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ORIGINAL RESEARCH



A 14-day follow-up of adult non-malarial fever patients seen by mobile clinics in rural Malawi

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Abstract

Background

While health providers consistently use malaria rapid diagnostic tests to rule out malaria, they often lack tools to guide treatment for those febrile patients who test negative. Without the tools to provide an alternative diagnosis, providers may prescribe unnecessary antibiotics or miss a more serious condition, potentially contributing to antibiotic resistance and/or poor patient outcomes.

Methods

This study ascertained which diagnoses and treatments might be associated with poor outcomes in adults who test negative for malaria. Adult patients for rapid diagnostic test of malaria seen in mobile health clinics in Mulanje and Phalombe districts were followed for 14 days. Participants were interviewed on sociodemographic characteristics, health-seeking behaviour, diagnosis, treatment and access to care. Mobile clinic medical charts were reviewed. Two weeks (± 2 days) following clinic visit, follow-up interviews were conducted to assess whether symptoms had resolved.

Results

Initially, 115 adult patients were enrolled and 1 (0.88%) was lost to follow-up. Of the 114 adult patients remaining in the study, 55 (48%) were seen during the dry season and 59 (52%) during the wet season. Symptoms resolved in 90 (80%) patients at the 14-day follow-up visit ($n=90$) with the rest ($n=24$) reporting no change in symptoms. None of the patients in the study died or were referred for further care. Almost all patients received some type of medication during their clinic visit (98.2%). Antibiotics were given to 38.6% of patients, and virtually all patients received pain or fever relief (96.5%). However, no anti-malarials were prescribed.

Conclusions

Mobile clinics provide important health care where access to care is limited. Although rapid tests have guided appropriate treatment, challenges remain when a patient's presenting complaint is less well defined. In rural areas of southern Malawi, simple diagnostics are needed to guide treatment decisions.

Key Words: Non-malarial fever, Malawi, febrile illness, mobile clinics, diagnostic tools, antibiotic resistance

Introduction

Malaria has long been thought to be the cause of fevers in many areas in sub-Saharan Africa. However, with the introduction of relatively accurate malaria rapid diagnostic tests (mRDT), fewer health providers are treating malaria solely on clinical symptoms¹. While this has improved outcomes for those who have malaria, those who test negative by mRDT are left without a clear diagnosis or treatment plan². Further, misdiagnosing fevers has important negative personal and public health consequences. The misdiagnosis of fever results in non-treatment of the actual etiology—upper respiratory tract infections (typically viral), lower respiratory tract infections (including bacterial/viral/fungal pneumonias, bronchiolitis and tuberculosis [TB]), gastroenteritis, schistosomiasis, intestinal nematode infections, genitourinary tract infections, meningitis, bacteraemia, and HIV and its associated illnesses—which may lead to significant morbidity and death³. Misdiagnosis thus leaves a patient vulnerable to worsening of the underlying illness, and allows for transmission of communicable diseases, such as TB. Consideration of the differential diagnoses of febrile illnesses is critical, though it is often difficult to differentiate the etiology of the fever in the absence of definitive diagnostic

tests⁴. Although there is research showing that fevers in hospitalised African adults and children are often associated with HIV-induced immunosuppression, upper respiratory illness and environmental exposures such as leptospirosis, rickettsiosis and brucellosis, little is known about causes of fever in primary health care facilities, where most patients are seen in low income countries⁵. Further, patients with mild respiratory symptoms and low-grade fever are often treated with antibiotics, despite the absence of signs/symptoms of a bacterial infection, and inappropriate use of antibiotics is known to contribute to the development of antibiotic resistance⁶. Such inappropriate treatment may indicate a lack of provider awareness of other causes of fever and how to manage them, a distrust in mRDT validity, or concern for patient well-being given limited access to health care services and follow-up.

Without another alternative diagnosis to offer or the ability to follow-up with a patient, health care providers may fail to prescribe appropriate treatment (or offer inappropriate treatment) for non-malarial fevers. The first step in improving management of febrile illness is to determine patient outcomes for mRDT-negative patients. This study explored clinical outcomes in mRDT-negative febrile adults

seen by mobile health clinics in southern Malawi. The overall objective of the study was to describe factors associated with incomplete symptom resolution in adult patients who test negative for malaria.

Methods

Overview

This study aimed to identify and describe mRDT-negative adult patients who are at risk for worsening illness in the 2-week period following a visit to a mobile health clinic. This article contains data from a larger prospective cohort study of people with non-malarial fevers who received care at the Global AIDS Interfaith Alliance (GAIA) mobile clinics in Muloza (Phalombe District) or Nkanda (Mulanje District), shown in Figure 1, during either the dry or wet seasons (August–September 2016 and November–December 2016, respectively). The methods and rationale of the parent study are reported on elsewhere⁷. Briefly, participants who presented at the clinics with a complaint of general illness and an axillary temperature of 37.5°C or reported a history of fever in the last 48 hours and were mRDT-negative were eligible for inclusion in the study. Patients were tested using the SDBioline mRDT which reports an overall sensitivity of 99.5% for *Plasmodium falciparum* and 92.6% for non-*P. falciparum* infection, and 98% specificity for *P. falciparum* and 100% for non-*P. falciparum* infections⁸. Patients in extremis or obtunded, already enrolled in the study, known to be unavailable or unreachable for follow-up, and who did not speak English or Chichewa fluently were excluded from recruitment.

Once identified, study staff fluent in Chichewa approached eligible patients to explain the purpose of the study and gain written consent. Patients who agreed to participate were then interviewed on sociodemographic characteristics, health-seeking behaviour, diagnosis, treatment, and access to care. Data were also extracted from GAIA mobile clinic charts. Two weeks (± 2 days) following the interview, local, trained field research staff went to the participants' homes to conduct follow-up interviews and assess the status of the presenting complaint and associated information.

This study reports on findings from all patients over the age of 14 years ($n=115$) who consented to participate during the course of the research. Assent was obtained for patients between the ages of 14 and 18 years. Presenting complaints were self-reported while clinical diagnoses, treatment and medication regimens were collected via clinical logbooks. HIV status was also self-reported because HIV diagnoses were tracked in separate logs that were unavailable to the researchers at the time of the study. Food insecurity was defined as having a family member or personal experience missing a meal when hungry in the past week. Medication non-compliance was defined as having, for any reason, not completed the entire course prescribed by GAIA, which was given to the patient in full as part of GAIA's service. Participant's medical status at follow-up was self-reported and defined as having resolved or got better compared with no change or worsening of symptoms.

Data analysis was completed using the R statistical package version 3.2.4 (The R Foundation, available from: <https://www.r-project.org/foundation/>). Initially, descriptive statistics were calculated to better understand cohort level demographic breakdowns. Next, bivariate analyses were run to explore the associations between participant characteristics

and their symptom status at follow-up (resolved/better vs no change/worsening). The Wilcoxon signed-rank test was used for continuous exposure variables as none were normally distributed. Fisher's exact test was used for categorical exposures due to the small sample size and occurrences of zero and cell counts of ≤ 5 . Measures of association include both P-values and unadjusted odds ratios (UOR) and 95% confidence intervals (95% CI) to show both direction and magnitude of associations.

Ethical approval

This study received ethical approval from the University of California San Francisco (UCSF) Human Research Protection Program (approval #16-19187) and the Malawi National Health Sciences Research Committee (Protocol #16/5/1594).

Results

During the study period, a total of 2893 patients (adults and children) were screened, 489 met inclusion criteria and 401 were enrolled. Of the enrolled subjects, 115 (28.6%) were over the age of 14 years. All 115 participants consented to join the study; however, 1 (0.88%) was lost to follow-up (Figure 2). This participant had no defining clinical or sociodemographic characteristics that stood apart from the rest of the cohort. Of the 114 patients over the age of 14 years remaining in the study, 55 (48%) were seen during the dry season and 59 (52%) during the wet season (Table 1). At enrolment, 80 (72%) of patients reported coming to the clinic because of fever and of those, 37 (46.3%) were febrile when tested by clinic staff. The remaining 34 participants (29.8%) did not report fever as a main reason for seeking care but were febrile when tested by clinic staff. Patient outcomes for those under the age of 14 years are reported elsewhere⁷. The majority of patients (80%) reported resolution of symptoms at 14-day follow-up ($n=90$) with the rest ($n=24$) reporting no change in symptoms. None of the patients in the study died or were referred for further care. Patients were primarily women (84%), and the median age of participants was 30.0 years (interquartile range [IQR]: 21.0–40.8), with a slight difference in age between patients who improved (28.5 years) compared with patients who reported no symptom resolution (35 years) (UOR: 1.03; 95% CI: 1.00–1.05). There were no differences between those who improved versus those who did not by season; roughly 50% of patients improved during each season, with the other half reporting no improvement during the follow-up period.

Only 16.7% of participants were new GAIA patients, with over 82% having already visited one of the clinics in the past year, though there was no association between these variables and patient outcomes. Patients were less likely to improve during the follow-up period if they reported commercial farming as their primary source of income compared with casual labour (UOR: 1.43; 95% CI: 1.43–16.44). Only four participants lived less than an hour from a government clinic, and participants were significantly more likely to have not gotten better if they lived more than 1 hour from the nearest GAIA clinic.

Acute respiratory illness was the most frequent diagnosis given by the mobile clinic clinicians (37.7%), followed by fever of unknown etiology (21.9%) and general body pain (16.7%) (Figure 3). The majority of patients (86%) received one diagnosis and 13.2% received two diagnoses (Table 2). There was no significant difference between the number of

Legend:

- Government hospital (Free)
- Government Health Facility (Free)
- CHAM Hospital (Fee)
- CHAM Health Facility (Fee)
- ▲ GAIA Offices
- Juma Clinic (est. 2014)
- Nhiramanja Clinic (est. 2014)
- Mlanda Clinic (est. 2010)
- Chitakale Clinic (est. 2008)
- Muloza Clinic (est. 2008)
- Nkhulambe Clinic (est. 2014)
- Chiwaliko Clinic (est. 2014)
- Namunda School

Towns and Villages in Phalombe District:

- TA JENATA
- TA CHIWALO
- TA KADUYA
- TA MKHUMBA
- TA NAZOMBE
- TA NKULAMBE
- TA NIEMA MGC-B VILLAGES
- TA MABUKA MULANJE MGC-A VILLAGES
- MULANJE MOUNTAIN (UNINHABITED)
- TA NTHIRAMANJA MULANJE D VILLAGES
- TA CHIKUMBU MULANJE D VILLAGES
- TA NKANDA MULANJE B VILLAGES MULANJE C VILLAGES
- TA JUMA

Phalombe District
POPULATION – 350,000

Villages Operational Dates:

- Mulanje A Villages (2008–2011)
- MGC-A Villages (2010–2013)
- MGC-B Villages (2010–2013)
- Mulanje B Villages (2001–2014)
- Mulanje C Villages (2013–2016)
- Mulanje D Villages (2014–2017)

Mulanje District
POPULATION – 550,000

```
graph TD; A[Screened for inclusion:  
n=2893] --> B[Met inclusion criteria  
n=489]; A --> C[Did not meet  
inclusion criteria  
n=2404]; B --> D[Included in overall study  
n=401]; B --> E[Declined or could not participate  
n=88]; D --> F[Over 14 years of age  
n=115]; D --> G[Under 14 years of age*  
n=286]; F --> H[Included in findings  
n=114]; F --> I[Lost to follow up  
n=1];
```

Screened for inclusion:
n=2893

Met inclusion criteria
n=489

Included in overall study
n=401

Over 14 years of age
n=115

Included in findings
n=114

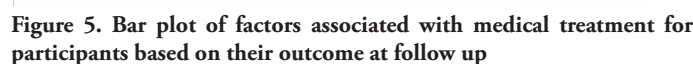
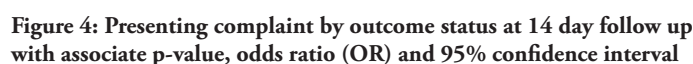
Did not meet
inclusion criteria
n=2404

Declined or could not participate
n=88

Under 14 years of age*
n=286

Lost to follow up
n=1

Those who did not report getting better were significantly more likely to self-report presenting to the clinic with non-specific ailment of 'looking or feeling poorly' (UOR: 12.71; 95% CI: 1.54–264.16) and receive a clinical diagnosis of a general neurological issue (Figure 4). Of those who did not get better, only 54.2% believed their diagnosis compared with 82.5% of those who did get better (UOR: 0.14; 95% CI: 0.04–0.45), and a quarter of those who did not get better also sought medication elsewhere during



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Table 1: Participant demographic characteristics overall and by symptom status at follow-up with associated(UOR), 95%CI, and p-value

	Total n (%)	Better n (%)	No change n (%)	UOR	95% CI	P-value
TOTAL	114 (100%)	90 (78.9%)	24 (21.1%)	–	–	
Season						
Dry	55 (48.2%)	44 (48.9%)	11 (45.8%)	0	Ref	0.82
Wet	59 (51.8%)	46 (51.1%)	13 (54.2%)	1.13	0.46–2.83	
Sex						
Female	96 (84.2%)	76 (84.4%)	20 (83.3%)	0	Ref	1
Male	18 (15.8%)	14 (15.6%)	4 (16.7%)	1.09	0.28–3.42	
Age						
Median (IQR)	30.0 (21.0–40.75)	28.5 (21.0–38.0)	35 (26.5–56.0)	1.03	1.00–1.05†	0.055
Marital status						
Married	66 (57.9%)	52 (57.8%)	14 (58.3%)	0	Ref	1
Unmarried	48 (42.1%)	38 (42.2%)	10 (41.7%)	0.98	0.38–2.42	
Level of education						
None	30 (26.3%)	23 (25.6%)	7 (29.2%)	0	Ref	0.93
Primary school	74 (64.9%)	59 (65.6%)	15 (62.5%)	0.84	0.31–2.43	
Secondary school	10 (8.8%)	8 (8.9%)	2 (8.3%)	0.82	0.11–4.32	
Source of income						
Casual labour	40 (35.1%)	36 (40.0%)	4 (16.7%)	0	Ref	0.16
Commercial farming	49 (43.0%)	33 (36.7%)	16 (66.7%)	4.36	1.43–16.44†	
Selling food or produce	6 (5.3%)	5 (5.6%)	1 (4.2%)	1.8	0.08–15.75	
Supported by family	3 (2.6%)	2 (2.2%)	1 (4.2%)	4.5	0.19–59.16	
Small shop or business	3 (2.6%)	3 (3.3%)	0 (0.0%)	0	–	
Tearoom/restaurant	1 (0.9%)	1 (1.1%)	0 (0.0%)	0	–	
Other	12 (10.5%)	10 (11.1%)	2 (8.3%)	1.8	0.23–10.72	
First visit to GAIA clinic						
Yes	19 (16.7%)	16 (17.8%)	3 (12.5%)	0	Ref	
No	95 (83.3%)	74 (82.2%)	21 (87.5%)	1.51	0.45–6.94	
Distance from GAIA clinic						
Less than 1 hour	82 (71.9%)	70 (77.8%)	12 (50.0%)	0	Ref	0.15
1–2 hours	26 (22.8%)	17 (18.9%)	9 (37.5%)	3.09	1.10–8.55†	
3–4 hours	0	0	0	0	–	
5–6 hours	1 (0.9%)	1 (1.1%)	0	8.75	1.32–71.99†	
Don't know	5 (4.4%)	2 (2.2%)	3 (12.5%)	0	–	

CI, confidence interval; GAIA, Global AIDS Interfaith Alliance; IQR, interquartile range; Ref, reference value; UOR, unadjusted odds ratio.

Table 2. Number of recorded diagnoses and medications prescribed by GAIA and UOR, 95% CI and P-values for measures of association

	Total n (%)	Better n (%)	No change n (%)	UOR	95% CI	P-value
Total number of recorded diagnoses						
0	1 (0.9%)	1 (1.1%)	0 (0.0%)	0	–	0.358
1	98 (86.0%)	75 (83.3%)	23 (95.8%)	0	Ref	
2	15 (13.2%)	14 (15.6%)	1 (4.2%)	0.23	0.01–1.26	
Medication(s) prescribed by GAIA						
Anti-epileptic						
No	111 (97.4%)	87 (96.7%)	24 (100.0%)	0	Ref	1
Yes	3 (2.6%)	3 (3.3%)	0 (0.0%)	–	–	
Anti-helminth						
No	113 (99.1%)	89 (98.9%)	24 (100.0%)	0	Ref	1
Yes	1 (0.9%)	1 (1.1%)	0 (0.0%)	–	–	
Anti-hypertensive						
No	109 (95.6%)	87 (96.7%)	22 (91.7%)	0	Ref	0.28
Yes	5 (4.4%)	3 (3.3%)	2 (8.3%)	2.64	0.33–6.86	
Antibiotic						
No	70 (61.4%)	54 (60.0%)	16 (66.7%)	0	Ref	0.64
Yes	44 (38.6%)	36 (40.0%)	8 (33.3%)	0.75	0.28–1.89	
Asthma drugs						
No	111 (97.4%)	87 (96.7%)	24 (100.0%)	0	Ref	1
Yes	3 (2.6%)	3 (3.3%)	0 (0.0%)	–	–	
ORS						
No	108 (94.7%)	86 (95.6%)	22 (91.7%)	0	Ref	0.6
Yes	6 (5.3%)	4 (4.4%)	2 (8.3%)	1.95	0.26–10.70	
Pain reliever						
No	4 (3.5%)	2 (2.2%)	2 (8.3%)	0	Ref	0.19
Yes	110 (96.5%)	88 (97.8%)	22 (91.7%)	0.25	0.03–2.18	
Supplement						
No	108 (94.7%)	86 (95.6%)	22 (91.7%)	0	Ref	0.6
Yes	6 (5.3%)	4 (4.4%)	2 (8.3%)	1.95	0.26–10.70	
Total medications given by GAIA						
1	20 (17.5%)	15 (16.7%)	5 (20.8%)	0	Ref	0.32
2	74 (64.9%)	57 (63.3%)	17 (70.8%)	0.89	0.30–3.07	
3	17 (14.9%)	16 (17.8%)	1 (4.2%)	0.19	0.01–1.34	
4	3 (2.6%)	2 (2.2%)	1 (4.2%)	1.5	0.06–19.35	
Did the patient seek care elsewhere after GAIA						
No	71 (62.2%)	59 (66.3%)	12 (50.0%)	0	Ref	0.22
Yes	42 (36.8%)	30 (33.7%)	12 (50.0%)	1.97	0.79–4.90	

CI, confidence interval; GAIA, Global AIDS Interfaith Alliance; ORS, oral rehydration salts; Ref, reference value; UOR, unadjusted odds ratio.

Discussion

Similar to findings from the paediatric subgroup in this parent study⁷, outcomes were excellent among adults in this setting, with 80% reporting resolution of symptoms at 14 days. However, occupation, distance from a mobile clinic, age and non-specific presenting complaint were associated with no change or worsening of symptoms among the participants in this study. The majority of participants who reported commercial farming as an occupation had symptoms that were unresolved or worsening during the 14-day follow-up period. In low- and middle-income countries, occupational hazards, including exposure to airborne particulates, are believed to contribute substantially to pulmonary disease⁹. This potential exposure among commercial farmers in the study may partially explain the delayed improvement or no change in condition reported especially in light of the fact that acute respiratory illness was the most frequent diagnosis.

Outcomes were not influenced by seasonality, but patients who lived far from the mobile clinics were less likely to improve, similar to other findings in the same area as well as results from the paediatric subgroup in this study^{7,10}. Although information on duration of symptoms prior to visiting the mobile clinics was not available, it is possible that those who live closer to the mobile clinic sites present earlier in an illness trajectory and, therefore, outcomes are better. Additionally, patients in this study who were older were less likely to improve, and although non-communicable diseases are thought to be increasingly responsible for morbidity in older adults¹¹, infectious disease remains a primary driver of illness in rural Malawi.

Similar to other non-malarial fever studies, this study found that most non-malarial fevers were assigned a diagnosis of respiratory illness and the majority of patients were given analgesics, followed by antibiotics^{12,13}. None of the adult patients in this study were given anti-malarials, unlike similar settings where providers may prescribe anti-malarials despite a negative mRDT¹⁴.

The significant association between distance to care and patient outcomes highlights the critical role that mobile clinics such as those of GAIA can play in the developing world. In a setting with no public transportation and where almost all rural inhabitants walk to clinics, only 3.5% of participants were less than an hour from a government clinic, with over 50% more than 3 hours away. In contrast, 71.9% of participants were less than an hour from a GAIA clinic. Interestingly, there was no significant association among participants between the number of prior visits to GAIA and their distance to the mobile clinics, potentially due to the fact that all but one participant lived within less than 2 hours of a clinic, making care accessible when needed. In underdeveloped regions where resources are scarce, patients may be unable to seek timely care due to financial or temporal reasons, allowing diseases to progress to a less easily treatable stage.

Despite being mobile clinics without the benefit of testing facilities associated with larger stationary facilities, the vast majority of patients seen at GAIA experienced symptom resolution following their visit. It is not surprising that those who did not get better were less likely to believe their diagnoses due to the perpetuation of their concerns. However, their ongoing complaints do not appear to be the result of incorrect diagnoses or treatment as they were no more likely to get better when seeking care elsewhere than

the mobile clinics. Instead, their lack of resolution may be associated with their less specific presenting complaint and/or generalised diagnosis. While these patients, due to their unresolved poor health, need to be considered when ramping up the diagnostic and treatment capabilities of mobile clinic facilities, they suggest that current testing and treatment algorithms used are successful and appropriate for the majority of patients served.

The study has several limitations worth noting. First, the follow-up period was 14 days, which may have prevented the researchers from capturing longer-term issues related to the underlying illnesses. In the future, an extended follow-up period may allow for a more nuanced understanding of non-malarial febrile illness, particularly around health-seeking behaviours outside the mobile clinics. Further, the majority of the patients in this study were women, limiting the ability to generalise to male populations. It is possible that men may seek care less often than women due to work responsibilities, because the mobile clinics visit sites during daytime working hours. It is interesting to note that this study did find commercial farming to be a risk for unresolved illness; therefore, future studies may seek to specifically enrol men with this occupation.

Conclusions

Despite many concerted efforts, the provision of equitable healthcare to rural populations in the developing world continues to face many structural, temporal and financial barriers. These are in addition to concerns that the care provided is of the highest quality. Rapid point-of-care testing kits and immediate provision of treatment for diseases such as HIV and malaria have been critical to addressing many of these issues. However, challenges remain when a patient's presenting complaint is less well defined. This study sought to assess the characteristics, diagnoses, treatment and outcomes in patients with mRDT-negative fevers presenting at mobile clinics in rural Malawi through a prospective cohort. The findings demonstrated that the clinics were by-and-large successful at addressing a wide variety of symptoms associated with a non-malarial fever leading to an improvement or complete resolution of symptoms. Although infectious disease remains an area where simple diagnostics facilitate sound treatment decisions, as the average lifespan increases in this setting, providers may need additional training in chronic diseases.

Availability of data and materials

The datasets analysed during the current study are available from the corresponding author on reasonable request.

Competing interests

The authors declare that they have no competing interests.

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