Genomic Copy Number Variations of the Complement Component \textit{C4B} Gene Are Associated With Chronic Central Serous Chorioretinopathy

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PURPOSE. Chronic central serous chorioretinopathy (cCSC) has recently been associated to variants in the complement factor \textit{H} gene. To further investigate the role of the complement system in CSC, the genomic copy number variations in the complement component 4 \textit{gene (C4)} were studied.

METHODS. \textit{C4A} and \textit{C4B} copy numbers were analyzed in 197 cCSC patients and 303 healthy controls by using a Taqman copy number determination assay. Copy numbers of \textit{C4A}, \textit{C4B}, and the total \textit{C4} load were compared between cases and controls, by using a Fisher exact test. For this analysis Bonferroni correction was performed for three tests, and \textit{P} values < 0.014 were considered to be significant. A logistic regression model was constructed to calculate the odds ratios (ORs) of each of the \textit{C4B} copy numbers, using two copies as a reference. For this model \textit{P} values < 0.05 were considered to be significant.

RESULTS. \textit{C4B} genomic copy numbers differed significantly between cCSC patients and healthy controls (\textit{P} = 0.0018). Absence of \textit{C4B} significantly conferred risk of cCSC (\textit{P} = 0.039, OR = 2.61 [95% confidence interval (CI) = 1.05–6.52]), whereas three copies of \textit{C4B} significantly decreased the risk of cCSC (\textit{P} = 0.014, OR = 0.45 [95% CI = 0.23–0.85]). The \textit{C4A} genomic copy numbers and total \textit{C4} load did not significantly differ between cases and controls.

CONCLUSIONS. This study showed that copy numbers of \textit{C4B} are significantly associated with cCSC. Carrying no copies of \textit{C4B} significantly increases the risk of cCSC, whereas carrying three \textit{C4B} copies is protective. These findings reinforce the hypothesis of a possible involvement of the complement system in the pathogenesis of cCSC.

Keywords: chronic central serous chorioretinopathy, cCSC, complement component 4, \textit{C4}, \textit{C4A}, \textit{C4B}
TABLE 1. Demographics of the Study Population

<table>
<thead>
<tr>
<th></th>
<th>cCSC Patients</th>
<th>Controls</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>No. of subjects</td>
<td>197</td>
<td>303</td>
<td>NA</td>
</tr>
<tr>
<td>Sex, male/female</td>
<td>154/43</td>
<td>226/77</td>
<td>0.392</td>
</tr>
<tr>
<td>Age, mean ± SD, y</td>
<td>53 ± 10</td>
<td>53 ± 11</td>
<td>0.755</td>
</tr>
<tr>
<td>Age range, y</td>
<td>29–74</td>
<td>29–77</td>
<td>NA</td>
</tr>
</tbody>
</table>

NA, not annotated; SD, standard deviation.

MATERIALS AND METHODS

Subjects

In this study, 197 patients diagnosed with cCSC who visited the outpatient clinic of the Department of Ophthalmology at the Radboud University Medical Center, Nijmegen, the Netherlands, were included (Table 1). The diagnosis cCSC was based on an extensive ophthalmologic examination including fundoscopy, spectral-domain optical coherence tomography, fluorescein angiography, and indocyanine green angiography. The definition of typical cCSC used in this study was based on the previously published subgroups by de Jong et al. and patients in this study undergone phenotyping by an experienced retina specialist (CJFB) (Fig. 1A–F). Additionally, a total of 303 control subjects were recruited from the blood bank of the Radboud University Medical Center (n = 154) and the European Genetic Database (EUGENDA, www.eugenda.org; provided in the public domain by the University Hospital of Cologne, Cologne, Germany and the Radboud University Medical Center) (n = 149) (Table 1). For this last group, fundus photographs were graded to rule out any ophthalmologic abnormalities at the moment of inclusion. Informed consent for the use of DNA for genetic studies was obtained from all subjects. This study followed the guidelines of the Declaration of Helsinki and was approved by the local ethics committee.

Copy Number Determination

DNA was isolated from peripheral blood by using standard procedures. C4A and C4B copy numbers were determined by real-time PCR using Taqman genotyping assays (Applied Biosystems, Thermo Fisher Scientific, Waltham, MA, USA). The FAM-labeled C4A (Hs07226349_cn) or C4B (Hs07226350_cn) Taqman copy number assay was combined with the VIC-labeled Ribonuclease P (RNaseP) reference assay (catalog No. 4403326), and Taqman genotyping mastermix (catalog No. 4381656). All samples were tested in duplicate for C4A and C4B on 384-wells plates by using 10 ng DNA in a total reaction volume of 10 μL.

Samples with known copy numbers for either C4A (0–4) or C4B (0–5) were kindly provided by C. Yung Yu. These samples were included as a reference on each plate to facilitate accurate copy number determination, using the method described previously. In each run the amplification efficiencies of the C4A/C4B and RNaseP probes were calculated by using a serial dilution (50–1.56 ng) of a sample with two C4A and C4B copies. The primer efficiencies of the probes were compared and deemed similar if they differed <2%; this was the case in all runs. Therefore, the efficiencies were not incorporated into the calculations of the copy numbers. Polymerase chain reaction was performed with a 7900HT thermocycler (Applied Biosystems, Thermo Fisher Scientific) using the following program: 2 minutes at 50°C, 10 minutes at 95°C, and 40 cycles of 15 seconds at 95°C and 60 seconds at 60°C. Data were analyzed with the Copycaller software (V2.0; Applied Biosystems, Thermo Fisher Scientific). Copy numbers determined by the Copycaller software were corrected by using the trend line based on the reference samples, as described before. If the results were inconsistent between the Copycaller output and the values corrected with the trend line, samples were retested on a new plate in triplicate.

Statistics

The comparison of C4A, C4B, and total C4 copy number distribution between cCSC patients and controls was performed by means of a Fisher exact test using SPSS Statistics (V20; IBM Corp., Armonk, NY, USA). Bonferroni correction for multiple testing was performed for three tests and P values < 0.017 were considered to be statistically significant. A logistic regression model was constructed to determine the odds ratios (ORs) for the various copy numbers of C4B. According to previously published studies, two genomic copy numbers of C4B are considered to be most common in the healthy population. We were able to confirm this in our cohort, and therefore this copy number was set as reference. In this model, P values < 0.05 were considered to be significant. Graphs were generated by using Graphpad Prism (V5; Graphpad Software, San Diego, CA, USA).

RESULTS

The copy numbers of C4A and C4B were successfully determined in 197 cCSC cases and 303 controls. No significant difference was observed between cases and controls for the C4A genomic copy number (range: 0–6, P = 0.649; Fig. 2A).
The C4B distribution was significantly different between cCSC patients and controls (range: 0–4, P = 0.0018; Fig. 2B). Overall, cases carried lower copy numbers of C4B than the control population. The total C4 genomic copy number was not different in cases compared to controls (P = 0.148; Fig. 2C). Age and sex were not associated with C4A, C4B, or total C4 genomic copy number (Table 2; data for C4A and total C4 not shown).

To assess the effect size of the different copy numbers of C4B on development of cCSC, a logistic regression was performed (Table 2). The logistic regression model based on the distribution of C4B between cases and controls was significant (P = 0.0035; Table 2). Carrying no copies of C4B conferred increased risk of cCSC (P = 0.039, OR = 2.61, 95% confidence interval [CI] = 1.05–6.52). A similar trend was observed for carriers of one copy of C4B, but the results were not significant (P = 0.080, OR = 1.47, 95% CI = 0.96–2.26). Carrying three C4B copies was associated with a significantly decreased risk of cCSC (P = 0.014, OR = 0.45, 95% CI = 0.24–0.85), whereas no significant association with cCSC was observed in individuals carrying four copies of C4B (P = 0.81).

**DISCUSSION**

Our study results demonstrated that cCSC patients have a significantly different C4B load as compared to healthy controls (P = 0.0018). Carrying no copies of C4B was associated with an increased risk of cCSC (OR = 2.61, 95% CI = 1.05–6.52), whereas carrying three C4B copies was associated with a decreased risk of cCSC (OR = 0.45, 95% CI = 0.24–0.85). No association with cCSC was observed in individuals carrying four C4B copies, which is likely due to the limited sample size of this group (cases, n = 4; controls, n = 2). No significant differences were observed between cases and controls for C4A and total C4 load.

The C4 gene lies within the RP-C4-CYP21-TNX (RCCX) locus located in the major histocompatibility complex (MHC) region III on chromosome 6 of the human genome. The MHC region contains an elevated level of genomic copy number variations that are presumably present to increase immunologic diversity. Duplications and deletions in the region have led to the formation of haplotypes containing variable copies of the RCCX locus in the human population (Fig. 3B). The C4 gene encodes the C4 protein, of which two variants have been described (C4A/C4B), differing in only four amino acids encoded by exon 26 (Fig. 3A). Copy number variations of either C4A or C4B have been associated with several systemic diseases with ocular involvement, such as Vogt-Koyanagi-Harada disease, Behçet’s disease, and systemic lupus erythematosus (SLE).13-15 Hou et al.13 have shown that a lower copy number of C4A and C4B increases the risk of Vogt-Koyanagi-Harada disease, an autoimmune disorder characterized by bilateral granulomatous panuveitis. The same group14 has also demonstrated that higher copy numbers of C4A confer risk of Behçet’s disease, an autoinflammatory disease, which presents with acute anterior uveitis. Several studies15,25,26 have shown an association between low copy numbers of C4A and an increased risk for SLE, an autoimmune disease that is typically mediated by immune complexes. In the past, CSC has been described in SLE patients, but it remains unclear whether this is a primary manifestation of SLE, or whether it is a consequence of corticosteroid treatment for SLE. Several studies have reported a positive linear correlation between serum C4 and C4 genomic copy number, suggesting that the lower number of C4B copies in cCSC patients leads to lower systemic C4B levels. This may indicate that an overall lower activity of the complement system might be present in cCSC patients.
Recent low copy numbers of \textit{C4B} have been associated with hyperreactivity of the HPA axis.\textsuperscript{16} Banlaki et al.\textsuperscript{16} have shown that in patients with adrenal incidentaloma and low (<2 copies) \textit{C4B} genomic copy number, baseline ACTH is significantly reduced as compared to high (≥2 copies) genomic copy number of \textit{C4B}. Moreover, a significantly higher cortisol response is observed after ACTH stimulation in the patients with low \textit{C4B} genomic copy number.\textsuperscript{16}

These results are of interest in the context of cCSC because of the described clinical associations with stress and the use of corticosteroids that both exert physiological effects at the level of the HPA axis.\textsuperscript{17} Various relatively small studies have studied cortisol levels in cCSC patients. Although 24-hour urine samples show elevated cortisol levels in cCSC patients in certain studies,\textsuperscript{34,35} these results are not observed in single samples show elevated cortisol levels in cCSC patients. Although 24-hour urine serum measurements during set times in other studies.\textsuperscript{36,37} These discrepancies could be explained by variable cortisol fluctuations between individuals during the day, and therefore changes in endogenous cortisol levels cannot be ruled out as a hallmark of cCSC. It is possible that patients with cCSC generally have normal cortisol levels but respond differently to stimulation of the HPA axis. Stress, which also appears to be associated with cCSC,\textsuperscript{2,38} stimulates the HPA axis and could lead to temporarily elevated cortisol levels in patients as compared to healthy individuals. How high levels of cortisol can lead to subretinal fluid accumulation is currently unknown. A study in rats suggests that the disease mechanism could be mediated by binding of corticosteroids to the mineralocorticoid receptor.\textsuperscript{39} In this study, activation of the mineralocorticoid receptor causes vascular effects similar to those observed in cCSC,\textsuperscript{39} but the underlying pathways still remain to be elucidated.

The mechanism through which low copy numbers of \textit{C4B} may lead to hyperresponsiveness of the HPA axis is unclear. It has been hypothesized that it is not the \textit{C4B} gene, but rather the neighboring \textit{CYP21A2} gene, that mediates this effect.\textsuperscript{16} The \textit{CYP21A2} gene encodes the enzyme 21-hydroxylase, which plays an important role in the steroid metabolism pathway by converting progesterone and 17-β-hydroxyprogesterone to 11-deoxycorticosterone and 11-deoxycortisol, respectively. Because of the genomic structure of the \textit{RCCX} locus, variation in the \textit{CYP21A2} gene is in high linkage disequilibrium with variation in the neighboring \textit{C4} gene (\textit{C4A} or \textit{C4B}).\textsuperscript{16} Therefore, further exploration of the precise structure and specific variations present in the \textit{RCCX} locus may reveal new insights into the pathogenesis of cCSC.

The current study and previous studies identified an association between complement genes and cCSC,\textsuperscript{7,8} suggesting that the complement system may be dysregulated in cCSC. Taken together, these findings may indicate that the immune system, influenced by environmental factors such as stress, could play a pivotal role in the pathophysiology of cCSC. Further studies are necessary to determine the physiological effects of genetic variation at the \textit{C4} gene and the \textit{RCCX} locus in cCSC.

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C4B in Chronic Central Serous Chorioretinopathy

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