Clinical utility of anti-C1q antibody in primary and secondary vasculitic conditions

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ABSTRACT

Objective: Anti-C1q antibodies (Anti-C1q Ab) are seen in hypocomplementemic urticarial vasculitis syndrome (HUVS), infection-associated vasculitis such as hepatitis C virus-related vasculitis and in autoimmune diseases such as rheumatoid vasculitis, polyarteritis nodosa, giant cell arteritis, vascular Behcet’s disease, and cryoglobulin associated vasculitis. Aim of this study is to evaluate the presence of Anti-C1q Ab in vasculitis and to determine if any difference exists between primary and secondary vasculitis in relation to this antibody.

Patients and Methods: Consecutive patients with diagnosis of either a primary or secondary vasculitis were recruited. Primary vasculitis were diagnosed by the American College of Rheumatology 1990 criteria. Clinical features and serological markers were noted. Anti-C1q Ab was assayed by commercially available ELISA kit (Demeditec Diagnostics GmbH, Germany).

Results: Sixty-four patients were recruited for the study comprising of 41 primary vasculitis and 23 secondary vasculitis cases. No difference in Anti-C1q Ab levels between primary and secondary vasculitis was noted. Four patients were positive for Anti-C1q Ab out of the 64 patients. Of the four, one patient was diagnosed as HUVS, 2 patients as systemic lupus erythematosus with vasculitis (16.7%) and another patient was diagnosed as rheumatoid arthritis with vasculitis (14.28%). Anti-C1q Ab negatively correlated with age and C3, but it correlated positively with erythrocyte sedimentation rate (ESR) in vasculitic patients.

Conclusion: Presence of anti-C1q Ab did not differ between the patients with primary and secondary vasculitis. Anti-C1q Ab titers correlated with younger age, high ESR, and low C3 in patients with vasculitis in our study.

Keywords: Anti-C1q antibody, complements, India, vasculitis

Introduction

C1q is specifically bound to early apoptotic belbs of keratinocytes and vascular endothelial cells, which subsequently leads to activation of complement system, thereby facilitating clearance of apoptotic bodies. Anti-C1q antibody (Anti-C1q Ab) develops in disorders characterized by immune complex mediated injury, more commonly in skin and glomerular microvasculature. Anti-C1q Ab are IgG antibodies bound to collagen like C1q. Anti-C1q Ab are seen in hypocomplementemic urticarial vasculitis syndrome (HUVS) (100%), mixed connective tissue disorder (94%), Felty’s syndrome (76%), systemic lupus erythematosus (SLE) (30-60%), and rheumatoid vasculitis (32%). Other than HUVS, Anti-C1q Ab is also seen in vasculitic disorders such as polyarteritis nodosa (PAN), giant cell arteritis, vascular Behcet’s disease and cryoglobulin associated vasculitis. Anti-C1q Ab is also seen in patients with vasculitis of infectious etiology, which includes hepatitis C virus related vasculitis and toxocariasis. Thus, vasculitis due to autoimmune conditions and infections can be associated with elevated Anti-C1q Ab. However, role of Anti-C1q Ab in pathogenesis of vasculitis is not clear. The aim of the present study is to determine if presence of ant-C1q antibody in primary and secondary vasculitic conditions has clinical implications.

Justification of undertaking the study

Ideal marker of disease activity in vasculitic diseases is in scarcity. As vasculitic diseases can be life-threatening, study of utility of Anti-C1q Ab in these diseases is worthwhile in the light of the biological basis explained above.
Patients and Methods

An observational study was carried out between January 2013 and January 2015 in the Department of Clinical immunology and Rheumatology of Christian Medical College, Vellore, India. This study was approved by the Institutional Review Board. Consecutive patients with diagnosis of either a primary or secondary vasculitis were recruited after getting written consent from the participants. Primary vasculitis were diagnosed by the American College of Rheumatology 1990 criteria. Various clinical parameters, laboratory and serological markers were noted at the time of recruitment. Clinical parameters included duration of disease before presentation, presence of organ system involvement (arthritis, skin symptoms, serositis, and central nervous system symptoms), presence of thromboembolic events and major infections.

Laboratory and serological markers

Laboratory and serological markers including routine hematology, ESR, complements C3 and C4, urine protein/urine creatinine ratio (UP/UC), presence of autoantibodies (dsDNA antibody, RF, and antiphospholipid antibodies), and histopathology reports were noted as done by some earlier workers.7 When these laboratory results were not available at the precise date of Anti-C1q measurement, values within 15 days of the measurement of Anti-C1q Ab were acceptable.7

Composite indices

Disease activity was assessed at the time of Anti-C1q estimation, by birmingham vasculitis activity score (BVAS).5 Vasculitis damage index (VDI) was also recorded.6

Estimation of Anti-C1q Ab

Blood samples for Anti-C1q Ab were collected during their hospital visits irrespective of disease duration or dose of immunosuppressant. Anti-C1q IgG antibody was assayed by commercially available ELISA kit (Demeditec Diagnostics GmbH, Germany). Results were expressed as unit/ml (U/ml) and serum Anti-C1q Ab level more than or equal to 10 U/ml (cutoff value), as recommended by the manufacturer, was considered to be a positive test value.

Statistical analysis

Sample size was calculated for the study assuming a 25% difference in positivity of Anti-C1q Ab between primary and secondary vasculitis. We, therefore needed a sample size of 40 primary and 20 secondary vasculitis participants with 80% power, 5% error, and an allocation ratio of 2 primary: 1 secondary vasculitis case.

The statistical analysis of data was done using STATA 13.1 I/C (StataCorp LP, Texas, USA, Version 13) statistical package. To test the normality of data distribution, K-S (Kolmogorov–Smirnov) test was done. Mann–Whitney t-test was used to compare Anti-C1q Ab between the groups. Correlation between Anti-C1q Ab and clinical features as well as laboratory markers were done by spearman correlation test. A value of $P < 0.05$ was considered as significance.

Results

Sixty-four patients were recruited for the study. Baseline characteristics of patients were shown in Table 1. Anti-C1q Ab did not differ between primary and secondary vasculitis. Anti-C1q Ab titer (>10 IU/ml) was elevated only in four patients in the total cohort of 64 patients. Three patients were diagnosed as secondary vasculitis and one was diagnosed as primary vasculitis. To be precise there was 1 patient with the diagnosis of HUVS (Primary vasculitis), 2 patients had SLE with vasculitis (16.7%) and the 4th patient with C1q antibody positivity was diagnosed as rheumatoid arthritis (RA) with vasculitis (14.28%). Serological markers of these four patients were shown in Figure 1.

Looking at the whole cohort, one patient out of 41 (2.4%) patients with primary vasculitis and 3 patients out of 23 (13%) patients with secondary vasculitis were found to be positive for the Anti-C1q Ab. However, this difference did not reach statistical significance ($P = 0.09$). Anti-C1q Ab, however, correlated with age ($r = −0.3952; P = 0.0013$), ESR ($r = 0.4628; P = 0.0001$), and C3 ($r = −0.360; P = 0.019$), which was shown in Figure 2. No other significant correlation was observed between the other parameters and Anti-C1q Ab.

Discussion

The present study found only four patients positive for Anti-C1q Ab out of 64 patients with vasculitis. The positivity of Anti-C1q Ab was higher in secondary vasculitis (13%) as compared to primary vasculitis (2.4%); however, this difference did not reach statistical significance. No definitive conclusion can be drawn due to small sample size.

One out of 4 patients with HUVS had high titer Anti-C1q Ab in our study. The presence of high titer of Anti-C1q Ab in patients with HUVS is consistent with earlier findings reported in the literature.7 Studies also documented positivity of Anti-C1q Ab...
in 21% of patients with antineutrophil cytoplasmic antibody (ANCA) associated vasculitis and 27% of patients with classic PAN.\textsuperscript{10,11} None of our ANCA-associated vasculitis or classic PAN patients had this antibody. This may be due to our small sample size.

Anti-C1q Ab was earlier reported in 5% and 16% of RA and rheumatoid vasculitis patients, respectively.\textsuperscript{12} In our study, 14% of patients with rheumatoid vasculitis had this antibody, thus matching with literature. The prevalence of Anti-C1q Ab levels in SLE varies between 30% and 70%. Anti-C1q Ab, however, is most commonly used in clinical practice to predict nephritis flare in SLE patients. In addition, raised Anti-C1q Ab is reported in lupus pneumonitis and nervous system involvement in patients with SLE.\textsuperscript{13}

Our cohort also had one SLE patient presenting with vasculitis, peripheral nervous system involvement, and gangrene who was
positive for anti C1q antibody. Yet another patient with lupus having urticarial vasculitis and renal involvement also tested positive for Anti-C1q Ab. Both lupus patients had low titers of anti-dsDNA antibody at time of Anti-C1q Ab measurement. Reason for elevated Anti-C1q Ab in these patients cannot be explained by vasculitis per se; is it likely that this antibody may be a marker of current or future development of lupus nephritis?

While studying our primary objectives, we have noted a few clinical correlations of Anti-C1q Ab in vasculitis, which are discussed below. Siegert et al. reported Anti-C1q IgG Ab to be more prevalent in younger patients with SLE; on the other hand, Anti-C1q IgG Ab titer increases with age and more frequent among healthy individuals in older age.14 Orbai et al. also demonstrated Anti-C1q Ab more commonly in younger individuals with lupus as compared to older patients using a cutoff age of 30 years.15 Thus, most of the studies mentioned above including our study demonstrated higher prevalence of Anti-C1q IgG Ab in younger patients with rheumatic diseases.

Anti-C1q Ab also positively correlated with ESR in our study, similar to observations in published reports involving patients with Behcet’s disease with vascular involvement. This correlation may implicate increased disease activity in vasculitic patients similar to lupus nephritis. Positive correlation of low C3 with Anti-C1q antibody reflects activation of complements in vasculitic patients in our study. Complement system is continuously activated by different autoantibodies in SLE patients which causes depletion of C3 and C4 in circulation.16 Anti-C1q Ab, therefore, negatively correlate with C3 and C4 in SLE patients17-19 and HUVS as noted in the present study.

Conclusion

This study found no difference in Anti-C1q Ab levels between our patients with primary and secondary vasculitis. Anti-C1q Ab titers correlated with younger age, high ESR, and low C3 in patients with vasculitis in our study. Therefore, Anti-C1q Ab may act as a potential marker of disease activity in a select subset of younger patients with vasculitis disorders. Therefore, Anti-C1q Ab is likely to have meaningful clinical implications.

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