Biochemical Analysis of the *Plasmodium falciparum* Erythrocyte-binding Antigen-175 (EBA175)-Glycophorin-A Interaction

**IMPLICATIONS FOR VACCINE DESIGN**

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**Background:** The GYPA-PfEBA175 interaction is important for erythrocyte invasion by the malaria parasite.

**Results:** The entire ectodomain of EBA175 interacted with GYPA with different biochemical parameters to the previously determined GYPA-binding fragment containing two DBL domains.

**Conclusion:** Regions outside of the tandem DBL domains contribute to GYPA binding by EBA175.

**Significance:** These findings may assist the design of an EBA175-based malaria vaccine.

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*Pf*EBA175 has an important role in the invasion of human erythrocytes by *Plasmodium falciparum* and is therefore considered a high priority blood-stage malaria vaccine candidate. *Pf*EBA175 mediates adhesion to erythrocytes through binding of the Duffy-binding-like (DBL) domains in its extracellular domain to Neu5Acα2–3Gal displayed on the O-linked glycans of glycophorin-A (GYPA). Because of the difficulties in expressing active full-length (FL) *P. falciparum* proteins in a recombinant form, previous analyses of the *Pf*EBA175-GYPA interaction have largely focused on the DBL domains alone, and therefore they have not been performed in the context of the native protein sequence. Here, we express the entire ectodomain of *Pf*EBA175 (*Pf*EBA175 FL) in soluble form, allowing us to compare the biochemical and immunological properties with a fragment containing only the tandem DBL domains (“region II,” *Pf*EBA175 RII). Recombinant *Pf*EBA175 FL bound human erythrocytes in a trypsin and neuraminidase-sensitive manner and recognized Neu5Acα2–3Gal-containing glycans, confirming its biochemical activity. A quantitative binding analysis showed that *Pf*EBA175 FL interacted with native GYPA with a *K* *D* ~0.26 μM and is capable of self-association. By comparison, the RII fragment alone bound GYPA with a lower affinity demonstrating that regions outside of the DBL domains are important for interactions with GYPA; antibodies directed to these other regions also contributed to the inhibition of parasite invasion. These data demonstrate the importance of *Pf*EBA175 regions other than the DBL domains in the interaction with GYPA and merit their inclusion in an EBA175-based vaccine.

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The human malaria parasite, *Plasmodium falciparum*, causes an estimated 300–500 million clinical cases and up to 800,000 deaths each year (1, 2). Widespread implementation of control measures within the last decade, including artemisinin combination therapy and the use of insecticide-treated bed nets, has resulted in a significant decrease in the incidence of *P. falciparum* malaria in endemic countries (3–5). In the face of emerging resistance to artemisinin in parasites and pyrethroid insecticides in mosquito vectors, however, the need for an effective vaccine for long term control and prevention of malaria remains an important global health objective (3, 4, 6). Vaccines that target the blood stage of the infection are conceptually attractive because the parasite is directly exposed to the host humoral immune system, and it is this stage of the life cycle that is responsible for all the clinical symptoms of malaria (7). The blood stage is initiated when an extracellular form of the parasite called the merozoite recognizes and invades host erythrocytes. Invasion is a complex process involving multiple interactions between host erythrocyte receptors and parasite ligands displayed on the merozoite surface. *Pf*EBA175 was the first *P. falciparum* invasion ligand identified and interacts with the highly abundant erythrocyte surface sialoglycoprotein, glycophorin-A (GYPA) (8–10). *Pf*EBA175 is considered a leading vaccine candidate because antibodies directed against *Pf*EBA175 are present in malaria-immune individuals, and antibodies raised against recombinantly expressed fragments of *Pf*EBA175 inhibit erythrocyte invasion by *P. falciparum in vitro* (11–16). EBA175 is a member of the erythrocyte-binding-like family of *Plasmodium* proteins, which include the Duffy-binding proteins of *Plasmodium vivax* and *Plasmodium knowlesi* (*Pv*DBP and *Pkd*DBP) and the *P. falciparum* paralogs EBA140, EBA181, and EBL1. The members of the erythrocyte-binding-
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like family share a similar gene structure, and this homology has been used to define six regions, RI–RV1, in their ectodomains (Fig. 1A) (17). A functional analysis of these regions showed that the RII fragment is capable of rosetting erythrocytes (10). The structure of RII, which comprises two tandem DBL domains (F1 and F2), has been determined by x-ray crystallography (18).

The PfEBA175 receptor, GYPA, is a dimeric type I transmembrane protein carrying 15 closely clustered O-linked tetrasaccharides capped with sialic acid/N-acetyleneuraminic acid (Neu5Ac) (19, 20). The “Neu5Acα2–3Gal” sequence of these glycans was shown to be essential for the recognition of GYPA by PfEBA175 (21). Co-crystallization of the RII fragment with a structural analog of this disaccharide, α-2,3-sialylactose, revealed six putative glycan-binding sites at the dimer interface of the parasite protein, with four of the sites located within two channels that span the dimer and another two in a deep groove accessible only through a cavity at the top of the dimer (18). Based on the locations of the glycan-binding sites in the RII fragment, and the predominance of the monomeric form of PfEBA175 RII in solution, Tolia and co-workers (22) have proposed a “receptor-induced dimerization model” for the erythrocyte binding of PfEBA175, which postulates monomeric PfEBA175 assembling into a dimer around the dimeric extracellular region of GYPA during invasion. This model is also consistent with recent structural studies investigating the mode of binding of the PvDBP RII to its sulfotyrosine-carrying receptor, DARC.

Although studies of the 68-kDa recombinant PfEBA175 RII have proven highly informative, native PfEBA175 is a much larger protein, being synthesized as a 190-kDa membrane-tethered precursor that is proteolytically cleaved during erythrocyte invasion to release the 175-kDa extracellular region (9). Because of the technical difficulties associated with expressing full-length *Plasmodium* proteins in a functionally active soluble recombinant form (23), much of the biochemical characterization of the PfEBA175-GYPA interaction has been performed using just the RII fragment.

In the study reported here, we expressed the full-length ectodomain of PfEBA175 (PfEBA175 FL) in a soluble recombinant form using a mammalian expression system. We confirmed that the recombinant protein is biochemically active and exhibited binding properties similar to those of native PfEBA175 isolated from parasite cultures. Using this protein, we investigated the biophysical parameters of the EBA175-GYP A interaction and compared them with those of the RII fragment to demonstrate that regions outside the DBL domains contributed to GYPA binding. These data have important implications for the rational design of an effective PfEBA175-based malaria vaccine.

**EXPERIMENTAL PROCEDURES**

*Recombinant Expression and Purification of Proteins*—The sequence encoding the entire extracellular domain of *P. falciparum* (3D7) EBA175 (PfEBA175 FL) except the signal peptide (amino acids 21–1424) was made by gene synthesis (GeneART). The codons were optimized for expression in human cells, and the potential N-linked glycosylation sites were removed as described (24). The sequence coding for RII of PfEBA175 (amino acids 142–764) was amplified from this construct by PCR. The coding sequences were cloned into pTT3-derived vectors using unique flanking NotI and Ascl restriction enzyme sites, between the leader sequence of the mouse variable κ light chain 7–33 (25) and a rat Cd4 (Ig-like domains 3 and 4) tag, followed either by an enzymatically biotinylatable peptide tag, the pentamerization domain of the rat cartilage oligomeric matrix protein (COMP), and ampicillin resistance protein β-lactamase, or a hexa-His tag (24, 26). The proteins were expressed by transient transfection of HEK293 cells grown in suspension culture as described (27, 28) and collected from the cell culture supernatant 6 days post-transfection. Biotinylation of proteins was achieved by co-transfection with a secreted form of the *Escherichia coli* biotin ligase, BirA (27), and excess unconjugated Ϝ-biotin was removed by extensive dialysis into HBS. His-tagged proteins were purified from the culture supernatants by affinity chromatography on HisTrap HP columns (GE Healthcare) using an AKTAxpress (GE Healthcare) according to the manufacturer’s instructions. Size exclusion chromatography (SEC) of nickel-purified proteins was carried out on a Superdex 200 Tricorn 10/600 column (GE Healthcare) in HBS-EP (HBS, 3 mM EDTA, 0.005% w/v Surfactant P20 (GE Healthcare)).

Western Blotting—Proteins were resolved under reducing conditions, blotted, and detected using horseradish peroxidase (HRP)-conjugated extravidin (Sigma) essentially as described (24).

Enzyme-linked Immunosorbent Assays (ELISA)—Biotinylated proteins were detected by ELISA essentially as described (24). Commercially available glycophorin-A preparations (catalog numbers: G7903 and A9791, Sigma) were biotinylated *in vitro* by incubation with a 20-fold molar excess of EZ-link sulfo-NHS-biotin (Thermo Scientific) for 30 min at room temperature and dialyzed into HBS prior to its use in the assays. When testing for immunoreactivity, proteins were immobilized with or without prior treatment at 80°C for 10 min.

Erythrocyte Binding Assays—Biotinylated PfEBA175 and Cd4 (negative control) were multimerized by immobilization on streptavidin-coated Nile Red fluorescent 0.4–0.6-μm microbeads (Spherotech Inc.) by incubation for 45 min at 4°C (29). The bead arrays were then presented to erythrocytes in flat-bottom 96-well microtiter plates at a density of ~3 × 10^6 cells/well (the ratio of cells/beads = 1:200). After incubation for 1 h at 4°C with the protein arrays, the cells were washed twice in HBS + 1% BSA (HBS/BSA) and analyzed by flow cytometry. The data were acquired on an LSRII cytometer (BD Biosciences) using the FACS Diva software (BD Biosciences). Nile Red was excited by a blue laser and detected with a 575/26 filter. Forward scatter and side scatter voltages of 430 and 300 V, respectively, and a threshold of 26,100 on forward scatter were applied when analyzing erythrocytes. To test for binding specificity, the erythrocytes were pretreated with either tosylsulfonfylphenylalanyl chloromethyl ketone-treated trypsin from bovine pancreas (Sigma) (at 0.25, 0.5, and 1 mg/ml), tosyl-lysyl chloromethyl ketone-treated trypsin from bovine pancreas (Sigma) (at 0.25, 0.5 and 1 mg/ml), or 0.1 milliunit/10^6 cells of *Vibrio cholerae* neuraminidase (Sigma), for 1 h at 37°C. Trypsin- and chymotrypsin-treated cells were washed once, treated with 0.5 mg/ml soybean trypsin-chymotrypsin inhibitor (Sigma) for 10 min at room temperature, and then washed twice more before incubation with PfEBA175-coated beads.
Neuraminidase-treated cells were washed twice prior to use in the binding assays.

**Modified Affinity-based Extracellular Interaction Screening (AVEXIS) Assays**—The ELISA-based AVEXIS methodology, as described previously (27), was adapted for detecting the interactions of recombinant PfEBA175 proteins, expressed as β-lactamase-tagged pentameric “preys” with biotinylated ‘bait’ forms of purified GYPA and synthetic carbohydrate probes (GlycoTech). Briefly, the baits and the preys were normalized to activities that have previously been shown to detect transient interactions (monomeric half-lives less than 0.1 s). The biotinylated baits were immobilized on streptavidin-coated 96-well microtiter plates (NUNC) at concentrations sufficient for the complete saturation of the available binding surface/well. The plates were then washed twice in HBST and blocked with HBS/BSA for 0.5–1 h, before addition of normalized β-lactamase-tagged prey proteins and incubation for 2 h. After washes in HBST and HBS, 125 μg/ml nitrocefin, a β-lactamase substrate, was added to the wells, and its hydrolysis was monitored by absorbance measurements at 485 nm on a Pherastar plus (BMG Laboratories). All steps were performed at room temperature.

**Lectin Binding Assay with Purified GYPA**—Biotinylated lectins (Vector Laboratories) were immobilized on streptavidin-coated 96-well microtiter plates (NUNC) at 10 μg/ml for 1 h. The plates were then washed twice in HBST and blocked with HBS/BSA for 30 min. The immobilized lectins were next incubated with 0.02 mg/ml purified GYPA (Sigma) for 2 h. After washes in HBST, the plates were incubated with 1 μg/ml of the anti-GYPA mouse monoclonal antibody BRIC256 (Abcam) for 1 h, followed by an alkaline phosphatase-conjugated antimouse secondary antibody (Sigma) for 30 min, before the addition of p-nitrophenyl phosphate (Sigma) at 1 mg/ml and measuring the absorbance at 405 nm on a Pherastar Plus (BMG Laboratories). All steps were performed at room temperature.

**Surface Plasmon Resonance (SPR)**—SPR studies were performed on a BIAcore T100 instrument (GE Healthcare) at 37 °C, using HBS-EP (GE Healthcare) as the running buffer. In each experiment, biotinylated baits were immobilized on streptavidin-coated sensor chips (GE Healthcare), with the negative control bait in the “reference” flow cell and an approximate molar equivalent amount of the “query” baits in the other flow cells. Purified analyte proteins were separated by size exclusion chromatography on a Superdex 200 Tricorn 10/600 column just before their use in the SPR experiments. Increasing concentrations of these proteins were injected over the immobilized baits at 20 μl/min for equilibrium measurements and at 100 μl/min for kinetic measurements. The surfaces were regenerated with a pulse of 5 M NaCl at the end of each injection cycle. Duplicate injections of the same concentration in each experiment were superimposable, demonstrating no loss of activity after surface regeneration. Reference-subtracted sensorgrams were analyzed using the BIAcore evaluation software version 1.1.1 (GE Healthcare). To determine the overall equilibrium binding affinity, binding responses in the steady-state region of the sensorgrams (R_{eq}) were plotted against analyte concentration (C) and fitted to the following equation: R_{eq} = CR_{max}/(C + K_D), where R_{max} is the maximum binding response, and K_D is the equilibrium dissociation constant. Kinetic constants were calculated by nonlinear regression fitting to the association and dissociation phases of the sensorgrams. To identify the mechanism of binding, the sensorgrams were globally fitted to three predefined interaction models as follows: simple 1:1 binding (A + B ↔ AB, where A is the soluble analyte and B is the immobilized ligand); conformational change (A + B ↔ AB ↔ AB*); and bivalent analyte (AA + B ↔ AAB; AAB + B ↔ AABB).

**Multiangle Light Scattering Measurements (MALS)**—Size exclusion chromatography was performed on Superose 6 10/30 (PfEBA175 FL) and Superdex 200 10/30 (PfEBA175 RII) columns (GE Healthcare) equilibrated in HBS (GE Healthcare) at 0.4 ml/min. The column was followed in line by a Dawn Heleos-II light scattering detector (Wyatt Technologies) and an Optilab-Rex refractive index monitor (Wyatt Technologies). Molecular mass calculations were performed using ASTRA 5.3.4.14 (Wyatt Technologies) assuming a dn/dc value of 0.186 ml/g.

**P. falciparum Culture and Invasion Assays**—The 3D7 and Dd2 strains of P. falciparum were cultured in human O+ erythrocytes at 5% hematocrit in complete medium (RPMI 1640 medium containing 10% human serum), under an atmosphere of 1% O_2, 3% CO_2, and 96% N_2. Invasion assays were performed as described previously (30).

**Polyclonal Antibodies**—To raise polyclonal sera against PfEBA175 FL and RII, the purified proteins were injected into rabbits (Cambridge Research Biochemicals,Billingham, UK). The sera were purified on HiTrap Protein G HP columns (GE Healthcare) using an ÄKTAxpress (GE Healthcare) and dialed into RPMI 1640 medium (Invitrogen) prior to their use. The anti-PfEBA175 FL and anti-PfEBA175 RII antibodies had similar anti-Cd4 activity, as determined by ELISA. An anti-AMA1 polyclonal (31) was used in the invasion assays as a control.

**RESULTS**

**Expression of a Soluble, Biochemically Active Full-length Ectodomain of PfEBA175 (PfEBA175 FL)**—The technical challenges associated with expressing full-length Plasmodium proteins in a recombinant, biochemically active form (23) have prevented a detailed biochemical investigation of the EBA175-GYPA interaction, with most studies limited to a fragment of EBA175 encompassing the two tandem DBL domains (RII). To investigate the biochemical properties and functional activity of a more physiologically relevant PfEBA175 protein, we took advantage of a strategy utilizing a mammalian expression system and codon-optimized gene sequences that had previously been successful for producing biochemically active Plasmodium proteins (24, 26). Using this method, we expressed the entire full-length ectodomain fragment of PfEBA175 (PfEBA175 FL) from the 3D7 isolate of P. falciparum as a C-terminally tagged soluble fusion protein (Fig. 1A). Western blotting of unpurified culture supernatants confirmed the presence of a protein at the expected size (Fig. 1B). To determine whether the recombinant PfEBA175 FL was correctly folded and biochemically active, its immunoreactivity to two mouse monoclonal antibodies, R217 and R218, raised against the RII fragment of EBA175 expressed in SF21 insect cells and known to bind to nonlinear heat-labile epitopes within
immunoreactivity was heat-labile suggesting that at least the 
PfEBA175 RII, three anti-EBA175 monoclonal antibodies to untreated and heat-treated 
PfEBA175 FL, were tested (32). Immunoreactivity to PfEBA175 FL was observed for both R217 and R218. Importantly, this 
immunoreactivity was heat-labile suggesting that at least the 
epitopes recognized by R217 and R218 are correctly folded in the recombinant protein (Fig. 1C). We were also able to de-
monstrate immunoreactivity to other parts of the EBA175 protein using a mouse monoclonal antibody raised against a 
yeast-expressed PfEBA175 RVI fragment (33). This antibody, however, showed similar binding to both untreated and heat-
treated samples of PfEBA175 FL, suggesting that it recognizes a nonheat-labile linear epitope (Fig. 1C).

PfEBA175 FL Is Primarily Monomeric with a Small Amount of Self-association—The crystal structure of RII has shown that 
EBA175 may function by dimerization (18), and so to assess the 
oligomeric state of PfEBA175 FL, the purified protein was ana-
lyzed by SEC. The elution profile of PfEBA175 consisted of two 
peaks, both containing the protein of interest as demonstrated by 
denaturing SDS-PAGE. The smaller peak eluted in the void 
volume of the column, and the size of the major peak was esti-
mated to be ~500 kDa. A similar mass for PfEBA175 FL was 
observed by native PAGE, consistent with it being primarily 

domimeric in solution (Fig. 1D). To further investigate the 
oligomeric state of PfEBA175 FL, it was subjected to MALS 
immEDIATELY FOLLOWING SEC. This analysis revealed that the 
main peak of PfEBA175 FL is predominantly monomeric with a 
certain degree of self-association (Fig. 1E). The early elution of 
PfEBA175 FL in SEC and its retarded mobility in native PAGE 
may therefore be due to a large hydrodynamic shape, possibly 
due to a high degree of conformational flexibility or an elon-
gated structure.

PfEBA175 FL Binds Human Erythrocytes in a Neuraminidase-
and Trypsin-sensitive but Chymotrypsin-resistant Manner—
To determine whether recombinant PfEBA175 FL is biochem-
ically active, we first asked whether it could bind human 
erthrocytes in a manner consistent with its recognition of 
GYP. Interactions between cell surface proteins typically have 
low binding affinities, and multimerized, highly avid binding 
reagents are often required to facilitate their detection (34). 
Therefore, to estimate the degree to which the PfEBA175 FL 
protein could associate with human erythrocyte surfaces, we 
first created a highly avid PfEBA175 FL binding reagent by clus-
tering the biotin-tagged monomeric EBA175 protein around 
fluorescent streptavidin-coated beads. The EBA175-coated 
beads were then presented to human erythrocytes, and the 
extent to which they bound was quantified by fluorescence-
activated cell sorting. We observed that PfEBA175 FL-coated 
beads were clearly able to bind to human erythrocytes at a 

dose-dependent manner (Fig. 2A), whereas a similar treatment with 
chymotrypsin had no significant effect, even at the highest con-
centration (Fig. 2B). Pretreatment of the erythrocytes with V. 
cholerae neuraminidase, which preferentially cleaves α2–3-
linked sialic acids of O-linked tetrasaccharides, was sufficient to 
prevent all EBA175 binding (Fig. 2C).
PfEBA175 FL binds human native GYPA and Neu5Acα2–3Gal-containing glycans—To determine whether recombinant PfEBA175 FL bound native GYPA directly, we used a preparation of GYPA extracted from human erythrocytes. First, we characterized the glycan profile of the native GYPA preparation using three lectins with known carbohydrate-recognition specificities, which revealed a predominance of Neu5Acα2–3Gal with a much smaller amount of Neu5Acα2–6-linked glycans (Fig. 2D). An asialylated form of GYPA included as a control appeared to mainly carry Galβ1–3GalNAc (T antigen), as expected. To detect the direct binding of PfEBA175 FL to native GYPA, we chemically biotinylated GYPA and captured it.
on streptavidin-coated plates before incubating this with PfEBA175 FL expressed as a highly avid β-lactamase-tagged pentamer (27). Using this approach, pentamerized PfEBA175 FL showed saturable binding to GYPA, but not to its asialylated derivative demonstrating that the recombinant PfEBA175 anti- gen associates directly with GYPA in a “Neu5Aca2–3Gal”-dependent manner (Fig. 2E).

The expression of the entire ectodomains of EBA175 as a highly avid recombinant protein enabled us to systematically assess the glycan binding properties of EBA175 by screening a large panel of 54 synthetic carbohydrates, primarily selected because they or their close derivates are known to be present at the surface of human erythrocytes (Fig. 2, F and G, and Table 1). All oligosaccharides containing the Neu5Aca2–3Gal determinant were recognized by PfEBA175 FL (Fig. 2G). Previous work has suggested that the glycan binding properties of PfEBA175 may be responsible for the restriction of P. falciparum to humans because it has been reported that PfEBA175 is unable to bind Neu5Gc (N-glycolylneuraminic acid) (35), the hydroxylated form of Neu5Ac that is absent from human erythrocytes but present in other great apes (36). Interestingly, we observed clear binding of PfEBA175 FL to the Neu5Gc monosaccharide in the screen, and this was subsequently confirmed by SPR (Fig. 2G). Together, these data demonstrate that the recombinant soluble protein consisting of the entire ectodomain of PfEBA175 is biochemically active and can directly bind native GYPA in a Neu5Aca2–3Gal-dependent manner.

PfEBA175 FL and GYPA Directly Interact with a K_d of ~0.26 μM—The ability to express and purify PfEBA175 FL enabled us to investigate its interaction with native GYPA in detail. We first determined the biophysical properties of the PfEBA175 FL-GYPA interaction using SPR as implemented in a BIACore instrument. Affinity purified PfEBA175 FL was first resolved by SEC, and fractions encompassing the main peak were pooled and serial dilutions injected over GYPA immobilized on a sensor chip. Binding of PfEBA175 FL to native GYPA was observed and quantified once equilibrium had been reached to derive an equilibrium dissociation constant (K_d) of ~0.26 μM (Fig. 3A). Although this interaction is relatively weak as expected, it is still ~4-fold stronger than the other two P. falciparum ligand-erythrocyte receptor interactions characterized in our laboratory, PFRH5-Basigin and PFMTRAP-Semaphorin 7A, each of which has a K_d of ~1 μM (24, 26).

Kinetic Analysis of the PfEBA175 FL-GYPA Interaction—Previous research using fragments of the PfEBA175 protein has hypothesized that EBA175 interacts with GYPA in a two-stage process, involving receptor-induced dimerization (18) and a conformational change in the EBA175 protein (37, 38). To characterize the mechanistic details of the interaction between GYPA and the entire PfEBA175 ectodomain, we performed a kinetic analysis using PfEBA175 FL. The association and disso-
Biochemical Analysis of the PfEBA175-GYPA Interaction

FIGURE 3. PfEBA175 FL binds sialylated GYPA with a $K_D$ of 0.26 μM. A, SPR analysis of the binding of PfEBA175 FL to GYPA. 2-Fold dilutions of purified PfEBA175 FL were injected over biotinylated GYPA immobilized on a streptavidin-coated sensor chip until equilibrium had been reached (shown in the inset). Reference-subtracted binding data were plotted as a binding curve, and the $K_D$ was calculated by global fitting to a steady-state 1:1 interaction model. $K_D$ is shown as mean ± S.E., based on two independent experiments. Biotinylated Cd4 tag was used as the reference. B, fits of a PfEBA175 FL-GYPA sensorgram to three predefined kinetic models. An experimental curve is shown overlaid with the fitted curves from the 1:1 binding ($\chi^2 = 7.98$), bivalent analyte ($\chi^2 = 0.642$), and conformational change ($\chi^2 = 0.763$) models. C, reference-subtracted sensorgrams obtained by injecting 0.33 μM of PfEBA175 FL over GYPA for various lengths of time (45–115 s) showed no alteration in the dissociation rate suggesting no detectable conformational changes of PfEBA175 upon binding. D, serial dilutions of soluble PfEBA175 FL were injected over biotinylated PfEBA175 FL immobilized on an SPR sensor chip and shown to interact. The dissociation phase data from the reference-subtracted sensorgrams is shown as mean ± S.D., $n = 3$; for clarity, only every 40th data point is shown.

PfEBA175 interacts with GYPA with a 10-Fold Lower Affinity than PfEBA175 FL—The fragment of EBA175 containing the tandem DBL domains (RII) is known to bind GYPA (10); however, whether other regions of EBA175 also contribute to erythrocyte recognition is not well established (12, 37, 38). To compare the binding properties of the RII fragment with the full-length PfEBA175, we also expressed RII as a soluble, recombinant form using HEK293 cells. A biochemical characterization of the PfEBA175 RII protein revealed that it was expressed at the expected size, was recognized by the conformation-sensitive monoclonal antibodies R217 and R218, and bound human erythrocytes in a neuraminidase-sensitive manner suggesting that it is correctly folded (Fig. 4, A–C). PfEBA175 RII eluted as a monodisperse peak at ~125 kDa when analyzed by SEC consistent with it adopting a primarily monomeric form in solution (Fig. 4D). Further analysis by MALS confirmed that PfEBA175 RII, similar to PfEBA175 FL, was primarily monomeric, but it showed some propensity to self-associate (Fig. 4E).

We used a pentamerized enzyme-tagged PfEBA175 RII protein to show that it could bind GYPA, although in comparison to interact. The dissociation phase data from the reference-subtracted sensorgrams is shown as mean ± S.D., $n = 3$; for clarity, only every 40th data point is shown.

ciation kinetic rate constants, $k_a$ and $k_d$, were determined using nonlinear curve fitting to a set of reference-subtracted sensorgrams using the initial binding and wash-out phases. Both the $k_a$ (8.6 ± 0.2 × 10^5 M$^{-1}$ s$^{-1}$) and $k_d$ (0.09682 ± 0.00008 s$^{-1}$) values were typical for a relatively weak protein-protein interaction. To gain a better mechanistic understanding of the interaction, the binding data were fitted to three models (Fig. 3B). The fits to the conformational change and bivalent analyte models were similar and better than that of the simple 1:1 binding model (Fig. 3B). To test whether a conformational change was involved in the binding of PfEBA175 FL to GYPA, we injected PfEBA175 FL until saturation was achieved for a range of different contact times. We observed that variations in contact time did not influence the dissociation phase, suggesting that the interaction of PfEBA175 FL with GYPA does not involve a slow, temporally resolvable conformational change that stabilizes the complex (Fig. 3C). Using SEC-MALS, we established that PfEBA175 FL was primarily monomeric in solution albeit with some self-association. To test the relevance of the bivalent analyte model for the PfEBA175-GYPA interaction, we analyzed the propensity of PfEBA175 FL to interact with itself by SPR. We quantified the observed homophilic binding using the dissociation phase of the interaction to avoid the confounding problem of analyte self-association, which would lead to an underestimate of the affinity in equilibrium analyses (Fig. 3D). PfEBA175 FL self-associated with a $k_d$ value (~0.04 s$^{-1}$), consistent with a role for EBA175 dimerization in its interaction with GYPA.

PfEBA175 RII Interacts with GYPA with a 10-Fold Lower Affinity than PfEBA175 FL—The fragment of EBA175 containing the tandem DBL domains (RII) is known to bind GYPA (10); however, whether other regions of EBA175 also contribute to erythrocyte recognition is not well established (12, 37, 38). To compare the binding properties of the RII fragment with the full-length PfEBA175, we also expressed RII as a soluble, recombinant form using HEK293 cells. A biochemical characterization of the PfEBA175 RII protein revealed that it was expressed at the expected size, was recognized by the conformation-sensitive monoclonal antibodies R217 and R218, and bound human erythrocytes in a neuraminidase-sensitive manner suggesting that it is correctly folded (Fig. 4, A–C). PfEBA175 RII eluted as a monodisperse peak at ~125 kDa when analyzed by SEC consistent with it adopting a primarily monomeric form in solution (Fig. 4D). Further analysis by MALS confirmed that PfEBA175 RII, similar to PfEBA175 FL, was primarily monomeric, but it showed some propensity to self-associate (Fig. 4E).

We used a pentamerized enzyme-tagged PfEBA175 RII protein to show that it could bind GYPA, although in comparison to interact. The dissociation phase data from the reference-subtracted sensorgrams is shown as mean ± S.D., $n = 3$; for clarity, only every 40th data point is shown.

ciation kinetic rate constants, $k_a$ and $k_d$, were determined using nonlinear curve fitting to a set of reference-subtracted sensorgrams using the initial binding and wash-out phases. Both the $k_a$ (8.6 ± 0.2 × 10^5 M$^{-1}$ s$^{-1}$) and $k_d$ (0.09682 ± 0.00008 s$^{-1}$) values were typical for a relatively weak protein-protein interaction. To gain a better mechanistic understanding of the interaction, the binding data were fitted to three models (Fig. 3B). The fits to the conformational change and bivalent analyte models were similar and better than that of the simple 1:1 binding model (Fig. 3B). To test whether a conformational change was involved in the binding of PfEBA175 FL to GYPA, we injected PfEBA175 FL until saturation was achieved for a range of different contact times. We observed that variations in contact time did not influence the dissociation phase, suggesting that the interaction of PfEBA175 FL with GYPA does not involve a slow, temporally resolvable conformational change that stabilizes the complex (Fig. 3C). Using SEC-MALS, we established that PfEBA175 FL was primarily monomeric in solution albeit with some self-association. To test the relevance of the bivalent analyte model for the PfEBA175-GYPA interaction, we analyzed the propensity of PfEBA175 FL to interact with itself by SPR. We quantified the observed homophilic binding using the dissociation phase of the interaction to avoid the confounding problem of analyte self-association, which would lead to an underestimate of the affinity in equilibrium analyses (Fig. 3D). PfEBA175 FL self-associated with a $k_d$ value (~0.04 s$^{-1}$), consistent with a role for EBA175 dimerization in its interaction with GYPA.

PfEBA175 RII Interacts with GYPA with a 10-Fold Lower Affinity than PfEBA175 FL—The fragment of EBA175 containing the tandem DBL domains (RII) is known to bind GYPA (10); however, whether other regions of EBA175 also contribute to erythrocyte recognition is not well established (12, 37, 38). To compare the binding properties of the RII fragment with the full-length PfEBA175, we also expressed RII as a soluble, recombinant form using HEK293 cells. A biochemical characterization of the PfEBA175 RII protein revealed that it was expressed at the expected size, was recognized by the conformation-sensitive monoclonal antibodies R217 and R218, and bound human erythrocytes in a neuraminidase-sensitive manner suggesting that it is correctly folded (Fig. 4, A–C). PfEBA175 RII eluted as a monodisperse peak at ~125 kDa when analyzed by SEC consistent with it adopting a primarily monomeric form in solution (Fig. 4D). Further analysis by MALS confirmed that PfEBA175 RII, similar to PfEBA175 FL, was primarily monomeric, but it showed some propensity to self-associate (Fig. 4E).

We used a pentamerized enzyme-tagged PfEBA175 RII protein to show that it could bind GYPA, although in comparison to interact. The dissociation phase data from the reference-subtracted sensorgrams is shown as mean ± S.D., $n = 3$; for clarity, only every 40th data point is shown.
Biochemical Analysis of the PfEBA175-GYPA Interaction

FIGURE 4. Biochemically active recombinant PfEBA175 RII is largely monomeric in solution. A, Western blot of unpurified cell culture supernatant containing biotinylated PfEBA175 RII. B, ELISA demonstrating immunoreactivity of two anti-EBA175 RII mouse monoclonal antibodies to untreated and heat-treated PfEBA175 RII and untreated PfEBA175 FL. C, median fluorescence intensities (MFI) associated with untreated and neuraminidase-treated erythrocytes incubated with fluorescent beads coated with Cad4, PfEBA175 FL, or PfEBA175 RII. D and C, bar charts show mean ± S.D., n = 3. D, SEC profile of purified His-tagged PfEBA175 RII and a Coomassie G-250-stained denaturing gel of the peak fraction (inset). E, multicycle light scattering of purified PfEBA175 RII (blue line) and a control monomeric protein of 30 kDa used as a reference (black line). The peaks correspond to SEC elution with the absorbance at 280 nm on the right y axis. The horizontal lines indicate the molecular mass (left y axis).

with the normalized PfEBA175 FL control, the interaction detected using this fragment was more sensitive to dilution of the protein, suggesting it had a lower affinity for GYPA (Fig. 5A). Using SPR, we quantified this binding and found that PfEBA175 RII bound GYPA with a $K_d$ of $\sim 2 \mu M$, 10-fold weaker in comparison with the PfEBA175 FL-GYPA interaction (Fig. 5B). Consistent with this, a comparative kinetic analysis showed that the weaker interaction strength of PfEBA175 RII was due to a faster dissociation rate, ruling out the possibility that a significant fraction of the PfEBA175 RII protein was functionally inactive (Fig. 5C). One reason for the weaker binding of the RII fragment to GYPA could be due to a reduced ability to self-associate. To examine this, we compared the binding affinity of the RII fragment and the full-length ectodomain to PfEBA175 FL and found that the former interacted with an $\sim 70$-fold weaker binding affinity (Fig. 5D). It has been previously shown that both the GYPA peptide backbone and sialic acid are required for EBA175 binding (10). To determine whether these binding properties could be distinguished and/or attributed to different regions of EBA175 protein, we directly compared the binding of both PfEBA175 FL and PfEBA175 RII analytes to GYPA and the oligosaccharide Neu5Acα2–3Gal. Binding of PfEBA175 RII to GYPA and Neu5Acα2–3Gal was indistinguishable from one another, whereas PfEBA175 FL bound GYPA with $\sim 2$-fold higher affinity than for the glycan alone (Fig. 5E). These results suggest that PfEBA175 RII interacts with the glycan moieties of GYPA but that the extracellular regions of EBA175 outside of RII also contact the polypeptide backbone of GYPA, contributing further binding energy to the interaction.

Comparison of PfEBA175 FL and PfEBA175 RII as Vaccine Candidates—Given the important role of EBA175 in erythrocyte invasion and the early success in expressing region II as an active recombinant protein, this fragment has been advanced as a potential malaria vaccine (12, 39). Because we found that regions of EBA175 outside of the tandem DBL domains influenced its ability to interact with GYPA, we asked whether PfEBA175 FL would be able to elicit a more potent invasion-blocking antibody response than the RII fragment alone. To address this, rabbits were immunized with an equal mass of both proteins to raise polyclonal antisera, which were subsequently purified and tested for their relative ability to inhibit erythrocyte invasion by P. falciparum in vitro. We used two different laboratory strains of P. falciparum that differed in their sensitivity to invade neuraminidase-treated erythrocytes as follows: Dd2, a strain that is dependent on sialic acid, and 3D7, which can invade through a sialic acid-independent route. The anti-PfEBA175 FL and anti-PfEBA175 RII sera, when tested at the same mass per volume (mg/ml) quantities, showed similar efficacy in inhibiting erythrocyte invasion by the two strains of P. falciparum (Fig. 6A). To assess whether antibodies targeting domains other than RII contributed to the inhibition of EBA175-dependent invasion, we first normalized the immunoreactivity of both antisera to RII, which revealed that anti-PfEBA175 FL contained $\sim 5$-fold less anti-RII antibodies than the anti-PfEBA175 RII serum (Fig. 6B). After normalizing for anti-RII immunoreactivity, we observed that antibodies elicited against PfEBA175 FL were able to inhibit parasite invasion more potently than those raised against the RII fragment alone, suggesting that antibodies directed against extracellular
regions of PfEBA175 outside of RII contribute to the ability to inhibit invasion (Fig. 6C).

**DISCUSSION**

*Plasmodium falciparum* EBA175 has long been considered an attractive anti-malarial vaccine target because of its important role in erythrocyte invasion mediated through its interactions with GYPA expressed on the surface of host erythrocytes. The technical difficulties associated with expressing *Plasmodium* proteins recombinantly have meant that most biochemical and vaccine research has relied on expressing subfragments of the EBA175 ectodomain, most commonly the tandem DBL domains known as region II. In this study, we successfully expressed the entire full-length ectodomain of PfEBA175 as a functionally active soluble recombinant protein that enabled us to perform a detailed biochemical analysis of its interaction with native GYPA, and we directly compared this with the region II subfragment.

One interesting finding from a systematic interaction screen against a panel of glycans was that the full-length PfEBA175 bound Neu5Gc-containing glycans as expected but also interacted with Neu5Ac. These results are consistent with the observations of Orlandi et al. (21) who reported that the binding of native PfEBA175 to erythrocytes was potently inhibited by Neu5Gc and oligosaccharides containing Neu5Ac2–3Gal. Neu5Gc is not present on human erythrocytes, due to the absence of the enzyme cytidine monophosphate-N-acetyleneuraminic acid hydroxylase, which is required...
for the conversion of Neu5Ac to Neu5Gc; however, it is the predominant form of sialic acid on the erythrocytes of other apes (36). This difference in sialic acid composition has been proposed to be responsible for the restriction of *P. falciparum* and the related parasite *Plasmodium reichenowi* to their respective human and chimpanzee hosts. Martin et al. (35) recombinantly expressed *Pf*EBA175 RII and *P. reichenowi* EBA175 RII (PrEBA175 RII) on the surface of monkey COS cells and observed binding only to human and chimpanzee erythrocytes, respectively, leading to the proposal that *Pf*EBA175 RII recognizes Neu5Ac-carrying GYPA, whereas PrEBA175 RII binds Neu5Gc-containing GYPA. The findings in our study do not support this hypothesis. Further investigation of the contribution of the EBA175-GYPA interaction to the restriction of *Plasmodium* spp. parasites to their natural hosts is therefore warranted, and this will be facilitated by the ability to express the entire ectodomain of EBA175 in an active form. This is of topical interest due to the recent discovery of new *Plasmodium* species that infect gorillas and that are the closest known relatives of *P. falciparum* (40, 41).

From the six regions of homology that constitute the extracellular domain of *Pf*EBA175, only RII is known to be essential and sufficient for binding to human erythrocytes (10). Several previous studies using short chemically synthesized peptides (37, 38, 42) or in vitro translated EBA175 fragments (38) have suggested additional erythrocyte binding determinants outside of region II, although a recent study did not support this (12). These findings have led to the proposition of two-step binding models that invoke conformational changes in the EBA175 protein (37, 38). Our quantitative binding analysis did not find any evidence to support a conformational change in EBA175 (at least a relatively slow one that could be detected using our SPR method) but did clearly show that the full-length ectodomain of EBA175 bound native GYPA with an affinity one order of magnitude higher than the region II fragment alone suggesting that extracellular regions of *Pf*EBA175 outside of RII do play some role in the interaction with GYPA.

The region II fragment of EBA175 crystallized as an anti-parallel dimer with the putative glycan-binding sites being formed at the dimer interface, leading to the suggestion that GYPA, which forms a dimeric complex at the erythrocyte surface, induces EBA175 dimerization upon binding (18). The SPR data we obtained for the interaction of *Pf*EBA175 with GYPA fitted better to a bivalent analyte model than to a simple 1:1 binding model, which is consistent with the dimerization of EBA175 playing a role in its interaction with GYPA. Although we showed that both *Pf*EBA175 RII and *Pf*EBA175 FL were primarily monomeric in solution, both were capable of self-association, and the full-length ectodomain of EBA175 could bind itself with ~70-fold higher affinity than the RII fragment. Extracellular regions outside of RII may therefore facilitate the
interaction with GYPA by promoting homodimerization of EBA175.

Our binding analysis also revealed that soluble PfEBA175 FL has a 2-fold lower affinity for Neu5Acα2–3Gal than for GYP A, but PfEBA175 RII bound both with similar affinities; therefore, RII probably interacts primarily with the glycan moieties of GYP A, whereas PfEBA175 FL forms contacts with both the oligosaccharides and the polypeptide backbone. We attempted to further investigate this by expressing GYP A as a recombinant protein so that it could be purified in large quantities and biochemically manipulated. Although we were able to express and purify the ectodomain of GYP A in a soluble form (rGYPA) and show that it was antigenically active by monoclonal antibody binding, it was unable to bind PfEBA175 FL (data not shown). We attributed the inability of rGYPA to bind PfEBA175 to under-sialylation because glycan profiling of rGYPA using a panel of lectins showed significantly lower levels of sialylation relative to native GYP A. Despite increasing the level of rGYPA sialylation by co-transfecting our GYP A expression construct with plasmids encoding an α-2,3-sialyltransferase and/or CMP-sialic acid transporter, this increase was not sufficient to confer binding to PfEBA175 (data not shown). In addition, we were unable to detect any binding to native GYP A using a fragment consisting of regions III to VI of EBA175 by AVEXIS (data not shown); among other possibilities, the interaction of EBA175 with the GYP A polypeptide backbone could therefore be dependent on the binding of RII to the glycan moieties on the receptor.

In conclusion, the 10-fold higher binding affinity of PfEBA175 FL for GYP A in comparison with the PfEBA175 RII fragment is likely due to the participation of the extracellular regions outside of RII in the homodimerization of EBA175 and the formation of additional contacts with GYP A.

The ability to express the entire ectodomain of P. falciparum EBA175 as a biochemically active recombinant protein and the finding that regions of EBA175 outside of the tandem DBL domains are important for GYP A binding suggested that PfEBA175 FL could be a better vaccine candidate than PfEBA175 RII alone. Consistent with these observations, when normalized for immunoreactivity to RII, the antisera to PfEBA175 FL was ~5-fold more potent than the antisera to PfEBA175 RII. This suggests that antibodies directed against extracellular regions of PfEBA175 outside of RII also contribute to inhibiting erythrocyte invasion. We envisage that the findings reported here will contribute to a more complete understanding of the molecular basis of erythrocyte invasion by P. falciparum and eventually lead to the development of an effective vaccine against this infectious disease.

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