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Blood group O protects against severe *Plasmodium falciparum* malaria through the mechanism of reduced rosetting


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Malaria has been a major selective force on the human population, and several erythrocyte polymorphisms have evolved that confer resistance to severe malaria. *Plasmodium falciparum* rosetting, a parasite virulence phenotype associated with severe malaria, is reduced in blood group O erythrocytes compared with groups A, B, and AB, but the contribution of the ABO blood group system to protection against severe malaria has received little attention. We hypothesized that blood group O may confer resistance to severe *falciparum* malaria through the mechanism of reduced rosetting. In a matched case-control study of 567 Malian children, we found that group O was present in only 21% of severe malaria cases compared with 44–45% of uncomplicated malaria controls and healthy controls. Group O was associated with a 66% reduction in the odds of developing severe malaria compared with the non-O blood groups (odds ratio 0.34, 95% confidence interval 0.19–0.61, P < 0.0005, severe cases versus uncomplicated malaria controls). In the same sample set, *P. falciparum* rosetting was reduced in parasite isolates from group O children compared with isolates from the non-O blood groups (P = 0.003, Kruskal–Wallis test). Statistical analysis indicated a significant interaction between host ABO blood group and parasite rosette formation that supports the hypothesis that the protective effect of group O operates through the mechanism of reduced *P. falciparum* rosetting. This work provides insights into malaria pathogenesis and suggests that the selective pressure imposed by malaria may contribute to the variable global distribution of ABO blood groups in the human population.

**Results and Discussion**

ABO blood group types were assessed on 567 blood samples from Malian children. Each severe malaria case (defined as unrousable by clinical criteria) was matched with two controls at similar age and sex. Gy6 and Gy7 were restricted to group O cells, and Gy6 was able to form in group A, B, or AB erythrocytes (14), and direct binding between the parasite rosetting ligand PfEMP1 and the A antigen has been demonstrated (15). The A and B trisaccharides are thought to act as receptors for rosetting on uninfected erythrocytes (14), and direct binding between the parasite rosetting ligand PfEMP1 and the A antigen has been demonstrated (15). *P. falciparum* rosettes are able to form in group O cells, but these rosettes tend to be smaller and more easily disrupted than rosettes formed in group A, B, or AB erythrocytes (10, 14).

Given the known role of the A and B trisaccharides in rosette formation, we hypothesized that blood group O may be a protective factor against severe malaria because of its rosette-reducing effects. Earlier studies have found no consistent effect of the ABO blood group on the incidence of uncomplicated malaria, parasite density, or levels of anti-malarial antibody (for review, see ref. 16). The effect of the ABO blood group on group O falciparum malaria has received little attention, although previous studies have suggested that in African children, blood group A may predispose to severe malaria (17, 18), and in Southeast Asian adults, blood group O may confer resistance to the multiorgan failure form of severe disease (19). A rigorous study of the effect of ABO blood group on susceptibility to severe malaria using a matched case-control design with adjustment for known host protective factors such as hemoglobin variants has not been performed (20). The possibility of a rosette-mediated protective effect of blood group O has been raised previously (17–21); however, the effects of ABO blood group on rosetting and susceptibility to severe malaria have not yet been examined in a single study. We therefore investigated the effect of ABO blood group on rosetting and malaria severity in a case-control study of African children.


The authors declare no conflict of interest.

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**Abbreviations:** CI, confidence interval; CR, complement receptor; OR, odds ratio.

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coma, severe anemia, neurological impairment, repeated seizures, or evidence of hepatic or renal failure; for full details see Materials and Methods) was matched by age, ethnicity, and place of residence to an uncomplicated malaria control and a healthy control child (22). Cases of nonsevere hyperparasitemia (>/H[11022]500,000 infected erythrocytes per l of blood) were also recruited and matched to uncomplicated malaria controls and healthy controls. Some researchers (e.g., refs. 18 and 23) include hyperparasitemia as a criterion for severe disease; however, we have found that children with hyperparasitemia and no other symptoms or signs of severe malaria have an excellent prognosis (0% mortality) and can thus be considered as having a form of uncomplicated malaria with very high parasite burdens (22, 24). The characteristics of the recruited children are shown in Table 1.

We found that blood group O was present in only 21% of the severe malaria cases compared with 44–45% of their matched uncomplicated malaria controls and healthy controls [Fig. 1A and supporting information (SI) Table 4]. The effect of the ABO blood group on susceptibility to severe malaria was analyzed by conditional logistic regression. The statistical model was tested for the potential confounding effect of variables occurring in the Malian population that may protect against severe malaria, including HbC and HbS (25, 26) and the CR1–Knops blood group allele Sl2 (Vil) (3, 27–29). The influence of Hb variant and CR1 genotype on the final model fit for the analysis of ABO blood group and malaria severity is described in detail in Materials and Methods (Statistical Analysis), and the frequencies of Hb variants and CR1 genotypes are shown in SI Table 5.

Conditional logistic regression analysis showed that blood group O confers significant protection against severe malaria compared with the non-O blood groups (Table 2; P < 0.0005). The protective effect of group O was seen when the severe malaria cases were matched to matched uncomplicated malaria controls and healthy controls.
compared with either the uncomplicated malaria controls or the healthy controls (Table 2). The level of protection afforded by group O [severe malaria cases versus uncomplicated malaria controls, odds ratio (OR) 0.34, 95% confidence interval (CI) 0.19–0.61, \( P < 0.0005 \) for group O versus the non-O blood groups] is equivalent to that seen for rosette-reducing CR1 deficiency polymorphism in Papua New Guinea (4) and is slightly lower than the protection against severe malaria conferred by HbS heterozygosity in previous studies (80–90% protection) (30, 31).

The above statistical analysis was carried out on severe malaria cases composed of patients with a spectrum of clinical syndromes (see Materials and Methods). This approach is valid for the hypothesis to be tested here because previous studies have shown an association between \( P. falciparum \) rosetting and multiple clinical forms of severe malaria, including cerebral malaria (7, 8), severe anemia (32), neurological impairment (8), and hepatic or renal dysfunction (33). Therefore, if blood group O does protect against severe malaria through the mechanism of reduced rosetting, we would expect this protective effect to be demonstrated across all forms of severe disease. Analysis of the effect of ABO blood group within the subcategories of severe malaria showed reduced odds ratios for group O versus non-O, ranging from 0.11 for severe anemia to 0.54 for neurological impairment (SI Table 6); however, CIs for each subcategory overlap, and a larger study would be required to address whether the protective effect of group O differs significantly between severe forms of severe malaria.

In contrast to the results for severe malaria, we found that the frequency of group O in the nonsevere hyperparasitemia cases did not differ markedly from that in their matched controls (Fig. 1B and SI Table 4). Conditional logistic regression analysis showed no significant effect of ABO blood group on susceptibility to the nonsevere hyperparasitemia form of clinical malaria (compared with uncomplicated malaria controls: OR 1.00, 95% CI 0.45–2.23, \( P = 1.0 \) for O versus non-O; compared with healthy controls: OR 0.94, 95% CI 0.46–1.90, \( P = 0.86 \) for O versus non-O). Interestingly, the ABO blood group frequencies of the controls matched to the nonsevere hyperparasitemia cases (Fig. 1B) appear to differ from the ABO blood group frequencies of the controls matched to the severe malaria cases (Fig. 1A). This result may be because the nonsevere hyperparasitemia cases and controls are older than the severe malaria cases and controls (Table 1), and they differ slightly in ethnic group composition (data not shown).

We also used conditional logistic regression to investigate the effect of ABO blood group on uncomplicated malaria compared with the healthy control children and found no significant effect of the ABO blood group on uncomplicated clinical disease (OR 0.97, 95% CI 0.58–1.61, \( P = 0.90 \) for O versus non-O using the severe malaria controls; OR 0.94, 95% CI 0.46–1.90, \( P = 0.86 \) for O versus non-O using the nonsevere hyperparasitemia controls). Therefore, blood group O only protects against severe, life-threatening malaria (Table 2) and not against uncomplicated clinical malaria with low or high parasite burdens. The number of deaths in the recruited children was too low for a reliable analysis of the effect of the ABO blood group on mortality from malaria. Fourteen children from the severe malaria group died, of which five (36%) were group A, one (7%) was group AB, six (43%) were group B, and two (14%) were group O.

To determine whether reduced rosetting could be the explanation for the protective effect of blood group O against severe malaria in Mali, parasite rosette frequency (percentage of infected erythrocytes binding two or more uninfected erythrocytes) was assessed in the same set of samples. The rosette frequency varies from one parasite isolate to another, and the level of rosetting in a given isolate will depend partly on the variant surface antigens expressed by that isolate (i.e., whether rosetting or nonrosetting PfEMP1 variants are being expressed) (3) and partly on the phenotype of the host erythrocytes (i.e., whether rosetting receptors such as the A or B trisaccharides are present). Rosette frequencies were found to be significantly lower in parasite isolates from patients with blood group O compared with isolates from patients with groups A, B, and AB (Fig. 1C, \( P = 0.003 \), Kruskal–Wallis test). In addition, rosetting was significantly associated with severe malaria in patients with blood groups A, B, and AB (Fig. 2); however, this relationship was not apparent in patients with blood group O (Fig. 2).

The results shown in Figs. 1 and 2 are consistent with the hypothesis that children with blood group O are protected against severe malaria through the mechanism of reduced \( P. falciparum \) rosetting. However, it remains possible that the ABO blood group polymorphism in Papuan children is a risk factor for severe malaria in children with the non-O blood groups, as is consistent with the results shown in Fig. 2. Further analysis showed that there was a significant interaction between the ABO blood group and parasite rosette frequency (interaction OR 0.11, 95% CI 0.02–0.58, \( P = 0.01 \)), which means that group O children infected with rosetting parasites (>5% rosette frequency) have a greatly increased risk of severe malaria compared with the reference group (OR 15.23, 95% CI 5.24–49.86, \( P < 0.0001 \)). This observation indicates that rosetting is a risk factor for severe malaria in children with the non-O blood groups, as is consistent with the results shown in Fig. 2. Further analysis showed that there was a significant interaction between the ABO blood group and parasite rosette frequency (interaction OR 0.11, 95% CI 0.02–0.58, \( P = 0.01 \)), which means that group O children infected with rosetting parasites (>5% rosette frequency) are still at risk of severe malaria (OR 1.63); however, this risk is

### Table 2. Effect of blood group O on resistance to severe malaria in Mali

<table>
<thead>
<tr>
<th>Blood groups</th>
<th>Severe malaria cases versus uncomplicated malaria odds ratio (95% CI)</th>
<th>P value*</th>
<th>Severe malaria cases versus healthy controls odds ratio (95% CI)</th>
<th>P value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>O vs. A</td>
<td>0.39 (0.21–0.75) P = 0.0048</td>
<td></td>
<td>0.28 (0.13–0.61) P = 0.0014</td>
<td></td>
</tr>
<tr>
<td>O vs. B</td>
<td>0.28 (0.14–0.57) P = 0.0005</td>
<td></td>
<td>0.29 (0.14–0.59) P = 0.0007</td>
<td></td>
</tr>
<tr>
<td>O vs. AB</td>
<td>0.31 (0.11–0.89) P = 0.0290</td>
<td></td>
<td>0.44 (0.17–1.17) P = 0.1000</td>
<td></td>
</tr>
<tr>
<td>O vs. non-O</td>
<td>0.34 (0.19–0.61) P = 0.0003</td>
<td></td>
<td>0.31 (0.17–0.58) P = 0.0003</td>
<td></td>
</tr>
</tbody>
</table>

*Conditional logistic regression analysis of matched cases and controls (\( n = 124 \) triplets).
Substantially reduced compared with the non-O children infected with rosetting parasites (OR 15.23). In summary, this analysis shows that group O only protects in the presence of rosetting parasites and that rosetting is a strong risk factor for severe disease in children with non-O blood groups but a much weaker risk factor in group O children. These data support the hypothesis that the protective effect of group O operates through the mechanism of reduced *P. falciparum* rosetting. The cutoff rule used in the above analysis (no/low rosetting is ≤5% rosette frequency) is arbitrary because the exact rosette frequency at which significant pathogenic effects start to occur is unknown. However, varying the cutoff point to ≤2% or ≤10% rosette frequency did not materially alter the conclusions of the analysis (see SI Table 7).

Parasite density did not differ between the ABO types in the Mali study, indicating that the protective effect of group O is not caused by reduced parasite burden \(\log_{10}\text{parasite density, median (±SD)}\) severe cases; group A, 4.94 (0.78); group AB, 4.80 (0.37); group B, 4.86 (0.91); and group O, 4.97 (0.93); uncomplicated malaria controls; group A, 3.87 (0.70); group AB, 3.71 (0.92); group B, 3.69 (0.71); and group O, 3.84 (0.68); \(P = 0.36\), ANOVA].

To determine whether the relationship between the ABO blood group and resistance to severe malaria is found elsewhere, we retrospectively examined the effect of ABO blood group on malaria severity by using samples collected during two previous studies in Kilifi, Kenya, in 1993 and 2003 (8, 34). Rosetting was shown to be significantly lower in parasite isolates from patients with blood group O compared with the non-O blood groups in the 1993 study (8). In both studies, the ABO blood group was assessed on samples from children with uncomplicated and severe malaria (for clinical definitions and inclusion criteria, see Materials and Methods). The patient characteristics for the samples included in the ABO analysis are shown in SI Table 8. The Kenyan data are derived from small unmatched, retrospective studies that are not adjusted for potential confounding variables; however, they do provide preliminary data addressing whether blood group O confers protection against severe malaria in another population. As can be seen in Table 3, the ORs for severe malaria were reduced in group O patients compared with non-O patients for both Kenyan studies, but only the 1993 study was statistically significant.

Other work also supports the protective effect of group O against severe malaria in multiple countries, although the level of protection varies between sites, and some of these studies do not use matched controls or take into account potential confounding variables (refs. 17–19 and A. Fry, M. Griffiths, S. Auburn, M. Diakite, J. Forton, et al., unpublished work). Previous work has also suggested that group A and AB are greater risk factors for severe disease than group B (17, 18), which was not apparent in our study in Mali (Table 2). It has been shown previously that *P. falciparum* rosetting strains are either A-preferring (forming larger rosettes with group A or AB erythrocytes) or B-preferring (forming larger rosettes with group B or AB erythrocytes) (10), therefore it is possible that variation in parasite rosetting phenotypes in distinct geographical areas could influence the effect of blood group on severe malaria.

Previous experimental work has provided evidence that rosetting may be a parasite virulence factor because it causes microvascular obstruction (9), which is thought to be a key process in the pathogenesis of severe malaria (35). In an ex vivo model, *P. falciparum* rosettes were disrupted by high shear forces in the arterial side of the circulation but reformed in the postcapillary venules by adhesion of uninfected erythrocytes onto infected erythrocytes that were bound to endothelial cells (9). This combination of rosetting and cytoadherence occurring simultaneously, resulted in rosetting parasites causing greater obstruction to microvascular blood flow than isogenic cytoadherent nonrosetting parasites (9). The ABO determinants are present on endothelial

**Fig. 2.** Rosetting and severe malaria in Mali in relation to ABO blood group. The distributions of rosette frequencies in patients with nonsevere hyperparasitemia (hyp), severe malaria (severe), and uncomplicated malaria (uncomp) are shown as box plots (as in Fig. 1) for each ABO blood group type. High levels of rosetting were seen most frequently in severe malaria isolates in groups A, B, and AB, but not in group O (Kruskal–Wallis test; \(P\) value shown in parentheses above each graph). Numbers in each category are as follows: group A (hyp, 8; severe, 27; and uncomp, 16); group AB (hyp, 2; severe, 4; and uncomp, 5); group B (hyp, 11; severe, 29; and uncomp, 25); group O (hyp, 17; severe, 15; and uncomp, 43).
cells and platelets as well as erythrocytes, and it seems likely that parasite isolates that bind to A or B determinants on erythrocytes to form rosettes may also bind A or B antigens on other cell types, which could enhance sequestration and increase pathogenic potential (20).

In this work we have shown that blood group O protects against severe malaria in a matched case-control study of Malian children, and we provided preliminary evidence for a similar protective effect of group O in Kenya. Statistical analysis of the Malian study indicates a significant interaction between the host ABO blood group and parasite rosette frequency that provides strong evidence to support the hypothesis that group O protects by the mechanism of reduced rosetting and sequestration. These findings indicate that blood group O provides a further example of an erythrocyte polymorphism that, similarly to CR1 deficiency, a-thalassemia, and HbC (4–6), is able to reduce the adhesion potential of P. falciparum-infected erythrocytes and consequently modify the virulence of the parasite.

One obvious question that arises from this work is why blood group O does not occur at higher frequency in all malaria endemic regions. It seems likely that this phenomenon represents an example of a balanced polymorphism in the human population because blood group O is thought to confer susceptibility to diseases such as cholera (36) and other diarrheal diseases (37–39) that may be a significant selective force in many malarious countries. The global distribution of ABO blood group types is complex and may be influenced by selection imposed by a variety of pathogenic microorganisms (40, 41). Our work indicates that malaria is likely to be a significant factor influencing ABO blood group frequencies in tropical and subtropical regions of the world.

Materials and Methods

Mali Study Site and Field Isolates. Blood samples were collected in Bandiagara, Mali, an area with intense seasonal transmission of P. falciparum (20–60 infected bites per person per month at the peak of the July–December transmission season) (42). The samples were collected as part of the Bandiagara Malaria Project case-control study that has been described in detail previously (22). Blood samples were collected after informed consent from children’s parents or guardians, and all protocols received institutional review board approval. The World Health Organization (WHO) criteria for severe malaria were applied (23), except that patients with board approval. The World Health Organization (WHO) criteria for severe malaria were applied (23), except that patients with severe malaria and no hyperparasitemia. The healthy control samples were collected after informed consent from children’s group. Uncomplicated malaria controls were children with malaria presenting at the outpatient department of the same hospital during the same study period with no signs of severe disease, who were successfully treated as outpatients with oral antimalarial therapy. ABO blood group data were collected for 2 months of the 1993 study (June and July) and throughout the study period in 2003. All samples fulfilling the clinical definitions above for which ABO data were available were included in the analysis.

ABO Blood Group Typing. Blood samples were typed for ABO blood group by standard hemagglutination techniques (44).

Hemoglobin Typing. Hemoglobin types (HbA, HbS, HbC, and HbF) of the Malian samples were analyzed by cellulose acetate electrophoresis (25). In 141 samples, electrophoresis results were confirmed by PCR (26).

CR1 Genotyping. Samples were genotyped for the CR1–Knops blood group alleles SlI and Sl2 by PCR and restriction digest as described (45).

P. falciparum Culture. After removal of lymphocytes (46), the Malian samples were washed and then frozen in glycerolcryo to −80°C and shipped to Edinburgh. The isolates were thawed and cultured for 18–36 h as described (46) to allow maturation from ring stage to the pigmented trophozoite stage at which rosetting occurs. No addi-

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**Table 3. Effect of blood group O on resistance severe malaria in Kenya**

<table>
<thead>
<tr>
<th>Category</th>
<th>Blood group</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>A</td>
</tr>
<tr>
<td>1993</td>
<td></td>
</tr>
<tr>
<td>Severe n = 19</td>
<td>5 (26.3)</td>
</tr>
<tr>
<td>Uncomplicated n = 36</td>
<td>9 (25.0)</td>
</tr>
<tr>
<td>2003</td>
<td></td>
</tr>
<tr>
<td>Severe n = 20</td>
<td>7 (35.0)</td>
</tr>
<tr>
<td>Uncomplicated n = 19</td>
<td>3 (15.8)</td>
</tr>
</tbody>
</table>

Shown are the numbers in each blood group with percentage in parentheses.

*Statistics for the comparison of group O vs. non-O patients: 1993: χ2, 5.15; OR, 0.24; 95% CI, 0.07–0.83; P = 0.043.

2003: χ2, 3.09; OR, 0.31; 95% CI, 0.09–1.14; P = 0.11.

Kenya Study Site and Field Isolates. Blood samples were collected from patients with a P. falciparum parasitemia of 0.5% or higher, attending Kilifi District Hospital, Kenya, in two studies carried out in 1993 and 2003. The effect of ABO blood group on rosetting in the 1993 study has been reported previously (8). The invasion phenotypes of some of the samples recruited in 2003 have also been reported (34). At this site in Kenya, malaria transmission is seasonal (June–August and December–February), with the average number of infected bites per person estimated at ~10–30 per year (43). In both 1993 and 2003, severe malaria was defined as cerebral malaria (as described above for Mali), prostration (inability to sit or in babies, to breast feed), or respiratory distress (abnormally deep breathing). This clinical definition identifies approximately the same group of children at risk of life-threatening malaria as those identified by the more comprehensive WHO criteria (23). Uncomplicated malaria controls were children with malaria presenting at the outpatient department of the same hospital during the same study period with no signs of severe disease, who were successfully treated as outpatients with oral antimalarial therapy. ABO blood group data were collected for 2 months of the 1993 study (June and July) and throughout the study period in 2003. All samples fulfilling the clinical definitions above for which ABO data were available were included in the analysis.
tional erythrocytes were added during this short culture step. Parasite maturity was monitored by Giemsa smear, and only those with normal morphology that matured to the pigmented tropho-
zoite stage were assessed for rosetting. Of a possible 378 P. falciparum-infected samples, 272 were put into culture, the others having been lost during a freezer malfunction. Of those cultured, 13 did not grow, and 51 were excluded because the parasitemia was too low to assess rosetting (<0.5%). Therefore, rosette frequency was assessed for 208 samples (91 uncomplicated malaria, 40 nonsevere hyperparasitemia, and 77 severe malaria).

**Rosetting Assays.** Rosette frequency was assessed in the first cycle of *in vitro* growth. An aliquot of culture suspension was stained with 25 µg/ml ethidium bromide for 5 min at 37°C. A wet preparation of the suspension (2% hematocrit) was viewed with a fluorescence microscope, and the number of mature infected erythrocytes binding two or more unfixed erythrocytes was counted. The rosette frequency is the percentage of infected erythrocytes in rosettes of 200 infected erythrocytes counted.

**Statistical Analysis.** A conditional logistic regression model (the survival package within the R statistical system) (47) was used to estimate ORs and 95% CIs for the effect of ABO blood group on malaria severity in Mali and to test for the potentially confounding effect of hemoglobin variant (HbS and HbC) and CR1 Sl genotype. Candidate variables were initially identified by univariate logistic modeling using P < 0.2 as an inclusion criterion. Data for some hemoglobin classes were sparse, so this variable was reclassified to wild-type/normal hemoglobin before analysis. The final logistic regression models for the ABO analyses were built by manual inspection of overall fit, significance of coefficients, and the effect of additional variables on estimated ORs. Hb variant and CR1 Sl genotype were not included in the final model for the comparison of severe cases and the uncomplicated malaria controls because their inclusion did not have a marked effect on ABO ORs, and their parameter estimates were not statistically significant. Hb variant did improve model fit for the ABO analysis using severe cases and healthy controls and was therefore included in the final model. CR1 Sl genotype was excluded because it did not improve model fit.

We are grateful to the Bandiagara Malaria Project team in Mali and the clinical, nursing, and laboratory staff at the KEMRI Unit in Kilifi for assistance with this work. We also thank the children and their parents/ guardians for their study site visit, and the KEMRI-Wellcome Trust, Ph.D. studentship (to A.-M.D.) and Senior Research Fellowship (67431 to J.A.R.), National Institutes of Health Contract N01-AI-85346 and Grant AI 42367, and Fogarty International Center Training Grant D43TW001589.