

Malaria Health Facility Survey SUMMARY REPORT

July 2008



















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Inquiries should be directed to:

Chief of Party Quality Health Partners 25 Senchi Street, PMB KIA Airport Residential Area Accra GHANA

++233 (21) 778-558 ++233 (21) 771-912 info@ghanaghp.org

The report will be available on-line at www.ghanaghp.org

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ACRONYMS

A/A Artesunate/Amodiaquine

ANC Ante Natal Care

CDC Centers for Disease Control
CHNs Community Health Nurses
CHPS Community Health Compound

FANC Focus Ante Natal Care
GHS Ghana Health Service

HBMF Home-based management of fever

IDSR Integrated Diseases Surveillance Response
IMCI Integrated Management of Childhood Illness

IPT Intermittent Preventive Treatment

IPTi Intermittent Preventive Treatment in Infants
IPTp Intermittent Preventive Treatment in Pregnancy

ITN Insecticide Treated Net

Jhpiego An Affiliate of John Hopkins University
LLIN Long-Lasting Insecticide Treated Nets

MA Medical Assistant
MIP Malaria in Pregnancy
MOH Ministry of Health

NGO Non Government Organisation

NHMS Nationwide Malaria Health Facility Survey NMCP: National Malaria Control Programme

OPD Out Patient Department

PMI President's Malaria Initiative

QHP: Quality Health Partners

RDT Rapid Diagnostic Test

SP Sulphadoxine Pyrimethamine
UNFPA United Nations Population Fund
UNICEF United Nations Children's Fund

USAID United States Agency for International Development WHO-AFRO World Health Organization Africa Regional Office

WHO World Health Organization

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A total of 36 surveyors collected the data for the survey. They were Justine Alornyo, Felix Doe, Simon Ahiataku, Grace Attimah, Mawuli Gyakobo, William Ansah, Mary Dartey, Emmanuel Ahiable, Mary Sagoe, Imoro Azumah, Beatrice Nyamekye, Harriet Charway, Ernestine Agyei Yobo, Faustina Asamoah, Kwasi Obeng, Kwaku Karikari, Florence Bedzrah, Lina Kyere, Theophilius Quaynor, William Akatsi, Beatrice Omenako, Rose Krah, Ananga Yamyolia, Vida Hervie, Boamah Boateng, Roseline Henry Udoh, Faustina Mwini, Rosina Asara, Patrick Atobrah, Agbanwa Gebiana, Margaret Azure, Sampson Sarsak, Adam Matthew, George Alira, Mery Amadu, Musa Awudu.

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EXECUTIVE SUMMARY

Introduction

The United States Government in December 2006, announced the selection of Ghana as one of the final eight countries in the five-year, \$1.2 billion President's Malaria Initiative (PMI) Program to rapidly scale up malaria prevention and treatment interventions in high-burden countries in sub-Saharan Africa. ¹ This initiative builds upon and collaborates with ongoing malaria control and eradication efforts of the National Malaria Control Program (NMCP) and its partners, including Roll Back Malaria and the Global Fund, as well as integrated maternal and child health strategies such as IMCI and Focused ANC, which address the problem of malaria in the high risk populations of children under age five and pregnant women. USAID, the WHO, UNICEF, UNFPA and others partner with the Ministry of Health/GHS and NMCP in these efforts.

The Quality Health Partners (QHP) project, managed by EngenderHealth, is one of four bilateral USAID Ghana health projects working to improve the health of Ghanaians, which is the overall goal of the Mission's strategic objective for health. QHP works from the policy level down to the health facility level to improve quality in the area of reproductive and child health. QHP assists nearly 200 facilities in 30 target districts in the seven southern regions of Ghana. To date, QHP has supported many malaria related activities, especially in the areas of Malaria in Pregnancy (MIP) and Case Management of Malaria, and therefore has data from the health facilities in the target area. However, QHP has neither collected data from "control districts" outside its target districts and facilities, nor worked in malaria related activities in the three Northern-belt regions. Since PMI is a nationwide program, there was the need to better understand the national situation with regard to malaria-related services in health care facilities. To inform QHP's PMI activity design and to assist partners, this nationwide health care facility survey was planned for the Year 1 Malaria Operational Plan.

The objectives of the study were to: 1) Describe the availability, functioning and quality of health and malaria treatment services in Ghana by assessing the quality of malaria case management for patients 2) Assess the preparedness (health system support) of health facilities to manage malaria and childhood illness, and 3) Assess the proportion of health workers who are treating malaria appropriately and have knowledge of the current protocols.

Methodology

The study was a cross-sectional survey. A nationally representative sample of facilities was selected for the survey that included all teaching and regional hospitals, and a sample of district hospitals, health centres and public as well as private clinics. A total of 60 facilities were assessed for the survey. Eight data collection tools adapted from those used in QHP facility baseline and mid-term surveys and WHO health facility survey were used. The tools comprised: Malaria Health Facility Audit, Patient Observation Tool, Provider Interview Tool, Re-examination of Patient and Client Exit Interview Tool, Assessment of Cases with Severe Malaria Tool, Laboratory Quality Control Tool, ANC Facility Audit Tool; and ANC Observation and Client Exit Interview Tool.

Data collectors were drawn from the GHS national and regional trainers and supervisors from QHP, WHO Country and Regional Office and the NMCP. A five day training including 2 days field practice was conducted. Data collection was completed in 2 weeks from 21 April – 3 May 2008. Field visits were undertaken by the primary investigators and other supervisors of the survey

during the fieldwork to ensure quality recording. The data were entered using EpiData and analyzed using SPSS. Data were weighted to reflect the probability of every patient having an equal chance of being selected.

Findings

Malaria Case Management

Children under 5 who should have been diagnosed with presumptive malaria during OPD visit on the day of the survey were correctly diagnosed in 80.1% of our observations-based on clinical assessment by the re-examiner. However, there was over diagnoses of presumptive malaria among 40% of patients in the over 5 population. In another vein, 42.2% of the under 5 who should have been presumptively diagnosed with malaria were not diagnosed as such, suggesting that many cases of the under 5's are being missed and not referred for treatment.

Overall, (91.4%) of all facilities are using Artseunate / Amodiaquine in the out patient dispensary. Encouragingly, all hospitals, be it teaching, regional, district or mission, are using Artesunate/Amodiaquine as the first line of drug for treating uncomplicated malaria. The figure drops to 92% of health centres and 82% of clinics doing same. Compared to government and mission facilities, fewer facilities (69.1%) from the private for profit sector are using this new regimen.

About 59% of all providers overall have knowledge of the correct dosage for Artesunate/Amodiaquine with variations from the different types of facilities. About 60% of providers from Government and mission hospitals as compared to 40% from maternity homes and 25% from CHPS zones have correct knowledge of the dosage. Only 1 in 5 of providers with correct knowledge was recorded for the private for profit facilities.

Treatment

Doctors were the least likely to prescribe Artesunate /Amodiaquine for patients diagnosed with uncomplicated malaria. For instance 48% of doctors as against 62% of Medical Assistants and 92% of nurses and midwives prescribed the regimen. However, patients seen by doctors were more likely than their other counterparts to have understood treatment information provided.

Malaria-focused counseling skills among providers were found to be weak. For instance, Only 10.1% of patients under 5 and 2.3% of patients over 5 diagnosed as having malaria were counseled on the management of malaria during consultations. Counseling was generally weak among all categories of staff with the nurses having some urge over Doctors, Medical Assistants and Midwives. None of the doctors observed gave counseling on management of malaria to caretakers of children under 5 seen.

Similarly, very few patients diagnosed as having malaria (10.8% for under 5 and 4.2% above 5) were counseled on ITN use.

Findings also show that less than 20% of patients or caregivers were told what illness their children had.

Also, only between 23%-28% of patients who were diagnosed with malaria were told what signs or symptoms should warrant a return to the facility.

Laboratory Capacity

22 out of the 59 facilities assessed had a lab. A 100 % of hospitals, 20% and 17% of clinics and maternity homes respectively had a lab.

Only 63.6% of facilities that had a lab also had all of the required materials (i.e functioning microscope, slides, giemsa stain, trained lab technician) needed for malaria microscopy.

Overall, less than 20% of patients (60 out of 314 patients) diagnosed with malaria had any laboratory tests ordered. Also, same day results were available for only 24% of patients for whom lab tests were ordered.

Assessment of Severe Malaria

Few patients (21% out of 32 cases) diagnosed as having severe malaria had a laboratory diagnosis to confirm malaria at admission. Compared to other facilities, regional hospitals were the most likely to have had laboratory confirmation of suspected severe malaria at admission.

Generally, patient records noted at least three signs of severe malaria or parasitemia in slightly more than half of the cases reviewed (50.5%).

On a positive note – patients who are being admitted for malaria generally get into care within one to three hours of arriving at the facility, suggesting the urgency of the cases is well understood

Preparedness of Health Facilities to Manage Malaria and Childhood Illnesses

About 23% of facilities had stock-out of co-packaged A/A on the day of the assessment. The privately run facilities were much less likely to have co-packaged A/A (50%).

Over 73% of facilities had copies of the standard treatment guidelines (2004). Also encouragingly over 76% of facilities had copies of the malaria treatment chart.

Effective tracking of malaria cases was a challenge due to poor filing systems for key data. Centralized data management systems where only one person in a facility controls data was also a limitation since in the absence of that person data cannot be accessed.

Encouragingly, about 70% of providers had benefited from a refresher training on malaria in the past three years. Fewer private sector facility providers (50%) also benefited refresher training in malaria within the same time frame. Not many providers (34.0%) however had had a refresher or training in IMCI in the last three years.

Malaria in Pregnancy

Encouragingly most facilities, including private sector facilities (94.1%), are providing SP for IPT.

Access to IPT2 and IPT3 remained a challenge in most facilities. Approximately, 62.0% of pregnant women received IPT1 in facilities during the past six months, whereas 38.1% received IPT2 and 36.6% received IPT3.

Overall, 27% of facilities experienced a stock out of SP in the past six months preceding the survey. Government facilities were the least likely to have stock-outs of SP (21%) as against 50% in mission hospitals and 50% private for profit facilities.

Recommendations

Malaria Case Management

- 1. Whereas there was over diagnoses of presumptive malaria among 40% of patients in the over 5 population, approximately 42.2% of the under 5 who should have been presumptively diagnosed with malaria were not diagnosed as such. As much as practicable, it is recommended that prescribers should order lab tests for all malaria suspected cases in patients over 5 years to ensure that patients with other diseases are not erroneously diagnosed and treated for malaria and secondly that malaria cases do not go undetected and hence untreated.
- 2. Since less than 21% of cases diagnosed as having severe malaria had a laboratory diagnosis to confirm malaria at admission, it is recommended that **laboratory** confirmation of severe malaria for both under 5 and adults be instituted and where microscopy is not feasible, RDT's use be employed.
- 3. Overall, 67% of patients diagnosed with uncomplicated malaria were treated with Artesunate Amodiaquine. Physicians are the category of providers found to be least compliant with use of AS/AQ with only 48% of those observed prescribing A/A for patients diagnosed with uncomplicated malaria. NMCP in collaboration with its stakeholders should develop and disseminate key messages targeting both providers, particularly physicians, and patients to provide education and support advocacy towards increasing provider and client compliance in the use of the combination therapy.
- 4. Although more than 90% of nurses and midwives treating uncomplicated malaria among patients prescribed Artesunate /Amodiaquine, fewer of the patients (37% and 51% of patients seen by nurses and midwives respectively) understood treatment information provided. Training programs and facilitative supervision activities would be useful in the various health facilities to help providers impart information to different categories of patients. Also, Malaria Job Aid and Treatment Chart (for wall) should be updated as needed, printed and widely disseminated, with appropriate orientation of providers.
- **5.** Correct knowledge of dosage of A/A was low among providers from maternity homes, CHPS zones and private for profit facilities. (See also # 4 above).
- 6. Fewer private for profit facilities (69%), compared to an overall figure of 92% of all facilities are prescribing Artesunate/Amodiaquine for the treatment of uncomplicated malaria. **NMCP should collaborate with the Association of private medical practitioners to**

- advocate and to strengthen capacity especially of private for profit facilities in treatment of uncomplicated malaria with the current treatment regimen by including them in training activities, continuing education and reporting.
- 7. Intensify public-private sector collaboration in malaria management generally. Advocate for intensive monitoring and facilitative supervision of private sector malaria control activities to ensure quality accurate diagnosis and treatment of cases.

Laboratory Capacity

- 7. Only 63.6% of facilities with lab had capacity for malaria microscopy. There is need to strengthen the capacities of the labs-particularly the regional and district hospital labs for effective malaria diagnosis. The labs should be assisted to procure needed materials and to hire lab technicians.
- 8. Same day lab results were available for 24% of lab requests made. In the long term, all hospital labs should be upgraded to offer 24 hour lab services. Within the short term, the use of RDTs that are highly sensitive and user friendly should be explored, particularly when lab facilities may not be open or otherwise fully operational.

Preparedness of Health Facilities to Manage Malaria and Childhood Illnesses

- 9. Stock-out of co-packaged A/A was recorded in a number of facilities, especially the privately run facilities. For instance, 73% of sites overall had co-packaged AS/AQ in stock on the day they were visited, with the figure at 33% for private for-profit facilities. Considering that A/ A is the main first line drug for the treatment of malaria and also that malaria is endemic in Ghana, it is important for facilities to reduce stock-outs of AA by working to improve stock management at the facility level, as well as overall commodity security.
- 10. Both public and private facilities continue to stock and prescribe monotherapies. NMCP should collaborate with the national and regional medical stores to take inventory and to withdraw from all facilities monotherapies for the treatment of malaria.
- 11. Close to a quarter of the facilities assessed did not have copies of the standard treatment guidelines (2004). Ensure that adequate copies of policies, standards and protocols are widely distributed to all facilities including the private sector facilities, since a number of facilities did not have copies Anti-Malaria drug policy and malaria treatment chart available in the OPDs (See also # 4 above).
- 12. Routine health data was found to be incomplete in a number of facilities and hence reports on malaria cases were not completely available. Management of health facilities should actively take keen interest in the use of routine data for decision making, so as to identify gaps and improve data capture, collation, dissemination and storage. Poor filing systems for key data was noted in a number of facilities.
- 13. Strengthen data management capacity at the facility level by training and updating Health Information Officers on indicators and their definitions and how data is related to overall regional and national goals. This will go a long way to help build a

- reputable national health management and information system (HMIS) to guide national policy decisions.
- 14. Data quality assessment (DQAs) should be planned and implemented to address the gaps in data capture and utilization within the facilities. Improve Monitoring and Evaluation Capacity of the NMCP to ensure that DQAs are done periodically- at least once in a year, feedback shared with facilities involved, timelines developed to address gaps and periodic monitoring instituted to ensure that gaps are addressed..

Malaria in Pregnancy

- 15. Whereas 62% of pregnant women received IPT1, less than 40% of them had IPT2 and IPT3. Identify and implement strategies to improve use of IPT2 and IPT3 among pregnant women. Distribution of doses of IPT2 and IPT3 at the community level by Community Health Nurses is a promising practice that has been identified in some districts of Ghana as a way of increasing IPT coverage in a facility- This could be explored further and up-scaled. Community Health volunteers could be useful in sensitizing women to access the services.
- 16. Emphasize other preventive measures of MIP apart from IPTp, ie use of ITN
- 17. Improve quality of counseling for case management and MIP (See also # 4 above).
- 18. Stock-outs of SP in facilities, especially private sector clinics and maternity homes remain fairly widespread. Reduce stock-outs of SP particularly in private sector clinics and maternity homes by improving stock management at the facility level, as well as overall commodity security.

I. Background:

In December 2006, the United States Government announced that Ghana had been selected as one of the final eight countries in a five-year, \$1.2 billion initiative to rapidly scale up malaria prevention and treatment interventions in high-burden countries in sub-Saharan Africa. ¹ This program to reduce the burden of malaria is called the President's Malaria Initiative or PMI.

PMI was officially launched in Ghana in December 2007 at a grand durbar in Agona Abodum, Central Region. PMI builds upon and collaborates with ongoing malaria control and eradication efforts of the National Malaria Control Program (NMCP) and its partners, including Roll Back Malaria and the Global Fund, as well as integrated maternal and child health strategies such as IMCI and Focused ANC, which address the problem of malaria in the vulnerable populations of children under age five and pregnant women. USAID, the WHO, UNICEF, UNFPA and others partner with the Ministry of Health/GHS and NMCP in these efforts.

The Quality Health Partners (QHP) project, managed by EngenderHealth, is one of four bilateral USAID health projects working to improve the health of Ghanaians, which is the overall goal of the Mission's strategic objective for health. QHP works from the policy level down to the health facility level to improve quality in the area of reproductive and child health. QHP does this through a strong collaboration with the Ministry of Health / Ghana Health Service in providing training, procuring of equipment and supplies, through facilitative supervision and by supporting the development, dissemination and implementation of standards and guidelines. Working through the regions and districts, QHP assists nearly 200 facilities in 30 target districts in the seven southern regions of Ghana. PMI is considered a nationwide program, so this will mean an expanded geographic focus for QHP beyond the present target districts, for certain activities such as this survey.

To support the work of PMI in Ghana, QHP was asked to undertake a number of tasks:

Malaria in Pregnancy (MIP)

- Update pre-service training curricula for MIP and for pre and in-service training for MIP.
- Work with ACCESS to support the provision of quality, comprehensive and integrated FANC services to enhance pregnant women's use of Long-Lasting Insecticide Treated Nets (LLIN), to complete Intermittent Preventive Treatment in Pregnancy (IPTp) and receive adequate education on malaria (including training materials and training health workers)

Case Management

- Update curricula, support pre and in-service training including private chemical sellers, as well as public and private sector facilities.
- Support NMCP in updating policy for HBMF (home-based management of fever), including advocacy for policy makers.
- Assess implementation of the new drug policy and malaria case management, IPTp use and health worker performance in sick child clinics, ANCs and in-patient facilities.

IPTi

 Sustain and build upon the IPTi programs in Tamale, Navrongo and or Kumasi and, consistent with evidence and decisions by the WHO and others, work with the MOH/NMCP to develop a written IPTi policy. NGO Collaboration and Capacity Building

• Improve NMCP capacity for program management and supervision at the zonal, regional and district levels in collaboration with in-country partners.

To date, QHP has supported many of these activities, especially in the areas of MIP and Case Management of Malaria, and therefore has good data on the health facilities in the target area. However the project has not collected data from "control districts" outside the target districts and facilities, nor has it had malaria work in the three Northern zone regions. Therefore – to inform QHP's PMI activity design and to assist partners to better understand the national situation with regard to malaria-related services in health care facilities, a nationwide health care facility survey was conducted as part of Year 1 Malaria Operational Plan.

1.1 Objectives of the Study:

- 1. To describe the availability, functioning and quality of health and malaria treatment services in Ghana.
 - a. the quality of malaria case management for patients (with emphasis on children under-five, in-patient and outpatient care)
 - b. the quality of care for malaria in pregnancy (prevention and management, with emphasis on IPTp)
 - c. the preparedness (health system support) of health facilities to manage malaria and childhood illness
 - d. by type of facility (CHPS Zone, Maternity Home, Clinic, Health Centre and Hospital)
- 2. To assess the proportion of health workers who are treating malaria appropriately and have knowledge of the current protocols
 - a. By type of provider (physician, MA, nurse, midwife)
 - b. By type of malaria (uncomplicated and complicated).

1.2. Study Design and Methodology:

The study was a cross-sectional survey consisting interviews with providers at the selected facilities in the sample, observations of service provision by providers, an audit of staffing, equipment and procedures routinely conducted at the facility level, exit interviews with caregivers of sick children and re-evaluation of clients (sick children) by expert supervisors (for comparison of actual care against the "gold standard"). There were observations of an ANC consultation, exit interview with ANC clients and in facilities where severe malaria is managed, there was observation of the care of these patients.

For this study, clinical case definition for malaria was based on national guidelines for adult and children under five years. The re-examiner based their diagnosis on patient history and examination. The criteria used in presumptively diagnosing malaria for adults were as follows: Patient with fever (history or temperature ≥ 37.5 or at least three non-fever symptoms (i.e. headache, joint pain, nausea/vomiting, chills/rigors, fatigue/general malaise, diarrhea, dizziness and anemia) then the patient is deemed to have suspected malaria. For patients under 5 years: fever (history or temperature ≥ 37.5) and other symptoms like diarrhea, inability to feed, vomiting everything and cough/difficult breathing (See under Appendices, Section 8; A2 Patient Observation form and A4 Interview and Re-examination of Patient Tool).

Data collection tools used for this study were adapted from those used in a QHP facility baseline and mid-term studies, which were also adapted from those used in the 2002 Service Provider Assessment. The WHO health facility survey tool was also reviewed as part of the tool development process. A working group comprising representatives from, GHS, the NMCP, QHP and WHO/Ghana developed the tools for the survey. In all eight tools were used. Further details on how the tools were administered in the field are given in (See under Appendices, Section 7 for details). The tools comprise the following:

- A-1: Malaria Health Facility Audit (review of staffing, supplies and equipment, Child Health and data management at the facility level).
- A-2: Patient Observation Form Observation of patient consultation
- A-3: Provider Interview Tool on malaria practice and knowledge
- A-4: Re-examination of patient and Client Exit Interview
- A-5: Assessment of Cases with Severe Malaria
- A-6: Laboratory Quality Control Tool
- B-1: ANC Facility Audit (review of staffing, supplies and equipment, procedures and data at the ANC clinic)
- B-2 ANC Observation and Client Exit Interview Tool

1.3 Training of Data Collectors and Field Work

Data collectors were health professionals (Doctors, Pharmacists, Medical assistants, Nurses and Midwives) drawn from among the GHS national and regional trainers and supervisors from QHP, and WHO Country Office and the NMCP.

A five day training of the data collectors was conducted from 14-19 April, 2008 and this included two days field practice. A reference guide for the instruments was used to ensure standard responses. Data collection was completed in 2 weeks from 21 April – 3 May 2008. Field visits were undertaken by the primary investigators and other supervisors of the survey during the fieldwork to ensure quality responses.

All interviews with providers were conducted in English, but the exit interviewers conducted with ALL patients whose care was observed were conducted in local languages of the caregivers' choice. The survey instruments did not have written translation because most people do not *read* the local language(s). The survey team was trained to simply formulate the questions in the local language for the benefit of respondents.

1.4 Sampling

The sample was selected to provide national representation of the health facilities that provide malaria treatment services. These included various types of hospitals, health centres and public as well as private clinics. Private pharmacies, psychiatric hospitals, specialist hospitals and clinics and dental clinics were not included in the sampling frame.

In order to have a nationally representative number of facilities for the survey, a total of 52 facilities was to be sampled. This number was calculated using the MEASURE Evaluation Project's Sampling Manual for Facility Surveys⁷ using the following formula and assumptions:

$$n = 3.84 f q$$

$$\frac{V^2 p}{V^2 p}$$

Where

n =the sample size we wish

p = the anticipated proportion of facilities with the attribute of interest (in the case of this survey we used .65 – which is less than the expected proportion of facilities that are expected to be using ACTs, but higher than the expect proportion of facilities where providers will be treating malaria appropriately. Given the numerous indicators of interest and the need for a reasonable sample size, this was the percentage finally decided upon by the working group.

q = is equal to (1-p) or in our case .35

f = is the design effect. In the case of this survey we used 1.0 – because our surveyed facilities will be drawn completely from the GHS list frame (list of health care facilities) and stratified and not subject to clustering.

 V^2 is the relative variance (square of the relative error). In the case of this survey the relative variance was set to .20 to yield relative error of $\pm 10\%$.

3.84 is the square of the normal deviate (1.96) needed to provide an estimate at the 95 percent level of confidence.

So:

n = 52

Within the national sample, the service delivery points were arranged by type of facility before a random start was chosen to systematically select the required sample using an equal probability sample (*epsem*) to the number of facilities of that type. A complete list of all health facilities in Ghana in 2007 was used to make the selection. In addition to this sample, a census of regional hospitals and teaching hospitals were planned for inclusion in the sample to ensure that these high volume facilities are adequately represented.

Table 1: Sample of Facilities for the Malaria Health Facility Survey

Background	Total Number of	Percentage of	Number of
Characteristics	Facilities in the	Total Sample	facilities proposed
	dataset		for the sample
CHPS	289	9.6	5
CLINIC	1287	43.0	22
HEALTH CENTRE	735	24.5	13
DISTRICT AND OTHER HOSPITALS	273	9.1	5
MATERNITY HOME	389	12.9	7
POLYCLINIC	10	.3	0
REGIONAL HOSPITAL	10	.3	10
TEACHING HOSPITAL	2	.06	2
Total	2998	100.0	64

1.5 Data Entry and Analysis

The data were entered using EpiData and double entry on all of the data was performed. All errors were checked and corrected. Data cleaning, merging of datasets resulting from the instruments and analysis was conducted using SPSS. Data were weighted to reflect the probability of every patient who visited the facility between 9.00 and 15.00 hours having an equal chance of having been selected. In addition, patient observations and findings from facilities were weighted to reflect the probability of having been selected from the overall set of health facilities in Ghana. Unless otherwise stated in the report, all results presented were weighted.

2.0. Findings

2.1. Health Facilities and Patients included in the Survey

A nationally representative sample of facilities was selected for the survey that included a census of all teaching and regional hospitals. A total of 60 facilities were assessed for the survey including a facility audit, patient observations and re-examinations, assessments of inpatient care for malaria and laboratory assessments.

Table 2: Actual Numbers Facilities and Patients in the MHFS

Type of Facility	Number of facilities surveyed	Number of OPD patients observed and re-examined		Number of providers interviewed	Number of ANC clients observed and
		Under 5	Above 5		interviewed
		Type of Fa	acility		
Teaching and Regional Hospitals	12	41	73	20	150
District and Other Hospitals	7	20	63	7	51
Health Centres	13	33	68	13	5
Clinics	15	37	107	17	31
Maternity Homes	6	4	15	7	11
CHPS Zones	7	4	8	7	10
		Operating A	L uthority		
Government	43	102	222	51	194
Mission	7	25	69	8	37
Private for Profit	10	12	63	12	23
Total Number	60	139	324	71	261

A total of 60 facilities were included in the survey including a census of all teaching and regional hospitals in the country and a sample of other types of facilities. A total of 633 patients (not shown in table) were observed, but only 464 were both observed and re-examined. Of patients who failed to return for a follow-up examination, the majority of these patients were being seen in the hospitals and larger facilities included in the survey and the majority of the patients (77.7%) were over 5 years of age. Most of those lost to follow up were in government facilities and were patients seen by either doctors or medical assistants. Those lost to follow up were not included in many parts of the final analysis.

All consenting providers who were observed were interviewed after the observations concluded (n=71).

Ante-natal care (ANC) patients were also observed and interviewed for the survey. Although a total of 261 clients were observed and interviewed, the majority of these clients were from hospitals, which are more likely to offer ANC services at least five days a week. Capturing observations of clients at health centres and other types of facilities was more challenging because these facilities still often have a specific day for services which may not have corresponded to the day the assessment was made.

2.1: Malaria Case Management

Key indicators for the survey were the percentage of patients who were diagnosed with malaria who were found to have malaria upon re-examination (Positive Predictive Value of diagnosis) and the percentage of patients who were not diagnosed with malaria who truly did not have malaria (specificity of the diagnosis). In both cases – these indicators were measured using presumptive diagnosis, because laboratory results were not often ordered for patients who were observed and were rarely available within the same day at facilities (see Table 9 for details). The results are presented in Table 3 disaggregated by age.

Table 3: Weighted Positive Predictive Value (PPV) and Negative Predictive Value (NPV) of Presumptive Malaria Diagnosis

% of patients diagnose actually had presumpting		% of patients not diagnosed with malaria that did not have presumptive malaria ¹		
Under 5	Under 5 Over 5		Over 5	
80.1	80.1 60.1		80.9	
n=139	n=139 n=324		n=324	

¹ Includes both uncomplicated and severe malaria. PPV was defined as the number of cases of malaria diagnosed by the provider that were also diagnosed by the re-examiner. Patients may have been diagnosed presumptively or with lab results if they were available. Most of the diagnoses were presumptive.

Children under 5 who should have been diagnosed with presumptive malaria by the provider at OPD were correctly diagnosed in 80.1% of our observations by trained expert 'Gold Standard'. However, there was over diagnoses of presumptive malaria among 40% of patients in the over 5 population, according to re-examination by expert. In another vein, 42.2% of the under 5 who should have been presumptively diagnosed with malaria were not diagnosed as such, suggesting that many cases of the under 5's are being missed and not referred for treatment. The positive predictive value of malaria diagnosis was high at 88.2% for children under 5 and 93.3% for those over five. The negative predictive values of diagnosis were also high in these groups at 86.2% for children under 5 and 91.3 for those over 5.

The majority of facilities in the country are using the new treatment regimen (91.4%) in the out patient dispensary. Of concern is that 31.9% of private for profit facilities are not using the new regimen. When providers in the OPD were asked the dosage of Artseunate / Amodiaquine for a child that weighs 20 kgs (and were allowed to refer to their references) only 58.6% of providers could answer the question correctly. Again, providers in private for profit facilities were the least likely to know the correct answer. Knowledge among providers in the CHPS Zones and maternity homes were also low (Table 4).

Table 4: OPD Use of New Artesunate/ Amodiaquine Treatment and Knowledge of Correct Dosage

	% of facilities u Artesunate / Ar for treatment o	modiaquine	% of providers that know the correct dosage (mg/kg) for this regimen ¹			
	%	n	%	N		
	Type of	Facility				
Teaching and Regional Hospitals	100.0	12	90.0	10		
District and Other Hospitals	100.0	7	66.7	6		
Health Centres	92.2	13	66.7	12		
Clinics	80.0	15	66.6	9		
Maternity Homes	100.0	6	25.1	4		
CHPS Zones	100.0	6	40.1	5		
	Operating	Authority				
Government	96.8	43	63.1	37		
Mission / Religious	100.0	7	79.6	5		
Private for Profit	69.1	10	20.1	5		
Total Number in Sample	91.4	59	58.6	46		

¹ Providers were asked if they knew the correct dosage for a 20 kg child.

2.3. Treatment

Prescription of Artesunate / Amodiaquine for uncomplicated malaria has become fairly commonplace in facilities throughout Ghana, with 66.8% of patients with malaria being prescribed the combination therapy (Table 5). Patients seen in government facilities were much more likely than those in the private sector to be prescribed the combination.

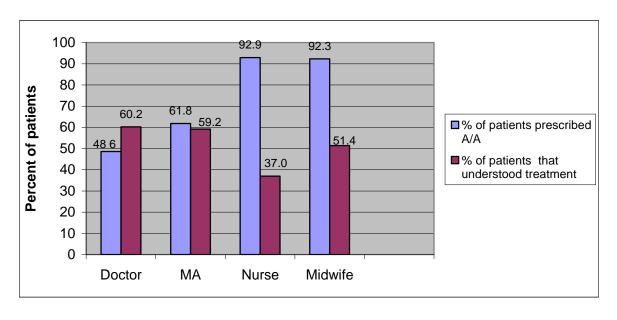
Also, the data show that patients seen and diagnosed as having uncomplicated malaria by midwives and nurses were in more than 90% of cases prescribed with Artesunate/Amodiaquine. Physicians are the least likely to be prescribed the new combination (Figure 1). During the provider interview, when asked, "how often do you prescribe Artesunate / Amodiaquine for malaria treatment?", 78.3% of physicians replied –always or often/sometimes. However their practice of prescribing for patients reveals a different scenario. For the five physicians who admitted that they rarely or never prescribed. Artesunate / Amodiaquine, three said they feared adverse reactions and two said they just hadn't adopted the new therapy yet.

Table 5: Percentage of patients diagnosed with malaria who received correct A/A and who correctly understood their treatment

Characteristic	Patients diagnosed and tre	Number of patients	
	% of patients with	% of patients	1
	uncomplicated malaria	diagnosed with	(n)
	prescribed Artesunate /	malaria that correctly	
	Amodiaquine treatment	understood their	
		malaria treatment	
	Type of Fa	acility	
Teaching and Regional	66.8	50.6	45
Hospitals	00.8	50.6	45
District and Other	59.0	61.0	44
Hospitals	33.0	01.0	77
Health Centres	95.1	61.0	56
Clinics	57.8	55.8	83
Maternity Homes	68.5	53.4	10
CHPS Zones	100.0	36.8	9
	Operating A	uthority	
Government	76.4	56.9	172
Mission/Religious	48.6	64.3	58
Private for profit	29.6	53.8	16
Total	66.8	58.5	247

¹ This includes all patients who were observed and who were diagnosed with uncomplicated malaria.

Figure 1: Percentage of malaria diagnosed patients treated with A/A and who understood their treatment by type of provider



The counseling skills of providers were also observed. The survey especially focused on basic communication about malaria. For patients that were diagnosed with malaria, overall counseling skills among providers were found to be very weak (Table 6). Providers rarely counseled patients on the management of fever during consultations, for children under 5

(10.1%) and for patients over 5 (2.3%). It is possible that counseling on the management of fever is done outside the consulting room; however it is also essential that this topic be addressed by providers. Similarly, very few patients diagnosed as having malaria were counseled on ITN use, for children under 5 (10.8%) and for patients above 5 (4.2%). Perhaps, of more concern, is the fact that less than 20% of patients or caregivers were even told what illness their children had. It will be difficult for health education messages on malaria to have an effect if patients are not even being told that they have the disease. Also, only between 23%-28% of patients who were diagnosed with malaria were told what signs or symptoms should warrant a return to the facility, meaning that few patients were counseled on the onset of severe malaria or severe disease and what to do if it occurs.

Table 6: Percentage of patients diagnosed with malaria who were counseled by providers on malaria related topics (all patients observed)

Type of		Pa	tients cou	nseled on	malaria r	elated top	ics	Patients counseled on malaria related topics Number of					
Facility	counse manage	tients eled on ement of ver	counse	atients eled on use	% of patients told what illness they have % of patients told signs and symptom to return to facility		Patients (n)						
Age of Patient	Under 5	Over 5	Under 5	Over 5	Under 5	Over 5	Under 5	Over 5	under 5	over 5			
	r	1		Туре	of Facility	/			_				
Teaching / Regional Hospitals	0.0	0.0	0.0	2.5	10.2	16.1	2.4	5.1	25	44			
District / Other Hospitals	5.6	3.2	0.0	1.3	5.6	8.1	27.1	9.7	21	47			
Health Centres	9.2	2.2	19.9	2.2	23.6	17.4	31.7	29.1	30	38			
Clinics	14.2	0.0	19.9	4.0	8.5	26.4	27.4	44.1	30	55			
Maternity Homes	0.0	9.7	0.0	29.0	0.0	71.1	0.0	0.0	2	9			
CHPS	71.4	0.0	28.6	40.0	71.4	0.0	28.6	53.3	3	7			
				Operati	ing Author	rity							
Govern- ment	8.7	0.6	7.7	2.6	13.7	13.8	35.7	27.3	83	137			
Mission/Rel igious	13.7	5.0	17.1	7.0	3.4	20.7	3.4	14.3	23	50			
Private for profit	17.7	0.0	35.1	0.0	35.1	71.3	0.0	40.2	5	12			
					of Provide								
Doctor	0.0	4.8	0.0	2.1	2.2	12.8	0.5	21.7	23	60			
MA	5.2	0.0	9.4	0.8	8.5	19.7	33.1	28.1	53	86			
Nurse	50.2	3.2	33.9	29.2	38.9	35.1	5.6	19.4	10	21			
Midwife	5.2	3.1	6.8	0.0	17.1	26.8	53.6	18.3	17	24			
Other	0.0	0.0	0.0	0.0	0.0	21.5	0.0	0.0	8	6			
Total	10.1	2.3	10.8	4.2	12.4	19.8	27.5	23.1	111	200			

2.4: Malaria Prevention in Pregnancy (IPT)

A record review on IPT practice was conducted in facilities that regularly use the Ghana Health Service ANC register to record IPT distribution. Service statistics on IPT in facilities were often not available for review on the day of the assessment. Of a total of 59 facilities audited, only 29 were able to provide complete service statistics for the previous 6 months on IPT distribution. Many midwives were not recording IPT use in the GHS ANC register. They explained that it is difficult to tabulate the IPT doses given during the month from the register for reporting— so they prefer to keep their IPT records separately, which means it is difficult for supervisors to assess how many missed opportunities there are and how systems might be improved.

From the service statistics that were available approximately 62.0% of pregnant women received IPT1 in facilities during the past six months, 38.1% received IPT2 and 36.6% received IPT3. Despite the limitation, these data provide a good estimate of the current status of IPT distribution in the country (Table 7).

Table 7: Service statistics on IPT from facilities (un-weighted)

% of pregnant women that received IPT1	% of pregnant women that received IPT2	% of pregnant women that received IPT3
62.0	38.1	36.6
(n=29)	(n=29)	(n=22)

Overall, most facilities (94.1%), including private sector facilities, are providing SP for IPT. Stock-outs of SP in facilities still remain fairly widespread, mainly focused in private sector clinics and maternity homes. The availability of the IPT training manual in most ANC units remains low with only 38.1% of facilities reporting that they had a copy. This could affect adherence to the national standard. Distribution of doses IPT2 and IPT3 at the community level by CHNs is a promising practice that has been identified in some districts of Ghana as a way of increasing IPT coverage in a facility. This community-based distribution practice was assessed during this survey and overall 39.8% of facilities are already doing this type of distribution. (Table 8).

Table 8: Number and Weighted Percentages for IPT practice in facilities surveyed.

Type of		IPT practice i	n facilities		Number of
Facility	% of facilities that	% of facilities	% of facilities	% of facilities	Patients
	offer ANC and	that have	that have the	use CHNs to	
	provide IPT with	experience a	IPT training	distribute	(n)
	SP.	stock out of	manual	IPT2 and	
		SP in the		IPT3 in the	
		past six		community	
		months	Cocility (
Tacabing and	T	Type of F	aciiity I	T	
Teaching and Regional	81.8	33.3	63.6	0.0	11
Hospitals	01.0	33.3	03.0	0.0	11
District and					
Other	100.0	16.8	50.0	33.3	6
Hospitals			33.3	33.3	
Health	04.0	40.0	44.7	50.0	44
Centres	91.6	18.2	41.7	58.3	11
Clinics	87.4	42.9	12.6	12.6	8
Maternity	100.0	39.9	60.1	0.0	6
Homes					
CHPS Zones	100.0	25.1	33.3	100.0	4
		Operating A			
Government	91.9	21.8	33.4	55.4	32
Mission/Religi	100.0	50.9	24.7	0.0	4
ous	100.0	50.5	27.1	0.0	7
Private for	100.0	33.0	67.0	0.0	6
profit		00.0	00	0.0	L
	T	T	T	T	
Total	94.1	27.4	38.1	39.8	46

2.5: Laboratory Capacity

All hospitals (Teaching, Regional and District hospitals) surveyed had laboratory facilities available and some clinics and maternity homes also had lab services available. However, of the twenty-two facilities that had labs, only 63.6% of them had all materials required for malaria microscopy, which included a functioning microscope, slides, giemsa stain and a trained laboratory technician. One facility did not have a functioning microscope and three of the facilities did not have a trained lab technician. Also on the day of the assessment two facilities did not have giemsa stain available (Table 9).

Of all patients who were diagnosed with malaria during observations, only 19.4% had any laboratory test for malaria ordered (Table 9). In the hospitals, only 29.1% of providers ordered any laboratory tests. Out of the patients that had blood film for malaria parasites ordered, few of them (23.6%) were able to receive their results on the same day. Most patients either did not return with their test results (39.9%) or did not have results available the same day (29.8%) (Not shown in Table)

.

Table 9 Percentage of Facilities with Labs and Patients Receiving Laboratory Services

Type of Facility	% of facilities that had a lab facility	% of facilities with labs that had all items for malaria microscopy ¹	% of patients observed & diagnosed with malaria who also had
	(n)	(n)	lab tests ordered
	Тур	e of Facility	
Teaching and Regional Hospitals	100.0 (11)	90.9 (11)	47.8 (68)
District and Other Hospitals	100.0 (7)	28.6 (7)	28.3 (68)
Health Centres	0.0 (13)	-	3.3 (68)
Clinics	20.0 (15)	33.1 (3)	15.3 (89)
Maternity Homes	16.7(6)	100.0 (1)	11.4 (11)
CHPS Zones	0.0 (6)	-	0.0 (10)
	Opera	ting Authority	
Government	38.1 (42)	68.8 (16)	19.0 (219)
Mission/Religious	28.6 (7)	50.0 (2)	17.9 (77)
Private for profit	40.0 (10)	50.0 (4)	33.9 (17)
Total	37.3 (n=59)	63.6 (n=22)	19.4 (n=314)

All items for malaria microscopy includes, functioning electric binocular microscope, Slides, Giemsa stain and a trained laboratory technician.

2.6 Assessment of Severe Malaria

In facilities that had in-patient capacity, an assessment of the care of severe malaria cases was made using a standardized form. The total number of cases reviewed was very small (n=32), so the results may not be generalizable. Findings show that very few suspected severe malaria cases had a laboratory confirmation of the condition at admission, and even so this was more likely to be done in the Regional hospitals. Generally, patient charts noted at least three signs of severe malaria or parasitemia in slightly more than half of the cases reviewed (50.5%). On a positive note – patients who are being admitted for malaria are generally getting into care within one to three hours of arriving at the facility, suggesting the urgency of the cases is well understood (Table 10).

Table 10: Percentages of patients reviewed for indicators on treatment of Severe Malaria

Type of	Patients reviewed on treatment of Severe Malaria				Number
Facility	% patients with lab	% of patients	% of patients	% of patients	of
	confirmed diagnosis	with three or	with severe	admitted in 1-	patients
ļ	at admission	more signs	malaria	3 hours of	(n)
		(or	prescribed	arriving at	
		parasitemia	correct	health care	
		noted on	treatment	facility.	
		chart)			
-	T	Type of Facili	ty		
Teaching	16.7	16.7	16.7	20.0	6
Hospitals	-	_	_		
Regional	56.3	81.3	31.3	46.7	16
Hospitals					
District /	00.0	50.0	0.0	400.0	40
Other	20.0	50.0	0.0	100.0	10
Hospitals		On a ratio a A sith a			
Operating Authority					
Government	16.7	16.7	16.7	20.0	6
Mission/Religi	18.5	51.4	1.4	97.7	22
ous					
Private for	25.1	50.0	0.0	100.0	4
profit	·				
Tatal	00.0	50.5	4.0	00.0	20
Total	20.9	50.5	1.0	98.0	32

2.7: Preparedness of Health Facilities to Manage Malaria and Childhood Illnesses

Availability of co-packaged Artesunate / Amodiaquine was good in the facilities with a total of 77.3% of facilities having stock on the day of the assessment. Of note is that clinics and privately run facilities were much less likely to have co-packaged Artesunate / Amodiaquine in stock, compared to the government or mission managed facilities.

Of the facilities that had co-packaged Artesunate / Amodiaquine available on the day of interview, a total of 12.4% had experienced a stock—out in the past six months prior to the survey. The problem of stock outs of the combination therapy appears to be centered in teaching and regional hospitals (Table 11).

Table 11: Availability and Stock outs of Artesunate / Amodiaguine at the facility level

Characteristic	Facilities with co-packaged Artesunate / Amodiaquine in stock		Facilities having A/A in stock on day of assessment that had a stock out of A/A in the last 6 months		
	%	N	%	N	
Type of Facility					
Teaching and Regional Hospitals	83.3	12	50.0	10	
District and Other Hospitals	71.5	7	0.0	4	
Health Centres	91.6	12	18.2	11	
Clinics	66.6	15	18.2	11	
Maternity Homes	60.1	5	0.0	2	
CHPS Zones	100.0	5	0.0	5	
Operating Authority					
Government	92.8	40	12.3	35	
Mission/Religious	71.0	7	0.0	3	
Private for profit	33.1	9	20.5	5	
Total	77.3	56	12.4	43	

Almost three quarters (73.3%) of facilities had current copies of the Standard Treatment Guidelines (2004) in their OPD areas, although the Standard Treatment Guideline is not the main reference for the new malaria drug treatment policy. Fewer (35.6%) facilities had copies of the latest Anti-Malaria drug policy available in the OPDs. Similarly, few (41.1%) facilities had copies of the IMCI chart booklet. More than three-quarters (76.3%) of facilities including half of the private sector facilities had copies of the malaria treatment chart. Generally private sector facilities had fewer of the standards and guidelines compared to the government facilities (Table 12).

¹

¹ Some concern has been expressed about the usefulness of this job aid, because the weight range for prescription is wide and may lead to overprescribing or underprescribing, especially in the lower weight categories. That this tool is widely available is good to know because the mechanisms for distribution could be replicated for future versions of the tool.

Table 12: Availability of Malaria Drug Policy and Treatment Protocols in Facilities

Type of Facility	Malaria Treatment Protocols in Facilities				Number of	
	% of facilities	% of facilities	% of facilities	% of facilities	Facilities	
	with the	with the Anti-	that had IMCI	that had the	(n)	
	Standard	Malaria Drug	Chart booklet	malaria		
	Treatment	Policy in	in OPD	treatment		
	Guidelines in	OPD		chart		
	the OPD					
Type of Facility						
Teaching and						
Regional	91.7	50.0	36.4	63.6	12	
Hospitals						
District and Other	85.7	50.0	33.3	71.4	7	
Hospitals	00.7	30.0	33.3	71.4	,	
Health Centres	76.9	30.8	61.5	84.6	13	
Clinics	60.0	13.3	40.0	73.3	15	
Maternity Homes	50.0	66.7	16.7	66.7	6	
CHPS Zones	71.4	28.6	42.9	100.0	6	
Operating Authority						
Government	81.4	34.9	47.6	81.0	43	
Mission/Religious	71.4	33.3	50.0	85.7	7	
Private for profit	40.0	40.0	10.0	50.0	10	
Total	73.3	35.6	41.1	76.3	76.3	

Overall, more than 70% of providers had some refresher training on malaria in the past three years, with a higher proportion of providers (85%) from Mission and religiously operated facilities leading the way. Providers from private sector facilities were the least likely (50%) to have had refresher training in malaria over the period (Table 13).

Only 34.0% of providers had had a refresher or training in IMCI in the last 3 years, with mission and religiously run institutions having the least access. Doctors were also less likely to have had any IMCI refresher training in the past three years (Table 13).

Table 13: Percentage of Providers with Training in Malaria Policy and IMCI

Characteristic	Providers with Training in	Number of			
	% of providers with	% of providers with	Number of Providers		
	refresher training in	training or refresher in			
	malaria in last 3 years	IMCI in last 3 years	(n)		
Type of Facility					
Teaching and Regional Hospitals	45.0	20.0	20		
District and Other Hospitals	57.1	28.5	7		
Health Centres	76.9	30.7	13		
Clinics	76.5	31.3	17		
Maternity Homes	57.1	28.5	7		
CHPS Zones	71.4	57.4	7		
Operating Authority					
Government	74.8	40.6	51		
Mission/Religious	85.1	16.5	8		
Private for profit	50.1	25.0	12		
Type of Provider					
Doctor	62.2	24.8	23		
MA	74.7	36.2	15		
Nurse	99.6	39.6	6		
Midwife	73.5	40.0	15		
CHN	60.0	60.0	5		
Other	51.0	0.0	6		
Total	70.5	34.0	71		

2.8: Malaria Burden in Health Facilities

During the survey, the service statistics from facilities surveyed were also collected. These service statistics reflect the number of reported cases (both presumptive and laboratory confirmed) of malaria as collected in the Out-Patient Department and In-Patient Department (in facilities that have that capacity). A total of 45 facilities (of the 60 surveyed) were able to provide complete basic service statistics on OPD attendance and malaria cases seen between April 2007 – March 2008. Of facilities that did not provide statistics, one was a teaching hospital, one was a regional hospital, two were district level hospitals, two were health centres, three were maternity homes, four were clinics and three were CHPS zones. Also, 57.3% of them were government operated, 7.3% mission operated and 35.3% were private for profit (weighted percentages).

,In total, the 45 facilities reporting their service statistics, are diagnosing roughly between 21,000 – 33,000 cases of malaria a month in OPDs, with average OPD attendance ranging between 88,000 – 119,000 patients a month (Figure 2).

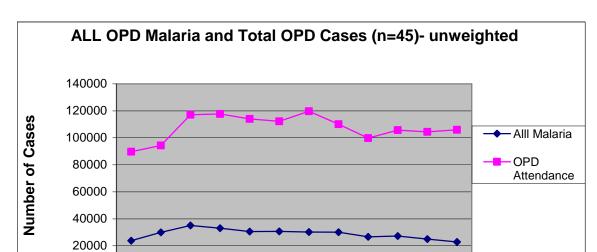


Figure 2: Total Burden of Malaria in Out Patient Dispensaries in Ghana – unweighted (n=45)

This burden of malaria in the OPDs as found during this survey, is less than what the Ghana Health Service (GHS) recently reported. Whereas malaria is reported as constituting 37.5% of the caseload in OPDs in the country (CHMS/GHS 2007), this survey found that the burden for all malaria cases ranged between 20.2% – 30.2% throughout one calendar year April 07- March 08). Of all malaria cases the burden of cases for children under five ranged between 23.0% - 26.1% for the same period (Figure 3). Since data were only collected for this one year period, comparisons with earlier time periods cannot be made for these facilities.

Oct- Nov- Dec-

07

07

Jan- Feb-

08

80

Mar-

80

0

Apr- May-

07

07

Jun-

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Jul-

07

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07

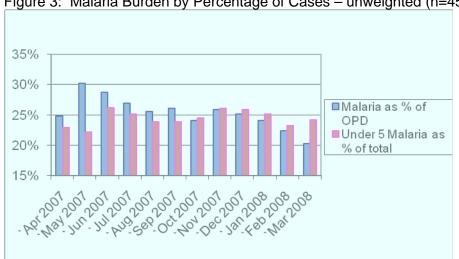
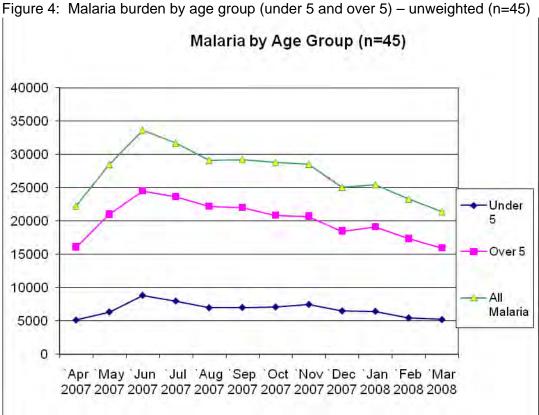


Figure 3: Malaria Burden by Percentage of Cases – unweighted (n=45)

May and June are the months with the highest rates of transmission in Ghana and the lowest months of transmission are February and March, which are traditionally the driest months of the year. (Figure 4).



The survey also looked the number of cases of malaria diagnosed in pregnant women. There were only 32 facilities for which data were available. There was a spike in cases in January

2008, but the reasons for this spike are unclear. Cases of malaria in pregnant women in these facilities as a whole average about 300 per month. The survey also collected information on the total number of anemia cases diagnosed in the facilities (all units) during each month for the past calendar year. Cases increase and peak immediately following the highest transmission months (June and July). See Figure 5 for details.

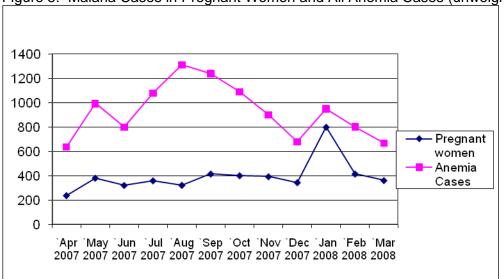
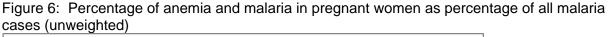
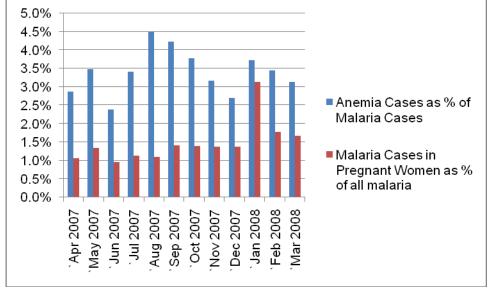


Figure 5: Malaria Cases in Pregnant Women and All Anemia Cases (unweighted) (n=32)

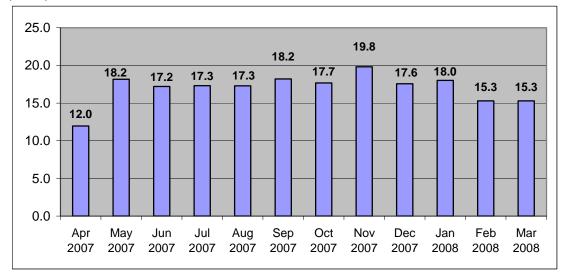
Overall the percentage of anemia cases as a percentage of the overall malaria cases remains below 5% nationwide and the percentage of malaria cases in pregnant women (from the limited data available) remain below 3% (Figure 6). However, these data must be interpreted with caution as they represent limited service statistics available at the facility level and may not necessarily correlate with actual prevalence in the community.





Finally the survey also looked at admissions for malaria as a percentage of the total admissions and found that malaria admissions do not have a spike following the trend in overall malaria cases, but generally remains between 17.2% - 19.8% of all admissions from May – January of each year. Between February and April of each year there is a brief lull in the percentage of cases admitted with malaria to between 12.0% - 15.9% of the total admissions (Figure 7).

Figure 7: Malaria admissions as a percentage of the overall total of admissions (unweighted) (n=13)



Recommendations

Malaria Case Management

- 1. While there was over diagnoses of presumptive malaria among 40% of patients in the over 5 population, approximately 42.2% of the under 5 who should have been presumptively diagnosed with malaria were not diagnosed as such. As much as practicable, prescribers should order lab tests for all malaria suspected cases in patients over 5 years to ensure that patients with other diseases are not erroneously diagnosed and treated for malaria and secondly that malaria cases do not go undetected and hence untreated.
- Since less than 21% of cases diagnosed as having severe malaria had a laboratory diagnosis to confirm malaria at admission, it is recommended that laboratory confirmation of severe malaria for both under 5 and adults be instituted and where microscopy is not feasible, RDT's use be employed.
- 3. Overall, 67% of patients diagnosed with uncomplicated malaria were treated with Artesunate Amodiaquine. Physicians are the category of providers found to be least compliant with use of AS/AQ with only 48% of those observed prescribing A/A for patients diagnosed with uncomplicated malaria. NMCP in collaboration with its stakeholders should develop and disseminate key messages targeting both providers, particularly physicians, and patients to provide education and support advocacy towards increasing provider and client compliance in the use of the combination therapy.
- 4. Although more than 90% of nurses and midwives treating uncomplicated malaria among patients prescribed Artesunate /Amodiaquine, fewer of the patients (37% and 51% of patients seen by nurses and midwives respectively) understood treatment information provided. Training programs and facilitative supervision activities would be useful in the various health facilities to help providers impart information to different categories of patients. Also, Malaria Job Aid and Treatment Chart (for wall) should be updated as needed, printed and widely disseminated, with appropriate orientation of providers.
- 5. Fewer private for profit facilities (69%), compared to an overall figure of 92% of all facilities are prescribing Artesunate/Amodiaquine for the treatment of uncomplicated malaria. NMCP should collaborate with the Association of private medical practitioners to advocate and to strengthen capacity especially of private for profit facilities in treatment of uncomplicated malaria with the current treatment regimen by including them in training activities, continuing education and reporting.
- 6. Intensify public-private sector collaboration in malaria management generally. Advocate for intensive monitoring and facilitative supervision of private sector malaria control activities to ensure quality accurate diagnosis and treatment of cases.

Laboratory Capacity

7. Only 63.6% of facilities with lab had capacity for malaria microscopy. There is need to strengthen the capacities of the labs-particularly the regional and district hospital labs for

- effective malaria diagnosis. The labs should be assisted to procure needed materials and to hire lab technicians.
- 8. Same day lab results were available for 24% of lab requests made. In the long term, all hospital labs should be upgraded to offer 24 hour lab services. Within the short term, the use of RDTs that are highly sensitive and user friendly should be explored, particularly when lab facilities may not be open or otherwise fully operational.

Preparedness of Health Facilities to Manage Malaria and Childhood Illnesses

- 9. Stock-out of co-packaged A/A was recorded in a number of facilities, especially the privately run facilities. For instance, 73% of sites overall had co-packaged AS/AQ in stock on the day they were visited, with the figure at 33% for private for-profit facilities. Considering that A/ A is the main first line drug for the treatment of malaria and also that malaria is endemic in Ghana, it is important for facilities to reduce stock-outs of AA by working to improve stock management at the facility level, as well as overall commodity security.
- 10. Both public and private facilities continue to stock and prescribe monotherapies. NMCP should collaborate with the national and regional medical stores to take inventory and to withdraw from all facilities monotherapies for the treatment of malaria.
- 11. Close to a quarter of the facilities assessed did not have copies of the standard treatment guidelines (2004). Ensure that adequate copies of policies, standards and protocols are widely distributed to all facilities including the private sector facilities, since a number of facilities did not have copies Anti-Malaria drug policy and malaria treatment chart available in the OPDs (See also # 4 above).
- 12. Routine health data was found to be incomplete in a number of facilities and hence reports on malaria cases were not completely available. Management of health facilities should actively take keen interest in the use of routine data for decision making, so as to identify gaps and improve data capture, collation, dissemination and storage. Poor filing systems for key data was noted in a number of facilities.
- 13. Strengthen data management capacity at the facility level by training and updating Health Information Officers on indicators and their definitions and how data is related to overall regional and national goals. This will go a long way to help build a reputable national health management and information system (HMIS) to guide national policy decisions.
- 14. Data quality assessment (DQAs) should be planned and implemented to address the gaps in data capture and utilization within the facilities. Improve Monitoring and Evaluation Capacity of the NMCP to ensure that DQAs are done periodically- at least once in a year, feedback shared with facilities involved, timelines developed to address gaps and periodic monitoring instituted to ensure that gaps are addressed..

Malaria in Pregnancy

15. Whereas 62% of pregnant women received IPT1, less than 40% of them had IPT2 and IPT3. Identify and implement strategies to improve use of IPT2 and IPT3 among

pregnant women. Distribution of doses of IPT2 and IPT3 at the community level by Community Health Nurses is a promising practice that has been identified in some districts of Ghana as a way of increasing IPT coverage in a facility- This could be explored further and up-scaled. Community Health volunteers could be useful in sensitizing women to access the services.

- 16. Emphasize other preventive measures of MIP apart from IPTp, ie use of ITN
- 17. Improve quality of counseling for case management and MIP (See also # 4 above).
- 18. Stock-outs of SP in facilities, especially private sector clinics and maternity homes remain fairly widespread. Reduce stock-outs of SP by improving stock management at the facility level, as well as overall commodity security.

APPENDICES

Section 1: Integrated Diseases Surveillance Response (IDSR)

Table 1A: Percentage of Facilities with Evidence of Use of Malaria Data

	% of facilities with a graph of malaria cases children <5 in last 2 qtrs of 2007	% of facilities with a graph of all malaria cases in last 2 qtrs of 2007	Total Number of facilities	% of facilities with graph of malaria deaths in last 2 qtrs of 2007	Total Number of facilities
	%	%	N	%	N
	Type of Facil	ity			
Teaching and Regional Hospitals	50.0	66.7	10	62.5	8
District and Other Hospitals	50.0	50.0	4	50.0	4
Health Centres	44.4	44.4	9	28.5	7
Clinics	39.9	20.1	5	0.0	4
Maternity Homes	50.0	50.0	2	50.0	2
CHPS Zones	0.0	0.0	3	0.0	3
	Operating Auth	ority			
Government	44.5	39.0	28	25.1	24
Mission / Religious	49.6	49.6	2	49.6	2
Private for Profit	0.0	0.0	3	0.0	2
Total Number in sample	39.1	34.8	33	25.0	28

Section 2: Laboratory

Of facilities that had labs 16 of 18 reporting maintained registers of the lab tests that they performed. However on the day of the assessment complete data were only available for blood films performed between April 2007 and March 2008 for 12 facilities (Table 2A).

Table 2A: Blood films done at the facilities between April 2007 and March 2008

	Apr 2007	May 2007	Jun 2007	Jul 2007	Aug 2007	Sep 2007	Oct 2007	Nov 2007	Dec 2007	Jan 2008	Feb 2008	Mar 2008
Total BF done (n=12)	4983	5242	5990	5993	5656	5376	6081	5489	4522	5023	5239	4908
Total Positive BF (n=12)	917	1203	1154	1382	1438	1317	1368	1114	1013	1132	997	946
Ave % positive BF (n=12)	18.4	22.9	19.3	23.1	25.4	24.5	22.5	20.3	22.4	22.5	19.0	19.3

There was only one facility in the dataset that was using RDTs. during the period.	There were no data on RDTs done

Section 3: Pharmacy

Table 3A: Percentage of Facilities with Availability of Essential Medicines for Malaria

	Artesu nate / Amodi aquine	Artesu nate Oral	Amodi aquine Oral	SP	Co- Artem	Quinin e injecta ble	Quinin e Oral	Chloro quine Oral	Alaxin	Artem os	Dara- prim
					Type of Fa	acility					
Teaching and Regional Hospitals	83.3	75.0	70.0	83.3	36.4	66.7	83.3	33.3	45.5	41.7.	18.2
District and Other Hospitals	71.5	85.6	71.5	71.5	33.3	60.1	42.9	33.3	42.9	14.4	33.3
Health Centres	100.0	27.3	41.7	75.0	10.1	66.6	54.5	20.0	22.2	0.0	0.0
Clinics	66.6	33.4	46.7	66.6	28.6	33.0	20.0	20.0	46.7	14.3	21.5
Maternity Homes	60.1	83.2	79.9	100.0	33.5	0.0	60.1	20.1	79.9	50.0	0.0
CHPS Zones	100.0	0.0	59.9	40.1	0.0	50.0	20.1	0.0	20.1	0.0	0.0
				0	perating A	uthority					
Gov't	96.2	29.8	46.6	64.4	4.4	47.5	33.5	12.0	32.2	4.3	4.2
Mission	71.0	71.0	71.0	85.4	49.4	39.1	56.3	42.4	33.7	0.0	19.7
Private	33.1	59.3	66.3	79.4	50.5	44.1	33.1	22.8	69.6	44.4	34.2
Totals	79.0 (n=55)	43.0 (n=56)	54.5 (n=54)	71.0 (n=57)	21.2 (n=49)	45.5 (n=42)	37.2 (n=55)	19.6 (n=53)	41.4 (n=52)	44.4 (n=52)	13.3 (n=52)

Other Malarial Medicines Available at facilities included; Artemeter 300mg suspension, Arthemeter Injection, Camosunate, Chloroquine Injection (1), Halfan Syrup (1)

Table 3B: Percentage of Facilities with a stock out of Essential Medicines for Malaria since Jan '08

	Artesu nate / Amodi aquine	Artesu nate Oral	Amodi aquine Oral	SP	Co- Artem	Quinin e injecta ble	Quinin e Oral	Chloro quine Oral	Alaxin	Arte- Mos	Dara- prim
					Type of F	acility					
Teaching and Regional Hospitals	50.0	11.1	40.0	20.0	80.0	28.6	30.0	66.7	50.0	66.7	80.0
District and Other Hospitals	0.0	0.0	0.0	0.0	33.5	0.0	39.9	0.0	0.0	100.0	0.0
Health Centres	20.0	74.9	39.9	33.4	50.0	0.0	0.0	100.0	74.9	100.0	100.0
Clinics	20.0	42.9	33.4	0.0	66.7	42.9	66.7	79.9	44.4	71.5	62.5
Maternity Homes	0.0	0.0	0.0	0.0	0.0	100.0	33.5	100.0	0.0	33.5	100.0
CHPS Zones	0.0	100.0	25.1	50.0	0.0	0.0	100.0	100.0	66.7	100.0	100.0
				0	perating A	uthority					
Gov't	13.3	57.7	28.6	26.2	85.7	18.7	46.4	99.3	57.1	99.4	85.9
Mission	0.0	0.0	25.5	0.0	34.3	50.9	34.3	0.0	0.0	100.0	50.9
Private	25.8	25.8	14.8	0.0	39.9	24.1	39.7	65.9	13.0	42.9	39.3
T	40.0	00.0	0.1.0	40.0	00.5	00.0	40.4	77.0	00.4	74.7	04.5
Totals	13.6 (n=40)	39.3 (n=32)	24.3 (n=35)	16.2 (n=41)	60.5 (n=25)	23.8 (n=24)	43.1 (n=31)	77.6 (n=15)	39.4 (n=29)	71.7 (n=29)	64.5 (n=29)

Section 4: Patient Observations and Re-examinations

Table 4A: Percentage Patients y Reasons given for visiting facility

Reason	Under 5	Over 5
Fever	74.5	28.0
Diarrhoea	23.2	4.8
Vomiting everything	23.2	1.0
Cough, difficult breathing	23.5	9.6
Ear problem	2.1	2.6
Skin problem	18.5	5.8
Total Number	n=131	n=322

Table 4B: Percentage of providers who asked about key symptoms for those presenting with fever - by age of patient

Symptom	Under 5	Over 5
Unable to eat or drink	44.0	37.9
Vomiting everything	45.3	14.1
Convulsions	7.6	5.5
Fever	41.2	63.3
Headache	3.9	56.8
Joint Pain	2.6	20.3
Difficult breathing/cough	58.2	35.5
Chills	4.3	14.1
Little or no urination in past 24 hours	0.9	4.6
Dark Urine	0.0	3.1
Abnormal bleeding	0.0	2.4
Difficult or painful urination	1.0	1.6
Diarrhea or abdominal pain	44.0	47.0
Bloody stools	29.1	4.1
Ear Pain	10.0	1.0
Total Numbers	n=98	n=87

Table 4C: Percentage of Providers who asked all danger signs in children under 5

Symptom	% who asked	% who are trained in IMCI
Unable to eat or drink	38.0	42.5
Vomiting everything	39.2	43.8
Convulsions	6.3	4.0
Fever	49.0	46.8
Asked all 4 above	2.0	4.0
Total number	n=134	n=116

Table 4D: Percentage of providers who performed the following examinations

Examination	Under 5	Over 5
Take temperature or refer to where it is written on card	91.4	81.8
Weigh patient or refer to written weight	84.1	64.4
Take blood pressure	0.9	63.3
Take pulse (for 60 seconds)	7.4	10.9
Check for anemia (palmor pallor)	47.9	27.2
Check for visible severe wasting	11.7	-
Count respiratory rate (for 60 seconds)	7.2	12.9
Check for dehydration (skin pinch) ¹	10.6	-
Check for lethargy	3.2	-
Check immunization status	9.6	-
Totals	n=134	n=322

Note: 10.6% of providers checked for dehydration in all patients observed and re-examined. However, 31.1% of providers checked for dehydration for patients under 5 whose reason for visiting the facility was diarrhea.

Table 4E: Percentage of providers who counseled patients

	All Patients Observed and Re-examined			diagnosed of vider at facility
Counselling Topic	Under 5	Over 5	Under 5	Over 5
Provide general information on feeding or breastfeeding	24.7		21.8	
Give extra fluids to patient during this illness	11.5		10.6	
Continue feeding / eating during this sickness	13.4		11.9	
Proper management of fever (sponging and paracetamol)	7.5		8.7	
Tell patient/caregiver what illness patient has	10.0	21.5	10.3	18.5
Talk to caregiver/patient about ITN use	12.1	5.0	14.2	8.5
Describe signs or symptoms in patient that they should return to the facility	23.2	20.9	26.0	18.6
If medicines are prescribed – tell patient how to take the medicine	27.9	24.8	27.0	24.8
If medicines are prescribed give the patient the first dose.	20.0		22.6	
Totala	n_124	N=322	n_00	n_162
Totals	n=134	N=322	n=88	n=163

Table 4F: Percentage of Providers using lab tests

	All pat	ients	Patients diagnosed with malaria		
	Under 5	Over 5	Under 5	Over 5	
Provider ordered lab tests	14.1	22.6	13.2	28.3	
Provider did not order tests	85.9	77.4	86.8	71.7	
Totals	n=131	N=319	n=86	n=162	

Table 4G: Percentage of Patients diagnosed with malaria for whom lab tests were ordered by type of test and the results

	Diagnosed with malaria				
	Under 5	Over 5			
Provider ordered lab tests	13.2 (n=88)	28.3 (n=163)			
Hematocrit <15%	0.0 (n=13)	0.0 (n=26)			
HB <5g/dl	0.0 (n=13)	3.9 (n=26)			
Parasitimia + or more	0.0 (n=13)	13.0 (n=26)			

Table 4H: Percentages of Patient Knowledge about How Malaria is Transmitted

	Mosquitoes / Mosquito bites	Insects (no mosquitoes mentioned)	Through the sun	Juju/Sorcery	Eating Certain Foods	Don't Know	Number of patients
Diagnosed with malaria	75.2	0.0	6.8	28.4	9.9	15.4	249
Not Diagnosed with malaria	68.7	1.2	8.4	36.7	13.3	14.1	197
Totals	72.4	0.5	7.5	32.0	11.4	14.9	446

Table 4I: Percentages of Patient Knowledge on how Malaria is Prevented

	Sleep under a net	Sleep under impregnated net	Give a drug	Use insecti- cide	Use coils or repellants	Use trad. Means	Prevent Mosq bites	Avoid fatigue	Don't Know	Number of patients
Diagnosed with malaria	52.1	14.5	2.9	10.1	9.8	1.8	4.4	0.3	12.1	248
Not Diagnosed with malaria	48.5	20.2	3.1	10.5	8.4	0.0	2.7	0.0	12.1	197
Totals	50.5	17.0	3.0	10.3	9.2	1.0	3.7	0.2	12.1	445

Table 4J: ITN Ownership, Use and Health Worker Interaction on ITNs

	Health worker talked to patient about ITN today	Patient has an ITN in the home	Number of Patients (n)	Patient slept under ITN last night	Number of Patients (n)
Diagnosed with malaria	11.2	59.3	224	53.4	158
Not Diagnosed with malaria	1.5	50.2	187	38.8	109
Under 5	10.1	77.3	138	58.8	105
Over 5	5.5	47.5	320	44.3	165
Totals	6.6	54.5	458	48.9	270

Section 5: Provider Knowledge on Treatment and Management of Malaria

Response and data in this section of the appendices are based on answers provided during the provider interview and may differ from the actual practice of provider reported in earlier sections of the report. This section reflects what providers say they do and may not reflect actual practice.

Table 5A: Percentage of Drugs providers say they commonly prescribe for treating uncomplicated malaria

	Type of drugs									Number of
	Artesunate / Amodiaquine	Artesunate Only	Amodia- quine Only	SP Only	Chloro-quine Only	Co-Artinate	Co-Artem	Alaxin	Quinine	Providers
		•		Ту	pe of Fac	ility	•	•	•	
Teaching and Regional Hospitals	80.0	15.0	5.0	0.0	0.0	10.0	45.0	10.0	25.0	20
District and Other Hospitals	71.5	14.4	28.5	0.0	0.0	14.4	0.0	28.5	0.0	7
Health Centres	100.0	0.0	15.4	15.4	0.0	7.8	0.0	15.4	23.1	13
Clinics	73.3	20.0	33.4	14.3	0.0	7.8	0.0	46.7	0.0	15
Maternity Homes	83.2	28.5	50.0	33.3	0.0	0.0	16.8	66.7	33.3	6
CHPS Zones	100.0	0.0	0.0	14.3	0.0	0.0	0.0	14.3	14.3	7
					rating Aut					
Govt	99.8	3.3	19.2	12.8	0.0	6.6	0.5	23.3	12.9	10
Mission	80.1	17.1	33.9	0.0	0.0	0.0	0.0	29.0	0.0	50
Private	44.8	33.4	35.9	27.1	0.0	9.4	8.9	63.2	17.7	11
		ı			e of Prov		1	1	1	
Doctor	50.7	13.0	0.2	12.4	0.0	12.8	1.5	37.0	0.6	23
MA	89.7	20.4	30.3	21.9	0.0	21.9	0.4	27.5	22.1	13
Nurse	100.0	20.1	20.1	0.0	0.0	0.0	0.0	0.0	0.0	6
Midwife	86.7	6.5	33.2	19.8	0.0	0.0	7.0	57.0	19.6	15
CHN	100.0	0.0	0.0	0.0	0.0	0.0	0.0	20.0	20.0	5
Other	100.0	16.3	59.6	20.2	0.0	0.0	0.0	19.5	0.0	6
Dune de la constitución				Pro	vider Trai	ning	1	ı	ı	
Provider had refresher training in malaria in last 3 years	91.3	11.2	20.0	17.0	0.0	5.8	0.3	25.7	14.1	68
Totals	85.3	12.3	24.8	14.7	0.0	6.6	2.5	33.2	12.7	68

Table 5B: Percentage of Providers Who Use Artesunate / Amodiaquine for Treatment of Malaria

	Always prescribe A/A	Often / Sometimes prescribe A/A	Rarely prescribe A/A	Never prescribe A/A	Number of providers
			Facility		
Teaching and Regional Hospitals	45.0	40.0	5.0	10.0	20
District and Other Hospitals	71.3	14.4	14.4	0.0	7
Health Centres	84.6	15.4	0.0	0.0	13
Clinics	58.7	17.7	17.7		17
Maternity Homes	57.1	42.9	0.0	0.0	7
CHPS Zones	85.7	14.3	0.0	0.0	7
		Operating	g Authority		
Govt	87.2	12.7	0.1	0.0	51
Mission	56.9	14.6	28.5	0.0	8
Private	32.7	41.6	17.1	8.6	12
			Provider		
Doctor	37.3	37.3	12.2	12.8	23
MA	82.8	0.3	16.9	0.0	15
Nurse	100.0	0.0	0.0	0.0	6
Midwife	80.1	13.1	6.9	0.0	15
CHN	80.0	20.0	0.0	0.0	5
Other	33.9	66.1	0.0	0.0	6
		Provider	Training		_
Had refresher training in malaria in last 3 years	74.7	16.8	8.4	0.0	71
,		1	ı	_1	
Totals	70.3	19.7	8.0	2.1	71

Table 5C: Percentage of Providers – Reasons Why prescribers do not prescribe the nationally mandated first line treatment for uncomplicated malaria – Artesunate/Amodiaquine.

	Fear of	Experience of	Other	Number of
	Adverse	patient		Providers
	Reaction			(n)
		Type of Facility		
Teaching and				3
Regional	33.3	0.0	66.7	
Hospitals				
District and	100.0	0.0	0.0	2
Other Hospitals	100.0	0.0	0.0	
Health Centres	-	-	-	-
Clinics	66.7	100.0	33.3	3
Maternity				-
Homes	-	-	-	
CHPS Zones	-	-	-	-
Totals	79.3	60.2	20.7	8

Table 5D: Percentage of Providers by practice of ordering laboratory tests

	Providers routinely order lab tests	Number of Providers	Orders Blood Film for Malaria Parasites	Orders Hemoglobin	Number of Providers
		Type of	Facility		
Teaching and Regional Hospitals	70.0	20	100.0	85.7	14
District and Other Hospitals	28.5	7	100.0	100.0	2
Health Centres	15.4	13	100.0	100.0	2
Clinics	43.8	16	100.0	100.0	7
Maternity Homes	50.0	6	100.0	100.0	3
CHPS Zones	0.0	7	-	-	-
			Authority		
Government	22.5	51	100.0	99.8	20
Mission	40.0	6	100.0	99.1	3
Private	41.6	12	100.0	100.0	5
		Provider	Training		
Refresher Training in Malaria in last 3 years	26.6	69	100.0	99.8	28
Totals	28.9	69	100.0	99.7	28

Table 5E: Percentage of Provider by practice of diagnosing severe malaria – Signs or Conditions Used for Diagnosis - Unprompted

	Hyper-parasitemia	Altered Consciousness	Convulsions	Hypo-glycemia	Difficult Breath-ing or pulmonary Oedema	Reduced Urine Output	Severe Anemia	Circulatory Collapse or Shock	Electrolyte Imba- lance	Jaundice	Hyper-pyrexia	Prostration	Number of providers
		I	I	I	Тур	e of Faci	lity	I.			ı		
Teaching and Regional Hospitals	35.0	30.0	70.0	5.0	10.5	40.0	80.0	15.8	10.5	45.0	65.0	75.0	20
District and Other Hospitals	57.1	71.5	85.6	0.0	0.0	14.4	71.5	14.4	0.0	14.4	100.0	71.5	7
Health Centres	7.8	30.7	61.5	0.0	15.4	23.1	15.4	15.4	7.8	0.0	92.2	76.9	13
Clinics	6.7	40.0	37.5	25.0	6.7	23.5	41.2	0.0	0.0	33.4	88.2	43.8	15
Maternity Homes	0.0	28.5	28.5	0.0	0.0	0.0	57.1	14.4	0.0	0.0	42.9	71.5	7
CHPS Zones	0.0	28.6	71.4	0.0	14.3	14.5	14.3	0.0	0.0	0.0	71.4	71.4	7
					Operating	Authority	/			•			
Governmen t	9.9	38.6	58.0	3.3	12.9	19.0	34.9	6.5	3.2	7.0	90.4	65.6	50
Mission	32.8	66.4	85.1	14.6	0.0	28.5	56.9	16.5	0.3	33.3	99.8	66.1	7
Private	8.6	25.0	25.3	17.1	0.0	8.6	33.0	8.2	0.0	17.1	50.8	58.0	12
	1		1		Type of		1	1	1			•	
Doctor	49.3	61.3	74.9	12.6	0.4	25.7	26.1	12.4	0.2	35.7	86.3	49.7	23
MA	10.1	59.9	72.2	18.5	0.0	25.3	50.4	19.7	0.2	0.2	91.3	64.0	14
Nurse	20.1	39.6	39.9	6.9	20.0	20.3	39.9	0.4	0.0	0.0	100.0	79.9	15
Midwife CHN	0.0	26.6 20.0	39.6 80.0	6.9 0.0	20.0	20.0	33.6 20.0	6.5 0.0	6.6 0.0	20.5	80.4 60.0	79.8 60.0	5 6
Other	0.0	17.2	17.2	0.0	0.0	0.0	49.6	0.0	0.0	17.2	67.5	33.0	6
Outer	0.0	11.4	11.2	0.0	Provider		43.0	0.0	0.0	17.2	07.5	33.0	U
Had malaria refresher training last 3 years	11.0	38.8	52.6	8.5	8.4	19.6	30.8	11.0	0.0	14.2	89.0	63.6	69
Totals	12.4	38.7	54.0	8.2	8.2	17.9	37.5	8.1	2.1	12.7	82.4	63.9	69

^{*}No providers mentioned using acidosis as a way to diagnose severe malaria.
** 6.2% of provider mentioned spontaneous bleeding as a way to diagnose severe malaria.

Table 5G: Percentage of Prescribers that Knew the Correct Treatment Drugs for Severe malaria - Unprompted

	Quinine Injection	Quinine Tablets	Artesunate Alone	Amodia- quine Alone	SP	Co- artinate	Co- Artem	Chloroquine	Number of Providers
			Тур	e of Facility					
Teaching and Regional Hospitals	75.0	20.0	5.0	0.0	0.0	5.0	5.0	0.0	20
District and Other Hospitals	28.5	16.8	0.0	0.0	0.0	0.0	0.0	14.4	7
Health Centres	23.1	7.8	15.4	15.4	0.0	0.0	0.0	0.0	13
Clinics	25.0	6.7	6.7	0.0	0.0	0.0	6.7	6.7	15
Maternity Homes	28.5	0.0	14.4	14.4	0.0	14.4	14.4	0.0	7
CHPS Zones	14.3	0.0	0.0	0.0	0.0	0.0	0.0	0.0	7
			Opera	ting Author	ity				
Govt	25.6	6.5	6.4	6.3	0.0	0.0	0.1	3.3	50
Mission	49.4	20.4	0.0	0.0	0.0	0.0	0.0	0.0	7
Private	8.2	0.0	16.7	8.2	0.0	0.0	16.7	8.3	12
				of Provide		1			
Doctor	14.3	0.5	12.6	0.0	0.0	0.0	12.6	12.0	23
MA	45.7	20.2	9.9	0.0	0.0	0.0	0.0	0.0	14
Nurse	39.9	20.5	0.0	0.0	0.0	0.0	0.0	0.0	15
Midwife	20.2	0.0	6.6	13.2	0.0	0.0	0.0	6.9	5
CHN	20.0	0.0	0.0 16.4	0.0 16.3	0.0	0.0	0.0	0.0	6 6
Other	0.0	0.0		ider Trainin	0.0	0.0	16.3	0.0	6
Provider had refresher training in last 3 years	27.8	8.6	2.7	2.7	0.0	0.0	0.0	2.8	69
Totals	24.3	6.4	8.1	5.9	0.0	0.0	4.1	4.1	69

Section 6: Ante-Natal Care and Malaria

Table 6A: Percentage of Facilities with ANC services and their organization

	% with daily ANC Services*	% offering FANC	% offering Adequate Lab Services**	% with functioning weighing scale	% with functioning BP apparatus	% with ANC Register	% with current ANC Record Entry
		•	Type o	f Facility		•	•
Teaching and Regional Hospitals	66.6	25.0	91.7	91.7	100.0	90.9	80.0
District and Other Hospitals	33.4	0.0	57.1	83.2	83.2	100.0	100.01
Health Centres	72.7	90.8	0.0	90.8	90.8	100.0	89.9
Clinics	62.8	85.8	14.3	100.0	100.0	100.0	85.5
Maternity Homes	66.0	67.0	16.6	100.0	100.0	83.4	100.0
CHPS Zones	100.0	100.0	0.0	100.0	100.0	100.0	74.9
			Operating	g Authority			
Govt	68.2	74.7	10.2	92.0	92.2	99.9	86.8
Mission	50.0	50.9	32.6	100.0	100.0	100.0	100.0
Private	66.0	60.1	20.1	100.0	100.0	83.4	100.0
Totala	65.7	69.5	15.2	94.1	94.2	97.1	90.4
Totals	65.7 (n=47)	69.5 (n=45)	15.2 (n=59)	94.1 (n=46)	94.2 (n=46)	97.1 (n=45)	90.4 (n=42)

^{* 5} days or more

** Blood mps, Hb and urine analysis

Record entry within the past 7 days

Table 6B: Percentage of Facilities by Type of SP in Stock

Facility type	Fansidar	Malafan	Other	Number of facilities
Teaching and Regional Hospitals	30.0	20.0	50.0	10
District and Other Hospitals	0.0	16.8	83.2	6
Health Centres	0.0	30.0	70.0	10
Clinics	43.0	14.4	42.5	7
Maternity Homes	33.0	33.0	33.9	6
CHPS Zones	25.0	50.0	25.0	4
Totals	18.4	27.2	54.5	43

Table 6C: Percentage of Facilities by Action When there is a Stock out

Facility type	Write a prescription	Nothing	Other	Number of facilities
Teaching and Regional Hospitals	20.0	40.0	40.0	5
District and Other Hospitals	0.0	50.0	50.0	2
Health Centres	20.1	20.1	59.9	5
Clinics	33.5	0.0	66.5	3
Maternity Homes	32.9	0.0	67.1	3
CHPS Zones	0.0	33.3	67.7	3
Totals	18.7	18.9	62.4	21

Of 59 facilities, 87.9% provide SP for free. Of the 5 facilities charging for SP, 1 is a government facility, 1 is a mission facility and 3 are private facilities.

Section 7: Instruments and Protocols for the Malaria Health Facility Survey

Instrument	# completed per facility	Person completing instrument	Methodology Inclusion / Exclusion Criteria
A-1. Malaria Health Facility Audit	1	Auditor	All facilities completed the facility audit in full, except where skip patterns indicated a section may be skipped
A-2 Patient Observation Form	ALL in selected consulting room	Observer	Random selection of consulting room in the OPD (or in a teaching hospital – the polyclinic). Observed ALL patient consultations from the first consultation to 2:00 PM
A-3 Provider Interview Tool	1 -2	Observer	Only the provider observed was interviewed
A-4 Re- examination of patient and Client Exit Interview	ALL from selected consulting room	Re-Examiner	All patients observed above.
A-5 Assessment of Cases with Severe Malaria	5	Auditor	Listed all patients currently admitted for severe malaria. Where less than five – observed all. Where more than five – randomly selected five patients.
A-6 Laboratory Quality Control Tool	ALL from the selected consulting room where the provider ordered lab tests in patients	Auditor sets up the process and monitors it throughout the day.	All patients who had lab work ordered by provider –second slide taken.
B-1 ANC Facility Audit	1	Midwife	All facilities completed audit section. For record review, determined how many patients had been seen between Oct 2007-Mar 2008. Divided the total number of consultations by 10. Then used this sampling interval to select records starting in Oct 2007 to Mar 2007 until 10 records were reviewed.
B-2 ANC Observation and Client Exit Interview Tool		Midwife	Random selection of consulting room done. Selection of ANC Clients: Sample of convenience – next woman was seen. Observed all consultations of the selected provider between 9:00 AM and 2:00 PM on the day of the assessment. Selection of providers: Randomly selected one provider from all the providers giving service on the day of the assessment.

Section 8: National Malaria Health Facility Survey Tools

A-1: Malaria – Health Facility Audit

FACILITY IDENTIFICATION				
Name of Region:`	FACILITY CODE			
Name of District:				
Name of the facility				
TELEPHONE:				
Interviewer Name:	FACILITY TYPE			
Date: Type of Health Facility: (1 = Teaching Hospital 2=Regional hospital; 3 = District Hospital; 4=Other Hospital 5=Polyclinic 6=Health Center 7= Maternity Home 8=Clinic 9=CHPS 10=RCH Unit 11=Other	OPERATING AUTHORITY			
Operating Authority: 1= Government; 2=Mission/Religious 3 = Private for profit 4=Quasi-Government 5 = Other)				

1. General Information				
NO.	QUESTIONS	CODE CLASSIFICATION	GO TO	
100	FOR OUTPATIENT SERVICES: FIND THE MANAGER OR MOST SENIOR HEALTH WORKER WHO IS PRESENT AT THE FACILITY. INTRODUCE YOURSELF AND READ THE FOLLOWING:			
	I am representing the Ghana Health Service and the survey. We are conducting research that will help us situation for malaria services. This information will be these services. The survey will assess case manage services for malaria in pregnancy. The tools include exit interview and collection of service statistics.	to understand the health service e used to design programs to improve ment of malaria and also assess		
	This part of the survey will ask questions about the services offered at this facility, the equipment that is available and staffing levels. It may take up to half a day to complete this part of the survey. All information from this survey is confidential and participation in answering questions for this survey is voluntary. You can refuse to answer any question or all the questions. There is no risk to you or your facility for participation. The results will be used to plan programming that may benefit your facility.			
	We are asking for your help to ensure that the information collected is accurate. If there are sections where someone else is the most appropriate person to provide information, we would appreciate your introducing us to that person.			
	Do you have any questions for me? Can you please sign the line below and put the date?			
	100			
	Signature of the interviewee indicates that consent w			
101	Is there a trained health provider present at the facility at all times (24 hours/day)	YES, TRAINED PROVIDER ALWAYS PRESENT1 NO, NOT ALWAYS PRESENT2 NO, NO TRAINED PROVIDER .3		
102	Is there a trained health provider available on call at all times after normal working hours? IF YES, ASK TO SEE A CURRENT DUTY ROSTER	YES, DUTY SCHEDULE SEEN 1 YES, NO DUTY SCHEDULE 2 NO 3		

Now I have some questions about the staff. We want to know the number of staff who are routinely assigned to provide services for sick people, including for malaria. This may include staff who provide both inpatient and outpatient services but NOT staff who function purely administratively.

COUNT STAFF IN ONLY ONE CATEGORY. DO NOT INCLUDE STAFF ON STUDY LEAVE.

QUALIFICATION	TOTAL NUMBER	
A) Medical Doctors	MEDICAL DOCTOR	
B) Medical Assistants	MEDICAL ASST	
C) Public Health Nurses	PH NURSE	
D) Midwives	MIDWIFE	
E) SRN	PROF. NURSE	
F) Community Health Nurses	CHN	
G) Enrolled Nurses	EN	
H) Ward Assistants / Ward Maid	WA	
I) Pharmacists	PHARMACIST	
J) Dispensing Technologists	DISPENSING TECH 1	
K) Dispensing Technicians	DISPENSING TECH 2	
L. Dispensing Assistants	DISPENSING Assist	
M) Lab Technicians		
N) Laboratory Assistants		
Now I have some questions about the staff that malaria. We want to know the number of staff w control and malaria monitoring		
O) Technical Officer/Field Technicians (Disease Control)	DCO	

P). Biostatistician / Records Officer	BIOSTATS
Are there any other staff who routinely provide n counted above?	nalaria-related services who were not
Q) Others: SPECIFY	OTHER

NO.	QUESTIONS	CODE CLASSIFICATION	GO TO
104	Does this facility have formal meetings to review	YES1	
	management or administrative issues?	NO2	→ 107
		DON'T KNOW8	→ 107
105	How often are formal meetings held to discuss	WEEKLY1	
	general management/administrative issues?	MONTHLY2	
		QUARTERLY3	
		SEMI-ANNUALLY4	
		OTHER5	
106	Is an official record of meetings maintained?	YES, DOCUMENT SEEN1	
		YES, DOCUMENT NOT	
	IF YES, ASK TO SEE SOME	SEEN2	
	DOCUMENTATION (MINUTES/NOTES) FROM	NO DOCUMENTATION	
	THE MOST RECENT MEETING	MAINTAINED3	
107	Does this facility have any system for	SUGGESTION BOX1	
	determining client opinion about the health	CLIENT SURVEY FORM2	
	facility or services?	CLIENT INTERVIEW3	
		COMMUNITY DURBAR4	
	IF YES, CIRCLE ALL METHODS FOR	PUBLIC FORUM5	
	ELICITING CLIENT OPINIONS THAT ARE	OTHER6	
	USED	(SPECIFY)	
	IF NO CLIENT FEEDBACK CIRCLE '7'	NO CLIENT FEEDBACK7 DON'T KNOW8	
	IF NO CLIENT FEEDBACK CIRCLE /	DON KNOW8	
108	Does this facility have a Quality Assurance	YES1	
	Team?	NO2	→ 110
		DON'T KNOW8	→ 110
109	Does the team have a Quality Assurance Action	YES, PLAN SEEN1	
	Plan?	YES, NO PLAN SEEN2	
		NO3	
	IF YES, ASK TO SEE THE PLAN OR		
	EVIDENCE OF RECENT ACTIVITY		

NO.	QUESTIONS	CODE CLASSIFICATION	GO TO
110	When was the last time a supervisor from	WITHIN PRIOR 6 MONTHS	
	OUTSIDE this facility came for a supervisory	1	→ 111
	visit?	MORE THAN 6 MONTHS	3 440
		AGO2	→ 112
		NEVER SUPERVISED FROM	→ 112
444	Within the meet Consents alide a consention from	OUTSIDE FACILITY3	
111	Within the past 6 months did a supervisor from outside the facility on a visit do any of the	YES NO) DK
	following activities?	CHECKED	
	A) Check some registers or service related	REGISTERS1 2	8
	books?	TREGIOTERO	o l
	booke.	DISCUSSED	
	B) Discuss problems?	PROBLEMS 1 2	8
	, ,		
	C) Discuss policy/administrative issues?	DISCUSSED POLICY1 2	8
	D) Discuss technical protocols, practices, or	DISCUSSED	
	service delivery technical issues?	TECHNICAL ISSUES 1 2	8
	T) Hold on official staff monting?	HELD STAFF	
	E) Hold an official staff meeting?	MEETING1 2	8
		WEETING	0
	F) Observe individual staff providing services?	OBSERVE SERVICE	
	Try esserve marriadar etan previanig cerviceer	PROVISION1 2	8
	G) Do anything else?	OTHER1 2	8
112	Is there a printed referral form which is sent with	YES, FORM SEEN1	
	referrals from this facility?	YES, FORM NOT SEEN2	
	IF YES, ASK TO SEE THE FORM.	NO FORM,	
		USE LETTERHEAD3	
	(IF THE FACILITY IS THE REFERRAL	NO FORM4	
	FACILITY, THEN CIRCLE "5" FOR REFERRAL	REFERRAL FACILITY5	
	FACILITY).	DON'T KNOW8	
113	If a decision is made to refer a patient to a	An ambulance1	
	higher level facility, what means of	A taxi	
	transportation is available	Private Vehicles3	
	CIDCLE ALL THAT ADDLY	Other4_	
	CIRCLE ALL THAT APPLY	specify	

114	Does this facility have electricity?	YES1	
		NO2	→ 116
115	Is the electricity always available during times when the facility is providing services or is it sometimes interrupted?	ALWAYS AVAILABLE1 SOMETIMES INTERUPPTED2 OFTEN INTERUPPTED3	

FOR EACH OF THE FOLLOWING ITEMS, CHECK WHETHER THE ITEM IS PRESENT AT THE FACILITY. IF YOU ARE NOT ABLE TO OBSERVE THE ITEM YOURSELF, ASK IF THE ITEM IS AVAILABLE. WHERE APPLICABLE, ASK IF THE ITEM IS IN WORKING ORDER OR NOT

	ITEM	a) Is the item present?		b) Is the item in working order?			
		Observed / RA	Not Available	Not Determined	Yes	No	Not Deter- mined
116	Generator	1	2 → 118	8 → 118	1	2	8
117	Fuel for generator	1	2	8			

NO.	QUESTIONS	CODE CLASSIFICATION	GO TO
118	What is the most commonly used source of	PIPED1	
	water for the facility at this time of the year?	PROTECTED WELL/BOREHOLE2	
		UNPROTECTED WELL3	
		RIVER/LAKE /POND4	
		OTHER6	
		(SPECIFY)	
		NO WATER SOURCE9	
119	Is this water source available on-site?	YES, ON-SITE 1	
		NO2	
120	How is water made available for use in	PIPED1	
	examination/consultation areas in the facility	BUCKET/BASIN2	
	today?	VERONICA BUCKET3	
		NO WATER PROVIDED	
	CIRCLE ALL THAT APPLY	IN SERVICE	
		DELIVERY AREAS4	
121	Is there a waiting area for clients, where they	YES1	
	are protected from sun and rain?	NO2	
122	Is there a toilet (latrine) in functioning	YES1	
	condition, which is available for clients' use?	NO2]
123	Is there cell phone or landline reception at this	No reception and no landline1	
	facility? If so on what networks?	MTN2	
		One Touch3	
	CIRCLE ALL THAT APPLY	Kasapa4	
		Tigo5	
		Landline available6	
		Radio phones (Motorola)7	
		Other8	

	Is the new combination therapy Artesunate/Amodiaquine being used to treat sick children at this facility?	YES1 NO2 → 128 DON'T KNOW8 → 128
	IF YES, What is the dosage for a child that weighs 20 kgs? (Answer Artesunate 4mg/kg (80mg) and Amodiaquine 10mg/kg (200 mg) for 3 days given in two divided doses.	CORRECT1 NOT CORRECT2
126	How many staff have been trained in the use of the new malaria drug policy in this facility?	Don't Know = 98
127	How many providers treat sick children at this facility?	Don't Know = 98
128	How many providers at this facility have been trained in IMCI?	Don't Know = 98

2. OPD Services

NO.	QUESTIONS	CODE CLASSIFICATION	GO TO
200	Does this facility offer OPD services or curative services for	Yes1	→ 200b
	malaria?	No2	→ 300
	FIND THE MANAGER OR MOST SENIOR HEALTH WORKER RESERVICES WHO IS PRESENT AT THE FACILITY. INTRODUCE THE FOLLOWING:		
	I am representing the Ghana Health Service and the Quality Health Partners Project for this survey. We are conducting research that will help us to understand the health service situation for malaria health services. This information will be used to design programs to improve these services. This part of the survey will ask questions about the services offered at this facility, the equipment that is available and staffing levels. It will take between 20-30 minutes to complete. All information from this survey is confidential and participation in answering questions for this survey is voluntary. You can refuse to answer any question or all the questions.		
	We are asking for your help to ensure that the information collected sections where someone else is the most appropriate person to proappreciate your introducing us to that person.		
	Do you have any questions for me?		
	Will you participate in this survey? If Yes, ask the person to sign the	e line below.	
200a			
	SIGNATURE OF INTERVIEWEE INDICATES AGREEMENT TO PA	ARTICIPATE	

NO.	QUESTIONS		CODING CLASSIF	GO TO						
201	adults. For each of the following s									
	HEALTH SERVICES	# Days per week service provided a facility		Service not offered						
	a). Consultation / curative care services for sick children under 5?	DAYS	11	95						
	b). Consultation / curative services for the Sick patient 5 years and above?	DAYS	11	95	IF BOTH 95 → 300					

ASK TO SEE WHERE CONSULTATION SERVICES FOR PATIENTS ARE OFFERED, AND EXPLAIN THAT YOU WANT TO ASK ABOUT AND SEE SOME OF THE MATERIALS USED FOR PROVIDING CURATIVE SERVICES. ASK TO SEE EACH OF THE FOLLOWING ITEMS. THE ITEMS SHOULD BE IN THE ROOM, OR IN AN IMMEDIATELY ADJACENT ROOM TO BE COUNTED AS PRESENT.

202	ITEMS REQUIRED FOR CONSULTATION SERVICES FOR SICK CHILDREN	(a) Is item present?				(b) Is item in working order?		
	Supplies	Observed	Not Available	ND	Yes	No	ND	
	a) Hand-washing Items (soap)	1	2	8				
	b) Single use towels	1	2	8				
	c) Water for hand-washing	1	2	8				
	d) Alcohol Hand Wash							
	e) Infant Scale	1	2	8	1	2	8	
	f) Child Scale	1	2	8	1	2	8	
	g) Adult Weighing Scale	1	2	8	1	2	8	
	h) Thermometer	1	2	8	1	2	8	
	i) Timer/Watch with second hand	1	2	8	1	2	8	
	j) Jar/Pitcher for ORS	1	2	8				
	k) Cup and spoon	1	2	8				

203	REFERENCES/ PROTOCOLS/ TEAC MATERIALS	HING	Ob	served	Not Available		ND
	a) Standard Treatment Guidelines 200)4		1	2		8
	b) Anti-Malaria Drug Policy for Ghana			1	2		8
	c) IMCI CHART BOOKLET			1	2		8
	d) Malaria Counseling Chart			1	2		8
	e) Malaria treatment chart			1	2		8
	f) Other visual aids for teaching careta	ker		1	2		8
204	Is there a ROUTINE system where sicl are measured/weighed/assessed <u>prior</u> consultation for the illness?	to the	NO DON'	T KNOW .			→205 →206 →206
205	IF YES, ASK TO SEE WHERE SICK (CONSULTATION AND INDICATE WH ARE ROUTINELY CARRIED OUT TH	HICH OF				;	
	PART OF ROUTINE SERVICES	Observed	b	Not Done Routinely	Don't Kno)W	
	a) Take Weight	1		2	8		
	b) Record weight	1		2	8		
	c) Take temperature	1		2	8		
	d) Assess immunization status			2	8		
	e) Sponge febrile children	1	2		8		
	f). Give paracetamol to febrile children	1		2	8	8	
	g) Other (SPECIFY)	1		2	8		

206	Do you have a stock of child health cards? If YES, ask to see one.	YES, CARD	RVED CARD 1 NOT SEEN 2 UAL CARDS 3		
	Is the new combination therapy Artesunate/Amodiaquine being used to treat sick children at this facility?	YES NO DON'T KNO	2	→ 209 → 209	
	IF YES, What is the dosage for a child that weighs 20 kgs? (Answer Artesunate 4mg/kg (80mg) and Amodiaquine 10mg/kg (200 mg) for 3 days given in two divided doses.	ORRECT NOT CORRI			
209	How many staff have been trained in the use of the r drug policy in this facility?	new malaria	Don't Know = 98		
210	How many providers treat sick children at this facility	?	Don't Know = 98		
211	How many providers at this facility have been trained	I in IMCI?	Don't Know = 98		

	3. Integrated Disease Surveil	lance and Response										
NO.	QUESTIONS	CODING CLASSIFICATION	GO TO									
300	Does this facility detect, track and report on	YES1										
	priority diseases (e.g. cholera, meningitis,	NO2	→ 400									
	guinea worm, malaria, etc) or use the IDSR											
	method?	INVOLVED IN THE DELIVERY OF	IDOD									
	FIND THE MOST SENIOR HEALTH WORKER SERVICES. IF DIFFERENT FROM INDIVIDUA											
	SECTIONS COMPLETE QUESTION BELOW.		1003									
	CONTINUE WITH 302.											
	FOR NEW RESPONDENTS READ AND COMPLETE QUESTION 301.											
	I am representing the Ghana Health Service and the Quality Health Partner for this survey. We are conducting research that will help us to better understand how malaria services are provided and											
	improve the quality of care. This information will be u											
	services. This part of the survey will ask questions a											
	Response. It will take about 10-20 minutes to complete	ete.										
	All information from this survey is confidential and pa	articination in answering questions for thi	e curvov									
	is voluntary. You can refuse to answer any question											
	to ensure that the information collected is accurate. I	f there are sections where someone else	e is the									
	most appropriate person to provide information, we v	would appreciate your introducing us to t	hat									
	person.											
	Do you have any questions for me?											
	Will you participate in this survey? If Yes, ask them	to sign the line below.										
301												
	CIONATURE OF INTERVIEWEE INDICATED DARK	COLDANIT ACREEMENT TO DARTICID	ATE AND									
	SIGNATURE OF INTERVIEWEE INDICATES PART THAT THE TIME IS CONVENIENT	ICIPANT AGREEMENT TO PARTICIP	ATE AND									
	THE TIME IS SOLVENIEN											
302	How many staff have been trained in the use											
	of surveillance forms?											
		Don't Know = 98										
303	How many staff have been trained in the use											
	of standard case definitions?	Dan't Know OC										
304	How many staff have been trained in the use	Don't Know = 98										
JU4	of surveillance data (i.e. training in surveillance											
	data analysis) to show trends of disease											
	morbidity and mortality?	Don't Know = 98										

305	Does this facility have protocols on the following: IF YES, ASK TO SEE A COPY.	Observ ed / Reporte d Availabl e	Not Available	Not Deter- mined
	a) National Technical Guidelines on IDSR	1	2	8
	b) Standard Case Definitions	1	2	8
	c) COMDAB	1	2	8
306	Data Analysis: Does the facility have monthly trend analysis of graphs for key diseases? (FOR LAST TWO QUARTERS OF 2007)	Observ ed	Not Available	Not Deter- mined
	a). Malaria cases in children < 5	1	2	8
	b). All Malaria cases reported at the facility	1	2	8
	c). Malaria deaths	1	2	8
	d). Other Malaria related graph (specify)	1	2	8
307	Does the facility have displayed demographic data (population, age/sex distribution) of the catchment area?	1	2	8
308	Do you have an estimate of the size of the catc population that is served by this facility? That is target population or total population lining in the served by this facility?	CATCHMENT PC		
			NO ESTIMATE DK 9999	

4. Laboratory

NO.	QUESTIONS	CODING CLASSIFICATION	GO TO
400	Does this facility offer laboratory services?	YES 1	→ 401
		NO 2	→ 500

FIND THE MOST SENIOR HEALTH WORKER INVOLVED IN THE DELIVERY OF LAB SERVICES. IF DIFFERENT FROM INDIVIDUAL(S) RESPONDING TO THE PREVIOUS SECTIONS COMPLETE QUESTION BELOW. IF THE PERSON IS THE SAME, CONTINUE WITH 402.

FOR NEW RESPONDENTS READ AND COMPLETE QUESTION 401.

I am representing the Ghana Health Service and the Quality Health Partners Project for this survey. We are conducting research that will help us to better understand how malaria services are provided. This information will be used to design programs to improve these services. This part of the survey will ask questions about laboratory service. It will take about 10-20 minutes to complete.

All information from this survey is confidential and participation in answering questions for this survey is voluntary. You can refuse to answer any question or all the questions. We are asking for your help to ensure that the information collected is accurate. If there are sections where someone else is the most appropriate person to provide information, we would appreciate your introducing us to that person.

Do you have any questions for me?

Will you participate in this survey? If Yes, ask the person to sign the line below.

401

SIGNATURE OF INTERVIEWEE INDICATES PARTICIPANT AGREEMENT TO PARTICIPATE AND THAT THE TIME IS CONVENIENT

	ITEMS REQUIRED FOR LABORATORY	(a) Is item pre	esent?	(b) If item/s available, in working order?									
	EXAMINATION	Observed / RA	Not Available	Not Deter- mined	Yes	No	ND						
402	Electric Binocular Microscope	1	2	8	1	2	8						
403	Hematocrit Centrifuge	1	2	8	1	2	8						
404	Slides and coverslips	1	2	8	1	2	8						
405	RAPID DIAGNOSTIC TEST (RDT) for Malaria	1	2	8									
406	Giemsa stain	1	2	8									

TES1	S FOR ANEMIA							
		(a) Is item pre	esent?		(b) If item/s available, in working order?			
		Observed / RA	Not Available	Not Deter- mined	Yes	No	ND	
407	Hemoglobinometer/ Colorimeter	1	2	8	1	2	8	
408	Drabkin's solution	1	2	8				
409	Capillary tubes (for hematocrit)	1	2	8	-			
410	Stericon strips or Tallquist test for measuring hemoglobin (w/ valid expiry date)	1	2	8				
	A	VAILABILITY	OF LAB TES	TS				
411	Are the following tests availa	ble at this facil	ity on a routine	basis?				
		Observed / RA	Not Available	Not Deter- mined				
	a) complete blood cell count	1	2	8				
	b) urine analysis	1	2	8	-			
	c) spot glucose level	1	2	8				
	d) electrolytes	1	2	8				
	e) liver function test	1	2	8				
	f) G6PD	1	2	8				

LAB	SERVICES STATISTICS	
Does the Lab maintain log book, register other record of tests done?	YES	→413 →500

413	APR	MAY	JUN	JUL	AUG	SEP	OCT	NOV	DEC	JAN	FEB	MAR
	2007	2007	2007	2007	2007	2007	2007	2007	2007	2008	2008	2008
a. Number of												
malaria blood												
films done												
b. Number of												
RDT for												
malaria done												
(if applicable)												
c. Number of												
positive												
malaria blood												
films												
d. Number of												
positive												
RDTs for												
malaria												

5. ESSENTIAL MEDICATIONS AND SUPPLIES

Does this facility have a pharmacy /dispensary or does it dispense drugs to patients?

Yes-----2

→500a **→600**

FIND THE PHARMACIST, DISPENSING TECHNICIAN OR OTHER HEALTH WORKER RESPONSIBLE FOR DRUG MANAGEMENT AT THE FACILITY. IF IT IS THE SAME PERSON YOU HAVE BEEN TALKING TO GO TO QUESTION 501

FOR NEW RESPONDENTS READ AND COMPLETE QUESTION 500.

I am representing the Ghana Health Service and the Quality Health Partners Project for this survey. We are conducting research that will help us to better understand how malaria treatment is provided. This information will be used to design programs to improve these services. This part of the survey will ask questions about the services offered at this facility, the availability of drugs and the equipment that is available. It will take about 30 minutes -1 hour to complete. All information from this survey is confidential and participation in answering questions for this survey is voluntary. You can refuse to answer any question or all the questions. We are asking for your help to ensure that the information collected is accurate.

Do you have any questions for me?

Will you participate in this survey? If Yes, ask the person to sign the line below.

500a

SIGNATURE OF INTERVIEWEE INDICATES PARTICIPANT AGREEMENT TO PARTICIPATE AND THAT THE TIME IS CONVENIENT.

	CHECK TO SEE IF EACH OF THE THE PHARMACY TODAY.	THEN CHEC								
		MEDICATIO	N SINCE 1	JAN 2008						
	MEDICATION	Stock-out	No-Stock	Not						
		Determined		Out	Determined					
	Anti-Malarial MEDICINES									
501	Artesunate/Amodiaquine oral	1	2	8	1	2	8			
	(co-packaged)									
502	Artesunate oral	1	2	8	1	2	8			
503	Amodiaquine oral	1	2	8	1	2	8			
504	Artemether-Lumefantrine	1	2	8	1	2	8			
	(Co-Artem or Lonart)									
505	Quinine oral	1	2	8	1	2	8			

	MEDICATION	Observed	Not Available	Not Determined	Stock-out	No-Stock Out	Not Determined
	Anti-Malarial MEDICINES						
506	Quinine Inj.						
507	Chloroquine oral	1	2	8	1	2	8
508	Sulphadoxine/pyrimethamine	1	2	8	1	2	8
	(FANSIDAR, Malafan)						
509	Alaxin tablets/Suspensions	1	2	8	1	2	8
510	Artemos tablets/suspension	1	2	8	1	2	8
511	Daraprim oral	1	2	8	1	2	8
512	Other Anti-Malarial (SPECIFY)	1	2	8			
	a						
	b						
	C						

	A).CHECK TO SEE IF EACH OF THESE THE PHARMACY TODAY.	B). THEN CHECK TO SEE IF THEY HAVE HAD A STOCKOUT OF THIS							
	THE PHARMACT TODAT.		TION SINCE 1						
	MEDICATION	Observe	Not	Not	Stock-	No-Stock	Not		
		d	Available	Determine	out	Out	Determined		
	Other MEDICINES			d					
513	Amoxicillin caps /Suspension	1	2	8	1	2	8		
514	Co-trimoxazole tabs/Susp.	1	2	8	1	2	8		
515	Ciprofloxacin tablets	1	2	8	1	2	8		
516	Iron preparations	1	2	8	1	2	8		
517	Multivitamin tabs/caps/syrups	1	2	8	1	2	8		
518	Paracetamol tablets	1	2	8	1	2	8		
519	Paracetamol Syrup /Suppositories	1	2	8	1	2	8		
520	Aspirin/aspirin containing drugs	1	2	8	1	2	8		
521	Diazepam tablets/Supp.	1	2	8	1	2	8		
522	ORS	1	2	8	1	2	8		
	Parenteral Preparations								
523	Diazepam Inj	1	2	8	1	2	8		
524	Penicillin/Ampicillin Inj	1	2	8	1	2	8		
525	Gentamycin Inj	1	2	8	1	2	8		
526	Chloramphenecol Inj	1	2	8	1	2	8		
527	Furosemide inj	1	2	8	1	2	8		
528	Dexamethasone Inj	1	2	8	1	2	8		
529	Normal Saline	1	2	8	1	2	8		
530	Dextrose and Saline	1	2	8	1	2	8		
531	Dextrose and Saline Infant Prep	1	2	8	1	2	8		
532	Dextrose	1	2	8	1	2	8		
533	Ringers Lactate	1	2	8	1	2	8		
534	Glucose 50%	1	2	8	1	2	8		
535	Glucose 20%	1	2	8	1	2	8		
536	Glucose 5%	1	2	8	1	2	8		
537	Furosemide	1	2	8	1	2	8		

505		41144414
535	During the past 6 months, have you	ALWAYS 1
	always, sometimes or almost never	SOMETIMES2
	received the amount of each medication	ALMOST NEVER3
	that you order (or that you are supposed	
	to routinely receive)?	
536	Were medicines organized according to	YES1
	expiry date "first in first out" on the	NO2
	shelves? (VERIFY WHEN	DON'T KNOW8
	COMPLETING ABOVE)	
537	Did the stock records indicate that FIFO is	YES1
	practiced? (VERIFY WHEN	NO2
	COMPLETING ABOVE)	DON'T KNOW8
538	Are the medicines off the floor and	YES1
	protected from water / dampness?	NO2
	ľ	DON'T KNOW8
539	Are medicines protected from the sun?	YES1
	· ·	NO2
		DON'T KNOW8
540	How will you describe the temperature of	Cool1
	your medicine storage room? At what	Warm2
	temperature(s) do you usually keep your	Very hot 3
	medicines?	
541	How do you decide how much of each	Order to bring stock to a fixed level1
	malaria medication to order? Do you:	Order the same quantity each time regardless
	a.aa	of how many of each medication remain in
	CIRCLE ALL THAT APPLY.	stock2
		Order different amounts based on calculation of
		prior utilization and expected future activity?3
		Order depending on what you think is needed
		without a specific method for calculating
		amounts4
		Need and amount determined elsewhere5
		Other – specify6
		Don't know8
543	How do you decide when to order	Place order whenever stock level falls to a pre-
	medications? Do you:	determined level1
	,	Have a fixed time when you are supposed to
		submit orders for medications2
	CIRCLE ALL THAT APPLY	The facility can place an order wherever there
		is believed to be a need
		It is necessary to wait until the official time for
		ordering4
		Other – specify5
		Don't know8

58

MONTHLY OUTPATIENT MORBIDITY (SERVICE STATISTICS)

600	۸DD	NANY	HIN	шш	ALIC	CED	OCT	NOV	DEC	IANI	EED	111
000	APR	MAY	JUN	JUL	AUG	SEP	OCT	NOV	DEC	JAN	FEB	MA
	2007	2007	2007	2007	2007	2007	2007	2007	2007	2008	2008	200
a. Malaria												
cases												
children<1yr												
b. Malaria												
cases												
children 1-												
4yrs												
c. Malaria												
cases												
OVER 5yrs												
d. Malaria												
cases												
Pregnant												
women												
e. Total												
Malaria												
cases												
f. Total												
Anemia												
cases												
g. Total												
OPD												
Attendance												

Lab Assessment Tool 59

	FACILITIES THAT HAVE INPATIENT CAPACITY ONLY											
601	APR	MAY	JUN	JUL	AUG	SEP	OCT	NOV	DEC	JAN	FEB	MAF
	2007	2007	2007	2007	2007	2007	2007	2007	2007	2008	2008	2008
A.												
Average												
Bed												
Occupancy												
B.												
Total												
Admissions												
(malaria)												
C .Total												
Admissions												
(all)												

Observation	Number
Observation	Number

A-2. Patient Observation Form

FACILITY IDENTIFICATION					
Name of Region:`	FACILITY CODE				
Name of District:					
Name of the facility	PROV TYPE				
·	PROV SEX				
Name of Observer	FACILTY TYPE				
Date:					
Type of Health worker Observed: Doctor =1, MA = 2 Nurse =3, Midwife =4, Other = 5	OPER AUTH				
Provider Sex Male =1 Female =2 Type of Health Facility: 1 = Teaching Hospital 2=Regional hospital; 3 = District Hospital; 4=Other Hospital 5=Polyclinic 6=Health Center 7= Maternity Home 8=Clinic 9=CHPS 10=RCH Unit 11=Other	PROVIDER NUMBER (For first provider observed in a consultation room write 1. If the provider in THAT consultation room changes write 2,3 etc) DO NOT CHANGE CONSULTATION ROOMS.				
Operating Authority: 1= Government;	OCHOCE IATHON ROOMS.				

NO.	QUESTIONS	CODING CLASSIFICATION		
O100a	How old is the patient?	Under 5 1→ O100c	Over 5 2→0100b	
O100b.	WRITE AGE IN FULL YEARS			
O100C	IF PATIENT IS <5 YEARS WRITE AGE IN MONTHS MONTHS			
O100D	Gender of the Patient	Male	Female	
		1	2	

O101	What reason/s does the Caregiver / Patient provide for coming to the facility	YES	NO
	A) Fever/ Hot body?	1	2
	B) Diarrhoea?	1	2
	C) Vomiting everything?	1	2
	D) Cough/difficult breathing	1	2
	E) Ear problem?	1	2
	F) Skin Problem	1	2
	G) OTHERS (SPECIFY)	1	2
	Assessment		
O102	Does the provider ask the patient/caregiver about the following symptoms since the beginning of the illness A) Unable to eat or drink?	YES 1	NO 2
	B) Vomiting everything	1	2
	C) Convulsions?	1	2
	,	1	2
	,	<u> </u>	
	,	1	2
	F) Joint Pain?	1	2
	G) Difficult breathing / Cough?	1	2
	H) Chills?	1	2
	I) Little or no urination in the past 24 hours?	1	2
	J) Dark Urine?	1	2
	K) Abnormal Bleeding?	1	2
	L) Difficult or painful urination?	1	2
	M) Diarrhea / Abdominal Pain?	1	2
	N) Bloody Stools?	1	2
	O) Ear Pain	1	2
	P) Other	1	2
O103	Does the provider or other clinical staff member perform any of the following examinations?	YES	NO
	A). Take the patient's temperature or refer to their card where it is written?	1	2
	B). Weigh the patient, refer to patients weight on the card or check weight for age	1	2
	C). Take blood pressure	1	2
	D) Take pulse (<u>>60</u> seconds, with timer or watch with second hand)	1	2
	E) . Check for anaemia (palmar pallor)	1	2
	F) Check for visible severe wasting	1	2 2
	G). Count respiratory rate (<u>>60</u> seconds, with watch or timer)	1	2
	H) .Check for dehydration (Skin pinch)	1	2
	I) Check for lethargy if child not awake	1	2
	J). Check immunization status	1	2
	K). Other	1	2
•			

	Communication / Counseling		
O104	Does the Provider provide any of the following advice when counseling the patient / caregiver?	YES	NO
	A) Provide general information about feeding or breastfeeding the patient?	1	2
	B) Give extra fluids to the patient during this sickness?	1	2
	C) Continue feeding the child / eating as a patient during this sickness and after?	1	2
	If the patient has fever does the provide discuss management of fever by sponging and giving paracetamol?	1	2
	D) Tell the caregiver/ patient what illness (es) the child/patient has?	1	2
	E). Talk to the caregiver/patient about ITN use?	1	2
	F) Describe signs or symptoms in the patient for which the caregiver /patient should return to the facility?	1	2
	G). If medicines are prescribed, does the provider describe how to take each medicine completely?	1	2
	H). If medicines are prescribed, does the provider ensure that the patient takes the first dose at the facility (administers himself or tells pharmacy to administer the medicine.)	1	2

	Laboratory	
O105	Does the provider order lab tests?	YES 1→106 NO 2→110
O106	What was the Hematocrit result?	< 15%1
		≥15%2
		Result not available the same day3
		Patient did not return with results4
		Test not done5
O107	What was the Hemoglobin result?	<5 g/dl1
		≥ 5 g/dl2
		Result not available the same day3
		Patient did not return with results4
		Test not done5
O108	What was the parasetimia level?	+1
		++2
		+++3
		++++4
		Result not available the same day5
		Patient did not return with results6
		Test not done7
O109	What was the Parasite species	p. Falciparum1
		p. Malariae2
		p. Ovale3
		Mixed4
		None specified5
		Result not available the same day6
		Patient did not return with results7
		Test not done8

AFTER THE CONSULTATION – REVIEW PATIENT'S CARD WITH THE PROVIDER and or ask him/her about his/her diagnosis on this case.

NO.	QUESTIONS	CODING CLASSIFICATION	GO TO
O110	How does the Provider classify the patient's illness with regard to fever? IF THE CHILD DID NOT HAVE FEVER ACCORDING TO THE PROVIDER'S CLASSIFICATION, CIRCLE CODE "4".	Uncomplicated Malaria Very Severe Febrile Disease incl. Seve Fever – other cause No Fever	re Malaria2 3
	ONLY FOR CHILDREN UNDER 5		
O111	What other diagnoses did the provider make? CIRCLE ALL APPLICABLE IF THERE IS NO ADDITIONAL DIAGNOSES CIRCLE CODE 22.	Severe Pneumonia	

O112	What oral / suppository treatments were	Pre-	Not	Prescription
	prescribed?	scribed	Prescribed	
	A) Artesunate + Amodiaquine antimalarial tablets? .	1	2	Formulation: a. Amount each time: b. number of times per day: c. total days:
	B) Artesunate/ Amodiaquine antimalarial syrup / suppository?	1	2	Formulation: a. Amount each time: b. number of times per day: c. total days:

C) Other antimalarial tablet/syrup? specify:	1	2	Formulation: a. Amount each time: b. number of times per day: c. total days:
D) Co-trimoxazole tablets/syrup?	1	2	Formulation: a. Amount each time: b. number of times per day: c. total days:
E) Amoxicilin tablets/syrup?	1	2	Formulation: a. Amount each time: b. number of times per day: c. total days:
F) Other antibiotic tablets/syrup? specify:	1	2	Formulation: a. Amount each time: b. number of times per day: c. total days:
G) Paracetamol?	1	2	
H) Antidiarrheal/antimotility drug?	1	2	
I) Vitamin A?	1	2	
J) Multi-vitamins?	1	2	
K) Other vitamins?	1	2	
L) Mebendazole?	1	2	
M) Iron tablets/syrup?	1	2	
N) Other Tablets/syrup	1	2	
O) ORS?	1	2	

0113	What injections were prescribed?	Prescribed	Not Prescribed	
	A) Antimalarial : specify	1	2	Formulation: a. Amount each time: b. number of times per day: c total days:
	B) Antibiotic: specify	1	2	Formulation: a. Amount each time: b. number of times per day: total days:

O114	Did the provider do any of the following?	Admit the patient1	
		Refer the patient2	

AT THE END OF THE CONSULTATION GIVE THE PATIENT A CARD WITH THEIR OBSERVATION NUMBER ON IT (FROM PAGE ONE).

ASK THEM TO GO TO THE LAB AND PHARMACY AS APPROPRIATE AND DIRECT THEM TO SEE YOUR COLLEAGUE BEFORE LEAVING THE FACILITY.

Comments:

A-3 Provider Interview Tool

FACILITY IDENTIFICATION				
Name of Region:`	FACILITY CODE			
Name of District:				
Name of the facility	PROV SEX			
Name of Interviewer(s)	FACILTY TYPE			
	OPER AUTH			
Date:				
Sex Male =1 Female =2	PROVIDER NUMBER (For first provider observed in a			
Type of Health Facility: (1 = Teaching Hospital 2=Regional hospital; 3 = District Hospital; 4=Other Hospital 5=Polyclinic 6=Health Center 7= Maternity Home 8=Clinic 9=CHPS 10=RCH	consultation room write 1. If the provider in THAT consultation room changes write 2,3 etc)			
Unit 11=Other)	DO NOT CHANGE CONSULTATION			
Operating Authority:	ROOMS.			
Operating Authority:				
1= Government; 2=Mission/Religious 3 = Private for profit 4=Quasi-Government 5 = Other)				

NOW I WOULD LIKE TO ASK YOU SOME QUESTIONS ABOUT YOU AND THE SERVICES YOU PROVIDE HERE IN RELATION TO MALARIA

	2. PROVIDER TRAINING AND EXPERIENCE					
NO.	QUESTIONS	CODING CLASSIFICATION				
O201	In what year did you start working in this facility?	YEAR				
O202	What is your current technical/professional qualification?	MEDICAL DOCTOR 1 MEDICAL ASST 2 NURSE 3 MIDWIFE 4 COMMUNITY HEALTH NURSE 5 OTHER 6				
O203	What year did you graduate with this qualification?	YEAR.				

	2. UNCOMPLICAT	ED MA	LARIA		
O204	What drugs do you usually prescribe for treating uncomplicated malaria: DO NOT READ, DO PROMPT – ASK ANY OTHERS?		Mentioned	Not Men	tioned
	a. Artesunate + Amodiaquine		1	2	
	b. Artesunate only		1	2	
	c. Amodiaquine only		1	2	
	d SP only		1	2	
	d. chloroquine only		1	2	
	e. Artesunate +S/P (Co-arinate)		1	2	
	f. Co-artem or Lonart		1	2	
	g. Alaxin		1	2	
	h. Quinine		1	2	
	i. Other? (specify)		1	2	
O205	How often do you prescribe Artesunate- Amodiaquine for malaria treatment?	OFTE RARE	AYS EN / SOMETIMES ELY ER	2 3	→ 207 → 207
O206	Why don't you prescribe the combination? CIRCLE ALL APPLICABLE	perso exper lack o exper	of adverse reaction onal experience rience of a patient of confidence rience of a colleague (specify	2 3 4	
O207	Do you routinely order laboratory tests in patients (> 5 years) with fever/ suspected malaria?	0.101	Yes	7.0	No
	If Yes: What laboratory tests do you routinely ask for?		1 → 207A		2 → 208
	A. blood film for malaria parasites		1		2
	B. haemoglobin		1		2
	C. Other (Specify)		1		2

		3. SEVERE/COMPLICAT	ED MAL	ARIA			
O208	severe	signs or conditions do you look for in order to diagno Malaria: DO NOT READ, DO NOT PROMPT – DTHERS?		Mentione	ed	Not N	Mentioned
		Hyperparasitaemia (High parasite Load)		1			2
	b)	Altered consciousness (confusion or drowsiness) o	r	1			2
	c)	Convulsions.		1			2
	d)	Hypoglycemia.		1			2
	e)	Acidosis		1			2
	f)	Difficulty in breathing or pulmonary oedema		1			2
	g)	Reduced urine output		1			2
	h)	Severe Anaemia or severe pallor (HB<5mg/dl)		1			2
	i)	Circulatory collapse or shock (low volume pulse/colextremities)	ld	1			2
	j)	Electrolyte imbalance		1			2
	k) Jaundice (yellowing of eyes)			1		2	
	I) Hyperpyrexia (axillary temperature ≥ 39.5 Oc)			1			2
	m) Spontaneous Bleeding (Disseminated Intravascular Coagulation)			1		2	
	n) Prostration i.e. generalized weakness (inability to stand or walk).		nd or	1		2	
		Other (SPECIFY)		1			2
O209	malaria facility? If they DO NO	nti-malarial drug/s do you use in treating severe a OR before referring a patient to a better equipped mention a drug – ask – what do you give? DT READ, DO NOT PROMPT – ASK – HING ELSE?**	Mentio Dose	ned Correct		ioned ng Dose	Not Mentioned
	a)			1		2	3
	b)			1		2	3
	c)	Artesunate alone a loading dose 4mg/kg (given in two doses) or 50mg suppository for child and 200 mg suppository for adult) and then 2mg/kg oral in two doses for 4 days		1		2	3
	d)			1		2	3
	e)	S/P 500mg/25mg		1		2	3
	f)	Artesunate –SP (e.g. Co-arinate) 1 tab each daily * 3days		1		2	3
	g)	Co-artem or Lonart (>35kg, 4tabs twice daily * 3days)		1		2	3
	h)			1		2	3
	i)	Other (SPECIFY)		1		2	3

^{**}NOTE –the provider may not know the exact dose – you may ask them to refer to their resources and tell you.

O210	What other supportive treatment do you sometimes find necessary to provide in cases of severe Malaria? DO NOT READ, DO NOT PROMPT	Mentioned	Not Mentioned
	a) Blood Transfusion for Severe Anaemia	1	2
	b) IV Infusions to correct fluid and electrolyte imbalance	1	2
	c) Anti-convulsants for convulsions	1	2
	d) Anti-pyretics for hyperpyrexia	1	2
	e) IV Dextrose for hypoglycaemia	1	2
	f) I refer all cases of severe or complicated malaria to a hospital or other facility for treatment	1 → 213	2
	g) Other (SPECIFY)	1	2
O211	What clinical monitoring do you do in cases of severe Malaria? DO NOT READ, DO NOT PROMPT		
	a) Level of consciousness (using Glasgow or Blantyre coma scale)	1	2
	b) Fluid intake/output and IV infusion/drip rate	1	2
	c) Blood Pressure, Body Temperature, Pulse.	1	2
	d) Breathing or Respiration rate	1	2
	e) Blood parasitemia	1	2
	f) Hemoglobin/hematocrit	1	2
	g) Other (SPECIFY)	1	2

	FOR FACILITIES THAT HAVE LABORAT	TORIES ONLY	
O212	What laboratory tests do you request in cases which you highly suspect to be complicated or severe malaria? DO NOT READ, DO NOT PROMPT – ASK ANYTHING ELSE?	Mentioned	Not Mentioned
	 a) Order Thick and/ thin blood film microscopy for malaria parasites. 	1	2
	b) Order/do Rapid Diagnostic Testing for MPs	1	2
	c) Order of check Haemoglobin or Hematocrit	1	2
	d) Order Blood Urea/Creatinine investigations	1	2
	e) Order Blood electrolytes investigations	1	2
	f) Order Blood Glucose assessment	1	2
	g) Do Lumber Puncture to exclude other causes of coma	1	2
	h) Other (SPECIFY)	1	2
	 i) I refer all cases of severe or complicated malaria to a hospital or other facility for treatment 	1	2

Now I would like to ask a few questions about training and standards and guidelines at this facility.

NO.	Question	Coding Classification			
O213	Do you have a copy of the following: (ask to see a copy),.	Yes seen or reported	No	Don't know	
		to have			
	a. National Malaria Drug Policy?	1	2	8	
	b. Standard treatment guidelines?	1	2	8	
	c. Malaria Treatment Guideline	1	2	8	
	d. Malaria Counseling Card?	1	2	8	
	e. IMCI guidelines (Chart booklet)?	1	2	8	
O214	Have you been trained or received refresher/ follow-up in last 3 years in Malaria	YES		1 2	
O215	Have you been trained or received refresher/ follow-up in last 3 years in IMCI	YES1 NO2			
O216	Do you feel you need additional training in order to be able to manage malaria better?			1 2	

Thank you very much for allowing us to observe your consultations today and thank you also for completing this interview. We hope that the results will lead to more malaria programming that will involve you.

INTERVIEWER COMMENTS

П		
П		
П		
П		
П		
П		
П		

A-4. Exit Interview and Re-Examination of Patient Tool

FACILITY IDENTIFICATION					
Name of Region:`	FACILITY CODE				
Name of District:					
Name of the facility	FACILTY TYPE				
Name of Interviewer	OPER AUTH				
Date:	SEX of PATIENT				
Type of Health Facility: (1 = Teaching Hospital 2=Regional hospital; 3 = District Hospital; 4=Other Hospital 5=Polyclinic 6=Health Center 7= Maternity Home 8=Clinic 9=CHPS 10=Other)	AGE of PATIENT YEARS AGE of CHILD MONTHS				
Operating Authority: 1= Government; 2=Mission/Religious 3 = Private for profit 4=Quasi-Government 5 = Other)					
SEX OF THE PATIENT 1 = MALE 2 = FEMALE					

I would like to ask you a few questions about your care at this facility today.

<u>N</u> O.	QUESTIONS	COI	DING CLASSIF	<u>ICATION</u>	<u>GO</u> <u>TO</u>
R100	Did the health worker give or prescribe any oral medicines (anti- malarials, antibiotics or ORS) for <patient> at the health facility today?</patient>	YES		→R108 →R108	
R101	If yes compare the caretaker's/Patient's medication with the oral medicines – (anti-malarials, antibiotics or ORS only)	samples for identification of the			
	Copy the information from the caretaker's medication or prescription – INCLUDE ORS: Does patient have the medicine(1) or do they have prescription to buy it outside the facility?(2)			r do they tion to buy it	
	A		1	2	
	B		1	2	
	C		1	2	
	D		1	2	
	E		1	2	
	F		1	2	
If yes,	ask the caretaker the following (record what you hear):				
R102	MEDICATION A				
	How much will you give <patient> each time: of medication <a></patient>		qua	antity	
		8 = DO1	N'T KNOW		

R102a	How many times will you give medication <a> to <patient> each</patient>		
	day?	times	
		8 = DON'T KNOW	
R102b	How many days will you give medication <a> to <patient> ?</patient>		
		days	
		8 = DON'T KNOW	
R103	MEDICATION B	NO MEDICATION B	→ R108
	How much will you give <patient> each time: of medication </patient>	quantity	
		8 = DON'T KNOW	
R103a	How many times will you give medication to <patient> each</patient>		
	day?	times	
		8 = DON'T KNOW	
R103b	How many days will you give medication to <patient>?</patient>		
		days	
		8 = DON'T KNOW	
R104	MEDICATION C	NO MEDICATION C	→ R108
	How much will you give <patient> each time: of medication <c></c></patient>		
		8 = DON'T KNOW	
R104a	How many times will you give medication <c> to <patient> each</patient></c>		
	day?	times	
		8 = DON'T KNOW	
R104b	How many days will you give medication <c> to <patient> ?</patient></c>		
		days	
		8 = DON'T KNOW	
R105	MEDICATION D	NO MEDICATION D1	→ R108
	How much will you give <patient> each time: of medication <d></d></patient>		
		8 = DON'T KNOW	
R105a	How many times will you give medication <d> to <patient> each day?</patient></d>		
	uay:	times	
DAOEL		8 = DON'T KNOW	
R105b	How many days will you give medication <d> to <patient> ?</patient></d>	davia	
		days	
D400	MEDICATION E	8 = DON'T KNOW	> D400
R106	MEDICATION E	NO MEDICATION E1	→ R108
	How much will you give <patient> each time: of medication <e></e></patient>	8 = DON'T KNOW	
R106a	How many times will you give medication at a spatiant, each	8 = DON 1 KNOW	
Kiloba	How many times will you give medication <e> to <patient> each day?</patient></e>	times	
		8 = DON'T KNOW	
R106b	How many days will you give medication <e> to <patient> ?</patient></e>	8 - DON I KNOW	
1000	Thow many days will you give medication <e> to <patients !<="" td=""><td>days</td><td></td></patients></e>	days	
		8 = DON'T KNOW	
R107	MEDICATION F	NO MEDICATION D1	→ R108
107	How much will you give <patient> each time: of medication <f></f></patient>	NO WEDIO/(TION D	21(100
	The magnitum you give spanishes each affect of medication si	8 = DON'T KNOW	
R107a	How many times will you give medication <f> to <patient> each</patient></f>	0 - DOINT INIOW	
I TOTA	day?	times	
	, in the second	8 = DON'T KNOW	
R107b	How many days will you give medication <f> to <patient> ?</patient></f>		
1075	Then many days will you give modication of 2 to spation 2:	days	
		8 = DON'T KNOW	
<u> </u>			

R108	Did the health worker give you a specific day when to come back to this facility?	1 = YES 2 = NO	DAYS	
	If the answer is Yes, but patient can't remember write 98 days	8 = DON'T KNOW		
R109	FOR CHILDREN <5 ONLY: Sometimes children's condition may worsen and they should be taken immediately to a health facility: What conditions in your patient would cause you to take your patient to a health facility right away? DO NOT PROMPT	Mentioned	Not Mentioned	
	a. patient not able to drink or breastfeed	1	2	
	b. patient becomes sicker	1	2	
	c. patient develops a fever	1	2	
	d. patient has fast breathing	1	2	
	e. patient has difficulty breathing / pneumonia	1	2	
	f. patient has blood in the stool	1	2	
	g. patient is drinking poorly	1	2	
	h. other (specify)	1	2	

	Inform	ation about the disease	
R110	When did your patient get sick? (CHECK ONLY ONE)	Today	
R111	Was the patient taken to some other place for care/treatment before being brought here today? CHECK ONLY ONE	YES	→R113 →R113
R112	IF YES: Where was the patient taken? CIRCLE ALL THAT APPLY	THIS FACILITY	
R113	Did you give any drugs to the patient since s/he got ill?	YES	→R116 →R116
R114	What drugs did you give the patient? CIRCLE ALL THAT APPLY	Anti-malarial treatment	
R115	Where did you get the drugs from? CIRCLE ALL THAT APPLY	PHARMACY/CHEMICAL SHOP .1 HEALTH FACILITY .2 NEIGHBOUR/RELATIVE .3 MARKET .4 OTHER (SPECIFY) .5 Don't Know .8	

	Knowledge of Malaria by the Caretaker/Patient				
NOW I A	M GOING TO ASK YOU SOME QUESTIONS				
R116	How does someone get malaria? DO NOT READ OPTIONS. CIRCLE ALL THAT THEY ANSWER.	MOSQUITOES OR MOSQUITO BITE			
		(SPECIFY) DON'T KNOW8 SLEEP UNDER NET1			
R117	How can someone protect himself or herself against malaria? DO NOT READ OPTIONS. CIRCLE ALL THAT THEY ANSWER.	SLEEP UNDER AN IMPREGNATED NET 2 GIVE A DRUG 3 USE INSECTICIDE 4 USE COILS OR REPELLANTS 5 USE OTHER TRADITIONAL MEANS 6 PREVENT MOSQUITO BITES 7 FATIGUE 8 OTHER 9 (SPECIFY)			
R118	Do you have an insecticide treated net in your home?	DON'T KNOW 10 YES 1 NO 2 DON'T KNOW 8	→R120 →R120		
R119	IF YES: Did you /your patient sleep under a net last night?	YES			
R120	Did any health worker talk to you about ITN today?	YES			
R121	What will you do if you / the patient does not get better? (CIRCLE ALL THAT APPLIES)	RETURN TO THIS FACILITY			
R122	How many days will you wait to see if the patient is not getting better before using the options mentioned above?	DAYS (IF DON'T KNOW = 98)			
R123	Could you tell me at least one thing that can be improved in the services delivered at this health facility?				

RE-EXAMINATION FOR PATIENTS >5 YEARS NOW I WOULD LIKE TO RE-EXAMINE YOU.

R200	Take patient's axillary temperature and record temperature		·_	c
R201	Assess for the following signs and symptoms since the beginning of the illness	YES	NO	
	A) Fever/ Hot body or history of fever?	1	2	
	B) Headache?	1	2	
	C) Joint Pain?	1	2	Count the number of
	D) Nausea / Vomiting?	1	2	non-fever
	E) Chills? / Rigors?	1	2	symptoms
	F) Poor Appetite?	1	2	and record the result
	G) Fatigue? General Malaise	1	2	uno roodii
	H) Diarrhea?	1	2	
	I) Dizziness?	1	2	
	J) Anemia (check for pale palms)	1	2	
R202	IF THE PATIENT HAS FEVER (HISTORY OR TEMPERATURE ≥37.5 OR AT LEAST THREE NON-FEVER SYMPTOMS THEN THE PATIENT HAS SUSPECTED MALARIA. GO TO QUESTION R204.	SUSPECTI MALARIA		→ R204
	IF THE PATIENT HAS NO FEVER <u>AND</u> LESS THAN 3 NON FEVER RELATED SYMPTOMS, THEN THE PATIENT DOES NOT HAVE SUSPECTED MALARIA. THANK THE CARETAKER/PATIENT AND END THE INTERVIEW.	NO SUSPE MALARIA		→END
	Assessment for Severe Malaria			
R204	Measure 60 second respiratory rate while patient is calm.	broo	tha nar mi	nuto
	(Normal breathing rate for an adult at rest is 8-16 breathes a minute)	breaths per minute		
R205	Measure patients pulse while patient is calm in a minute.	beats per minute		er minute
R206	Ask the patients caregiver about the following symptoms since the beginning of the illness	YES	NO	Not APP
	A) Little or no urine in the past 24 hours	1	2	8
	B) Dark urine (Cola colored urine)	1	2	8
	C) Convulsions	1	2	8
	D) Prostration or abnormal behavior	1	2	8
	E) Abnormal Bleeding	1	2	8
R207	Assess the patient for the following signs			
	A. Respiratory Distress	1	2	8
	B. Unable to take liquids, profuse, repeated vomiting	1	2	8
	C. Prostration or abnormal behavior	1	2	8
	D. Weak Pulse or > 110 beats/min	1	2	8
	E. Cyanosis	1	2	8
	E la cracia	4		
	F. Jaundice	1	2	8
	G. Severe Palmor Pallor	1	2	8
7222	G. Severe Palmor Pallor H. Bruising or Bleeding	1	2 2	8
R208 R209	G. Severe Palmor Pallor	1	2	8

R211	Hypoglycaemia (Blood sugar <40 mg/dL)	<40 mg/dL	≥40	Test not
		<2.2mmol/	mg/dL	done
		1	or	
			>2.2mm	
			ol/l	
R212	Parasitemia	No parasite	s seen	1
		+		2
		++		3
		+++		4
		++++		5
		Test not do	ne	6
R213	Determine if the patient has severe malaria.	Severe	Uncom	Unable to
	If the patient was assessed positively for ANY 1 of the indicators in the	Malaria	plicated	determine
	gray boxes – the patient has severe malaria. Ensure the patient receives		malaria	
	adequate pre-referral dose of an anti-malarial and counsel the	1	2	3
	patient/caretaker to seek inpatient care urgently.			
	If the patient has none of the signs in the shaded gray boxes then the patient has uncomplicated malaria. Ensure the patient received the correct dose of antimalarial treatment.			
]		

USE THE FOLLOWING FOR CHILDREN < 5

R300	What reason/s does the Caregiver / Patient provide for coming facility	to the	YE	S	NO
	H) Fever/ Hot body?		1		2
	I) Diarrhoea?		1		2
	J) Vomiting everything?		1		2
	K) Cough/difficult breathing		1		2
	L) Ear problem?		1		2
	M) Skin Problem		1		2
	,		•		
	N) OTHERS (SPECIFY)	1		2
R301a	Take patient's temperature and record temperature				C
R301b	Record Patient's weight				Kg
R302	Does the child have any of the general DANGER SIGNS?:	Yes	No	Classifi	cation / DIAG
	A) Is the child unable to drink/breastfeed?	1	2		
	B) Is the child lethargic or unconscious?	1	2		
	C) Has the child had convulsions with this sickness?	1	2		
	D) Does the child vomit everything?	1	2		
Dooo	E) Is the Child Convulsing now?	1	2	0	
R303	Does the child have a cough or difficult breathing?	1	2 → 304		Pneumonia1
	A). For how long Days		0		onia2 or Cold – no
	B). Count the breaths in 1 minute Fast breathing?	1	2		onia3
	C). Look for Chest In-drawing	1	2	pricum	Jilla
	D). Stridor?	1	2		
R304	Does the child have Diarrhoea?	1	2 → 305	Severe	dehydration1
11004	A). For how long days	•	2 2 303	Some dehydration2	
	B). Does the child have sunken eyes?	1	2		ydration3
	C). Is the child able to drink?	1	2		persistent
	D). Is the child dehydrated? (skin pinch goes back only after 2	1	2	diarrhe	a4
	seconds).				ent diarrhea5 ery6
R305	Does the child have fever (by history or above 37.5)	1	2 → 307	Lincom	policated
11303	A). For how long Days	'	2-7-301	Malaria	
	B). If for more than 7 days – fever present every day?	1	2		evere Febrile
	C). Look or feel for stiff neck	1	2		e incl. Severe
R306	Does child have any of the signs of severe malaria? (other signs		_		a
	covered elsewhere in this form)			2	4
	A). Altered consciousness or behavior	1	2	Fever	– other
	B). Severe Pallor (severe anemia) / Palmor Pallor	1	2	cause.	
	C). Patient passing little or no urine?	1	2	No	0
	D). Patient passing dark or cola colored urine?	1	2		4
	E). Child having abnormal bleeding (from mouth, nose skin)?	1	2		
	C). Prostration (unable to eat or drink at all) D) Weak / Rapid Pulse	<u>1</u> 1	2		
	E). Jaundice	1	2		
R307	Does the child have measles now or in the last three months?	1	2 →	Severe	complicated
1307	2003 the office friedsies flow of itt the last tiffee months!	1	R308		s1
	A). Is there a generalized rash?	1	2	Į.	s with eye/mouth
	B). Does the child have one of these cough, runny nose or red	1	2		ation2
	eyes?				s3
	C). Does the child have deep and extensive mouth ulcers?	1	2		
	D). Does the child have pus draining from the eye?	1	2		
	E). Does the child have clouding of the cornea?	1	2		

R308	Does the child have an ear problem?	1	2→	Acute ear infection1
			R309	Chronic ear infection.2
	A). Is there ear pain?	1	2	No ear infection3
	B). Is there ear discharge? If Yes how long?	1	2	
	C). Is there tender swelling behind the ear?	1	2	
R309	Malnutrition and Anemia			Severe malnutrition1
	A. Is there severe visible wasting?	1	2	Severe anaemia2
	B). Is there oedema – of both feet?	1	2	Anaemia3
	C). Is the child very low weight for age?	1	2	Very low weight4
	D) Severe Palmar Pallor	1	2	No anaemia and not
				very low weight5
R310	Other diagnoses			
		i		
		i		

COMMENTS:

A-5 Assessment of Cases with Severe Malaria

FACILITY IDENTIFICATIO	N
Name of Region:`	FACILITY CODE
Name of District:	
Name of the facility	FACILTY TYPE
Name of Interviewer	OPER AUTH
Date:	
Type of Health Facility: (1 = Teaching Hospital 2=Regional hospital; 3 = District Hospital; 4=Other Hospital 5=Polyclinic 6=Health Center 7= Maternity Home 8=Clinic 9=CHPS 10=Other)	
Operating Authority: 1= Government; 2=Mission/Religious 3 = Private for profit 4 = Other)	

NO.	QUESTIONS	CODING CLASSIFICATION		
	Are there any cases of severe malaria on admission at the facility on the day of the assessment?	Yes1 No2	→S001 →END	

OBSERVATION #1

CONSENT – READ THIS STA CONSULTATION:	ATEMENT TO EACH CAREGIVER/PATIENT BEFORE (CONDUCTING THE
Project. We are conducting your treatment and case hist participation will help improve	and I am representing the Ghana Health Service and gresearch on the treatment of malaria at this facility. story for our study. We will not collect any personal invertible quality of care at health facilities in Ghana. You way affect your ability to receive health care at this form	We would like to observe nformation about you. Your ou do not have to participate.
Do you have any questions?	? Will you participate in our study? If Yes, can you p	lease sign the line below?
\$001. SIGNATURE OR THUM AGREEMENT TO PARTICIPAT	MBPRINT OF PATIENT/CAREGIVER INDICATES CARE ATE	E GIVER'S/PATIENTS

If you have any questions about this observation you may contact the people on this card (give them a card).

NO.	QUESTIONS	CODIN	G CLASSIFI	CATION
S100a	Gender of the Patient	Male (1)	Female (2)	
S100b1	Age years			
S100b2	If Child Under Five indicate months			
	If Child Under Five indicate months			DON'T
		YES	NO	KNOW
S101	Was this case laboratory-confirmed at the admission?	1	2	8
S102	Does the patient have at least one malaria smear per day of hospitalization until the parasitemia is negative?	1	2	8
S103	Assess whether the following signs or assessments were recorded	on the chart	-	
	A. Respiratory Distress	1	2	8
	B. Fast breathing is ≥ 50/min for ages <12 months and ≥ 40/min for ages 1-5 years)	1	2	8
	C. Temperature ≥ 38.5 C	1	2	8
	D. Convulsion	1	2	8
	E. Lethargy, unconsciousness, or change in behavior	1	2	8
	F. Weak but fast Pulse or > 110 beats/min	1	2	8
	G. Cyanosis	1	2	8
	H Jaundice	1	2	8
	I Capillary Refill > 3 seconds	1	2	8
	J Severe Palmor Pallor / Severe Anemia	1	2	8
	K. Bruising or Bleeding	1	2	8
S104	Are Lab Results Available?	1 → S105	2 → S109	8
S105	Hematocrit	<u><</u> 15%	≥15%	Test not done
S106	Hemoglobin	<5 g/dl	≥5g/dl	Test not done
S107	Hypoglycaemia (Blood sugar <40 mg/dL or 2.2mmol/L)	<40 mg/dL <2.2mmol/	≥40 mg/dL or >2.2mmol/l	Test not done
S108	Parasite Density/Count	++ +++ ++++ Result not a	available the	2 4 same day .5

S109	Was a lumbar puncture test done?	Yes (1)	No (2)	Don't Know(8)
S110	Note other lab tests conducted for this patient's stay:			

S111	Are the following monitored?	Yes	No	Frequency		
	A. Temperature	1	2			
	B. Blood Pressure	1	2			
	C. Respiratory Rate	1	2			
	D. Fluid intake/output	1	2			
	E. Lumbar Puncture done	1	2			
S112	How long after the start of symptoms was patient hospitalized?	One day or	less	1		
		One to thre	e days	2		
				33		
		Not clear fr	om the ch	art or patient.4		
S113	How long after arriving at the FIRST health facility was the patient	1-3 hours		1		
	hospitalized?	3-6 hours		2		
				3		
		Not clear from the chart or patient4				
S114	What is the patient current treatment?	Medication	n dose pe	r day?		
	A .					
	B.					
	C.					
	D.					
S115						
20	What is patient's weight?	·	Kg	3		

Observer Comments About the Case:

OBSERVATION #2

CONSENT – READ THIS STATEMENT TO EACH CAREGIVER/PATIENT BEFORE CONDUCTING THE CONSULTATION:
My name is and I am representing the Ghana Health Service and Quality Health Partners
Project. We are conducting research on the treatment of malaria at this facility. We would like to observe your treatment and case history for our study. We will not collect any personal information about you. Your participation will help improve the quality of care at health facilities in Ghana. You do not have to participate. If you decline, this will in no way affect your ability to receive health care at this facility.
Do you have any questions? Will you participate in our study? If Yes, can you please sign the line below?
\$002. SIGNATURE OR THUMBPRINT OF PATIENT/CAREGIVER INDICATES CARE GIVER'S/PATIENTS AGREEMENT TO PARTICIPATE

If you have any questions about this observation you may contact the people on this card (give them a card).

NO.	QUESTIONS		CODING CLASSIFICATION			
S200a	Gender of the Patient	Male (1)	Female (2)			
S200b1	Age years		` '			
S200b2	If Child Under Five indicate months					
	If Child Under Five indicate months	VEC	NO	DON'T		
		YES	NO	KNOW		
S201	Was this case laboratory-confirmed at the admission?	1	2	8		
S202	Does the patient have at least one malaria smear per day of hospitalization until the parasitemia is negative?	1	2	8		
S203	Assess whether the following signs or assessments were recorded	on the chart	•			
	A. Respiratory Distress	1	2	8		
	B. Fast breathing is ≥ 50/min for ages <12 months and ≥ 40/min for ages 1-5 years)	1	2	8		
	C. Temperature ≥ 38.5 C	1	2	8		
	D. Convulsion	1	2	8		
	E. Lethargy, unconsciousness, or change in behavior	1	2	8		
	F. Weak but fast Pulse or > 110 beats/min	1	2	8		
	G. Cyanosis	1	2	8		
	H Jaundice	1	2	8		
	I Capillary Refill > 3 seconds	1	2	8		
	J Severe Palmor Pallor / Severe Anemia	1	2	8		
	K. Bruising or Bleeding	1	2	8		
S204	Are Lab Results Available?	1 → S105	2 → S109	8		
S205	Hematocrit	<u><</u> 15%	≥15%	Test not done		
S206	Hemoglobin	<5 g/dl	≥5g/dl	Test not done		
S207	Hypoglycaemia (Blood sugar <40 mg/dL or 2.2mmol/L)	<40 mg/dL <2.2mmol/	≥40 mg/dL or >2.2mmol/l	Test not done		
S208	Parasite Density/Count	++ ++ +++ Result not a	available the s			

S209	Was a lumbar puncture test done?	Yes (1)	No (2)	Don't Know(8)
S210	Note other lab tests conducted for this patient's stay:			

S211	Are the following monitored?	Yes	No	Frequency		
	F. Temperature	1	2			
	G. Blood Pressure	1	2			
	H. Respiratory Rate	1	2			
	I. Fluid intake/output	1	2			
	J. Lumbar Puncture done	1	2			
S212	How long after the start of symptoms was patient hospitalized?	One day or	less	1		
				2		
		More than	three days	s3		
		Not clear fr	om the ch	nart or patient.4		
S213	How long after arriving at the FIRST health facility was the patient					
	hospitalized?		3-6 hours2			
				3		
		Not clear from the chart or patient4				
S214	What is the patient current treatment?	Medication	n dose pe	er day?		
	A.					
	B.					
	C.					
	D.					
C045						
S215	What is patient's weight?		Kg	S		

Observer Comments About the Case:

OBSERVATION #3

CONSENT – READ THIS STATEMENT TO EACH CAREGIVER/PATIENT BEFORE CONDUCTING THE CONSULTATION:
My name is and I am representing the Ghana Health Service and Quality Health Partners Project . We are conducting research on the treatment of malaria at this facility. We would like to observe your treatment and case history for our study. We will not collect any personal information about you. Your participation will help improve the quality of care at health facilities in Ghana. You do not have to participate. If you decline, this will in no way affect your ability to receive health care at this facility.
Do you have any questions? Will you participate in our study? If Yes, can you please sign the line below?
\$003. SIGNATURE OR THUMBPRINT OF PATIENT/CAREGIVER INDICATES CARE GIVER'S/PATIENTS AGREEMENT TO PARTICIPATE

NO.	QUESTIONS	CODIN	G CLASSIFI	CATION
S300a	Gender of the Patient	Male (1)	Female (2)	
S300b1	Age years			
S300b2	If Child Under Five indicate months			
	II Child Order Five indicate months	YES	NO	DON'T KNOW
S301	Was this case laboratory-confirmed at the admission?	1	2	8
S302	Does the patient have at least one malaria smear per day of hospitalization until the parasitemia is negative?	1	2	8
S303	Assess whether the following signs or assessments were recorded	on the chart		
	A. Respiratory Distress	1 1	2	8
	B. Fast breathing is ≥ 50/min for ages <12 months and ≥ 40/min for ages 1-5 years)	1	2	8
	C. Temperature ≥ 38.5 C	1	2	8
	D. Convulsion	1	2	8
	E. Lethargy, unconsciousness, or change in behavior	1	2	8
	F. Weak but fast Pulse or > 110 beats/min	1	2	8
	G. Cyanosis	1	2	8
	H Jaundice	1	2	8
	I Capillary Refill > 3 seconds	1	2	8
	J Severe Palmor Pallor / Severe Anemia	1	2	8
	K. Bruising or Bleeding	1	2	8
S304	Are Lab Results Available?	1 → S105	2 → S109	8
S305	Hematocrit	<u><</u> 15%	≥15%	Test not done
S306	Hemoglobin	<5 g/dl	≥5g/dl	Test not done
S307	Hypoglycaemia (Blood sugar <40 mg/dL or 2.2mmol/L)	<40 mg/dL <2.2mmol/	≥40 mg/dL or >2.2mmol/l	Test not done
S308	Parasite Density/Count	+ ++ +++ Result not available the same Test not done		2 3 4 same day .9

S309	Was a lumbar puncture test done?	Yes (1)	No (2)	Don't Know(8)
S310	Note other lab tests conducted for this patient's stay:			

S311	Are the following monitored?	Yes	No	Frequency		
	K. Temperature	1	2			
	L. Blood Pressure	1	2			
	M. Respiratory Rate	1	2			
	N. Fluid intake/output	1	2			
	O. Lumbar Puncture done	1	2			
S312	How long after the start of symptoms was patient hospitalized?	One day or less				
				2		
		More than	three days	s3		
		Not clear fr	om the ch	eart or patient.4		
S313	How long after arriving at the FIRST health facility was the patient					
	hospitalized?		3-6 hours2			
				3		
		Not clear from the chart or patient4				
S314	What is the patient current treatment?	Medication	n dose pe	er day?		
	A.					
	B.					
	C.					
	D.					
S315						
2010	What is patient's weight?		Kg	S		

Observer Comments About the Case:

OBSERVATION #4

	TATEMENT TO EACH CAREGIVER/PATIENT BEFORE CONDUCTING THE
CONSULTATION:	
Project. We are conducting your treatment and case his participation will help improve.	and I am representing the Ghana Health Service and Quality Health Partners gresearch on the treatment of malaria at this facility. We would like to observe story for our study. We will not collect any personal information about you. Your ove the quality of care at health facilities in Ghana. You do not have to participate. o way affect your ability to receive health care at this facility.
Do you have any questions	s? Will you participate in our study? If Yes, can you please sign the line below?
\$004. SIGNATURE OR THI AGREEMENT TO PARTICIF	JMBPRINT OF PATIENT/CAREGIVER INDICATES CARE GIVER'S/PATIENTS PATE

If you have any questions about this observation you may contact the people on this card (give them a card).

NO.	QUESTIONS	CODIN	CATION	
S400a	Gender of the Patient	Male (1)	Female (2)	
S400b1	Age years	. ,		
S400b2	If Child Under Five indicate months			
	II Child Olider Five indicate months			DON'T
		YES	NO	KNOW
S401	Was this case laboratory-confirmed at the admission?	1	2	8
S402	Does the patient have at least one malaria smear per day of hospitalization until the parasitemia is negative?	1	2	8
S403	Assess whether the following signs or assessments were recorded	on the chart	-	
	A. Respiratory Distress	1	2	8
	B. Fast breathing is ≥ 50/min for ages <12 months and ≥ 40/min for ages 1-5 years)	1	2	8
	C. Temperature ≥ 38.5 C	1	2	8
	D. Convulsion	1	2	8
	E. Lethargy, unconsciousness, or change in behavior	1	2	8
	F. Weak but fast Pulse or > 110 beats/min	1	2	8
	G. Cyanosis	1	2	8
	H Jaundice	1	2	8
	I Capillary Refill > 3 seconds	1	2	8
	J Severe Palmor Pallor / Severe Anemia	1	2	8
	K. Bruising or Bleeding	1	2	8
S404	Are Lab Results Available?	1 → S105	2 → S109	8
S405	Hematocrit	<u><</u> 15%	≥15%	Test not done
S406	Hemoglobin	<5 g/dl	≥5g/dl	Test not done
S407	Hypoglycaemia (Blood sugar <40 mg/dL or 2.2mmol/L)	<40 mg/dL <2.2mmol/	≥40 mg/dL or >2.2mmol/l	Test not done
S408	Parasite Density/Count	++ +++ ++++ Result not a	available the	

S409	Was a lumbar puncture test done?	Yes (1)	Don't Know(8)
S410	Note other lab tests conducted for this patient's stay:		

S411	Are the following monitored?	Yes	No	Frequency		
	P. Temperature	1	2			
	Q. Blood Pressure	1	2			
	R. Respiratory Rate	1	2			
	S. Fluid intake/output	1	2			
	T. Lumbar Puncture done	1	2			
S412	How long after the start of symptoms was patient hospitalized?	One day or	· less	1		
				2		
		More than	three days	33		
		Not clear fr	om the ch	art or patient.4		
S413	How long after arriving at the FIRST health facility was the patient					
	hospitalized?		3-6 hours2			
				3		
				art or patient4		
S414	What is the patient current treatment?	Medication	n dose pe	r day?		
	A .					
	B.					
	C.					
	D.					
S415						
	What is patient's weight?		Kg	3		

Observer Comments About the Case:

OBSERVATION #5

CONSULTATION:	IS STATEMENT TO EACH CAREGIVER/PATIENT BEFORE CONDUCTING THE
Project. We are cond your treatment and ca participation will help	and I am representing the Ghana Health Service and Quality Health Partners ucting research on the treatment of malaria at this facility. We would like to observe se history for our study. We will not collect any personal information about you. Your improve the quality of care at health facilities in Ghana. You do not have to participate. I in no way affect your ability to receive health care at this facility.
Do you have any ques	tions? Will you participate in our study? If Yes, can you please sign the line below?
S005. SIGNATURE OF AGREEMENT TO PAR	R THUMBPRINT OF PATIENT/CAREGIVER INDICATES CARE GIVER'S/PATIENTS TICIPATE

NO.	QUESTIONS	CODING	CODING CLASSIFICATION			
S500a	Gender of the Patient	Male (1)	Female (2)			
S500b1	Age years					
S500b2	If Child Under Five indicate months					
	II Child Offder Five indicate months			DON'T		
		YES	NO	KNOW		
S501	Was this case laboratory-confirmed at the admission?	1	2	8		
S502	Does the patient have at least one malaria smear per day of hospitalization until the parasitemia is negative?	1	2	8		
S503	Assess whether the following signs or assessments were recorded	on the chart	-			
	A. Respiratory Distress	1	2	8		
	B. Fast breathing is ≥ 50/min for ages <12 months and ≥ 40/min for ages 1-5 years)	1	2	8		
	C. Temperature ≥ 38.5 C	1	2	8		
	D. Convulsion	1	2	8		
	E. Lethargy, unconsciousness, or change in behavior	1	2	8		
	F. Weak but fast Pulse or > 110 beats/min	1	2	8		
	G. Cyanosis	1	2	8		
	H Jaundice	1	2	8		
	I Capillary Refill > 3 seconds	1	2	8		
	J Severe Palmor Pallor / Severe Anemia	1	2	8		
	K. Bruising or Bleeding	1	2	8		
S504	Are Lab Results Available?	1 → S105	2 → S109	8		
S505	Hematocrit	<u><</u> 15%	≥15%	Test not done		
S506	Hemoglobin	<5 g/dl	≥5g/dl	Test not done		
S507	Hypoglycaemia (Blood sugar <40 mg/dL or 2.2mmol/L)	<40 mg/dL <2.2mmol/	≥40 mg/dL or >2.2mmol/l	Test not done		
S508	Parasite Density/Count	++ +++ ++++ Result not a	available the	24 same day		

				Know(8)
S510	Note other lab tests conducted for this patient's stay:			
S511	Are the following monitored?	Yes	No	Frequency
	U. Temperature	1	2	
	V. Blood Pressure	1	2	
	W. Respiratory Rate	1	2	
	X. Fluid intake/output	1	2	
	Y. Lumbar Puncture done	1	2	
S512	How long after the start of symptoms was patient hospitalized?	One to three	e days three days	
S513	How long after arriving at the FIRST health facility was the patient hospitalized?	1-3 hours 3-6 hours More than	 6 hours	
S514	What is the patient current treatment?	Medication	n dose pe	r day?
	A.			
	B.			
	C.			
	D.			

Yes (1)

No (2)

_ Kgs

Don't

Observer Comments About the Case:

What is patient's weight?

S509

S515

Was a lumbar puncture test done?

A-6 Laboratory Quality Control Tool

FACILITY IDENTIFICATION					
Name of Region:`	FACILITY CODE				
Name of District:					
Name of the facility	FACILTY TYPE				
Name of Interviewer	OPER AUTH				
Date:					
Type of Health Facility: (1 = Teaching Hospital 2=Regional hospital; 3 = District Hospital; 4=Other Hospital 5=Polyclinic 6=Health Center 7= Maternity Home 8=Clinic 9=CHPS 10=Other Operating Authority:					
1= Government; 2=Mission/Religious 3 = Private for profit					
4 = Other)					
CONSENT – READ THIS STATEMENT TO EACH CAREGIVER BEFORE CONDUCTING THE LAB SLIDES COLLECTION: My name is and I am representing the Ministry of Health/GHS and Partners, USAID/Quality Health Partners Project. We are conducting research on the treatment of malaria at this facility. We would like to collect information about the lab diagnoses selected patients at this facility. We are asking you to conduct any malaria tests as ordered by the provider for patients in our study. While collecting a sample for your slide, we are asking you to collect a second thin and thick blood smear on a second slide for us. We will not collect any personal information about you. Your participation will help improve the quality of care at health facilities in Ghana. You do not have to participate. If you decline, this will in no way affect your ability to receive future training or assistance at this facility. If you have any questions about this quality control process you may contact the people on this card (give them a card).					
Do you have any questions for me?					
Do you agree for record your findings and collect specimens for the s	tudy?				
100a. SIGNATURE OF INTERVIEWEE INDICATES CARE GIVER'S AGR	EEMENT TO PARTICIPATE				

	Observation Number	M/F	Type of specie Pf, Pm, Po, Pv	Parasite count or density	Comment
Slide 001					
Slide 002					
Slide 003					
Slide 004					
Slide 005					
Slide 006					
Slide 007					
Slide 008					
Slide 009					
Slide 010					
Slide 011					
Slide 012					
Slide 013					
Slide 014					
Slide 015					
Slide 016					
Slide 017					

Slide 018			
Slide 019			
Slide 020			

B-1: Ante-Natal Care Facility Audit

	FACILITY IDENTIFICATION								
Name	of Region:`		FACILITY CODE						
Name	of District:								
Name	of the facility		FACILITY TYPE						
Date:			OPERATING AUTHORITY						
Name	of Interviewer:								
hospita 6=Hea	of Health Facility: (1 = Teaching Hospital 2=Regional; 3 = District Hospital; 4=Other Hospital 5=Polyclialth Center 7= Maternity Home 8=Clinic 9=CHPS her)								
	ting Authority: vernment; 2=Mission/Religious 3 = Private for pro	ofit							
	her)								
NO.	QUESTIONS		DDING CLASSIFICATION		GO TO				
700	Does this facility offer antenatal care?	NON	ANTENATALE OF THESE SERVICE	S.2					
WITH T	O GO TO WHERE THE MATERNAL HEALTH SE THE MOST SENIOR HEALTH WORKER INVOLVI								
	'H SERVICES. presenting the Ghana Health Service and the Qual	itv Hea	Ith Partners for this surve	v. We	are				
collection	ng information that will help us to better understand	d how m	nalaria treatment services	are pro	ovided.				
This inf	ormation will be used to design programs to improve	ve thes	e services. This part of th	e surve	ey will				
	estions about the services offered at this facility, the ke about 45 minutes – 1 hour to complete. All infor								
particip	ation in answering questions for this survey is volu	ntary. `	You can refuse to answer	any qu	uestion or				
	questions. We are asking for your help to ensure the								
	tions where someone else is the most appropriate ate your introducing us to that person.	person	to provide information, w	e would	u				
sign be		ırticipat	e in our survey? If Yes do	o you p	lease				
701	OLONATURE OF INTERVIEWEE INDICATE		TIOIDANT ACCESS	 IT TO					
	SIGNATURE OF INTERVIEWEE INDICATES PARTICIPATE	5 PAR	HCIPANT AGREEMEN	11 10					

NO.	QUESTIONS		CODING CLASSIFICATION		GO TO		
702	How many days in a week are anten	atal care					
	services provided at the facility?		# DAY	S			
703	Are antenatal care services being pro-	ovided	YES1		1		
	at the facility today?		NO		2		
704		s focus-ANC practiced at this facility?					
				KNOW			
705	How many midwives are assigned to Al services at this facility?	NC					
	WRITE IN THE NUMBER. FOR NUMBER AT THE RESERVE AT T						
	LESS THAN 10 – WRITE A ZERO BEF	ORE					
706	THE NUMBER.	ly ovoile	blo ot th	sic facility?			
706	Are the following services routinely available at this facility?						
	PART OF ROUTINE SERVICES	Routine	ly done	Not Done	Don't Know		
		/ Availa		Routinely			
	a. Routine laboratory services		1	2	8		
	Blood _ mps						
	b. Blood HB		1	2	8		
	c. Urine for RE		1	2	8		
	d. Stool for RE		 1	2	8		
	e. G6PD testing		1	2	8		
	f. Iron folate routinely distributed?		1	2	8		
707	Are the following equipment availa	able at t	his facil	ity?	•		
	<u> </u>		ilable	Available not	Not		
	APPARATUS/TOOL	Func	ioning	Functioning	Available		
	g. Adult weighing scale		1	2	8		
	h. Blood Pressure apparatus		1	2	8		
708	Is Intermittent Preventive Treatmen	t	Yes		1		
	(IPT) with SP provided for pregnant	t l	No		2	→ 713	
	women at this facility?		Don't Kr	now	8	→ 713	
709	What type of SP do you keep in sto	ck?	Fansida		4		
			Don't kn	ow	3		
			Other (S	Specify)	4		
710	Has there been a stock out of SP in	this	Yes		1		
' 0	facility in the past 6 months ?						
	isomy in the paor o months:			now			
711	What is done when there is a shorta	age?		prescription for			
				specify)			
712	How much does SP cost in this faci	lity?	Nothing		1		
			5 Gp - 1	0 Gp	3		
				s			

713	Does this facility have copies of the following: If yes, ask to see a copy	•		Not available	Not determined
	a. Standard treatment guidelines	a. Standard treatment guidelines		2	9
	b. New anti malaria drug policy (Nov	v 2004)	1	2	9
	c. IPT Training Manual	•	1	2	9
714	How many staff have been trained in the use of IPT?	Don't kr	now9	9	
715	Do CHNs dispense SP in the community for IPT2 and IPT3?	NO			2 → 717
		DON'T	KNOW		8 →717
716	How are doses given in the community reflected in the ANC register?	LATER	TO ANC REGIS	OOK AND TRA	1
	CIRCLE ALL THAT APPLY	A TALLY FORM IS USED AND TRANSFERRED TO ANC REGISTER2			
	OTHER:	ANC RE	EGISTER (FOR	OUTREACH) US	SED3
		A LIST	OF MOTHERS I	AIN REGISTER S TAKEN TO TH CKED AGAINS	HE FIELD
		OTHER	(SPECIFY)		6
717	Has there been a supervisory visit for ANC services in the past 6 months at this facility?	YES NO DON'T	KNOW		1 2 8
718	Was the supervisor from this facility or from outside this facility?	OUTSIE	DE THIS FACILI	TY	2
719	Have you (person providing services) received any in-service training in ANC in the last 3 years ?	YES			1 2

Statistics (from monthly returns)

720	Month(1)	ANC Registra nts(2)	ANC Attendan ce(3)	IPT1(4)	IPT2(5)	IPT3(6)	# with HB at 36 weeks	# with HB < 7gm/dl at 36 weeks
А	October 2007							
В	November 2007							
С	December 2007							
D	January 2008							
E	February 2008							
F	March 2008							

ANC Record Review

721	Is there a register where client information from ANC visits is recorded? IF YES, ASK TO SEE REGISTER.	YES, REGISTER SEEN1 YES, REGISTER NOT SEEN	→END →END
722	How recent is the date of the most recent entry for ANC?	WITHIN THE PAST 7 DAYS1 > 7 DAYS2	
723	INTERVIEWER: Does the ANC register in this facility have records on IPT services offered?	YES1 NO2	→END

ANC RECORD REVIEW METHODOLOGY

Take the total number of ANC registrants for the period Oct 2007 – Mar 2008 from Question 718G above. Then divide this number by 10. This is your sampling interval and will give you exactly 10 records from the register.

Then select a random number between 1 and your sampling interval. That is where you start. Record the information from that record. Then from that record, count down the number of records in your sampling interval and record the information from that record. Then count down again the number in your sampling interval and record the information from that record. Continue until 10 records from the period Oct – Dec 2007 have been sampled.

Fill in the information from your facility in the charts

Month	Number of ANC	Number of ANC registrants
	registrants	at your facility
Oct 2007	450	
Nov 2007	275	
Dec 2007	125	
Jan 2008	250	
Feb 2008	312	
Mar 2008	157	
Total	1569	
Total / 10	1569/10 = 157	
Sampling Interval	157	

	Number to be selected in example	Your number to be selected
Record 1	(Random number between 1 and 157) = 87	
Record 2	87+157 = 244	
Record 3	244+157 = 401	
Record 4	401+157=558	
Record 5	558+157=715	
Record 6	715+157=872	
Record 7	872+157=1029	
Record 8	1029+157=1186	
Record 9	1186+157=1343	
Record 10	1343+157=1500	

No	No Date of Age ANC1 dd/mm /yy		Age Parity	Parity Gest (in weeks)	Gest (in Total number of ANC visits to date	mber last ANC ANC visit sits to	IPT1 If no writing indicate by		IPT2 If no writing indicate by	IPT3 If no writing indicate by		ITN if there is a date or a tick in this column circle 1 for Yes if no date 2 for no.		
							Date	FHt	Date	FHt	Date	FHt	Yes	No
1													1	2
2													1	2
3													1	2
4													1	2
5													1	2
6													1	2
7													1	2
8													1	2
9													1	2
10													1	2

B-2: Ante-Natal Care Observation and Client Exit Interview

FACILITY IDENTIFICATION					
Name of Region:`	FACILITY CODE				
Name of District:					
Name of the facility	FACILITY TYPE				
Date:					
Name of Interviewer:	OPERATING AUTHORITY				
Type of Health Facility: (1 = Teaching Hospital 2=Regional hospital; 3 = District Hospital; 4=Other Hospital 5=Polyclinic 6=Health Center 7= Maternity Home 8=Clinic 9=CHPS	PROVIDER				
10=Other)	OBSERVATION				
Operating Authority: 1= Government; 2=Mission/Religious 3 = Private for profit 4 = Other)	(Number the patients sequentially as they are observed)				
Provider Type: Midwife = 1, Community Health Nurse =2, CHO = 3, Other =4.					

	OBSERVE TO SEE WHETHER THE PROVIDER CARRIES OUT THE FOLLOWING ACTIVITIES					
801	Is this the Client's first visit?	Yes 1	No 2			
802	Routine Treatment and services for ANC related to malaria	Observed	Not Observed			
	a. Medical History Taken at first visit	1	2			
	b. Obstetrical History Taken	1	2			
	c. Physical examination done	1	2			
803	Which laboratory investigations are ordered during the consultation you observed?	Observed	Not Observed			
	A. Blood film for malaria parasites	1	2			
	B. Haemoglobin level	1	2			
	C. Sickling	1	2			
	D. G6PD	1	2			
	E. HIV status	1	2			
	F. Electrophoresis					
	G. Urine Protein	1	2			

Educa	tion on Malaria in Pregnancy	Observed	Not Observed
804	Is Client educated on following effects of malaria on pregnan	cy during the c	onsultation?
	A. Malaria can cause Anaemia in the woman and the	1	2
	effects of anemia on pregnancy		
	B. That malaria may cause a spontaneous abortion	1	2
	C. That severe malaria can endanger the life of the mother	1	2
	and the unborn child?		
	D. That low birth weight can result	1	2
IPT US		Observed	Not Observed
805	Does service provider determine if Client is qualified for IPT b	y asking if clie	nt
	A. has received treatment of SP in less than one month	1	2
	B. is taking co-trimoxazole to treat other infections	1	2
	C. is pregnant at least 16 weeks	1	2
	D. is pregnant less than 36 weeks	1	2
	E. If woman has history of allergy or is G6PD deficient	1	2
806	Does the provider counsel the woman on the possible side effects of IPT?	1	2
807	Does the provider counsel the client on what to do if she has side effects from IPT medication?	1	2
808	Does the provider give IPT1, IPT2 or IPT3 as appropriate (using DOT)?	1	2
809	Does the provider record the IPT status on the Client ANC	1	2
	card?		
Other	Preventive Measures	Observed	Not Observed
810	Does the provider discuss client use of ITN, indoor residual	1	2
	spray, mosquito repellants or that she should avoid staying outside from dusk until dawn?		
811	Does the provider tell the client when to return for her next visit?	1	2

When the consultation concludes – politely ask the client if you can talk with her outside the consulting room.

CLIENT EXIT INTERVIEW

	ASK CLIENT THE FOLLOWING QUESTIONS			
900	How old is this pregnancy?	·		
		weeks		
901	Can you tell me what effects malaria can have on	Anaemia1		
	pregnancy?	Spontaneous Abortion		
		Severe malaria		
	DO NOT PROMPT	Low Birth Weight//		
		Don't know/Can't remember		
	CIRCLE ALL THAT APPLY	Other (Specify	8	
000	De very elega via den en ITNI ell the time	All the stice of	₹005	
902	Do you sleep under an ITN all the time,	All the time1 Sometimes2	→ 905	
	sometimes, or not at all?	Not at all3		
903	If Sometimes or Not at all. What do you do to	Indoor residual Spraying1		
903	If Sometimes or Not at all, What do you do to avoid getting malaria?	Mosquito Repellents2		
	DO NOT PROMPT	Avoids staying outside after		
	CIRCLE ALL THAT APPLY	dark3		
		Has screen in the house4		
904	What are the reasons why you do not sleep under	Too hot1		
001	ITN all the time?	Prefer to put my kids under		
		net instead2		
		Don't have a net3		
		No room to hang a net4		
		Don't like nets5		
		Other6		
905	Do you know that there is a medicine that can	Yes1		
	prevent pregnant women from getting malaria?	No2	→ 910	
906	Do you know how many doses of the medicine	Correct (3 Doses)1		
	(SP) you are supposed to take during your	Incorrect (Not equal to 3		
	pregnancy to prevent malaria?	doses)2		
		Don't know8		
907	Were you given the medicine (SP) to prevent	Yes1	3 0 1 5	
	malaria today?	No2	→ 910	
908	Did you take it in front of the midwife or were you	Given water to take in front of		
	given it to take home or something else?	midwife1		
		Given to take at home2		
		Asked to buy at pharmacy and		
		take3 Other4		
		(Specify)		
		(Opecity)		
909	What were you told about the side effects from IPT	Rare, but serious skin		
505	medication?	reactions1		
		Itching2		
	_	g		

	<u>-</u>		
		Nausea or vomiting3	
		Blurred vision4	
		Not told anything5	
		Other6	
910	May I see your ANC card please? Note the dose	IPT11	
010		IPT22	
	the patient has been given to date.		
		IPT33	→ 911
		Not Noted on Card4	→ 911
		No dose given5	→ 911
911	Ask Only If client has had only IPT1 or IPT2.	In a month's time1	-
511	1		
	When were you asked to return for your next dose	More than a month2	
	of IPT?	Can't remember3	
		Was not told4	
912	In general, were you satisfied with the care you	Yes1	→END
•	received in this facility today?	No2	→ 913
040	, , ,	140	2313
913	If no, what were you unsatisfied about?		

Thank Client for Her Time

IF THIS IS THE LAST CONSULTATION OBSERVED. HOW MANY REMAINING ANC PATIENTS REMAIN TO BE SEEN BY THIS PROVIDER AT THE TIME YOU CLOSED?