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Multiple *Plasmodium falciparum* Erythrocyte Membrane Protein 1 Variants per Genome Can Bind IgM via Its Fc Fragment Fcγμ

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The *Plasmodium falciparum* erythrocyte membrane protein 1 (PfEMP1) adhesive proteins expressed on the surfaces of infected erythrocytes (IEs) are of key importance in the pathogenesis of *P. falciparum* malaria. Several structurally and functionally defined PfEMP1 types have been associated with severe clinical manifestations, such as cerebral malaria in children and placental malaria in pregnant women. PfEMP1 that can bind the Fc part of IgM (Fcγμ) characterizes one such type, although the functional significance of this IgM binding to PfEMP1 remains unclear. In this study, we report the identification and functional analysis of five IgM-binding PfEMP1 proteins encoded by *P. falciparum* NF54. In addition to the VAR2CSA-type PFL0030c protein, already known to bind Fcγμ and to mediate chondroitin sulfate A (CSA)-specific adhesion of IEs in the placenta, we found four PfEMP1 proteins not previously known to bind IgM this way. Although they all contained Duffy binding-like (DBLε) domains similar to those in VAR2CSA-type PfEMP1, they did not mediate IE adhesion to CSA, and IgM binding did not shield IEs from phagocytosis of IgG-opsonized IEs. In this way, these new IgM-binding PfEMP1 proteins resemble the rosette-mediating and IgM-binding PfEMP1 HB3VAR06, but none of them mediated formation of rosettes. We could map the capacity for Fc-specific IgM binding to DBLε domains near the C terminus for three of the four PfEMP1 proteins tested. Our study provides new evidence regarding Fc-dependent binding of IgM to PfEMP1, which appears to be a common and multifunctional phenotype.

Many microorganisms express molecules that can bind immunoglobulins independently of the antigen specificity of the antibodies. A prominent example is the antibody-binding proteins found in the cell wall of the bacterium *Staphylococcus aureus* (1, 2). These proteins have high affinities for the conserved elements in the Fc and Fab parts of various antibody classes, and they appear to serve an immunoevasive function, as binding of antibodies to these proteins interferes with phagocytosis of antibody-opsonized bacteria (3).

Some erythrocytes infected by the malaria parasite *Plasmodium falciparum* bind IgM, but not IgG, independently of the specificity of the antibodies (4, 5). This Fc-mediated binding of IgM has been described for infected erythrocytes (IEs) that bind to the sulfated glycosaminoglycan chondroitin sulfate A (CSA) (5) and for IEs capable of forming rosettes (several uninfected erythrocytes adhering to a central IE) (4). Both IE phenotypes are related to expression of particular types of *P. falciparum* erythrocyte membrane protein 1 (PfEMP1). Thus, adhesion of IEs to CSA requires expression of the atypical PfEMP1 type VAR2CSA, which has nanomolar affinity for CSA and is responsible for placental IE sequestration (6–9). Rosetting can be mediated by several different PfEMP1 proteins that have a semiconserved N-terminal head structure made up of certain subtypes of Duffy binding-like α (DBLα)—cysteine-rich interdomain region γ/δ (CIDRγ/δ) domains (10–13), and it appears to depend mainly on relatively low-affinity interactions with a range of host carbohydrates (14–17).

The function of Fc-dependent binding of IgM to *P. falciparum* IEs is not fully understood (reviewed in reference 18). In the case of VAR2CSA-type PfEMP1, it appears to be mainly immunoevasive, as it can protect IEs from specific IgG recognition and immune destruction without compromising the CSA-adhesive function of the antigen (19). However, such masking is ineffective in the case of rosette-mediating PfEMP1 antigens (15), where binding of IgM to PfEMP1—in combination with other serum factors—seems to function to increase the low-affinity adhesive interactions involved in rosetting (15, 20). Given the apparent clinical importance of IgM binding (4, 21), it is of interest to know how many IgM-binding PfEMP1 variants exist within the PfEMP1 repertoire of a single *P. falciparum* clone and how IgM binding is related to the structural and functional characteristics of the involved PfEMP1 proteins. We therefore set out to identify IgM-binding PfEMP1 proteins in *P. falciparum* NF54. We show that the genes for at least five IgM-binding PfEMP1 variants exist in the genome of this parasite. In addition to PFL0030c, which is the VAR2CSA-type antigen in *P. falciparum* NF54, we found four others (PFL0020w, PF07_0139, MAL6P1.4, and MAL6P1.316). Surprisingly, these did not mediate rosetting in functional assays and do not possess structural features indicative of being rosette mediating. Our study shows that Fc-mediated binding of IgM to
PIEMP1 proteins is not limited to those that can adhere to CSA or mediate formation of rosettes.

MATERIALS AND METHODS

Recombinant PIEMP1 proteins and specific antisera and monoclonal antibodies. Recombinant proteins representing full-length PFL0030c and single- and triple-domain constructs of MAL6P1.4, MAL6P1.316, PFL0020w, and PFL0003c were produced in a baculovirus expression system, essentially as described previously (15, 22). The domain nomenclature proposed by Rask et al. in 2010 is used throughout (23). Antisera against MAL6P1.4, MAL6P1.316, and PFL0020w were raised in rats (24), and the human monoclonal antibody PAM1.4, specific for several VAR2CSA-type PIEMP1 proteins, including PFL0030c, was generated as described elsewhere (25, 26). Nonimmune IgM binding to recombinant PIEMP1 constructs was quantified by enzyme-linked immunosorbent assay (ELISA) as described previously (15).

Malaria parasite cultivation and in vitro selection procedures. P. falciparum NF54 parasites (27) were grown in vitro in O Rh+ erythrocytes at 37°C in a controlled atmosphere, using complete culture medium (RPMI 1640 supplemented with 0.5% AlbuMax II [Life Technologies BV, Nerum, Denmark]), essentially as described previously (28). The P. falciparum NF54-derived and pBVH-transfected clone G6 was generated as described elsewhere (29–31). It was maintained in the same way as the parental strain P. falciparum NF54, except that blastocidin (10 mg/ml; Life Technologies) was added to shut down transcription of endogenous var genes and to erase the epitogenic memory. IEs were selected for surface expression of defined PIEMP1 proteins by immunomagnetic selection using (i) PIEMP1-specific rat antisera followed by biotinylated anti-rat antibody (Dako) and streptavidin-coupled Dynabeads (Fisher Scientific), or (ii) PAM1.4 followed by protein A-coupled Dynabeads (Fisher Scientific), essentially as described previously (32). For selection of IgM-binding IEs, we used human IgM (Sigma) coupled to M-450 epoxy beads (Life Technologies) according to the manufacturers’ instructions. The genotypic identity of the parasites and the absence of Mycoplasma contamination were verified regularly as described previously (33).

var gene transcription analysis. For analysis of var gene transcription by quantitative real-time PCR, we used cDNA generated from ring-stage parasite RNA and P. falciparum 3D7 var gene-specific primers as described in detail elsewhere (6, 34, 35). Transcription levels relative to those of the seryl-tRNA synthetase housekeeping gene were calculated by the 2−ΔΔCT method.

Antibody labeling of IEs. The binding of nonimmune IgM and PIEMP1-specific antibody to IEs was quantified by flow cytometry, essentially as described previously (19). In brief, we used magnetically activated cell sorting (MACS) to purify late-stage IEs, labeled with ethidium bromide as described above, and IE surface expression of defined PfEMP1 proteins is not limited to those that can adhere to CSA or mediate formation of rosettes.

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Statistical analysis. One-way analysis of variance followed by Holm-Sidak multiple pairwise comparisons was used to evaluate the capacity of Fcγ-specific antibodies to inhibit binding of IgM to IEs, and Student’s t test was used to evaluate the ability of IgM and IgA to inhibit phagocytosis of IgG-opsonized IEs. P values of <0.05 were considered statistically significant.

RESULTS
Identification of candidate IgM-binding PfEMP1 proteins in P. falciparum NF54. We used the P. falciparum NF54-derived clone G6 to identify IgM-binding PfEMP1 proteins. This clone, which is transfected with a pVBH plasmid containing a blasticidin resistance gene controlled by a var promoter, initially transcribed the var gene pfd1015c (31). Culturing this parasite under high blasticidin selection pressure results in a complete shutdown of endogenous var transcription and erasure of epigenetic memory, and removal of drug pressure allows in vitro var switching to begin (31). We labeled IEs with IgM and single-cell sorted IgM-positive IEs into 96-well plates. Four cell-sorting control lines with 50 IEs per well were also established. After in vitro expansion for 3 weeks, we had 19 growing cultures originating from individual single-cell-sorted IEs. var gene transcription was assessed in all lines containing IgM-positive IEs (eight from subclones and two from the 50-cell control sublines). Each of the subclones transcribed multiple var genes as a result of shifts in var transcription during the period of in vitro expansion (Fig. 1A). Nevertheless, all but one subclone (G6.2.5) and one 50-cell subline (G6.1.13) showed dominant expression of a single var gene. One subclone (G6.3.16) and one 50-cell subline (G6.1.14) mainly transcribed pfl0030c. Four clones (G6.1.57, G6.2.20, G6.2.56, and G6.2.79) mainly transcribed pfl0020w, two (G6.3.40 and G6.3.76) predominantly transcribed pf07_0139, and both the 50-cell sublines showed prominent transcription of mal6p1.316. The relative transcription levels of the identified candidate var genes were reflected in the proportions of erythrocytes infected by the subclones and sublines that bound IgM (Fig. 1A).
The PfEMP1 protein encoded by the pfl0030c var gene in *P. falciparum* NF54 is the CSA-adhering VAR2CSA-type protein PFL0030c (6, 7). This protein is known to bind IgM via Fc (19, 39, 40), and several DBL domains have been implicated in this binding (39, 40). Only the three C-terminal DBL domains in PFL0030c are of a type (DBLε) that is also found in non-VAR2CSA-type PfEMP1 proteins. All the non-VAR2CSA-type candidate IgM-binding PfEMP1 proteins we identified above contained C-terminal DBLε domains (Fig. 1A). However, we recently mapped the Fc-ε-binding domain in the rosetting PfEMP1 protein HB3VAR06 to its C-terminal DBLε_8 domain, and two of our candidate var transcripts (pfl0020w and mal6p1.316) additionally contain a DBL (domain each (Fig. 1B)). Together, these data point to C-terminal DBLε and DBLζ domains as being relevant for Fc-mediated IgM binding.

**Verification of the IgM-binding properties of candidate PfEMP1 proteins.** To verify the predicted IgM-binding capacity of our candidate PfEMP1 proteins, we next subjected previously unselected *P. falciparum* NF54 parasites to repeated selection with either the VAR2CSA-specific monoclonal antibody PAM1.4 or rat antisera to PFL0020w and MAL6P1.316. In addition, we selected IEs by using a rat antisera to the PfEMP1 protein MAL6P1.4, because two of our subclones (G.6.256 and G.6.2.79) showed prominent transcription of mal6p1.4 (Fig. 1A). MAL6P1.4 contains three C-terminal DBLε domains (Fig. 1B), making it a strong IgM-binding candidate. We did not have an antisera specific for PF07_0139 and instead subjected subclone G6.3.40 to repeated rounds of selection for IgM binding. As a negative control, we used *P. falciparum* NF54 selected by an antisera specific for the intercellular adhesion molecule 1 (ICAM-1)-binding PfEMP1 protein PFD1235w (33, 41). These selection protocols all resulted in parasites with dominant transcription of the expected var gene (Fig. 2A), and all the selected IEs bound IgM to various degrees, except for the IEs selected for expression of PFD1235w (Fig. 2B and C).

**Determination of the IgM domain involved in binding to PfEMP1.** We previously showed that the Cα3-Cα4 domains of IgM bind to VAR2CSA-type PfEMP1 (19) and to the rosetting PfEMP1 HB3VAR06 (15). We therefore measured the ability of monoclonal antibodies specifically recognizing the Cα2, Cα3, and Cα4 domains of human IgM to interfere with IgM binding to IEs expressing each of the IgM-binding PfEMP1 proteins studied here. In each case, all three Fcα-specific antibodies significantly (P < 0.001 in all cases) reduced the binding of IgM to the IEs (Fig. 3), although the Cα2-specific antibody was less effective than the other two antibodies at inhibiting IgM binding to the VAR2CSA-type PFL0030c protein, in accordance with previous findings (19).

**Determination of the PfEMP1 domains involved in Fc-specific binding of IgM.** Multiple domains have been implicated in Fc-dependent binding of IgM to PfEMP1 (15, 39, 40, 42, 43). We recently used multiple recombinant single- and multi-DBL-domain constructs to map this type of IgM binding to the penultimate C-terminal DBL (DBLζ_8) domain in the rosette-mediating PfEMP1 HB3VAR06 (15). Taking a similar approach here, we used ELISA to identify the domains involved in IgM binding, employing an array of recombinant proteins representing single and triple domains of PFL0030c, PFL0020w, MAL6P1.4, and MAL6P1.316, as well as a construct representing the full ectodomain of PFL0030c (see Fig. 1B for an overview of the recombinant antigen constructs used). With this approach, we unequivocally identified DBLεPAM5 as the IgM-binding domain in the VAR2CSA-type PFL0030c protein (Fig. 4A). This is the penultimate C-terminal DBL domain in that protein and was the most prominent IgM-binding domain in an earlier study of VAR2CSA-type PfEMP1 (39). Similarly, we mapped the IgM-binding capacity of PFL0020w and MAL6P1.4 to their C-terminal domains (DBLε_4_6 and DBLε_3_9, respectively) (Fig. 4B). MAL6P1.4 contained an additional IgM-binding domain (DBLε_2_7). We were unable to map the IgM-binding domain of MAL6P1.316 with our recombinant constructs.

We used surface plasmon resonance analysis to show that the binding affinity of IgM for each of the domains identified to bind IgM by ELISA was high, with *K*~D~ values in the nanomolar range (Table 1), which is the same range as that reported previously (15). Triple-domain constructs had higher affinities (lower *K*~D~ values) than single-domain constructs, which may be related to the construct conformation but likely also reflects the fact that two IgM-binding domains (DBLε_2_7 and DBLε_4_9) were present in our MAL6P1.4 triple-domain construct (Fig. 4 and Table 1).
All constructs displayed rapid association kinetics, with $k_a$ values of $1 \times 10^6$ to $10 \times 10^6$ M$^{-1}$ s$^{-1}$. The dissociation kinetics were more variable but were slower for the triple-domain constructs than the corresponding single-domain constructs, suggesting that the higher-order conformation of the PfEMP1 proteins contributes to retention of IgM once it is bound.

**CSA-binding and rosetting properties of the new IgM-binding PfEMP1 proteins.**

Fc-dependent IgM binding to PfEMP1 has previously been shown for parasites expressing CSA-adherent VAR2CSA-type or rosette-mediating PfEMP1 variants (reviewed in reference 18). We therefore tested the abilities of erythrocytes infected by *P. falciparum* expressing either PFL0030c or each of the four new non-VAR2CSA-type PfEMP1 proteins to bind to CSA and to form rosettes. We included parasites expressing the PfEMP1 protein HB3VAR06 as a positive control in the rosetting assays, as this PfEMP1 is known to mediate IgM-dependent rosette formation (15). IEs expressing the VAR2CSA-type PFL0030c protein adhered strongly to CSA-expressing BeWo cells, in accordance with previous reports (6, 37), while none of the other PfEMP1 proteins mediated significant adhesion to this receptor (Fig. 5A). None of the five IgM-binding PfEMP1 proteins studied here was able to mediate formation of rosettes (Fig. 5B). We previously reported that HB3VAR06-mediated rosetting requires soluble serum factors (15, 20), but serum did not lead to rosetting in IEs expressing any of our new IgM-binding PfEMP1 proteins (Fig. 5B).

**IgM-mediated interference with phagocytosis of IgG-opsonized infected erythrocytes.**

Fc-dependent IgM binding to...
VAR2CSA-type PfEMP1 markedly inhibits the binding of specific IgG to the IE surface and subsequent phagocytosis of opsonized IEs (19). However, this is not the case for parasites expressing the rosette-mediated HB3VAR06 protein, as IgM binding has a limited effect on the phagocytosis of antibody-opsonized IEs (15). To test whether IgM could inhibit binding of PfEMP1-specific IgG to IEs positive for each of the new IgM-binding PfEMP1 proteins identified here, we opsonized the IEs with immune human plasma and subsequently measured phagocytosis by using a robust in vitro assay (38). Phagocytosis of IEs expressing the VAR2CSA-type PFL0030c protein was markedly (about two-thirds) and significantly reduced in the presence of IgM, in accordance with previous data (19) (Fig. 6). A smaller or nonsignificant impact of IgM on IgG-dependent phagocytosis was seen for each of the other PfEMP1 proteins (Fig. 6). In this respect, they therefore resembled the rosette-mediating PfEMP1 HB3VAR06 (15), although none of them appear to be involved in rosetting (Fig. 5B).

### DISCUSSION

The particular virulence of *P. falciparum* parasites is related to their ability to express members of the clonally variant protein family PfEMP1 on the surfaces of the erythrocytes they infect (reviewed in reference 44). The PfEMP1 proteins mediate adhesion of IEs to different host receptors in various tissues to avoid IE destruction in the spleen (45). Each parasite genome contains about 60 PfEMP1-encoding *var* genes that are transcribed in a mutually exclusive manner (46–48). Thus, normally only one PfEMP1 is expressed on the IE surface at any given time, but the parasites can switch transcription among the different *var* genes from one asexual 48-h multiplication cycle to the next, thereby in the absence or presence of IgM and subsequently measured phagocytosis by using a robust in vitro assay (38). Phagocytosis of IEs expressing the VAR2CSA-type PFL0030c protein was markedly (about two-thirds) and significantly reduced in the presence of IgM, in accordance with previous data (19) (Fig. 6). A smaller or nonsignificant impact of IgM on IgG-dependent phagocytosis was seen for each of the other PfEMP1 proteins (Fig. 6). In this respect, they therefore resembled the rosette-mediating PfEMP1 HB3VAR06 (15), although none of them appear to be involved in rosetting (Fig. 5B).

### TABLE 1 Surface plasmon resonance analysis of IgM binding to recombinant PfEMP1 domains

<table>
<thead>
<tr>
<th>PfEMP1</th>
<th>Domain(s)</th>
<th>$k_a$ (M$^{-1}$ s$^{-1}$)</th>
<th>$K_d$ (s$^{-1}$)</th>
<th>$K_D$ (nM)$^b$</th>
</tr>
</thead>
<tbody>
<tr>
<td>MAL6P1.4</td>
<td>DBLe2_7-DBLe7_8-DBLe3_9</td>
<td>$1.21 \times 10^6 \pm 1.02 \times 10^6$</td>
<td>$7.19 \times 10^{-4} \pm 2.73 \times 10^{-4}$</td>
<td>2.07 $\pm$ 2.78</td>
</tr>
<tr>
<td></td>
<td>DBLe2_7</td>
<td>$7.14 \times 10^5 \pm 5.32 \times 10^5$</td>
<td>$5.03 \times 10^{-3} \pm 1.24 \times 10^{-3}$</td>
<td>14.80 $\pm$ 7.43</td>
</tr>
<tr>
<td></td>
<td>DBLe7_8</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td></td>
<td>DBLe3_9</td>
<td>$8.58 \times 10^5 \pm 6.18 \times 10^5$</td>
<td>$9.63 \times 10^{-3} \pm 0.58 \times 10^{-3}$</td>
<td>66.06 $\pm$ 56.22</td>
</tr>
<tr>
<td>PFL0020w</td>
<td>DBL414_4-DBL45_5-DBL46_6</td>
<td>$3.25 \times 10^6 \pm 0.63 \times 10^6$</td>
<td>$3.71 \times 10^{-3} \pm 1.15 \times 10^{-3}$</td>
<td>2.36 $\pm$ 2.31</td>
</tr>
<tr>
<td>PFL0030c</td>
<td>DBLF_PAM5_6</td>
<td>$5.96 \times 10^5 \pm 3.1 \times 10^5$</td>
<td>$3.98 \times 10^{-2} \pm 1.97 \times 10^{-2}$</td>
<td>140.02 $\pm$ 77.91</td>
</tr>
</tbody>
</table>

$^a$ NA, not available because the domain did not bind IgM.

$^b$ The $K_D$ values were calculated based on a 1:1 binding model from 3 to 6 independent analyses, with 4 to 8 different concentrations tested in duplicates per analysis, and the values are shown with composite standard deviations.

**FIG 5** Functional properties of erythrocytes infected by *P. falciparum* parasites expressing defined PfEMP1. The graphs show the ability of IE surface-expressed PfEMP1 proteins to mediate adhesion to CSA on BeWo cells (A) or to mediate rosetting (B) in the presence (white) or absence (black) of serum. Means (bars) and standard deviations (error bars) for triplicate measurements from a representative experiment (of three) are shown.

**FIG 6** IgM-dependent interference with phagocytosis of IgG-opsonized infected erythrocytes. The graph shows the ability of IgM (white) and IgA (black) to interfere with IgG-specific opsonization and phagocytosis of erythrocytes infected by *P. falciparum* parasites expressing defined PfEMP1. Overall means (bars) and standard deviations (error bars) for three independent experiments are shown. ***, significant differences between the two bars.
changing the adhesive and antigenic properties of the IEs (46, 47). By now, it is clear that certain structurally defined and functionally related PfEMP1 subfamilies are involved in particular types of severe malaria (reviewed in reference 49). The role of endothelial protein C receptor (EPCR)-adhering PfEMP1 proteins sharing domain cassette 8 (DC8) and DC13 motifs (50, 51) in the pathogenesis of cerebral malaria is a recent example. The similar role of DC4-containing PfEMP1 proteins adhering to ICAM-1 is another (33). Finally, VAR2CSA-type PfEMP1 proteins adhering to CSA have a well-established key role in the pathogenesis of placental malaria (reviewed in reference 52). The tissue-specific adhesion mediated by these types of PfEMP1 makes it easy to appreciate their role in pathogenesis. It is much less obvious why the ability of some PfEMP1 proteins to bind IgM via Fc should be linked to malaria severity, as has repeatedly been shown (reviewed in reference 18). Several PfEMP1 domains have been implicated in IgM binding (15, 36, 39, 40, 42, 43), but a unifying picture has not yet emerged. Furthermore, it is unclear whether Fc-specific binding of IgM to IEs is restricted to parasites expressing PfEMP1 proteins that can adhere to CSA or lead to the formation of rosettes. Finally, it is not known how many PfEMP1 proteins encoded by any single P. falciparum genome possess this phenotype. Thus, the purpose of the present study was to address these unanswered questions.

Individual P. falciparum parasites transcribe only a single var gene at a time, and this mutually exclusive transcription ensures that normally only a single PfEMP1 variant is present on the surface of any given IE (53–55). Even at the population level, the diversity of PfEMP1 expression is often limited due to the presence of epigenetic memory (reviewed in references 56 and 57). To overcome this difficulty, we selected pVBH-transfected P. falciparum G6 clonal parasites for resistance to plasticidin, which effectively erases the epigenetic var gene transcription memory (29–31). Following induction of var switching, we single-cell sorted IEs from this population based on their capacity to bind IgM at the IE surface, and we obtained subclones and sublines with dominant transcription of four different var genes (Fig. 1). One was pf0030c (6), which encodes the VAR2CSA-type PfEMP1 in the P. falciparum 3D7 clone (from which the G6 clone is derived) and was already known to bind IgM via Fc (19, 39, 40). The other three (mal06p1.316, pf0020w, and pf070139) encode PfEMP1 proteins not previously reported to bind IgM. However, all contain at least one DBL-type domain, similar to the C-terminal domains previously implicated in the IgM-binding capacity of VAR2CSA-type PfEMP1 (39, 40). Finally, the var gene mal06p1.4, which encodes a PfEMP1 with three C-terminal DBL domains in tandem, was also prominently transcribed in several subclones/sublines. These structural characteristics supported the IgM-binding capacity of these PfEMP1 variants. Two of the genes (pf0020w and mal06p1.316) additionally encode a C-terminal DBL domain. This further reinforced their candidature, as we recently mapped Fc receptor binding of IgM to a domain of that type (15). Indeed, we experimentally confirmed their ability to bind IgM (Fig. 2).

The Fc-specific binding of IgM to PfEMP1 proteins has consistently been shown to involve the Cµ3–Cµ4 domains in the Fc part of pentameric IgM. Our findings confirm this location of the PfEMP1-binding site in IgM (Fig. 3). More ambiguity exists with respect to the corresponding PfEMP1 domains involved. Thus far, at least three of the DBL domains in VAR2CSA-type PfEMP1 (DBL0AM2_2, DBL0AM5_6, and DBL0AM10_7) have been implicated (39, 40). We resolved this uncertainty by using recombinant constructs of each of the DBL domains in PFL0030c to map the IgM-binding site to the penultimate N-terminal domain DBL0AM5_6 (Fig. 4). Similar confusion has existed with respect to IgM binding to non-VAR2CSA-type PfEMP1 proteins, in which N-terminal cysteine-rich interdomain region α (CIDRα), central DBLβ, and C-terminal DBLγ domains have all been implicated (15, 42, 43). We found that DBLγ domains near the C terminus were responsible for mediating binding to IgM in the PfEMP1 proteins studied here. One of them (MAL6P1.4) even contained two IgM-binding domains (DBL0AM2_7 and DBL0AM3_9), something which has not been observed previously. Overall, presently available evidence points to DBLγ and DBLδ domains near or at the C terminus as the key IgM-binding elements in PfEMP1 proteins.

The functional significance of Fc-mediated binding of IgM to PfEMP1 proteins remains unclear (18). In the case of VAR2CSA-type PfEMP1, it appears to serve an immunoevasive role that interferes with phagocytosis of IgG-opsonized IEs without compromising the adhesive function of PfEMP1 (19). Our findings here support these earlier results. Thus, PFL0030c-expressing IEs adhered strongly to CSA, whether IgM was present or not (Fig. 5A), and IgM markedly inhibited phagocytosis of IgG-opsonized IEs expressing this PfEMP1 (Fig. 6). In contrast to these characteristics of VAR2CSA-type PfEMP1, we recently reported that Fc-dependent binding of IgM does not protect parasites expressing the rosette-mediating HB3VAR06 protein from phagocytosis of IgG-opsonized IEs but rather may serve to augment low-affinity adhesive interactions between the PfEMP1 head structure and as yet undefined carbohydrate moieties (15). Fc-specific binding of IgM to the new PfEMP1 proteins identified here did not markedly protect them from phagocytosis following opsonization by antigen-specific IgG (Fig. 6). In this respect, they resemble the rosette-mediating PfEMP1 protein HB3VAR06, which suggests that they have an elongated conformation. Only one of them (MAL6P1.316) contains an N-terminal DBL domain of a type previously associated with rosetting (DBLβ1.5, DBLβ1.6, DBLβ1.8, or DBLβ2) (10–13, 15), and none of them mediated rosetting (Fig. 5B). These findings establish for the first time that the ability to bind IgM via Fcµ is not restricted to VAR2CSA-type and rosette-mediating PfEMP1 variants. They also indirectly support our hypothesis that an important function of IgM binding is to augment low-affinity adhesive interactions between the PfEMP1 head structures and host endothelial receptors (15). Only when such receptors are also found on erythrocytes would this lead to rosetting.

In conclusion, we have provided evidence that each P. falciparum genome encodes several PfEMP1 proteins with the capacity to bind Fcµ. We cannot formally rule out that other parasite-encoded IE proteins, e.g., rosette-mediating RIFINs (58, 59), also have affinity for Fcµ, but there is presently no evidence to support that possibility. We identified five PfEMP1 proteins in P. falciparum 3D7/NF54 parasites, but this is almost certainly an underestimation considering the limited number of subclones we investigated. Our findings confirm that Fc-specific binding of IgM to VAR2CSA-type PfEMP1 has an immunoevasive function. However, they also support recent data pointing to alternative roles for Fc-mediated IgM binding to other types of PfEMP1 (15, 18, 20). In any case, the IgM-binding phenotype is likely to be both common and important, as we showed that Fc-dependent binding of
IgM is not restricted to PfEMP1 proteins mediating either adherence to CSA or rosetting. Although rosetting has been associated with expression of PfEMP1 proteins causing severe malaria, the relationship is not absolute (reviewed in reference 60). Indeed, several PfEMP1 proteins that mediate adhesion to EPCR and ICAM-1, phenotypes that have repeatedly been associated with severe malaria, do not mediate formation of rosettes (50). Future analysis of the interrelationship among various PfEMP1 phenotypes that have been individually associated with malaria severity may well produce important new insights. This prediction is supported by the fact that one of the PfEMP1 proteins studied here (MAL8P1.316), which bound Fcμ but did not mediate formation of rosettes, contains a CIDR of a type (CIDRA1.8) that is involved in adhesion to EPCR (51).

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