Severe malaria in Battambang Referral Hospital, an area of multidrug resistance in Western-Cambodia: a retrospective analysis of cases from 2006–2009

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Abstract

Background: Despite recent malaria containment and control efforts leading to reduced incidence, Cambodia remains endemic for both Plasmodium vivax and multidrug-resistant Plasmodium falciparum malaria. Little has been reported in the peer-reviewed literature regarding the burden of severe malaria (SM) in Cambodia.

Methods: Medical records for all patients admitted to the Battambang Referral Hospital (BRH) with an admitting or discharge diagnosis of SM from 2006 to 2009 (suspected SM cases) were reviewed. Those meeting the case definition of SM according to retrospective chart review and investigator assessment of probable cases, based on published national guidelines available at the time, were analysed for trends in demographics, mortality and referral patterns.

Results: Of the 537 suspected SM cases at BRH during the study period, 393 (73%) met published WHO criteria for SM infection. Despite limited diagnostic and treatment facilities, overall mortality was 14%, with 7% mortality in children 14 and under, but 19% in adults (60% of cases). Cerebral malaria with coma was relatively rare (17%), but mortality was disproportionately high at 35%. Mean time to hospital presentation was five days (range one to 30 days) after onset of symptoms. While patients with delays in presentation had worse outcomes, there was no excess mortality based on treatment referral times, distance travelled or residence in artemisinin-resistance containment (ARC) Zone 1 compared to Zone 2.

Conclusions: Despite limitations in diagnosis and treatment, and multiple confounding co-morbidities, mortality rates at BRH were similar to reports from other countries in the region. Interventions to improve access to early diagnosis and effective treatment, combined with modest improvements in intensive care, are likely to reduce mortality further. Patients referred from Zone 1 did not have excess mortality compared to Zone 2 ARC areas. A steep decrease in SM cases and deaths observed in the first half of 2009 has since continued, indicating some success from containment efforts despite the emergence of artemisinin resistance in this area.
Background

By definition, patients with severe malaria are in critical condition and require emergency therapy with parenteral drugs [1]. Untreated, mortality approaches 100%. Little has been reported in the peer-reviewed literature about the burden of severe malaria in Western Cambodia, located in World Health Organization's (WHO) Zone 1 artemisinin resistance containment (ARC) area.

Through implementation of a successful national malaria control programme, officially reported malaria incidence in Cambodia decreased by more than 50% between 2001–2010, from 11.03 cases per 1,000 in 2000 to 4.07 cases per 1,000 in 2010. Malaria mortality steadily decreased over the last decade from 5.2 deaths (in 2001) to 0.98 deaths per 100,000 in 2010 [2]. Despite these successes, 60% of Cambodia’s territory continues to be endemic for malaria, though much of this is sparsely populated. The majority of cases occur in eastern provinces, with artemisinin resistance emerging in the west.

In 2007 an estimated 2.13 million Cambodians (about 15% of the total population) living within 2 km of an endemic forested area were at risk of malaria, with 42,124 confirmed cases. There were 2,648 severe cases admitted to referral hospitals, of which 219 died. Data from the National Center for Parasitology, Entomology and Malaria Control, Cambodia (CNM) in 2007 indicated that Battambang Province had the second highest case-fatality rate (CFR) nationally (17.4%) – double the national CFR (Figure 1).

Study objectives were to assess incidence of severe malaria at a tertiary referral hospital, and further characterize the presentation of severe malaria in this resource-limited setting based on WHO recommended case definitions. A priori, it was suspected that access to pre-hospital care, drug availability and compliance, and delayed presentation would be important factors in

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**Figure 1** Morbidity, mortality (number of deaths per 100,000 per year) and case fatality rates (CFR, % deaths) from severe malaria in referral hospitals between 2006–9, Cambodia versus Battambang Referral Hospital.
outcomes and survival. Differences in mortality from the
two separate ARC zones served by Battambang Referral
Hospital (BRH) were assessed. The operational utility of
published guidelines for the diagnosis and management
of severe malaria were also evaluated. The authors re-
port here the first data on the burden of severe mala-
ria at the government referral hospital in Battambang,
Cambodia's second largest city, and the first area in
the world to report artemisinin-resistant malaria [3].

Methods
Study location
The study was conducted at the BRH, located in
Battambang City in Battambang Province, Western
Cambodia (Figure 2). BRH is a 270-bed facility and the
main hospital for Battambang city and surrounding dis-
tricts and admits on average 2,000 patients per month.

Identification of cases
Medical records for all patients admitted to the BRH
from January 2006 to June 2009 with an admission and/
or discharge diagnosis of severe malaria (irrespective of
confirmation by slide or rapid diagnostic test) were
reviewed to analyse morbidity, mortality, demography
and referral patterns. Since no separate log for severe
malaria cases existed, all hospital records for the study
period were searched in order to identify the severe mala-
ria cases. Individual patient records were reviewed by a
study physician, and data systematically extracted onto
case report forms. Discharge diagnoses were reviewed
independently for adherence to the Cambodian national
diagnosis and treatment guidelines in use at the time.
Severe malaria was defined as a complicated case of ma-
laria infection (Plasmodium falciparum and/or Plasmo-
dium vivax) presenting with one or more criteria of
severe disease [1]. WHO criteria for severe malaria were
adopted by the Ministry of Health as part of the
National Malaria Treatment Guidelines in Cambodia
and all clinicians would have had access to these guide-
lines at the time of the study. The availability of means
to assess the various diagnostic and prognostic param-
eters is outlined in Table 1.

Figure 2 Map of Cambodia with study site location and case fatality rates according to province of residence/origin, with inset
showing a map of Battambang Province with case fatality rates according to district of residence.
For the purpose of this paper, referral to ‘smear-positive’ patients indicates those with a presence of asexual malaria parasites as detected by microscopy or rapid diagnostic test. Overall CFR and number of cases who died included those dying in hospital, as well as those who were discharged to home in worsening (moribund) condition. The retrospective hospital record review was carried out in compliance with the Declaration of Helsinki 2000, following protocol approval by the Walter Reed Army Institute of Research IRB and the National Ethical Committee for Health Research (NECHR) in Cambodia. Personal identifying information was not collected on the case report forms.

Data analysis
Data were translated where needed from French and/or Khmer into English, and transcribed from medical records onto paper case report forms, and analysed using SPSS 12.0.0 for Windows. Double data entry was performed using Microsoft Excel to minimize transcription errors. Descriptive statistical analyses were performed for demography, referral patterns, symptoms, signs, admitting laboratory parameters, complications with respect to survival outcomes, and case management data. Fisher’s exact test (two-sided) was used to compare proportions from non-parametric tests when the expected values were less than five. Mortality risk was expressed with odds ratios (OR) and 95% confidence intervals (95% CI). Multiple logistic regression analysis was used to calculate adjusted odds ratios (aOR) to evaluate associations of mortality with presenting signs and symptoms, and complications occurring during hospital admission. Variables with crude OR significant at the level of p <0.2 in univariate analyses were included in a multivariate analysis.

Results
Between January 2006 and June 2009, a total of 537 records from suspected severe malaria cases were reviewed from patients admitted or discharged from BRH with a diagnosis of severe malaria. After review by study physicians, 393 patients met published Cambodian severe malaria case definitions and were considered to be ‘probable severe malaria cases’ (Figure 3).

Table 1 Availability of means to assess diagnostic and prognostic parameters, and provide treatment of severe malaria at Battambang Referral Hospital

| WHO severe malaria criteria | Description as published in national guidelines | Availability at BRH | Treatment available |
|----------------------------|-------------------------------------------------|---------------------|-------------------|
| Prostration                | Inability to sit upright without support or to drink | Yes | Supportive only |
| Impaired consciousness     | Modified Glasgow score ≤ 9 in adult and children >5 years old or Blantyre coma score ≤ 2/5 in children who have not learned to speak | Yes | Supportive only |
| Respiratory distress       | Sustained nasal flaring, intercostals recession and deep or fast breathing (RR >25/min in adults and >40/min in children) | Yes | Limited - no ventilators, low flow oxygen only |
| Multiple convulsions       | > 1/24 h | Yes | Parenteral or NG diazepam |
| Circulatory collapse       | BP <50 mmHg before 5 years old or BP <80 mmHg above 5 years old, cool peripheries, weak pulses | Yes | Peripherally administered IV pressors, fluids |
| Pulmonary oedema           | Radiological features | Yes | Limited - diuretics, oxygen, morphine |
| Abnormal bleeding          | (No definition provided) | Yes | Blood transfusion (limited) |
| Jaundice                   | (No definition provided) | Yes | Supportive |
| Haemoglobinuria            | (No definition provided) | Yes | Blood transfusion (limited) |
| Frequent vomiting          | Vomiting everything with inability to retain food or medicines | Yes | Antiemetic drugs, parenteral route of drug administration |
| Laboratory finding         | | | |
| Severe anaemia (or pallor) | Hb <5g/dl or Ht <15% | Yes | Blood transfusion (limited) |
| Hypoglycaemia              | Glycaemia <2.2 mmol/l or 0.4 g/l | Yes | IV dextrose |
| Metabolic acidosis         | Plasma bicarbonate <15 mmol/l | No | IV fluids |
| Hyperlactataemia           | Plasma lactate >5 mmol/l | No | IV fluids |
| Hyperparasitaemia          | *P. falciparum* "+++" or >200 000 parasites/μl | Clinical only - rare lab tests | Intra muscular artemether |
| Renal failure (or oliguria)| Urine output <400 ml/24 h or plasma creatinine>265 μmol/l in adult or urine output <12 ml/kg/24 h or plasma creatinine above the age-related normal range, persisting after rehydration in children | Clinical only - rare lab tests | Peritoneal dialysis occasionally available; supportive care |
Demographic and clinical characteristics and mortality

Overall CFR among the 393 probable severe malaria cases was 14.0% (55 out of 393), with the majority dying (38 or 9.7%) in hospital. The majority were male (224 or 57.0%), aged between five and 30 (median age 18 years), and were school-aged children (107 or 27%) or farmers (101 or 26%). Most patients (92%) were self-referred, arrived by private vehicle, and resided in Battambang Province (Table 2). Among the 393 probable severe malaria cases, 265 (67%) had malaria diagnoses confirmed by blood smear or rapid diagnostic test (RDT). Malaria species identified in the 265 smear/RDT confirmed cases

Figure 3 Study flow diagram.
Out of 55 total deaths reported, there were 36 deaths among the 265 smear-positives (mortality rate 13.6%), versus 19 deaths among the 128 smear-negatives (14.9% mortality). Lowest case fatalities were found in the five to 14 year age group (6.2%), while most fatalities were 15 to 30 years old (19%) or older than 50 (19%). The under-five mortality rate was 13%. Occupations with the

Table 2 Demographic and clinical characteristics of patients that met severe malaria criteria (n=393)

|                         | Slide positive |               | Slide negative/Not done |               |
|-------------------------|----------------|---------------|-------------------------|---------------|
|                         | Total          | Survived      | Died                    | Survived      | Died                    |
|                         | n  | (%)                | n  | (%)                | n  | (%)                |
| **Age category**        |    |                    |    |                    |    |                    |
| 0-4                     | 30 | 28 (93)            | 2  | (7)                | 23 | 18 (78)            | 5  | (22) |
| 5-14                    | 72 | 70 (97)            | 2  | (3)                | 41 | 36 (88)            | 5  | (12) |
| 15-30                   | 89 | 70 (79)            | 19 | (21)               | 27 | 24 (89)            | 3  | (11) |
| 31-49                   | 53 | 44 (83)            | 9  | (17)               | 26 | 22 (85)            | 4  | (15) |
| >=50                    | 21 | 17 (81)            | 4  | (19)               | 11 | 9 (82)             | 2  | (18) |
| **Sex**                 |    |                    |    |                    |    |                    |
| Male                    | 152| 133 (88)           | 19 | (13)               | 72 | 62 (86)            | 10 | (14) |
| Female                  | 113| 96 (85)            | 17 | (15)               | 56 | 47 (84)            | 9  | (16) |
| **Occupation**          |    |                    |    |                    |    |                    |
| Labour                  | 40 | 30 (75)            | 10 | (25)               | 7  | 6 (86)             | 1  | (14) |
| Government              | 3  | 2 (67)             | 1  | (33)               | 2  | 2 (100)            | 0  | (0)  |
| Housewife               | 11 | 8 (73)             | 3  | (27)               | 6  | 4 (67)             | 2  | (33) |
| Farmer                  | 66 | 52 (79)            | 14 | (21)               | 35 | 30 (86)            | 5  | (14) |
| Business                | 2  | 2 (100)            | 0  | (0)                | 1  | 1 (100)            | 0  | (0)  |
| Monk                    | 1  | 1 (100)            | 0  | (0)                | 1  | 1 (100)            | 0  | (0)  |
| School-aged child       | 74 | 69 (93)            | 5  | (7)                | 33 | 29 (88)            | 4  | (12) |
| Unemployed              | 15 | 14 (93)            | 1  | (7)                | 6  | 5 (83)             | 1  | (17) |
| Other                   | 48 | 46 (96)            | 2  | (4)                | 36 | 30 (83)            | 6  | (17) |
| **Referral by**         |    |                    |    |                    |    |                    |
| Self                    | 173| 152 (88)           | 21 | (12)               | 79 | 72 (91)            | 7  | (9)  |
| Private sector          | 1  | 1 (100)            | 0  | (0)                | 11 | 7 (64)             | 4  | (36) |
| Government Referral Hospital | 9 | 6 (67)             | 3  | (33)               | 37 | 29 (78)            | 8  | (22) |
| Government Health Centre| 82 | 70 (85)            | 12 | (15)               | 1  | 1 (100)            | 0  | (0)  |
| **Arrived by**          |    |                    |    |                    |    |                    |
| Private vehicle         | 181| 157 (87)           | 24 | (13)               | 76 | 69 (91)            | 7  | (9)  |
| Government ambulance    | 78 | 67 (86)            | 11 | (14)               | 46 | 35 (76)            | 11 | (24) |
| **Parasite species**    |    |                    |    |                    |    |                    |
| Pf                      | 217| 185 (85)           | 32 | (15)               | 110| 94 (85)            | 16 | (15) |
| Pv                      | 33 | 29 (88)            | 4  | (12)               |    |                    |    |      |
| Mixed Pf+Pv             | 15 | 15 (100)           | 0  | (0)                |    |                    |    |      |
| Negative                |    |                    |    |                    |    |                    |
| ND                      | 18 | 15 (83)            | 3  | (17)               |    |                    |    |      |
| **Parasitaemia time 0** |    |                    |    |                    |    |                    |
| No or low (1+ to 3+)    | 145| 122 (84)           | 23 | (16)               | 110| 94 (85)            | 16 | (15) |
| Hyperparasitaemia       | 113| 101 (89)           | 12 | (11)               | 0  | 0 (0)              | 0  | (0)  |
| (4+ to 6+)              |    |                    |    |                    |    |                    |

included 217 *P. falciparum* (81.9%), 33 *P. vivax* (12.4%); and 15 documented mixed infections (5.7%) positive for both *P. falciparum* and *P. vivax* (Table 2).
highest CFR included labourers (23%), government workers (20%), farmers (19%), and housewives (29%). A higher death rate was found in referred patients (19.2%), in particular those referred from government health facilities, versus patients who came directly to the hospital (8%). There were two fatalities among the 13 pregnant severe malaria cases (both *P. falciparum* infections), as well as one spontaneous abortion and three stillbirths (all *P. falciparum*). The only pregnant *P. vivax* case survived with normal delivery at term during hospital admission.

**Signs and symptoms on admission**

The most common presenting signs and symptoms among 393 probable severe malaria cases included fever (95%), headache (63%), dark urine (56%), chills (53%), and impaired consciousness (42%). There were 65 patients who presented in a coma (17%). The only presenting symptoms, adjusted for age group (aOR age >15 years=2.0, 95% CI 0.9-4.1) that remained significantly associated with a fatal outcome after a multiple logistic regression analysis were coma (aOR=5.8, 95% CI 2.9-11.5), pallor (aOR=4.2, 95% CI 1.9-9.0), dyspnea (aOR= 2.5, 95% CI 1.2, 5.4) and dark urine (aOR=0.4, 95% CI 0.2-0.8) (Table 3).

**Laboratory data**

Laboratory assessment at BRH was limited and included microscopy for parasitaemia (94%), white blood cell count (92%), haematocrit (63%) or haemoglobin (6%), platelets (48%) and plasma glucose (32%). Assessment of liver enzymes (3%) were done infrequently, and other commonly used tests including blood urea nitrogen, creatinine, bilirubin, albumin, plasma lactate and bicarbonate were unavailable. Higher parasitaemia based on the qualitative scoring system used was not associated with higher mortality. Although severe anaemia was common, with approximately 29% of patients having haematocrit values less than 15% at initial presentation, mean presenting haematocrit values between non-survivors and survivors were similar (25.2% versus 24.0%), and no patient had a presenting glucose level less than 40 mg/dL. Non-survivors had higher mean leukocyte counts than survivors (10.6 versus 8.0 x 10^3 cells/ml, p-value = 0.02). Of 191 patients tested, the majority had a platelet count greater than 150,000/μL, with the mean platelet count significantly lower in non-survivors (183,000/μL versus 208,000/μL, p-value = 0.04) (Table 4).

**Clinical course and complications**

Features of the clinical course, and complications that developed during hospital admission are shown in Table 5 (all cases) and Table 6 (children <15 years old), along with their association with mortality. In patients who met the severe malaria case definition (all age groups), the most common clinical features included prostration (68%), impaired consciousness (65%), and respiratory distress (48%), and all three were associated with a substantially increased risk of mortality in a univariate analysis. Circulatory collapse (15%), renal failure (27%), and pulmonary oedema (2%), though less common, were also associated with increased mortality, while severe anaemia defined as a haematocrit of less than 15 (55%) and hyperparasitaemia (33%) were not. In multivariate analysis, only circulatory collapse (aOR67.5; 95% CI 23.2-196) and renal failure (aOR19.7; 95% CI 4.6-83.9) remained statistically significantly associated with a higher mortality risk independent of age. Clinical features were similar in the under 15 age group, although the majority (60%) developed multiple convulsions during their hospital stay. Circulatory collapse (adjusted OR= 186, 95% CI 30.4, 1,143) was the only complication significantly associated

| Table 3 Signs and symptoms on admission in probable severe malaria patients (n= 393) |
|---------------------------------------------------------------|
| **Sign/ symptoms** | **Total Died OR (95% CI)** | **p-value** |
|---------------------------------------------------------------|
| Fever | 372 (95) 50 (13) | 0.5 (0.2, 1.4) | 0.19 |
| Headache | 247 (63) 32 (13) | 0.8 (0.4, 1.4) | 0.46 |
| Dark urine | 220 (56) 24 (11) | 0.6 (0.3, 1.0) | 0.06 |
| Chills | 207 (53) 26 (13) | 0.8 (0.4, 1.4) | 0.47 |
| Impaired consciousness (excl coma) | 166 (42) 23 (14) | 1.0 (0.6, 1.8) | 1 |

*Fisher’s Exact Test (2-sided).

Adjusted for Dark Urine, Coma, Dyspnea and Pallor and Age Group, aOR (95% CI) of Dark Urine= 0.4 (0.2, 0.8), aOR of Coma = 5.8 (2.9, 11.5), aOR of Dyspnea = 2.5 (1.2, 5.4), aOR of Pallor = 4.2 (1.9, 9.0) and aOR of Age > 15 = 2.0 (0.9, 4.1).
with mortality in multivariate logistic regression analysis in the younger age group.

**Treatment**

Nearly 90% (345) of suspected severe malaria cases received parenteral therapy on admission which included either an anti-malarial drug alone (238 or 61%), or a combination of anti-malarial and an antibiotic (107 or 27%) (Table 7). Intramuscular artemether was the most frequently administered anti-malarial, given to 324 or 82% of suspected severe malaria patients on admission. Aminopenicillins, ciprofloxacin and cephalosporins were the most frequently given antibiotics.

**Table 4 Laboratory values reported on admission in probable severe malaria patients (n= 393)**

| Laboratory test | Total evaluated | Died | Survived | p-value* | Reference range |
|-----------------|-----------------|------|----------|----------|-----------------|
| WBC (x 10^3cells/ml) | 368 (94) | 1.6-35.2 | 10.6 ± 7.3 | 8.0 ± 4.8 | 0.02 | ≤ 12 | 303 |
| Hematocrit (%) | 253 (64) | 5.0-50 | 25.2 ± 10.4 | 24.0 ± 10.4 | 0.5 | ≤ 15 | 74 |
| Hemoglobin (g/dl) | 22 (6) | 3.0-16.2 | 9.8 ± 2.6 | 9.1 ± 4.2 | 0.8 | < 10 | 12 |
| Platelets (x 10^3cells/ml) | 191 (49) | 10-450 | 183 ± 49.4 | 208 ± 47.4 | 0.04 | ≤ 100 | 2 |
| Glucose (mg/dl) | 129 (33) | 50-459 | 102 ± 45.9 | 113 ± 73.4 | 0.4 | ≥ 100 | 2 |
| SGOT (IU/L) | 11 (3) | 19-390 | 144 ± 154.5 | 123 ± 126 | 0.8 | ≥ 3 times ULN | 6 |
| SGPT (IU/L) | 11 (3) | 20-430 | 66±45 | 110 ± 137 | 0.6 | ≥ 3 times ULN | 7 |

p< 0.05, *t-test (2-sided) or **One-Way ANOVA.

Treatment delay, cost and residence according to ARC Zone

Mean time to hospital presentation in severe malaria patients was five days (range one to 30 days) after onset of symptoms. No clear relationship was observed between treatment delay and outcome. Overall CFR of patients originating from ARC Zone 1 districts was lower than those from Zone 2 districts (11% versus 15%) (Table 8). For the majority (45%) of the study population, treatment was subsidized by local NGOs with a fixed US $20 payment, while 38% (150/393) paid for treatment out-of-pocket at a cost of between US$2.5 and 75 (median US$12.5). For 67 patients or 17%, no cost data were available, meaning these patients did not pay or no payment records could be found.

**Discussion**

This is one of the few studies to describe the presentation and epidemiology of severe malaria in Western Cambodia, a low transmission setting with multidrug-resistant *P. falciparum*, and an increasing predominance of *P. vivax* malaria. The study provides baseline information on the burden of severe malaria in a referral hospital near the epicentre of global anti-malarial drug resistance (WHO Zone 1) during the 2.5 years preceding the Artemisinin Resistance Containment Project and supports other preliminary evaluations reporting success in containment and reduction of mortality rates [4]. These intense efforts led by WHO and others provided bed net distribution, and trained local village malaria workers to provide earlier case detection and treatment...
free of charge, resulting in declines in falciparum malaria cases [5,6]. Following the downward trend in other Containment Project provinces, severe malaria incidence and CFR in Battambang Province reported to the National Malaria Control Program continued to decline significantly after 2009 with only two deaths out of 116 severe cases in 2010 and one death out of 101 severe cases in 2011 [7]. Other likely contributors to the reduction in severe disease in the interim have included malaria case management training, a ban on artemisinin monotherapy, a government crack-down on counterfeit drugs and illegal pharmacies, improved supply of good quality anti-malarials, an increased programme focus on migrant and mobile populations (including the military), and

| Table 5 Clinical course and complications experienced during hospital admission in probable severe malaria patients (n=393) |
|---|---|---|---|---|---|---|---|---|
| Neg | Pf | Pv | Pf+Pv | Not recorded | Total | Died | OR (95% CI) | p-value* |
|---|---|---|---|---|---|---|---|---|
| Prostration | 82 | 150 | 16 | 5 | 12 | 265 (68) | 53 (20) | 15.6 (3.7, 65.2) | 0 |
| Impaired consciousness | 81 | 143 | 14 | 5 | 11 | 254 (65) | 53 (21) | 18.1 (4.3, 75.4) | 0 |
| Respiratory distress | 55 | 105 | 12 | 5 | 9 | 186 (48) | 39 (21) | 3.9 (2.0, 7.5) | 0 |
| Severe anaemia | 55 | 90 | 11 | 3 | 10 | 169 (55) | 27 (16) | 1.1 (0.6, 2.1) | 0.75 |
| Hyperparasitaemia | 1 | 101 | 7 | 11 | 8 | 128 (33) | 13 (10) | 0.6 (0.3, 1.1) | 0.12 |
| Jaundice | 21 | 38 | 4 | 1 | 4 | 68 (17) | 14 (21) | 1.7 (0.9, 3.5) | 0.12 |
| Abdominal bleeding | 23 | 29 | 5 | 0 | 5 | 62 (16) | 12 (19) | 1.6 (0.8, 3.2) | 0.23 |
| Frequent vomiting | 17 | 34 | 6 | 3 | 1 | 61 (16) | 5 (8) | 0.5 (0.2, 1.3) | 0.23 |
| Circulatory collapse | 17 | 32 | 7 | 1 | 3 | 60 (15) | 43 (72) | 67.5 (30.2, 150.8) | 0** |
| Renal failure | 8 | 16 | 2 | 0 | 1 | 27 (7) | 16 (59) | 16.8 (7.1, 39.7) | 0** |
| Multiple convulsion | 7 | 11 | 2 | 2 | 2 | 24 (6) | 7 (29) | 2.8 (1.1, 7.1) | 0.03 |
| Macroscopic haemoglobinuria | 6 | 6 | 0 | 0 | 0 | 12 (3) | 3 (25) | 2.5 (0.7, 9.7) | 0.17 |
| Hypoglycemria | 1 | 7 | 0 | 2 | 0 | 10 (7) | 3 (30) | 1.5 (3.6, 6.2) | 0.7 |
| Pulmonary oedema | 2 | 2 | 1 | 0 | 0 | 5 (2) | 4 (80) | 423 (4.5, 398) | 0 |

*Fisher’s Exact Test (2-sided).
**Adjusted for age group, circulatory collapse and renal failure, aOR (95% CI) of circulatory collapse=67.5 (23.2, 196) and aOR of renal failure=19.7 (4.6, 83.9).

| Table 6 Clinical course and complications experienced during hospital admission in probable severe malaria patients for only those aged 0 to 15 years (n=130) |
|---|---|---|---|---|---|---|---|---|
| Neg | Pf | Pv | Pf+Pv | Not recorded | Total | Died | OR (95% CI) | p-value* |
|---|---|---|---|---|---|---|---|---|
| Prostration | 51 | 65 | 8 | 2 | 4 | 130 (78) | 13 (10) | 3.8 (0.5, 29.9) | 0.31 |
| Impaired consciousness | 51 | 65 | 8 | 2 | 4 | 130 (78) | 13 (10) | 3.9 (0.5, 30.8) | 0.31 |
| Multiple convulsion | 3 | 83 | 12 | 1 | 1 | 100 (60) | 1 (1) | 2.9 (0.3, 27.4) | 0.36 |
| Severe anaemia | 36 | 51 | 5 | 2 | 3 | 97 (58) | 10 (10) | 1.4 (0.4, 4.8) | 0.77 |
| Respiratory distress | 26 | 34 | 4 | 3 | 3 | 70 (42) | 11 (16) | 8.7 (1.9, 40.5) | 0 |
| Hyperparasitaemia | 1 | 33 | 1 | 5 | 1 | 41 (25) | 1 (2) | 0.2 (0.03, 1.6) | 0.12 |
| Frequent vomiting | 14 | 17 | 3 | 2 | 1 | 37 (22) | 1 (3) | 0.2 (0.03, 2.0) | 0.2 |
| Abdominal bleeding | 12 | 9 | 0 | 0 | 0 | 21 (13) | 3 (14) | 2.0 (0.5, 7.7) | 0.4 |
| Circulatory collapse | 8 | 3 | 1 | 7 | 0 | 19 (11) | 10 (53) | 186 (30.4, 1143) | 0** |
| Jaundice | 6 | 4 | 0 | 0 | 0 | 10 (6) | 2 (20) | 3.0 (0.6, 15.8) | 0.2 |
| Macroscopic haemoglobinuria | 3 | 0 | 0 | 0 | 0 | 3 (2) | 1 (33) | 5.7 (0.5, 66.6) | 0.24 |
| Pulmonary oedema | 0 | 0 | 1 | 0 | 0 | 1 (1) | 0 (0) | 1.0 (1.0, 1.1) | 1 |
| Renal failure | 0 | 0 | 0 | 0 | 0 | 0 (0) | 0 (0) | - | - |
| Hypoglycaemia | 0 | 0 | 0 | 0 | 0 | 0 (0) | 0 (0) | - | - |

*Fisher’s Exact Test (2-sided).
**Adjusted for Circulatory Collapse, aOR (95% CI) of Circulatory Collapse = crude OR = 186 (30.4, 1143).
Table 7 Anti malarial and antibiotic treatment on admission versus outcome in probable severe malaria cases (n=393)

| Anti-malarial and antibiotic use | Total n (%) | Died n (%) |
|----------------------------------|-------------|------------|
| Only anti malarials              | 238 (61)    | 25 (11)    |
| Only antibiotics                  | 18 (5)      | 2 (11)     |
| Antibiotics and anti malarials    | 107 (27)    | 22 (21)    |
| No anti-malarials or antibiotics  | 30 (8)      | 6 (20)     |

Number and type of anti malarials

| Anti malarials | n (%) | n (%) |
|----------------|-------|-------|
| At least one anti-malarial | 345 (88) | 47 (14) |
| Two or more anti-malarials | 15 (4) | 2 (13) |
| Artemether | 324 (82) | 44 (14) |
| Artesunate + Mefloquine | 13 (3) | 0 (0) |
| Chloroquine | 10 (3) | 1 (10) |
| Quinine | 9 (2) | 3 (33) |
| Mefloquine | 4 (1) | 1 (25) |
| DHA+Piperaquine | 2 (1) | 0 (0) |
| DHA | 2 (1) | 1 (50) |

Number and type of antibiotics

| Antibiotics | n (%) | n (%) |
|-------------|-------|-------|
| At least one antibiotic | 125 (32) | 24 (19) |
| Two or more antibiotics | 15 (4) | 4 (27) |
| Aminopenicillins | 61 (16) | 10 (16) |
| Cephalosporins | 37 (9) | 14 (38) |
| Ciprofloxacin | 27 (7) | 0 (0) |
| Gentamycin | 10 (3) | 4 (40) |
| Co-trimoxazole | 4 (1) | 0 (0) |
| Metronidazole | 4 (1) | 0 (0) |
| Tetracycline | 1 (0) | 0 (0) |

basic improvements in infrastructure such as road access in Western Cambodia. The relationship between level of artemisinin tolerance and pathogenicity of malaria parasites is unclear. High levels of differentiation in artemisinin-resistant *P. falciparum* subpopulations have recently been demonstrated in Cambodia [8] and future research may or may not link these to different levels in pathogenicity.

On review of the hospital records, it was often apparent that malaria was a co-existing infection of another serious condition, such as meningitis, sepsis and gastrointestinal haemorrhage which is not unexpected, given the recent evidence of *falciparum* malaria infection as a risk factor for bacteraemia in African children [9]. Bacterial cultures that could have ruled out bacteraemia, were not available at BRH for this population.

Despite substantial limitations, diagnostic and treatment guidelines were reasonably well adhered to where possible at this facility, likely contributing to an overall CFR of 14.2% which was intermediate compared to other reports from the region. CFR in other retrospective studies in the region have varied from 1.8% in Thailand [10], 6.3% in the Philippines [11], to 35% in Malaysia [12]. This compares to recent mortality rates in African studies, largely in children, ranging from 4.5% in Cameroon [13] to 11.5% in Madagascar [14] to up to 22% in children with bacteraemia in Mozambique [6,15]. Despite the emergence of artemisinin resistance in Western Cambodia during the study period, cases of severe malaria and deaths declined in BRH and nationally. Overall CFR was lower in severe malaria patients residing in districts with confirmed artemisinin resistance (Zone 1), likely resulting from increased active case detection and early treatment. Overall malaria incidence in Cambodia increased from 4.1 per 1,000 in 2008 to 6.22 per 1,000 in 2009 attributed to increased migrational movements, while CFR of severe malaria cases continued to decline [2].

Although life-threatening malaria complications can affect patients of all ages, disease presentation and mortality have been found to be closely related to transmission intensity and age of the patient [16]. In a low transmission setting such as this, where immunity may never be acquired, older patients may have higher mortality rates owing to delayed presentation. The most frequent complications associated with mortality, particularly in older patients, were prostration, respiratory distress and coma, with 35% of comatose patients dying on or soon after admission. Coma and metabolic acidosis have been found to be the most significant risk factors for death in both children and adults in prior studies in Asia [17,18]. Others have demonstrated that renal function monitoring in these patients may be crucial for survival [16,19]. The lack of metabolic and chemistry testing inhibited the ability to identify metabolic acidosis and renal dysfunction early, making prognosis and management more challenging. Of the limited diagnostic testing available, mortality did appear to correspond with laboratory abnormalities for both leukocyte and platelet counts, lending some limited prognostic value to these tests. Elevated or depressed leukocyte counts often indicate concomitant bacterial infections, and patients may benefit from empiric antibiotic treatment [20]. Thrombocytopenia, one of the most common laboratory abnormalities in severe malaria [21] also had some prognostic value for severe disease though it did not predict bleeding complications.

In addition to diagnostic and prognostic limitations, challenges in management included lack of mechanical ventilation units, renal haemodialysis, centrally administered pressors, exchange transfusion capabilities or real time haemodynamic and metabolic data to guide resuscitation. Limited peritoneal dialysis was available at times, but rarely employed. As a result, renal failure and
pulmonary oedema, while rare at this facility (reported in only 7% and 2% patients), were associated with the highest CFR in this group.

Recent descriptions of treatment-seeking behaviour, cultural practices and socio-economic barriers have revealed causes for delays in appropriate malaria treatment in Cambodia, representing an important potential area for intervention to reduce mortality [22]. Several factors appear to have contributed to mortality from severe malaria at the BRH including delayed presentation, delayed admission referral, cultural beliefs, and difficulty accessing health care facilities in this austere setting. The average time between onset of symptoms and admission to BRH was 5.5 days, greater in some of the outlying districts. Although Cambodia has an extensive peripheral health centre network, lack of public sector resources may prompt patients to seek initial treatment directly at the highest level of care possible, bypassing outlying networks. The majority of severe cases (70%) presented directly to the BRH, and were admitted directly for treatment to the intensive care unit, with only the minority (approximately 30%) being referred from outlying centres. Referred patients (especially from the public sector) had worse outcomes, suggesting increased treatment delay and lack of pre-referral treatment such as parenteral artemisinins, suppositories and/or transfusion capability.

Roughly 90% of all patients discharged with a severe malaria diagnosis at BRH received intramuscular artemether on the day of admission, which at the time of publication remains the current national standard of care for severe malaria in Cambodia. A five to seven-day course is used, followed by treatment with mefloquine, but is sometimes unavailable, prompting patients to obtain drugs from private-sector pharmacies risking counterfeit or substandard treatment. Parenteral artesunate was not available in Cambodia during the study period. Parenteral artesunate currently remains the most effective treatment remaining for severe malaria [23], and was found to be more effective and less expensive than quinine in children in sub-Saharan Africa [24] and adults in Southeast Asia [25]. Though

| Province | District | Onset to presentation (days) |
|----------|----------|-----------------------------|
|          | Died     | Survived | Total | Mean | SEM |
| Battambang | 122 | 11 (10) | 101 (90) | 112 | 5.1 | 0.3 |
| Phnom Preuk | 0 | (0) | 5 (100) | 5 | 6.0 | 0.6 |
| Kam Reang | 1 (25) | 3 (75) | 4 | 5.5 | 3.2 |
| Sampov Loun | 0 (0) | 1 (100) | 1 | 3.0 | - |
| Pailin | 16 | 3 (19) | 13 (81) | 16 | 6.7 | 1.8 |
| Pursat | 2 | 1 (50) | 1 (50) | 2 | 5.0 | 2.0 |
| Total ARC Zone 1 | 140 | 16 (11) | 124 (89) | 5.3 | 0.3 |

| Province | District | Onset to presentation (days) |
|----------|----------|-----------------------------|
|          | Died     | Survived | Total | Mean | SEM |
| Battambang | 241 | 10 (15) | 55 (85) | 65 | 6.0 | 0.7 |
| Mondol | Banan | 8 (18) | 37 (82) | 45 | 6.2 | 0.6 |
| Battambang | 4 (11) | 32 (89) | 36 | 5.4 | 0.6 |
| Thmar Koul | Sangker | 5 (24) | 16 (76) | 21 | 4.6 | 0.4 |
| Moung Russey | Ek Phnom | 3 (17) | 15 (83) | 18 | 7.8 | 1.6 |
| Other | Bavel | 3 (43) | 4 (57) | 7 | 6.7 | 1.0 |
| Koh Kralor | 1 (14) | 6 (86) | 7 | 5.3 | 1.2 |
| Total ARC Zone 2 | 253 | 39 (15) | 214 (85) | 5.8 | 0.3 |

*Banteay Meanchey(4), Pursat(4), Siem Reap(1), Sihanoukville(1), Svay Rieng(1), Takeo(1).
limited head to head comparisons between intravenous artesunate suggest some degree of inferiority for intramuscular artemether [26], both were found to be effective, and the relative simplicity of rapid intramuscular administration in critical illness may support its continued use in Cambodia. Availability and frequent use of this life-saving drug may explain the relatively low overall mortality rates seen, particularly considering the limited array of diagnostic and treatment facilities available. Quinine continues to be the first-line agent for pregnant women in Cambodia.

There has been recent recognition of the substantial proportion of severe malaria cases attributable to P. vivax in tropical regions [27]. Observed rates of severe P. vivax in Southeast Asia and the Pacific have been high in several reports, accounting for up to 25% of severe malaria cases. While some have shown reduced risk of severe disease attributable to P. vivax and/or mixed infection in low transmission settings [28], others have shown risks of severe disease to be increased, particularly in young children and in areas with high grade chloroquine resistance, though the latter has not been reported in Cambodia [29]. Though P. vivax is typically thought of as a relatively benign disease, 14.5% of patients meeting the case definition of severe malaria at BRH during the study period had slide proven P. vivax, and there were four documented deaths. There were no deaths reported among 17 patients with documented mixed infection. HIV-1 infection has been reported to be an important risk factor for severe malaria and death [30], but there was only one confirmed case of HIV co-infection in this study population.

The study's retrospective design placed limits on accurately interpreting the underlying disease burden. Accurately defining the terms ‘impaired consciousness’ and ‘prostration/coma’ retrospectively were challenging. WHO defines impaired consciousness as a ‘rousalable mental condition’, and prostration as the inability to sit upright without support or drink. However, varying terms were used by clinicians at BRH, including descriptions such as “somnolence”, “obnubilation”, “agitation”, “delirium”, “irritation”, “precoma”, and in some cases a Glasgow Coma Score (GCS) was given. In the retrospective analysis, the study team adopted the term ‘coma’ for those found to have either a clearly documented coma state, or understood to have an unarousable coma based on GCS scores. Limitations on diagnostic capabilities at this facility leave open the possibility that cases and/or complications may have been missed. The study does not reflect cases that never made it to medical attention, died out of hospital, or were treated at local facilities, in the private sector or at home. As such, the in-patient data are only representative of the referral centre and not of the overall severe malaria burden for this region of Cambodia. Likewise, the non-availability of important diagnostic measurements may have resulted in under-reporting of complications.

A large proportion of patients were slide-negative on admission, though nearly 50% of these patients still responded to anti-malarials alone. In such cases, preliminary microscopic diagnosis in hospital may have been obscured or simply not performed due to pre-treatment out of hospital prior to transfer. Many cases were diagnosed by rapid diagnostic kits or simply treated on clinical grounds. Patients with negative malaria smears in this cohort had higher crude mortality rates at 16% versus 11% in the patients with high parasitaemia, possibly due to delays in diagnosis and appropriate treatment, or earlier parenteral therapy in the higher parasitaemia patients. In this setting, a high index of suspicion for severe malaria remains warranted and may prove lifesaving. However, an empiric treatment approach to severe malaria should not preclude investigation of co-morbidities, which were common.

Conclusions

BRH continued to have a significant number of severe malaria cases between 2006–2009, suggesting that interventions to improve access to early diagnosis and treatment of malaria remain priorities for this region. Diagnostic ambiguities were common in this setting, with multiple confounding co-morbidities and limited patient management resources. The key to preventing mortality remains early detection, treatment and referral where necessary, and maintaining a high index of suspicion for severe malaria cases is essential. This includes not only adherence to guidelines, but consideration of the individual patients’ risks based on local endemicity, clinical history, pre-hospital care, and severity of presenting illness syndromes. Co-administration of broad-spectrum antibiotics in this resource-poor setting with frequent co-infection and septicemia may improve outcome. A subset of technologically appropriate core guidelines for diagnosis and management of severe malaria may be useful in this setting, and monitoring of the patient’s renal and acid–base balance through simple chemistry panels may be a useful and cost-effective addition to management at this and similar facilities. The drop in both numbers of severe cases and mortality during the period immediately following this study suggests some improvement in malaria control, likely to due to large-scale containment efforts in the region.

Competing interests

The authors declared that they have no competing interests.

Authors’ contributions

DS, CL, AT, DB, JL, YS, NS, KL, ST, MF, DW and CMC conceived, designed and supported the study. NB, SN, SC, CL, DS and YS collected the data. AT, CL, DS, SK, TW, DC and SJ analysed the data. DS, AT, DB and CL wrote the paper. YS passed away during the writing of this manuscript. All authors read and approved the final manuscript.
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