Properdin binding to complement activating surfaces depends on initial C3b deposition

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Two functions have been assigned to properdin; stabilization of the alternative convertase, C3bBb, is well accepted, whereas the role of properdin as pattern recognition molecule is controversial. The presence of nonphysiological aggregates in purified properdin preparations and experimental models that do not allow discrimination between the initial binding of properdin and binding secondary to C3b deposition is a critical factor contributing to this controversy. In previous work, by inhibiting C3, we showed that properdin binding to zymosan and Escherichia coli is not a primary event, but rather is solely dependent on initial C3 deposition. In the present study, we found that properdin in human serum bound dose-dependently to solid-phase myeloperoxidase. This binding was dependent on C3 activation, as demonstrated by the lack of binding in human serum with the C3-inhibitor compstatin C440, in C3-depleted human serum, or when purified properdin is applied in buffer. Similarly, binding of properdin to the surface of human umbilical vein endothelial cells or Neisseria meningitidis after incubation with human serum was completely C3-dependent, as detected by flow cytometry. Properdin, which lacks the structural homology shared by other complement pattern recognition molecules and has its major function in stabilizing the C3bBb convertase, was found to bind both exogenous and endogenous molecular patterns in a completely C3-dependent manner. We therefore challenge the view of properdin as a pattern recognition molecule, and argue that the experimental conditions used to test this hypothesis should be carefully considered, with emphasis on controlling initial C3 activation under physiological conditions.

Properdin, also referred to as factor P, was first described in 1954 by Pillemer and colleagues as a component that, in an antibody-independent manner, is able to promote complement activation on zymosan particles and on other carbohydrates (1). These claims were controversial, and properdin-dependent complement activation was dismissed by the scientific community (2–4); however, the “properdin system” was reborn as the alternative pathway (AP) more than 20 y later (3), with properdin described as a stabilizer and positive regulator of the AP C3 convertase (5, 6). Properdin and its possible different roles in complement activation have been a basis for further studies in this area (7–11).

In the current conception, although yet to be proven in vivo, the AP of the complement system is slowly autoactivated via spontaneous or induced formation of fluid-phase AP C3 convertase (12, 13). The C3 moiety within this convertase is C3(H2O) formed on exposure and subsequent hydrolysis of the internal thioester, which is normally protected inside native C3 (14–16). C3(H2O) is “C3b-like”; it still contains C3a, but is conformationally similar to C3b. C3(H2O) can bind factor B, which is cleaved by factor D into Ba and Bb. Bb remains bound to C3(H2O), forming the enzymatic complex that cleaves C3 into C3b and C3a. Surface-bound C3b can form additional AP C3 convertase molecules with Bb, which rapidly cleave more C3 resulting in self-amplification and generation of the C5 convertase C3bBbC3b. The degree of amplification on a surface is determined by the rate of the C3b feedback (i.e., C3 cleavage) and breakdown (i.e., C3b degradation) cycles (17).

We previously reported that amplification via the AP on an unprotected surface contributes to more than 80% of terminal pathway activation after specific initial classical pathway or lectin pathway activation (18, 19). The C3bBb complex is relatively unstable, with a half-life of 90 s under physiological conditions (6, 20); however, properdin can associate with C3bBb and create the more stable C3bBbP complex that is essential for effective AP amplification (5, 21). Recently published electron microscopy images of the C3bBbP complex have shown how properdin is associated with the convertase near the C345C domain of C3b and the von Willebrand factor type A domain of factor B (22).

Discussion of the role of properdin as a pattern recognition molecule and initiator of the AP was renewed with experiments showing that purified unfraccionated properdin covalently attached to a biosensor surface could serve as a platform for in situ assembly of the AP C3 convertase (23). This was done in a relatively artificial system by using purified components in buffer milieu. That study was followed by several reports of biological substrates suggested to serve as patterns for the

Significance

The role of properdin in stabilization of the alternative pathway C3 convertase is indisputable, whereas its role as pattern recognition molecule remains controversial. Properdin lacks the structural homology shared by other pattern recognition molecules of the complement system, and has its major function in stabilizing the C3bBb convertase. We found that properdin binding was completely abolished by C3 inhibition after the exposure of human serum to myeloperoxidase, human umbilical vein endothelial cells, and Neisseria meningitidis, showing that properdin is not a pattern recognition molecule for these targets. We therefore challenge the view of properdin as a pattern recognition molecule, and argue that properdin typically binds a complement-activating surface substrate to C3b to stabilize the alternative pathway C3 convertase.


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direct recognition by properdin for AP complement activation. Reported patterns include exogenous microorganisms (7, 24), endogenous cells (25–27), and various biological substrates (9, 28, 29); however, many of these experiments were performed in systems permitting either C3 activation with initial C3b deposition or in buffer systems with purified properdin. In the presence of intact C3, it is virtually impossible to demonstrate whether properdin acts in a recognition manner or subsequently binds to C3b. On the other hand, purified properdin is sensitive to aggregation into sticky multimers, and these aggregates, referred to as activated properdin, do not behave as the native form; thus, results with purified properdin must be judged with care (30, 31), especially in unfractionated preparations (32). Early preparations of properdin, such as those used in the initial studies by Pillemer et al. (1), were later shown to contain impurities and high molecular aggregates of properdin, which were lost when adsorption to zymosan was omitted from the purification protocols (30, 33, 34). Furthermore, it is crucially important to distinguish between passive binding of properdin to a surface and subsequent activation of AP as a result of this binding. For this purpose, the methodological conditions should be under strict control.

In previous work, by inhibiting C3 cleavage with the peptide inhibitor compstatin Cp40, we showed that properdin binding to zymosan and Escherichia coli is not a primary event, but is completely dependent on initial C3 deposition (35). In the present study, we aimed to investigate whether the binding of properdin to other tentative targets for pattern recognition, i.e., granulocyte myeloperoxidase (MPO), human umbilical vein endothelial cells (HUVECs), and Neisseria meningitidis, is a primary event or dependent on C3 deposition. For this purpose, we compared normal human serum (NHS) with and without compstatin Cp40, as well as C3-depleted human serum with and without reconstitution of purified C3.

Results

Properdin Binding to MPO-Coated Wells After Incubation with Human Serum. Owing to the retained peroxidase activity of immobilized MPO, the binding of properdin to MPO-coated wells was not possible to evaluate in our regular ELISA with HRP-labeled detection antibody and substrate dependent on peroxidase activity. Thus, a detection system based on alkaline phosphatase-labeled anti-mouse IgG antibody was developed. A pool of NHS incubated in serial dilutions on the MPO coat led to a dose-dependent binding of properdin (Fig. 1A). In contrast to properdin-containing NHS, there was minimal binding of purified properdin using a concentration corresponding to that in NHS (OD = 1.35 vs. 0.17), with a signal similar to that of the buffer control (OD = 0.12) (Fig. 1B).

MPO-Induced Terminal Pathway Activation and Dependency on C3. MPO was previously shown to activate complement via the AP (28). MPO coated onto microtiter plates activated complement, as detected by binding of the monoclonal antibody aE11, specific for a C9 neoepitope in the terminal C5b-9 complement complex (36). No generation of C5b-9 on the surface by MPO occurred in human serum depleted of C3 (OD = 0.13); however, on reconstitution with purified C3, a strong deposition of C5b-9 was found (OD = 0.81), which was efficiently blocked by compstatin Cp40 (OD = 0.13) (Fig. 2A).

C3-Dependent Binding of Properdin to MPO. We next examined C3 dependence for the binding of properdin to MPO. Binding of properdin in NHS to MPO was completely abolished when C3 cleavage was inhibited with compstatin Cp40 (OD = 2.92 vs. 0.15) (Fig. 2B, Left). Similarly, no binding of properdin occurred from C3-depleted human serum, but binding was restored by reconstitution with purified C3 (OD = 0.10 vs. 1.44) (Fig. 2B, Middle). This binding was abrogated in the presence of compstatin Cp40 (OD = 0.10).

We then assessed binding in properdin-depleted serum before and after reconstitution with purified properdin in the presence and absence of compstatin Cp40 (Fig. 2B, Right). Properdin binding was achieved when properdin-depleted human serum was reconstituted with purified properdin (OD = 1.33), but was again completely dependent on C3 cleavage, as demonstrated by the fact that compstatin Cp40 reduced the binding to background level (OD = 0.17). The purified properdin used was stored at −70 °C and thawed only once for the experiments. Because the composition and quality of the purified properdin preparation used in these experiments are crucial for correct interpretation of the data, we performed a careful characterization of the purified properdin preparation by exclusion chromatography,
SDS/PAGE, and Western blot analysis, as described in SI Materials and Methods and shown in Fig. S1.

**C3-Dependent Binding of Properdin to HUVECs.** We tested HUVECs incubated with NHS for the binding of C3d and properdin. Incubation for 4 h on HUVECs resulted in substantial deposition of C3d fragments detected by flow cytometry, with a median fluorescence intensity (MFI) of 813 (Fig. 3A and B). In the presence of compstatin Cp40, C3d deposition was virtually abolished (>97%). Properdin binding on HUVECs followed the same trend and was largely dependent on C3 (Fig. 3C and D). In the presence of compstatin Cp40, properdin binding from NHS was reduced by 90%, from an MFI of 40.6 to an MFI of 16.7, a value in the same range as the isotype control (MFI = 13.9). Controls with properdin- and C3-depleted human serum showed comparably low values with a distinct increase in properdin binding after reconstitution with purified properdin and C3, respectively (Fig. 3E).

**C3-Dependent Binding of Properdin to N. meningitidis.** Binding of properdin from NHS to *N. meningitidis* was evaluated by flow cytometry. Incubation of *N. meningitidis* in NHS diluted 1:2 yielded a distinct signal for deposition of properdin on the bacteria (MFI = 31.2) (Fig. 4A and B). The addition of compstatin Cp40 abrogated this deposition completely (MFI = 3.6), showing values similar to the isotype control (MFI = 3.8). Virtually identical findings were seen with C3-depleted human serum (Fig. 4C and D). After exposure of *N. meningitidis* to C3-depleted human serum, properdin deposition was in the range of the isotype control (MFI = 4.3). Reconstitution of C3-depleted

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**Fig. 2.** C3 dependence for complement activation and properdin binding on MPO. (A) NHS and C3-depleted human serum diluted 1:5 with and without purified C3 (130 μg/mL) were incubated on MPO-coated wells (gray bars) or uncoated wells (white bars) in the presence or absence of compstatin Cp40. Complement activation triggered by MPO was detected as surface deposition of C5b-9 using the monoclonal antibody aE11, specific for a C9 neoepitope in C5b-9. (B) The same experimental setup as in A, but also including properdin-depleted human serum diluted 1:5 with and without properdin (2 μg/mL), was tested for the binding of properdin to MPO. Properdin was detected as in ELISA with mouse anti-human properdin. Data are presented as mean with 95% CI OD values of six experiments. ***P < 0.001.

**Fig. 3.** C3 deposition and binding of properdin to the surface of HUVECs. Undiluted NHS with and without compstatin Cp40 (A–D) or human serum depleted and reconstituted for C3 or properdin (E) was incubated for 4 h at 37 °C on monolayers of confluent HUVECs. Deposition of C3d fragments (A and B) and binding of properdin (C–E) were analyzed by flow cytometry. A and C show representative curves for binding of C3d and properdin (continuous lines), respectively, to HUVECs in NHS (gray background) or NHS with compstatin Cp40 (white background), or in NHS binding of the isotype control (dotted line). B and D show individual values and mean with 95% CI of six experiments. *P < 0.05; **P < 0.01. Stippled lines represent the range for binding of the isotype control. E shows the mean ± SD value of six experiments.
human serum with purified C3 showed binding approaching the levels in NHS (MFI = 15.1).

**Discussion**

In the present study, we have demonstrated that properdin is unable to bind directly to granulocyte MPO, to the surface of HUVECs, or to *N. meningitidis*, but that the interaction is fully dependent on initial C3 activation and thus does not act as a pattern recognition molecule for these targets. We investigated these targets because some interesting observations regarding the interaction between them and properdin have been reported. MPO was recently reported to directly attract serum properdin for initiation of the AP (28). *N. meningitidis* and endothelial cells were studied based on their central role in properdin pathophysiology. Properdin is significant in the defense against *Neisseria* species, and *Neisseria gonorrhoeae* was claimed to directly bind properdin (7), findings that have been extrapolated to *N. meningitidis* (37). Endothelial cells are a source of serum properdin (38), and properdin has been shown to interact with various glycosaminoglycans on endogenous cells (25, 39, 40).

In a previous study, we showed that inhibition of C3 cleavage could serve to distinguish between the primary binding of properdin (i.e., pattern recognition function) and properdin binding secondary to initial C3 fragment deposition (i.e., convertase stabilization) (35). Using the C3 inhibitor compstatin Cp40, we consistently found that properdin in human serum and properdin released from activated polymorphonuclear leukocytes did not bind directly to zymosan or *E. coli*, but that binding of properdin was fully dependent on the initial deposition of C3b. By using the same principle here, properdin binding to MPO, HUVECs, and *N. meningitidis* was found to be fully dependent on accessible C3. Thus, the addition of a C3 inhibitor, compstatin Cp40, to NHS or the use of C3-depleted serum did not show any direct binding of properdin to these five different tested surfaces and ligands that we tested, of which three were exogenous and two were endogenous, supporting the notion that properdin is not a promiscuous molecule binding to various “danger” ligands.

The background of the present study was the controversy surrounding properdin acting as a pattern recognition molecule in activation of the AP. Properdin lacks the structural homology shared by other pattern recognition molecules of the complement system. C1q and the recognition molecules of the lectin pathway, including mannose-binding lectin, ficolins, and collectins, all have a common structure with a bouquet of globular heads built up of a cysteine-rich N-terminal stretch, a collagen-like domain, and a carbohydrate-recognition domain or fibrinogen-like domain (41). These molecules bind their respective targets with high avidity using multiple sites for interaction (42, 43). This is distinctly different from properdin, which is a linear molecule composed of repeating thrombospondin type I domains (44), a noteworthy difference from the typical fluid-phase pattern recognition molecules of innate immunity. Although this does not completely exclude properdin for recognizing certain structures besides the AP convertase, it is in line with our findings, making it unlikely that properdin can be defined as a recognition molecule.

Indications of the role of properdin as pattern recognition molecule started with experiments showing that purified properdin covalently bound to a biosensor surface could serve as a platform for the in situ assembly of AP C3 convertases (23). Initial data introduced the attractive concept of how properdin could not only stabilize the C3bBb convertase, but also promote surface attraction of preformed fluid-phase C3bBb; however, studies using surface plasmon resonance depend on the use of purified proteins in a buffer system, which introduces a certain degree of artificiality. Properdin in purified preparations is prone to aggregation, and extrapolating findings on covalently bound properdin to native serum properdin interacting on a biological substrate is virtually impossible. Nonetheless, later data indicated that AP is indeed initiated by the noncovalent attachment of properdin on surfaces of different microorganisms, including yeast cell walls, *N. gonorrhoeae*, and LPS-negative *E. coli* (7). However, these experiments were performed in a system permitting autoactivation of C3 (12, 45) and allowing slow, continuous surface C3b deposition, thus making it difficult to determine whether properdin reacts primarily to the activating surface as a true pattern recognition molecule or binds secondarily to C3b.

Agarwal et al. (46) later extensively studied the ability of several strains of *N. gonorrhoeae* and *N. meningitidis* to directly bind properdin. None of the tested strains bound native properdin, but binding did occur with purified preparations of properdin containing high-order oligomers of aggregated properdin, a phenomenon later demonstrated on zymosan and *Streptococcus pneumoniae* as well (46, 47). Aggregates of properdin were shown to appear spontaneously in certain purified preparations on preparation or during long-term storage or freezing and thawing (30, 31). These aggregates are artifactuals with different properties than serum properdin; for example, they promote fluid-phase complement activation and C3 depletion, as illustrated by the old term “activated properdin” (30, 31). Ali et al. (47) recently showed that recombinant highly polymerized properdin can be a powerful agent to therapeutically combat infections by *N. meningitidis* and *S. pneumoniae*. Mice infected with these bacteria were protected when given recombinantly expressed properdin containing high-order oligomers, which, in
contrast to low-order oligomers of properdin in plasma, directly interact with the bacterial surface and focus C3b deposition (47).

In contrast to our findings with exogenous ligands, previously shown with zymosan and E. coli (35) and in the present work with *N. meningitidis*, there is evidence of C3-independent properdin binding to ligands on endogenous cells, such as DNA on late apoptotic or necrotic cells (26), and presumably various glycosaminoglycans on early apoptotic T cells, but not necrotic T cells (25), activated platelets (27), and renal tubular epithelial cells (39, 40). To the best of our knowledge, no data for endothelial cells and properdin binding have been reported previously, even though endothelial cells are a source of serum properdin and can express heparin sulfate proteoglycans, which have been shown to bind properdin (39, 48). In the present study, after incubation of human serum on HUVECs for 4 h, surface complement activation was detected as deposition of C3d fragments on the cells. Properdin was also found on the cells, but, like for *N. meningitidis*, this binding was dependent on initial C3 activation, because compstatin Cp40 almost entirely inhibited properdin binding.

Recently, MPO was added to the list of suggested endogenous molecules capable of binding properdin directly (28). Thus, MPO could be the molecule that links properdin binding to the surface of activated neutrophils, as has been reported previously (49). By studying properdin binding to immobilized MPO, the authors proposed that MPO promotes complement activation by direct binding of properdin (49). We questioned those findings, however, because they were based on a detection system using HRP-labeled antibodies in which we found it impossible to distinguish substrate turnover caused by peroxidase activity by the antibody-linked HRP and that caused by the immobilized MPO. By changing the enzyme and substrate in the detection system, we showed that properdin was not attracted to MPO in the absence or inhibition of C3; however, immobilized MPO did serve as an excellent complement activator by promoting the formation of C5b-9 on the surface and, in the presence of C3, the binding of properdin.

In conclusion, we challenge the view of properdin as a pattern recognition molecule by providing evidence that it binds to different exogenous and endogenous molecular patterns in only a C3-dependent manner. To support the claim that properdin is a pattern recognition molecule, it should be clearly shown that the initial binding and subsequent activation are C3-independent, and the use of purified properdin preparations should be critically evaluated. Taken together, our data provide further support for the role of properdin as a specific stabilizer of the AP C3 convertase that generally does not act as a recognition molecule.

**Materials and Methods**

**Antibodies and Other Reagents.** Two mouse anti-human properdin IgG1 antibodies (P1: A233 and P2: A235) were obtained from Quidel, and anti-C3d from Jackson ImmunoResearch. The compstatin analog Cp40 (FITC was obtained from Southern Biotech, and goat anti-mouse IgG-FITC was obtained from GE Healthcare UK, goat anti-mouse IgG1-phatase-labeled anti-mouse IgG was obtained from Sigma-Aldrich, F(ab′)2 anti-mouse IgG (whole molecule) (1:10,000). Deposition of C5b-9 onto MPO surfaces or controls after incubation with NHS with or without compstatin Cp40 or with C3-depleted serum with or without purified C3 (130 μg/mL), and on properdin-depleted serum (1:5) with or without purified properdin (2 μg/mL). Binding of properdin was detected with mAb anti-properdin (1:1,000), followed by alkaline phosphatase-labeled sheep F(ab′)2, anti-mouse IgG (whole molecule) (1:1,000). Deposition of C5b-9 was evaluated with mAb anti-C5 (diluted 1:500) or isotype control clone IS7, all followed by goat anti-mouse IgG-FITC (1:50). Samples were washed twice, resuspended in PBS with 1% BSA, and analyzed on a FACSSilmar flow cytometer (BD Biosciences). *N. meningitidis* was gated on forward-scatter and side-scatter, and 20,000 events were collected. Data were evaluated using FlowJo version 10.

**Binding of Properdin to MPO.** Microtiter plates (Costar high-binding; Corning) were coated with MPO (20 μg/mL, dissolved in 0.05 M NaClO4, pH 9.5) by overnight incubation at room temperature, or incubated solely with 0.05 M NaClO4 for 1 h. All wells were subsequently blocked with 3% BSA (2 h). Two-fold dilutions of properdin (in PBS pH 7.4) and Tween (0.1%) for 60 min. NHS, serially diluted in 2.5 mM veronal-buffered saline (VBS), pH 7.2, with MgEGTA (5 mM Mg2+ and 10 mM EGTA) or purified properdin (2 μg/mL in VBS/MgEGTA) were incubated on MPO-coated surfaces or controls for 60 min at 37 °C. In separate sets of experiments, incubation was done on MPO-coated surfaces with NHS (1:5) with or without compstatin Cp40 (8 μg/mL) on C3-depleted serum (1:5) with or without purified C3 (130 μg/mL) and on properdin-depleted serum (1:5) with or without purified properdin (2 μg/mL). Binding of properdin was detected with mAb anti-properdin (1:1,000), followed by alkaline phosphatase-labeled sheep F(ab′)2, anti-mouse IgG (whole molecule) (1:1,000). Deposition of C5b-9 was determined with mAb anti-C5 (diluted 1:500) or isotype control clone IS7, followed by F(ab′)2, anti-mouse IgG-FITC (1:50). Samples were washed twice, resuspended in PBS with 1% BSA, and analyzed on a FACSSilmar flow cytometer (BD Biosciences). *N. meningitidis* was gated on forward-scatter and side-scatter, and 20,000 events were collected. Data were evaluated using FlowJo version 10.

**Binding of Properdin to *N. meningitidis***. *N. meningitidis* were washed by repeated centrifugation at 3,220 × g for 10 min and then resuspended in PBS with 0.1% BSA. Bacteria were then resuspended in VBS with MgCl2 (0.5 mM), CaCl2 (2 mM), and gelatin (0.1%), pH 7.5. NHS was preincubated or with without compstatin Cp40 (40 μM) for 5 min and then added to the bacteria (final concentration 1:2) for 1 h at 37 °C. After incubation, bacteria were washed by re-suspension in PBS, serially diluted in 2.5 mM veronal-buffered saline (VBS), pH 7.2, with MgEGTA (5 mM Mg2+ and 10 mM EGTA) or purified properdin (2 μg/mL in VBS/MgEGTA) were incubated on NHS (1:5) with or without compstatin Cp40 (8 μg/mL) on C3-depleted serum with or without purified C3 (130 μg/mL), and on properdin-depleted serum (1:5) with or without purified properdin (2 μg/mL). Binding of properdin was detected with mAb anti-properdin (1:1,000), followed by alkaline phosphatase-labeled sheep F(ab′)2, anti-mouse IgG (whole molecule) (1:1,000). Deposition of C5b-9 was determined with mAb anti-C5 (diluted 1:500) or isotype control clone IS7, followed by F(ab′)2, anti-mouse IgG-FITC (1:50). Samples were washed twice, resuspended in PBS with 1% BSA, and analyzed on a FACSSilmar flow cytometer (BD Biosciences). *N. meningitidis* was gated on forward-scatter and side-scatter, and 20,000 events were collected. Data were evaluated using FlowJo version 10.

**Statistics.** GraphPad Prism version 5 was used for statistical analyses. Statistical differences were analyzed on normally distributed data using the Student t test or, when more than two conditions were compared, with one-way ANOVA followed by the Bonferroni posttest. A P value <0.05 was considered statistically significant.

**Ethics Statement.** The study was designed and performed according to the ethical guidelines from the Declaration of Helsinki. Informed written consent
The down-stream effects of mannan-induced lectin comple-
ment activation.

The quantitative role of al-
ternative pathway C3 convertase.


