The Major Autoantibody Epitope on Factor H in Atypical Hemolytic Uremic Syndrome Is Structurally Different from Its Homologous Site in Factor H-related Protein 1, Supporting a Novel Model for Induction of Autoimmunity in This Disease*

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Background: It is unknown why patients with autoantibodies against complement factor H (CFH) lack homologous CFHR1 protein.

Results: The autoantibody epitope on CFH was identified, and the structure of the corresponding part of CFHR1 was solved.

Conclusion: The autoantigenic epitope of CFH and its homologous site in CFHR1 are structurally different.

Significance: A plausible explanation for formation of autoantibodies due to CFHR1 deficiency in autoimmune atypical hemolytic uremic syndrome was obtained.

Atypical hemolytic uremic syndrome (aHUS) is characterized by complement attack against host cells due to mutations in complement proteins or autoantibodies against complement factor H (CFH). It is unknown why nearly all patients with autoimmune aHUS lack CFHR1 (CFH-related protein-1). These patients have autoantibodies against CFH domains 19 and 20 (CFH19–20), which are nearly identical to CFHR1 domains 4 and 5 (CFHR14–5). Here, binding site mapping of autoantibodies from 17 patients using mutant CFH19–20 constructs revealed an autoantibody epitope cluster within a loop on domain 20, next to the two buried residues that are different in CFH19–20 and CFHR14–5. The crystal structure of CFHR14–5 revealed a difference in the conformation of the autoantigenic loop in the C-terminal domains of CFH and CFHR1, explaining the variation in binding of autoantibodies from some aHUS patients to CFH19–20 and CFHR14–5. The autoantigenic loop on CFH seems to be generally flexible, as its conformation in previously published structures of CFH19–20 bound to the microbial protein OspE and a sialic acid glycan is somewhat altered. Cumulatively, our data suggest that association of CFHR1 deficiency with autoimmune aHUS could be due to the structural difference between CFHR1 and the autoantigenic CFH epitope, suggesting a novel explanation for CFHR1 deficiency in the pathogenesis of autoimmune aHUS.

Atypical hemolytic uremic syndrome (aHUS) is a rare and often fatal systemic disease characterized by hemolytic anemia, thrombocytopenia, microvascular thrombosis, and kidney failure (1). It is associated with dysregulation of complement activation via mutations, polymorphisms, or rearrangements in genes coding for various complement proteins (2). The mutations are found mainly in the gene coding for complement factor H (CFH) (3, 4), which mediates elimination of the central complement activation component C3b. We and others (5–8) have shown how mutations in CFH domains 19 and 20 (CFH19–20) cause impaired regulation of C3b on host cells, leading to complement attack against red blood cells, platelets, and endothelial cells as seen clinically in aHUS. However, some aHUS cases are caused by autoantibodies against CFH (CFH-AAs). These antibodies have been identified in 5–11% of aHUS patients in different cohorts (9–13), but even 56% of 246 HUS patients have been reported with CFH-AAs in India (14). In nearly all of the cases, the patients have autoantibodies against the C terminus of CFH, although usually in addition to such antibodies, some

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The atomic coordinates and structure factors (code 4MUC) have been deposited in the Protein Data Bank (http://wwpdb.org/).

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3 The abbreviations used are: aHUS, atypical hemolytic uremic syndrome; CFH, complement factor H; CFH-AA, anti-CFH autoantibody; r.m.s.d., root mean square deviation.
patients have autoantibodies against other parts of CFH as well (10, 15). Patients with the autoimmune form of aHUS nearly always lack certain CFH-related proteins, primarily CFHR1 (CFH-related protein-1) (10, 16).

CFH and CFHR proteins are encoded by adjacent genes and form a protein family (Fig. 1) (17). CFHR proteins are composed of four to nine complement control protein domains, also called short consensus repeats, and the two most C-terminal domains have relatively high sequence homologies to the C-terminal domains of CFH (Fig. 1) (17). There are only two residues that are different in the last two C-terminal domains of CFH and CFHR1 (domains 19–20 and 4–5, respectively). The functional importance of these minor differences is obvious because the hybrid CFH/CFHR1 genes producing fusion proteins CFH1–18/CFHR11–5 and CFHR11–3/CFH19–20 have been found in aHUS patients in the absence of other mutants or CFH-AAs (4, 18–20). Domains 19 and 20 of CFH are responsible for directing its complement regulatory activity to cell and extracellular matrix surfaces by binding simultaneously to both C3b and C3d (25). However, CFH-AAs have also been described, although rarely, in patients with two normal copies of CFHR1 and CFHR3 but mutations in CFH, CFI, CD46, or C3 genes (12, 13). CFH-AAs often cross-react with CFHR1 (13, 15, 26), but the exact location of the autoantibody site on CFHR1 has not been determined. On the basis of inhibition of autoantibody binding to CFHR1 by mAb C18 (26) and the sequence homology to the C terminus of CFH, it is likely, however, that the autoantibody-binding site is within the last two domains of CFHR1, i.e. far away from its N-terminal dimerization site (27).

To date, the reason for the association between CFH-AAs and CFHR1 deficiency has been unknown. In this study, we aimed to solve why a deficiency of one molecule (CFHR1) predisposes to autoimmunity against another, highly homologous molecule (CFH) in aHUS. We mapped the binding sites of CFH-AAs within CFH19–20 and compared the CFH-AA-binding sites with the previously reported ligand-binding sites on CFH19–20. Because the autoantibody epitopes formed a cluster next to the residues that are different in the two C-terminal domains of CFH and CFHR1, we decided to solve and analyze the structure of CFHR11–5 and to study the potential differences in antigenicity of those two molecules. We found structural differences in the autoantibody-binding site of CFH domain 20 and the corresponding homologous site of CFHR1 domain 5. Based on these data, a novel model is proposed, suggesting how immunization against CFH domain 20 could be linked to CFHR1 deficiency.

**EXPERIMENTAL PROCEDURES**

**Proteins**—Cloning, expression, and purification of WT CFH19–20 and mutant proteins with the 14 single-point mutations have been described previously (5, 7, 28). Proper folding of the constructs was verified for three mutants (Q1139A, R1203A, D1119G/Q1139A) by solving the structures by x-ray crystallography (6, 28, 29) and for five mutants (R1182A, W1183L, K1188A, E1198A, and R1206A) by circular dichroism (30). CFHR14–5 was generated by site-directed mutagenesis of CFH19–20 encoding DNA in the pPICZαB vector (Invitrogen). The primer used to introduce the S1191L and V1197A mutations in CFH was CAG AAG CTT TAT TTG AGA ACA TCA GGT GAA GAA and the sequence homology to the C terminus of CFH, it is likely, however, that the autoantibody-binding site is within the last two domains of CFHR1, i.e. far away from its N-terminal dimerization site (27).

**Patients and Blood Samples**—The studies were approved by the Research Ethics Committee of the Medical Faculty of Friedrich Schiller University Jena and were performed in accordance with the Declaration of Helsinki. Patients with aHUS were screened for CFH-AAs using ELISA as described (9, 11). All patients lacked the CFHR1 and CFHR3 genes and proteins except for patient 2, who had two copies of both genes, and patient 10, who carried a homozygous CFHRI deletion and was heterozygous for CFHR3. Three of the patients (patients 3, 7, and 8) have been described previously (23). The characteristics

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**Structural Insight on CFHR1 Deficiency in Autoimmune aHUS**

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In this study, we aimed to solve why a deficiency of one molecule (CFHR1) predisposes to autoimmunity against another, highly homologous molecule (CFH) in aHUS. We mapped the binding sites of CFH-AAs within CFH19–20 and compared the CFH-AA-binding sites with the previously reported ligand-binding sites on CFH19–20. Because the autoantibody epitopes formed a cluster next to the residues that are different in the two C-terminal domains of CFH and CFHR1, we decided to solve and analyze the structure of CFHR11–5 and to study the potential differences in antigenicity of those two molecules. We found structural differences in the autoantibody-binding site of CFH domain 20 and the corresponding homologous site of CFHR1 domain 5. Based on these data, a novel model is proposed, suggesting how immunization against CFH domain 20 could be linked to CFHR1 deficiency.
of the sera containing CFH-AAs from the 17 patients are summarized in Table 1.

**Microtiter Plate Assays**—WT or mutant CFH<sub>19–20</sub> proteins (5 μg/ml) were coated onto Nunc MaxiSorp plates (Thermo Scientific). Binding of patient sera (1:50–1:200 depending on the antibody titer) was analyzed as described (11). Binding of mAb C18 (5 μg/ml) was detected using peroxidase-conjugated swine anti-mouse IgG.

Binding of patient CFH-AAs (sera diluted 1:100) to CFH, its recombinant fragments, CFHR1, and CFHR1<sub>4–5</sub> was compared

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**FIGURE 2.** Mapping of the CFH-AA-binding region on CFH<sub>19–20</sub>. A, binding of IgG from 17 aHUS patients and mAb C18 to 14 CFH<sub>19–20</sub> constructs with various single-point mutations in relation to binding to WT CFH<sub>19–20</sub>. Error bars indicate S.E., and the level of WT binding is indicated by dotted lines. B, comparison of the autoantibody-binding epitopes by identification of the mutations that impaired binding of patient IgG by at least 30% (indicated by ). C, locations of the residues involved in binding of autoantibodies to CFH<sub>19–20</sub> as indicated in dark gray and annotated on a previously published structure of CFH<sub>19–20</sub> (Protein Data Bank code 2G7I) (7). The location of Trp-1183 is indicated by stripes. D and E, for comparison, the locations of residues involved in binding of CFH<sub>19–20</sub> to heparin (5, 40) (D) and in the common microbe-binding site on CFH<sub>19–20</sub> (30) (E) are indicated in dark gray.
as described above using 250 nm immobilized recombinant proteins, plasma-purified CFH (65 nm), or recombinant CFHR4B and albumin as negative controls. Relative binding was calculated from representative data sets performed in triplicates. Equal binding of anti-CFH polyclonal antisem and negligible binding of anti-IgG to the CFH19–20 mutants were confirmed by ELISA.

The assay comparing the binding avidity of CFH19–20 and CFHR14–5 with purified IgG from patient 11 was done after coating CFH19–20 (10 µg/ml) onto Nunc MaxiSorp plates. After blocking (0.05% Tween 20 in PBS) for 120 min and washing (0.02% Tween 20 in PBS), 40 µl of patient IgG (3.8 µg/ml) and CFH19–20, CFHR14–5, or CFH5–7 were added in different dilutions and incubated for 120 min at 37 °C, followed by measuring IgG binding as described (11). The experiment was performed three times in triplicates, and the background subtracted data were normalized using the values obtained without an inhibitor (100% binding).

Crystallization and Solving the CFHR14–5 Structure—CFHR14–5 was crystallized at 293 K from hanging drops in the presence of 2 M ammonium sulfate and 0.1 M sodium acetate at pH 4.6. The cube-shaped crystals appeared within 5 days and were cryoprotected with 25% glycerol (supplemented with the mother liquor). The diffraction data (to 2.9 Å) were collected at European Synchrotron (ESRF) beamline ID14-4 (34) at 100 K on an ADSC Q315r charge-coupled device detector at 0.979520 Å. The data were indexed and scaled using XDS (35). The structure of CFH19–20 mutant R1203A (Protein Data Bank code 3KZJ (28)) was used as a search model in Phaser (36), and two molecules of CFHR14–5 were identified in the asymmetric unit. After successive rounds of model building with Coot (37) and refinement using phenix.refine software (38), we could refine the structure to $R_{work}/R_{free} = 0.20/0.26$ (see Table 2). The last refinement cycles were done using TLS parameters (10 TLS groups). In the Ramachandran plot, 95% of the amino acid structures were within the most favored region.

The superpositions of different structures and structural illustrations were prepared using PyMOL software (Schrödinger, Portland, OR). The surface charge distribution of both the CFHR14–5 and CFH19–20 molecules was calculated using APBS (39), and the potentials on the solvent-accessible surfaces were displayed in PyMOL.

Statistical Analyses—Values are expressed as means ± S.E. using GraphPad Prism software (version 6). All curves and bar diagrams were made using the same software.

RESULTS

Mapping the Autoantibody-binding Residues on CFH19–20—The binding of IgG from sera of 17 aHUS patients with CFH-AAs to 14 different CFH19–20 mutants was tested (Fig. 2A). The binding data indicated a congruent tendency as follows. The binding of IgG from all patients to the L1189R mutant was diminished by at least 30% compared with WT CFH19–20. IgG from 14 patients showed at least 30% impaired binding to the E1198A mutant, and IgG from 10 or 11 patients showed impaired binding to CFH19–20 mutants T1184R, K1186A, and K1188A compared with WT CFH19–20 (Fig. 2B). IgG from one to three patients showed impaired binding to mutants D1119G, Q1139A, R1182A, W1183L, and R1210A, whereas IgG from all patients showed similar binding to WT CFH19–20 and mutants W1157L, R1203A, R1206A, and R1215Q (Fig. 2B).

We also found that binding of mAb C18, which has previously been shown to have an epitope overlapping with that of several CFH-AAs (11, 23), to the K1186A mutant was diminished (Fig. 2A). This antibody efficiently inhibited binding of CFH-AA to CFH19–20. This inhibition was also observed in those cases (patients 10, 11, 12, and 14) where the autoantibodies showed only moderately impaired binding to the five CFH mutants within the common autoantibody binding site (data not shown). This indicates that the main binding site of the autoantibodies from those patients is likely to be overlapping.

The Autoantibody-binding Site Overlaps with the Heparin- and Common Microbe-binding Sites—Our previously solved crystal structure of CFH19–20 (7) was used to visualize the location of the five residues of CFH19–20 (Thr-1184, Lys-1186, Lys-1188, Leu-1189, and Glu-1198) that were found to be involved in binding of CFH-AAs from at least 10 of 17 patients. These residues form a tightly packed cluster (diameter of ~11 Å) on one side of CFH domain 20 termed the CFH-AA site (Fig. 2C).

Compared with the previously described functional sites on CFH19–20, the CFH-AA site is clearly distinct from the two sites for C3b or C3d on domains 19 and 20 (6, 21) but adjacent to and partially overlapping with the site involved in binding of CFH to heparin and endothelial cells (Fig. 2D) (5, 40), as well as with the recently described common microbe-binding site on CFH domain 20 (Fig. 2E) (29, 30).

Binding of CFH-AAs to the CFHR1 C Terminus—On the basis of the previously solved structure of CFH19–20, the two residues that are different in CFH19–20 and CFHR14–5 (Ser-1191 and Val-1197 of CFH) are buried beneath the identified CFH-AA site. Therefore, we studied whether the difference of the two amino acids in the C-terminal domains of CFH and CFHR1 has an influence on the binding of autoantibodies from patient sera. The level of binding of antibodies from patient sera to both CFH19–20 and CFHR14–5 varied from patient to patient, and this seemed to correlate to the antibody titer in the sera (Table

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**TABLE 1**

| Patient | CFHR1 genotype | Sampling times | Ig subtype | Light chain type | CFH-AA titer |
|---------|----------------|---------------|-----------|-----------------|--------------|
| Patient 1 | /-/- | Convalescence | IgG3 | A | 456 |
| Patient 2 | +/- | Convalescence | IgG1, IgG3 | k | 868 |
| Patient 3 | -/- | Acute phase | IgG1, IgG3 | k | 1325 |
| Patient 4 | /-/- | Convalescence | IgG3 | | 1067 |
| Patient 5 | -/- | Acute phase | IgG1, IgG3 | k | 1043 |
| Patient 6 | -/- | Acute phase | IgG3 | | 1136 |
| Patient 7 | /-/- | Convalescence | IgG3 | k | 1132 |
| Patient 8 | -/- | Convalescence | IgG3 | k | 883 |
| Patient 9 | /-/- | Convalescence | IgG1 | | 1063 |
| Patient 10 | /-/- | Convalescence | IgG3 | k | 769 |
| Patient 11 | -/- | Acute phase | IgG3 | | 3548 |
| Patient 12 | /-/- | Convalescence | IgG1, IgG3 | k | 1239 |
| Patient 13 | /-/- | Acute phase | IgG1, IgG3 | k, k | 1260 |
| Patient 14 | /-/- | Convalescence | IgG3 | k | 737 |
| Patient 15 | /-/- | Convalescence | IgG3 | k | 444 |
| Patient 16 | /-/- | Convalescence | IgG1, IgG3 | k, k | 671 |
| Patient 17 | /-/- | Convalescence | IgG3 | | 852 |
IgG from 8 of the 10 CFH-AA patient samples of which we had enough available for analysis bound similarly to CFHR14–5, CFH, and CFH19–20, whereas binding of autoantibodies from two of the samples (patients 2 and 9) to CFHR14–5 was diminished (Fig. 3, A–C). Next, a purified IgG fraction from patient 11 was used to compare the affinities of CFH-AAs to CFH19–20 and CFHR14–5 by inhibition assay. Binding of IgG to CFH was somewhat stronger in this assay (Fig. 3, D and E), although a clear difference in binding to CFH19–20 and CFHR14–5 could not be detected in the ELISA assay (Fig. 3, B and C). Taken together, the results on the difference in binding of CFH-AAs to CFH19–20 and CFHR14–5 indicated that the conformation of the CFH-AA site needs to be slightly different in CFH domain 20 and CFHR domain 5.

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Structures of the C Termini of CFHR1 and CFH Are Nearly Identical—To detect possible differences in the CFH-AA site on CFH19–20 and the corresponding site on CFHR1, we used x-ray crystallography to solve the structure of CFHR14–5. The structure was obtained as a homodimer at 2.9 Å resolution from a different space group (P622) than the previously published CFH19–20 structures (I4122) (Table 2) (7, 28), but the structures aligned very well with each other (Fig. 4, A). Superimposing the CFHR14–5 structure with the previously published structures of CFH19–20 solved as homotetramers (7), in complex with C3d (6), or in complex with the borrelial OspE protein (29) gave a root mean square deviation (r.m.s.d.) of 0.5–1.1 Å for the 113 aligned Cα atoms, indicating that the structures are nearly identical. Also, the charge potentials on the solvent-accessible
Structural Insight on CFHR1 Deficiency in Autoimmune aHUS

Autoimmune aHUS is an unusual autoimmune disease because it is associated with a deficiency of a protein (CFHR1) homologous to the autoantigen (CFH). Therefore, it offers an exceptional opportunity to study phenomena leading to antibody-associated autoimmunity. This study shows that the amino acid residues contributing to the binding sites of CFH-AAs from 17 patients with autoimmune aHUS form a cluster on domain 20 of CFH adjacent to the common microbe-binding site. The differential binding of CFH-AAs from two aHUS patients to CFH19–20 and CFHR14–5 as also suggested previously (13, 26), and the small but clear differences in the x-ray crystal structures of the loop forming the autoantigenic epitope on CFH and CFHR1 indicate that the C-terminal domain of CFH and CFHR1 can have slightly different conformations. In addition, the conformation of the autoantigenic loop on CFH seems flexible because we noticed that the loop conformation is slightly different after binding of ligands to domain 20 of CFH in the previously published structures (22, 29). The reason for the association of CFHR1 deficiency with aHUS is unexplained, but our results enabled us to evaluate previous results and explanation models and to generate a new hypothesis of an induced autoantigenic neoepitope. This explains the association between CFHR1 deficiency and autoantibody formation against the common CFH-AA epitope on domain 20.

aHUS-associated mutations in the C-terminal domains of CFH have been shown to cause reduced binding of CFH to C3b or host cell-surface structures such as glycosaminoglycans/heparin (5–7, 41–43). aHUS-associated autoantibodies to CFH19–20 cause uncontrolled complement attack against host cells, and autoantibodies from some patients have been shown to impair CFH binding to C3b or to host cells (11). The location of the CFH-AA-binding site we identified on domain 20 indicates that the autoantibodies are likely to block binding of CFH at least to glycosaminoglycans/heparin due to the proximity of the CFH-AA site to the heparin-binding site. In addition, the location of the hemolysis-inducing aHUS mutation W1183L (26, 44) next to the CFH-AA site (Figs. 2C and 4D) indicates the importance of the site in protecting host cells from comple-

### TABLE 2

Data collection and refinement statistics for CFHR14–5

| Statistic | Value |
|-----------|-------|
| Resolution range (Å) | 48.53–2.897 (3.001–2.897) |
| Space group | P21 |
| Unit cell dimensions | a, b, c (Å) 143.406, 143.406, 77.784 |
| α, β, γ (°) | 90°, 90°, 120° |
| Unique reflections | 10,935 (1048) |
| Completeness (%) | 99.78 (98.59) |
| Mean I/σI | 15.30 (2.46) |
| Wilson B-factor | 72.41 |
| Rwork | 0.2070 (0.3078) |
| Rfree | 0.2683 (0.3642) |
| No. of atoms | 1939 |
| Macromolecules | 1904 |
| Ligands | 35 |
| Water | 3 |
| Protein residues | 248 (both molecules in the asymmetric unit) |
| r.m.s.d. Bond lengths (Å) | 0.009 |
| Bond angles | 1.16° |
| Ramachandran favored (%) | 95 |
| Ramachandran outliers (%) | 0 |
| Average B-factor | 96.4 |
| Macromolecules | 95.5 |
| Solvent | 80.4 |

Surface displayed at the ±2 kT/e level on CFHR14–5 and CFH19–20 were similar all around the molecules (Fig. 4B).

Differences Noticed in the Structure of the CFH-AA Site of CFH19–20 and CFHR14–5—The structures of CFH19–20 and CFHR14–5 are clearly different (Fig. 4). Also, the hydrogen bonds stabilizing the loop in CFH and CFHR1 are distinctly different, as Arg-281–Leu-288 of CFHR14–5 are distinct from that on CFH19–20 (r.m.s.d. of the backbone of the loop region = 3 Å). Both the backbone and loops forming the CFH-AA site (Arg-281–Leu-288 of CFHR1 and Arg-1182–Leu-1189 of CFH, respectively) have a different orientation (Fig. 4D). The main difference in the backbone is the formation of a short α-helix in the loop of CFHR14–5, whereas no prominent helix is seen in this region in any of the solved structures of the CFH19–20. The orientation of some of the side chains is also different, as Arg-281–Leu-288 of CFHR14–5 are distinctly different from that on CFHR1 (Fig. 4). The differential binding of CFH-AAs from two aHUS patients to CFH19–20 and CFHR14–5 as also suggested previously (13, 26), and the small but clear differences in the x-ray crystal structures of the loop forming the autoantigenic epitope on CFH and CFHR1 indicate that the C-terminal domain of CFH and CFHR1 can have slightly different conformations. In addition, the conformation of the autoantigenic loop on CFH seems flexible because we noticed that the loop conformation is slightly different after binding of ligands to domain 20 of CFH in the previously published structures (22, 29). The reason for the association of CFHR1 deficiency with aHUS is unexplained, but our results enabled us to evaluate previous results and explanation models and to generate a new hypothesis of an induced autoantigenic neoepitope. This explains the association between CFHR1 deficiency and autoantibody formation against the common CFH-AA epitope on domain 20.

Because the data indicated that the Arg-281–Leu-288 loop of CFHR1 is flexible (or has two conformations), we next analyzed whether flexibility is observed in the CFH-AA site of the previously published structures of CFH19–20. The conformation of the Arg-1182–Leu-1189 loop is somewhat different in free CFH19–20 compared with the mutually similar conformation of CFH19–20 in complex with either the microbial protein OspE (r.m.s.d. of the backbone of the loop residues = 2.7 Å) or the natural ligands C3d (r.m.s.d. of the backbone of the loop residues = 3.0 Å) or sialic acid glycan (r.m.s.d. of the backbone of the loop residues = 2.9 Å) (Fig. 4H). This conformation is also different from that of CFHR1 (r.m.s.d. of the backbone of the loop residues = 3.0 Å). This indicates the structural flexibility of the CFH-AA site upon binding of a ligand to the same domain (Fig. 4H).

To exclude potential misinterpretation of the x-ray diffraction data of the CFH-AA site, we next compared the electron density maps (2mFo–DFc) of Arg-1182–Leu-1189 of CFH19–20 (7) and Arg-281–Leu-288 of CFHR14–5 (Fig. 5). Clearly, the model of CFHR14–5 (but not CFH19–20) fits very well with the electron density map of CFHR14–5 in this region (Fig. 5, A and B), whereas the corresponding region in the model of CFH19–20 (but not CFH14–5) fits well with the electron density map of CFH19–20 (Fig. 5, C and D).
FIGURE 4. Crystal structure of CFHR1\textsubscript{4-5} and comparison with the previously solved structure of CFH\textsubscript{19-20}. A, structural superposition of the two molecules of CFHR1\textsubscript{4-5} (orange and yellow) found in the asymmetric unit along with CFH\textsubscript{19-20} (gray) shown in a cartoon representation. B, comparison of the surface charge potentials of CFHR1\textsubscript{4-5} and CFH\textsubscript{19-20}. Potentials on the solvent-accessible surfaces were calculated and displayed at the \(\pm 2 \text{kT/e} \) level on both structures after modeling all of the missing side chains of the previously published structure of CFH\textsubscript{19-20} (Protein Data Bank code 2G7I) (7). C, close-up view of the two residues that are different in the amino acid sequences of these two protein constructs with the \(2mF_0-F_D\) electron density map of CFHR1\textsubscript{4-5} shown. D, close-up view of the region in which the tertiary structures of CFHR1\textsubscript{4-5} and CFH\textsubscript{19-20} are dissimilar (Arg-1182–Leu-1189 of CFH\textsubscript{19-20} and the corresponding Arg-281–Leu-288 of CFHR1\textsubscript{4-5}) with backbone and side chain atoms shown as a stick model. This region corresponds to the CFH-AA-binding site shown in Fig. 2C. The hydrogen bonds found in the autoantigenic loop of CFH (E) and its homologous region in CFHR1 (F) that stabilize the structure. G, comparison of the real space correlation constants and \(B\)-factors of the residues of CFHR1\textsubscript{4-5}. The \(B\)-factors are indicated on the right \(y\) axis, and the real space correlation coefficients are indicated on the left \(y\) axis. The CFH-AA site (Arg-281–Leu-288) is indicated. H, cartoon representation of the structural superposition of CFHR1\textsubscript{4-5} (orange; Protein Data Bank code 4MUC) with CFH\textsubscript{19-20} (gray; code 2G7I (7)), CFH\textsubscript{19-20} in complex with a sialic acid glycan and C3d (slate; code 4ONT (22)), and CFH\textsubscript{19-20} in complex with OspE (turquoise; code 4J38 (29)).
CFH19–20 (6, 7, 21, 22, 40, 45) shows minor differences in the loop of CFH in the published crystal and NMR structures of patients (1). C3b can lead to compromised protection of the plasma-exposed binding of CFH to either the cell-surface structures or found (11) to bind to the same region as the CFH-AAs. Diminished binding of CFH to either the cell-surface structures or C3b can lead to compromised protection of the plasma-exposed host cells, which has been reported widely in aHUS.

It seems possible that the corresponding site on CFHR1 is also altered conformation only upon binding of a ligand. However, we do not suggest that binding of self-molecules (such as heparin, C3b, or sialic acids) leads to autoimmunity, but the flexibility of the autoantigenic loop upon binding of these ligands to CFH domain 20 is obvious.

Close comparison of the conformation of the autoantigenic loop of CFH in the published crystal and NMR structures of CFH19–20 (6, 7, 21, 22, 29, 40, 45) shows minor differences in the Arg-1182–Leu-1189 region of CFH19–20 in complex with OspE (29), C3d (6), or a sialic acid glycan (22). The r.m.s.d. of the backbone atoms of the loop region is 2.7–3.0 Å compared with CFH19–20 alone (Fig. 4H). Binding of CFH19–20 to heparin tetrasaccharide has also been shown to cause chemical shift perturbations in NMR at the CFH-AA site (22, 40). Thus, it is likely that the conformation of the region is somewhat flexible. It seems possible that the corresponding site on CFHR1 is also structurally flexible because of the two monomers in the unit cell of the CFHR14–5 crystal lattice, conformation of the CFH-AA site of one monomer was similar to CFH, and the other one was different (Fig. 4A). The key difference between CFH and CFHR1 could therefore be that CFHR1 takes the alternative conformation spontaneously, whereas on the basis of various crystal and NMR structures, CFH19–20 takes a slightly altered conformation only upon binding of a ligand. However, we do not suggest that binding of self-molecules (such as heparin, C3b, or sialic acids) leads to autoimmunity, but the flexibility of the autoantigenic loop upon binding of these ligands to CFH domain 20 is obvious.

The observed difference between the CFHR14–5 and the CFH19–20 structures is unlikely to be a crystallographic artifact due to the four following reasons. First, only one of the monomers in the unit cell containing two CFHR14–5 molecules shows a structure that is considerably different from CFH domain 20 (see Protein Data Bank code 4MUC), indicating structural flexibility in that loop of CFHR1 domain 5. Second, there are no direct contacts between the residues of the CFH-AA site (Arg-281–Leu-288) and the molecule in the neighboring crystal cell. Third, CFH-AAs from two of the studied 10 patients bound differently to CFH19–20 and CFHR4–5 (Fig. 3C), indicating that there is a difference within the CFH-AA-binding site of these molecules. Fourth, CFHR14–5 and CFH19–20 have been reported to have functional differences (45), which is obvious because the fusion proteins CFH19–20/CFH14–5 and CFHR14–5/CFH19–20 are associated with aHUS and have different functions compared with normal full-length CFH and CFHR1 (4, 18–20). Domain 19 of CFH and domain 4 of CFHR1 are identical, it has been deduced that the difference leading to the clinical disease is within the terminal domain of the fusion proteins, leading to the inability to control complement on self-surfaces (19). Our results suggest that the reason for the functional difference between the most C-terminal domains of CFH and CFHR1 is their varied ability to bind to heparin or glycosaminoglycans on self-cells because the loop that has a different conformation in CFH and CFHR1 contains several of the heparin-binding residues (Fig. 2, C and D; and Fig. 4D) (5, 40).

Nearly all patients with CFH-AAs lack CFHR1 (12, 13); thus, it is likely that the absence of CFHR1 imparts the risk of CFH-AA generation. The risk for anti-CFH autoimmunity in the absence of CFHR1 is very high, as the odds ratio is 442 (16). In this study, we provided data for a structure-based molecular explanation of the phenomenon. The explanation is based on four observations from our study. First, the binding sites of CFH-AAs from the 17 patients analyzed clearly formed a cluster, the CFH-AA site. Second, the CFH-AA site in CFH...
domain 20 is adjacent to the two buried residues (Ser-1191 and Val-1197) that are different between CFH\textsubscript{19–20} and CFHR1\textsubscript{4–5}. Third, although the crystal structures of CFHR1\textsubscript{4–5} and CFH\textsubscript{19–20} are similar due to the 98.5% sequence identity between them, there is a small structural difference exactly at the CFH-AA site on the Arg-1182–Leu-1189 loop of CFH (Fig. 4D). Fourth, a small conformational change has been detected within the autoantigenic loop upon binding of CFH\textsubscript{19–20} to a microbial protein (29), a sialic acid glycan (22), heparin, or C3d.

The two usual models to explain autoantibodies in general, an analogous epitope (molecular mimicry) (46) and a co-epitope (formed by adjacent molecules) (47), are unable to explain the association between CFHR1 deficiency and CFH-AA binding to the CFH-AA site on domain 20. Thus, on the basis of the new data, we propose a novel explanation for the association of CFHR1 deficiency with the autoimmune disease. In this model, called the “induced neoepitope model,” binding of a microbial protein to CFH domain 20 (adjacent to the autoantigenic Arg-1182–Leu-1189 loop, thus not masking it from B-cells) induces a conformational change in the Arg-1182–Leu-1189 loop of domain 20, thereby making its conformation similar to that of CFHR1 domain 5.

Infections can precede autoimmune aHUS, and this is concordant with our model because several microbial molecules are known to bind close to the autoantigenic epitope of CFH (29), (30), and we observed from previously published data that CFH\textsubscript{19–20} in complex with OspE (29) has a slightly altered conformation in this region.

It is possible that binding of certain microbial ligands to the autoantigenic loop or close to it leads to masking of the autoantigenic epitope. We reported previously that several microbes bind to the autoantigenic loop (29, 30). Therefore, all of these may induce the autoantigenic neoepitope, but some of them could simultaneously mask the loop. This is the case at least with borreliosis (29). It is noteworthy that borreliosis has never been reported preceding autoimmune aHUS. Therefore, we do not suggest that all microbial molecules that bind to CFH domain 20 could lead to autoimmunity in CFHR1-deficient individuals. However, it is possible that some of the microbial molecules that bind to the domain do not mask the autoantigenic loop, thereby leading to the risk of autoimmunity in the absence of CFHR1 (Fig. 6).

Our model provides an explanation as to why immunization against the CFH-AA site could occur only in CFHR1-deficient individuals.
individuals because, in normal individuals, the presence of CFHR1 with an epitope similar to the hypothetical induced autoantigenic conformation of CFH would have guaranteed tolerance to that conformation of CFH. The model proposed here also explains the other four key biological phenomena described in autoimmune aHUS. First, the association with infections (14, 48). Second, the clustering of the autoimmune epitopes on CFH domain 20 (Fig. 2). Third, the high prevalence of IgG or IgA class autoantibodies (9, 26) because the foreign peptide needed for class switch of B-cells by T-cell help could be provided by the microbial protein bound to CFH 19–20. And finally, the model explains the polyclonality of the autoimmune response (15) because different epitopes on the autoantigenic loop could be recognized by various B-cell receptors.

In this study, we have shown that CFH-AA s bind to a common site on the Arg-1182–Leu-1189 loop of CFH next to the two buried residues that are different in CFH1 9–20 and CFHR1 4–5. The crystal structure of CFHR1 4–5 presented here shows that the conformation of the autoantigenic loop is different in CFH and CFHR1. Taken together, these data provide the basis for the suggested novel model (Fig. 6) to explain how CFHR1 deficiency is linked to CFH-AA formation.

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REFERENCES
1. Noris, M., and Remuzzi, G. (2009) Atypical hemolytic-uremic syndrome. N. Engl. J. Med. 361, 1676–1687
2. Sánchez-Corral, P., and Melgosa, M. (2010) Advances in understanding the aetiology of atypical haemolytic uraemic Syndrome. Br. J. Haematol. 150, 529–542
3. Zipfel, P. F., Edey, M., Heinen, S., Józsi, M., Richter, H., Misselwitz, J., Hoppe, B., Routledge, D., Strain, L., Hughes, A. E., Goodship, J. A., Licht, C., Goodship, T. H., and Skerka, C. (2007) Deletion of complement factor H-related genes CFHR1 and CFHR3 is associated with atypical haemolytic uraemic syndrome. PLoS Genet. 3, e41
4. Venables, J. P., Strain, L., Routledge, D., Bourn, D., Powell, H. M., Warricker, P., Diaz-Torres, M. L., Sampson, A., Mead, P., Webb, M., Pirson, Y., Jackson, M. S., Hughes, A., Wood, K. M., Goodship, J. A., and Goodship, T. H. (2006) Atypical haemolytic uraemic syndrome associated with a hybrid complement gene. PLoS Med. 3, e431
5. Lehtiniemi, M. J., Rops, A. L., Isenman, D. E., van der Vlag, J., and Jokiranta, T. S. (2009) Mutations of factor H impair regulation of surface-bound C3b by three mechanisms in atypical haemolytic uraemic syndrome. J. Biol. Chem. 284, 15650–15658
6. Kajander, T., Lehtiniemi, M. J., Hyvärinen, S., Bhattacharjee, A., Leung, E., Isenman, D. E., Meri, S., Goldman, A., and Jokiranta, T. S. (2011) Dual interaction of factor H with C3d and glycosaminoglycans in host-nonhost discrimination by complement. Proc. Natl. Acad. Sci. U.S.A. 108, 2897–2902
7. Jokiranta, T. S., Jaakola, V. P., Lehtiniemi, M. J., Pärepolo, M., Meri, S., and Goldman, A. (2006) Structure of complement factor H carboxyl-terminus reveals molecular basis of atypical haemolytic uraemic syndrome. EMBO J. 25, 1794–1794
8. Ferreira, V. P., Herbert, A. P., Cortés, C., McKee, K. A., Bluma, B. S., Esswein, S. T., Uhrin, D., Barlow, P. N., Pangburn, M. K., and Kavanagh, D. (2009) The binding of factor H to a complex of physiological polyanions and C3b on cells is impaired in atypical hemolytic uremic syndrome. J. Immunol. 182, 7009–7018
9. Dragon-Durey, M. A., Loirat, C., Cloarec, S., Macher, M. A., Blouin, J., Nivet, H., Weiss, L., Fridman, W. H., and Frémeaux-Bacchi, V. (2005) Anti-Factor H autoantibodies associated with atypical hemolytic uremic syndrome. J. Am. Soc. Nephrol. 16, 555–563
10. Józsi, M., Licht, C., Strobel, S., Zipfel, S. L., Richter, H., Heinen, S., Zipfel, P. F., and Skerka, C. (2008) Factor H autoantibodies in atypical hemolytic uremic syndrome correlate with CFHR1/CFHR3 deficiency. Blood 111, 1512–1514
11. Józsi, M., Strobel, S., Dahse, H. M., Liu, W. S., Hoyer, P. F., Oppermann, M., Skerka, C., and Zipfel, P. F. (2007) Anti-factor H autoantibodies block C-terminal recognition function of factor H in hemolytic uraemic syndrome. Blood 110, 1516–1520
12. Abharragueui-Garrido, C., Martínez-Barricarte, R., López-Trascasa, M., de Córdoba, S. R., and Sánchez-Corral, P. (2009) Characterization of complement factor H-related (CFHR) proteins in plasma reveals novel genetic variations of CFHR1 associated with atypical hemolytic uremic syndrome. Blood 114, 4261–4271
13. Moore, J., Strain, L., Pappworth, I., Kavanagh, D., Barlow, P. N., Herbert, A. P., Schmidt, C. Q., Staniforth, S. J., Holmes, L. V., Ward, R., Morgan, L., Goodship, T. H., and Marchbank, K. I. (2010) Association of factor H autoantibodies with deletions of CFHR1, CFHR3, CFHR4, and with mutations in CFH, CF, CD46, and C3 in patients with atypical hemolytic uremic syndrome. Blood 115, 379–387
14. Sinha, A., Gulati, A., Saini, S., Blanc, C., Gupta, A., Gurjar, B. S., Saini, H., Kotresh, S. T., Ali, U., Bhatia, D., Ohri, A., Kumar, M., Agarwal, I., Gulati, S., Anand, K., Vijayakumar, M., Sinha, R., Sethi, S., Salmona, M., George, A., Bal, V., Singh, G., Dinda, A. K., Hari, P., Rath, S., Dragon-Durey, M. A., and Bagga, A. (2014) Prompt plasma exchanges and immunosuppressive treatment improves the outcomes of anti-factor H autoantibody-associated hemolytic uremic syndrome in children. Kidney Int. 85, 1151–1160
15. Blanc, C., Roumenina, L. T., Ashraf, Y., Hyvärinen, S., Sethi, S. K., Ranchin, B., Niaudet, P., Loirat, C., Gulati, A., Bagga, A., Fridman, W. H., Sautès-Fridman, C., Jokiranta, T. S., Frémeaux-Bacchi, V., and Dragon-Durey, M. A. (2012) Overall neutralization of complement factor H by autoantibodies in the acute phase of the autoimmune form of atypical hemolytic uremic syndrome. J. Immunol. 189, 3528–3537
16. Dragon-Durey, M. A., Blanc, C., Marliot, F., Loirat, C., Blouin, J., Sautès-Fridman, C., Fridman, W. H., and Frémeaux-Bacchi, V. (2009) The high frequency of complement factor H related CFHR1 gene deletion is restricted to specific subgroups of patients with atypical haemolytic uraemic syndrome. J. Med. Genet. 46, 447–450
17. Zipfel, P. F., Skerka, C., Hellwage, J., Jokiranta, S. T., Meri, S., Brade, V., Kraicy, P., Noris, M., and Remuzzi, G. (2002) Factor H family proteins on complement, microbes and human diseases. Biochem. Soc. Trans. 30, 971–978
18. Heinen, S., Sánchez-Corral, P., Jackson, M. S., Strain, L., Goodship, J. A., Kemp, E. J., Skerka, C., Jokiranta, T. S., Meyers, K., Wagner, E., Robitaille, P., Esparza-Gordillo, J., Rodríguez de Córdoba, S., Zipfel, P. F., and Goodship, T. H. (2006) De novo gene conversion in the RCA gene cluster (1q32) causes mutations in complement factor H associated with atypical haemolytic uraemic syndrome. Hum. Mutat. 27, 292–293
19. Eyler, S. J., Meyer, N. C., Zhang, Y., Xiao, X., Nester, C. M., and Smith, R. J. (2013) A novel hybrid CFHR1/CFH gene causes atypical hemolytic uremic syndrome. Pediatr. Nephrol. 28, 2221–2225
20. Roumenina, L., Strain, L., Deury, D., Merle, N., Halbwachs-Mecarelli, L., Goodship, T., and Frémeaux-Bacchi, V. (2013) A prevalent CFHR1/FH reverse hybrid gene in aHUS patients induces deregulation of the alternative pathway. Mol. Immunol. 56, 248
21. Morgan, H. P., Schmidt, C. Q., Guarente, M., Blau, B. S., Gillespie, D., Herbert, A. P., Kavanagh, D., Mertens, H. D., Svergun, D. I., Johansson, C. M., Uhrin, D., Barlow, P. N., and Hannan, J. P. (2011) Structural basis for engagement by complement factor H of C3b on a self surface. Nat. Struct.
Structural Insight on CFHR1 Deficiency in Autoimmune aHUS

Mol. Biol. 18, 463–470

22. Blaum, B. S., Hannan, J. P., Herbert, A. P., Kavanagh, D., Uhrin, D., and Stehle, T. (2015) Structural basis for sialic acid-mediated self-recognition by complement factor H. Nat. Chem. Biol. 11, 77–82

23. Strobel, S., Hoyer, P. F., Mache, C. J., Sulyok, E., Liu, W. S., Richter, H., Oppermann, M., Zipfel, P. F., and Jösi, M. (2010) Functional analyses indicate a pathogenic role of factor H autoantibodies in atypical haemolytic uraemic syndrome. Nephrol. Dial. Transplant. 25, 136–144

24. Hofer, J., Janecek, A. R., Zimmerhackl, L. B., Riedl, M., Rosales, A., Giner, T., Cortina, G., Haindl, C. J., Petzelberger, B., Pawlik, M., Jeller, V., Vester, U., Gadner, B., van Husen, M., Moritz, M. L., Würzner, R., Jungraithmayr, T., and German-Austrian HUS Study Group (2013) Complement Factor H-related protein 1 deficiency and Factor H antibodies in pediatric patients with atypical hemolytic uremic syndrome. Clin. J. Am. Soc. Nephrol. 8, 407–415

25. Holmes, L. V., Strain, L., Staniforth, S. J., Moore, I., Marchbank, K., Kavanagh, D., Goodship, J. A., Cordell, H. J., and Goodship, T. H. (2013) Determining the population frequency of the CFHR3/CFHR1 deletion at 1q32. PLoS ONE 8, e60352

26. Oppermann, M., Manuelian, T., Jósi, M., and Jósi, M. (2012) Factor H-related protein 1 activates complement C3 convertase via its interaction with C3b protein. J. Biol. Chem. 287, 19525–19536

27. Castilblanco-Musa, M. F., Fraga, T. R., Silva, L. B., Monaris, D., Abreu, P. A., Strobel, S., Jósi, M., Gómez, L., and Barbosa, A. S. (2012) Leptospiral immunoglobulin-like proteins interact with human complement regulators factor H, FHL-1, FHR-1, and C4BP. J. Infect. Dis. 205, 995–1004

28. Oppermann, M., Manuelian, T., Jósi, M., Brandt, E., Jósi, M., Heinen, S., Meri, S., Skerka, C., Götz, O., and Zipfel, P. F. (2006) The C-terminus of complement regulator Factor H mediates target recognition: evidence for a compact conformation of the native protein. Clin. Exp. Immunol. 144, 342–352

29. McCarthy, A. A., Brockhauser, S., Nurizzo, D., Theveneau, P., Mairs, T., Spruce, D., Guijarro, M., Lesourd, M., Ravelli, R. B., and McSweeney, S. (2009) A decade of user operation on the macromolecular crystallography MAD beamline ID14-4 at the ESRF. J. Synchrotron Radiat. 16, 803–812

30. Kabsch, W. (2010) XDS. Acta Crystallogr. D Biol. Crystallogr. 66, 125–132

31. Strobel, S., Abarrategui-Garrido, C., Fariza-Requejo, E., Seeberger, H., Sánchez-Corral, P., and Jósi, M. (2011) Factor H-related protein 1 neutralizes anti-factor H autoantibodies in atypical hemolytic uremic syndrome. Kidney Int. 80, 397–404

32. Goicoechea de Jorge, E., Caesar, J. J., Malik, T. H., Patel, M., Colledge, M., Johnson, S., Hakobyan, S., Morgan, B. P., Harris, C. L., Pickering, M. C., and Lea, S. M. (2013) Dimerization of complement factor H-related proteins modulates complement activation in vivo. Proc. Natl. Acad. Sci. U.S.A. 110, 4685–4690

33. Bhattacharjee, A., Lehtinen, M. J., Kajander, T., Goldman, A., and Jokiranta, T. S. (2010) Both domain 19 and domain 20 of factor H are involved in binding to complement C3b and C3d. Mol. Immunol. 47, 1686–1691

34. Bhattacharjee, A., Oeming, J. S., Kolodziejczyk, R., Meri, T., Kajander, T., Lehtinen, M. J., Iwai, H., Jokiranta, T. S., and Goldman, A. (2013) Structural basis for complement evasion by lyme disease pathogen Borrelia burgdorferi. J. Biol. Chem. 288, 18685–18695

35. Meri, T., Amdahl, H., Lehtinen, M. J., Hyvärinen, S., McDowell, J. V., Bhattacharjee, A., Meri, S., Marconi, R., Goldman, A., and Jokiranta, T. S. (2013) Microbes bind complement inhibitor factor H via a common site. PLoS Pathog. 9, e1003308

36. Hebecker, M., and Jósi, M. (2012) Factor H-related protein 4 activates complement by serving as a platform for the assembly of alternative pathway C3 convertase via its interaction with C3b protein. J. Biol. Chem. 287, 19525–19536

37. Castilblanco-Musa, M. F., Fraga, T. R., Silva, L. B., Monaris, D., Abreu, P. A., Strobel, S., Jósi, M., Gómez, L., and Barbosa, A. S. (2012) Leptospiral immunoglobulin-like proteins interact with human complement regulators factor H, FHL-1, FHR-1, and C4BP. J. Infect. Dis. 205, 995–1004

38. Oppermann, M., Manuelian, T., Jósi, M., Brandt, E., Jósi, M., Heinen, S., Meri, S., Skerka, C., Götz, O., and Zipfel, P. F. (2006) The C-terminus of complement regulator Factor H mediates target recognition: evidence for a compact conformation of the native protein. Clin. Exp. Immunol. 144, 342–352

39. McCarthy, A. A., Brockhauser, S., Nurizzo, D., Theveneau, P., Mairs, T., Spruce, D., Guijarro, M., Lesourd, M., Ravelli, R. B., and McSweeney, S. (2009) A decade of user operation on the macromolecular crystallography MAD beamline ID14-4 at the ESRF. J. Synchrotron Radiat. 16, 803–812

40. Kabsch, W. (2010) XDS. Acta Crystallogr. D Biol. Crystallogr. 66, 125–132

41. McCoy, A. J. (2007) Solving structures of protein complexes by molecular replacement with Phaser. Acta Crystallogr. D Biol. Crystallogr. 63, 32–41

42. Pangburn, M. K. (2002) Cutting edge: localization of the host recognition functions of complement factor H at the carboxyl-terminal: implications for hemolytic uremic syndrome. J. Immunol. 169, 4702–4706

43. Jósi, M., Heinen, S., Hartmann, A., Ostrowicz, C. W., Häblich, S., Richter, H., Kunert, A., Licht, C., Saunders, R. E., Perkins, S. J., Zipfel, P. F., and Skerka, C. (2006) Factor H and atypical hemolytic uremic syndrome: mutations in the C-terminus cause structural changes and defective recognition functions. J. Am. Soc. Nephrol. 17, 170–177

44. Sánchez-Corral, P., González-Rubio, C., Rodríguez de Córdoba, S., and Lópe–Trascasa, M. (2004) Functional analysis in serum from atypical hemolytic uremic syndrome patients reveals impaired protection of host cells associated with mutations in factor H. Mol. Immunol. 41, 81–84

45. Herbert, A. P., Kavanagh, D., Johansson, C., Morgan, B. S., Blaum, H. S., Hanon, P. P., Barlow, P. N., and Uhrin, D. (2012) Structural and functional characterization of the product of disease-related factor H gene conversion. Biochemistry 51, 1874–1884

46. Fritzler, M. J., and Salazar, M. (1991) Diversity and origin of rheumatologic autoantibodies. Clin. Microbiol. Rev. 4, 256–269

47. Freddo, L., Hays, A. P., Nickerson, K. G., Spatz, L., McGinnis, S., Lieberson, R., Vedeler, C. A., Shy, M. E., Autilio-Gambetti, L., and Grauss, F. C. (1986) Monoclonal anti-DNA IgM kappa in neuropathy binds to myelin and to a conformational epitope formed by phosphatidic acid and gangliosides. J. Immunol. 137, 3821–3825

48. Dragón-Durey, M. A., Sethi, S. K., Bagga, A., Blanc, C., Blouin, J., Ranchin, B., André, J. L., Takagi, N., Cheong, H. I., Hari, P., Le Quintrec, M., Niaudet, P., Loirat, C., Fridman, W. H., and Frémeaux-Bacchi, V. (2010) Clinical features of anti-factor H autoantibody-associated hemolytic uremic syndrome. J. Am. Soc. Nephrol. 21, 2180–2187