Epistasis between the haptoglobin common variant and $\alpha^+\text{thalassemia}$ influences risk of severe malaria in Kenyan children

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Key Points

- Epistasis between the haptoglobin (Hp) common variants and $\alpha^+\text{thalassemia}$ predicts the risk of severe malaria in Kenyan children.
- The inconsistent relationship between Hp type and malaria reported in previous studies may be explained by this epistatic interaction.

Haptoglobin (Hp) scavenges free hemoglobin following malaria-induced hemolysis. Few studies have investigated the relationship between the common Hp variants and the risk of severe malaria, and their results are inconclusive. We conducted a case-control study of 996 children with severe Plasmodium falciparum malaria and 1220 community controls and genotyped for Hp, hemoglobin (Hb) S heterozygotes, and $\alpha^+\text{thalassemia}$. Hp S heterozygotes and $\alpha^+\text{thalassemia}$ homozygotes were protected from severe malaria (odds ratio [OR], 0.12; 95% confidence interval [CI], 0.07-0.18 and OR, 0.69; 95% CI, 0.53-0.91, respectively). The risk of severe malaria also varied by Hp genotype: Hp2-1 was associated with the greatest protection against severe malaria and Hp2-2 with the greatest risk. Meta-analysis of the current and published studies suggests that Hp2-2 is associated with increased risk of severe malaria compared with Hp2-1. We found a significant interaction between Hp genotype and $\alpha^+\text{thalassemia}$ in predicting risk of severe malaria: Hp2-1 in combination with heterozygous or homozygous $\alpha^+\text{thalassemia}$ was associated with protection from severe malaria (OR, 0.73; 95% CI, 0.54-0.99 and OR, 0.48; 95% CI, 0.32-0.73, respectively), but $\alpha^+\text{thalassemia}$ in combination with Hp2-2 was not protective. This epistatic interaction together with varying frequencies of $\alpha^+\text{thalassemia}$ across Africa may explain the inconsistent relationship between Hp genotype and malaria reported in previous studies. (Blood. 2014;123(13):2008-2016)

Introduction

Severe malaria ranks third among infectious causes of childhood death worldwide.1 Cell-free hemoglobin (Hb), released following intravascular hemolysis, is highly toxic and may contribute to the pathogenesis of severe malaria by increasing inflammation, vasoconstriction, and oxidant damage and by promoting adhesion of parasite-infected red blood cells to molecules such as intercellular adhesion molecule-1.2,3 Haptoglobin (Hp) is an acute-phase plasma protein that binds rapidly and irreversibly with cell-free Hb and the resultant Hp-Hb complexes are cleared via CD163 receptors on monocytes and tissue macrophages.4,5 Hp exists in 3 common variants in humans: the Hp1-1 dimer, the Hp2-1 linear polymer, and the large Hp2-2 circular multimer.4 Hb binding to the various Hp in humans: the Hp1-1 dimer, the Hp2-1 linear polymer, and the Hp2-2 circular multimer.4,5 Hb binding to the various Hp proteins may vary, as seen in the Hp2-2-Hb complex, which has an unstable heme moiety and is itself redox active.5 Moreover, the Hp2-2 type is associated with Hb-mediated oxidant damage,6-9 inflammation,10 endothelial cell dysfunction,11,12 and vasospasm following hemolysis.13,14

Whether the risk of severe malaria is altered by Hp type remains unclear. Four studies that have investigated this question have produced conflicting results.15-18 Early studies, conducted in Sudan and Ghana using plasma electrophoresis to determine Hp phenotype, suggested that the Hp1-1 type was associated with an increased risk of severe malaria.15,16 However, malaria-induced hemolysis depletes plasma Hp, and in the Ghanaian study, 45% of children had an unknown or Hp0 phenotype,16 whereas the prevalence of Hp0 was not reported in the Sudanese study.15 More recent studies, conducted in The Gambia and Northern Ghana using polymerase chain reaction (PCR)-based genotyping methods, found no association between Hp genotype and severe malaria and an increased risk of severe malaria in children with the Hp2-2 genotype, respectively.17,18 Recent studies investigating the relationship between Hp genotype and the risk of uncomplicated malaria are similarly inconclusive, finding variously an increased risk of symptomatic malaria in Hp2-2 individuals,19 a reduced incidence among older children with the Hp2-2 genotype,20,21 and an increased placental parasite density among pregnant mothers with the Hp1-1 genotype.22 The effect of Hp concentration on the risk of severe malaria is also not known. Hb was toxic to malaria parasites in vitro,23 and increased parasite burden was demonstrated in an Hp knockout mouse.24 The A-61C single nucleotide polymorphism (SNP) in the Hp promoter region,


S.H.A. and S.M.U. contributed equally to this study.

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which causes very low Hp concentrations,\textsuperscript{25,26} was associated with a reduced risk of uncomplicated malaria.\textsuperscript{21}

Because the common $HP^2$ allele arises from a large gene duplication\textsuperscript{27} and cannot be uniquely identified by a SNP, it has not been included in recent genome-wide association studies of severe malaria.\textsuperscript{28,29} Here we report the results of the largest case-control study conducted to date ($n = 996$ severe malaria cases and $1220$ controls) that has investigated the association between Hp genotype and susceptibility to severe Plasmodium falciparum malaria. We further considered for the first time possible interactions between Hp and two other malaria resistance genes: Hb S heterozygotes (HbAS) and $\alpha^+\text{ thalassemia}$.

### Methods

#### Study population

The study was conducted in the Kilifi district on the coast of Kenya, where the majority of the population are rural dwellers of the Mijikenda ethno-linguistic group. The clinical characteristics and epidemiology of malaria in the study area have been described in detail previously.\textsuperscript{30,31} Case patients were children ($<14$ years of age) who were residents of the study area served by the Kilifi Health and Demographic Surveillance System\textsuperscript{32} and who were admitted to the High Dependency Unit at Kilifi District Hospital between January 7, 2001 and January 6, 2010 with severe malaria. Severe malaria was defined as a positive blood film for P falciparum parasites in association with $\geq 1$ of the following: prostration (Blantyre Coma Score 3–4), coma (Blantyre Coma Score 2), or respiratory distress (intercostal recession or deep or labored breathing).\textsuperscript{33} Children with uncomplicated severe malarial anemia (ie, Hb $<5$ g/dL, but with no signs of respiratory distress, prostration, or any other complications) are not routinely admitted to our High Dependency Unit and were not included in our study, because uncomplicated severe malarial anemia (SMA), when treated with transfusion, does not require intensive supportive care and is associated with a mortality of $<1\%$.\textsuperscript{30} All case patients were treated according to standard guidelines as described in detail previously.\textsuperscript{33} The collection of case samples is summarized in Figure 1.

Controls were children born within the same study area as cases between August 14, 2006 and August 20, 2010 and were sampled as part of a cohort study of genetic susceptibility to infectious diseases during home visits at 3 to 11 months of age.\textsuperscript{34} Individual written informed consent was provided by all study participants or their parents/guardians. The study was approved by the Kenya Medical Research Institute (KEMRI)/National Ethical Review Committee in Nairobi. The study was conducted in accordance with the Declaration of Helsinki.

#### Laboratory methods

Routine hematologic, biochemical, and malaria parasite data were collected for all case children using standard methods as described previously.\textsuperscript{35} DNA was extracted by standard methods using proprietary kits (Qiagen DNA Blood Mini Kit; Qiagen, West Sussex, United Kingdom; or Puregene; Gentra Systems, Minneapolis, MN), and Hp genotypes were determined by allele-specific PCR as previously described.\textsuperscript{36} Hb typing for HbAA, HbAS, and HbSS was conducted by either electrophoresis using cellulose acetate gels (Helena) or by high-performance liquid chromatography (Variant analyzer; BioRad, Hercules, CA) using the $\beta$-thalassemia short program, whereas the $\alpha$-thalassemia deletion in the $\alpha$-globin gene, the common form of $\alpha$-thalassemia in Africa, was typed by PCR as described previously.\textsuperscript{36} The Hp promoter polymorphism, A-61C (rs5471), was typed at the Wellcome Trust Centre for Human Genetics, Oxford, United Kingdom, by matrix-assisted laser desorption/ionization time-of-flight mass spectrometry using the SEQUENOM system (San Diego, CA).\textsuperscript{37}

#### Statistical analyses

All analyses were conducted using Stata V12.0 (StataCorp, College Station, TX). Non-normally distributed variables were log-transformed prior to analysis. Categorical data were analyzed using the $\chi^2$ test, whereas the distribution of continuous variables by Hp genotype was investigated using linear regression. The association between Hp genotype and severe malaria was determined by analysis of variance. Logistic regression was used to derive odds ratios (ORs) for risk of severe malaria overall, severe malaria subgroups, and in-patient death according to Hp genotype in univariable and multivariable models adjusted for potential confounders including gender, ethnicity, Hb type (HbAS), and $\alpha$-thalassemia genotype. Interactions between Hp genotype and other explanatory variables were assessed using the likelihood-ratio test.

#### Meta-analysis of studies of Hp type and severe malaria

The following criteria were applied to potential studies for inclusion in the meta-analysis: (1) indexed in PubMed (National Center for Biotechnology Information) under the search terms “severe malaria” and “ Hp phenotype” or “ Hp genotype”; (2) original studies in which both case patients and controls were children; and (3) a case-control study design with clear comparative groups of cases with severe malaria and controls without severe malaria. Four studies were identified,\textsuperscript{15-18} of which 2 were excluded: one because the study also included adult cases and a control group that consisted only of adults recruited among the staff, technicians, workers, and students at the institute;\textsuperscript{15} and a second because 45% of cases and controls were of unknown (Hp0) phenotype.\textsuperscript{16} Data from the 2 remaining studies\textsuperscript{17,18} were extracted and summarized, and meta-analysis was performed using STATA V.12.0.
Table 1. Characteristics of severe malaria patients by Hp genotype

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Hp 1-1</th>
<th>Hp 2-1</th>
<th>Hp 2-2</th>
</tr>
</thead>
<tbody>
<tr>
<td>No. (%)</td>
<td>312 (31.3)</td>
<td>423 (42.5)</td>
<td>261 (26.2)</td>
</tr>
<tr>
<td>Median age, mo (IQR)</td>
<td>27.4 (15.2-43.6)</td>
<td>26.8 (15.2-41.4)</td>
<td>27.5 (14.7-41.6)</td>
</tr>
<tr>
<td>Male gender, no. (%)</td>
<td>156 (50.0)</td>
<td>203 (48.0)</td>
<td>133 (51.0)</td>
</tr>
<tr>
<td>Ethnic group, no. (%)</td>
<td>Girama 168 (53.9)</td>
<td>225 (53.2)</td>
<td>157 (60.2)</td>
</tr>
<tr>
<td></td>
<td>Chonyi 82 (26.3)</td>
<td>119 (28.1)</td>
<td>56 (21.5)</td>
</tr>
<tr>
<td></td>
<td>Kauma 35 (11.2)</td>
<td>41 (9.7)</td>
<td>21 (8.1)</td>
</tr>
<tr>
<td></td>
<td>Other Mijikenda 19 (6.1)</td>
<td>21 (5.0)</td>
<td>24 (9.2)</td>
</tr>
<tr>
<td></td>
<td>Non-Mijikenda 8 (2.6)</td>
<td>17 (4.0)</td>
<td>3 (1.2)</td>
</tr>
<tr>
<td>HbS type no. (%) (n = 990)</td>
<td>Hb AA 300 (96.5)</td>
<td>408 (97.6)</td>
<td>254 (97.3)</td>
</tr>
<tr>
<td></td>
<td>Hb AS 10 (3.2)</td>
<td>7 (1.7)</td>
<td>5 (1.9)</td>
</tr>
<tr>
<td></td>
<td>Hb SS 1 (0.3)</td>
<td>3 (0.7)</td>
<td>2 (0.8)</td>
</tr>
<tr>
<td>Hb type [HbAS], and α thalassemia genotype no. (%)</td>
<td>α-/α- 126 (40.4)</td>
<td>170 (40.2)</td>
<td>102 (39.1)</td>
</tr>
<tr>
<td></td>
<td>−α/α 150 (48.1)</td>
<td>203 (48.0)</td>
<td>128 (49.0)</td>
</tr>
<tr>
<td></td>
<td>α/αa 36 (11.5)</td>
<td>50 (11.8)</td>
<td>31 (11.9)</td>
</tr>
<tr>
<td>Body mass index (n = 893)*</td>
<td>10.6 (10.2-11.0)</td>
<td>10.2 (9.9-10.6)</td>
<td>10.3 (9.9-10.7)</td>
</tr>
<tr>
<td>Parasitemia, log10 value, μL (n = 988)</td>
<td>4.58 (4.45-4.71)</td>
<td>4.64 (4.53-4.75)</td>
<td>4.67 (4.54-4.81)</td>
</tr>
<tr>
<td>Hb, g/dL (n = 993)</td>
<td>6.6 (6.3-6.8)</td>
<td>6.6 (6.4-6.8)</td>
<td>6.7 (6.4-7.0)</td>
</tr>
</tbody>
</table>

Numbers (with percentages) are presented for categorical variables. Geometric means are presented with 95% confidence intervals (CIs) for continuous variables.

*aDue to severity of illness, a proportion of children did not have height measured.

Results

Characteristics of severe malaria patients

Our study included 996 case patients with severe malaria and 1220 controls. The median age of patients was 27.2 months (range, 0-147 months), 49.8% were boys, and 97.2% were of the Mijikenda ethnic group. The characteristics of case patients, stratified by Hp genotype, are summarized in Table 1. Overall, the geometric mean parasite density and Hb concentrations were 57 435/μL (95% CI, 49 405-66 770) and 6.6 g/dL (95% CI, 6.5-6.8), respectively, whereas the proportions of severe malaria patients with HbAS, −α/α, thalassemia, −α/−α thalassemia, and Hp-61C were 2.2%, 48.3%, 11.8%, and 20.2%, respectively. The demographic characteristics of severe malaria patients did not vary according to Hp genotype (Table 1). Cerebral malaria was diagnosed in 528 children (53.1%), severe anemia in 224 children (22.6%), and 104 children (10.5%) had both cerebral malaria and severe anemia. Other nonexclusive categorizations of severe malaria, stratified by Hp genotype, are summarized in Table 2. Ninety-four severe malaria patients (9.4%) died.

Hp genotype and risk of severe malaria

Hp genotype was associated with risk of severe malaria by analysis of variance (P = .01 before and after adjustment for gender, ethnicity, Hb type [HbAS], and α thalassemia genotype). We therefore used logistic regression models to determine which of the 3 2-way comparisons between Hp genotypes accounted for this statistically significant variation in risk (Table 3).

Hp2-2 was associated with the greatest risk andHp2-1 with the least risk of severe malaria. The risk of severe malaria did not differ significantly between Hp1-1 and Hp2-2 genotypes (P = .32) and varied with marginal significance between Hp1-1 and Hp2-1 genotypes (P = .06; Table 3). However, the risk of severe malaria was significantly increased in the Hp2-2 genotype compared with the Hp2-1 genotype (OR, 1.35; 95% CI, 1.10-1.67; P = .005) or compared with Hp2-1 and Hp1-1 genotypes combined (OR, 1.26; 95% CI, 1.03-1.53; P = .02). Conversely, the Hp2-1 genotype was associated with 26%
protection against severe malaria compared with Hp2-2 (OR, 0.74; 95% CI, 0.60-0.91; $P = .005$) or 21% protection compared with the Hp2-2 and Hp1-1 genotypes combined (OR, 0.79; 95% CI, 0.67-0.93; $P = .006$), indicating a heterozygote advantage.

ORs were significantly raised in the Hp2-2 genotype compared with the Hp2-1 genotype for all categories of severe malaria except for symptomatic SMA, in which there was a nonsignificantly increased risk ($P = .10$; Table 3). Hp2-2 was also associated with a marginally significant increased OR for death compared with Hp1-1. Compared with children of the Hp2-1 genotype, those of the Hp1-1 genotype were at significantly increased risk of deep breathing and metabolic acidosis (Table 3). Parasite densities did not differ significantly between the Hp genotypes.

**Hp A-61C promoter polymorphism**

To investigate the effect of Hp concentration on the risk of severe malaria, we typed the Hp A-61C promoter SNP, which is known to be associated with very low Hp concentrations,23 in a subset of 519 case patients and 876 controls for whom DNA samples were available in the United Kingdom. The characteristics of the subset typed did not differ from the nontyped group in any respect except with regard to subgroups within the Mijikenda ethnic group (the typed did not differ from the nontyped group in any respect except available in the United Kingdom. The characteristics of the subset case patients and 876 controls for whom DNA samples were

### Table 3. ORs for severe malaria by Hp genotype

<table>
<thead>
<tr>
<th>Groups</th>
<th>Hp1-1 vs Hp2-1 OR (95% CI)*</th>
<th>P†</th>
<th>Hp1-1 vs Hp2-2 OR (95% CI)‡</th>
<th>P‡</th>
<th>Hp2-2 vs Hp1-1 OR (95% CI)*</th>
<th>P*</th>
</tr>
</thead>
<tbody>
<tr>
<td>All severe malaria</td>
<td>1.21 (0.99-1.47)</td>
<td>.06</td>
<td>0.89 (0.71-1.12)</td>
<td>.32</td>
<td>1.35 (1.1-1.67)</td>
<td>.005</td>
</tr>
<tr>
<td>All severe malaria adjusted§</td>
<td>1.24 (1.01-1.53)</td>
<td>.04</td>
<td>0.92 (0.72-1.16)</td>
<td>.47</td>
<td>1.36 (1.09-1.70)</td>
<td>.007</td>
</tr>
<tr>
<td>Severe malaria syndromes¶</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Coma</td>
<td>1.30 (0.99-1.71)</td>
<td>.06</td>
<td>1.01 (0.76-1.36)</td>
<td>.90</td>
<td>1.33 (1.03-1.70)</td>
<td>.03</td>
</tr>
<tr>
<td>Prostration</td>
<td>1.35 (0.96-1.89)</td>
<td>.09</td>
<td>0.90 (0.61-1.33)</td>
<td>.60</td>
<td>1.49 (1.04-2.14)</td>
<td>.03</td>
</tr>
<tr>
<td>Deep breathing</td>
<td>1.35 (1.03-1.77)</td>
<td>.03</td>
<td>1.0 (0.73-1.36)</td>
<td>.99</td>
<td>1.35 (1.01-1.81)</td>
<td>.04</td>
</tr>
<tr>
<td>Coma and deep breathing</td>
<td>1.38 (0.97-1.97)</td>
<td>.08</td>
<td>0.91 (0.61-1.35)</td>
<td>.65</td>
<td>1.51 (1.04-2.21)</td>
<td>.03</td>
</tr>
<tr>
<td>Metabolic acidosis</td>
<td>1.46 (1.14-1.90)</td>
<td>.003</td>
<td>1.03 (0.76-1.36)</td>
<td>.85</td>
<td>1.42 (1.08-1.85)</td>
<td>.01</td>
</tr>
<tr>
<td>Symptomatic SMA</td>
<td>1.22 (0.87-1.72)</td>
<td>.25</td>
<td>0.90 (0.61-1.33)</td>
<td>.59</td>
<td>1.36 (0.95-1.97)</td>
<td>.10</td>
</tr>
<tr>
<td>Fatal outcome</td>
<td>0.71 (0.41-1.23)</td>
<td>.22</td>
<td>0.54 (0.30-1.0)</td>
<td>.05</td>
<td>1.29 (0.78-2.16)</td>
<td>.32</td>
</tr>
</tbody>
</table>

* OR (95% CI)†
** OR (95% CI)‡
§Adjusted for gender, ethnic group, sickle status, and $\alpha^+\text{thalassemia}$ type.
¶Defined as in Table 2.

#### $\alpha^+\text{thalassemia}$

The heterozygous $\alpha^+\text{thalassemia}$ ($-\alpha/\alpha\alpha$) was carried by 481 cases (48.3%) and 591 controls (48.4%), whereas homozygous $\alpha^+\text{thalassemia}$ ($-\alpha/-\alpha$) was carried by 117 cases (11.8%) and 187 controls (15.3%). The $-\alpha/-\alpha$ genotype was associated with 31% protection against severe malaria (OR, 0.69; 95% CI, 0.53-0.91; $P = .008$), whereas the $-\alpha/\alpha\alpha$ thalassemia type was not significantly associated with protection ($P = .27$). However, we found statistically significant evidence for an interaction between $\alpha^+\text{thalassemia}$ and Hp type in predicting the risk of severe malaria ($P = .02$ in likelihood-ratio testing). The risk of severe malaria was lowest for $-\alpha/-\alpha$ thalassemia inherited in combination with Hp2-1, but $\alpha^+\text{thalassemia}$ was not protective when inherited in combination with Hp2-2. Hp1-1 was associated with a nonsignificant trend toward protection against severe malaria when inherited with $\alpha^+\text{thalassemia}$ (Figure 2). ORs for the risk of severe malaria stratified by $\alpha^+\text{thalassemia}$ are shown in Table 4, illustrating the effect of $\alpha^+\text{thalassemia}$ in modifying risk associated with Hp genotype.

#### Meta-analysis

In addition to our current study, we identified 2 eligible additional studies relating to Hp type and the risk of severe malaria17,18 for inclusion in a meta-analysis. Our analysis suggested that the Hp2-2 genotype is associated with an increased risk of severe malaria compared with the Hp2-1 genotype (OR, 1.20; 95% CI, 1.02-1.42; $P = .028$; Figure 3) but that there is no significant variation in risk of severe malaria between Hp2-2 and Hp1-1 or between Hp1-1 and Hp2-1. We further stratified the current study by $\alpha^+\text{thalassemia}$ and showed significant heterogeneity between these groups in predicting risk of severe malaria for Hp2-2 vs Hp2-1 ($P = .006$); the OR for severe malaria was 0.93 (95% CI, 0.67-1.30) in the group without $\alpha^+\text{thalassemia}$ and 1.71 (95% CI, 1.30-2.24) in the group with $\alpha^+\text{thalassemia}$ (Figure 3).

#### Discussion

In a large case-control study of 996 severe malaria case children and 1220 controls, we found that in Kilifi, Kenya, the risk of severe malaria is clearly associated with the common Hp variants. The highest risk for severe malaria was found in the Hp2-2 genotype and...
the lowest risk in the Hp2-1 genotype. This finding was supported by a meta-analysis of this and 2 earlier studies that suggested an increased risk of severe malaria in Hp2-2 compared with Hp2-1 individuals. Hp1-1 individuals may have had an increased risk of severe malaria compared with Hp2-1 individuals, but this was not supported on meta-analysis. In subgroup analyses, we found that the increased risk of severe malaria among Hp2-2 subjects was seen in all the severe malaria syndromes, although this did not reach statistical significance in the symptomatic SMA group. Finally, we found evidence to suggest that the effect of Hp on the risk of severe malaria is modified by a1 thalassemia.

Our finding of an increased risk of severe malaria among Hp2-2 individuals is in apparent contradiction to 2 earlier studies conducted in Sudan and Ghana,15,16 both of which reported a reduced risk of severe malaria among children of the Hp2-2 phenotype. This might be explained by the fact that in these previous studies Hp type was based on phenotyping rather than genotyping, because Hp2-2 is present at lower plasma concentrations and is more likely to be depleted during malaria-related hemolysis and therefore lead to biased misclassification. Similarly, our findings are not in complete agreement with 2 studies conducted by genotyping. In 1 study, conducted in Northern Ghana, the authors found evidence for a marginally increased risk of severe malaria in Hp2-2 individuals,18 whereas in a second study, conducted in The Gambia, the authors found no association between Hp genotype and severe malaria risk.17 It is possible that the clinical characteristics of case patients included in these studies may have differed from those included in our current study. In our study, we did not include children presenting with uncomplicated severe malaria anemia because this condition is not associated with significant mortality.30 Differences in ethnicity and genetic diversity may be another explanation. Finally, the varying observations made in different populations might also be explained

<table>
<thead>
<tr>
<th>α1 Thalassemia</th>
<th>Hp</th>
<th>Severe malaria (n)</th>
<th>Controls (n)</th>
<th>OR (95% CI)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>aa/aa</td>
<td>Hp1-1</td>
<td>126</td>
<td>133</td>
<td>1.0 (Reference)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Hp2-1</td>
<td>170</td>
<td>188</td>
<td>0.93 (0.67-1.30)</td>
<td>0.70</td>
</tr>
<tr>
<td></td>
<td>Hp2-2</td>
<td>102</td>
<td>121</td>
<td>0.89 (0.61-1.30)</td>
<td>0.55</td>
</tr>
<tr>
<td>α/α and -α/-α</td>
<td>Hp1-1</td>
<td>186</td>
<td>228</td>
<td>0.87 (0.63-1.20)</td>
<td>0.39</td>
</tr>
<tr>
<td></td>
<td>Hp2-1</td>
<td>253</td>
<td>402</td>
<td>0.63 (0.47-0.86)</td>
<td>0.003</td>
</tr>
<tr>
<td></td>
<td>Hp2-2</td>
<td>159</td>
<td>148</td>
<td>1.10 (0.78-1.55)</td>
<td>0.60</td>
</tr>
</tbody>
</table>

ORs were derived by logistic regression analyses adjusted for gender, ethnicity, and sickle status. Hp1-1 inherited with aa/aa was the reference type with an OR for severe malaria of 1.0.
by epistasis, which holds that the effect of a genotype at a particular locus depends on the genotype co-inherited at a second unrelated locus. Epistasis is likely to be more common than previously thought.

We looked for possible epistasis between Hp genotype and 2 known malaria-resistance genes: HbAS, which results from a structural abnormality of the β-globin chains of Hb, and α1 thalassemia, which results from the underproduction of normal α-globin chains of Hb. The number of children with HbAS and severe malaria were few (n = 22), and our study was underpowered to investigate any potential epistasis between HbAS and Hp genotype. However, we did find evidence for an epistatic interaction between α1 thalassemia and Hp genotype (P = .02). Our data suggest that α1 thalassemia inherited in combination with Hp-2 is strongly protective against severe malaria (37% protection, P = .003) but that protection is considerably reduced when inherited in combination with Hp-1 (13% protection, P = .39) and is lost altogether when inherited in combination with Hp-2 (Table 4). Similarly, although the Hp genotype inherited in combination with α1 thalassemia significantly altered the risk of severe malaria, the Hp genotype in the absence of α1 thalassemia did not appear to alter risk in subgroup analyses (Table 4). Although this possible epistasis should be viewed with caution, it seems plausible that varying frequencies of α1 thalassemia across Africa might explain apparent differences in the results of previous studies investigating the association between Hp type and severe malaria. For example, the frequency of the α2 allele, the most common cause of α1 thalassemia in sub-Saharan Africa, is 0.12 in The Gambia, 0.33 in Ghana, and 0.40 in our population. Varying HP2 allele frequencies could similarly explain differing findings for α1 thalassemia association studies.

If epistasis exists between the common Hp variants and α1 thalassemia, by what mechanism might it act? Despite high population frequencies of α1 thalassemia and the HP2 allele in many malaria-endemic areas, little is known about the pathophysiological consequences of their coinheritance. A number of mechanisms for an epistatic interaction seem plausible, a few of which are summarized here. Individuals with α1 thalassemia are under increased oxidative stress, a fact that likely relates to an intraerythrocytic excess of unmatched β chains and reactive free thiols. Hp-2, unlike Hp-1 and Hp-2, is significantly less able to quench Hb-iron mediated oxidant stress, which may account for reduced ferroxidase and vitamin C levels in Hp-2 individuals. Additionally, methemoglobin, an endothelial cell activator, is increased in thalassemia and may be further increased in Hp-2 individuals following hemolysis.
due to an oxidatively unstable heme moiety in the Hp2-2–Hb complex. The Hp2-2 variant is further associated with a shift toward a proinflammatory Th1 cytokine response. Thus, it is possible that the protection conferred by α+ thalassemia is lost when coherited with Hp2-2 due to excessive oxidant damage, inflammation, and endothelial cell activation, especially within the brain and deep tissues where the large Hp2-2 protein is less able to penetrate. Indeed, subarachnoid hemorrhage in patients with the Hp2-2 genotype results in more severe vasospasm, tissue ischemia, and inflammation. Additionally, the reduced rosetting and endothelial cell adherence observed in α+ thalassemia may be countered by the up-regulation of vascular cell adhesion molecules and endothelial cell dysfunction observed in Hp2-2 individuals, resulting in loss of protection in individuals coheriting both genotypes.

Conversely, Hp-2 coherited with α+ thalassemia appeared to confer significant protection from severe malaria indicating heterozygote advantage. In addition to reduced rosetting, infected α+ thalassemia red cell membranes bind significantly more malaria immune globin and are more susceptible to phagocytosis compared with control cells, a process likely mediated by oxidative damage to the red cell membrane from unbound β chains and accelerated acquisition of aggregated band 3 proteins. It is possible that the Hp2-1 variant optimizes the protective effect of α+ thalassemia by providing a balanced plasma environment with optimal oxidant/antioxidant levels and Th1 vs Th2 cytokines compared with Hp1-1 and Hp2-2. Hp2-1 has a distinct molecular structure compared with the other variants; for example, Vibrio vulnificus is unable to acquire iron from Hp2-1–Hb complexes, despite iron acquisition from Hp1-1–Hb and Hp2-2–Hb complexes. Heterozygote advantage from Hp2-1 is further observed with protection from Kaposi’s sarcoma and reduced mortality in HIV patients. A similar trend toward protection was seen in individuals coheriting Hp1-1 and α+ thalassemia, although this did not reach statistical significance. Epistasis may provide valuable insights into the mechanisms by which malaria resistance genes provide protection.

We did not find an interaction between the common Hp variants and the −61C allele or any influence of the −61C allele on the risk of severe malaria. It might be expected that low Hp concentrations associated with the −61C allele would expose children to the deleterious effects of cell-free Hb and therefore predispose such children to severe malaria. However, in our previous study of children with uncomplicated malaria, although the −61C allele was associated with reduced Hp concentrations during convalescence when malaria parasites had been cleared, it had no influence on Hp concentration at the time of a malaria episode. Hp is up-regulated during the acute phase response, and this may counter the reduced transcription that might result from the −61C allele outside the acute phase response. Alternatively, it is also possible that the structure or function of the common Hp variants may be more clinically relevant than their concentrations. Our finding differs from that made in our earlier study conducted in The Gambia, in which a reduced incidence of uncomplicated malaria was found only in children >3 years of age carrying the −61C allele. Because this protection was only seen in older children, we proposed that an immune mediated mechanism, possibly mediated by low Hp concentration, oxidative damage to the red cell membrane, and accelerated acquisition of aggregated band 3 proteins or by other immune modulation, might have been involved. In the current study, the majority of children were <3 years of age and as such will have had less time to acquire immunity. The patient mix was also different between the studies. Children in the current study had severe life-threatening malaria requiring high dependency care in hospital, whereas the previous study involved children in the community with a mild febrile episode associated with malaria parasites on blood film. Moreover, although Hp concentration at convalescence might influence acquisition of immunity following uncomplicated malaria, Hp concentration at the time of malaria may be more critically important in severe malaria. It also seems possible that the apparently conflicting findings between these 2 studies could be explained by a range of other genetic, ethnic, or environmental differences between the children involved. We also previously reported a lower incidence of uncomplicated clinical malaria in older children (>3 years) carrying the Hp2-2 genotype, postulating that this might be mediated by oxidative damage to the red cell membrane and accelerated acquisition of antibodies to aggregated band 3 proteins. Similarly, in the current study, the majority of children were <3 years old and thus would have had less time to acquire immunity.

In summary, this large case control study and meta-analysis suggest that the common Hp 2-2 genotype confers an increased risk of severe malaria and that the Hp2-1 genotype is protective. Moreover, our data are compatible with an epistatic interaction with regard to susceptibility to severe malaria between Hp and α+ thalassemia. It seems plausible that heterozygote advantage, balancing selection pressure, and epistasis might explain the varying gene frequencies of the Hp1 allele and α+ thalassemia in different populations and explain why the relationship between malaria and Hp has proved difficult to confirm. Further studies investigating these associations in other malaria-endemic populations are indicated.

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Authorship

Contribution: S.H.A. conceived the study, performed experiments, analyzed data, and wrote the manuscript; S.M.U. performed experiments, analyzed data, and contributed to writing the manuscript; E.N. and A.W.M. performed experiments and contributed to data analysis; G.N. and C.N. managed databases and analyzed data; D.P.K. conceived the study and performed experiments; K.A.R. performed experiments, contributed to data analysis, and wrote the manuscript;
and T.N.W. conceived the study, obtained funding, contributed to data analysis, and wrote the manuscript.

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References


Epistasis between the haptoglobin common variant and α⁺thalassemia influences risk of severe malaria in Kenyan children

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