

**ADDIS ABABA UNIVERSITY
SCHOOL OF GRADUATE STUDIES**

**ANTI-MALARIAL DRUG AND MOSQUITO NET USE
PATTERN IN PAWE SPECIAL WOREDA: A
COMMUNITY BASED SURVEY**

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Table of contents	Page
Acknowledgements	i
List of tables	iv
List of figures	v
List of annexes	vi
List of abbreviations	vii
Summary	ix
1. INTRODUCTION	1
2. LITERATURE REVIEW	3
2.1. Overview	3
2.1.1. History of the disease	3
2.1.2. Biology of the malaria parasite and vector	3
2.1.3. Burden of malaria	5
2.2. Malaria control	7
2.2.1. Malaria prevention	8
2.2.2. Malaria treatment & trends in Ethiopia	9
2.3. Self-medication with anti-malarials	12
2.4. Drug hoarding	14
2.5. The rationale of the study	15
3.OBJECTIVES OF THE STUDY	16
3.1. General objective	16
3.2. Specific objectives	16
4. METHODOLOGY	17
4.1. Study area and population	17
4.2. Study design	18
4.3. Sample size determination	18
4.4. Sampling procedures	19
4.5. Data collection	20
4.5.1. Quantitative study	20
4.5.1.1. Training and supervision	21
4.5.1.2. Pre-testing	21

4.5.2. Qualitative study	21
4.6. Study variables	22
4.6.1. Independent variables	22
4.6.2. Dependent variables	22
4.7. Data processing and analysis	22
4.8. Data quality management	23
4.9. Operational definitions	23
4.10. Ethical considerations	25
5. RESULTS	26
5.1. Findings from cross sectional study	26
5.1.1. Socio-demographic characteristics of the respondents	26
5.1.2. Reported illnesses	29
5.1.3. Perceived malaria illness	30
5.1.4. Knowledge about malaria: cause, transmission, signs & symptoms, prevention and diagnosis	31
5.1.5. Actions taken against perceived malaria/symptom complexes	34
5.1.5.1. Self-medication practices	35
5.1.5.2. Health institution visits	37
5.1.6. Knowledge about malaria treatment and anti-malarial drugs	38
5.1.7. Anti-malarial drug hoarding	41
5.1.8. Mosquito net coverage	42
5.1.9. Traditional remedies used to treat malaria	45
5.2. Qualitative results	46
5.2.1. Findings of FGDs with mothers	46
5.2.2. Findings of key informants' interviews	56
6. DISCUSSION	60
7. STRENGTHS AND LIMITATIONS OF THE STUDY	73
8. CONCLUSION	75
9. RECOMMENDATIONS	76
REFERENCES	78
ANNEXES	

List of tables	Pages
Table 1: Socio-demographic characteristics of respondents of cross-sectional survey in Pawe Special Woreda, 2008	28
Table 2: Reported illnesses by different socio-demographic variables in Pawe Special Woreda, 2008	29
Table 3: Reported illnesses in Pawe Special Woreda, 2008	30
Table 4: Distribution of reported malaria illness among study participants in Pawe Special Woreda, 2008	31
Table 5: Level of awareness about malaria: its cause, signs & symptoms, transmission and prevention in Pawe Special Woreda, 2008	33
Table 6: Anti-malarials used for self-medication practices in Pawe Special Woreda, 2008	36
Table 7: Factors associated with self-medication with anti-malarials in Pawe Special Woreda, 2008	37
Table 8: The different methods used to differentiate anti-malarial drugs from others in Pawe Special Woreda, 2008	40
Table 9: Reported consequences of non-adherence to anti-malarial medication in Pawe Special Woreda, 2008	40
Table 10: Factors associated with possession of more than one reported mosquito net in Pawe Special Woreda, 2008	44

List of figures	pages
Figure 1: Schematic presentation of sampling procedure	19
Figure 2: Knowledge of respondents about the diagnosis of malaria in Pawe Special Woreda, 2008	34
Figure 3: Actions taken against reported malaria/symptom complexes in Pawe Special Woreda, 2008	35
Figure 4: Diagnosis results of ill individuals who visited health institutions in Pawe Special Woreda, 2008	38
Figure 5: Knowledge of respondents about the names of the different anti-malarial drugs in Pawe Special Woreda, 2008	39
Figure 6: Commonly hoarded anti-malarial drugs at household level in Pawe Special Woreda, 2008	41
Figure 7: Mosquito net possession in Pawe Special Woreda, 2008	42
Figure 8: One of the FGD sessions conducted in Pawe Special Woreda, 2008	46

List of abbreviations

AADAC	Alberta Alcohol and Drug Abuse Commission
ACT	Artemisinin-based combination therapy
AHS	Alberta Health Services
AL	Artemether-lumefantrine
AMREF	African Medical and Research Foundation
AOR	Adjusted odds ratio
Asl	Above sea level
BGR	Beneshangul-Gumuz Region
BGRHB	Beneshangul-Gumuz Regional Health Bureau
CDC	US Center for Disease Control and Prevention
CI	Confidence interval
COR	Crude odds ratio
CSA	Central Statistical Agency of Ethiopia
DALYS	Disability adjusted life years
DDT	Di chloro diphenyl trichloro ethane
EC	Ethiopian calendar
EDHS	Ethiopian Demographic and Health Survey
ETB	Ethiopian birr
FGD	Focus Group Discussion
Fig	Figure
GNP	Gross National Product
HEW	Health-Extension Workers
HH	Household
IRS	Indoor residual spraying
ITNS	Insecticide treated bed nets
KII	Key informant interview
Kms	Killo meters
LLIN	Long-lasting insecticidal net

M	meter
MOH	Federal ministry of health of Ethiopia
MOP	Malaria operational plan
NEHC	Navy Environmental Health Center
OPD	Out Patient Department
OR	Odds ratio
PF	Plasmodium falciparum
PMI	US president's malaria initiative
PV	Plasmodium vivax
SD	Standard deviation
SPSS	Statistical Package for Social Sciences
SP	Sulfadoxine-Pyrimethamine
SSA	Sub-Saharan Africa
TV	Television
UNICEF	United Nations International Children's Fund
USAID	United States Agency for International Development
WHO	World Health Organization
Yr(s)	Year(s)

Thirty six, (8.3%) of the households hoarded anti-malarials at home. The survey shows that 427(98.8%) respondents reported to own at least one mosquito net. Care givers' occupation and household family size were associated with reported possession of more than one mosquito net. Being a housewife (OR=2.96, CI=0.92, 9.59, P=0.04) was highly associated with possession of more than one mosquito net. Households with a family size of less than or equal to 5 persons (OR= 0.05, CI=0.02, 0.17, P=0.000) were found to have a significantly less chance of possessing more than one mosquito net in comparison with those having a family size of more than 5 persons .

Conclusion and Recommendations: The prevalence of perceived malaria in the study area which is 6.1%, had declined by more than 50% when compared with previous studies reported in Ethiopia, and also it is very smaller than the results obtained from other developing countries. Though there is no available comparison data, the practice of self-medication with anti-malarials for perceived malaria illnesses/symptom complexes in the study place looks high. The study revealed that the percentage of households stockpiling anti-malarial drugs to be similar or lower than other finding in Africa. The coverage of mosquito nets in the area is encouraging compared to most findings in this country and other countries around the world, although the set goal (100%) was not yet achieved.

Continuous education and awareness creation of residents: on the cause, transmission, prevention and treatment of malaria; on the consequences of rampant self-medication; on the problems associated with drug hoarding; on the advantages and nature of mosquito nets distributed should be maintained if the malaria control efforts need to be successful within a short period of time.

Key Words: Anti-malarial drugs, Mosquito net, Self-medication, Perceived malaria, Drug hoarding, Pawe.

1. INTRODUCTION

Malaria is a global major disease caused by infection with any of four species of the protozoan parasite *Plasmodium* (i.e., *P. falciparum*, *P. vivax*, *P. ovale*, *P. malariae*) (Desai and Parise, 2002), but one, *Plasmodium falciparum*, accounts for the majority of instances of morbidity and mortality (White, 2004).

Anopheles arabiensis, a member of the *An. gambiae* complex, is the primary malaria vector, with *An. funestus*, *An. pharoensis* and *An. nili*, as secondary vectors (PMI, 2008).

Malaria continues to be one of the most important and devastating infectious diseases in developing areas of the world (CDC, 2007). Its transmission occurs in more than 100 countries. Regions include Africa, Asia, Islands of the South, West, and Central Pacific Ocean, Latin America, certain Caribbean Islands, and Turkey (NEHC, 2000). Around 3.2 billion people are at risk of malaria each year, with around 500 million people proceeding to clinical disease, and 2-3 million deaths occurring. Over 90% of these deaths occur in Sub-Saharan Africa (SSA) (Barry, 2005).

Malaria has been a major challenge to both public health and socio-economic development particularly in countries of SSA (Deressa, 2004). It is endemic in 42 of the 46 countries of the WHO African region and ranks in the top five causes of illness and death (WHO, 2004). In the 30 seconds it takes to read this sentence and the next, malaria will kill another African child (AMREF, 2005). The burden of morbidity and mortality is biased towards young children, not yet immune to clinical symptoms, and pregnant women where parasites are sequestered in the placenta (Barry, 2005).

In Ethiopia, malaria epidemics generally occur immediately after the long rains in September to November and in some places also after the short-lived shower rains in March to May (Ramos et al., 2003). Malaria is a major public health problem & the leading communicable disease in Ethiopia. It has been consistently reported as one of the three leading causes of morbidity and

2. LITERATURE REVIEW

2.1. Overview

2.1.1. History of the disease

Malaria is a life-threatening disease caused by the *Plasmodium* parasite. It was previously thought to come from fetid marshes, hence the name *male aria*, from the Italian, 'bad air'. During the past 100 years, an estimated 150-300 million people have died from malaria. This translates to 2-5% of all deaths globally in the 20th century having malaria-related causes (WEF, 2006).

It is now generally held that malaria arose in our primate ancestors in Africa and evolved with humans, spreading with human migrations first throughout the tropics, subtropics, and temperate regions of the Old World and then to the New World with explorers, missionaries, and slaves (Cox, 2002).

Although the disease has been eradicated in most of North America and Europe, malaria persists in many developing countries, primarily in Sub-Saharan Africa (SSA), Latin America and Asia (WEF, 2006).

2.1.2. Biology of the malaria parasite and vector

Malaria is transmitted through the bite of an infected female Anopheles mosquito (Oberlander & Elverdan, 2000; Lo Re III and Gluckman, 2003) and it is caused by the protozoan parasite *Plasmodium* (WHO, 2007^e) which spends its life cycle both in humans and certain species of mosquitoes (WHO, 2002). The Four species of *Plasmodium* that cause malaria in humans are *P. falciparum*, *P. vivax*, *P. malariae* and *P. ovale* (Desai and Parise, 2002; Lo Re III and Gluckman, 2003). Similar parasites are common in monkeys and apes (Cox, 2002). *P. falciparum* and *P. vivax* are the two most common types of malaria (WEF, 2006) and of these, *Plasmodium falciparum* is the most important in most parts of the tropics, being responsible for most severe illnesses and deaths (White, 2004; WHO, 2002). Occasional infections with monkey malaria parasites, such as *P. knowlesi*, also occur (WHO, 2006).

Ethiopia has adopted the use of ITNs as one of its vector control strategies primarily in selected malarious areas with the view to a gradual scaling-up of the intervention. The use of mosquito nets is, however, limited and there are a number of possible explanations for this low coverage. These may be due to absence of a sustainable mechanism for the distribution of ITNS, lack of cultural exposure to the use of mosquito nets, lack of awareness, low acceptance by the community, and concerns regarding its high cost (Jimma et al., 2005).

Only about 6 percent of households in Ethiopia own a mosquito net whether treated or untreated. The percentage of households having more than one net is about 1 percent. Five percent of households own at least one ever-treated net. Regarding Benishangul-Gumuz regional state, 15.4 percent of households had at least one bed nets (whether treated or untreated), and only 2.3% of households possess more than one (CSA and ORC Macro, 2006).

In some areas, malaria transmitted by vectors that rest indoors can be prevented or controlled by indoor residual spraying with DDT or Malathion, as per WHO recommendations, is one of the major malaria vector control interventions (WHO, 2002; WHO, 2005^a; CSA and ORC Macro, 2006).

Eliminating breeding sites and killing larvae, pupae and adult mosquitoes will help to reduce the number and longevity of vectors (WHO, 2002). In areas where it is a viable option, larval control must be carried out thoroughly if it is to have an impact (WHO, 2005^a).

2.2.2 Malaria treatment & trends in Ethiopia

Malaria is a curable disease and not an inevitable burden. Effective medicines are available (WHO, 2005^b). The first and foremost malaria control strategy promoted by the World Health Organization (WHO) and adopted by most African countries emphasizes the need for treatment of malaria episodes with an efficacious drug within 24 hours after onset of symptoms (Hetzl et al., 2006).

During decades, chloroquine, a 4-aminoquinoline, was largely used for treatment and prophylaxis of acute malaria with an amazing impact on malaria control (Krettli, 2001). Chloroquine resistance, which first appeared in East Africa in the late 1970s, has now spread throughout most of the continent, and resistance to sulfadoxine–pyrimethamine (Fansidar) has followed rapidly (Collins, 2000). *P.falciparum* is now highly resistant to chloroquine in most malaria-affected areas (White, 2004).

The emergence and rapid spread of *P. falciparum* resistance to commonly used anti-malarial drugs poses a serious challenge to the effectiveness of early diagnosis and prompt treatment as a priority strategy within the current malaria control. Effective treatment, as an intervention, depends highly on anti-malarial drugs which should be safe, effective, available, affordable and acceptable to populations at risk (WHO, 2003). Chloroquine and SP were affordable drugs at approximately US\$0.10–0.20 per adult course (Yeung et al., 2004; WHO, 2004), and their safety and efficacy as oral regimens means they have generally been readily accessible (Shunmay et al., 2004; Yeung et al., 2004). As a response to the anti-malarial drug resistance situation, WHO now recommends that treatment policies for falciparum malaria in all countries experiencing resistance to monotherapies, such as chloroquine, sulfadoxine/pyrimethamine and amodiaquine, should be combination therapies(CT), preferably those containing an artemisinin derivative (WHO, 2005^b). The rationale for using drugs in combination is well established in the treatment of tuberculosis, infection with human immunodeficiency virus, and cancer (Yeung et al., 2004). The basic tenet of combination therapy is that the probability of resistance developing simultaneously to two chemotherapeutic agents with independent mechanisms of action is extremely low, of the order of once in 10^{12} treatments. The effect of combination therapy is enhanced by the inclusion of an artemisinin derivative (Bloland et al., 2000). Artemisinin anti-malarials decrease parasite density more rapidly than other anti-malarial drugs (Lalloo et al., 2007; Bloland et al., 2000). When used alone, the short half-life of the artemisinin derivatives minimizes the period of parasite exposure to sub-therapeutic blood levels. In combination with another drug with a longer half-life, the short half-life and rapid parasite clearance time of artemisinin derivatives mean that many fewer parasites are exposed to the companion drug alone after elimination of the artemisinin component. Furthermore, exposure occurs when blood levels

of the drug close to the maximum are still present. Another benefit of artemisinin combinations is the 90% reduction in gametocyte levels in treated patients. These characteristics minimize the probability that a resistant mutant will survive therapy and may also reduce overall malaria transmission rates (Bloland et al., 2000).

Although combination therapy is accepted as the rational approach to case management in Africa (WHO, 2001), there are a number of concerns about widespread deployment of ACT, the chief one being cost. ACTs are relatively expensive (Coleman et al., 2004), currently cost more than US\$1 for an adult course (although this cost is decreasing), so for them to be widely deployed as first-line therapy, substantial subsidy will be required to ensure that they are available to everyone, including those who cannot afford the market price (Yeung et al., 2004). A second concern is lack of post-marketing surveillance data on the new therapies (WHO, 2004) and thirdly, there is little or no information on the safety and efficacy of combination treatment in pregnant women and young children, which are specific high-risk groups in Africa (WHO, 2001). The fourth concern is that by deploying the artemisinin derivatives now, we risk losing our most valuable anti-malarial, a potentially catastrophic event (Yeung et al., 2004).

In Ethiopia the first-line anti-malarial drug for the treatment of malaria has been changing over the past decade. The main reason for change was the level of efficacy of the drugs. Chloroquine was the first-line anti-malarial drug for the treatment of uncomplicated malaria until 1998. However, because of the high level of failure (65%) of chloroquine for the treatment of uncomplicated falciparum malaria that was detected through a nationwide study conducted in 1997/98, the drug was replaced by SP/Fansidar (MOH, 2004; CSA and ORC Macro, 2006).

At the time of the introduction of SP/Fansidar as the first-line drug, the level of treatment failure observed was about 7%. In subsequent years, however, unpublished reports from isolated studies indicated higher treatment failure rates. As a result, a nationwide study on the therapeutic efficacy of SP/Fansidar for the treatment of uncomplicated falciparum malaria was conducted in 2003 and a mean treatment failure rate of 36% (ranging from 20-54%) was reported (MOH, 2004).

Cognizant of the high treatment failure rates of SP/fansidar and the need to shift to more effective anti-malarial drugs, the Ministry of Health-after a series of consultative meetings with experts in the field and based on WHO recommendations-decided to introduce the Artemisinin-based Combination Therapy (ACT) drug Artemether-lumefantrine in July 2004 (MOH,2004; CSA and ORC Macro, 2006). The introduction of the new ACT drug and the phasing out of the old drug was estimated to take up to two years given the limited supply of the new drug and the size of the country (CSA and ORC macro, 2005). Quinine is the second line anti-malarial drug and these drugs are supposed to be provided by health professionals (i.e. prescription drugs) (MOH, 2004).

As mentioned above, treatment failures and mortality due to malaria has risen in recent years, probably due to increasing resistance to anti-malarial medicines (WHO, 2006). The main factor that leads to the aforementioned problem is irrational use of anti-malarial drugs, such as inappropriate self-medication. Irrational use is the use of medicines in a way that is not compliant with rational use as defined below:

Rational use of drugs requires that patients receive medications appropriate to their clinical needs, in doses that meet their own individual requirements for an adequate period of time, and at the lowest cost to them and their community (Bhatnagar et al., 2003).

In the absence of a clear alarming evidence of resistance, the treatment of choice for non-falciparum malaria is a 3-day course of oral chloroquine, to which only a limited proportion of *P. vivax* strains have gained resistance (WHO, 2005^a; Lalloo et al., 2006).

2.3. Self-medication with anti-malarials

In developing countries, most illnesses are treated by self-medication (Awad et al., 2005). It is also recognized that treatment of malaria usually starts at home and households would seek care outside if home treatment failed (Deressa, 2007^a). The decision for self-medication may be influenced by different factors (cultural, economical, psychosocial, etc) that culminate in the utilization of various therapeutic approaches in searching for a relief (Abula and Worku, 2001). A major shortfall of self-medication is the lack of clinical evaluation of the condition by a trained medical professional, which could result in missed diagnosis and delays in timely and appropriate

treatment, which may end up in tragic consequences (Abula and Worku, 2001; Awad et al., 2005).

In addition, self-treatment of malaria is common following self-diagnosis mainly based on presumptive symptoms of malaria. Inappropriate self-medication results in wastage of resources, increases resistance of pathogens, and generally entails serious health hazards such as adverse reaction & prolonged suffering (Worku and G/Mariam, 2003). Antimicrobial resistance is a current problem world-wide particularly in developing countries, where antibiotics are often available without a prescription (Awad et al., 2005).

The type and extent of self-medication and the reasons for it may vary from country to country. In developing countries, both modern drugs and traditional medicines are commonly used for self-medication. It was also noted that prescription-only-medications could easily be obtained without prescriptions for self-medication in developing countries like Ethiopia. The use of such drugs without the knowledge of physicians can be less beneficial or even be dangerous for the patient. The efficacy and safety of most traditional medicines used in Ethiopia is not scientifically proven, and there is lack of precision in dosage by traditional healers. However, the role of traditional medicines cannot be undermined provided that large proportion of the population rely on it (Abula and Worku, 2001). According to WHO, up to 90% of the population in developing countries uses TM (traditional medicine), including medicinal plants to help meet their primary health care needs (Vanderbroek et al., 2004). Thus, an extensive research work is required to ensure its safety and efficacy (Abula and Worku, 2001).

Self-treatment of common illnesses by lay people is common in economically deprived countries (Shankar et al., 2003). In part, this arises through poor geographic access to facilities with a trained health worker, but the popularity of self-treatment is strengthened by the frequent drug shortages, long waiting times and user fees common in public health facilities (Marsh et al, 2004), previous experience of illness, economic constraints (Shankar et al., 2003) and the issue of curative stance of drugs are worth mentioning (Worku and G/Mariam, 2005). In all areas where self-treatment practices have been studied, they have been typified by the use of inappropriate

children. On the other hand, the availability of anti-malarials in places other than the health facilities like retail pharmacies, drug stores and ordinary shops where over the counter dispensing is practiced have emerged contributing to the factors that promote home drug storage. Improper drug storage and rampant self-medication are some of the factors that may contribute to an increase in the development of drug resistance by malaria parasites towards anti-malarials. Home stocked drugs may lose potency due to poor storage, as a result of exposure to heat, light, humidity and air. Expired drugs pose toxicity risk as in some cases allow the active substance to undergo degradation leading to formation of toxic products. The administration of such drugs may lead to unsuccessful treatment of malaria and onset of complications such as cerebral malaria and anemia as a result of delayed appropriate treatment (Temu et al., 2006).

2.5 The rationale of the study

For there has not yet been any systematic research conducted as to the distribution and use of drugs at the community level, little is known about self-medication in the third world. Thus overall self-medication in modern pharmaceuticals seems to be a field in which information is scarce (Worku and G/Mariam, 2005).

Since studies on the prevalence of perceived malaria, the prevalence and factors associated with self-medication with anti-malarials, anti-malarial drugs hoarding and the extent of mosquito net coverage in Pawe Special Woreda are lacking, this study will fill the gap. In addition, this study will give insight about the commonly used traditional anti-malarial remedies in the study community.

It is expected that the findings generated from this study will contribute to knowledge and understanding of anti-malarial drug use and hoarding patterns at the community level, and also assesses mosquito net coverage which will be useful in developing interventions that will be undertaken to address their proper use.

3. OBJECTIVES OF THE STUDY

3.1. General Objective

To assess the utilization pattern of anti-malarial drugs, and mosquito net coverage in Pawe Special Woreda, Beneshangul-Gumuz National Regional State.

3.2. Specific Objectives

- To determine the prevalence of perceived malaria
- To determine the extent of anti-malarial drug use for self-care
- To determine the extent of anti-malarial drug hoarding at house hold level
- To assess mosquito net coverage and use

4. METHODOLOGY

4.1. Study area and population

Benishangul-Gumuz National Regional State is located in the North-Western part of Ethiopia, bordering the Sudan in the West, Amhara region in the North and North-East, and Oromia region in the South, South-East and East (JRM, 2004). The region has an estimated surface area of 49,289.46 km² and a population of 670,847 (CSA, 2007; CSA, 2008). Forty five percent of the population is below 15 years of age (JRM, 2004). Regarding ethnic composition: 25.9% are Berta, 21.3% Amhara, 21.1% Gumuz, 13.2% Oromo, 7.6% Shinasha, 4.2% Agew-Awi and the remaining 6.7% are from other different ethnic groups. 85.4% of the region's population lives in rural areas (CSA, 2008).

The study was conducted in Pawe special Woreda. Pawe is one of the 20 woredas in the Benishangul-Gumuz Region (CSA, 2008). It is bordered on the South and West by Metekel zone, and on the East and North by the Amhara Region (JRM, 2004). Based on figures from the 2007 Population and Housing Census, the woreda had a total population of 37,711, with male to female ratio of nearly 1:1 (CSA, 2008). The rural population accounts for 71.1% of the Woreda's total population (CSA, 2008). The Woreda is found at an altitude of 1050-1250m a.s.l and has mainly one climatic zone, *kola* and the average temperature is 32.2°C. Pawe is situated 573 kms North-West of Addis Ababa. Among the top ten diseases in the Woreda, malaria is the leading cause of morbidity and mortality. The Woreda is comprised of 20 kebeles which are subdivided in to 48 villages (*Gottes*); of which 3 are urban and the remaining are rural.

The woreda had fifteen governmental health institutions: one hospital, twelve clinics, one health center and one health post. In addition, the woreda comprises of the following private health institutions: two medium clinics, two drug stores, five rural drug vendors and two diagnostic laboratories. Twelve of the twenty kebeles had at least one health-extension workers (HEWs). As is true in many areas, there could also be a number of unregistered drug outlets in the district.

4.2. Study design

A community based cross-sectional household survey and qualitative study were conducted in late August and early September 2008 to collect data on perceived illnesses, perceived malaria illnesses, possession and usage of mosquito net, household anti-malarial drug stockpiling and, self-medication with anti-malarials and as well as the sources of treatment.

In many populations (in Africa), the wellbeing of the family devolves on the mother (Pilkington et al., 2004) and they are the primary care givers at home (Paulander et al., 2009). In view of this, data was collected primarily from mothers in both the quantitative and qualitative part of the study.

4.3. Sample size determination

The required sample size for the study was calculated using the formula for a single population proportion (Chadha, 2006):

$$n = \frac{Z_{\alpha/2}^2 \times p(1-p)}{d^2}$$

Since there is no available perceived malaria prevalence data for the study area, prevalence of perceived malaria from a study conducted in Adamitulu Jido Kombolcha Woreda of Oromia Region was used (Deressa, 2007^c). Assuming that 14.3% of the households would have at least one perceived malaria case over the 2-weeks recall period with a 5% margin of error at 95% confidence level (so $Z_{\alpha/2}=1.96$), the study required a minimum sample size of 433 households, considering a 15% non-response rate and a factor of 2 for the design effect.

$$n = \frac{(1.96)^2 \times 0.143 \times (1-0.143)}{(0.05)^2} = 188.3$$

Since the non-response rate is 15%; $0.15 \times 188.3=28.2$

Total sample size = $n + 15\%n = 188.3 + 28.2=216.5$

The overall total sample size becomes: $2 \times 216.5=433$.

4.4. Sampling procedures

The woreda is administratively organized into 48 villages (*Gottes*), 45 rural villages and 3 towns. Fifteen rural villages and 1 town were selected by simple random sampling technique using the lottery method. The recent registration lists of households was used to determine the number of households to be selected from each study sites. The share of households for each village (town) was divided by the total number of households in a given village (town) to determine a sampling interval and then households were selected by systematic random sampling (since Pawe was a settlement area during the *Derg* regime, the houses were arranged straight to the straight). To identify the first household in each site, bottle spinning technique was used.

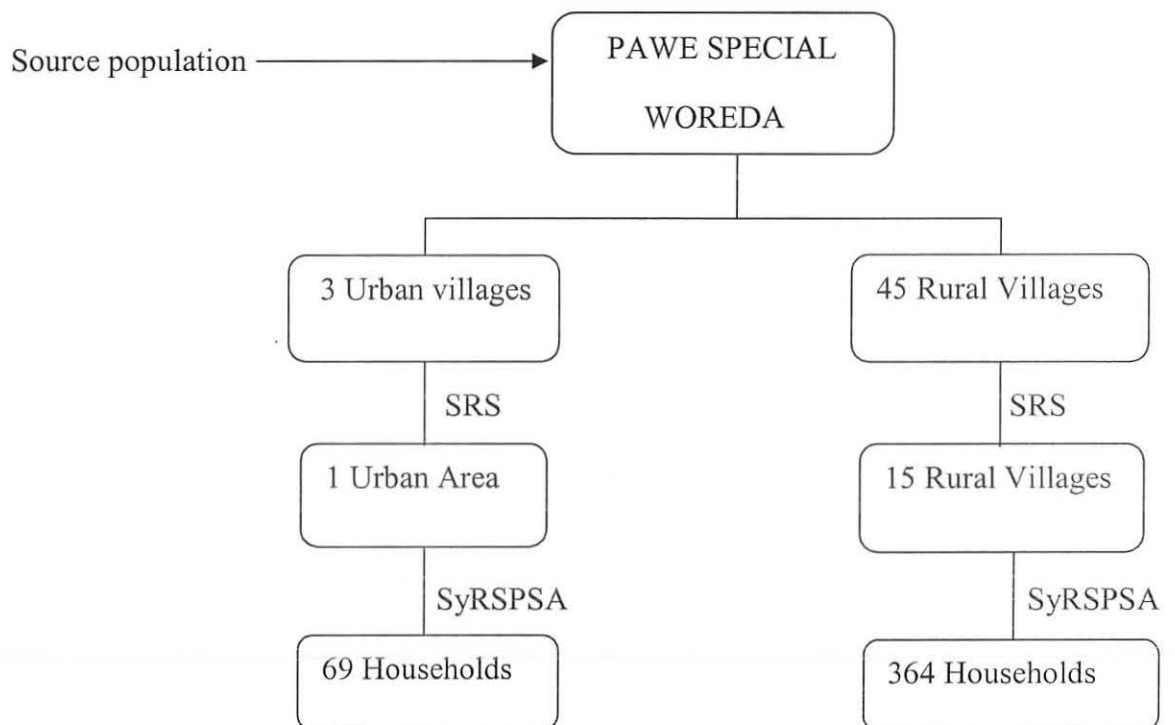


Figure 1: Schematic presentation of sampling procedure

SRS-simple random sampling

SyRSPSA-systematic random sampling, proportion to size allocation

4.5.1.1. Training and supervision

Ten enumerators who were 12th grade complete or university students and who speak Amharic were selected and trained for three days. Training was given by the principal investigator on methods of approaching, interviewing, recording and handling of the documents. The enumerators were monitored & assisted in the field by four trained supervisors and the principal investigator.

4.5.1.2. Pre-testing

The questionnaire was pre-tested for content and design in one of the neighbouring villages, called village 30, which was not selected for the study. Data collectors were exposed to practical situation before the start of actual data collection, and in the presence of principal investigator, the interviewers assessed clarity, understandability, flow and completeness of questions and the time needed to fill them. Finally, based on the pre-test, slight modifications were done.

4.5.2. Qualitative study

Qualitative research methods (key informant interviews and focused group discussions) were also employed in part as a primary data collection methods for some topics and in part for augmenting the findings of the quantitative survey and to elaborate more on the deep-rooted perceptions about malaria, anti-malarial drugs, self-medication, drug hoarding, and mosquito net possession and use etc. All the discussions of the qualitative study were held in Amharic.

The FGDs were conducted in a silent and comfortable place inside the villages, under a tree, at a time convenient to all of them and was held by the principal investigator. The three FGDs were conducted in three successive days and each was transcribed on the same day of gathering the information. Transcription of one FGD took about 2½ hours. Topic guides were developed based on the research questions to facilitate and keep the discussion on track.

The key-informant interviews were conducted at different quiet places within the villages thought to be conducive to the respondents. One key-informant interview took an average of 54.5 minutes and the transcription of the information collected was done on the same day of the data collection

and took about 1½ hour. In this case too, appropriate topic guides were developed to facilitate and keep the interview on track. All of the KIIs were handled by the principal investigator.

4.6. Study Variables

4.6.1. Independent

A. Socio-demographic variables such as age, sex, marital status, family size, occupation, religion, ethnicity, residence locality, & educational status etc.

B. Anti-malarial drug related variables

- Knowledge of: malaria symptoms, medication for malaria, the consequences of non-adherence to anti-malarial medication, advantages of obtaining drugs from legal sources etc.
- Knowledge about commonly used drugs for self-care

C. Knowledge about mosquito nets (benefit, utilization, prioritization in case of shortage etc.)

4.6.2. Dependent

- Perceived malaria
- Self-medication with anti-malarial drugs
- Anti-malarial drug hoarding
- Mosquito net possession
- Use of traditional medicines for malaria treatment

4.7. Data processing and analysis

After the collection of all the necessary data, it was coded on pre-arranged coding sheet. Quantitative data entry and analysis was done using EPI-Info Version 6.04 and SPSS version 11 statistical packages, respectively. Tables and graphs are used to present frequencies of pertinent findings. The association between the independent and dependent variables was measured and tested using Odds Ratio and 95 % CI. The relative contribution of each selected variables to the outcome of interest was assessed using logistic regression.

Kebele: the lowest administrative unit with a population of 1000 to 3000, further subdivided in to villages (*Gottes*).

Metata: a cheese well mixed with many spices such as *Zingiber officinale* (*Zingible*), *Alium sativum* (*nech shinkurt*), *Brassica nigra* (*senafitch*), *Lipidium sativum* (*feto*), *Ruta chalpensis* (*tena-adam*), *Aframomum melegueta* (*korerima*), *Coriandrum sativum* (*Coriander*) etc. and then kept for seven years in cool dry place.

Mich: a febrile illness which is believed to be caused by excessive sunlight and manifested by swelling and/or formation of sore on part the human body.

Mothers: a caregiver woman who is sixteen years of age and above and stays much of her time with and is responsible to give daily care for children and other members of the family in the sampled household was labeled as a mother and included as respondent in the study.

“Other sources” (of treatment): treatment of malaria/symptom complexes with modern anti-malarial drugs at home which may be obtained by the following means: left overs from a previous visit to a formal health provider, purchased and kept at home, borrowed/shared from relatives and/or neighbours etc.

Self-medication: taking modern drugs at home without any consultation with a health professional. It includes taking drugs bought from the retail sector (rural drug vendors, drug stores and pharmacies), and by using drugs obtained from “other sources” of treatment.

Village/Gotte: a small sub-component of kebele which usually consists of people very related in in one way or the other, and with a population of usually less than 1000.

Woreda: the lower level government administrative unit subdivided in to kebeles, and serves as the basic unit of planning.

5. RESULTS

5.1. Findings from cross sectional study

5.1.1. Socio-demographic characteristics of the respondents

Among the 433 visited households in 15 rural villages and 1 urban area, data was collected from 432(99.8%) households. With respect to residence, the majority, 363(84%) were from rural villages and the remaining 69(16%) were from urban areas.

In terms of marital status, 313(72.5%) of the respondents were married, 61(14.1%) were widowed, 36(8.3%) were divorced, and 22(5.1%) were single. The median age of the respondents was 25 years and the mean (\pm SD) age was 34.7(\pm 12.98) years.

As depicted in Table 1, 278(64.4%) of the respondents were unable to read and write, 76(17.6%) had attended class in the primary education, 37(8.5%) had gone through intermediate schools or above (Grades 7+), 33(7.6%) were able to read and write and 8(1.9%) were able to read only.

As far as household family size is concerned, 321(74.3%) households had a family size of less than or equal to 5 persons and 111(25.7%) had more than 5 persons. The average (\pm SD) family size was 4.5(\pm 1.85) people.

With regard to household monthly income, 95(22%) households had a monthly income of less than or equal to 112.5 ETB (Ethiopian birr), 86(19.9%) had from 112.5-300.0 ETB, 79(18.3%) had from 300.0-495.8 ETB, 86(19.9%) had from 495.8-901.7 ETB and 86(19.9%) had from 901.7-5062.5 ETB.

Most of the study participants, 336(77.8%) were from Amhara ethnic group followed by Kambata and Agew ethnic groups each comprising of 32(7.4%). Hadiya, Tigrie and Oromo constituted 18(4.2%), 7(1.6%) and 6(1.4%) of the total, respectively.

Concerning the religion of the respondents, 285(66.0%) were followers of Orthodox Christianity followed by Islam 96(22.2%). Protestant and Catholic constituted 44(10.2%) and 7(1.6%), respectively.

The farmers accounted more than three fourth of the study participants, 337(78%). This was followed by housewives 55(12.7%).

With regard to the educational status of the spouse, from the 313 married respondents, the husbands of 148(47.4%) were unable to read and write, while 62(19.9%) have attended class somewhere in the primary grades, 45(14.4%) had gone through intermediate or secondary school, 41(13.1%) were able to read and write, 12(3.8%) were able to read only, while only 3(0.96%) had attended post secondary school.

Table 1: Socio-demographic characteristics of respondents of cross-sectional survey, Pawe Special Woreda, North-West Ethiopia, August 2008(N=432).

Characteristics	Frequency	Percentage
Age Group		
<20	37	8.6
20-29	142	32.9
30-39	106	24.5
40-49	74	17.1
>49	73	16.9
Ethnicity		
Amhara	336	77.8
Kambata	32	7.4
Agew	32	7.4
Hadiya	18	4.2
Tigrie	7	1.6
Oromo	6	1.4
Marital status		
Married	313	72.5
Widowed	61	14.1
Divorced	36	8.3
Single	22	5.1
Religion		
Orthodox	285	66.0
Muslim	96	22.2
Protestant	44	10.2
Catholic	7	1.6
Educational status		
Illiterate	278	64.4
Read only	8	1.9
Read and write	33	7.6
Grades 1-6	76	17.6
Grades 7+	37	8.5
Occupation		
Farmer	337	78.0
House wife	55	12.7
Merchant	15	3.5
Government employee	8	1.9
Student	8	1.9
Daily labourer	7	1.6
Others	2	0.4
Family size		
≤5	321	74.3
>5	111	25.7
HH monthly income		
<112.5 ETB	95	22.0
112.5-300.0 ETB	86	19.9
300.0-495.8 ETB	79	18.3
495.8-901.7 ETB	86	19.9
901.7-5062.5 ETB	86	19.9

With regard to the possession of information sources (radio and TV) of households, 256(56.9%) had a radio, 12(2.8%) had a television.

5.1.2 Reported illnesses

The household survey covered a total population of 1943, residing in 432 households. 165(38.2%) of the respondents had at least one of their household members who had been ill in the past two weeks preceding the interview date. Accordingly, illness episodes were reported for 209(10.8%) individuals and some households reported more than one ill person. The average age (\pm SD) of persons with reported illnesses was 27.3(\pm 19.2) years with male to female ratio of 1:1.13. Concerning ill persons' status in the HH, Children 95(45.5%) predominated followed by mothers 68(32.5%) and husband 39(18.7%) (Table 2).

Table 2: Reported illnesses by different socio-demographic variables, Pawe Special Woreda, North-West Ethiopia, August, 2008(N=209).

Characteristics	Frequency	Percentage
Number of ill Persons Per HH		
One	134	81.2
Two	19	11.5
Three	11	6.7
Four	1	0.6
Sex of the ill persons		
Male	98	51.1
Female	111	48.9
Age of the ill persons		
<5	31	14.8
5-9	14	6.7
10-14	11	5.3
\geq 15	153	73.2
Ill persons' status in the HH		
"Child"	95	45.5
The mother	68	32.5
The Husband	39	18.7
Others	7	3.3

In those households where ill individuals were found, many illnesses/symptom complexes were reported, among which the top five were fever 128(61.2%) followed by malaria 119(56.9%), headache 107(51.2%), loss of appetite 82(39.2%) and chills 78(37.3%) (Table 3).

Table 3: Reported illnesses, Pawe Special Woreda, North-West Ethiopia, August 2008 (N=209).

Reported Illness	Frequency	Percentage*
Fever	128	61.2
Malaria	119	56.9
Head ache	107	51.2
Loss of appetite	82	39.2
Chills	78	37.3
Vomiting	63	30.1
Joint pain	58	27.8
Shivering	54	25.8
“Bird”	16	7.7
Spasm	14	6.7
Rheumatoid arthritis	14	6.7
Diarrhea	10	4.8
Typhoid	5	2.4
Asthmatic	4	1.9
Heart diseases	4	1.9
Colds & Coughing	4	1.9
Others	18	8.6

* Sum of percentages may exceed 100% because of multiple responses.

5.1.3 Perceived malaria illness

Malaria illness was reported by 119(6.1%) of individuals among 1943 people assessed during 14 days prior to the interview date. The proportion of reported malaria was higher for females 66(55.5%) than males 53(44.5%). Most of the reported malaria occurred in “children” 55(46.2%) followed by mothers 42(35.3%) and the husband 18(15.1%).

The mean (\pm SD) and median ages for the reported malaria cases were 25.3(\pm 18.7) and 22 years, respectively, with a range of 1 month–67 years. The majority of them were greater than or equal to 15 years old 84(70.6%), in comparison with those less than 15 years-old 35(29.4%) (Table 4).

Spring (*Tsedey*), followed by summer (*kiremt*), were recognized by 345(79.9%) and 95(22%) of the study participants, respectively, as the major seasons in which malaria transmission hits the highest point. Winter (*Bega*) and Autumn (*Belg*) were also mentioned by 13(3%) and 8(1.9%) respondents respectively.

Knowledge about signs and symptoms of malaria

As shown in table 5, knowledge about signs and symptoms of malaria was relatively high and almost all of the respondents, 430(99.5%), knew at least one of the common symptoms of malaria; namely, fever, chills & shivering, headache, vomiting, loss of appetite, thirst, muscle, back and joint pain.

Knowledge about malaria prevention

From a total of 432 respondents, only 380(88%) considered malaria as a preventable disease. Among these, bed net utilization, 325(75.2%) and draining areas of water nearby house, 278(64.4%) were mentioned by most of the respondents. Significant number of them also believed that blocking mosquito entry holes using local materials, smoking (by burning cow dung or leaves), closing windows & doors early, spraying insecticide inside the house as malaria prevention methods (Table 5).

Knowledge about the vulnerability to malaria

A large proportion of the respondents, 400(92.6%), believed that malaria could potentially attack people of all age groups and both sexes. However, very few respondents 31(7.2%) believed that certain groups are exempted (not vulnerable to malaria) and 1(0.2%) of the respondents did not know whether malaria could potentially attack people of all age group and sex. From those which perceived that certain groups are exempted, 21(67.7%) said children less than 15 years of age, 7(22.6%) said people of more than 65 years, 6(19.4%) said people within the age range of 15-65 years, 3(9.7%) said pregnant women and 4(12.9%), though they are sure that certain groups are exempted, they were unable to name the groups. Some respondents cited more than one “exempted groups.”

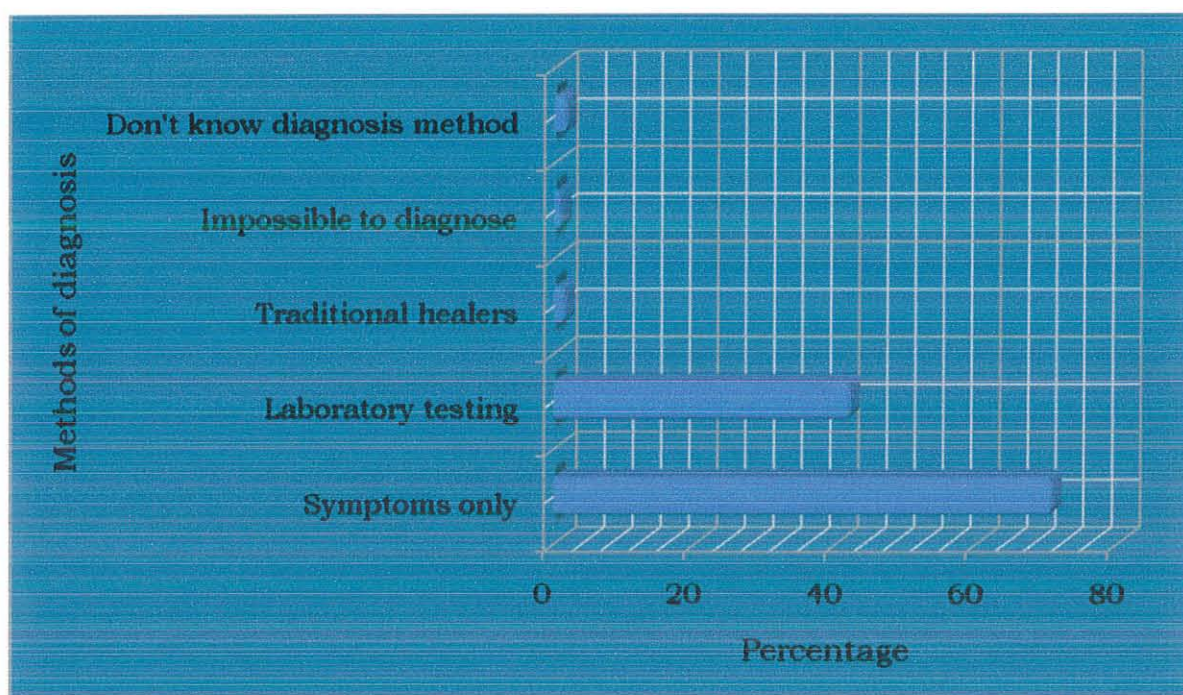
Table 5: Level of awareness about malaria: its cause, signs & symptoms, transmission and prevention, Pawe Special Woreda, North-West Ethiopia, August 2008 (N=432).

Characteristics	Frequency	Percentage*
Perceived Cause		
Mosquito bite	262	60.6
Lack of basic sanitation around houses	242	56.0
Lack of personal hygiene	149	34.5
Hunger	83	19.2
Cold weather	51	11.8
<i>Mich</i>	31	7.2
Cloudy weather	27	6.3
Excessive work load	7	1.6
Evil spirit	5	1.2
Do not know	17	3.9
Others	12	2.8
Symptoms		
Fever	334	77.3
Chills & Shivering	326	75.5
Headache	290	67.1
Vomiting	275	63.7
Loss appetite	187	43.3
Joint pain	85	19.7
Thirst	82	19.0
Back pain	79	18.3
Muscle pain	19	4.4
Nausea	10	2.3
Don't know	2	0.5
Others	24	5.7
Mode of malaria transmission (N=118)		
Bite of infective mosquitoes	83	70.3
Breathing	19	16.1
Sleeping together with the malaria patient	18	15.3
Physical contact with the malaria patient	14	11.9
Mother to child (placenta)	11	9.3
Sweat	7	5.9
By flies	6	5.1
Infected blood donation	5	4.3
Others	8	6.8
Malaria preventable		
Yes	380	88.0
No	52	12.0
Prevention methods (N=380)		
Mosquito net use	325	75.2
Drain areas of water nearby house	278	64.4
Closing holes with different materials	145	33.6
Smoke (burning cow dung or leaves)	126	29.2
Closing windows & doors early	89	20.6
Spraying insecticide inside the house	28	6.5
Keeping personal and environmental hygiene	14	3.2
Others	10	2.3

* Sum of percentages may exceed 100% because of multiple responses.

Knowledge about diagnosis of malaria

Regarding the diagnosis of malaria, 305(70.6%) of the respondents had the understanding that malaria could be diagnosed certainly by the signs and symptoms only, 181(41.9%) of the respondents pointed out the need of laboratory testing for diagnosis and a even few of the respondents 6(1.4%) said that malaria could only be diagnosed by experienced traditional healers. 7(1.6%) of the respondents, though they are sure that malaria can be diagnosed, they were unable to cite the diagnosis method. A very small, 4(0.9%) of the respondents said that it was impossible to diagnose malaria (Fig. 2).



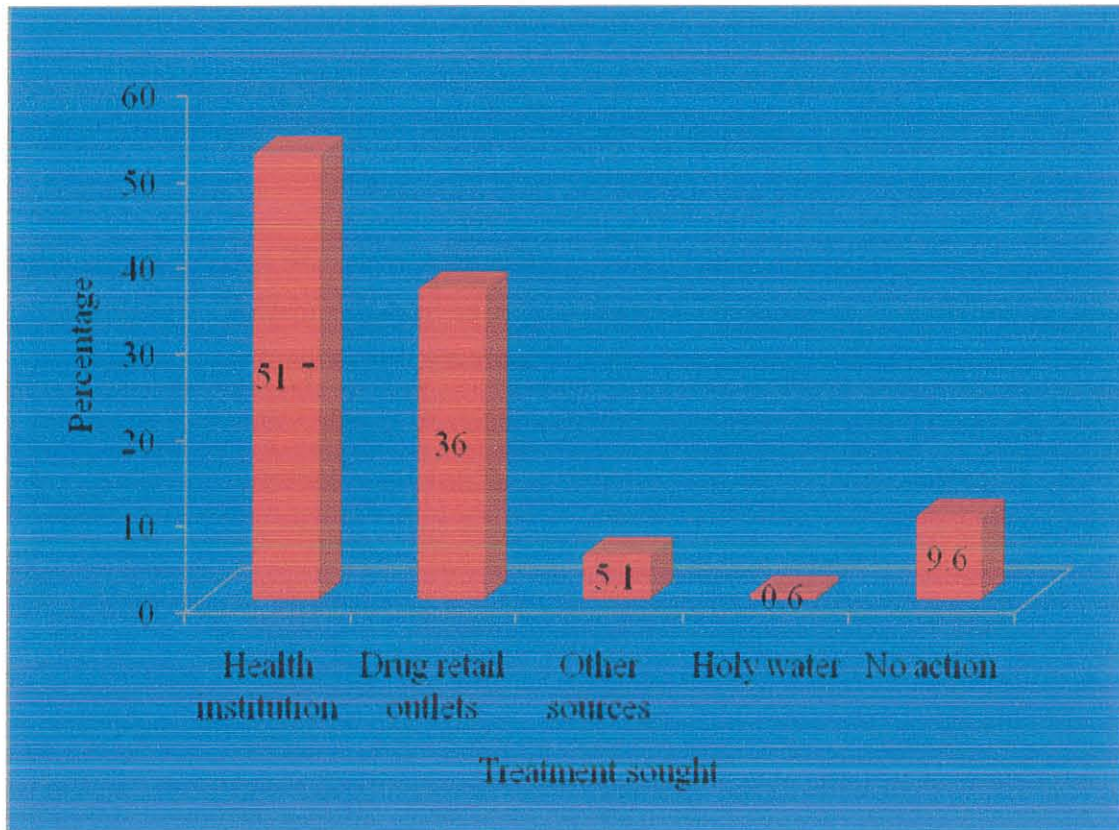
* Sum of percentages may exceed 100% because of multiple responses.

Figure 2: Knowledge of respondents about the diagnosis of malaria, Pawe Special Woreda, North-West Ethiopia, August 2008 (N = 432).

5.1.5 Actions taken against perceived malaria/symptom complexes

From a total of 209 reported illness episodes, only 178(85.2%) reported malaria illness and/or malaria symptom complexes. Out of the 178 persons which reported to have malaria and/or symptoms of malaria, 92(51.7%) went to health institutions, 64(36%) obtained drugs from drug retail outlets, 9(5.1%) obtained drugs from “other sources” and 1(0.6%) individual had used holy water. 17(9.6%) of the individuals were neither taken to health facilities nor received any form of

home treatment, the main reasons being thinking that the illness was minor. Some patients 5(2.8%) sought treatment from two sources (Fig. 3).



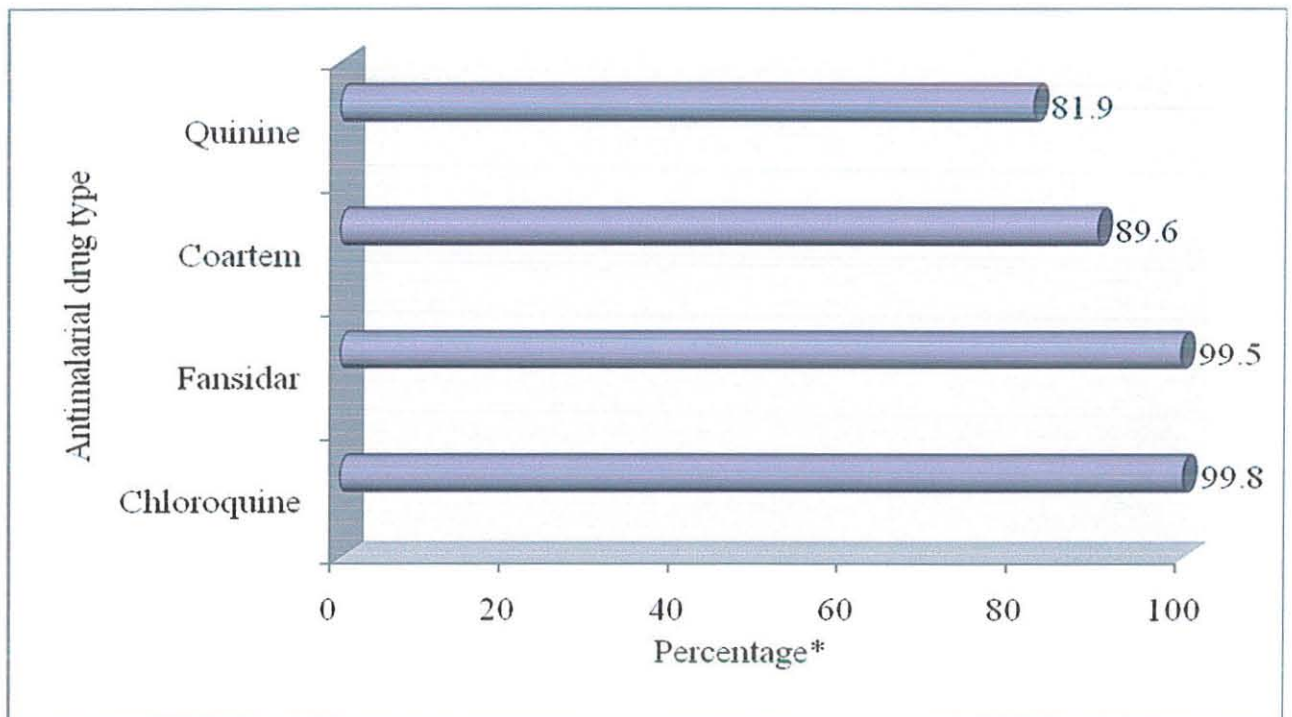
* Sum of percentages may exceed 100% because of multiple responses.

Figure 3: Actions taken against reported malaria/symptom complexes, Pawe special woreda, North-West Ethiopia, August 2008 (N = 178).

5.1.5.1 Self-medication practices

Out of 178 persons with reported malaria/related symptoms, 69(38.8%) self-medicated themselves with anti-malarial drugs obtained either from private drug retail outlets or from “other sources.” In short, out of the 209 ill people, 69(33%) self-medicated with anti-malarial drugs.

From the 69 reported malaria/symptom complexes that self-medicated themselves, 29(42%) took Chloroquine, 26(37.7%) took Coartem, 14(20.3%) took Fansidar and 7(10.1%) took Quinine. 10(15.6%) individuals had taken antibiotics along with anti-malarials (Table 6).



*sum of percentages may exceed 100% because of multiple responses.

Figure 5: Knowledge of respondents about the names of the different anti-malarial drugs, Pawe Special Woreda, North-West Ethiopia, August 2008(N=432).

Concerning the methods people employ to identify anti-malarial drugs from others, half of the interviewed mothers 216(50%) said by looking the color of the anti-malarial drug and/or packaging, 189(43.8%) of them said by reading the label or by asking others to read the label, 63(14.6%) said by making signs on the containers or packaging, 60(13.9%) said by storing in different places and 30(6.9%) said by smelling the drug. Four (0.9%) of the caretakers do not know how to differentiate/identify anti-malarial drugs from others.

Table 8: Different methods used to differentiate anti-malarial drugs from others, Pawe Special Woreda, North-West Ethiopia, August 2008 (N= 432).

Methods	Frequency	Percentage*
Color	216	50.0
Reading the label or by asking others to read	189	43.8
Making signs on the containers (packaging)	63	14.6
By storing them in different places	60	13.9
Smell (odour)	30	6.9
Do not know	4	0.9
Other methods	8	1.9

* Sum of percentages may exceed 100% because of multiple responses.

When asked about the consequences of non-adherence to anti-malarial treatment, more than half of the respondents, 229(53%) said not being cured from the illness (risk of relapse), 87(20.1%) said death follows, 86(19.9%) said severe worsening of the disease condition, 12(2.8%) said it leads to other diseases, 10(2.3%) did not know anything about the consequences of poor adherence to anti-malarial medication and 18(4.2%) mentioned other reasons. Some respondents mentioned more than one consequences of non-adherence.

Table 9: Reported consequences of non-adherence to anti-malarial medication, Pawe Special Woreda, August 2008 (N = 432).

Consequences	Frequency	Percentage*
Not being cured from the illness (relapse)	229	53.0
Death	87	20.1
Severe worsening of the disease condition	86	19.9
Leads to other diseases	12	2.8
Do not know	10	2.3
Others	18	4.2

*Sum of percentages may exceed 100% because of multiple responses

In this study, almost one-twentieth, 19(4.4%) of the respondents claimed that either the mother herself or any family member has consumed alcohol while on anti-malarial treatment.

container/package in open spaces and 1(3.3%) was stored in torn container/package in dry area, such as on the dry mud shelves.

5.1.8 Mosquito net coverage

The survey shows that 427(98.8%) respondents reported to have had at least one mosquito net in their household.

Regarding the number of mosquito nets they possess, 118(27.6%) had only one, 262(61.4%) had two, 41(9.6%) had three and only 6(1.4%) had four. Figure 7 shows the percentage of households along with the number of mosquito nets they own.

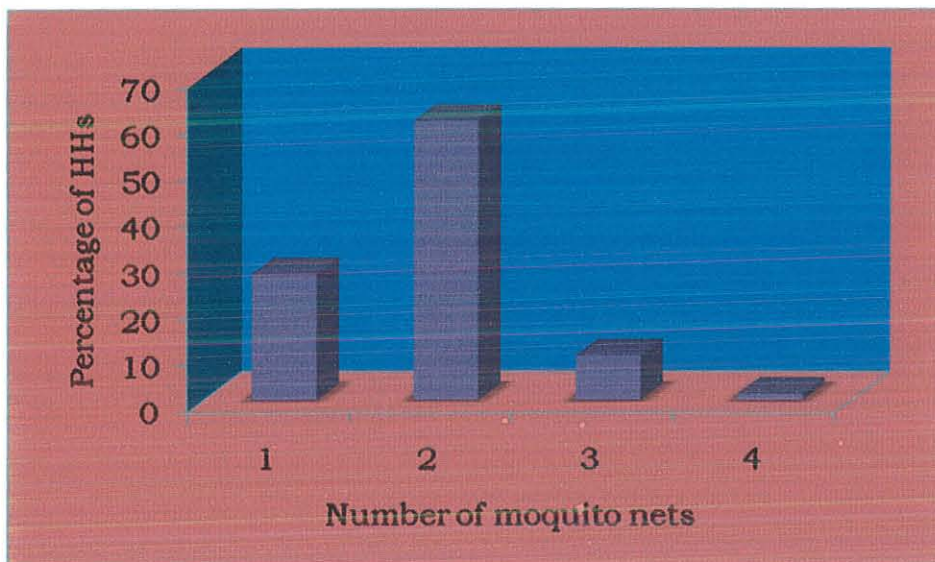


Figure 7: Mosquito net possession in Pawe Special Woreda, North-West Ethiopia, August 2008. (N=427)

Factors associated with reported possession of more than one mosquito net

In multivariate logistic regression analysis, among the socio-demographic variables, the mothers' (care givers') occupation and household family size were associated with reported possession of more than one mosquito net. Being a housewife (OR=2.96, CI=0.92, 9.59) was highly associated with possession of more than one mosquito net. Households with a family size of less than or equal to 5 persons (OR=0.05, CI=0.02, 0.17) were found to have a significantly less chance of possessing more than one mosquito net. Ethnicity, age, level of schooling, possession of

information sources (radio and television), respondents knowledge that mosquito net prevent from malaria, mosquitoes transmit malaria, and perceived that mosquito as cause of malaria, and residence place (urban or rural) were not associated with reported possession of more than one mosquito net (Table 10).

Table 10: Factors associated with possession of more than one reported mosquito net, Pawe Special Woreda, North-West Ethiopia, August 2008(N=427).

Variables	Possession of >1 mosquito net		COR (95% CI)	AOR (95% CI)
	Yes	No		
Education of the Mother				
Illiterate	198	77	1.45(0.70, 3.014)	1.46(0.62, 3.43)
Read and write/Read only	31	9	1.95(0.71, 5.33)	2.20(0.71, 6.85)
Grade 1-6	57	19	1.70(0.72, 4.00)	1.89(0.73, 4.93)
Grade 7+	23	13	1	1
Family size				
≤5	201	115	0.05(0.02, 0.16)**	0.05(0.02, 0.17)**
>5	108	3	1	1
HH Income				
<112.5 ETB	55	35	0.39(0.20, 0.76)*	0.54(0.25, 1.18)
112.5-300.0 ETB	58	28	0.51(0.25, 1.02)	0.65(0.30, 1.45)
300.0-495.8 ETB	62	17	0.90(0.42, 1.91)	0.99(0.43, 2.26)
495.8-901.7 ETB	65	21	0.76(0.37, 1.57)	1.88(0.40, 1.93)
901.7-5062.5 ETB	69	17	1	1
Radio				
Yes	185	60	1.44(0.94, 2.21)	1.17(0.72, 1.88)
No	124	58	1	1
Occupation of the mother				
Housewife	45	9	3.85(1.29, 11.46)*	2.96(0.92, 9.59)*
Merchant	10	5	1.54(0.40, 5.96)	1.19(0.26, 5.36)
Farmer	241	94	1.97(0.84, 4.65)	1.33(0.51, 3.49)
Others***	13	10	1	1
Respondents perceived that mosquito is a cause of malaria				
Yes	189	70	1.08(0.70, 1.67)	1.06(0.65, 1.74)
No	120	48	1	1
Respondents know that mosquitoes transmit malaria				
Yes	60	22	1.05(0.61, 1.81)	0.78(0.42, 1.45)
No	249	96	1	1
Respondents know that mosquito net prevents from malaria				
Yes	237	86	1.22(0.76, 1.99)	0.90(0.52, 1.54)
No	72	32	1	1

*P-value<0.05, **P-value<0.005

***Others include: civil servants, daily labourers and students

5.1.9 Use of traditional remedies to treat malaria

By interviewing mothers, information was collected on the different traditional remedies used to treat malaria. It was found that 23(5.3%) of the visited households used traditional remedies to treat malaria. The route of administration mentioned for all the traditional drugs was oral (100%) and in formulating liquid preparations, water was reported to be used as a solvent.

Medicinal Plants used to treat malaria were reported in 19(4.4%) of the households studied and the most commonly mentioned herbs were, papaya (*Carica papaya*), neem (*Azadirachta indica*), zingible (*Zingiber officinale*), damakesse (*Ocimum lamifolium*), bisana (*Croton macrostachys*), garlic (*Alium sativum*), lemon (*Citrus limon*), pepper (*capsicum annum*), senafitch (*Brassica nigra*). From these, the most commonly mentioned were *Carica papaya* and *Azadirachta indica*. The doses and duration of administration of herbs varied from household to household.

Animal origin traditional remedies were reported in 4(0.93%) of the households studied. All four of the households mentioned “*Metata*”, a cheese well mixed with different spice species, kept for seven years and then will become a hard solid mass. Though there is difference regarding the dose, there is no difference regarding the duration of treatment. All the four respondents said the liquid dosage of “*metata*” is prepared by combining a part of it with water and taken for seven days.

5.2 Qualitative results

5.2.1 Findings of FGDs with mothers

Three FGDs, with an average 78 minutes of interesting discussion with each group, one group comprising of seven women and the other two groups comprising of eight & nine women were conducted. The FGDs were conducted with different women who were thought to represent the whole women community: rural and urban residents, young and old, from all ethnic groups and considering all religions practiced in the study area.



Figure 8: One of the FGDs sessions, Pawe Special Woreda, North-West Ethiopia, September 2008.

All participants were willing to participate in the discussion to share their views and also to be tape-recorded. The average age of participants was 39.8 years (range 25-65 years). Almost all

were housewives, married and regarding their educational status, majority of the participants were illiterate (not able to read and write), some of them were able to read and write, and a very few of them attended grades 1-8.

Results the FGDs were analyzed by thematic approach. The transcriptions of the FGDs were reviewed to classify the primary categories within the themes. Finally, four themes were selected for presentation of findings. These include knowledge about malaria, actions taken against perceived malaria/symptom complexes, knowledge about malaria treatment and anti-malarial drugs, and mosquito net possession and use.

Knowledge about malaria: cause, transmission, signs & symptoms, prevention and diagnosis

Though the extent of emphasis among discussants varied, all of the women involved in FGDs knew malaria called “*weba*” by the local language and all recognized malaria as a major health problem in the study area followed by typhoid and tuberculosis (TB). The perception was that nearly everyone in the community has already experienced malaria episodes. One of the FGD participants said:

“Here in Pawe we are husband and wife with malaria” i.e. to show the severity and endemic nature of malaria in the study area.

There were interesting variations in the perception about the cause of malaria within the FGD participants. Some put the cause of malaria as something that follows when people fail to keep their personal and environmental hygiene, like lack of personal cleanliness, formation of swampy areas and accumulation of dirt around houses etc. Others said eating food contaminated by “*tinign*” as the cause of malaria: a participant in one of the FGDs stated that:

“When immature children eat food contaminated with “tinign”, they will be infected by malaria. Adults will not eat such foods as they are very aware of the consequences!”

Most participants understood malaria as a diseases caused by “*bimbii*” (group of night time flying insects reproduced and lived in garbage and swampy areas). When a “*bimbii*” bites people, it causes the disease. Asked if all “*bimbiis*” cause malaria, the FGD discussants said no. As one mother said:

“The vomit due to malaria is yellowish-green in color but the vomit due to other diseases, for example due to typhoid is white and full of froth but there is no froth in malaria vomit.”

Most participants reported that ‘farcifarum’ (=falciparum) malaria is very dangerous and leads to unconsciousness if treatment is not sought immediately. One participant narrated that:

“There is continuous diarrhea and vomiting in ‘good’ types of malaria illnesses but the one without such symptoms is a very dangerous type of malaria called farcibarum (=falciparum), it immediately rises to your head and kills you within a short period of time.”

The study revealed that the Pawe community largely believed malaria is preventable. And mosquito nets, draining stagnant water around home, keeping personal and environmental hygiene were mentioned as important methods of prevention.

The majority of the FGD participants showed their preference to bed nets instead of insecticidal aerosols for mosquito bite prevention. A young discussant stated that:

“The latter is used for shorter period of time and cost wise it needs ‘huge’ investment, but the former can be used for a year or more.”

All of the FGD participants mentioned chemical spraying (i.e. IRS) as a means of malaria prevention. However, some participants expressed their discontent with this method as it is tedious. One participant elaborated that:

“It is very beneficial since it kills almost every small insect including mosquitoes but not let my house to be sprayed as it is tiresome to move in and out household utensils/materials.”

Some of the participants in all the FGDs underlined eating better foods as an important method of malaria prevention and they justified this by saying it is mainly the poor that usually get attacked by malaria since they do not get quality food. One woman stated that:

Self-medication can be seen as a first step in the attempt to solve a health problem. It is mainly after self-medication failed to solve the perceived malaria/symptom complex illness that an ill individual is taken to the health care facility. One participant from the urban area admitted that:

“If any of my family members is febrile or has malaria illness, I will buy Chloroquine from the private pharmacy for the febrile or malaria ill individual and Paracetamol if there is head ache. If improvement is not seen, I will take the individual to the hospital.”

In relation to non-compliance with a clinician’s prescription, a negative assessment of a drug’s efficacy is the most frequently cited reason for deciding to abandon treatment and to start what they believed to be effective. One of the FGD partakers described:

“The nurse in our clinic prescribed Coartem, and I took them, but I didn’t get any better. Then I bought and took Quinine that cured me soon.”

Regarding hoarding of anti-malarials most of the FGD participants did not hoard, but some said they stockpiled anti-malarials for different reasons. Frequent episodes of illness, to save time (for plowing or harvesting), distance from the source of anti-malarial drugs etc. were given as the main reason for hoarding of anti-malarials at household. Some respondents relied on the fact that they had previous experience with similar ailments therefore giving them the confidence to self-medicate. A woman stated that:

“When I and my family members got sick, I used to give them anti-malarial drugs bought and kept at home without going to the clinic which is situated around three-four kilometers away.” And another participant further elaborated this as: *“Instead of going to the town during farming or harvesting time, we purchase and keep the drugs in case someone becomes ill.”*

Knowledge about malaria treatment and anti-malarial drugs

FGD participants were asked to mention the commonly used anti-malarial drugs. Accordingly, almost all the participants were able to mention Chloroquine, Fansidar, Coartem and Quinine. In order to assess whether they could be able to identify the different anti-malarial drugs, samples of the packages of Coartem, Chloroquine, Fansidar and Quinine were shown to the participants.

Most of the participants recognized Coartem, Chloroquine & Fansidar and some identified Quinine.

Moreover, they said that they are accustomed to the drugs very well because they are using them in everyday life for them/their family members and identified each, as they said, by color of the tablets, packing design and special illustrations/figures on the pack. When they were showed to identify a pack of tablet (Coartem) and asked for what purpose it is used, one of the participants said:

“Really it is curtum(=Coartem) taken for the treatment of malaria; because of the picture of a child on the pack, the yellow color of the tablets and the special arrangement of the tablets on the pack.”

Though many FGD participants were able to name, and identify when shown the currently used first line anti-malarial drug for falciparum malaria-Coartem, they were unable to comprehend how to administer the drug by just studying the drawing on the pack.

One FGD participant when showed to identify a blister pack of tablet (Fansidar) and to tell for what purpose it is used, she stated:

“Bring it! May I see it closely...oh! It is Fansidar, a drug used for malaria treatment! Because mosquito, which is the cause of malaria, is drawn on the pack.”

For most case in the first line of malaria treatment, anti-malarial drug is given twice a day for three consecutive days (Coartem) and once per day for three consecutive days (Chloroquine) as per Ethiopian malaria treatment guideline. Having this idea in mind, the respondents were asked about how many times the drugs were taken per day and for how many days in the first line of malaria treatment.

In all FGDs, majority of the participants know the correct adult dose of Chloroquine and Fansidar and almost all recognized:

“Chloroquine once per day for three consecutive days, Fansidar all the three tablets at once.”

However, most of the participants did not know the appropriate dose of the newly introduced anti-malarial drug-Coartem: some say four tablets per day for six days, others say four tablets twice per day for three days etc. An old mother in one of the FGDs elaborated this as:

“I was given Coartem before a week; I took four tablets (all once) per day for six days.”

Almost all the participants explained that Coartem was very effective and that they were satisfied with it. A mother was particularly impressed with the rapid improvement observed in the health of the ill following administration of the Coartem and she said:

“Whenever sick persons are given the drug, the persons recover promptly and by the second day of use many of the sick persons were active and healthy.”

Although majority of the participants have acknowledged Coartem, few have doubts over its efficacy. Even though Fansidar is prohibited for the treatment of malaria based on the country’s malaria treatment guideline, some of the FGD participants claimed to use it for the treatment of malaria. Their main reason was lack of confidence in the efficacy of the newly introduced drug, Coartem-from their experience. Since it is not prescribed for malaria treatment in government institutions, they claim to get it by purchasing from private drug retail outlets. One mother said:

“My son was ill before some days & I took him to the clinic in our kebele & was given coartem but did not show improvement. I went to the nearby town and bought Fansidar and Chloroquine from the drug retail outlets and felt better within two days of treatment initiation.”

Concerning the perception about the efficacy of anti-malarials, some participants associate efficacy with taste: bitter or sour or other non-sweet tasting drugs were perceived to be more effective than other medications. One mother lamented that:

“Coartem is not effective, even it is not bitter and if so how do I expect it to cure my malaria illness?!”

When asked about the most effective anti-malarial drug for the treatment of *P.falciparum* malaria, almost all of the FGD participants said Quinine. One mother put it as:

“Though it is dangerous for body, Quinine is the most effective drug for treating malaria.”

A majority of the FGD participants reflected the concept that taking alcohol while on anti-malarial treatment as a dangerous act which may lead to treatment failure and even death. However, a very few of the participants had the opposite belief, i.e. it leads to rapid cure and serves as a long term prophylactic mechanism against malaria infection. One elderly mother who participated in one of the FGDs said:

“I was ill because of malaria three years back. My daughter bought Chloroquine and Fansidar from the nearby private drug shop. I took the first dose in the morning and I drunk much alcohol at the evening and vomited a lot. I have never been infected for the past three years till the first week of the previous month.”

Reasons given by most of the FGD partakers for non-adhering to anti-malarial treatment regimens include: feeling recovered, failure to improve in expected time, feeling worse, forgetfulness, unpleasant side effects (especially vomiting), unclear instructions from drug providers, sharing of drugs at home etc. One participant stated that:

“After taking three or four doses, I would feel better and start going to work and I forget taking the drug. Even if I take it, I may take it after missing one or more doses. Sometimes, the left over drugs from my treatment will be kept in case my children get sick.”

Asked if they purchase drugs from ordinary groceries, almost all of the FGD participants said no and their reasons also vary. Some participants said they did not purchase drugs from shops because the drugs might be expired, and others said though drugs are available in few shops, drugs from shops are costly. One participant strongly expressed that:

“How can I purchase a drug from a shop? The shopkeeper may be selling drugs which are expired & discarded by somebody/health institutions; I know medicines are different

6. DISCUSSION

The study was population based and the aim was to determine the prevalence of perceived malaria, treatment-seeking behaviour for reported malaria illness/symptom complexes and about anti-malarial drug hoarding at household level in an area of seasonal malaria transmission in rural Ethiopia. Furthermore, it aimed to determine the coverage and use of mosquito nets in the area, and factors influencing its possession and usage. The study discovered essential information about factors associated with self-medication with anti-malarials, anti-malarial drug hoarding and possession of mosquito nets among the study population.

A number of malaria-related symptoms particularly fever, shivering, chills, loss of appetite, headache, vomiting and joint pain were considered and presented as symptom complexes, which taken together may approximate a clinical diagnosis of malaria, although using these symptoms, particularly fever, as a proxy for malaria appears to be neither sensitive nor specific when compared to parasitologically confirmed diagnosis (Font et al., 2001; Deressa et al., 2007^a). Nevertheless, fever along with other symptom complexes serves as a proxy for malaria both at household level for home management of malaria and peripheral health facilities where diagnosis is made presumptively upon the presence or history of fever and other symptom complexes. However, a recent change of first-line treatment of malaria to artemisinin-based combination therapies (ACTs) in many countries of SSA has highlighted the potential cost implications of malaria over diagnosis based upon clinical signs and symptoms. Use of a rapid diagnostic test would help in identifying malaria parasites especially in areas with seasonal malaria transmission where presumptive diagnosis of the disease may be inaccurate but the magnitude of asymptomatic carriers is assumed to be low (Deressa et al., 2007^a).

In the present study area, laboratory tests were unavailable at peripheral public health facilities such as clinics and health posts. At health center and hospital levels, malaria diagnosis through blood smear examination is usually recommended, but there are occasions that these facilities might not provide blood testing services due to patient overload or shortage of supplies.

The present study revealed that the prevalence of perceived malaria was 6.1%, which was lower than the report from Adamitulu Jido Kombolcha Woreda in central Ethiopia and other studies done in Kenya and Papa New Guinea (Deressa et al., 2007^c; Yé et al., 2007; Pluess et al., 2009). The main reasons for this difference may be the fact that the present study was conducted on the eve of a high malaria transmission season in the area, and its sensitivity might not be high considering the transmission season. In addition to this, nowadays the Ethiopian government is focusing on prevention of malaria. Because of this, there is huge distribution of mosquito nets, relatively better access to IRS in comparison with olden days and better environmental hygiene keeping practices (e.g. removal of mosquito breeding sites). All these may partly explain the decrease in the reported malaria illness. Moreover, the variation in the study areas and methodology might partly explain the disparity in the reported rates of malaria illness episodes.

The decline in perceived malaria is also substantiated by a decline in the number of malaria cases year by year in Pawe as evidenced by reports from the woreda. The number of malaria cases in Pawe were 44094, 27162, 23985 and 15063 in 2004/05, 2005/06, 2006/07 and 2007/08 respectively (unpublished report from Pawe Special Woreda Health Office). This vividly indicates that there is a step decline in the rate of malaria infection during the past four years. Though the impact of malaria is declining, it is still the number one cause of morbidity in the study area in each of the respective four years. The decline in the malaria infection trend was also explained by FGDs & KIIs participants.

In the present study, higher number malaria episodes were reported in females than in males. This might be due to the fact that mostly in Ethiopian culture, particularly in such remote and undeveloped" regions, males are favored in most aspects and in this case too, males in general and male children in particular, may be given priority to use the different malaria prevention methods, including mosquito nets. Moreover, females may not get appropriate treatment on time in comparison with males. These factors all may have contributed their effect for the higher number of reported malaria cases in females.

More malaria cases were also reported in "children" than in mothers and husbands. This might be due to the fact that since there is report of mosquito net shortage and priority is given for

presented evidence on the types of anti-malarials used and sources of treatment for reported malaria illness/symptom complexes in an area of seasonal transmission in rural Ethiopia. In this study, from those persons who reported malaria illnesses/symptom complexes, 69(38.8%) practiced self-care with anti-malarials. Because of the general incomparability of surveys due to differences in definitions, methodology and study settings of different studies, and due to scarcity of documented data for self-medication with anti-malarials for perceived malaria illnesses/symptom complexes, it was difficult to make a comparative analysis with the results of other surveys conducted in Ethiopia or elsewhere.

Individual health-seeking behaviour is driven by perceptions about illness and so does the choice of treatment. With self-reported malaria, one would expect a high use of anti-malarial drugs both in self-treatment and at health facilities, which if wrongly treated may in turn create unnecessary drug pressure leading to resistance (Ye et al., 2008). In the present study most malaria ill people self-cared with Chloroquine 29(42%) and 26(37.7%) Coartem. This trend might be due to what they have observed from the health institutions in their locality i.e. from previous experiences in their illness. These two drugs are commonly administered together based on sign and symptom diagnosis in the peripheral health institutions as is stated in the Ethiopian malaria treatment guideline (MOH, 2004). In Pawe, as evidenced by most FGD and KII participants, some people did not follow the appropriate dosing system while taking anti-malarials. This inappropriate use will result not only in treatment failures, but also it theoretically increases the risk of drug resistance to the currently available effective anti-malarials like Coartem. Many studies done have shown that artemether–lumefantrine (Coartem) is the most effective of the anti-malarial compounds in shortening the fever and malaria parasite clearance times. However, several treatment failures have been reported. Even in those appropriately adhered patients, treatment failures have been reported as is the case in one Japanese patient and these failures are believed to be a consequence of poor bioavailability of the lumefantrine component when ingested without fatty food (Mizuno et al., 2009). In Ethiopia in general and in Pawe in particular, the residents are poor and unable to get enough fatty food while on artemether–lumefantrine treatment. If this food shortage is further enhanced by poor treatment adherence, the risk of treatment failure and anti-malarial drug resistance will be further aggravated.

Quinine was also used by few patients 7(10.1%) for self-medication purpose. Since Quinine is taken for seven to ten days, there is high chance that the patient may not complete the treatment regimen leading to treatment failure and even loss of life and drug resistance. This also puts warning on loss of important drugs used for severe malaria treatment. Though Fansidar is prohibited for malaria treatment in Ethiopia (MOH, 2004), people in the study area still use the drug and in this study 14(20.3%) of those who self-medicated took Fansidar. As evidenced by most FGD and KII participants, the main reason they cite for using Fansidar is loss of confidence in the efficacy of Coartem. To the knowledge of the principal investigator, in Ethiopia there is no documented data about the development of malaria parasite resistance to Coartem. Probably, the treatment failures reported might be due to lack of fatty food while on Coartem treatment and problems associated with adherence. People must be informed about fatty food issue while drug dispensing and health professionals must be trained to do so.

Household family size was associated with self-medication practice in the present study. In comparison with households which have a family size of more than 5 persons, households with family size of less than or equal to 5 persons were found to be at lower risk for self-medication with anti-malarials. Although the interpretation is difficult, potential explanations of the finding might be economic factors. The fact that households with larger family size may not have sufficient amount of money even to cover clothing and food consumption costs, such households will not have the capacity to cover transportation and other costs to take each ill individual to health institutions. Considering such problems, they instead buy and keep drugs in their home in case a family member becomes ill, or they may have leftovers from previous treatment etc. This in turn opens the door for self-medication. In contrary to this, most households with smaller family size will have the capacity to cover not only clothing and food consumption costs but also transportation and other costs to take each ill individual to health institutions. Because of this, such households will have less chance of self-medication.

Ability to meet the costs associated with a particular treatment is an important factor influencing access and choice of treatment (Chuma et al., 2008). Even where malaria treatment is supposedly free (i.e., in government health facilities) as is in Pawe, individuals still incur costs, such as

transport and laboratory services. In agricultural communities, the seasonality of cash income may prevent households from seeking treatment on time. Households make a significant amount of their annual income in the busy wet season, and people are unlikely to take time off to seek treatment for themselves or to accompany an ill person to a health facility, unless an episode is perceived to be serious enough (Chuma et al., 2008). And this leads people to practice self-medication either by buying drugs or by using left over or drugs obtained by other means. This practice was particularly stressed and reported by some FGD partakers.

In developing countries, self-medication should be considered a public health problem owing to lack of appropriate medical education to patients. Although a significant number of people throughout the world practice self-medication, only very low proportion get information about medicines from sources in the community because very little appropriate information is available at this level (Abdo-Rabbo, 2003). In the present study area, as reported by FGD and KII participants, under dosing and over dosing were common which may lead to development of drug resistance. Therefore, mere knowledge of anti-malarial drug is not a guarantee by itself unless the knowledge is internalized and put in to practice correctly by caregivers and the patients. It is speculated that internalizing anti-malarial drug related knowledge predicts rational use.

Different modalities of self-medication were mentioned by most FGD and KII participants: consumption of medicines acquired without a professional prescription, shared use of medicines, non-compliance with professional prescriptions, and use of leftover medicines stored at home. Similar results were also reported by Filho et al., (2004).

Seeking care from the health facility mainly comes after the failure of care at home, or malaria patients might directly seek treatment from health care facilities without the initiation of treatment at home (Deressa et al., 2007^a). In the present study, from those who reported malaria/symptom complexes, 92 sought treatments from health institutions and from these only 71(77.2%) were diagnosed as malaria. The remaining were diagnosed as typhoid, *birdd*, hypertension and others. Most of these diseases share one or more common symptoms with malaria. Imagine the over usage of anti-malarial drugs that may result if these patients were self-diagnosed and self-medicated with anti-malarials. From the total of 119 malaria/symptom

complex ill persons, some (69 persons) were self-medicated with anti-malarials and if these patients were diagnosed at health facility, parts of their illnesses might have been diagnosed as other diseases. Because of this evidence, it is speculated that there is irrational usage of anti-malarials among the study community, which will lead to loss of life and emergence of anti-malarial drug resistance.

Over the past two decades, there has been an increase in public awareness of the existence of counterfeit and substandard drugs. The potential risk of counterfeit anti-infectious agents for individual and community health includes clinical aggravation leading to complications and even mortality from either the disease itself or possible toxic components in the product, increased health expenditures to attain cure, and selection of drug-resistant bacteria and parasites (Basco, 2004). In the present study area, the main sources of anti-malarials for self-medication were private drug retail outlets. Key-informants working in government health institutions claimed to come across expired drugs dispensed from private drug retail outlets. Some of the FGD partakers who practiced self-care also reflected the same idea. Because of this evidence, while they obtain drugs from private drug retail outlets, the accessed drugs might be counterfeit and/or substandard which in turn jeopardizes the health of the patient. In view of this, the worst side of self-medication practice in Pawe is manifested not only by inappropriate use of anti-malarials but also by the huge risk of counterfeit and substandard drugs.

Implementation of pharmaceutical care in community pharmacies could help alleviate problems associated with rampant self-medication. Community pharmacists can play an active role in the provision of primary health care by attending to minor ailments and refer patients to physicians where patients require further investigation (Awad et al., 2005).

Regarding the knowledge of respondents about the names of the currently used different anti-malarials, Chloroquine 431(99.8%) and Fansidar 430(99.5%) were the two commonly known anti-malarial drugs and this is higher than what has been reported in a study done in North-Western Ethiopia by Tilaye and Deressa (2007). In the present study, the least known (named) anti-malarial, Quinine, was named by 354(81.9%) respondents still which is higher than a result

reported from Tigray (Habtesion, 2007). Since malaria is very endemic in Pawe and nearly everyone had experienced the disease and has taken the drugs, it is natural to expect such a high percentage of family caretakers to know or mention the names of anti-malarials that they knew and use. In most FGDs conducted in this study, participants stressed that they are well acquainted with anti-malarials as they are using them in everyday life for themselves and/or their family members.

Concerning the methods people employ to identify/differentiate anti-malarial drugs from others, half of the interviewed mothers 216(50%) said by looking the color of the anti-malarial drug and/or packaging from experience. Unless appropriate interventions are designed, since different tablets used for different ailments have almost related (pack) colour, patients may take drugs other than anti-malarials which will lead to severe disease complications and even death. Surprisingly, 30(6.9%) caretakers claim to use smell as a means of differentiating anti-malarials from others. Smell is really a very unspecific method and may again lead to drug misuse.

Though most of the respondents, 412(95.3%), are knowledgeable in one way or the other about the consequences of non-adherence to anti-malarials i.e. not being cured from the illness (relapse), death, severe worsening of the disease condition or leads to other diseases (e.g. anemia, splenomegaly etc.), some did not have the knowledge or have unrelated beliefs about the consequences of poor-adherence to anti-malarials which may aggravate the problem. FGD participants also reflected the same beliefs and mentioned the above consequences. The respondents understanding about the consequences of poor adherence to anti-malarials is promising. It is speculated that good knowledge of the consequences of poor-adherence to anti-malarials is related with rational use.

Many medications including anti-malarials can interact with alcohol, leading to increased risk of illness, injury, or death (AADAC, 2003). Individuals who drink alcoholic beverages should be aware that simultaneous use of alcohol and medications--both prescribed and over-the-counter--has the potential to cause problems (AHI, 2007). In this study, almost one-twentieth, 4.4%, of the respondents claim that either the mother herself or any family member has consumed alcohol

while on anti-malarial treatment. A few FGD participants also reported that drinking alcohol while on anti-malarial treatment to have a long term prophylactic effect. Unless immediate interventions are implemented, such misconceptions are dangerous as they lead to treatment failure and loss of life.

In remote areas of Ethiopia, like Pawe, private pharmacies, drug stores and rural drug vendors dispense anti-malarials over the counter without prescription. This condition favours people to buy and keep drugs at home. It is doubtful if the aforementioned drug sources give the appropriate dose, meaning that patients either do not take a required course, which leads to drug resistance or might even be overdosed resulting in unwanted toxic effects.

In the present study, it was found out that 8.3% of household hoarded anti-malarial drugs. This is similar with a study done in rural Tanzania, Bagamoyo by Temu et al., (2006). However, the present study result differs significantly from the study done in Dar es Salaam by Mnyika et al., (1995), where as much as 71.7% of households were reported to have been storing and self-caring with anti-malarials in their homes. This difference in findings may be attributed to the fact that Mnyika did the study in urban area while this study was done in remote rural area. This would mean that people living in urban areas have much more access to the drug sources, purchasing power and education level is high than those residing in rural areas.

Unlike the findings of Temu et al., (2006) where Amodiaquine and Quinine were the most stored anti-malarial drugs, in the present study it was noted that Chloroquine and Coartem were the most commonly stored anti-malarials, while Quinine & Fansidar were the least stocked. This difference in the kinds of drugs stored might be attributed to the policy difference in Tanzania and Ethiopia, since Amodiaquine is not approved to be used and marketed in Ethiopia, as per the malaria treatment guideline. Coartem, relatively being expensive hence reducing the probability of storage at home in comparison with Chloroquine. To a lesser extent policy change from Fansidar as first line drug for malaria treatment (since 2004) to Coartem could have played a role in the observed decline in the amount of Fansidar stored, as people may be told in the health institutions about this policy change issue. Moreover, in comparison with Chloroquine or

Quinine, Fansidar is taken as a single dose and as a result people will prefer to take Fansidar instead of a three day treatment course of Chloroquine or a seven to ten day Quinine, thus creating a room for not finishing the treatment course; as a result these drugs were remaining and stored at home. Despite being forbidden for the treatment of malaria, Fansidar was still stockpiled and used as anti-malarial. Some people have the perception that Fansidar is more efficacious than Coartem for *P. falciparum*, as explained in the FGDs and KIIs.

With respect to different places where anti-malarials were stored in the households, it was revealed that most drugs were stored in hot areas. Since heat aggravates the degradation of drugs in to useless or some times to toxic products, these drugs if taken may not cure the patient or even may cause hazardous consequences, including death and drug resistance. Unless such issues are closely monitored, they may lead to loss of the “golden” anti-malarial drugs.

Ethiopia has adopted the use of ITNs as one of its vector control strategies (Jimma et al., 2005). To this effect level of net ownership was assessed in this study area. Overall net ownership was very high with almost all, (98.8%) of the care givers' in the surveyed households had self-reported to have at least one bed net, and most owning more than one. The total number of nets owned by households, as reported by household respondents was 789 – an average of 1.8 nets per net-owning household. At least one net ownership is better when compared with other studies done in Ethiopia (Jimma et al., 2005; Tilaye and Deressa, 2007; Legesse et al., 2007; Paulander et al., 2009). At least one net ownership per household was also found to be higher in the present study area when compared with results from studies done in other African countries such as Gambia, Senegal, Zambia and Nigeria (Wiseman et al., 2007; NetMark, 2004^a; NetMark, 2004^c; NetMark, 2004^b).

Though overall bed net ownership per household was very encouraging in the study area, this study noted both differences and similarities when the average number of nets owned per household among net-owning households was compared with those identified in other studies. The finding in this study was lower than the findings in Gambia and Senegal (Wiseman et al., 2007; NetMark, 2004^a), but was found to be similar with the findings from Nigeria (NetMark,

With regard to occupation of respondents, being a housewife was significantly associated with possession of more than one mosquito net. Although the interpretation is difficult, the possible explanation might be that in comparison with those mothers which are civil servants, merchants etc., those which are housewives may not have the knowledge of family planning and/or access to family planning education, which leads to giving birth to more children. Since net distribution was based on family size, these groups of mothers will have better chance of getting more than one mosquito net during distribution.

In line with the family size based distribution issue, households with less than or equal 5 persons were found to have a significantly less chance of possessing more than one mosquito net. Particularly, this association was in accordance with what has been reported from the woreda health office, key-informants and FGD participants. Since the nets were distributed based on family size, those households with smaller family size are not expected to have higher number of mosquito nets.

The refusal of household to IRS in many parts of the world has been recorded either as due to ignorance of mosquito control or to rigid folk social beliefs that vary by degrees (Yadav et al., 2007). In the present study, some people admitted to be reluctant as far as spraying is concerned. The main reason mentioned is that spraying causes inconvenience and waste of time in shifting household goods. Unless such misconceptions are circumvented and minor inconveniences tolerated, the malaria control strategy will be hampered.

The constructive and substantial role of traditional medicine among African societies was recognized during the post-colonial period when the Western-oriented health care system failed to effectively meet the health care needs of the African broad masses. This unquestionably shows that the populations of developing countries heavily rely on traditional medicine to cope up with their health problems. The only medical care for the people of Ethiopia in the past was traditional medicine. In traditional medical practice, different types of medicines including plants, animal products and minerals are used (Teshome-Bahiru, 2006). In the present study area, it was found that 23(5.3%) of the respondents used traditional remedies to treat malaria. The route of

administration mentioned for all the traditional drugs was oral (100%). In formulating liquid preparations, water was reported to be used as a solvent. In this survey, in most plant traditional remedies used for malaria, single plant species remedies and leaf remedies predominate. Concurrent to a report in central Ethiopia (Gedif and Hahn, 2002^b), liquid preparations were reported to be the predominant dosage forms. The doses and duration of administration of herbs varied from household to household and the two most commonly mentioned herbs were *Carica papaya* (papaya) and *Azadirachta indica* (neem). From animal products, "metata" was found to be used by some respondents and repeatedly reported that it is effective in the treatment of *P.vivax* malaria.

There are multiple reasons why people turn to herbal medicine. The most common reasons are cultural acceptability, perceived efficacy and inaccessibility of modern health services in terms of geography, cost, or time (Gedif and Hahn, 2002^a). In accord with these arguments, a very few of the FGD participants in the present study revealed that because of problems associated with cost (indirect cost), perceived efficacy etc., they preferred to use traditional anti-malarial drugs unless the perceived malarial illness was serious.

7. STRENGTHS AND LIMITATIONS OF THE STUDY

This study was not conducted without a limitation. It was carried out in August, the time when malaria problem in the area is not that much high. This could lead the respondents to underestimate the risk for malaria and chance to self-medicate with anti-malarials. It could have been better to undertake such kind of studies during the peak transmission months of malaria to elucidate the heartfelt needs of the community. Nevertheless, this study provides useful information about perceived malaria, and anti-malarial drug use for self-care.

The study data was based on 15-days self-reports, and therefore was subject to reporting or recall bias and respondents may not remember the exact drug that they used for self-medication, the number of ill individuals and symptom complexes before weeks. In addition to these, limitations of the study design and the methods of data collection might create some potential for biases in this study. The cross-sectional design gives information about a certain point of time and so answers might have been varied in different malaria seasons. Data collection relied on information given by the interviewees because of this, practices such as presence and use of ITN and treatment seeking could not be verified. Not many studies conducted particularly in Ethiopia, with regard to self-medication with anti-malarials for perceived malaria/symptom complexes and about anti-malarial drug hoarding trends made it difficult in comparing results.

Not revealing the truth by the respondents was also one of the limitations of this study. Since the interviewees were not willing to show part of the anti-malarial drugs hoarded, and to report all the mosquito nets they own, this may create difference from the reality.

Despite the above limitations, the present study is the first of its kind to investigate self-medication with anti-malarials for perceived malaria/symptom complexes, and about anti-malarial drug hoarding. Considering these, the present study is vital as it had provided at least baseline information about the above critical issues. Moreover, the study included both quantitative and qualitative parts, which enabled to understand the community perceptions and knowledge in depth. This research had included the needs of the current information by

governmental and non-governmental bodies at different levels. And also respondents were represented from all residential settings; peri-urban, nearest to peri-urban and farthest rural communities of the study area.

9. RECOMMENDATIONS

Considering the complexity of behavioral and deep-rooted convictions, there are no cheap 'magic bullet' solutions for the problems highlighted above. However, the following recommendations are forwarded:

- Though reported malaria in the study place is lower in comparison with other areas, further efforts like training and distributing more health extension workers, and closely supervising them etc., must be done in order to coordinate and strengthen control activities, and then completely eradicate malarial.
- Although it would not be possible or even desirable to try to eliminate self-medication completely, it is important to find ways of using this practice to strengthen primary health care through educating consumers in how to avoid the irrational use of anti-malarial drugs. People should be informed about alternatives to self-medication, i.e. seeking qualified advice. Moreover, a proper anti-malarial drug policy and adequate flow of reliable drug information is essential.
- Communities' perceptions of different types of treatment may hinder prompt access to the most effective treatment. These findings demonstrated the need to create awareness at the community level when new drugs are introduced to the market and to inform the community and health professionals about the reasons behind the drug policy change. Awareness creation is also necessary on the consequences of drug hoarding and rampant self-medication, and further studies to investigate these practices have to be done.
- Education of health professionals, particularly those working in the private sector not to sell prescription drugs without prescription and enforcement of the law on those who transgressed the rules and laws governing the pharmaceutical sector.
- The community should be informed now and then about the nature of mosquito nets distributed, whether it is Long-lasting insecticidal net (LLIN) or the usual Insecticide-treated bed net (ITN) as

REFERENCES

- AADAC (Alberta Alcohol and Drug Abuse Commission) (2003). BEYOND the ABCs: Alcohol-Medication Interactions. URL: http://www.aadac.com/documents/beyond_abcs_alcohol_medical_interaction.pdf. (Last accessed on May 01/2009).
- Abdo-Rabbo A (2003). Household survey of treatment of malaria in Hajjah, Yemen. *East Mediterr Health J*, 4(9): 601-606.
- Abuaku BK, Koram KA, Binka FN (2005). Anti-malarial Prescribing Practices: A Challenge to Malaria Control in Ghana. *Med Princ Pract*, 14:332–337.
- Abula T and Worku A (2001). Self-medication in three towns of North-West Ethiopia. *Ethiop J Health Dev*, 15(1):25-30.
- AHS (Alberta Health Services) (2007). BEYOND the ABCs-Alcohol and Medication Interaction. URL: http://www.aadac.com/87_143.asp. (Last accessed on May 01/2009).
- AMREF (2005). Malaria Prevention and Control Strategy 2006-2010, Roll Back Malaria Partnership. URL: <http://www.amref.org>. (Last accessed on December 03/2008).
- Awad A, Eltayeb I, Matowe L, Thalib L (2005). Self-medication with Antibiotics and Anti-malarials in the community of Khartoum State, Sudan. *J Pharm Pharmaceut Sci*, 8(2):326-331.
- Barry AE (2005). Malaria Epidemiology: Insights from the genome of the malaria parasite. *J Mol Genet Med*, 1(2):76- 86.
- Basco Lk (2004). Molecular epidemiology of malaria in Cameroon. XIX. Quality of anti-malarial drugs used for self-medication. *Am J Trop Med Hyg*, 70(3):245–250.

Collins FH, Kamau L, Ranson HA, Vulule JM (2000). Molecular entomology and prospects for malaria control. *Bull World Health Organ*, 78(12): 1412–1423.

Cox FEG (2002). History of Human Parasitology. *Clin Microbiol Rev*, 15(4): 595–612.

CSA [Ethiopia] (2007). Social Statistics. URL: http://www.csa.gov.et/text_files/national%20statistics%202006/Population.pdf. (Last accessed on December 14/2008).

CSA [Ethiopia] and ORC Macro (2006). Ethiopian Demographic and Health Survey 2005. Addis Ababa, Ethiopia and Calverton, Maryland, USA: CSA and ORC Macro.

CSA [Ethiopia] (2008). Summary and statistical report of the 2007 population and housing census results. Addis Ababa: Ethiopia.

Dash AP, Adak T, Raghavendra K, Singh OP (2007). The biology and control of malaria vectors in India. *Current Science*, 92(11): 1571-1578.

Davis JC, Clark TD, Kemble SK, Talemwa N, Njama-Meya D, Staedke SG, Dorsey G (2006). Longitudinal study of urban malaria in a cohort of Ugandan children: description of study site, census and recruitment. *Malar J*, 5:18. doi:10.1186/1475-2875-5-18.

Deressa W, Ali A, Berhane Y (2006). Review of the interplay between population dynamics and malaria transmission in Ethiopia. *Ethiop J Health Dev*, 20(3): 137-144.

Deressa W, Ali A, Hailemariam D (2007^a). Malaria-related health-seeking behavior and challenges for care providers in rural Ethiopia: Implications for the future. *J biosoc Sci*, 40(1): 115-135.

Deressa W, Chibsa S, Olana D (2004). The distribution and magnitude of malaria in Oromia, Ethiopia. *Ethiop J Health Dev*, 18(3):164-181.

Gedif T and Hahn HJ (2002^b). Herbalists in Addis Ababa and Butajira, Central Ethiopia: Mode of service delivery and traditional pharmaceutical practice. *Ethiop J Health Dev*, 16(2):191-197.

Gonzalez MA, Menendez C, Font F, Kahigwa E, Kimario J, Mshinda H, Tanner M, Bosch-Kapblanch, Alonso PL (2000). Cost-effectiveness of iron supplementation and malaria chemoprophylaxis in the prevention of anemia and malaria among Tanzanian infants. *Bull World Health Organ*, 78(1):97-107.

Habtetsion T (2007). How Local Community Perception about Malaria Affects the Utilization of Insecticide Treated Mosquito Nets in Hawzien Woreda, Eastern Tigray, Ethiopia. [MPH Thesis], Department of Community Health, Faculty of Medicine, Addis Ababa University.

Hetzel MW, Msechu JJ, Goodman C, Lengeler C, Obrist B, Kachur SP, Makemba A, Nathan R, Schulze A, Mshinda H (2006). Decreased availability of anti-malarials in the private sector following the policy change from chloroquine to sulphadoxine-pyrimethamine in the Kilombero Valley, Tanzania. *Malar J*, 5: 109. doi:10.1186/1471-2458-8-317.

Jima D, Tesfaye G, Deressa W, Woyessa A, Kebede D, Alamirew D (2005). Baseline survey for the implementation of insecticide treated mosquito nets in Malaria control in Ethiopia. *Ethiop J Health Dev*, 19(1): 16-23.

JRM (Joint Review Mission) (2004). Education Sector Development Programme II, Benishangul-Gumuz Regional Report. URL:[http://www.dagethiopia.org/DOC/2004/Benishangul Gumuz %20Report%20JRM04.doc](http://www.dagethiopia.org/DOC/2004/Benishangul%20Gumuz%20Report%20JRM04.doc). (Last accessed on December 12/2007).

Krettli AU (2001). Anti-malarial chemotherapy: Mechanisms of action, resistance, and new directions in drug discovery. *Mem Inst Oswald Cruz*, 96(8): 1185-1186.

Krzych U and Schwenk RJ (2005). The Dissection of CD8 T Cells During Liver-Stage Infection. In: Langhorne J (Ed). *Immunology and Immunopathogenesis of Malaria*. Germany: Springer-Verlag Berlin Heidelberg, PP.11-24.

Lalloo DG , Shingadia D, Pasvol G, Chiodini PL, Whitty CJ, Beeching NJ, Hill DR, Warrell DA, Bannister BA (2007). Practice Guidelines: UK malaria treatment guidelines for the HPA Advisory Committee on Malaria Prevention in UK Travellers. *J Infect*, 54: 111-121.

Legesse Y, Tegegn A, Belachew T and Tushune K (2007). Knowledge, Attitude and Practice about Malaria Transmission and Its Preventive Measures among Households in Urban Areas of Assosa Zone, Western Ethiopia. *Ethiop J Health Dev*, 21(2): 157-165.

LO RE III V and Gluckman SJ (2003). Prevention of Malaria in Travelers. *Am Fam Physician*, 68(3): 509-514.

Mabaso MLH, Vounatsou P, Midzi S, Da Silva J, Smith T (2006). Spatio-temporal analysis of the role of climate in inter-annual variation of malaria incidence in Zimbabwe. *Int J Health Geogr*, 5:20. doi:10.1186/1476-072X-5-20.

Marsh VM, Mutemi WM, Willetts A, Bayah K, Were S, Ross A, Marsh K (2004). Improving malaria home treatment by training drug retailers in rural Kenya. *Trop Med Int Health*, 9(4): 451-460.

Mizuno Y, Kato Y, Kudo K, Kano S (2009). First Case of Treatment Failure of Artemether-Lumefantrine in a Japanese Traveler with imported falciparum malaria. *Jpn J Infect Dis*, 62: 139-141.

Mnyika KS, Killewo JZ, Kabalimu TK (1995). Self-medication with anti-malarial drugs in Dar es Salaam, Tanzania. *Trop Geogr Med*, 47(1): 32-4.

- MOH (2008). The Ethiopia National Malaria Indicator Survey 2007. Addis Ababa: Ethiopia.
- MOH (2002). Guidelines for malaria vector control in Ethiopia. Malaria and other vector-borne diseases control unit, Epidemiology and AIDS control department. Addis Ababa: Ethiopia.
- MOH (2004). Malaria diagnosis and treatment guidelines for health workers in Ethiopia. Second edition. Addis Ababa: Ethiopia.
- NEHC (Navy Environmental Health Center) (2000). Navy Medical Department Pocket Guide to Malaria Prevention and Control. URL: <http://www.nehc.med.navy.mil/downloads/prevmed/malariapocketsep2000.pdf>. (Last accessed on December 10/2007).
- Ndugwa RP, Ramroth H, Müller O, Jasseh M, Sié A, Kouyaté B, Greenwood B, Becher H (2008). Comparison of all-cause and malaria-specific mortality from two West African countries with different malaria transmission patterns. *Malar J*, 7:15. doi:10.1186/1475-2875-7-15.
- NetMark (2004^a). NetMark 2004 Survey on Insecticide-Treated Nets (ITNs) in Senegal. Washington DC: USA.
- NetMark (2004^b). Awareness, Ownership, and Use of Mosquito Nets in Nigeria, Senegal, Zambia, Ghana, and Ethiopia. Cross-country results from the 2004 NetMark surveys. Washington DC: USA.
- NetMark (2004^c). NetMark 2004 Survey on Insecticide-Treated Nets (ITNs) in Zambia. Washington DC: USA.
- Oberlander L and Elverdan B (2000). Malaria in the United Republic of Tanzania: cultural considerations and health-seeking behavior. *Bull World Health Organ*, 78 (11): 1352–1357.

- Shunmay Y, Pongtavornpinyo W, Ian MH, Anne JM, Nicholas JW (2004). Anti-malarial Drug Resistance, Artemisinin-Based Combination Therapy and the Contribution of Modeling to Elucidating Policy Choices. *J Trop Med Hyg*, 71(2): 179–186.
- Sultana A (2001). Malaria, Knowledge, Attitude & Practice; in a semi-urban population of Rawalpindi, Pakistan. *The Professional*, 8(4):433-438.
- Teklehaimanot A, McCord GC, Sachs JD (2007). Scaling Up Malaria Control in Africa: An Economic and Epidemiological Assessment. *Am J Trop Med Hyg*, 77(Suppl 6): 138–144.
- Temu MJ, Kaale E, Marawiti M (2006). Storage of anti-malarials at household level and associated factors in Kiromo ward, Bagamoyo Tanzania. *Afr Health Sci*, 6(1): 43–48.
- Teshome-Bahiru W (2006). Impacts of Urbanisation on the Traditional Medicine of Ethiopia. *Anthropologist*, 8(1): 43-52.
- Tilaye T and Deressa W (2007). Community perceptions and practices about urban malaria prevention and control in Gondar town, North-West Ethiopia. *Ethiop Med J*, 45(4):343-51.
- UNICEF (2007). Malaria & children. Progress in intervention Coverage. USA: New York.
- Vanderbroek I, Clewaert JB, De Johnckheere, Sanca S, Semo L, Van Dame P, Van Puyelde L, De Kimoe N (2004). Use of medicinal plants and pharmaceuticals by indigenous communities in the Bolivian Andes and Amazon. *Bull World Health organ*, 82(4): 243–250.
- WEF(World Economic Forum) (2006). Guidelines for Employer-Based Malaria Control Programmes. CH-1223 Cologny/Geneva: Switzerland.
- White NJ (2004). Antimalarial Drug Resistance. *J Clin Invest*, 113 (8): 1083-1092.

WHO (2005^a). Malaria controls in complex emergencies: An Inter-Agency Field Handbook. Geneva: Switzerland.

WHO (2001). Antimalarial drug combination therapy report of a WHO technical consultation. Geneva: Switzerland.

WHO (2002). Malaria Entomology and Vector Control: Learner's Guide. Social Mobilization and Training-Control, Prevention and Eradication Department; Communicable Diseases Cluster. Trial Edition. Geneva: Switzerland.

WHO (2003). Combination Therapies and Formulation of Antimalarial Drug Policy. Tutor's Guide. HIV/AIDS, Tuberculosis and Malaria Roll Back Malaria. Trial Edition. Geneva: Switzerland.

WHO (2004). Combination therapy in malaria: orientations and options for the african region, regional office for Africa. Brazzaville: Republic of the Congo.

WHO (2005^b). Roll Back Malaria Department. Malaria Control Today: Current WHO Recommendations. *Working Document*. Geneva: Switzerland.

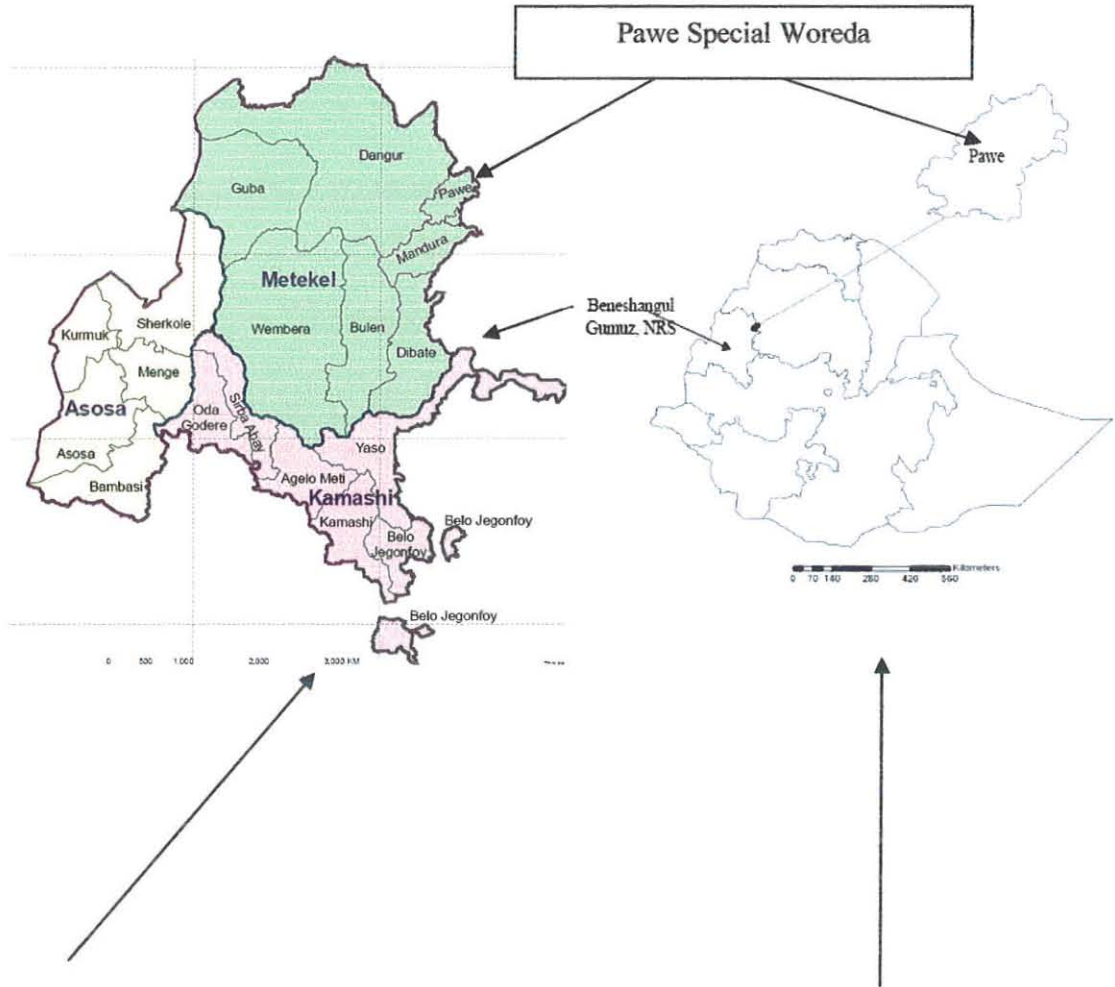
WHO (2006). Guidelines for the Treatment of Malaria. Geneva: Switzerland.

WHO (2007). Malaria elimination: A field manual for low and moderate endemic countries. Geneva: Switzerland.

WHO (2008). World malaria report. Geneva: Switzerland.

Wiseman V, Scott A, Mcelroy B, Conteh L, Stevens W (2007). Determinants of bed net use in the Gambia: Implications for malaria control. *Am J Trop Med Hyg*, 76(5):830–836.

Annex 1: Map of pawe special woreda



Map of Beneshangul-Gumuz region

Map of Ethiopia

Adapted from: <http://www.ethiodemographyandhealth.org/Benishangul.html>

106	Household family size (including yourself)	(Write in numbers)_____		
107	From the total household members			
	107.1	Number of females (Write in numbers)_____		
	107.2	Number of males (Write in numbers)_____		
108	Occupation	Housewife 01 Merchant (trader) 02 Farmer 03 Government employee 04 Daily labourer 05 _____Others, specify 06	[][]	
109	House hold income in a month	Monthly income in birr _____ or Income in kind _____ quintals per year (specify type)		
110	Educational status of spouse (for those who claim to have husband)	Illiterate 01 Read only 02 Read & write 03 Grades 1-6 04 Grades 7-8 05 Grades 9-12 06 Grades 12+ 07 Technical Education 08	[][]	
Part II: Information sources				
201	Do you have the following properties?			
201.1	Functional Radio	Yes 01 No 02	[][]	
201.2	Functional Television	Yes 01 No 02	[][]	
Part III: Measures taken during perceived malaria illness/symptom complexes				
301	From your family members was there any ill person in the previous two weeks?	Yes 01 No 02	[][]	If the response is 02, skip to Q 401

302	How many of your family members?	_____ (mention in numbers)		
303	For each ill person("patient"), please ask the following questions (if the number of ill persons is more than one, use another similar copy for data collection)			
303.1	Sex of the patient	Male 01 Female 02	[] []	
303.2	Age of the patient in years	_____ (years)		
303.3	Patient's status in the family	Child(of any age group as far as he/she is a "child" in the household) 01 Servant 02 mother 03 Husband 04 _____ Others, specify 05	[] []	
303.4	What was the illness for each individual? (Read for the respondents)			
303.4.1	Fever 01	1. Yes 2. No	[] []	If malaria and/or malaria symptom complexes are not mentioned, directly go to Q 401.
303.4.2	Chills 02	1. Yes 2. No	[] []	
303.4.3	Shivering 03	1. Yes 2. No	[] []	
303.4.4	Head ache 04	1. Yes 2. No	[] []	
303.4.5	Joint pain 05	1. Yes 2. No	[] []	
303.4.6	Vomiting 06	1. Yes 2. No	[] []	
303.4.7	Loss of appetite 07	1. Yes 2. No	[] []	
303.4.8	Thirst 08	1. Yes 2. No	[] []	
303.4.9	Malaria 09 _____ (others specify, all the encountered illnesses)	1. Yes 2. No	[] []	
303.5	What measure was taken for the ill person(s)? <i>(write the respective numbers of all applicable)</i>	Taken to health institutions (hospital, health center, health post) 01 Medicines were given at home 02 Used drugs bought from private drug retail outlets 03 No measure was taken 04 _____ Others, specify 05	[] [] [] []	If the response is 04, skip to Q 303.5.2
303.5.1	If the response to question 303.5 is 01, ask the following questions:			If the response to Q 303.5 is not 01, skip to Q303.5.2

409	How many mosquito nets do you have?	Specify in numbers _____		
410	In a given year, when is the probability of getting malaria high? <i>(write the respective numbers of all applicable)</i>	Winter 01 Spring 02 Summer 03 Autumn 04	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	
411	Can malaria infect people of all age group?	Yes 01 No 02	<input type="checkbox"/> <input type="checkbox"/>	If the response is 01, skip to Q 413
412	Which age group is/are exempted? <i>(write the respective numbers of all applicable)</i>	children (<15 years) 01 adults (≥15-65 years) 02 elders(>65 years) 03 pregnant women 04 Non-pregnant women 05 _____ Others, specify 06	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	

Part V: Questions to assess general knowledge about anti-malarial drugs

501	Is malaria curable by modern medicines available in your area?	Yes 01 No 02	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	If the response is 02, skip to Q 503
502	Please mention the anti-malarial drugs that you know? <i>(write the respective numbers of all applicable)</i>	Chloroquine 01 Quinine 02 Coartum 03 Fansidar 04 _____ Others, specify 05	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	
503	What Methods do you use to differentiate/identify anti-malarial drugs from others? <i>(write the respective numbers of all applicable)</i>	Color of the drug/package 01 Smell (odour) of the drug 02 reading /letting others to read the label 03 making signs on the containers (packaging) 04 preparing a different storing place for each kind of drug 05 Don't know 06 _____ Others, Specify 07	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	
504	What do you think is the consequence(s) of non-adherence to anti-malarial	Specify _____		

	medication?			
505	Is there any of your family member (including yourself) who has ever taken alcoholic drink while on anti-malarial drug treatment?	Yes 01 No 02	[] []	
Part VI : Questions about hoarding of anti-malarial drugs at household level				
601	Is there any hoarded anti-malarial drug in your home?	Yes 01 No 02	[] []	If the response is 02, skip to Q 701
602	Are you willing to show me the hoarded anti-malarials?	Yes 01 No 02	[] []	If the response is 02, skip to Q 701
603	Write the names of the hoarded anti-malarials.	write the names _____ (if the label is illegible, please write it as <i>label illegible</i>)		
604	What are the storage conditions of the anti-malarials?	In a torn container/package in dry area 01 without container/package in dry area 02 In untorn containers/packages in hot area 03 In untorn containers/packages in moist area 04 In untorn container/package in dry area 05 _____ Others, Specify 06	[] []	
Part VII: Questions to assess general knowledge & practice of traditional anti-malarial drugs				
701	Is there any traditional drug that you use for malaria treatment?	Yes 01 No 02	[] []	If the response is 02, thank the respondent and end the interview
702	What type of traditional anti-malarial treatment do you know?	Plant product 01 Animal product 02 _____ Others, Specify 03		If the response is 02, skip to Q 704

703	If the response to question 702 is Plant product (01), please fill the following table (if the respondent did not said plant product, go to question 704)				
No	Name of the plant	Part of the plant used for treatment (leaf, root, stem, etc.)	Method of preparation	How many times is it administered per day?	For how long (how many days) is it administered?
1					
2					
3					
4					
704	If the response to question 702 is animal product(02), please fill the following table				
	Name of the animal	Part of the animal used for treatment	Method of preparation	How many times is it administered per day?	For how long (how many days) is it administered?
1					
2					
3					
4					

Finally, thank the respondent and complete the interview

Supervisors' code _____ signature _____

Annex 4: Verbal Consent Form and topic guides for FGD participants

(to be read to the FGD participants)

Good morning (afternoon), Thank you for your coming here

1. Introduction

My name is Seid Mussa. I am a student at AAU, School of Pharmacy. My colleague besides me is called

_____ He came from _____

2. Title of the study

Anti-malarial drug and mosquito net use pattern in Pawe Special Woreda: A community based survey.

3. Purpose of the study:

The study will help in identifying and enhancing the knowledge and understanding of anti-malarial drug and mosquito net use at the community level, factors contributing to non-adherence to anti-malarial drugs, preference among anti-malarial drugs, anti-malarial drug hoarding, and mosquito net use pattern and be useful in developing interventions that will be undertaken to address the proper use of anti-malarial drugs, and mosquito nets in the woreda in particular and in the region in general. The data collected could also be used by organizations involved in this area to deal with the problems associated with non-adherence and anti-malarial drug resistance and also about the misperception about malaria, anti-malarials, and mmosquito nets which could adversely affect the outcome of malaria control program.

4. How you were identified:

You were selected to participate in FGD part of the study as the woman who has lived in the area for long years(≥ 10 Years) and knew the drug and mosquito net use trends and the principal investigator thinking that you represent the whole women community and you could reflect the prevalent drug and mosquito net use trends in the community. When you express your opinion or idea, you are encouraged to be honest in your view about malaria, anti-malarial drugs and mosquito net.

5. Consent for participation and tape –recording

Your presence in this session indicates that you are volunteer and agreed to participate for the discussion. Your participation in this study is completely based on your will and there is no penalty for refusing to take part. You have the right to refuse to answer any questions and end the discussion if you find it necessary to do so. For the sake of accuracy and efficiency, we will tape-record the sessions, unless there is objection.

6. Issues of confidentiality

Please be certain that any information collected here is completely confidential. The recorded voice will be erased after transcribing the information and your name will never be used in connection with any information you provide in the results of this research.

7. Importance of total /group responses

In this group and session everybody should feel free and talk freely. Each and every opinion /idea/ is important and wanted. It is vital that all the group participants get a chance to express their idea (opinion). In this session, there is no wrong or right answer. Everybody can express the opinion or attitude pertinent to him/her. When you express your opinion or idea, you are encouraged to be honest in your view about malaria, anti-malarial drugs and mosquito net. We want you to focus your comments on the program and not towards each other or member of the group.

8. Introducing each other

At this moment, we would like you to introduce yourself to the rest of the group.

9. Questions for Discussion

Next, we would like to hear a little about your experience or knowledge about malaria, anti-malarial drug and mosquito net use pattern at the household level.

- ♣ Discussion if malaria is a major health problem in the study area & on the cause of malaria
- ♣ Discussion on the transmission of malaria, including seasonal peaks
- ♣ Discussion on the risk groups to malaria
- ♣ Discussion on the common symptoms of malaria
- ♣ Discussion on the prevention methods of malaria
- ♣ Discussion on the ITN or mosquito net-benefit, disadvantage, usage, etc.
- ♣ Discussion on the treatment of malaria
- ♣ Discussion on the knowledge and usage practice of anti-malarial drugs (drug centered approach)
- ♣ Discussion on the sources of anti-malarial drugs and associated factors
- ♣ Discussion on the type of treatment employed in case when any family member is ill due to malaria and/or symptom complexes
- ♣ Discussion on the anti-malarial drug hoarding and associated factors
- ♣ Discussion on the types of traditional modes of treatment for malaria

Up to now we have talked about malaria, anti-malarial drug and mosquito net use pattern at the household level. We thank you for giving us your time and for your cooperation. We do appreciate all your suggestions and ideas. At this point, is there anything we forgot to ask or anything you would like to mention before we close the session?

Personal information of Key-informant interview participants

1. Date _____
2. Age _____
3. Marital Status _____
4. Religion _____
5. Educational Status _____
6. Occupation _____
7. Number of participants _____
8. Duration of Session _____

Annex 5: Verbal consent form and topic guides for key-informant interview Participants

(to be read to the key informants)

Good morning (afternoon), Thank you for your coming here

1. Introduction

My name is Seid Mussa. I am a student at AAU, School of Pharmacy.

2. Title of the study

Anti-malarial drug and mosquito net use pattern in Pawe Special Woreda: A community based survey

3. Purpose of the study:

The study will help in identifying and enhancing the knowledge and understanding of anti-malarial and mosquito net use at the community level, factors contributing to non-adherence to anti-malarial drugs, preference among anti-malarial drugs, anti-malarial drug hoarding, and mosquito net use pattern and be useful in developing interventions that will be undertaken to address the proper use of anti-malarial drugs and mosquito nets in the woreda in particular and in the region in general. The data collected could also be used by organizations involved in this area to deal with the problems associated with non-adherence and anti-malarial drug resistance and also about the misperception about malaria, anti-malarials, and bed nets which could adversely affect the outcome of malaria control program.

4. How you were identified:

You were selected to participate in Key-informant interview part of the study as the health workers at the health facility, administrative body in the woreda health office, formal village leaders etc., and the principal investigator thinking that you represent the community and you could reflect the prevalent drug and mosquito net use trends in the community. When you express your opinion or idea, you are encouraged to be honest in your view about malaria, anti-malarial drugs and mosquito net.

5. Consent for participation and tape –recording

Your presence in this session indicates that you are volunteer and agreed to participate for the discussion. Your participation in this study is completely based on your will and there is no penalty for refusing to take part. You have the right to refuse to answer any questions and end the discussion if you find it necessary to do so. For the sake of accuracy and efficiency, I will tape-record the sessions, unless there is objection.

6. Issues of Confidentiality

Please be certain that any information collected here is completely confidential. The recorded voice will be erased after transcribing the information and your name will never be used in connection with any information you provide in the results of this research.

7. Questions for Discussion

Next, I would like to hear a little about your experience or knowledge about malaria, anti-malarial drug and mosquito net use pattern at the house hold level.

- ♣ Discussion on the cause of malaria
- ♣ Discussion on the transmission of malaria, including seasonal peaks
- ♣ Discussion on the risk groups to malaria
- ♣ Discussion on the Common symptoms of malaria
- ♣ Discussion on the prevention methods of malaria
- ♣ Discussion on the ITN or mosquito net- benefit, disadvantage, usage, etc.
- ♣ Discussion on the treatment of malaria
- ♣ Discussion on the knowledge and usage practice of anti-malarial drugs (drug centered approach)
- ♣ Discussion on the sources of anti-malarial drugs and associated factors
- ♣ Discussion on the type of treatment employed in case when any family member is ill due to malaria and/or symptom complexes
- ♣ Discussion on the anti-malarial drug hoarding and associated factors
- ♣ Discussion on the types of traditional modes of treatment for malaria
- ♣ Discussion on the reason behind the decision of the MOH to change the first line anti-malarial drug for the treatment of *P.falciparum* from fansidar to coartem in 20004 (Question to be presented mainly for health professionals)

Up to now we have talked about malaria and anti-malarial drug use pattern at the house hold level. We thank you for giving us your time and for your cooperation. We do appreciate all your suggestions and ideas. At this point is there anything we forgot to ask or anything you would like to mention before we close the session?

Personal information Key-informant interview participants

1. Date _____
2. Age _____
3. Marital Status _____
4. Religion _____
5. Educational Status _____
6. Occupation _____
7. Duration of Session _____

በአዲስ አበባ ዩኒቨርሲቲ ፣ ፋርማሲ ትምህርት ቤት

ፋርማሲዮቲክስ ትምህርት ክፍል

የፋርማኮኢፒዲሚያሎጂይ እና ሶሻል ፋርማሲ ድህረ ምረቃ ፕሮግራም

በጥናቱ የሚሳተፉት (ተጠያቂዎቹ) እናቶች ናቸው

ሰላም ነዎት? እኔ ዛሬ እዚህ የተገኘሁት በአዲስ አበባ ዩኒቨርሲቲ ተማሪ የሆኑት አቶ ሰኢድ ሙሳ በቤተሰብ ደረጃ ስለ ወባ ፀረ-ወባ መድሃኒት እና ከአልጋ አጎብር አጠቃቀም ለሚሰሩት ጥናት መረጃ ለመሰብሰብ ነው ። ከዚህ ጥናት የሚገኙት ውጤቶች ከፀረ-ወባ መድሃኒትና ከአልጋ አጎብር አጠቃቀም ጋር ተያያዥ የሆኑ ችግሮችን ለመዳሰስና ችግሮቹን ለመቅረፍ ጠቃሚ የሆኑ መንገዶችን ለመቀየስ ይረዳሉ። ስለሆነም የተወሰኑ ከወባ ። ፀረ-ወባ መድሃኒትና ከአልጋ አጎብር ጋር የተያያዙ ጥያቄዎች እንጠይቀዎታለን፤ እዚህ ላይ ልናረጋግጥልዎት የምንወደው ነገር ለጥያቄዎቻችን የሚሰጡን መልስ ሚስጥራዊ መሆኑን ነው ምክንያቱም መጠይቁ ላይ ስምዎን አንመዘግብም። ከዚህ በተጨማሪ በጥናቱ ላለመሳተፍ መወሰን ወይም ያልፈለጉትን ጥያቄ በማንኛውም ሰዓት ያለመመለስ መብት አለዎት።

በዚህ ጥናት ውስጥ ላለመሳተፍ ቢወስኑ ምንም ችግር አይደርስብዎትም።

እዚህ ላይ ስለ ጥናቱ ሊጠይቁኝ የሚፈለጉት ጥያቄ አለ?

በጥናቱ ላለመሳተፍ ይስማማሉ? _____ (እሺ ወይም እምቢ)

እሺ (1) ካሉ ቀጥል

እምቢ (2) ካሉ አመሰግንና አቁም

ማስታወሻ:- የተጠያቂዎ ቤት ዝግ ከሆነ በሌላ ጊዜ ለመመለስ መንደሩን እና የቤት ቁጥሩን መዝገበው መጠይቁን ለተቆጣጣሪው ይመልሱ።

2ኛ ጊዜ ጉብኝት _____

3ኛ ጊዜ ጉብኝት _____

Annex 7: መጠይቅና የመረጃ ማሰባሰቢያ ቅጽ

1. ስለ ቤተሰብ እና መረጃ ሰብሳቢ ጠቅላላ መረጃ

- የመዝገብ ቁጥር _____
- የመረጃ ሰብሳቢው ኮድ _____
- ቃለ መጠይቅ የተካሄደበት ቀን _____
- የተጠያቂው መኖሪያ 1. ከተማ _____ 2. ገጠር _____
- የቀበሌው (የመንደሩ) ስም _____
- የቤት ቁጥር _____

ክፍል I: አጠቃላይ የተጠያቂው መሰረታዊ የግል መረጃዎች				
መለ/ቁ	ጥያቄ	መልስ	ኮድ	ይታለፍ
101	እድሜዎ ስንት ነው?	_____ ዓመት		
102	ብሔርዎ ምንድን ነው?	<p>አማራ 01</p> <p>ከምባታ 02</p> <p>ሀድያ 03</p> <p>አሮሞ 04</p> <p>ትግሬ 05</p> <p>አገው 06</p> <p>ወላይታ 07</p> <p>-----ሌላ ካለ ይጠቀስ 08</p>	[]	
103	ሀይማኖትዎ ምንድን ነው?	<p>ሙስሊም 01</p> <p>ኦርቶዶክስ 02</p> <p>ካቶሊክ 03</p> <p>ፕሮቴስታንት 04</p> <p>-----ሌላ ካለ ይጠቀስ 05</p>	[]	

104	የትምህርት ደረጃዎችን በነግሩኝ	ማንበብና መጻፍ የሚችሉ 01 ማንበብ ብቻ የሚችሉ 02 ማንበብና መጻፍ የማይችሉ 03 1-6 ክፍል 04 7-8 ክፍል 05 9-12 ክፍል 06 12+1 እና ከዚያ በላይ 07 የሞያ ትምህርት 08	[]	
105	አሁን ያለዎትን የትዳር ሁኔታ በነግሩኝ?	ያላገቡ 01 ባለትዳር 02 የተፋቱ 03 የትዳር ጓደኛዎ በሞት የተለየዎት 04 -----ሌላካለ ይጠቀስ 05	[]	
106	የቤተሰብ ብዛት (እርስዎን ጨምሮ)	(በቁጥር ይገለጥ)-----		
107	ከአጠቃላይ የቤተሰቡ አባል:	107.1 የሴቶች ቁጥር ምን ያህል ነው?(በቁጥር ይገለጥ)----- 107.2 የወንዶች ቁጥር ምን ያህል ነው?(በቁጥር ይገለጥ)-----		
108	ስራዎ ምንድን ነው?	የቤት እመቤት 01 ነጋዴ 02 ገበሬ 03 የመንግስት ሰራተኛ 04 የቀን ሰራተኛ 05 -----ሌላ ካለ ይጠቀስ 06	[]	
109	የቤተሰብዎ ገቢ በወር ምን ያህል ነው?	ወርሀዊ ገቢ ብር_____ ወይም ገቢ በአይነት በአመት_____		

		ኩንታል(የምርቱ አይነት ይጻፍ)		
110	የቤተሰቡን አባዎራ የትምህርት ደረጃ ቢነግሩን	ማንበብና መጻፍ የሚችሉ 01 ማንበብ ብቻ የሚችሉ 02 ማንበብና መጻፍ የማይችሉ 03 1-6 ክፍል 04 7-8 ክፍል 05 9-12 ክፍል 06 12+1 እና ከዚያ በላይ 07 የሞያ ትምህርት 08	[] []	ተጠያቂዎ እማውራ ከሆኑ በቀጥታ ወደ ጥያቄ ቁጥር 201 ይሂዱ

ክፍል-የቤተሰብ የመረመረጃ ምንጭዎች መጠየቂያ

መለ/ቁ	ጥያቄ	መልስ	ኮድ	ይታለፍ
201	የሚከተሉት ንብረቶች አሏችሁ?			
201.1	ሬዲዮ	አዎ 01 የለንም 02	[] []	
201.2	ቴሌቪዥን	አዎ 01 የለንም 02	[] []	

ክፍል-III በወባ ህመም/ /በወባ ምልክቶች ጊዜ ስለተወሰዱ እርምጃዎች መረጃ መጠየቂያ

መለ/ቁ	ጥያቄ	መልስ	ኮድ	ይታለፍ
301	ከቤተሰብዎ ውስጥ ባለፉት ሁለት ሳምንታት ውስጥ ታሞ የነበረ ሰው ነበርን?	አዎ 01 የለም 02	[] []	መልስዎ 02 ከሆነ በቀጥታ ወደ ጥያቄ 401 ይሂዱ
302	መልሱ አዎ ከሆነ ከቤተሰብዎ አባላት ስንቶቹ ናቸው?	በቁጥር ይገለጥ_____		
303	ለእያንዳንዱ ታማሚ የሚከተለውን መረጃ ይጠይቁ:- (ታማሚው ከአንድ በላይ ከሆነ ተመሳሳይ የሆነ ለመረጃ መሰብሰቢያ ተጨማሪ ሌላ ኮፒ ይጠቀሙ)			
303.1	የታማሚው ስያ	ወንድ 01 ሴት 02	[] []	

303.5	<p>ለህመምተኛው ምን ተደረገለት?</p> <p>(ከአንድ በላይ መልስ መስጠት ይቻላል)</p>	<p>ቤት ውስጥ መድሀኒት ተሰጠው 01</p> <p>ከመድሀኒት ቤት መድሀኒት በመግዛት ተጠቀመ 02</p> <p>ወደ ጤና ድርጅት (ጤና ጣቢያ፣ ጤና ኬላ ፣ ሆስፒታል) ተወሰደ 03</p> <p>ምንም አልተደረገለትም 04</p> <p>-----ሌላ ካለ ይጠቀስ 05</p>	<p>[] []</p> <p>[] []</p>	<p>መልስዎ 04 ከሆነ በቀጥታ ወደ ጥያቄ ቁጥር 401 ይሂዱ.</p>
303.5.1	<p>የጥያቄ 303.5 መልስ ጤና ድርጅት ከሆነ የሚከተሉትን ጥያቄዎች ይጠይቁ :-</p>			<p>የጥያቄ 303.5 መልስ 03 (ጤና ድርጅት) ካልሆነ በቀጥታ ወደ ጥያቄ ቁጥር 303.5.2 ይሂዱ.</p>
303.5.1.1	<p>ጤና ድርጅት ሄዳችሁ ህመሙ ምንድን ነው አሏችሁ?</p>	<p>_____ይገለጥ</p>		
303.5.1.2	<p>የጤና ድርጅት ከጎበኙ ከጣትዎ (ከክንድዎ) የደም ናሙና ተወስዶ በጤና ተቋሙ በደም መመርመሪያ (በማይክሮስኮፕ) ተመርምሯል?</p>	<p>አዎ 01</p> <p>አልተመረመረም 02</p>	<p>[] []</p>	
303.5.2	<p>የጥያቄ 303.5 መልስ ቤት ውስጥ መድሀኒት ተሰጠው ከሆነ ምን ዓይነት መድሀኒት ተሰጠው?</p>	<p>_____ (የመድሀኒቱ ስም ይገለጥ)</p>		<p>የጥያቄ 303.5 መልስ 01 ወይም 03 ካልሆነ በቀጥታ ወደ ጥያቄ ቁጥር 303.5.3 ይሂዱ.</p>
303.5.2.1	<p>መድሀኒቱን ወስደው ጨርሰዋል ወይስ ገና</p>	<p>ጨርሰዋል 01</p> <p>አልጨረሱም 02</p>	<p>[] []</p>	

	እየወሰዱ ነው?			
303.5.3	የጥያቄ 303.5 መልስ ከመደሀኒት ቤት መደሀኒት በመግዛት ተጠቀመ(02) ከሆነ ምን ዓይነት መደሀኒት ምን ገዝቶ ተጠቀመ?	_____ (የመደሀኒቱ ስም ይገለጥ)		
303.5.3.1	መደሀኒቱን ወስደው ጨርሰዋል ወይስ ገና እየወሰዱ ነው?	ጨርሰዋል 01 አልጨረሱም 02	[] []	

ክፍል IV ስለ ወባ ጠቅላላ ዕውቀትን ለመዳሰስ የቀረቡ ጥያቄዎች

401	የወባ መንስኤው ምንድን ነው ብለው ያስባሉ ? (ከአንድ በላይ መልስ መስጠት ይቻላል)	የግል ንጽህና አለመጠበቅ 01 ጭጋጋማ የአየር ጠባይ 02 ቀዝቃዛ የአየር ጠባይ 03 ቆሻሻ ነገሮች 04 በወባ ትንኝ መነደፋ 05 መራብ 06 ርኩስ መንፋት 07 -----ሌላ ካለ ይጠቀስ 08	[] [] [] [] [] [] [] []	
402	አንድ ሰው በወባ በሽታ እንደተያዘ በእርግጠኝነት እንዴት ማወቅ ይቻላል? (ከአንድ በላይ መልስ መስጠት ይቻላል)	በምልክቶቹ ብቻ 01 በላቦራቶሪ ምርመራ 02 በባህላዊ ህኪሞች 03 ማዎቅ አይቻልም 04 -----ሌላ ካለ ይጠቀስ 05	[] [] [] []	
403	የወባ በሽታ ዋና ዋና ምልክቶች ምን ምን ናቸው? (ከአንድ በላይ መልስ መስጠት ይቻላል)	ትኩሳት 01 ብርድ ብርድ ማለትና ማንቀጥቀጥ 02 እራስ ምታት 03 የጀርባ ህመም 04 የመገጣጠሚያ ህመም 05	[] [] [] [] [] [] [] []	

407	<p>የትኞቹን የወባ መከላከያ ዘዴዎች ያውቃሉ?</p> <p>(ከአንድ በላይ መልስ መስጠት ይቻላል)</p>	<p>የወባ ትንኝ ወደ ቤት እንዳትገባ የቤቱን ቀዳዳዎች በተለያዩ ነገሮች በመድፈን 01</p> <p>ከመኖሪያ ቤት አካባቢ ያሉ ውሃ ያቆሩ ኩራዎችን በማድረቅ 02</p> <p>የቤት በሮችንና መስኮቶችን ምሽት ላይ በጊዜ በመዝጋት 03</p> <p>የኩብት ኩብትን ወይም የተለያዩ እንጨቶችን ጭስ በማጨስ 04</p> <p>ፀረ-ወባ ትንኝ ኬሚካል ወይም ሌሎች መንገዶችን በመጠቀም ወባ ትንኝን በመግደል 05</p> <p>በፀረ-ወባ ትንኝ የተነከረ የአልጋ አጎበር በመጠቀም 06</p> <p>የግልና የአካባቢ ንጽህናን በመጠበቅ 07</p> <p>ምንም አይነት የወባ መከላከያ መንገድ አላውቅም 08</p> <p>-----ሌላ ካለ ጥቀስ 09</p>	<p>[] []</p> <p>[] []</p> <p>[] []</p> <p>[] []</p>	
408	<p>አሁን ቤትዎ ውስጥ የአልጋ አጎበር አለዎትን?</p>	<p>አለን 01</p> <p>የለንም 02</p>	<p>[] []</p>	<p>መልስዎ 02 ከሆነ በቀጥታ ወደ ጥያቄ 410 ይሂዱ</p>
409	<p>አሁን በአጠቃላይ ቤታችሁ ውስጥ ስንት የአልጋ አጎበር አላችሁ?</p>	<p>(በቁጥር ይገለጥ)-----</p>		
410	<p>በአመቱ ውስጥ በወባ የመያዝ እድል ከፍተኛ የሚሆነው መች ነው?</p> <p>(ከአንድ በላይ መልስ መስጠት ይቻላል)</p>	<p>ከታህሳስ-የካቲት 01</p> <p>ከመስከረም-ህዳር 02</p> <p>ከሰኔ-ነሐሴ 03</p> <p>ከመጋቢት-ግንቦት 04</p> <p>አመቱን በሙሉ እንታመማለን 05</p>	<p>[] []</p>	

		አላውቅም 06		
411	የወባ በሽታ በሁሉም የእድሜ ክልል የሚገኙ ሰዎችን ያጠቃልላል?	አዎ 01 አያጠቃም 02 አላውቅም 03	[] []	መልስዎ 01 ወይም 03 ከሆነ በቀጥታ ወደ ጥያቄ 501 ይሂዱ
412	በየትኛው የእድሜ ክልል የሚገኝ/የሚገኙት በወባ አይያዙም ወይም የመያዝ እድላቸው ትንሽ ነው ይላሉ ? (ከአንድ በላይ መልስ መስጠት ይቻላል)	ልጆች (ሀፃናት)(<15 አመት) 01 አዋቂ ሰዎች (≥15-65 አመት) 02 በእድሜ የገፋ ሰዎች(>65 አመት) 03 ነፋሰጡር ሴቶች 04 ነፋሰጡር ያልሆኑ ሴቶች 05 -----ሌላ ካለ ይጠቀስ 06	[] [] [] []	
ክፍል V: ስለ ፀረ-ወባ መድሃኒት ጠቅላላ ዕውቀትን ለመዳሰስ የቀረቡ ጥያቄዎች				
501	በአካባቢዎ በሚገኙ ዘመናዊ ፀረ-ወባ መድሃኒቶች ከወባ መፈወስ ይቻላል ብለው ያምናሉ?	አዎ 01 አይደለም 02	[] []	
502	እባክዎ የሚያውቋቸውን ዘመናዊ ፀረ-ወባ መድሃኒቶችን ስማቸውን ይጥቀሱልኝ? (የሚናገሩትን ሁሉ ቁጥሩን በመልስ መስጫው ይጻፉ)	ክሎሮኪን 01 ኩኒን 02 ኮአርተም 03 ፋንሲደር 04 ሌላ ካለ ይጠቀስ 05	[] [] [] [] [] [] [] [] [] []	

503	<p>ዘመናዊ ፀረ-ወባ መድሃኒቶችን ከሌሎች ቤትዎ ውስጥ ከሚገኙ/ሊገኙ ከሚችሉ ዘመናዊ መድሃኒቶች ለመለየት ምን አይነት መንገድ ይጠቀማሉ?</p> <p>(ከአንድ በላይ መልስ መስጠት ይቻላል)</p>	<p>በማሸጊያው ወይም በመድሃኒቱ መልክ (ከለር) 01 በሽታ 02 [] [] መያዣው ላይ የተጻፈውን በማስነበብ ወይም በማንበብ 03 ማሸጊያው (መያዣው) እቃ ላይ ምልክት በማድረግ 04 ለተለያዩ መድሃኒቶች የተለያዩ ማስቀመጫቦታዎችን በማዘጋጀት 05 አላውቀውም 06 -----ሌሎች(ይገለጹ) 07</p>		
504	<p>የፀረ-ወባ መድሃኒትን በስርዓቱ (በተገቢው ሁኔታ) አለመውሰድ ምን ችግር ያመጣል ብለው ያስባሉ?</p>	<p>----- (ይገለጥ)</p>	<p>[] [] [] []</p>	
505	<p>እርስዎ ወይም የቤተሰብዎ አባል ታመው በነበሩበት ጊዜ የፀረ-ወባ መድሃኒቶችን በአልኮል መጠጦች (ጠላ፣አረቄ፣ቢራ ወዘተ) ውጠው ወይም ወስደው ያውቃሉ?</p>	<p>አዎ 01 አላውቅም 02</p>	<p>[] []</p>	
<p>ክፍል-VI: በቤት ውስጥ የጸረ-ወባ መድሃኒት ማከማቸትን ለማወቅ የቀረቡ ጥያቄዎች</p>				
601	<p>አሁን በዚህ ሰዓት ቤትዎ ውስጥ ፀረ-ወባ መድሃኒት አለ?</p>	<p>አዎ 01 የለም 02 [] []</p>	<p>[] []</p>	<p>መልስዎ 02 ከሆነ በቀጥታ ወደ ጥያቄ 701 ይሂዱ.</p>
602	<p>ፀረ-ወባ መድሃኒቱን ሊያሳዩኝ ይችላሉን?</p>	<p>አዎ 01 አይቻልም 02 [] []</p>	<p>[] []</p>	<p>መልስዎ 02 ከሆነ በቀጥታ ወደ ጥያቄ 701 ይሂዱ.</p>
603	<p>የፀረ-ወባ መድሃኒቶቹን</p>	<p>----- (ሌብሉ የማይነበብ)</p>		

	ስም ይጻፉ	ከሆነ በባዶ ቦታው ላይ የማይነበብ ብለው ይጻፉ)			
604	የፀረ-ወባመድሃኒቶች የተቀመጡበት ሁኔታ ምን ዓይነት ነው	በተቀደደ ማሸጊያ ውስጥ በደረቅ ቦታ 01 [] [] በደረቅ ቦታ ያለማሸጊያ 02 [] [] በሞቃታማ ቦታ ባልተቀደደ በማሸጊያ 03 [] [] በርጥበት ቦታ ባልተቀደደ በማሸጊያ 04 ባልተቀደደ ማሸጊያ ውስጥ በደረቅ ቦታ ባልተቀደደ ማሸጊያ ውስጥ የተቀመጠ 05 -----ሌላ ካለ ይጠቀስ 06			
ክፍል VII: ስለ ባህላዊ ፀረ-ወባ መድሃኒቶች ጠቅላላ ዕውቀትን ለመዳሰስ የቀረቡ ጥያቄዎች					
701	ባህላዊ የሆነ መድሃኒት □ ለወባ ህክምና የሚጠቀሙበት አለን?	አዎ 01 [] [] የለም 02		መልስዎ 02 ከሆነ አመስግነው ይሰናበቱ	
702	ምን ዓይነት የባህል ህክምና ያውቃሉ?	የአታክልት ክፍሎች (የእጽዋት ተዋፅኦ) ህክምና 01 የእንሰሳት ተዋፅኦ ህክምና 02 -----ሌላ ካለ ይጠቀስ 03	[] []		
703	በጥያቄ ቁጥር 702 መልስዎ 01 (የእጽዋት ተዋፅኦ የሆነ) ብለው ከሆነ የሚከተለውን ሳጥን ይሙሉ:- (ተጠያቂው የእጽዋት ተዋፅኦ የሆነ (01) ካላሉ ወደ ጥያቄ ቁጥር 704 ይሂዱ)				
ተ.ቁ	የእጽዋቱ ስም (የተክሉ ስም)	ጥቅም ላይ የሚውለው የተክሉ ክፍል ስም (ቅጠል፣ ስር፣ ግንድ ወዘተ)	አዘገጃጀቱ እንዴት ነው? (ያዘገጃጀቱ ሂደት ይጻፍ)	በቀን ስንት ጊዜ ነው ህክምናውን የሚከታተሉት?	ለምን ያህል ጊዜ ነው ህክምናውን የሚከታተሉት?

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704	በጥያቄ ቁጥር 702 መልስዎ 02 (የእንሰሳት ተዋፅኦ የሆነ) ብለው ከሆነ የሚከተለውን ሳጥን ይሙሉ። (የእንሰሳት ተዋፅኦ የሆነ (02) ካላሉ አመስግነው ይሰናበቱ)				
ተ.ቁ	የእንሰሳው ስም	ጥቅም ላይ የሚውለው እንሰሳት ተዋፅኦው ስም	አዘገጃጀቱ እንዴት ነው? (ያዘገጃጀቱ ሂደት ይጻፍ)	በቀን ስንት ጊዜ ነው ህክምናውን የሚከታተሉት?	ለምን ያህል ጊዜ ነው ህክምናውን የሚከታተሉት?
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2					
3					

በመጨረሻም ተጠያቂዎን አመስግነው ይሰናበቱ::

የተቆጣጣሪው ኮድ _____ ፊርማ _____

Annex 8: በቡድን ውይይት ለመሳተፍ የፍቃደኝነት መግለጫ ና የመወያያ ሃሳቦች

ሰላም ዋላችሁ/አደራችሁ እዚህ ስለተገኛችሁ አመሰግናለሁ

1. መግቢያ

እኔ ሰኢድ ሙሳ እባላለሁ በአዲስ አበባ ዩኒቨርሲቲ ፋርማሲ ት/ቤት ተማሪ ነኝ። እዚህ ከእኔ አጠገብ ያሉት _____ ይባላሉ የመጡትም ከ _____ ነው።

2. የውይይቱ ርዕስ

በቤተሰብ ደረጃ የጸረ-ወባ መድሃኒትና የአልጋ አጎበር አጠቃቀም ምን እንደሚመስል መወያየት

3. የጥናቱ ዓላማ

ይህ ጥናት (ውይይት) በቤተሰብ ደረጃ ስለ ወባ በሽታ ጠቅላላ ግንዛቤን የጸረ-ወባ መድሃኒት አጠቃቀምን፣ አመራረጥን፣ መድሃኒቱን በአግባቡ ካለመውሰድ ጋር ተያያዥ የሆኑ ጉዳዮችን ለመዳሰስና ጠቃሚ የሆኑ ከዚህ ውይይት የተሰበሰቡ መረጃዎችን ከላይ ያየናቸውን ችግሮች ለማስተካከል ይቻል ዘንድ ጠቃሚ የማሻሻያ መንገዶችን ለወረዳው አልፎም ለክልሉ ለመቀየስ ይረዳል።

ከዚህ ውይይት የሚገኘው መረጃ ከላይ የተገለጹትን ችግሮች ለማቃለል ለሚሰሩ ድርጅቶችም ጠቀሜታው የጎላ ነው።

4. እርስዎ በቡድን ውይይት እንዲሳተፍ እንዴት እንደተመረጡ

እናንተ በዚህ ጥናት እንድትሳተፍ የተመረጣችሁት ጥናቱን የሚያካሂደው ግለሰብ እናንተ እዚህ አካባቢ ለረጅም ጊዜ (≥ 0 ዓመት) ስለኖራችሁ ማህበረሰቡን በተለይም የአካባቢውን ሴቶች ትውክላላችሁ ብሎ ስለገመተና በማህበረሰቡ ውስጥ ያለውን የጸረ-ወባ መድሃኒትና የአልጋ አጎበር አጠቃቀም ልማድ ታውቃላችሁና እዚህ ታወጉናላችሁ ብሎ በመገመት ነው።

5. ሚስጥር ስለመጠበቅ

እርግጠኛ ሆኖ የምንነግራችሁ ነገር ቢኖር እናንተ እዚህ የምትነግሩኝ (የምታወጉኝ) ሀሳብ ሁሉ ሚስጥራዊ ሆኖ ይያዛል። ከዚህም በተጨማሪም የተቀረጸው ድምጽም ሀሳቡን (ፍሬ ነገሩን) ወደ ወረቀት በእርጋታ ከተገለበጠ በኋላ የተቀረጸው ድምጽ በሙሉ ይደመስሳል (እንዲጠፋ ይደረጋል)። ሌላው ላረጋግጥላችሁ የምወደው ስማችሁ ካወራችሁት (ከገለጣችሁት) ሀሳብ ጋር ተያያዥ ሆኖ ከዚህ ጥናት ውጤት ጋር አይገለጽም።

6. በውይይት ለመሳተፍና ድምጽ ለመቅረጽ ስምምነት

Annex 8: በቡድን ውይይት ለመሳተፍ የፍቃደኝነት መግለጫ ና የመወያያ ሃሳቦች

ሰላም ዋላችሁ/አደራችሁ እዚህ ስለተገኛችሁ አመሰግናለሁ

1. መግቢያ

እኔ ሰኢድ ሙሳ እባላለሁ በአዲስ አበባ ዩኒቨርሲቲ ፋርማሲ ት/ቤት ተማሪ ነኝ። እዚህ ከእኔ አጠገብ ያሉት _____ ይባላሉ የመጡትም ከ _____ ነው።

2. የውይይቱ ርዕስ

በቤተሰብ ደረጃ የጸረ-ወባ መድሃኒትና የአልጋ አጎበር አጠቃቀም ምን እንደሚመስል መወያየት

3. የጥናቱ ዓላማ

ይህ ጥናት (ውይይት) በቤተሰብ ደረጃ ስለ ወባ በሽታ ጠቅላላ ግንዛቤን የጸረ-ወባ መድሃኒት አጠቃቀምን፣ አመራረጥን፣ መድሃኒቱን በአግባቡ ካለመውሰድ ጋር ተያያዥ የሆኑ ጉዳዮችን ለመዳሰስና ጠቃሚ የሆኑ ከዚህ ውይይት የተሰበሰቡ መረጃዎችን ከላይ ያየናቸውን ችግሮች ለማስተካከል ይቻል ዘንድ ጠቃሚ የማሻሻያ መንገዶችን ለወረዳው አልፎም ለክልሉ ለመቀየስ ይረዳል።

ከዚህ ውይይት የሚገኘው መረጃ ከላይ የተገለጹትን ችግሮች ለማቃለል ለሚሰሩ ድርጅቶችም ጠቀሜታው የጎላ ነው።

4. እርስዎ በቡድን ውይይት እንዲሳተፍ እንዴት እንደተመረጡ

እናንተ በዚህ ጥናት እንድትሳተፍ የተመረጣችሁት ጥናቱን የሚያካሂደው ግለሰብ እናንተ እዚህ አካባቢ ለረጅም ጊዜ (≥0 ዓመት) ስለኖራችሁ ማህበረሰቡን በተለይም የአካባቢውን ሴቶች ትወክላላችሁ ብሎ ስለገመተና በማህበረሰቡ ውስጥ ያለውን የጸረ-ወባ መድሃኒትና የአልጋ አጎበር አጠቃቀም ልማድ ታውቃላችሁና እዚህ ታወጉናላችሁ ብሎ በመገመት ነው።

5. ሚስጥር ስለመጠበቅ

እርግጠኛ ሆኖ የምንነግራችሁ ነገር ቢኖር እናንተ እዚህ የምትነግሩኝ (የምታወጉኝ) ሀሳብ ሁሉ ሚስጥራዊ ሆኖ ይያዛል። ከዚህም በተጨማሪም የተቀረጸው ድምጽም ሀሳቡን (ፍሬ ነገሩን) ወደ ወረቀት በእርጋታ ከተገለበጠ በኋላ የተቀረጸው ድምጽ በሙሉ ይደመስሳል (እንዲጠፋ ይደረጋል)። ሌላው ላረጋግጥላችሁ የምወደው ስማችሁ ካወራችሁት (ከገለጣችሁት) ሃሳብ ጋር ተያያዥ ሆኖ ከዚህ ጥናት ውጤት ጋር አይገለጽም።

6. በውይይት ለመሳተፍና ድምጽ ለመቅረጽ ስምምነት

እርስዎ እዚህ ቦታ ከላይ ለተገለጸው ጉዳይ እንዲገኙልኝ በጠየቅኩዎ መሰረት እዚህ ቦታ መገኘትዎ በውይይት ለመሳተፍ ፈቃደኛ መሆንዎንና መስማማትዎን ያሳያል።

የእርስዎ በዚህ ውይይት ለመሳተፍ መወሰን በእርስዎ ፈቃደኝነት ላይ ብቻ የተመሰረተ ከመሆኑም በላይ ላለመሳተፍ ቢወስኑም ምንም የሚደርስብዎት ችግር የለም። በፈለጉበት ጊዜ ከፈለጉና መስሎ ከታየዎት ውይይቱን ማቋረጥ ይችላሉ።

ተቃውሞ ከሌለ ለጥንቃቄና ስራን ለማቀላጠፍ ሲባል ውይይቱን በቴፕ እንዲቀዳ (እንዲቀረጽ) እደርጋለሁ።

7. የቡድን ተሳትፎ ውይይት ጠቃሚነቱ

በዚህ ውይይት ስህተት ወይም ትክክል የሚባል መልስ የለም። ማንኛውም ተሳታፊ የመሰለውን የሚያውቀውን እና የሚያደርገውን ነገር መናገር ይችላል።

በዚህ ውይይት ማንኛውም ተሳታፊ ነፃነት እንዲሰማውና በነጻነት እንዲያወራ ያስፈልጋል። ማናችሁም እዚህ የተገኛችሁ የውይይቱ ተሳታፊዎች ሁሉ ስለ እያንዳንዱ የውይይት ርዕስ የመሰለውን የሚያውቀውንና የሚያደርገውን ሃሳብ እንዲሰጥ ይበረታታል።

እያንዳንዱ ተሳታፊ ሃሳቡን ሲገልጥ ስለ ወጣ፣ ፀረ-ወጣ መድሃኒት እና ስለ አልጋ አጎበር አጠቃቀም እውነታኛውን ነገር እንድትገልጹልን እናሳስባለን። በውይይቱ ጊዜ የምትሰጡት ሃሳብ ተሳታፊዎች በሚሰጡት አስተያየት ሳይሆን ከላይ በገለጽኩላችሁ የውይይት አላማዎች ላይ እንድታተኩሩ እጠይቃለሁ።

8. እርስ በርስ መተዋወቅ

ከዚህ በመቀጠል እያንዳንዱ የውይይቱ ተሳታፊ እራሱን እንዲያስተዋውቅ እጠይቃለሁ።

9. የመወያያ ጥያቄዎች

ውይይቱ የሚያነጥነው በሚከተሉት ሃሳቦች (ጥያቄዎች) ላይ ነው።

- እዚህ አካባቢ ወጣ ዋና የጤና ችግር ነውን?
- የወጣ መንስኤ ፣ መተላለፊያ መንገዶችና ወጣ የሚበዛባቸው ወቅቶች
- ለወጣ ተጋላጭና ተጠቂ የሆኑ የህብረተሰብ ክፍሎች
- የወጣ ዋና ዋና ምልክቶች
- የወጣ መከላከያ መንገዶች
- የአልጋ አጎበር፡- ጥቅሙ፣ ጉዳቱ፣ አጠቃቀም ወዘተ
- የወጣ ህክምና

- ስለ ዘመናዊ ጸረ-ወባ መድሃኒቶች ጠቅላላ ዕውቀት
- የጸረ-ወባ መድሃኒቶችን ከየት እንደሚያገኙና ተያያዥ ጉዳዮች
- ወባ ነው ተብሎ የሚገመት በሽታ ወይም የወባ ምልክቶች ሲታዩ የሚወሰዱ እርምጃዎች ወይም ህክምና
- የጸረ-ወባ መድሃኒትን ቤት ውስጥ ገዝቶ ሰው ሲታመም ለመስጠት ማስቀመጥ (ማከማቸት) እና ተያያዥ ጉዳዮች
- ባህላዊ የጸረ-ወባ መድሃኒቶች

እስካሁን ስለ ወባ፣ ፀረ-ወባ መድሃኒት እና ስለ አልጋ አጎበር አጠቃቀም ወዘተ ተወያይተናል። ጊዜያቸውን መስዋእት አድርጋችሁ እዚህ ተገኝታችሁ ለሰጣችሁን አስተያየትና ጥቆማ እንዲሁም ላደረጋችሁልን ትብብር በጣም እናመሰግናለን። እዚህ ላይ ይኸ ቀረ የምትሉት ካለ ውይይታችንን ከማጠናቀቃችን በፊት ልታነሱትና ልንወያይበት እንችላለን።

በመጨረሻም በጣም እናመሰግናለን

ውይይቱ ተጠናቋል።

የቡድን ውይይት ተሳታፊዎች የግል ና ሌሎች አስፈላጊ መረጃዎች

ቀን _____

እድሜ _____

የጋብቻ ሁኔታ _____

ሀይማኖት _____

የት/ት ደረጃ _____

ስራ _____

የተሳታፊዎች ብዛት _____

ውይይቱ የፈጀው ጊዜ _____

Annex 9: በቃለመጠይቅ ለመሳተፍ የፍቃደኝነት መግለጫ ና የመወያያ ሃሳቦች

ሰላም ዋላችሁ/አደራችሁ እዚህ ስላተገኙልኝ አመሰግናለሁ

1. መግቢያ

እኔ ሰኢድ ሙሳ እባላለሁ በአዲስ አበባ ዩኒቨርሲቲ ፋርማሲ ት/ቤት ተማሪ ነኝ።

2. የውይይቱ ርዕስ

በፖዌ ልዩ ወረዳ በቤተሰብ ደረጃ የፀረ-ወጣ መድሃኒትና የአልጋ አጎበርን አጠቃቀም ምን እንደሚመስል መዳሰስ

3. የጥናቱ አላማ

ይህ ውይይት በቤተሰብ ደረጃ ስለ ወጣ በሽታ ጠቅላላ ግንዛቤን፣ የፀረ- ወጣ በሽታ መድሃኒት አጠቃቀምን፣ አመራረጥን፣ መድሃኑን በአግባቡ ካለመውሰድ ጋ ተያያዥ የሆኑ ጉዳዮችን እንዲሁም የአልጋ አጎበር አጠቃቀምን እና ከአልጋ አጎበር ጋ የተያያዙ ችግሮችን ለመዳሰስ ና ጠቃሚ የሆኑ ከዚህ ውይይት የተሰበሰቡ መረጃዎች እነዚህን ችግሮች ለማስተካከል ይቻል ዘንድ ጠቃሚ የማሻሻያ መንገዶችን ለወረዳው አልፎም ለክልሉ ለመቀየስ ይረዳል።

ከዚህ ውይይት የሚገኘው መረጃ ከላይ የተገለጹትን ችግሮች ለማቃለል ለሚሰሩ ድርጅቶችም ጠቀሜታው የጎላ ነው።

4. እርስዎ በቃለመጠይቁ እንዲሳተፉ እንዴት እንደተመረጡ

እርስዎ በዚህ ጥናት ውስጥ በቃለ መጠይቁ እንዲሳተፉ የተመረጡት በጤና ተቋም ውስጥ በጤና ባለሙያነት ስለሚሰሩ፣ በወረደው ጤና ቢሮ በጋላፊነት ስለሚሰሩ፣ የሃገር ሽማግሌ ወዘተ. ስለሆኑ ጥናቱን የሚያካሂደው ግለሰብ እርስዎ ማህበረሰቡን ያውቃሉ ብሎ ስለገመተና በማህበረሰቡ ውስጥ ያለውን የመድሃኒትና የአልጋ አጎበር አጠቃቀም ልማድ እዚህም ይነግሩኛል ብሎ በማሰብ ነው።

5. ሚስጥር ስለመጠበቅ

እርግጠኛ ሆኜ የምንገርዎት ነገር ቢኖር እርስዎ እዚህ የሚነግሩኝ ሀሳብ ሁሉ ሚስጥራዊ ሆኖ ይያዛል። ከዚህ በተጨማሪም የተቀረጸው ድምጽ ሀሳቡን (ፍሬ ነገሩን) ወደ ወረቀት በእርጋታ ከገለበጥኩ በኋላ፣ የተቀረጸው ድምጽ በሙሉ ይደመሰሳል (እንዲጠፋ ይደረጋል)። ሌላው ላረጋግጥሎት የምወደው ነገር ስምዎ ካወሩኝ(ከገለጡ ልኝ) ሃሳብ ጋር ተያያዥ ሆኖ ከዚህ ጥናት ውጤት ጋር አይገለጽም።

6. በቃለመጠይቁ ለመሳተፍና ድምጽ ለመቅረጽ ስምምነት

እርስዎ እዚህ ቦታ ከላይ ለተገለጸው ጉዳይ እንዲገኙልኝ በጠየቅኩዎ መሰረት እዚህ ቦታ መገኘትዎ በቃለመጠይቁ ለመሳተፍ ፍቃደኛ መሆንዎንና መስማማትዎን ያሳያል።

የእርስዎ በዚህ ቃለመጠይቅ ለመሳተፍ መወሰን በእርስዎ ፍቃደኝነት ላይ የተመሰረተ ከመሆኑም በላይ ላለመሳተፍ ቢወስኑ ምንም የሚደርስቦት ችግር የለም። በፈለጉበት ጊዜ መስሎ ከታይዎት ውይይቱን ማቋረጥ ይችላሉ።

ተቃውሞ ከሌለዎት ለጥንቃቄ እና ሥራን ለማቀላጠፍ ሲባል ቃለመጠይቁን በቴፕ እንዲቀዳ (እንዲቀረጽ) አደርጋለሁ።

7. በቃለ መጠይቁ ጊዜ የሚነሱ ጥያቄዎች

ቃለ መጠይቁ የሚያጠነጥነው በሚከተሉት ሀሳቦች ላይ ነው።

- እዚህ አካባቢ ወባ ዋና የጤና ችግር ነውን?
- የወባ መንስኤ፣ መተላለፊያ መንገዶች እና ወባ የሚበዛባቸው ወቅቶች
- ለወባ ተጋላጭ እና ተጠቂ የሆኑ የህብረተሰብ ክፍሎች
- የወባ ዋና ዋና ምልክቶች
- የወባ መከላከያ መንገዶች
- የአልጋ አጎበር ጥቅሙ ፣ ጉዳቱ፣ አጠቃቀም ወዘተ.
- የወባ ህክምና
- ስለዘመናዊ የወባ መድሃኖች ጠቅላላ እውቀት (መድሃኒቶችን በማሳየት)
- የጸረ-ወባ መድሃኒቶችን ከየት እንደሚያገኙባቸውና ተያያዥ የሆኑ ጉዳዮች
- ወባ ነው ተብሎ የሚገመት በሽታ ወይም የወባ ምልክቶች ሲታዩ የሚወሰዱ እርምጃዎች ወይም ህክምና
- የፀረ-ወባ መድሃኒትን ቤት ውስጥ ገዝቶ ሰው ሲታመም ለመስጠት ማስቀመጥ (ማከማቸት) እና ተያያዥ ጉዳዮች
- ባህላዊ ፀረ-ወባ መድሃኒቶች
- እኤአ በ2004 የጤና ጥበቃ ሚኒስቴ ለፋርሲፋርም ወባ ህክምና የሚውል አዲስ መድሃኒት (ኳርተም የሚባል) በፋንሲደር ምትክ ጥቅም ላይ እንዲውል ማድረግ ይታወቃል። ለመሆኑ ለሰው ምክንያቱ ምን ነበር?(በተለይ ለጤና ባለሙያዎች የሚቀርብ የመወያያ ጥያቄ)

እስካሁን ስለወባ፣ ፀረ-ወባ መድሃኒት፣ ስለ አልጋ አጎበር አጠቃቀም ወዘተ ተወያይተናል። ጊዜዎትን መስዋዕት አድርገው እዚህ ተገኝተው ለሰውኝ አስተያየት እና ጥቆማ እንዲሁም ላደረጉልኝ ትብብር

በጣም አመሰግናሁ። እዚህ ላይ ቃለ መጠይቃችንን ከማጠናቀቃችን በፊት ይኸ ቀረ የሚሉት ካለ ሊያነሱትና ልንነጋገርበት እንችላል።

የቃለ መጠይቅ ተሳታፊዎች የግልና ሌሎች አስፈላጊ መረጃዎች

- ቀን _____
- እድሜ _____
- የጋብቻ ሁኔታ _____
- ሀይማኖት _____
- የት/ት ደረጃ _____
- ስራ _____
- ቃለ መጠይቁ የፈጀው ጊዜ _____

በጣም አመሰግናሁ። እዚህ ላይ ቃለ መጠይቃችንን ከማጠናቀቃችን በፊት ይኸ ቀረ የሚሉት ካለ ሊያነሱትና ልንነጋገርበት እንችላል።

የቃለ መጠይቅ ተሳታፊዎች የግልና ሌሎች አስፈላጊ መረጃዎች

ቀን _____

እድሜ _____

የጋብቻ ሁኔታ _____

ሀይማኖት _____

የት/ት ደረጃ _____

ስራ _____

ቃለ መጠይቁ የፈጀው ጊዜ _____

I, the undersigned, declare that this thesis is my original work and has not been prese
degree in any other university.

Name: Seid Mussa Ahmed

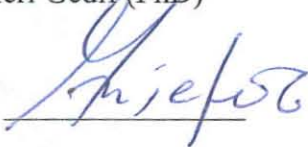
Signature: _____



This thesis has been submitted for examination with my approval as university advisor

Name: Teferi Gedif (PhD)

Signature: _____



Place and date of submission: Addis Ababa, Ethiopia

July 2009