Hypocomplementemia at Diagnosis of Pauci-immune Glomerulonephritis Is Associated With Advanced Histopathological Activity Index and High Probability of Treatment Resistance

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Introduction: Recent evidence suggests that complement activation is important in the pathogenesis of pauci-immune (PI) vasculitis. This is a retrospective investigation of the frequency of hypocomplementemia at pauci-immune glomerulonephritis (PIGN) diagnosis, in relation to vasculitic manifestations, renal histopathology, and treatment outcomes.

Methods: A total of 115 patients with biopsy-proven PIGN were categorized based on their serum complement C3 (sC3). Histopathology evaluation included activity and chronicity indexes. The primary outcome of interest was treatment resistance, defined as a progressive decline in kidney function, with persistently active urine sediment, leading to dialysis dependency or vasculitis-related death.

Results: In all, 20.9% of patients had low sC3 levels associated with more advanced renal impairment (P < 0.01), requiring acute dialysis (P < 0.01) more frequently compared to patients with normal sC3. Within 1 year, 85.7% of patients with normal sC3 responded to therapy, versus 58.3% of those with low sC3 (P = 0.001). The probability of treatment resistance was strongly associated with low sC3 (P = 0.004), high serum creatinine (P < 0.001), acute dialysis requirement (P < 0.001), and high histopathological score of chronicity (P < 0.01). Advanced histopathological activity was related to more intense interstitial leukocyte infiltration (P = 0.005) and higher likelihood of fibrinoid necrosis documentation in a vessel wall (P = 0.02). The probability of treatment resistance was higher in patients with low sC3 (odds ratio [OR] = 6.47, 95% confidence interval [CI] 1.47–28.35, P = 0.013), oliguria (OR = 29.57, 95% CI = 4.74–184, P < 0.0001), and high chronicity score (OR = 1.77, 95% CI = 1.23–2.54, P = 0.002).

Conclusion: Low sC3 is emerging as an independent predictor of treatment resistance in patients with PIGN associated with higher index of histopathological activity at diagnosis compared to normal sC3.

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KEYWORDS: complement; glomerulonephritis; histopathology; outcome; pauci-immune

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Evidence from animal models and histopathological findings suggest that complement plays a critical role in the pathogenesis of pauci-immune (PI) vasculitis and glomerulonephritis (GN). Pauci-immune vasculitis is most often characterized by acute inflammatory lesions that are extremely destructive, with influx of neutrophils, leukocytoclasis, and necrosis. Observations in human beings and in experimental animal models of PIGNs clearly implicate activation of the alternative complement pathway, with the presence of hypocomplementemia at the onset and increased levels of factors Bb, C3a, and C5a. In an anti-myeloperoxidase anti-neutrophil cytoplasmic antibody (MPO-ANCA) vasculitis murine model, C5a and its
receptor C5aR/CD88 were proved to be critical actors in the development of PIGN.6–8 Wu et al.8 showed that complement activation occurs in both MPO-ANCA and PR3-ANCA vasculitis, but the profile of activation seems to differ between active disease versus disease in remission. Typically, lesions of PI vasculitis are characterized by a paucity of immune deposits, but low-intensity deposition of complement has been reported to be common, especially at focal sites of inflammation and necrosis.8 From a clinical viewpoint, hypocomplementemia in PIGN has been associated with more advanced renal involvement and significantly worst prognosis.10–12

The aim of this study was to investigate what proportion of patients with PIGN have low serum complements at diagnosis, if any, and whether they have different clinical and/or histopathological features or experience different treatment outcomes compared to patients with normal serum complement levels.

MATERIALS AND METHODS

Study Population and Definitions

Patients with PIGN who were recruited by nephrologists or referred from rheumatologists to nephrologists in Laiko Hospital (Athens, Greece) between 2006 and 2019 were studied retrospectively. All patients were identified at or near the initial diagnosis and were eligible to be included in this study if they had the following prerequisites: diagnosis of PIGN in a native kidney biopsy sample with the combined use of light microscopy and immunofluorescence parameters, serum complement measurements at diagnosis (prior to initiation of immunosuppressive therapy), age 16 years or more, and signed informed consent for review of medical charts. For the purpose of histopathological analysis, the included patients were required to have a minimum of 10 glomeruli in the diagnostic kidney biopsy sample. Exclusions were eosinophilic granulomatosis with polyangiitis, overlap syndromes with other diseases (i.e., anti–glomerular basement membrane or lupus), non-adherence issues, and inadequate follow-up time. All patients were tested for anti-neutrophil–associated autoantibodies (ANCA) by immunofluorescence or enzyme-linked immunosorbent assay13 or both. Clinical phenotypes of PI vasculitis were assigned according to the Chapel Hill Vasculitides Nomenclature Consensus Conference.14 Thus, a diagnosis of GPA was defined by the presence of necrotizing granulomatous inflammation in any tissue by histology, and/or imaging showing pulmonary nodules or cavities (noninfectious) and/or bony erosions, and/or subglottic stenosis in the upper respiratory tract. Eosinophilic granulomatosis with polyangiitis was defined by the presence of asthma, eosinophilia, and necrotizing granulomatous inflammation. Microscopic polyangiitis was defined by systemic necrotizing small vessel vasculitis without evidence of granulomatous inflammation or asthma. Organ involvement was defined based on previously described criteria.15

The primary outcome of interest was treatment resistance, which was assessed by the end of the first year, and was determined after a minimum of 3 months of therapy with immunosuppressants. It was defined as a progressive decline in kidney function, with persistently active urine sediment leading to dialysis dependency (with or without new or persistent extrarenal vasculitic manifestations) or vasculitis-related death.

Secondary outcomes of interest included treatment response, remission, relapse, end-stage kidney disease (ESKD) in the long term, and death. Treatment response, which was assessed by the end of the first year after diagnosis and initiation of immunosuppressive treatment, was defined as avoidance of death due to vasculitis, along with dialysis independency (i.e., glomerular filtration rate >15 ml/min per 1.73 m²) with no clinical signs of active systemic vasculitis.16

Remission, following response to immunosuppressive treatment, was defined as the stabilization or improvement of kidney function, as measured by serum creatinine levels, with resolution of hematuria or other manifestations of systemic vasculitis for more than 1 month. Persistent proteinuria with bland urine sediment was not considered indicative of active renal vasculitis. Relapse could be recorded only among patients who had achieved remission, and was characterized by recurrent or new signs and symptoms of active vasculitis in any organ.17,18 End-stage kidney disease was characterized by the initiation of chronic dialysis.

Data Collection

Clinical, laboratory, and serological variables, which were recorded at diagnosis of PIGN, included the following: demographics, clinical phenotype, organ involvement, serum complements C3 and C4 (sC3 and sC4), serum creatinine, peak serum creatinine, estimated glomerular filtration rate (eGFR), oliguria, acute dialysis requirement, and ANCA type. The eGFR was determined using the 4-variable Modification of Diet in Renal Disease equation19 and disease activity was estimated using the Birmingham Vasculitis Score (BVAS).20

Histopathological Evaluation

Histopathological evaluation was performed by an experienced renal pathologist (GL), by review of the
data in diagnostic pathology reports, following a detailed scoring system according to the concept presented by Lee et al. Specifically, the number of normal glomeruli was presented as a percentage of the total (grade 4 if <10%, 3 if ≥10 <25%, 2 if ≥25% to <50%, and 1 if ≥50%). Glomerular and tubulointerstitial lesions were scored separately and as part of the activity index and chronicity index scores (Table 1). The activity index score represented the sum of scores from 6 categories: glomerular necrosis, cellular crescents, interstitial leukocyte infiltration, fibrinoid necrosis in vessel walls, red blood cell (RBC) casts, and circumferential crescents. Each of the first 3 categories was scored semi quantitatively from 1 (none/mild) to 3 (severe), and the other 3 categories were scored as 1 if the characteristic was present in the specimen or as 0 if it was absent, for a maximum of 12 points. The feature of cellular crescents was defined as the percentage of the involved glomeruli out of the total number of glomeruli in the biopsy sample (grading: 1 for 1%–25%, 2 for 26%–50%, and 3 for >50%). Glomerular necrosis was defined as the percentage of the involved glomeruli out of the total number of glomeruli in the biopsy sample (grading: 1 for 1%–25%, 2 for 26%–50%, and 3 for >50%). Interstitial leukocyte infiltration was defined as the percentage of the inflamed interstitium out of the total (grading: 1 for 1%–25%, 2 for 26%–50%, and 3 for >50%). Red blood cell casts were noted as 0 for absent and 1 for present). Circumferential crescents were noted as 0 for absent and 1 for present. Necrosis in the wall of arterioles, interlobular or arcuate arteries were noted as 0 for absent and 1 for present. The chronicity index score represented the sum of scores from 4 categories, namely, global glomerulosclerosis, interstitial fibrosis, tubular atrophy and fibrotic crescents/segmental sclerosis, which were scored semiquantitatively from 1 to 3 (grade 1 for 0%–25%, grade 2 for 26%–50%, and grade 3 for >50%) for a maximum of 12 points. Fibrotic crescents/segmental sclerosis was defined as the sum of fibrotic crescents (crescents containing >50% fibrosis) and segmental glomerular sclerosis (unspecified sclerosis, probably due to a repair process after necrosis or crescent formation but without cellular remnants), reflected as a percentage of the total number of the glomeruli (grading from 1 to 3). Immunofluorescence examination on frozen tissue was applied in all cases for immunoglobulins (IgG, IgA, IgM), complement components (C3 and C1q), fibrinogen, and kappa and lambda light chains (DAKO FITC, polyclonal rabbit 1/50 dilution). Traces were defined as stain intensity of <1+, and all others stain intensities were recorded on a 4-grade scale (0–3).

**Statistical Analyses**

Continuous variables were expressed as the mean and standard deviation, whereas categorical variables as frequencies and percentages on the basis of the serum C3 level at the time of diagnosis of PIGN. To investigate the differences between clinical and histopathological variables between patients with normal and low C3 levels, the t test and Mann–Whitney U test for independent samples for continuous variables and the χ² and Fisher exact test for categorical variables were applied. Univariate logistic regression analyses were performed for different outcomes (remission, response to immunosuppressive therapy, ESKD, death) and time periods (at the first year after diagnosis, from diagnosis to end of follow-up), and Cox regression analysis for investigating the association between the renal survival time of the patients and their clinical and histopathological characteristics. Variables that were found to be significant in the univariate analyses were included in the multivariate models. The estimated odds ratios (ORs) and hazard ratios (HRs) of both the univariate and multivariate models, as well as the 95% confidence intervals (CIs) of the ORs/HRs and the related P values, are presented. Given the short time period after diagnosis (12 months) that was set for estimations of the first outcome, only a small number of deaths were reported. For this reason, when implementing models of logistic regression analysis for the estimation of a possible relation between the measured variables and the outcomes in the first year, those 4 patients were included in the group of patients with an “adverse outcome,” which included patients who developed ESKD or died during the same period of time. Time to ESKD (in the long-term) was calculated using the Kaplan–Meier estimates, and categories were compared using the log-rank test. The low sC3 variable, albeit not significant from the univariate analysis, was included in the multivariate model for the first-year outcome, as it was the primary explanatory variable for this study. Data were analyzed using Stata 13.0 software (Stata Corporation, College Station, TX), and significance was set at α = 0.05. All tests proceeded as 2 tailed.

**RESULTS**

**Baseline Characteristics of the Cohort**

A total of 115 patients with biopsy-proven PIGN were included in the study, 24 (20.9%) of whom had sC3 values below the lower limit of normal range (i.e., <90 mg/dl). One patient (0.8%) had sC4 below the lower
Table 1. Comparison of demographics, disease-related characteristics, and histopathological parameters of patients with pauci-immune glomerulonephritis with low or normal serum C3 complement at diagnosis

| Characteristic                          | Normal C3 n = 91 | Low C3 n = 24 | P value |
|----------------------------------------|------------------|---------------|---------|
| Patients                               | 91 (79.1%)       | 24 (20.9%)    |         |
| Sex, male                              | 46 (50.5)        | 14 (58.3)     | 0.49    |
| Race, Caucasian                        | 88 (96.7)        | 23 (95.8)     | 1       |
| Age, yr                                | 56.8 (17.5)      | 57.6 (17.5)   | 0.82    |
| ANCA type                              |                  |               |         |
| Negative                               | 5 (5.5)          | 2 (8.3)       | 0.69    |
| C/ PR3-ANCA                            | 30 (30)          | 9 (37.5)      |         |
| P/MPO-ANCA                             | 56 (61.5)        | 13 (54.2)     |         |
| Clinical phenotype                     |                  |               |         |
| Renal limited disease                  | 29 (31.8)        | 11 (45.8)     | 0.46    |
| Granulomatosis with polyangiitis       | 22(24.2)         | 4 (16.7)      |         |
| Microscopic polyangiitis               | 40 (44)          | 9 (37.5)      |         |
| Oliguria                               | 35 (38.5)        | 12 (50)       | 0.30    |
| Acute dialysis requirement             | 20 (22)          | 13 (54.2)     | < 0.01  |
| Serum creatinine, mg/dl                | 3.5 (2.2)        | 5.3 (3.4)     | < 0.01  |
| Estimated GFR, ml/min/1.73 m²          | 24.7 (22.8)      | 18.3 (15.0)   | 0.11    |
| Peak serum creatinine, mg/dl           | 4.5 (2.7)        | 6.4 (3.5)     | 0.02    |
| BVAS score                             | 17.4 (6.7)       | 17.5 (8.2)    | 0.83    |
| Immunosuppressive therapy (initial)    |                  |               |         |
| Cyclophosphamide + glucocorticoids     | 91 (100)         | 24 (100)      |         |
| Plasma exchange                        | 21 (29.6)        | 9 (37.5)      | 0.45    |
| Rituximab                              | 18 (19.7)        | 7 (29.2)      | 0.25    |
| Histopathological parameters           |                  |               |         |
| Normal glomeruli (%)                   | N=85             | N=24          | 0.51    |
| Grade 1, >50%                          | 12 (14.1)        | 3 (12.5)      |         |
| Grade 2, 25%-50%                       | 21 (24.7)        | 9 (37.5)      |         |
| Grade 3, 10%-24%                       | 22 (25.8)        | 9 (37.5)      |         |
| Grade 4 <10%                           | 30 (35.3)        | 8 (33.3)      |         |
| Normal glomeruli (>10%)                | 55 (64.7)        | 16 (66.7)     | 0.8     |
| EUVAS categories                       | 0.5              |               |         |
| Focal class                            | 21 (24.7)        | 4 (16.7)      |         |
| Crescentic class                       | 21 (24.7)        | 6 (25)        |         |
| Mixed class                            | 32 (37.7)        | 8 (33.3)      |         |
| Sclerotic class                        | 11 (12.9)        | 6 (25)        |         |
| Global glomerulosclerosis              | 0.8              |               |         |
| Mild (0%-25%)                          | 52 (61.2)        | 15 (62.5)     |         |
| Moderate (25%-50%)                     | 18 (21.2)        | 4 (16.6)      |         |
| Severe (>50%)                          | 15 (17.6)        | 5 (20.9)      |         |
| Fibrillar crescents/segmental sclerosis| 0.8              |               |         |
| None/mild (0%-25%)                     | 59 (70.2)        | 19 (79.2)     |         |
| Moderate (25%-50%)                     | 14 (16.7)        | 3 (12.5)      |         |
| Severe (>50%)                          | 12 (14.1)        | 2 (8.3)       |         |
| Tubular atrophy                        | 0.08             |               |         |
| None/mild (0%-25%)                     | 42 (48.4)        | 16 (66.7)     |         |
| Moderate (25%-50%)                     | 39 (45.9)        | 6 (25)        |         |
| Severe (>50%)                          | 4 (4.7)          | 2 (8.3)       |         |
| Interstitial fibrosis                  | 0.65             |               |         |
| None/mild (0%-25%)                     | 37 (42.5)        | 13 (54.2)     |         |
| Moderate (25%-50%)                     | 42 (48.4)        | 10 (41.7)     |         |
| Severe (>50%)                          | 6 (7.1)          | 1 (4.1)       |         |
| Chronicity (1–12)                      | 5.5 (2.4)        | 5.2 (2.5)     | 0.58    |
| Cellular crescents                     | 0.17             |               |         |
| Grade 1, 0%-25%                        | 42 (49.4)        | 7 (29.2)      |         |
| Grade 2, 25%-50%                       | 21 (24.7)        | 10 (41.6)     |         |
| Grade 3, >50%                          | 22 (25.9)        | 7 (29.2)      |         |

Data are n (%) or mean (SD). EUVAS, European Vasculitis Society; GFR, glomerular filtration rate.

normal limit and 1 patient had sC4 above the upper normal limit. Patients were categorized according to the measurement of sC3 into 2 groups, 1 group with low sC3 and 1 group with normal sC3. No differences were identified in terms of demographics (Table 1 and Supplementary Table S1) and disease-related characteristics. Patients with low sC3 had more severe renal involvement, that is, higher serum creatinine and lower eGFR, and they were more likely to experience oliguria and to require acute dialysis around the initial diagnosis (P < 0.01). Data from 109 renal biopsy samples were reviewed. The pattern of glomerular injury according to the European Vasculitis Society (EUVAS) classification schema was similar between groups. The mean histopathological activity score, assessed in the diagnostic biopsy sample, was significantly higher in patients with low sC3 than in patients with normal sC3 (6.33 vs. 4.98, P = 0.01) (Table 1). Advanced histopathological activity was related to more intense interstitial leukocyte infiltration (P = 0.005) in the renal tissue and higher likelihood of documented fibrinoid necrosis in a vessel wall in a certain specimen (P = 0.02) (Figure 1a, b). Histopathological parameters related to chronic renal damage were similar between groups, as were immunofluorescence findings (Figure 1c, d). All patients were treated with a combination of cyclophosphamide (intravenous or oral) and glucocorticoids, and similar proportions of patients in both groups also received plasma exchange or rituximab as part of the initial treatment (Table 1).
Incidence and Predictors of Treatment Failure by the End of the First Year

By univariate analysis, the composite outcome of ESKD or death (Supplementary Table S2) was significantly associated with a low sC3; that is, patients with low sC3 at diagnosis were 4 times more likely to experience a worst outcome (OR = 4.28, 95% CI = 1.57–1.67, \( P = 0.004 \)). A higher baseline serum creatinine value was associated with higher probability of ESKD or vasculitis-related death, and patients with oliguria at presentation were found to have a 10.85 times higher likelihood of ESKD or vasculitis-related death. In terms of histopathology, patients with <10% normal glomeruli in the diagnostic biopsy, severe arteriosclerosis, moderate or severe global glomerulosclerosis, and cellular crescents in >50% of the glomeruli in the biopsy sample were more likely to develop ESKD or death. An activity score of \( \geq 8 \) in the diagnostic biopsy sample was associated with a 4-fold higher probability of ESKD or death at the end of the first year, compared

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**Figure 1.** Histopathological parameters in the diagnostic kidney biopsy samples of patients with pauci-immune glomerulonephritis. (a) Histopathological parameters of activity in patients with low serum C3. (b) Histopathological parameters of activity in patients with normal serum C3. (c) Histopathological parameters of chronicity in patients with low serum C3. (d) Histopathological parameters of activity in patients with normal serum C3. (Continued)
with an activity score of \(<8\) (Supplementary Table S2). There was a strong correlation between the chronicity score and the probability of a poor outcome, as patients with a chronicity score of \(\geq 8\) were shown to be 4 times more likely to experience a poor outcome compared to patients with a lower chronicity score \((P = 0.004)\).

After adjustments, multivariable analysis showed that low sC3 at diagnosis was independently associated with a worse first-year outcome (i.e., ESKD or death), as well as oliguria, moderate or severe global glomerulosclerosis, and cellular crescents in more than 50% of the glomeruli examined. Moreover, low sC3 at diagnosis conferred a 6.47-fold likelihood of ESKD or death within the first 12 months of diagnosis (compared to normal sC3) \((OR = 6.47, 95\% CI = 1.47–28.35, P = 0.013)\); significant acute renal dysfunction with

![Figure 1. (Continued)](image-url)
oliguria conferred an almost 30-fold higher likelihood of ESKD or vasculitis-related death (compared to absence of oliguria) (OR = 29.57, 95% CI = 4.74–184, P < 0.0001), and a chronicity score of ≥8 conferred a 1.77-fold higher likelihood of a worse outcome (compared to lower chronicity scores) (OR = 1.77, 95% CI = 1.23–2.54, P = 0.002) (Table 2). After adjusting for covariates, a sC3 value below normal at diagnosis conferred a 81% higher probability of treatment resistance, compared to normal sC3 value (OR = 0.19, 95% CI = 0.03–0.96, P = 0.04) (Supplementary Table S3).

Incidence and Predictors of Treatment Response

Treatment response was more common in patients with normal sC3 at diagnosis (87.9% vs. 58.3%, P = 0.001). At the end of first year, dialysis independency was significantly more frequent in patients with normal sC3 compared to those with normal sC3 values (41.7% vs. 9.8%, P = 0.002). During the first year, 4 deaths occurred, all in the normal sC3 group, with 2 of them attributed to treatment-resistant vasculitis, 1 attributed to a cardiovascular event, and 1 to sepsis. By univariate analysis, treatment response was associated with normal sC3 (OR = 0.19, 95% CI = 0.06–0.53, P = 0.002), lower serum creatinine, absence of oliguria, no need of acute dialysis near diagnosis, percentage of normal glomeruli >10%, EUVAS focal class (vs. sclerotic class), and chronicity score per unit increase (Supplementary Table S4). Each unit of increase in the chronicity score was associated with a 66% reduction in the probability of responding to treatment (P = 0.006) (Table 3). No correlation was observed between sC3 and the intensity of C3 in the renal tissue by immunofluorescence or between patients with normal and low sC3 (P = 0.31), whereas differences were noted with respect to immunoglobulins IgA, IgM, and IgG.

Incidence and Predictors of ESKD Among Responders

Among patients who responded to initial treatment and achieved remission by the end of the first year, the probability of ESKD in the long term (median follow-up of 57 [45 + 12] months) was higher for patients with low sC3 levels at diagnosis (16.7% vs. 35.7%, P = 0.1) (Figure 2). Moreover, the median renal survival time after remission was significantly shorter for patients who had low sC3 (6.5 years vs. 11.5 years, P = 0.04). Relapse rates among patients who responded to therapy and achieved remission were similar between groups, although there were 7 deaths in the normal sC3 group and 1 death in the low sC3 group during the total observation period.

By multivariate analysis, C3 hypocomplementemia at diagnosis (HR = 1.99, 95% CI = 0.51–7.70, P = 0.31), was not associated with the probability of requiring chronic dialysis (Table 4). Lower levels of serum creatinine, EUVAS focal class (vs. sclerotic), and the presence of RBC casts in the diagnostic biopsy sample were the factors that were associated with lower probability of requiring chronic dialysis as shown by the multivariate analysis.

**DISCUSSION**

In this retrospective study, we compared outcomes of patients with biopsy-proven PIGN who were grouped based on the level of sC3 measurements at diagnosis.
Low serum C3 was associated with a higher probability of ESKD or disease-related death within the first year.12,30 In this cohort, the higher histopathological activity index compared to patients with normal sC3, as reflected in more severe interstitial leukocyte infiltration and advanced cellular crescent formation. Overall, treatment resistance was also associated with a higher percentage of normal glomeruli, an advanced chronicity score (especially a chronicity score ≥8), crescentic class by the EUVAS categorization,21 and severe arteriosclerosis. Importantly, low sC3 at diagnosis was also associated with a higher probability of requiring chronic dialysis in the long term.

Treatment-resistant PI vasculitis has been reported in randomized clinical trials22-27 to occur in 10% to 40% of enrolled patients, as well as in clinical practice.28,29 Certain risk factors for treatment resistance were previously recognized, including older age,28,29 severe kidney disease, female sex, African American race, and the presence of MPO-ANCA. Low baseline sC3 is emerging as an independent predictor of treatment resistance, as it conferred a 6.47-fold likelihood of ESKD or disease-related death within the first 12 months. Patients with low sC3 were found to have a higher histopathological activity index compared to patients with normal sC3, as reflected in more severe interstitial leukocyte infiltration and advanced cellular crescent formation. Overall, treatment resistance was also associated with a higher percentage of normal glomeruli, an advanced chronicity score (especially a chronicity score ≥8), crescentic class by the EUVAS categorization,21 and severe arteriosclerosis. Importantly, low sC3 at diagnosis was also associated with a higher probability of requiring chronic dialysis in the long term.

Table 4. Risk factors for ESKD in the long term among patients with pauci-immune glomerulonephritis who responded to initial immunosuppressive therapy and achieved remission by the end of the first year

| Variable (at diagnosis) | Hazard ratio (95% CI) | P value |
|------------------------|----------------------|---------|
| Multivariable model    |                      |         |
| Low serum C3           | 1.99 (0.51–7.70)     | 0.31    |
| Serum creatinine       | 1.31 (1.03–1.66)     | 0.02    |
| Normal glomeruli >10% (versus ≤10%) | 0.61 (0.13–2.85) | 0.53 |
| EUVAS category         |                      |         |
| Focal class versus sclerotic class | 0.06 (0.005–0.75) | 0.03 |
| Crescentic class versus sclerotic class | 0.44 (0.08–2.39) | 0.34 |
| Mixed class versus sclerotic class | 0.62 (0.11–3.40) | 0.6 |
| Interstitial fibrosis (moderate or severe versus none/mild) | 1.06 (0.28–4.01) | 0.92 |
| RBC casts present in specimen | 0.11 (0.01–0.66) | 0.01 |

CI, confidence interval; ESKD, end-stage kidney disease; GFR, glomerular filtration rate; RBC, red blood cell; EUVAS, European Vasculitis Society.

Patients with low sC3 at baseline represented one-fifth of the total cohort, had more severe renal impairment at presentation, and required acute dialysis more frequently. Low sC3 at diagnosis of PI vasculitis has been previously described,8-12 with Fukui et al.10 reporting a rate similar to the one found in this study. More importantly, low sC3 at presentation was shown from our study to be an independent predictor of treatment resistance, as it conferred a 6.47-fold likelihood of ESKD or disease-related death within the first 12 months. Patients with low sC3 were found to have a higher histopathological activity index compared to patients with normal sC3, as reflected in more severe interstitial leukocyte infiltration and advanced cellular crescent formation. Overall, treatment resistance was also associated with a higher percentage of normal glomeruli, an advanced chronicity score (especially a chronicity score ≥8), crescentic class by the EUVAS categorization,21 and severe arteriosclerosis. Importantly, low sC3 at diagnosis was also associated with a higher probability of requiring chronic dialysis in the long term.

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Figure 2. Renal survival of patients with pauci-immune glomerulonephritis (PIGN), who responded to initial therapy and achieved remission, stratified by serum C3 at diagnosis.
might be related to enhanced activation of the alternative complement pathway. First, the complement system can be engaged by activation of the classic, lectin, and/or alternative pathway, leading to C3 to divide into C3a and C3b. C3b allows for the formation of C5 convertase that cleaves C5 into C5a and C5b and the assembly of the membrane attack complex C5b-9. Importantly, in a murine model of PIGN, the knockout of C5 or factor B but not the C4 or C6 abrogated PIGN formation, which indicates that the alternative complement pathway and not the classic one or membrane attack complex C5b-9 is implicated in renal injury. 

Subsequently, C5a derived from complement activation was shown to cause priming of neutrophils, resulting in the release of small amounts of ANCA antigens, which interact with ANCA with further neutrophil activation. Properdin secretion by activated neutrophils enhances the activation of the alternative pathway of the complement system, whereas activated neutrophils and monocytes activate complement, generating C5a. C5a functions as a chemoattractant to neutrophils and monocytes. Xiao et al. showed that knocking out C5 resulted in strong attenuation of glomerulonephritis, whereas knocking out C4 had no effect in the disease phenotype. We did not find any correlation between sC3 and the intensity of C3 in the renal tissue by immunofluorescence, between patients with normal and low sC3. In contrast, a study including 85 patients with PIGN in which renal histopathology was studied using the Berden categorization found that C3d-positive staining, which was observed in 49.4% of the biopsy samples, was associated with the severity of renal impairment and a lower rate of response to treatment. 

The main limitation of this study is its retrospective design. However, it has also important strengths, including the fact that all patients had biopsy-proven disease, and that histopathological evaluation followed a comprehensive scoring system for all kidney biopsy specimens for assessment of activity and chronicity, similar to the report by Lee et al., with excellent correlations with clinical outcomes, and especially response to initial therapy. In conclusion, in this study one-fifth of patients with PIGN had C3 hypocomplementemia at diagnosis. These patients had advanced activity indexes by histopathology and were more likely to experience treatment resistance compared with patients with normal sC3. If hypocomplementemia was a result of activation of the alternative complement pathway in these patients, and considering that increased activity was associated with enlarged interstitial leukocyte infiltration and significant crescent formation, one might speculate that the intensity and/or duration of complement activation correlates with the severity of the clinical picture and response to treatment. Identification of patients with PIGN with significant complement activation reflected in low sC3 at diagnosis might be helpful in terms of planning the type of immunosuppressive regimen, which might include agents targeting the complement system at the first place at one or multiple sites, to attenuate stimulation of neutrophils, to control inflammation, and to avoid treatment resistance.

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