

Iron Status Predicts Malaria Risk in Malawian Preschool Children

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Abstract

Introduction: Iron deficiency is highly prevalent in pre-school children in developing countries and an important health problem in sub-Saharan Africa. A debate exists on the possible protective effect of iron deficiency against malaria and other infections; yet consensus is lacking due to limited data. Recent studies have focused on the risks of iron supplementation but the effect of an individual's iron status on malaria risk remains unclear. Studies of iron status in areas with a high burden of infections often are exposed to bias. The aim of this study was to assess the predictive value of baseline iron status for malaria risk explicitly taking potential biases into account.

Methods and materials: We prospectively assessed the relationship between baseline iron deficiency (serum ferritin <30 µg/L) and malaria risk in a cohort of 727 Malawian preschool children during a year of follow-up. Data were analyzed using marginal structural Cox regression models and confounders were selected using causal graph theory. Sensitivity of results to bias resulting from misclassification of iron status by concurrent inflammation and to bias from unmeasured confounding were assessed using modern causal inference methods.

Results and Conclusions: The overall incidence of malaria parasitemia and clinical malaria was 1.9 (95% CI 1.8–2.0) and 0.7 (95% CI 0.6–0.8) events per person-year, respectively. Children with iron deficiency at baseline had a lower incidence of malaria parasitemia and clinical malaria during a year of follow-up; adjusted hazard ratio's 0.55 (95%-CI:0.41–0.74) and 0.49 (95%-CI:0.33–0.73), respectively. Our results suggest that iron deficiency protects against malaria parasitemia and clinical malaria in young children. Therefore the clinical importance of treating iron deficiency in a pre-school child should be weighed carefully against potential harms. In malaria endemic areas treatment of iron deficiency in children requires sustained prevention of malaria.

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Introduction

Childhood iron deficiency is prevalent globally with highest estimates in children in developing countries. In sub-Saharan Africa, 33–66% of children are iron deficient, most of whom are under the age of five years [1,2]. The high prevalence of iron deficiency in these populations is a consequence of low nutritional iron intake due to low economic status [2], decreased absorption of dietary iron due to a high prevalence of infections [3] or intestinal blood loss in the case of helminth infections [4]. Iron deficiency causes anemia and developmental dysfunction [2,3] and the current consensus is to treat all iron deficient children [5]. Conversely, evidence of a protective effect of iron deficiency against malaria and other infections exists [6–12].

In recent years concerns have focused mainly on the possible harmful effects of treating iron deficiency rather than on the effects

of iron status itself [13–19]. A recent Cochrane meta-analysis reviewing iron supplementation in children in malaria endemic areas concluded that iron supplementation did not adversely effect children in malaria endemic areas when adequate malaria surveillance is provided, although the authors considered their data were limited by various factors including lack of data on baseline iron status [20]. There is little conclusive data on the protective effect of iron deficiency against malaria. The reliability of current data on iron status is limited since available iron biomarkers are affected by inflammation which makes interpretation of results from malaria endemic areas problematic [21]. In addition, the complexity of factors influencing both malaria susceptibility and iron status are not always taken into account, potentially leading to confounded results. To improve control and management of malaria as well of iron deficiency, reliable data about the influence of iron status on malaria risk are needed. The

aim of this study was to assess the predictive value of baseline iron status in children for subsequent malaria risk, explicitly taking misclassification of iron deficiency due to concurrent inflammation and confounding bias into account by utilizing modern causal inference methods in the analyses of a dataset of Malawian pre-school children.

Methods

Study Design

This is a secondary analysis of follow-up data from a cohort of pre-school children recruited as the control group for the *Severe Anemia Study (SevAna study)*, a large case-control study investigating the etiology, pathophysiology and outcome of severe anemia in southern Malawi [22–24]. The analysis is restricted to the non-severely anemic control population (hemoglobin level ≥ 5.0 g/dL), because the cases all received a blood transfusion. Follow-up data of the first year after enrolment were used, allowing the observation of a full year of malaria transmission per child.

Ethical Statement

The *SevAna study* was approved by the Ethics Committees of the College of Medicine, Malawi (ref. no P.00/01/116, and the Liverpool School of Tropical Medicine, United Kingdom (ref. no 01.29). Prior to enrolment in the study, written informed consent was obtained from the parent or guardian of all study participants.

Study population

The *SevAna study* recruited 381 cases and 757 controls between July 2002 and July 2004 (Figure S1). Every child (6–60 months old) presenting at the hospital with severe anemia (hemoglobin < 5.0 g/dL) was defined as a case. For each case, two controls were enrolled; a community control living within 100–1000 meter of the case, and a hospital control, which was the first child presenting at the outpatient department on the day following presentation of the case patient. Controls were eligible for recruitment if aged 6–60 months and if their hemoglobin level was at least 5.0 g/dL. All children were visited at 1, 3, 6 and 12 months from date of recruitment (active follow-up). If children did not report for follow up, families were visited by the study team. In addition, guardians were asked to present their child to a study-clinic whenever the child was sick (passive follow-up).

Study setting

The *SevAna study* took place in Malawi, a land-locked country in the south-eastern part of Africa of which 20% is covered by Lake Malawi. The climate is tropical, but prevalence of malaria and other infectious diseases vary with proximity to the lake and altitude. Rough estimates of the prevalence of iron deficiency vary from 20–40% [2,25]. Study participants were recruited in two settings; Blantyre district (study site: Queen Elizabeth Central Hospital) and Chikwawa District (study site: Chikwawa District Hospital), both in the southern region of Malawi. Blantyre is the main commercial town of Malawi with a predominantly urban population of half a million. At an altitude of 800 m above sea level malaria is mainly seasonal (approximately 1 infectious bite per person per year) [26]. Chikwawa District Hospital, which caters for a predominantly rural population of approximately 400 thousand people, is situated in the lower Sire Valley, 50 km south of Blantyre. With an altitude of 250 m above sea level malaria transmission is year round (approximately 170 infectious bites per person per year) (Milahowa T, personal communication).

Clinical procedures

At recruitment a detailed medical and socio-economic history was recorded and a physical examination was performed. Samples of blood, stool and urine were collected. Children were treated if indicated using local treatment guidelines. Clinical malaria was defined as a positive blood slide with concurrent fever (axillary temp $> 37.5^{\circ}\text{C}$) or history of fever (caregiver recall of fever in the last week). Severe malaria was defined as a positive blood slide with either severe anemia (hemoglobin < 5.0 g/dl) or coma [27]. Malaria was treated with sulfadoxine-pyrimethamine (SP) and if the child was unable to take oral medication parenteral quinine was administered. As a standard procedure at recruitment all study participants received presumptive malaria treatment (25.0/1.25 mg/kg SP) and according to local guidelines all children received iron supplementation 2 mg/kg/day for 28 days. Follow-up procedures consisted of a medical history, physical examination and a blood sample to determine hemoglobin and malaria parasitemia, and if indicated children were treated using local treatment guidelines. Deaths were recorded and if they occurred outside the study clinics they were investigated as completely as possible using a verbal autopsy form.

Laboratory procedures

Hemoglobin (Hb) concentrations were measured by HemoCue B-Hemoglobin analyzer (HemoCue, Ångelholm, Sweden) to judge eligibility-criteria. Hb measured by Coulter counter analyzer (Beckman Coulter, Durban, South Africa) was used for statistical analyses. Moderate and severe anemia were defined as an Hb of < 11.0 g/dL and < 5.0 g/dL, respectively. Plasma levels of C-reactive protein (CRP) and serum ferritin were analyzed on Modular P800 and Monular Analytic E170 systems (Roche Diagnostics, Switzerland). Iron deficiency was defined as serum ferritin < 30 $\mu\text{g/L}$ [25]. Inflammation was defined as $\text{CRP} > 10$ g/L. For malaria diagnoses thin-film blood slides were read by two independent readers; with a third reader being used if results differed by 25% or more. The number of *Plasmodium falciparum* asexual parasites per 200 white cells was counted and expressed as the number per microliter of blood. Malaria parasitemia was defined as the presence of one or more *Pl. falciparum* asexual parasites [28]. Human immunodeficiency virus (HIV) infection was assessed using two rapid tests (Determine, Abbott Laboratories; and Uni-Gold, Trinity Biotech). Discordant or reactive rapid-test results in children less than 18 months of age were resolved by PCR [29].

Statistical Analyses

The incidence of clinical malaria and parasitemia was calculated for a follow-up period of one year. Multiple events per child were registered. A child was considered not to be at risk for incident malaria for the 28 days following malaria treatment (SP), as the prophylactic effect of malaria treatment was assumed [30]. This 28 day period was excluded as person-time and malaria events during this period were not counted for analyses. Malaria parasitemia or clinical malaria occurring during this period were defined as treatment-failure and were excluded from analysis.

The main predictor, iron deficiency, was defined as serum ferritin < 30 $\mu\text{g/L}$. Possible confounders considered for inclusion were: sex, age, area of residence, season, socio economic status, genetic predisposition of malaria, cellular immunity, humoral immunity, cumulative malaria exposure, malaria immunity, history of other infections than malaria, HIV infection, hookworm infection, nutritional status and zinc deficiency (figure 1). We used DAGitty[®], a web-based application based on the *Directed Acyclic Graph theory*, to model the causal relationship between iron status,

malaria risk and any potential (measured and unmeasured) confounders [31,32]. This DAG-program was used to identify sets of confounders that together fully adjust for confounding in multivariable modeling [31,33,34]. Several of these sets of confounders (*minimally sufficient adjustment sets*) were identified from figure 1. The set containing HIV-status, socio economic status, age, nutritional status and study site was used for analysis, as these variables were available in our data set. Socio economic status was scored on parents' education (1-4), job (1-4) and number of assets (0-6). Chronic malnutrition defined as height-for-age <-2 SD was used as the marker for nutritional status [35].

Due to inadequate sample volume in 29% of the children serum ferritin was not available. The missing data were assumed to be missing at random; basic characteristics between children with and without available ferritin assessments were comparable (Table S1). Because complete-cases analysis with a large amount of missing values may lead to loss of efficiency and bias [36], missing data were imputed using multiple imputation by chained equations in the R software version 2.14.0 utilizing the MICE package [37,38]. The imputation model included all baseline covariates in table 1 plus soluble transferrin receptor (sTfR) and mean corpuscular hemoglobin concentration (MCHC) as additional markers of iron

status. The outcome variable was included in the imputation model via the event indicator and the Nelson-Aalen estimator of baseline hazard, as studies suggest that these specific components lead to lower bias in estimation [39]. Regression estimates and their standard errors from analyses of the 25 separate imputed data sets were summarized using Rubin's rules [40].

To compare the incidence of malaria during follow-up between iron deficient and iron replete individuals, a marginal structural Cox proportional hazards model was used to adjust for confounding. Marginal structural models are a new class of statistical models to estimate causal effects [41,42], and are equivalent to non-parametric multivariate standardization methods [41,43,44]. Effect estimates can be adjusted for measured confounders using inverse probability of exposure weighting. In addition, marginal structural models allow the quantitative evaluation of unmeasured confounding [42,45,46]. Stabilized inverse probability weights were estimated using the 'ipw' package in R [30,47]. To adjust for the correlation between recurrent malaria events in a single person, robust standard errors were computed. All covariates were treated as time-independent. The proportional hazards assumption was checked by plotting the scaled *Schoenfeld* residuals against time and tested using *cox.zph*

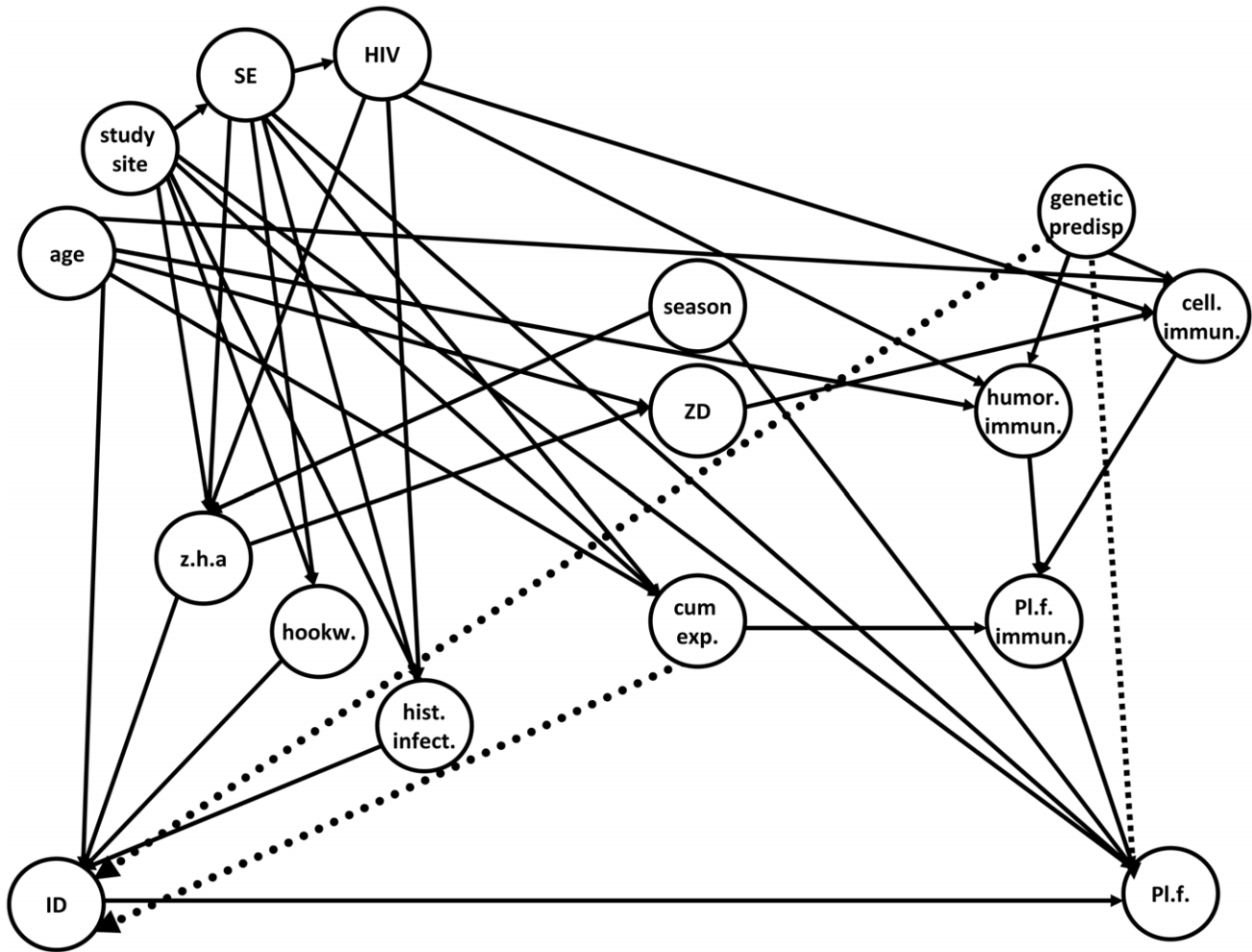


Figure 1. Directed acyclic graph for the relation between iron deficiency and malaria risk. cell. immun: cellular immunity; cum exp: cumulative malaria exposure; genetic predisp: genetic predisposition; humor immun: humoral immunity; hookkw: hookworm infection; hist infect: history of other infections than malaria; HIV: human immunodeficiency virus; ID: iron deficiency; Pl.f.immun: immunity for *Pl.falciparum*; *Pl.f*: *Plasmodium falciparum* infection; SE: socio economic score; ZD: zinc deficiency; z.h.a: z-score height for age. doi:10.1371/journal.pone.0042670.g001

Table 1. Baseline characteristics of the study population.

Characteristic	Result
Female	357/727 (49.1%)
Age in months: mean (s.d)	23.9 (12.6)
Age<2 years	408/727 (56.1%)
Living in an urban area	376/727 (51.7%)
SE-score: mean (s.d)	7.2 (2.1)
Hospital control	357/727 (49.1%)
Hemoglobin: mean (s.d)	9.7 (2.1)
Iron deficiency	146/513 (28.5%)
Malnourished	275/669 (41.1%)
Fever (>37.5°C axillary)	205/727 (28.4%)
CRP>10 mg/L	373/630 (59.2%)
Malaria parasitemia	308/720 (42.8%)
Clinical malaria	108/720 (14.9%)
HIV-infected	39/670 (5.8%)

Hemoglobin in g/dl; CRP: C-Reactive Protein; malnourished defined as height-for-age <-2 SD. HIV: Human Immunodeficiency Virus. Iron deficiency defined as serum ferritin <30 ug/L. SE-score: Socio Economic Score, a sum of the following scores: parents' education (1-4), job parents (1-4) and number of assets (0-6).

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procedure in R [48]. We assessed whether the effect of iron deficiency on malaria was modified by age or study site by testing their interaction terms with the Wald statistic. Interaction terms were considered significant if $p < 0.1$.

We assessed to which extent the outcomes could have been biased because of possible misclassification of the iron deficiency indicator (serum ferritin), by reanalyzing the data using a more sensitive and a more specific definition of iron deficiency [49]; cut-offs for serum ferritin that varied by inflammation status; children without inflammation (CRP<10 g/L) were defined as iron deficient having a serum ferritin of <12 µg/L; for children with current inflammation (CRP>10 g/L) serum ferritin cut-offs of 30 and 70 µg/L were explored [25,50].

In addition, to evaluate how large the average difference in susceptibility to malaria at baseline between the iron deficient and iron-replete group would need to have been to partly or fully explain our observed differences in malaria incidence, we used a method for sensitivity analysis for unmeasured confounding developed by *Robins et al.* [42,45,46]. The method requires that unmeasured confounding is quantified as a bias parameter $u = \alpha^{(2 \times ID - 1)}$, with ID = 1 or 0 as indicator of iron deficiency and α the degree of unmeasured confounding or, in other words, the average difference in malaria risk at baseline on a ratio-scale and conditional on the covariate distribution.

Results

Study population

Follow-up data were available for 96.0% of all study participants (727/757). Baseline characteristics are presented in table 1. Baseline characteristics of children with and without follow up data were comparable (data not shown). Of the remaining 727 children the attendance at active follow-up visits ranged from 86–89%.

Incidence of malaria parasitemia and clinical malaria

During the follow-up year the overall incidence of malaria parasitemia and clinical malaria was 1.9 (95% CI 1.8–2.0) and 0.7 (95% CI 0.6–0.8), respectively. This incidence was higher in the rural areas and in the children less than 24 months of age (Table 2). There were four cases of severe malaria, three of which were iron replete. During follow-up 0.7% (1/141) of the iron deficient children and 1.3% (5/372) of iron replete children died.

Baseline iron status and incidence of malaria

The marginal structural Cox model showed that iron deficiency was an independent predictor of malaria parasitemia and clinical malaria; HR (95%CI) 0.55 (0.41–0.74) and 0.49 (0.33–0.73) respectively (Table 3). The effect of iron deficiency on malaria parasitemia and clinical malaria was consistent across different sub-populations; no significant effect measure modification by study site (urban vs. rural) or age (linear term) was seen (all p -values>0.9). In addition, the effects were sustained throughout the study period; the proportional hazards assumption was not refuted and when analysis was done on consecutive periods of 4 months, consistent effect measures were seen (data not shown). Analyses conducted with the complete cases data set (including only children with a ferritin result) showed comparable results (HR (95%CI) 0.61 (0.44–0.81) and HR (95%CI) 0.53 (0.36–0.78) for malaria parasitemia and clinical malaria respectively).

Sensitivity analyses

Analyses assessing sensitivity to misclassification of iron status are presented in Table 3 and show consistent results using different definitions of iron deficiency. Hazard ratio estimates for clinical malaria from the marginal structural Cox model including adjustment for varying amounts of potential unmeasured confounding are presented in Figure 2. Two distinct distributions of unmeasured confounding were considered. Figure 2a presents results from analyses in which the net amount of bias per group is specified. The a priori clinical malaria risk in the iron deficient group should have been at least 0.72 times the a priori risk in the iron replete group to remove significance of our findings, and 0.48 times to fully explain our findings (Figure 2a). Figure 2b presents results from analyses in which the amount of unmeasured confounding varied as function of age; increasing log-linearly up to a peak at the age of 3 years and then decreasing to no confounding effect. The ratio of priori risks would have to have been at least 0.49 at its peak at 3 years-of-age to remove significance and 0.18 to fully explain our finding (Figure 2b). Baseline malaria is often included as a proximate measure of susceptibility to malaria in multivariable analyses. However, since our hypothesis was that baseline iron status influences malaria risk, baseline malaria would technically not be a confounder, i.e. the exposure may not affect the confounder (Causal knowledge as a prerequisite for confounding evaluation [33]). We therefore chose to assess the effect of differences in malaria susceptibility at baseline through sensitivity analysis. Notwithstanding, when baseline malaria parasitemia respectively clinical malaria was included, effect estimates were slightly attenuated though still significant HR (95%CI) 0.66 (0.48–0.91) and HR (95%CI) 0.51 (0.33–0.79) for malaria parasitemia and clinical malaria respectively.

Discussion

In this study, children with iron deficiency at baseline (serum ferritin <30 µg/L) had a lower incidence of malaria parasitemia and clinical malaria during a follow up period of one year. This

Table 2. Incidence of malaria parasitemia and clinical malaria per person year.

		All		Iron deficient group		Iron replete group	
		Events	Incidence	Events	Incidence	Events	Incidence
Malaria parasitemia	All	896	1,9	180	1,3	716	2,2
	Urban	253	1,0	38	0,5	215	1,1
	Rural	643	3,1	142	2,2	501	3,6
	Age<24 months	557	2,1	139	1,4	418	2,5
	Age≥24 months	339	1,6	41	0,9	298	1,8
Clinical malaria	All	318	0,7	58	0,4	260	0,8
	Urban	65	0,2	11	0,1	54	0,3
	Rural	253	1,2	47	0,7	206	1,5
	Age<24 months	207	0,8	44	0,5	163	1,0
	Age≥24 months	111	0,5	14	0,3	97	0,6

Incidence of malaria parasitemia and clinical malaria per person year stratified per area, age group and iron status. Malaria parasitemia defined as a positive malaria blood slide; Clinical malaria: positive malaria blood slide with concurrent fever (axillary temp >37.5°C) or history of fever.

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finding suggests that iron deficiency protects against malaria parasitemia and clinical malaria in pre-school children living in malaria endemic areas. The results were consistent across study sites with varying transmission intensity and across age-groups despite changing susceptibility with age. In addition, the difference in risk between baseline iron status groups persisted throughout the follow-up period.

Our findings are in line with results of the few other studies assessing influence of iron status on malaria risk [10,51]. The protective effects of iron deficiency on malaria risk shown in these two studies, were of similar magnitude, yet Gwamaka *et al* also showed an effect on all cause and malaria associated mortality [51].

As with any observational study, the results of this study are subject to bias. Therefore, we explicitly evaluated the potential impact that misclassification and confounding bias could have had on our results. Firstly, because serum ferritin levels are influenced by concurrent inflammation it may not adequately reflect true iron status. As a consequence, misclassification of baseline iron status may have occurred and thus resulted in biased effect measures. We evaluated whether use of a more sensitive and a more specific definition, taking concurrent inflammation into account, would substantially change the results. As expected, use of a more specific definition slightly increased the magnitude of effect and the use of a more sensitive definition led to a slightly weaker effect.

Importantly, this did not change our conclusions, suggesting that the findings are robust to misclassification bias.

In the directed acyclic graph (DAG) in Figure 1 all potential confounders of the effect of iron status on malaria risk as conceptualized by the authors, were evaluated. We considered two additional yet unmeasured variables which could theoretically confound our results by both protecting against malaria and causing iron deficiency. We assessed how strong these potential confounding effects should have been to explain the observed difference in malaria risk between iron status groups. The first unmeasured factor was 'cumulative exposure to malaria'. Recent studies have suggested that chronic infection, including malaria, may reduce iron absorption in the gut by inducing an increase of hepcidin [52–54] and that malaria infection may lead to iron loss through hemosiderin deposition. In addition, repeated malaria infections (cumulative exposure) result in development of immunity to malaria. The age at which clinical immunity is attained is influenced by local infection pressure represented by the number of infectious bites per year (EIR) [55]. If it is true that increased exposure to malaria simultaneously induces iron deficiency and increased malaria immunity, cumulative exposure is a potential confounder in our analyses. In other words, the lower incidence of malaria in the iron deficient group may have been due to higher malaria exposure and subsequently better immunity to malaria, rather than directly due to mechanisms involving iron status. We conducted sensitivity analyses allowing unmeasured confounding

Table 3. Hazard ratios (95% CI) of iron deficiency for the risk on malaria parasitemia en clinical malaria.

Definition iron deficiency	univariate Cox model		marginal-structural Cox model	
	Malaria parasitemia	Clinical malaria	Malaria parasitemia	Clinical malaria
	HR (95% CI)	HR (95% CI)	HR (95% CI)	HR (95% CI)
SF<30	0.59 (0.44–0.78)	0.54 (0.36–0.80)	0.55 (0.41–0.74)	0.49 (0.33–0.73)
SF<30(CRP>10)/SF<12(CRP<10)	0.57 (0.40–0.82)	0.53 (0.32–0.89)	0.46 (0.31–0.71)	0.43 (0.25–0.73)
SF<70(CRP>10)/SF<12(CRP<10)	0.66 (0.52–0.85)	0.59 (0.41–0.84)	0.60 (0.46–0.79)	0.52 (0.35–0.75)

Iron deficiency is defined with different cut-offs for serum ferritin (SF) in ug/L, depending on presence of inflammation, defined as C-reactive protein (CRP) >10 mg/L. The marginal-structural Cox model included HIV-infection, socio economic score, age, nutrition and study site.

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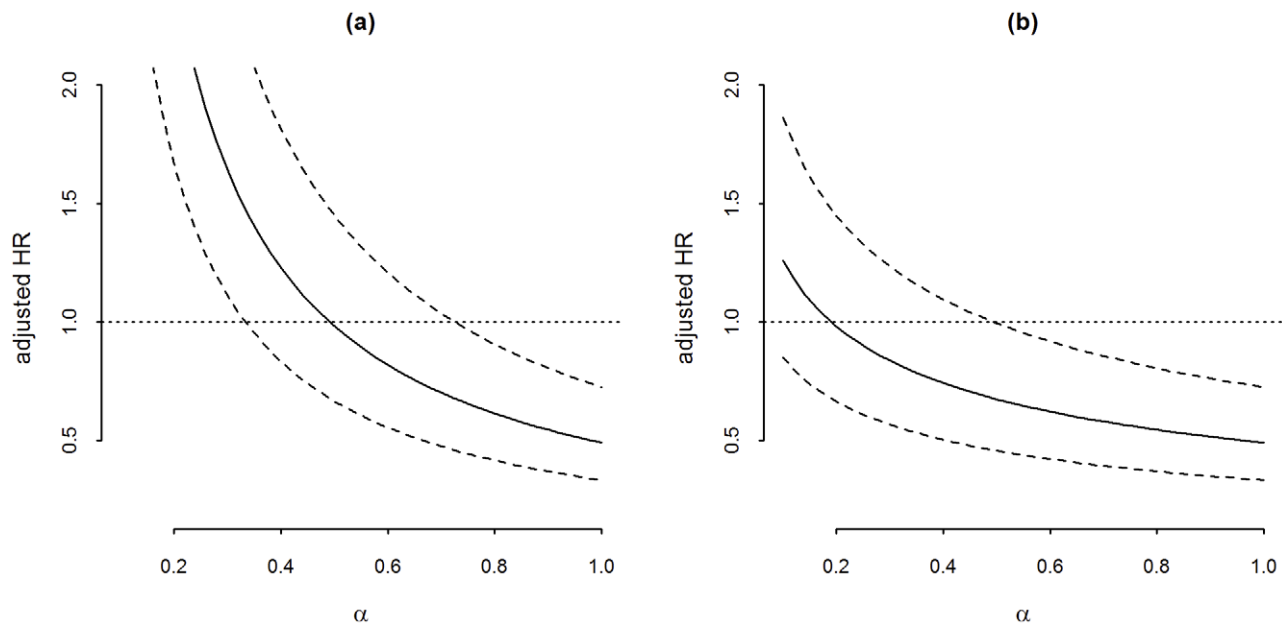


Figure 2. Sensitivity analysis of the impact of unmeasured confounding on the effect of baseline iron deficiency on risk of clinical malaria. The y-axis represents the hazard ratio for malaria infection comparing iron deficiency to iron replete after adjustment for average differences in prior risk represented by α on the x-axis. Figure 2a represents analyses in which net amount of confounding is specified for both iron status groups (bias parameter $u = \alpha^{(2 \times ID - 1)}$). Figure 2b represents analyses in which the amount of confounding was varied as a function of age (bias parameter $u = \alpha^{(2 \times ID - 1) \times (1 - \text{abs}(\text{age} - 36) / 30)}$). doi:10.1371/journal.pone.0042670.g002

bias to vary as a function of age; reflecting the theoretical confounding effect of cumulative exposure, where the difference in immunity (and thus the maximum prior risk difference) gradually changes with increasing age, reaching a peak at the age of 36 months [56] to subsequently decrease after that. The average prior risk of malaria should then have been at least half the risk in the iron replete group at the peak age of 36 months to have led to a spurious significant finding (Figure 2b). Although theoretically a plausible confounder, it is unlikely that in reality ‘cumulative exposure to malaria’ would result in such large differences in prior malaria risk, suggesting that our measured effects are unlikely due to confounding by differences in cumulative exposure. As a second potential confounding factor we considered a (yet undiscovered) genotype simultaneously causing iron deficiency and increasing immunity to malaria, for example a variant in iron metabolism that inhibits maturation of *Plasmodium* in the red cells [57]. Our analysis showed that to be responsible for our statistically significant finding, the average prior risk of malaria in the iron deficient group should have been at least 0.72 times the risk compared to that of the iron replete group (Figure 2a). A genetic predisposition causing a 28% lower prior risk is imaginable (i.e. sickle cell trait) and if such a genotype is discovered this may change the interpretation of our results.

In the interpretation of our analyses we assume that iron status measured at baseline remained relatively stable throughout the year of follow-up. Unfortunately, iron status measurements were not repeated after baseline, so we are not able to check this assumption. It is possible that children changed from the iron deficient group to the iron replete group and vice versa. However, because our study population was closely followed and adequately treated for infections it is more likely that iron deficient children became iron replete than the other way around. The resulting misclassification would have resulted in an underestimation of the effect.

For the underlying mechanism of the protective effect of iron deficiency on susceptibility to malaria infection, several hypotheses have been raised. It is suggested that iron deficiency alters the membrane structure of parasitized erythrocytes which facilitates their elimination by phagocytic cells, yet further research is needed to further support this hypothesis [58]. Another hypothesis is that the increased zinc protoporphyrin in iron deficient parasitized red blood cells binds to heme crystals and thus inhibits the formation of hemozoin, in analogue to antimalarial quinolines [59].

We cannot fully exclude that the observed effect was the result of increased susceptibility in the iron replete group resulting from the month of iron supplementation, rather than a protective effect of iron deficiency. In 2006 a large supplementation trial in Zanzibar was stopped prematurely after increased morbidity and mortality was observed in iron replete children receiving iron and folic acid [19]. However it remains unclear whether the increased morbidity was due to the effect of the iron, or the folic acid component of the supplements [60,61]. Although the month of iron supplementation in this study complicates the interpretation of our study results, it is a reflection of current practice in this such settings in which short courses of iron supplementation are routinely given. In addition we think its influence was minimal for several reasons. Firstly, since the difference in risk was still observable towards the end of follow-up, the influence of the one month of iron supplementation was likely to be negligible. Secondly 59% of the children had an infection at baseline and this was likely to inhibit the absorption of iron in the first two weeks of supplementation [52–54].

In conclusion, iron deficiency seems to protect against malaria parasitemia and clinical malaria. However, because we lack direct measures of baseline immunity to malaria our results should be confirmed by well conducted randomized controlled trials or observational birth cohort studies. Our findings support the theory of nutritional adaptation to infectious diseases [62,63]. To avoid

increased malaria risk due to “neutralizing” the protective effect of iron deficiency, treatment of iron deficiency should occur under simultaneous and sustained control of prevalent infections. We recommend that treatment of iron deficiency occurs within the context of the Integrated Management of Childhood Illness (IMCI) and will benefit from the addition of Integrated Vector Management (IVM) [64,65]. However, because oral iron is often poorly absorbed in individuals with concurrent inflammation [52–54], prevention of iron deficiency may be more effective. Treatment and especially prevention of infections, in particular hookworm infections [4,66], will contribute to prevention of iron deficiency through decreased loss and increased absorption of dietary iron. Whichever strategy is chosen, concurrent and sustained prevention of malaria through Integrated Management of Childhood Illness is imperative in malaria endemic areas.

Supporting Information

Figure S1 Flow chart study population. This flowchart presents number of children enrolled in the main study, a case

control study investigating etiology of severe anemia (I); number of children enrolled in the cohort study (II) and the number of children attending the follow-up visits (III).

(PDF)

Table S1 Baseline characteristics of the study population.

(DOC)

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Author Contributions

Conceived and designed the experiments: MBvH, JJC, KP, BJB. Performed the experiments: JJC, KP, MBvH. Analyzed the data: FAMJ, TL, RG. Contributed reagents/materials/analysis tools: TL, RG. Wrote the paper: FAMJ, TL, RG, JJC, BJB, MBvH. Guarantors of the paper: FAMJ, TL.

References

- Unicef website. Available: www.unicef.org/sowc98/slight5.htm. Accessed 1998.
- World Health Organization website. Available: http://whqlibdoc.who.int/publications/2008/9789241596657_eng.pdf. Accessed 2008.
- Trey JE, Kushner I (1995) The acute phase response and the hematopoietic system: the role of cytokines. *Crit Rev Oncol Hematol* 21: 1–18.
- Jonker FA, Calis JC, Phiri K, Brienen EA, Khoffi H, et al. (2012) Real-time PCR Demonstrates *Ancylostoma duodenale* Is a Key Factor in the Etiology of Severe Anemia and Iron Deficiency in Malawian Pre-school Children. *PLoS Negl Trop Dis* 6: e1555. 10.1371/journal.pntd.0001555 [doi];PNTD-D-11-00973 [pii].
- WHO (2007) Conclusions and recommendations of the WHO Consultation on prevention and control of iron deficiency in infants and young children in malaria endemic areas. World Health Organization website Available: <http://www.who.int/nutrition/publications/micronutrients/FNBvol28N4supdec07.pdf>. Accessed 2006 Jun 01.
- Franchini M, Targher G, Capra F, Montagnana M, Lippi G (2008) The effect of iron depletion on chronic hepatitis C virus infection. *Hepatology* 47: 335–340. 10.1007/s12072-008-9076-z [doi].
- Gangaidzo IT, Moyo VM, Mvundura E, Aggrey G, Murphree NL, et al. (2001) Association of pulmonary tuberculosis with increased dietary iron. *J Infect Dis* 184: 936–939. 10.1086/323203 [doi].
- Murakawa H, Bland CE, Willis WT, Dallman PR (1987) Iron deficiency and neutrophil function: different rates of correction of the depressions in oxidative burst and myeloperoxidase activity after iron treatment. *Blood* 69: 1464–1468.
- Murray MJ, Murray AB, Murray MB, Murray CJ (1978) The adverse effect of iron repletion on the course of certain infections. *Br Med J* 2: 1113–1115.
- Nyakeriga AM, Troye-Blomberg M, Dorfman JR, Alexander ND, Back R, et al. (2004) Iron deficiency and malaria among children living on the coast of Kenya. *J Infect Dis* 190: 439–447. 10.1086/422331 [doi];JID32289 [pii].
- Weinberg ED (1996) Iron withholding: a defense against viral infections. *Biomaterials* 17: 393–399.
- Kabyemela ER, Fried M, Kurtis JD, Mutabingwa TK, Duffy PE (2008) Decreased susceptibility to *Plasmodium falciparum* infection in pregnant women with iron deficiency. *J Infect Dis* 198: 163–166. 10.1086/589512 [doi].
- Doherty CP, Weaver LT, Prentice AM (2002) Micronutrient supplementation and infection: a double-edged sword? *J Pediatr Gastroenterol Nutr* 34: 346–352.
- Iron Deficiency Project Advisory Service Website (Document Number: 163660) Available: www.idpas.org/pdf/330SafetyOfIron.pdf. Accessed 1999.
- Mebrahtu T, Stoltzfus RJ, Chwaya HM, Jape JK, Savioli L, et al. (2004) Low-dose daily iron supplementation for 12 months does not increase the prevalence of malarial infection or density of parasites in young Zanzibari children. *J Nutr* 134: 3037–3041. 134/11/3037 [pii].
- Oppenheimer SJ (2001) Iron and its relation to immunity and infectious disease. *J Nutr* 131: 616S–633S.
- Iannotti LL, Tielsch JM, Black MM, Black RE (2006) Iron supplementation in early childhood: health benefits and risks. *Am J Clin Nutr* 84: 1261–1276. 84/6/1261 [pii].
- Richard SA, Zavaleta N, Caulfield LE, Black RE, Witzig RS, et al. (2006) Zinc and iron supplementation and malaria, diarrhea, and respiratory infections in children in the Peruvian Amazon. *Am J Trop Med Hyg* 75: 126–132. 75/1/126 [pii].
- Sazawal S, Black RE, Ramsan M, Chwaya HM, Stoltzfus RJ, et al. (2006) Effects of routine prophylactic supplementation with iron and folic acid on admission to hospital and mortality in preschool children in a high malaria transmission setting: community-based, randomised, placebo-controlled trial. *Lancet* 367: 133–143. S0140-6736(06)67962-2 [pii];10.1016/S0140-6736(06)67962-2 [doi].
- Okebe JU, Yahav D, Shbita R, Paul M (2011) Oral iron supplements for children in malaria-endemic areas. *Cochrane Database Syst Rev* CD006589. 10.1002/14651858.CD006589.pub3 [doi].
- Beesley R, Filteau S, Tomkins A, Doherty T, Ayles H, et al. (2000) Impact of acute malaria on plasma concentrations of transferrin receptors. *Trans R Soc Trop Med Hyg* 94: 295–298.
- Calis JC, Phiri KS, Faragher EB, Brabin BJ, Bates I, et al. (2008) Severe anemia in Malawian children. *N Engl J Med* 358: 888–899. 358/9/888 [pii];10.1056/NEJMoa072727 [doi].
- Phiri KS, Calis JC, Kachala D, Borgstein E, Waluza J, et al. (2009) Improved method for assessing iron stores in the bone marrow. *J Clin Pathol* 62: 685–689. 62/8/685 [pii];10.1136/jcp.2009.064451 [doi].
- Phiri KS, Calis JC, Siyasiya A, Bates I, Brabin B, et al. (2009) New cut-off values for ferritin and soluble transferrin receptor for the assessment of iron deficiency in children in a high infection pressure area. *J Clin Pathol* 62: 1103–1106. 62/12/1103 [pii];10.1136/jcp.2009.066498 [doi].
- World Health Organization website Available: http://www.who.int/nutrition/publications/micronutrients/anaemia_iron_deficiency/9789241596107/en/index.html. Accessed 2007.
- Slutsker L, Taylor TE, Wirima JJ, Steketee RW (1994) In-hospital morbidity and mortality due to malaria-associated severe anaemia in two areas of Malawi with different patterns of malaria infection. *Trans R Soc Trop Med Hyg* 88: 548–551.
- (2000) Management of the child with a serious infection or severe malnutrition; guidelines for care at the first referral level in developing countries, WHO department of child and adolescent health and development, 2000 Hong-kong/ geneva. ISBN 9241545313. 57–61. WHO/FCH/COSH/00.1.
- Sullivan AD, Ittarat I, Meshnick SR (1996) Patterns of haemozoin accumulation in tissue. *Parasitology* 112 (Pt 3): 285–294.
- Molyneux EM, Tembo M, Kayira K, Bwanaisa L, Mwenyichanya J, et al. (2003) The effect of HIV infection on paediatric bacterial meningitis in Blantyre, Malawi. *Arch Dis Child* 88: 1112–1118.
- World Health Organization website. Available: www.who.int/malaria/publications/atoz/9789241500470/en/index.html. Accessed 2010.
- Pearl J (2000) *The art and science of cause and effect*, Cambridge University. 331–358.
- Textor J, Hardt J, Knuppel S (2011) DAGitty: a graphical tool for analyzing causal diagrams. *Epidemiology* 22: 745. 10.1097/EDE.0b013e318225c2be [doi];00001648-201109000-00022 [pii].
- Hernan MA, Hernandez-Diaz S, Werler MM, Mitchell AA (2002) Causal knowledge as a prerequisite for confounding evaluation: an application to birth defects epidemiology. *Am J Epidemiol* 155: 176–184.
- Shrier I, Platt RW (2008) Reducing bias through directed acyclic graphs. *BMC Med Res Methodol* 8: 70. 1471-2288-8-70 [pii];10.1186/1471-2288-8-70 [doi].
- Gorstein J, Sullivan K, Yip R, de Onis M, Trowbridge F, et al. (1986) Use and interpretation of anthropometric indicators of nutritional status. *Bulletin of the World Health Organization*, 64(6):929–941, 1986.
- White IR, Carlin JB (2010) Bias and efficiency of multiple imputation compared with complete-case analysis for missing covariate values. *Stat Med* 29: 2920–2931. 10.1002/sim.3944 [doi].
- Cran-R website (R manuals) Available: <http://cran.r-project.org/manuals.html>. Accessed 2010.

38. van Buuren S, Groothuis-Oudshoorn K (2011) MICE: Multivariate Imputation by Chained Equations.
39. White IR, Royston P (2009) Imputing missing covariate values for the Cox model. *Stat Med* 28: 1982–1998. 10.1002/sim.3618 [doi].
40. Rubin DB (1987) Multiple imputation for nonresponse in surveys. *Handbook of Statistical Modeling for the Social and Behavioral Sciences*. New York: John Wiley & Sons.
41. Hernan MA, Brumback B, Robins JM (2000) Marginal structural models to estimate the causal effect of zidovudine on the survival of HIV-positive men. *Epidemiology* 11: 561–570.
42. Robins JM, Rotnitzky A, Scharfstein DO (1999) Sensitivity analysis for selection bias and unmeasured confounding in missing data and causal inference models. In: Halloran ME, Berry DA, eds. *Statistical Models in Epidemiology*. New York: Springer Verlag. 1–92.
43. Cole SR, Hernan MA (2008) Constructing inverse probability weights for marginal structural models. *Am J Epidemiol* 168: 656–664. kwn164 [pii];10.1093/aje/kwn164 [doi].
44. Sato T, Matsuyama Y (2003) Marginal structural models as a tool for standardization. *Epidemiology* 14: 680–686. 10.1097/01.EDE.0000081989.82616.7d [doi].
45. Brumback BA, Hernan MA, Haneuse SJ, Robins JM (2004) Sensitivity analyses for unmeasured confounding assuming a marginal structural model for repeated measures. *Stat Med* 23: 749–767. 10.1002/sim.1657 [doi].
46. Klungsoyr O, Sexton J, Sandanger I, Nygard JF (2009) Sensitivity analysis for unmeasured confounding in a marginal structural Cox proportional hazards model. *Lifetime Data Anal* 15: 278–294. 10.1007/s10985-008-9109-x [doi].
47. Van der Wal WM, Geskus RB (2011) An R Package for Inverse Probability Weighting. 43 (13): 1–23.
48. Grambsch PM, Therneau TM (1994) Proportional Hazards Tests and Diagnostics Based on Weighted Residuals. Vol. 81, No. 3: 515–526.
49. Lash TL, Silliman RA (2000) A sensitivity analysis to separate bias due to confounding from bias due to predicting misclassification by a variable that does both. *Epidemiology* 11: 544–549.
50. Coenen JL, van Diejen-Visser MP, van Pelt J, van Deursen CT (1991) Measurements of serum ferritin used to predict concentrations of iron in bone marrow in anemia of chronic disease. *Clin Chem* 37: 560–563.
51. Gwamaka M, Kurtis JD, Sorensen BE, Holte S, Morrison R, et al. (2012) Iron Deficiency Protects Against Severe Plasmodium falciparum Malaria and Death in Young Children. *Clin Infect Dis* 54: 1137–1144. cis010 [pii];10.1093/cid/cis010 [doi].
52. Cercamondi CI, Egli IM, Ahouandjinou E, Dossa R, Zeder C, et al. (2010) Afebrile Plasmodium falciparum parasitemia decreases absorption of fortification iron but does not affect systemic iron utilization: a double stable-isotope study in young Beninese women. *Am J Clin Nutr* 92: 1385–1392. ajcn.2010.30051 [pii];10.3945/ajcn.2010.30051 [doi].
53. Doherty CP, Cox SE, Fulford AJ, Austin S, Hilmers DC, et al. (2008) Iron incorporation and post-malaria anaemia. *PLoS One* 3: e2133. 10.1371/journal.pone.0002133 [doi].
54. Nweneka CV, Doherty CP, Cox S, Prentice A (2010) Iron delocalisation in the pathogenesis of malarial anaemia. *Trans R Soc Trop Med Hyg* 104: 175–184. S0035-9203(09)00294-6 [pii];10.1016/j.trstmh.2009.08.007 [doi].
55. Smith DL, Dushoff J, Snow RW, Hay SI (2005) The entomological inoculation rate and Plasmodium falciparum infection in African children. *Nature* 438: 492–495. nature04024 [pii];10.1038/nature04024 [doi].
56. Barry AE, Trieu A, Fowkes FJ, Pablo J, Kalantari-Dehaghi M, et al. (2011) The stability and complexity of antibody responses to the major surface antigen of Plasmodium falciparum are associated with age in a malaria endemic area. *Mol Cell Proteomics* 10: M111. M111.008326 [pii];10.1074/mcp.M111.008326 [doi].
57. Iolascon A, De Falco L, Beaumont C (2009) Molecular basis of inherited microcytic anemia due to defects in iron acquisition or heme synthesis. *Haematologica* 94: 395–408. haematol.13619 [pii];10.3324/haematol.13619 [doi].
58. Matsuzaki-Moriya C, Tu L, Ishida H, Imai T, Suzue K, et al. (2011) A critical role for phagocytosis in resistance to malaria in iron-deficient mice. *Eur J Immunol* 41: 1365–1375. 10.1002/eji.201040942 [doi].
59. Raiten DJ, Namaste S, Brabin B (2011) Considerations for the safe and effective use of iron interventions in areas of malaria burden - executive summary. *Int J Vitam Nutr Res* 81: 57–71. 10.1024/0300-9831/a000051 [doi].
60. Carter JY, Loolpapit MP, Lema OE, Tome JL, Nagelkerke NJ, et al. (2005) Reduction of the efficacy of antifolate antimalarial therapy by folic acid supplementation. *Am J Trop Med Hyg* 73: 166–170. 73/1/166 [pii].
61. English M, Snow RW (2006) Iron and folic acid supplementation and malaria risk. *Lancet* 367: 90–91. S0140-6736(06)67939-7 [pii];10.1016/S0140-6736(06)67939-7 [doi].
62. Denic S, Agarwal MM (2007) Nutritional iron deficiency: an evolutionary perspective. *Nutrition* 23: 603–614. S0899-9007(07)00149-9 [pii];10.1016/j.nut.2007.05.002 [doi].
63. Wander K, Shell-Duncan B, McDade TW (2009) Evaluation of iron deficiency as a nutritional adaptation to infectious disease: an evolutionary medicine perspective. *Am J Hum Biol* 21: 172–179. 10.1002/ajhb.20839 [doi].
64. Mutero CM, Schلودder D, Kabatereine N, Kramer R (2012) Integrated vector management for malaria control in Uganda: knowledge, perceptions and policy development. *Malar J* 11: 21. 1475-2875-11-21 [pii];10.1186/1475-2875-11-21 [doi].
65. Rowe AK, Onikpo F, Lama M, Osterholt DM, Deming MS (2011) Impact of a malaria-control project in Benin that included the integrated management of childhood illness strategy. *Am J Public Health* 101: 2333–2341. AJPH.2010.300068 [pii];10.2105/AJPH.2010.300068 [doi].
66. Smith JL, Brooker S (2010) Impact of hookworm infection and deworming on anaemia in non-pregnant populations: a systematic review. *Trop Med Int Health* 15: 776–795.