Anthropometry of fetal growth in rural Malawi in relation to maternal malaria and HIV status

B F Kalanda, S van Buuren, F H Verhoeff, B J Brabin

Objective: To describe fetal growth centiles in relation to maternal malaria and HIV status, using cross sectional measurements at birth.

Design: A cross sectional study of pregnant women and their babies. Data on maternal socioeconomic status and current pregnancy, including HIV status and newborn anthropometry, were collected. Malaria parasitaemia was assessed in maternal peripheral and placental blood, fetal haemoglobin was measured in cord blood, and maternal HIV status was determined.

Setting: Two district hospitals in rural southern Malawi, between March 1993 and July 1994.

Outcome variables: Newborn weight, length, Rohrer’s ponderal index.

Results: Maternal HIV (adjusted odds ratio (AOR) 1.76 (95% confidence interval 1.04 to 2.98)) and first pregnancy (AOR 1.83 (1.10 to 3.05)) were independently associated with low weight for age. Placental or peripheral parasitaemia at delivery (AOR 1.73 (1.02 to 2.88)) and primigravidae (AOR 2.13 (1.27 to 3.59)) were independently associated with low length for age. Maternal malaria at delivery and primiparity were associated with reduced newborn weight and length but not with disproportionate growth. Maternal HIV infection was associated only with reduced birth weight. The malaria and parity effect occurred throughout gestational weeks 30–40, but the HIV effect primarily after 38 weeks gestation.

Conclusion: Fetal growth retardation in weight and length commonly occurs in this highly malarious area and is present from 30 weeks gestation. A maternal HIV effect on fetal weight occurred after 38 weeks gestation.

Methods

Study area
This study was undertaken between March 1993 and July 1994 in Chikwawa District, Shire Valley, southern Malawi. This rural area with endemic malaria transmission covers about 4800 km² and is 10–300 m above sea level. Average rainfall in the study period was 520 mm/year mostly falling in November–March. Small scale agriculture of maize, sorghum, cotton, and sugar cane are the primary sources of food and income. Estimated population size in 1998 was 356 682, of which 77 701 were women of child bearing age. The study was located in Chikwawa District Hospital, a government hospital with free services, and Montfort Hospital, 30 km away, a fee paying mission hospital.

Enrolment
All women attending the antenatal facilities at the two hospitals were screened at their first antenatal visit after verbal consent had been obtained. A questionnaire, completed by a project nurse, included information on age, obstetric history, and antimalarial use in pregnancy. Literacy status was assessed from the reading of a simple sentence in the local language.

Delivery
It was not possible to obtain delivery information from home or health centre deliveries. Birth weight was measured on a Salter scale (nearest 10 g), and the baby was examined at 6–24 hours for gestational age using a modified Ballard method. Birth length was measured (nearest 0.1 cm) with the baby supine using a standardised measuring mat. Head circumference (occipitofrontal) was measured by tape (mm) (Talc, Guildford, Surrey, UK), and so was the circumference at the mid-point of the left upper arm. A blood sample was collected at delivery by venepuncture from the mother and umbilical cord. A malaria slide was made from blood collected deep between the placenta villi. These were stained with Giemsa and read counting asexual Plasmodium falciparum parasites against 200 white blood cells. Maternal HIV status was determined using two different enzyme linked immunosorbent assays: ICES HIV-1.0.2 (Murex, Dartford, Kent, UK) and, for confirmation, VIDAS HIV-2 new test (bioMerieux, Lyon, France). This procedure was in accordance with WHO recommendations for HIV diagnosis in asymptomatic patients from regions with prevalence above
Counselling was provided by trained nurses before and after the test. A subsample of sera, selected randomly (n = 651), was tested for HIV.

**Definitions**

Cord haemoglobin of less than 125 g/l defined fetal anaemia. Adolescence was defined as 12–19 years of age. Malaria at delivery was present if the mother had placental malaria and/or peripheral parasitaemia. Babies less than 37 weeks at delivery were preterm and less than 2500 g low birth weight. Wasting (low weight for age (W/A)), thinness (low weight for length (W/L)) and stunting (low length for age (L/A)) were defined as a standard deviation score of less than or equal to −2. Rohrer’s ponderal index was computed as birth weight (g)/birth length3 (cm).

**Analysis**

Data were analysed using SPSS for windows, version 11.0 (2001). The χ² and Fisher’s exact tests were used for comparison of discrete variables, and analysis of variance for continuous variables. Logistic regression was used to analyse risk factors related to wasting, stunting, and thinness. For adjusted odds ratios (AORs), p<0.05 was taken as significant. Growth curves were drawn using smoothed centile standards derived by the LMS program, version 1.16 (2000). The LMS method assumes that the data can be normalised by using a power transformation, which stretches one tail of the distribution and shrinks the other to remove skewness. The optimal power to obtain normality depends on age and is summarised by a smooth age dependent L curve. Trends in the median (M) and coefficient of variation (S) are similarly smoothed. Curve smoothness was decided by varying the L, M, and S values and observing the detrended Q-Q plots. Standard deviation scores were calculated using the EPI info 2002, nutrition program, using the CDC 2000 reference as the standards. Twins and stillbirths were excluded from the analysis.

**Ethical approval**

The study was granted ethical approval by the College of Medicine Research and Ethics Committee.

### Table 1

<table>
<thead>
<tr>
<th>Index</th>
<th>Parasitaemia –ve</th>
<th>Parasitaemia –ve</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth weight (g)</td>
<td>n</td>
<td>Mean (SD)</td>
<td>n</td>
</tr>
<tr>
<td>Birth length (cm)</td>
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<td>2843 (533)</td>
<td>983</td>
</tr>
<tr>
<td>Head circumference (cm)</td>
<td>355</td>
<td>47.1 (2.2)</td>
<td>971</td>
</tr>
<tr>
<td>PI (kg/cm³)</td>
<td>353</td>
<td>27.5 (7.2)</td>
<td>970</td>
</tr>
</tbody>
</table>

PI, Rohrer’s ponderal index.

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**Figure 1** Birth weight for gestational age centiles (10th, 50th, 90th) in relation to maternal malaria. Broken line, positive for malaria parasites in peripheral or placental blood; continuous line, negative for parasites.

**Figure 2** Birth length for gestational age centiles (10th, 50th, 90th) in relation to maternal malaria at delivery. Broken line, positive for malaria parasites in peripheral or placental blood; continuous line, negative for parasites.

**Figure 3** Rohrer’s ponderal index centiles (10th, 50th, 90th) by gestational age in relation to maternal malaria at delivery. Broken line, positive for malaria parasites in peripheral or placental blood; continuous line, negative for parasites.
RESULTS
A total of 4104 pregnant mothers were recruited at antenatal clinics. Of these, 1571 (38%) delivered in the study hospitals. There were 86 twins and 55 stillbirths. Four twins were stillborn, and for three infants there was missing information. These were excluded, leaving 1423 infants for the analysis. Of these, 49.7% (707) were male, 21.7% (309) had fetal anaemia, 17.1% (243) were preterm, and 14.8% (211) were low birth weight. Details of the gestation age distribution in this sample have been previously published.21 Of the mothers, 24.3% (344) were primigravidae, 22.4% (248) adolescent, and 26.9% (362) had placental or peripheral parasitaemia at delivery. From a sample of 621 screened for HIV, 25.6% were positive for HIV antibody.

Table 1 shows anthropometric indices for infants of mothers with and without malaria at delivery. Birth weight, length, and head circumference were significantly reduced if mothers were parasitaemic at delivery. Figures 1–3 present the centile curves for the same variables stratified by the presence or absence of malaria parasitaemia. Centile values for both weight and length were consistently lower in babies of parasitaemic compared with non-parasitaemic mothers. The 50th centile ponderal index of babies did not differ by maternal malaria status, indicating that growth was proportionate (fig 3). Gestational age did not differ by maternal malaria status (38.4 v 38.6 weeks, p = 0.07).

Figure 4 presents the centile curves for birth weight for primigravidae and multigravidae separately. Mean birth weight was lower (p=<0.001) at all gestational ages in infants of primigravidae. Similarly, birth length, head circumference, and mid-upper arm circumference were lower at all gestational ages in infants of primigravidae (data not shown). Figure 5 presents birth weight centile curves in relation to maternal HIV status. There was little difference between growth patterns of babies born to HIV positive or negative mothers, before 38 weeks gestation, although the fall off in centiles after 38 weeks in babies of HIV positive mothers was significant (p<0.05). Centiles also did not differ between babies with or without fetal anaemia (p>0.1) (data not shown).

Mean z scores for W/A and L/A were significantly different between babies of adolescents and non-adolescents, between primigravidae and multigravidae, and in relation to placental or peripheral parasitaemia at delivery (table 2). Differences in anthropometric indices in relation to the infant’s sex were consistent for different maternal characteristics, except for maternal HIV status. This showed that female infants of HIV positive mothers were likely to be lighter (p = 0.055) and shorter (p = 0.095). Although the effects are not statistically significant, there is a trend that girls born of HIV positive mothers are lighter and shorter.

Univariate analysis showed placental and/or peripheral malaria at delivery was associated with low W/A and L/A, but not with low W/L. Maternal HIV infection was associated with low W/A but not low L/A. Primigravidae adolescence and fetal anaemia were associated with low W/A and L/A.

The following risk factors were included in the multivariate analysis: infant sex, fetal anaemia, adolescent status, parity, maternal malaria and HIV status, and gestational age. In this analysis maternal HIV infection (AOR 1.76 (95% confidence interval 1.04 to 2.98)) and primigravidae (AOR 1.83 (1.10 to 3.05)) were associated with low W/A controlling for the covariates. Placental or peripheral parasitaemia at delivery (AOR 1.73 (1.02 to 2.88)) and primigravidae (AOR 2.13 (1.27 to 3.59)) were independently associated with reduced L/A.

DISCUSSION
Mean birth weight and length were significantly lower in babies of mothers with delivery malaria. This reduction is probably associated with placental insufficiency resulting from malaria, which leads to intrauterine growth restriction.22 23 This is illustrated in fig 1, which shows that the mean birth weights of babies born to mothers with malaria at delivery were lower across all gestational ages. Malarial lesions probably lead to inadequate uteroplacental24 and umbilical blood flow, and alterations in the transplacental transfer of glucose or production of fetal insulin. Placental lactogen, somatomedin, and somatomostatin-like substances, which influence fetal growth, may be affected.25 26 Disturbed placental folate metabolism may be important.27 Gestational age did not differ significantly between mothers with or without delivery malaria. This suggests that, in this population, malaria was associated with low birth weight primarily through growth restriction, leading to proportionate fetal growth retardation. This is supported by the similarity of the ponderal index centiles for babies of mothers with and without malaria at delivery (fig 3) and by the univariate analysis, as malaria at delivery was independently associated with low W/A and L/A but not with low W/L (table 2).

Mean birth weight and length were significantly lower from about 30 weeks gestation in babies whose mothers were primigravidae. Although it is universally recognised that the mean birth weights of primigravidae are lower,28–30 far less has been reported on the association between parity and other anthropometric measurements. Parity differences have been considered to be enhanced by greater efficiency.
of the uterine circulation in second and subsequent pregnancies. However, in malarious areas, primigravidae have a much higher incidence of falciparum malaria than multigravidae, and this exposure must increase the magnitude of the normal parity difference observed for fetal size between parity groups as has been reported in this population. Fetal size for all newborn anthropometric variables for this population, including head and arm circumference, is considerably below that of a Swedish reference population (unpublished work).

Anthropometric measurements did not differ significantly between babies with or without fetal anaemia. In Brazil, Rondo and Tomkins assessed the effects of fetal anaemia on intrauterine growth restriction and also found no significant association between fetal anaemia and growth restriction (OR 0.66 (95% confidence interval 0.46 to 0.95)). We have previously reported no significant association between fetal anaemia and growth restriction in intrauterine growth restriction and also found no significant association of HIV with growth restriction for both sexes combined, birth weight (p = 0.04) and W/A (p = 0.03) differed significantly according to maternal HIV status, and this was more apparent from 38 weeks gestation. This may indicate that mother to child transmission of HIV primarily occurs in late gestation.

CONCLUSIONS
In this area of Malawi, chronic fetal growth restriction occurred primarily due to falciparum malaria and resulted in proportionate changes in fetal weight and length. HIV infection may have a late gestational effect on fetal size. To improve fetal growth in developing countries, health interventions should prioritise malaria control in pregnancy, as well as preventing mother to child transmission of HIV.

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REFERENCES
Fetal growth, malaria, and HIV