Complement has emerged as a likely driver of the immune response and end-organ damage in COVID-19. In patients with severe disease, deposition of terminal complement and microthrombosis have been observed in the lung, skin, kidney, and heart.1-4 Recently, we demonstrated that the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) spike protein leads to amplification of the alternative pathway of complement on cell surfaces through competition with complement factor H (CFH) for binding heparan sulfate.5 Thus, in vitro, the SARS-CoV-2 spike protein can convert an inactivator surface to an activator surface on nucleated cells.

Two messenger RNA (mRNA)-based vaccines that lead to transient expression of the SARS-CoV-2 spike protein are highly efficacious in preventing severe infection.6,7 Reactions to these vaccines are generally mild; however, increased complement amplification could theoretically lead to more severe effects in diseases like paroxysmal nocturnal hemoglobinuria (PNH), where blood cells lack complement regulatory proteins.8 Here, we describe significant adverse reactions to COVID-19 vaccines in 4 patients with PNH. We also present 2 patients with PNH who received the vaccine without significant adverse effects or hemolysis.

Patient characteristics and reactions to the COVID-19 vaccines are shown in Table 1. Patients were identified based on self-report of receiving the vaccine. Patients were aged 25 to 63 years, had PNH granulocyte clones of ≥80%, and had not received transfusions in the past year. Reactions occurred from the day of administration to 5 days later and lasted 1 to 6 days. Four of 6 patients reported fever. Patients 2, 3, and 4 experienced severe hemolysis with 2 to 4 g/dL hemoglobin decrease. Patient 1 had a presumptive thrombotic manifestation. Patients 5 and 6 received both doses of the Pfizer-BioNTech BNT162b2 (Pfizer-BioNTech) COVID-19 vaccine without clinical or laboratory signs of hemolysis.

Patient 1 is a 25-year-old man diagnosed with PNH 6 months prior in the setting of hemoglobinuria and had limited disease manifestations not requiring PNH-directed therapy. Five days after receiving the second dose of the Pfizer-BioNTech vaccine, he developed abdominal pain leading to evaluation in an emergency department. Computed tomography with contrast showed peripancreatic fat stranding with normal lipase, concerning for possible small bowel microvascular thrombosis. D-dimer was elevated to 0.73 μg/mL fibrinogen-equivalent units from 0.21 μg/mL. His symptoms resolved after 5 days. He subsequently was initiated on ravulizumab, a terminal complement inhibitor.

Patient 2 is a 45-year-old man with a 20-year history of PNH. His last dose of ravulizumab was 4 weeks prior to vaccination. On the evening after his first dose of the Pfizer-BioNTech vaccine, he experienced fever, headache, myalgia, and severe fatigue, which lasted 6 days. He also developed hemoglobinuria on postvaccination days 1 and 2, associated with 2.7 g/dL hemoglobin decrease.

Patient 3 is a 32-year-old woman with a 10-year history of PNH on ravulizumab with her last dose 4 weeks prior to vaccination. She was also part of a clinical trial with danicopan, an oral complement factor D inhibitor. She missed 2 doses of danicopan immediately after her second vaccination due to concerns that the drug may interfere with vaccine potency; therefore, danicopan was not at therapeutic levels at the time of her reaction. Notably, she took danicopan throughout her first vaccination and did not experience breakthrough hemolysis. Approximately 12 hours after receiving the second dose of the Moderna mRNA-1273 COVID-19 vaccine, she had a fever (39°C) and rigors. She presented to a local hospital, where she was noted to have a 3 g/dL hemoglobin decrease and received 2 units of packed red blood cells.

Patient 4 is a 63-year-old man diagnosed with PNH 30 years ago, currently treated with ravulizumab. He experienced fatigue and darkening of his urine following his first dose of the Moderna vaccine. He had an ~1 g/dL hemoglobin decrease on laboratory tests 3 days following his first vaccination. In addition, his total bilirubin rose to 7.1 mg/dL from baseline 2.4 mg/dL. Following his second dose, he noted fevers, diarrhea, vomiting, severe fatigue, and dark urine. Laboratory tests 1 week later, as symptoms were resolving, showed >4 g/dL hemoglobin decrease from his baseline.

Based on these observations, we sought to evaluate whether the SARS-CoV-2 spike protein directly leads to increased hemolysis. Recent data suggest the SARS-CoV-2 spike protein binds heparan sulfate on nucleated cells9 and amplifies the alternative pathway of complement through interference with the binding of CFH, an alternative pathway inhibitor.5 However, CFH primarily binds sialic acid on human erythrocytes,10 and mature erythrocytes express little heparan sulfate.11
Table 1. Characteristics of patients with PNH and COVID-19 vaccine reactions

| Patient | 1   | 2   | 3   | 4   | 5   | 6   |
|---------|-----|-----|-----|-----|-----|-----|
| Sex     | Male| Male| Female| Male| Female| Female|
| Age, y  | 25  | 45  | 32  | 63  | 51  | 59  |
| Age at diagnosis, y | 24  | 24  | 23  | 31  | 43  | 46  |
| Disease history | Hemoglobinuria | Aplastic anemia, transfusion dependence, microvascular small bowel thrombosis, renal failure, smooth muscle dystonia | Aplastic anemia, transfusion dependence | Transfusion dependence, hemoglobinuria, smooth muscle dystonia | Hemoglobinuria, fatigue | Hemoglobinuria, fatigue, extravascular hemolysis with transfusion dependence on C5 inhibition |
| Complement inhibitor treatment | No | Ravulizumab | Ravulizumab* | Ravulizumab | Ravulizumab | Ravulizumab, Danicopan |
| PNH clone | 11% RBC, 80% granulocytes | >99% RBC, >99% granulocytes | 39% RBC, 99% granulocytes | 99% RBC, 99% granulocytes | 77% RBC, 95% granulocytes | 76% RBC, 81% granulocytes |
| Blood type | A positive | A positive | A positive | O positive | A negative | O positive |
| COVID-19 vaccine | Pfizer-BioNTech | Pfizer-BioNTech | Moderna | Moderna | Pfizer-BioNTech | Pfizer-BioNTech |
| Vaccine dose | 2 | 1 | 2 | 2 | 2 | 2 |
| Last dose ravulizumab prior to vaccination | N/A | 4 wk | 4 wk | 4 wk (dose 1) | 7 wk (dose 2) | 5 wk | 6.5 wk (dose 2) |
| Time to symptom onset postvaccine | 5 d | Same day | Same day | Same day | 1 d | N/A |
| Duration of symptoms | 5 d | 6 d | 1 d | 6 d | 1 d | N/A |
| Hemoglobin prevaccination, g/dL | 10.9 | 11.6 | 11.3 | 11.9 | 10.9 | 11.4 |
| Hemoglobin postvaccination, g/dL | 10.4 | 8.9 | 8.4 | 10.7 (dose 1) | 7.1 (dose 2) | 11.1 | 11.0 (dose 1) | 10.7 (dose 2) |
| LDH prevaccination, U/L | 1127 | 211 | 255 | 305 | 207 | 263 |
| LDH postvaccination, U/L | 1633 | — | — | 312 (dose 1) | 342 (dose 2) | 220 | 276 (dose 1) | 258 (dose 2) |
| Total bilirubin prevaccination, mg/dL | 0.8 | 1.4 | 0.7 | 2.4 | 0.6 | 1.5 |
| Total bilirubin postvaccination, mg/dL | 2.5 | — | 2.7 | 7.1 (dose 1) | 3.0 (dose 2) | — | 2.1 (dose 1) | 1.2 (dose 2) |

*denotes no information is available; AST, aspartate amino transferase; LDH, lactate dehydrogenase; N/A, not applicable; RBC red blood cell.

*Additionally on danicopan; however, 2 doses were missed immediately following vaccination.

†None of these patients were previously transfused in the past year.
To test the effect of the SARS-CoV-2 spike protein on hemolysis, we performed erythrocyte lysis using erythrocytes from a patient with PNH and acidified normal human serum (aNHS) with addition of the SARS-CoV-2 spike protein subunit 1 (S1) (see supplemental methods, available on the Blood Web site). Briefly, type O-positive red blood cells from 1 patient with PNH and 1 control were collected. aNHS preincubated with and without S1 was added to the erythrocytes. Following incubation at 37°C for 1 hour, absorbance at 405 nm was measured in the cell-free supernatants. Hemolysis in each sample was compared with total water-induced lysis of the erythrocytes. S1 did not increase hemolysis of PNH erythrocytes as compared with aNHS alone (Figure 1). Furthermore, S1 does not appear to bind erythrocytes (data not shown). This suggests that postvaccination hemolysis is not mediated by the direct effect of the spike protein. We postulate that strong complement amplification as a byproduct of the inflammatory response is responsible for the clinically observed hemolysis, as has been reported with other vaccines, infections, and surgeries.

In patients with PNH on complement inhibition, strong complement-amplifying conditions such as infection, surgery, or pregnancy may trigger pharmacodynamic breakthrough (as previously defined by Ristano et al).12 Complement-amplifying conditions lead to C3b accumulation on the cell surface; at high densities of C3b, C5 can assume a conformational change, disrupting the ability of eculizumab to inhibit terminal complement.13,14 Hemolysis following COVID-19 vaccination, which occurred in 3 patients on ravulizumab, suggests pharmacodynamic breakthrough. Although many vaccines can lead to hemolysis and thrombosis in PNH, this effect is mitigated in most patients on complement inhibitors.15 Ravulizumab, a new C5 inhibitor with a half-life 4 times longer than eculizumab, is reported to have significantly fewer instances of pharmacokinetic breakthrough hemolysis.16 Furthermore, 3 instances of breakthrough hemolysis occurred 4 weeks from the last ravulizumab infusion, making suboptimal C5 inhibition unlikely. As seen in patients 3 and 6, a proximal complement inhibitor, such as
Danicopan, may prevent breakthrough hemolysis precipitated by the vaccine; however, it is equally possible that the stronger immune response after the second vaccine dose was primarily responsible for the breakthrough hemolysis in patient 3.

This study provides insight into the mechanism of pharmacodynamic breakthrough precipitated by COVID-19 vaccination in patients with PNH on ravulizumab. Similar disease flares may be anticipated in other complement-mediated disorders, such as complement-mediated hemolytic uremic syndrome, cold agglutinin disease, catastrophic antiphospholipid syndrome, and HELLP syndrome. As SARS-CoV-2 leads to a severe inflammatory state, the benefits of vaccinating patients with PNH likely outweigh the risks; however, clinicians and patients should be aware of this serious adverse effect, and patients should be educated to report any symptoms postvaccination. We recommend vaccination within 4 weeks of the last ravulizumab infusion and 1 week of eculizumab infusion and that patients maintain optimal hydration. Adverse reactions appear time-limited and can be managed with supportive care and transfusions as needed.

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Authorship

Contribution: G.F.G. performed research, collected and analyzed data, and wrote the first draft of the manuscript; X.Y. designed and performed research; J.Y., B.A.Y.C., and E.M. contributed reagents or analytic tools; and all authors reviewed the manuscript and approved the final version.

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ORCID profiles: G.F.G., 0000-0001-9855-0075; X.Y., 0000-0002-7304-9927; J.Y., 0000-0001-9773-5377; B.A.Y.C., 0000-0002-2035-5496; E.M. B., 0000-0001-5072-7636; S.C., 0000-0001-6832-2389; R.A.B., 0000-0001-5741-1255.

Correspondence: Robert A. Brodsky, Division of Hematology, School of Medicine, Johns Hopkins University, 720 Rutland Ave, Ross Research Bldg, Room 1025, Baltimore, MD 21205; e-mail: brodso@jhmi.edu.

Footnotes

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