidence of more severe disease by clinical criteria than patients with lower levels of GBM-binding IgG (Table I). The former group of patients

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<th>Parameter</th>
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<th>GBM &lt; 0.285 (n = 46)</th>
<th>P</th>
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<td>0.3000</td>
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**Table I. Clinical Parameters in Lupus Nephritis Patients Subdivided by GBM-binding IgG Level**

Lupus nephritis patients were subdivided into two groups, depending on whether or not the GBM-binding IgG was greater or less than the range defined by the inactive lupus patients (> or < 0.285 OD).
IgG after therapy in the responders was significantly decreased (two groups before and after therapy are shown. The GBM-binding responders and nonresponders). GBM-binding IgG values for these whether or not they exhibited a successful response to therapy (i.e., patients. Patients with lupus nephritis were subgrouped depending on class III) were seen in the two groups.

and focal/segmental proliferative glomerulonephritis (WHO patients with diffuse proliferative nephritis (WHO class IV) were similar in the two groups (Table I). Proportionate numbers of exhibited evidence of more substantial complement activation in vivo with lower serum levels of C3 and C4, and a more severely depressed CH50 than the latter group. These patients also exhibited higher erythrocyte sedimentation rates, higher levels of IgG, and lower hematocrits on average. Renal function was diminished in both groups of patients, with a trend toward more severe renal dysfunction in the former group. Despite these differences, the overall histopathologic severity was similar in the two groups (Table I). Proportionate numbers of patients with diffuse proliferative nephritis (WHO class IV) and focal/segmental proliferative glomerulonephritis (WHO class III) were seen in the two groups.

We additionally segregated the cohort of patients into subgroups, depending on their subsequent response to therapy after 1 yr to analyze the relationship between the amount (and consequently type) of GBM-binding IgG and prognosis (Fig. 6). There was no significant difference in the average level of GBM-binding IgG before therapy in those patients responding to therapy vs. those patients who were refractory to therapy. Patients who were successfully treated exhibited a significant decrease in GBM-binding IgG, however, while those patients who were not successfully treated exhibited a more modest (and not statistically significant) decrease in GBM-binding IgG.

Discussion

This study demonstrates the potential heterogeneity in glomerulotrophic antibodies in SLE patients. As we observed, one subgroup of patients with lupus nephritis exhibited GBM-binding IgG that bound largely in a DNA- and nucleosome-dependent fashion. A subset of these patients also had GBM binding that was histone dependent. A second subgroup of patients exhibited GBM-binding IgG that bound largely in a nucleosome-dependent fashion. These data support our contention derived from an analysis of murine lupus that lupus nephritis is immunologically complex, specifically that multiple antibodies (including but not limited to anti-DNA antibodies) may contribute to nephritis (24, 25).

Recent observations suggest that SLE is characterized by a breakdown of tolerance to chromatin (14, 15, 17). Apropos this issue, the type of glomerular-binding antibodies present in the patients who were analyzed in the current study recapitulated their underlying antichromatin response. Our observations that lupus nephritis patients can be divided into two groups, depending both on their GBM-binding IgG and their overall antichromatin response, parallel other recent studies on the underlying autoimmune response in lupus (15). In both studies, SLE patients could be segregated into two groups: one with predominant reactivity with chromatin, subnucleosomes, and dsDNA, and one with predominant reactivity with chromatin and subnucleosomes. The former group serologically resembles MRL lpr mice with established autoimmunity, whereas the latter group resembles young MRL lpr mice at the inception of autoimmunity, suggesting that these groups of patients may represent different stages in the evolution of the breakdown of tolerance (14, 16).

In addition to supporting the concept that multiple autoantibodies contribute to lupus nephritis, the current data are most consonant with either the planted antigen or immune complex hypotheses for lupus nephritis (1). Most of IgG binding to GBM in lupus nephritis patients was abolished by DNase pretreatment of GBM extracts, reconstituted with nuclear antigens after DNase, and inhibited by exogenous nuclear antigens similar to what we have observed in murine lupus (24, 25). These data demonstrate that the autoantibodies detected by the GBM ELISA can form immune complexes with nuclear antigens trapped by the GBM. It is nonetheless possible that GBM-binding IgG may complex with nucleosomes in the circulation and then deposit in the GBM. Nucleosome/antinucleosome complexes have been shown to bind avidly to the glomerulus in vivo and to the GBM in vitro (18, 20, 25). Although our data suggest that most GBM-binding IgG is not comprised by such complexes (GBM binding was stable to DNase/high ionic strength), we cannot exclude a contribution of these complexes to GBM binding. In fact, our murine data suggest the presence of small amounts of immune complexes with glomerular-binding activity (22). The relatively low level of such complexes, however, may be a function of rapid clearance.

The current data do not support a major contribution of antibodies that bind directly to glomeruli (e.g., cross-reactive anti-DNA antibodies, autoantibodies to GBM proteins [38–41]), although the GBM ELISA may not detect autoantibodies to intrinsic glomerular cells (42). Of relevance to this issue are recent studies that have cast doubt on the existence of such autoantibodies. The binding of cross-reactive anti-DNA mAbs to nonnuclear antigens or cells appears to be indirect in many circumstances and mediated by immune complexes that are formed in vitro with nucleosomes (20, 43). Nuclear antigens also bind avidly to basement membrane components isolated from biological sources (as we have noted with our GBM preparations [24]), and cells appear to release nucleosomes during in vitro culture that may adhere to cell surfaces and the pericellular matrix (44, 45). These adherent nuclear antigens may be responsible for the binding interactions observed with autoantibodies and various basement membrane proteins or cells.
It is of note that lupus sera bound differently to GBM compared with placental ECM (without or with nucleosomes), despite the overall similarity in protein composition. This divergence may be a function of finer differences in the composition of the GBM relative to other ECMs (e.g., in the α chain composition of type IV collagen), as well as the nuclear antigens that adhere to GBM relative to other ECMs. Further investigation of how nuclear antigens interact with the ECM thus would appear relevant to the pathogenesis of lupus nephritis.

Germane to this issue, the present study specifically suggests that GBM collagen (the major GBM protein [46]) is important for autoantibody binding to the glomerulus in lupus nephritis, as we have noted with murine lupus sera and glomerulotropism of mAbs [24, 25]. Unlike murine lupus, however, we noted substantial variability between patients, suggesting that GBM proteins other than collagen (e.g., heparan sulfate proteoglycan [20]) probably play an important role in autoantibody binding to the glomerulus. The collagen dependence of GBM binding is likely to be a function of the avidity of histones for type IV collagen (35) and does not appear to be caused by the binding of sera to collagen per se. We also observed that the degree of collagen dependence of binding relates to the level, and consequently, type of GBM binding IgG (i.e., DNA/nucleosomal vs. nucleosomal) that also correlates with the patient’s clinical status. Thus, nuclear antigen specificity, GBM binding properties, and pathogenicity of autoantibodies appear to be interrelated variables.

The current work further supports the conclusion that GBM-binding IgG in lupus is pathogenic and necessary for the development of nephritis. Elevated levels were noted in virtually all lupus patients with nephritis, adequate responses to therapy in these patients were accompanied by significant reductions in antibody levels, and particularly high levels of GBM-binding IgG were associated with more severe clinical disease. The presence of GBM-binding IgG, however, is clearly not sufficient for the development of nephritis. As noted, GBM-binding IgG was observed in patients with active nonrenal lupus, symptomatic drug-induced lupus, and occasionally in inactive lupus. In addition, neither the level nor the overall characteristics of GBM-binding IgG in nephritis patients could be related to the renal histopathology or prognosis in a simple fashion. Similar disparities between the autoantibody response and both the presence and severity of nephritis, while not apparent within the inbred strains of murine lupus, are nonetheless apparent in interspecific back-crosses of MRL/lpr mice and in the graft vs. host model of murine lupus when induced across varying MHC class II differences (47, 48).

Therefore, we would conclude that glomerular binding IgG is necessary but not sufficient for the development of nephritis. The development of nephritis and specific histopathologic presentations, as well as the propensity of glomerular inflammation to progress to glomerulosclerosis, must result from a more complex array of factors. Conjecturally, specific antibodies within the broad categories defined by our GBM assay may be more nephritogenic than others. This capacity may simply be a function of the IgG subclass (and the ability to fix complement) or may be a function of the epitopes of chromatin that are most often accessible when adherent to the GBM in vivo. Additionally, factors relating to the release and clearance of chromatin, as well as the formation of circulating immune complexes, may contribute to the development of nephritis and specific pathologic variants. Finally, genetic variables controlling the extent of the local inflammatory response may contribute to the severity and long-term outcome of nephritis. Resolution of these issues will be critical in refining current diagnostic tests, improving prognostication, and targeting therapy in a more specific fashion.

In conclusion, the current study demonstrates the heterogeneity inherent in glomerulotropism in lupus nephritis and argues against an exclusive role for anti-DNA antibodies in this disorder. Specifically, the data suggest that the pathogenesis of nephritis results from the binding of antibodies to various epitopes on chromatin that bind to GBM largely (though not exclusively) via collagen. Although GBM-binding IgG levels are correlated to the concurrent clinical status of the patient and vary with the therapeutic response, the genesis of nephritis and its various pathologic forms and the factors controlling the long-term outcome of nephritis remain to be clarified.

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References


