

ORIGINAL ARTICLE**Five Years Malaria Trend Analysis in Woreta Health Center, Northwest Ethiopia****Awoke Derby^{1*}, Megbaru Alemu¹****OPEN ACCESS**

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ABSTRACT

BACKGROUND: An estimated 68% of the Ethiopian population, living in 75% of the landmass, is at risk of contracting malaria at any time making it the leading public health problem. The temporal analysis of malaria data could be important to evaluate the performance of malaria prevention programmes. Thus, the aim of this study was to determine the trend of malaria at Woreta Health Center (WHC) over a period of five years.

METHODS: We analyzed the records of 8,057 presumptive malaria patients registered in 2012 to 2016. The following patient data were retrieved from laboratory registration logbook for analysis: sex, age, residence, blood film (BF) microscopy result, type of malaria parasite identified, year and month when the patients visited WHC. Logistic regression was employed to assess the association between potential associated factors and positive BF result; $p < 0.05$ was considered significant.

RESULTS: Among the total presumptive individuals, 4447(55.2%) were females. The prevalence of malaria in each year ranged from 4.1% to 6.7%. The overall prevalence of malaria was 5.4% (95%CI: 4.9%-5.9%). The two most important species of malaria parasite identified were *P. falciparum* at 233(53.7%) and *P. vivax* at 184(42.4%). Relatively higher proportions of cases were documented in the months of November, December and June (11.1%, 8.1% and 7.2%, respectively). Patients who visited the health center in the month of December were >4 times more likely to be infected as compared with those who came to the health center in September [AOR: 4.2, 95%CI (2.374-7.560)]. Females were 1.3 times more likely to be infected than males, [AOR: 1.3, 95%CI (1.101-1.638)]. Similarly, patients in the age group above 15 were 1.9 times more likely to be infected than individuals < 5, [AOR: 1.9 95%CI (1.498-2.455), p value 0.000].

CONCLUSION: In the studied area, malaria remains a major public health challenge. Hence, interventions to decrease the impact of the disease have to be evaluated and strengthened.

KEYWORDS: Malaria, trend analysis, Ethiopia

INTRODUCTION

Malaria is one of the major public health problems around the world. Despite being preventable and treatable, malaria continues to have a devastating impact on people's health and livelihoods around the world (1-5). According to the latest available data, about 3.2 billion people, nearly half the world's population, remained at risk of malaria in 97 countries, territories and areas in 2013, and an estimated 198 million cases occurred. In the same year, the disease killed about 584,000 people, mostly children aged under 5 years, in sub-Saharan Africa (6). In 2015 alone, there were 214 million new cases of the disease and more than 400,000 malaria-related deaths around the globe (5). More than 90% of clinical cases and death due to malaria occur in Africa (7). Of those Africans who die from malaria each year, most are children under 5 years of age (8) and pregnant women due to low level of immunity (4). In 2015, the African region accounted for approximately 9 in 10 malaria cases and deaths globally (5).

About 75% of the landmass of Ethiopia is malarious, an estimated 68% (~52million) of the population are at risk of contracting malaria (9,10). According to the world malaria report of the year 2016, the number of people living in high transmission (>1 case per 1000 population) and low transmission (0-1 cases per 1000 population) areas in Ethiopia was estimated to be 27,000,000 (27%) and 40,600,000 (41%), respectively. According to this report, there were 662 reported deaths due to malaria in the country (11). The exact number of people getting sick with and dying of malaria every year in Ethiopia is not well known due to poor documentation. However, it is known that millions of people get sick and tens of thousands die due to malaria every year, and that rates of mortality and morbidity dramatically increase during epidemics (9,10). *Plasmodium falciparum* and *P. vivax* are the two most dominant malaria parasites in Ethiopia (12,13). They are prevalent in all malaria endemic areas in the country with *P. falciparum* representing about 65-75% of the total reported malaria cases, relative frequency varying in time and space within given geographical ranges. *P. malariae* and

P. ovale are rare and account for <1% of all confirmed malaria cases. According to the world malaria report of the year 2016, the predominant Plasmodium species identified in Ethiopia were: *P. falciparum* (64%) and *P. vivax* (36%) (11). The major malaria vector incriminated in Ethiopia is *Anopheles arabiensis* (12) with *A. funestus*, *A. pharoensis* and *A. nili* being secondary vectors (13).

The distribution of malaria in Ethiopia is not uniform (9) and is seasonal in most parts of the country with variable transmission and prevalence patterns affected by the large diversity in altitude, rainfall and population movement. In most parts of Ethiopia, the major transmission periods of malaria occurs from September to December, following the main rainy seasons (June-September). The minor one is from March to May, following small showers of rain in autumn (12, 14). Since peak malaria transmission often coincides with the planting and harvesting season, and the majority of malaria burden is among older children and working adults in rural agricultural areas, there is a heavy economic burden in Ethiopia (13).

Ethiopia has achieved a remarkable progress in the fight against malaria during the most recent decade through strong preventive and case management interventions with large engagement of the health extension workers (HEWs) and the health development army (HAD) volunteers providing community based care at the household level. The country is also one of the few sub-Saharan African countries that have shown progress in the fight against malaria (15). Because of the inadequacy of malaria case data from many sub-Saharan African countries including Ethiopia (16), health facility based prevalence studies, like this study, can be used to enhance understanding of the level of malaria and how it is changed over time. The findings would also help policy makers to assess their malaria prevention strategies and their degree of interventions. Moreover, the epidemiological picture of malaria is not yet determined in the study area. Hence, the aim of this study was to determine the trend of malaria at Woreta Health Center (WHC), Northwest Ethiopia, over a period of five years.

MATERIALS AND METHODS

Study design, setting and data collection: In this study, the authors analyzed the records of 8,057 clinically suspected malaria patients who had blood film (BF) examinations for malaria parasites. The study was conducted at WHC. Woreta is a town in Northwestern Ethiopia. According to figures from the Central Statistical Agency of Ethiopia in 2005, the town had an estimated total population of 26,317. The town is located approximately 610 km Northwest of Addis Ababa, the Capital of Ethiopia, having an elevation of about 1828 meters above sea level. The town is wet land with annual temperature range of 18-28°C.

All patients who visited the health center from 1 September 2011 to 30 August 2016 and had the following registered data were included for analysis: sex, age, residence, BF microscopy result (+, -), type of malaria parasite identified (*P. falciparum*, *P. vivax* or mixed), year and month when the patients visited the health center. Patients were suspected for malaria when they presented with fever and related clinical symptoms. Data were retrieved directly from laboratory registration logbook using prepared data extraction sheet.

Blood film microscopy procedure: Microscopic examination remains the "gold standard" method for laboratory confirmation of malaria in developing nations, like Ethiopia. The Woreta Health Center is using BF as a standard procedure to diagnose malaria among suspected patients. Capillary blood samples were collected from each presumptive patient for microscopic examination. The finger tip or heel of the child patient was cleaned with swab moistened with 70% v/v alcohol. After air-drying, a sterile lancet was used to prick the finger or heel, and then squeezed gently to obtain the blood (17). Two blood samples for thick and thin films were prepared on clean slides from each participant according to the standard WHO approved protocol. Slides were labeled properly and air-dried horizontally on a slide tray. Thin films were fixed with absolute methanol immediately after drying, and both thin

and thick blood films were stained with 3% Giemsa solution for 30 minutes. During examination, blood slides were read as either negative, *P. falciparum* positive, *P. vivax* positive, or mixed infection. Two hundred fields (the equivalent of 0.5µl of thick blood film) were examined at a magnification of 1000x before identifying a slide as negative. If positive, the thin film was read to determine the species (17,18).

Data analysis: All data were entered, cleaned and analyzed using Statistical Package for Social Science (SPSS) software (*IBM Corp. Released 2011. IBM SPSS Statistics for Windows, Version 23.0. Armonk, NY: IBM Corp.*). Descriptive statistics, like proportion and mean, was used to present data. Logistic regression was employed to assess the association between potential predictors and malaria infection. *P* value less than 0.05 was considered to indicate statistically significant difference.

Ethical approval and consent: Permission and consent from WHC administrators was obtained to use the data for the research purpose. No patient details that may link to the patient identity, like names, was used and confidentially was maintained.

RESULTS

Demographic characteristics of study participants: The authors assessed BF microscopic results of 8,057 malaria suspected patients from laboratory reporting logbook processed over a 5 years' period in WHC. Of the total patients registered, 4447(55.2%) were females. The median age of the patients was 25 years, ranging from 1 month to 85 years. Among the participants, 5062 (62.8%) and 6176 (76.7%) were from rural settings and in the age group of above 15 years, respectively (Table 1).

Prevalence of malaria and its trend: The number of clinically suspected malaria patients who visited the WHC over the five years' period was comparable, ranging from 1473(17.5%) in 2012 to 1766(21.9%) in 2015 (Figure 1).

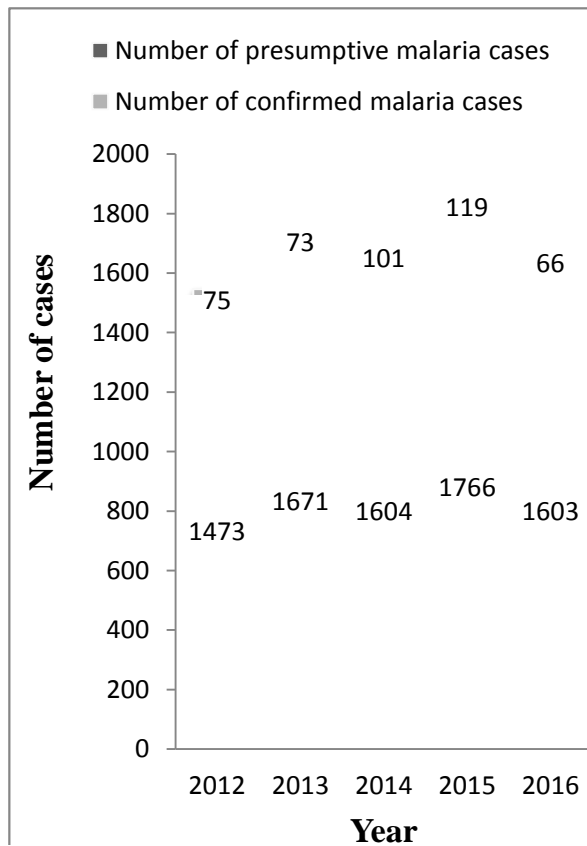


Figure 1: Number of BF confirmed and presumptive malaria cases over the five years period at WHC, 2012-2016

Similarly, the prevalence of BF confirmed malaria cases in each year ranged from 4.1% in 2016 to 6.3% in 2014 (Figure 2). The overall prevalence of malaria among suspected cases was 5.4% (434/8057, 95%CI: 4.9%-5.9%) over the five years' period. When we look at the annual distribution of malaria in the five years' period, relatively higher proportions of confirmed cases were documented in the months of November, December and June at 11.1%, 8.1% and 7.2%, respectively (Figure 3). The two most important species of malaria identified were *P. falciparum*, 233(53.7%, 95%CI: 49%-58.3%), and *P. vivax*, 184(42.4%, 95%CI: 37.8%-47.1%). There were 17(3.9%, 95%CI: 2.5%-6.6%) reports of mixed infection with both species (Table 1).

Table 1: Socio-demographic characteristics of the study participants and distribution of malaria at WHC, 2012-2016.

Variables	n (%)
Sex	
Male	3610 (44.8)
Female	4447 (55.2)
Total	8057 (100)
Age category	
<5	822 (10.2)
5-15	1059 (13.1)
>15	6176 (76.7)
Mean age 26.1 years	
SD 15.4 years	
Median age 25.0 years	
Range: 1 month to 85 years	
Residence	
Rural	5062 (62.8)
Urban	2995 (37.2)
BF result	
Positive	434 (5.4)
Negative	7623(94.6)
Identified malaria parasites	
<i>P. falciparum</i>	233 (53.7)
<i>P. vivax</i>	184 (42.4)
Mixed	17 (3.9)
Month	
January	605 (7.5)
Feb	585 (7.3)
Mar	467 (5.8)
Apr	552 (6.9)
May	668 (8.3)
Jun	846 (10.5)
Jul	504 (6.3)
Aug	492 (6.1)
Sep	885 (11.0)
Oct	995 (12.3)
Nov	660 (8.2)
Dec	798 (9.9)
Year	
2012	1473 (17.5)
2013	1671 (20.7)
2014	1604 (19.9)
2015	1766 (21.9)
2016	1603 (19.9)

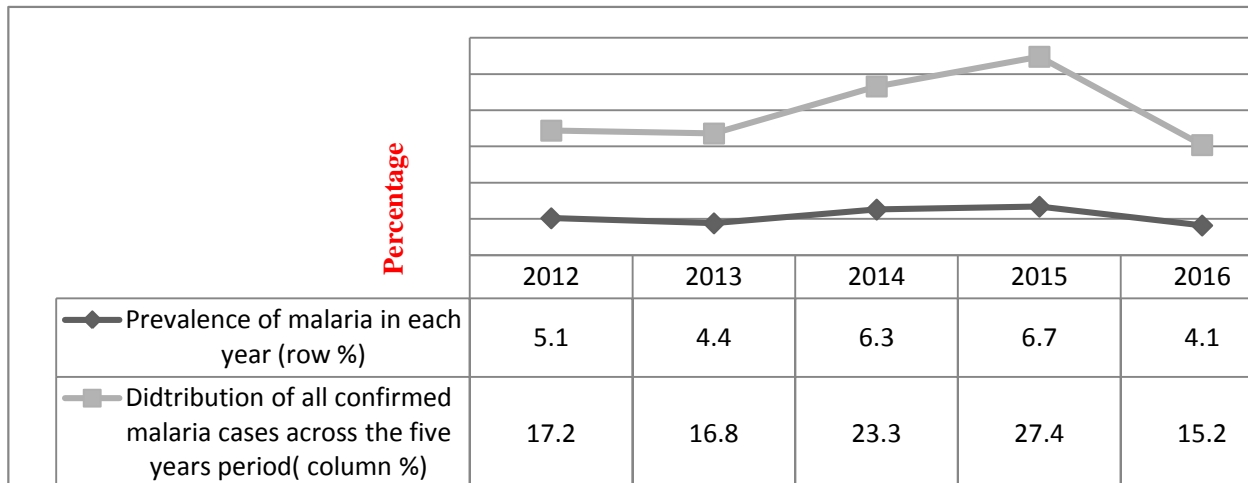


Figure 2: Prevalence of malaria over the five years period at WHC, 2012-2016.

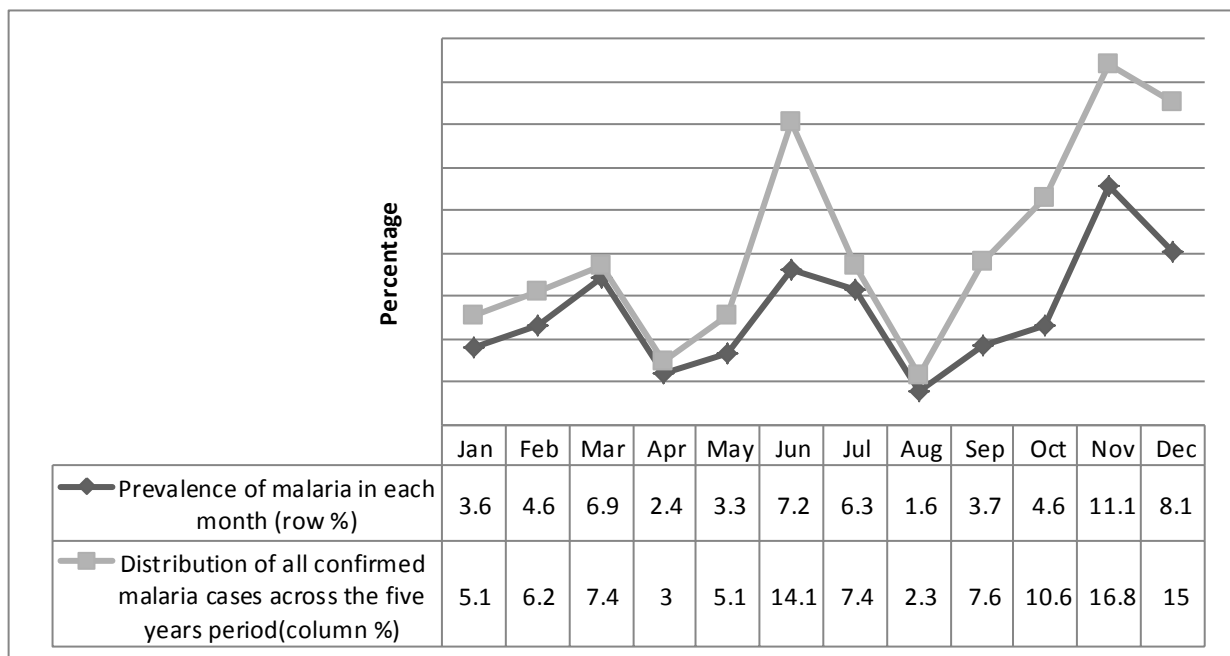


Figure 3: Annual distribution of malaria over the five years period at WHC, 2012-2016.

Factors associated with BF confirmed malaria:

In the logistic regression model, it was found that females were 1.3 times more likely to be infected with malaria than males, [AOR: 1.3, 95%CI (1.101-1.638), *p* value 0.004]. Similarly, patients in the age group above of 15 were 1.9 times more likely to be positive for malaria than individuals under the age of 5, [AOR: 1.9 95%CI (1.498-

2.455), *p* value 0.000]. Moreover, patients who visited the health center in 2016 were 1.6 times more likely to be infected than those who came in 2012 to the center, [AOR: 1.6, 95%CI (1.133-2.214), *p* value 0.007]. On top of this, patients who visited the WHC in the month of December were more than 4 times more likely to be infected as compared with those who came in September

[AOR: 4.2, 95%CI (2.374-7.560), *p* value 0.000]. The distribution of malaria also showed a significant difference in the months of November,

January, April, July and August (*p* value <0.05) (Table 2).

Table 2: Logistic regression analysis of factors associated with BF confirmed malaria cases among suspected cases in WHC, 2012-2016.

Variables	n (%) of Bf positives	n (%) of BF negatives	<i>COR (95%CI)</i>	<i>AOR (95%CI), p value</i>
Sex				
Male	232 (6.4)	3378 (93.6)	1.0	1.0
Female	202 (4.5)	4245 (95.5)	1.4 (1.189-1.752)	1.3 (1.101-1.638), 0.004
Age group in years				
<5	47 (5.7)	775 (94.3)	1.0	1.0
5-15	96 (6.3)	963 (93.7)	0.6 (0.424-0.873)	1.2 (0.845-1.606), 0.352
>15	291 (4.7)	5885 (95.3)	1.2 (0.893-1.684)	1.9 (1.498-2.455), 0.000
Residence				
Urban	141 (4.7)	2854 (95.3)	1.0	1.0
Rural	293 (5.8)	4769 (94.2)	0.8 (0.653-0.987)	0.8 (0.660-1.003), 0.053
Month				
Sep	33 (3.7)	852 (96.3)	1.0	1.0
Oct	46 (4.6)	949 (95.4)	0.8 (0.506-1.261)	0.5 (0.224-1.288), 0.164
Nov	73 (11.1)	587 (88.9)	0.3 (0.204-0.476)	1.9 (1.027-3.535), 0.041
Dec	65 (8.1)	733 (91.9)	0.4 (0.284-0.671)	4.2 (2.374-7.560), 0.000
Jan	22 (3.6)	583 (96.4)	1.0 (0.592-1.778)	3.3 (1.871-5.717), 0.000
Feb	27 (4.6)	558 (95.4)	0.8 (0.476-1.346)	1.4 (0.705-2.816), 0.332
Mar	32 (6.9)	435 (93.1)	0.5 (0.319-0.868)	1.6 (0.840-3.206), 0.174
Apr	13 (2.4)	539 (97.6)	0.3 (0.134-0.494)	2.0 (1.122-3.543), 0.019
May	22 (3.3)	646 (96.7)	0.5 (0.224-0.896)	0.8 (0.385-1.559), 0.474
Jun	61 (7.2)	785 (92.8)	0.2 (0.107-0.360)	1.5 (0.782-2.822), 0.226
Jul	32 (6.3)	472 (93.7)	0.2 (0.117-0.433)	2.6 (1.448-4.508), 0.001
Aug	8 (1.6)	484 (98.4)	0.9 (0.380-2.243)	2.2 (1.148-4.275), 0.018
Year				
2012	75 (5.1)	1338 (94.9)	1.0	1.0
2013	73 (4.4)	1598 (95.6)	1.2 (0.882-1.707)	1.6 (0.980-2.722), 0.059
2014	101 (6.3)	1503 (93.7)	0.8 (0.613-1.135)	1.0 (0.965-1.459), 0.971
2015	119 (6.7)	1647 (93.3)	0.8 (0.576-1.045)	1.2 (0.817-1.811), 0.335
2016	66 (4.1)	1537 (95.9)	1.3 (0.930-1.832)	1.6 (1.133-2.214), 0.007

DISCUSSION

Malaria is the world's deadliest mosquito-borne disease. The African continent continues to bear the greatest burden of malaria and the greatest diversity of parasites, mosquito vectors and human victims (19). Ethiopia's complex topography and seasonal rainfall largely support its transmission, and it makes one of the public health challenges in the country.

This study area is one of the potentially malarious localities in Ethiopia. The number of malaria suspected individuals and tested in WHC over the five years' period ranged from 1473 (17.5%) in 2012 to 1766 (21.9%) in 2015. Moreover, the prevalence of confirmed malaria cases in each year ranged from 4.1% in 2016 to 6.7% in 2015, making the overall prevalence of malaria 5.4%, (95%CI: 4.9% - 5.9%). According to the world malaria report of 2016, the number of malaria

suspected and tested individuals in public health facilities of Ethiopia showed increment from about 9% in 2005 to 38% in 2015. However, the prevalence of BF confirmed malaria among these suspected cases over these periods ranged from 1% to 4%; the highest, (4%), was recorded in the year 2013 (11). The finding of this study is in agreement with the nationwide estimate although a minor difference was observed which might be related to the difference in sample size. Similarly, according to the Ethiopian 2011 malaria indicator survey, malaria prevalence was reported to be 1.3% (13,15) which makes our finding a bit higher than this report. However, other similar studies in Ethiopia demonstrated 14.8-25.6% (20,21) prevalence which is quite higher than our finding and the national estimate. The disparity might be related to the difference in geographical location, season and the employed sample size.

The two most important species of malaria identified in this study were *P. falciparum* at 233 (53.7%) and *P. vivax* at 184 (42.4%). There were 17(3.9%) reports of mixed infections. This result was comparable with the national prevalence study where about 57% and 43% of malaria cases in Ethiopia are accounted to *P. falciparum* and *P. vivax*, respectively (1). Similar findings were also reported in other studies in Ethiopia (20,22). Furthermore, different literatures indicated that *P. falciparum* is responsible for the most common form of malaria in Ethiopia (15). According to the WHO malaria report (7), *P. falciparum* is most prevalent in the African continent, and is responsible for most deaths. The report also indicated that *P. vivax* accounts for about 38% of the reported cases in Ethiopia which is concurring with the finding of this study.

In this study, the annual distribution of malaria in the five years' period showed that higher proportions of confirmed cases were documented in the months of November, December and June at 11.1%, 8.1% and 7.2%, respectively. It was also found that patients who visited the health center in the month of December were more than 4 times likely to be malaria infected as compared with those who came in the first month of Ethiopian calendar, September, [AOR: 4.2, 95% CI (2.374-7.560), *p* value 0.000].

The distribution of malaria also showed a significant difference in the month of November, January, April, July and August (*p* value <0.05). According to different reports, in most parts of the country, the peak periods of malaria transmission occur from September to November, following the main rainy seasons, and from May to June, following the small rainy season (12,14,15). This makes our report consistent with the national report. A similar finding was also reported by Yewhalaw *et al.* in 2013 in the country (22). The unstable malaria transmission patterns make Ethiopia prone to focal and multifocal epidemics that have on occasion caused catastrophic public health emergencies (15). The prevalence of malaria over the five years' period in the present study did not show typical trend/direction, instead there were ups and downs ranging from 4.1% to 6.3% which might indicate inconsistent intervention measures taken to reduce the burden of the disease in Ethiopia.

In this study, it was found that females were 1.3 times more likely to be positive for malaria than males [AOR: 1.3, 95%CI (1.101-1.638), *p* value 0.004]. This difference may be related to their relative low immune status compared with males. In this case, this finding is in line with the reports of the world malaria report (16). Similarly, it was found that patients in the age group above 15 were 1.9 times more likely to be infected with malaria than individuals under the age of 5, [AOR: 1.9 95%CI (1.498-2.455), *p* value 0.000]. This might be related with their frequent outdoor activities.

Due to its retrospective nature, the study lacks detailed clinical picture of patients which might have a determinant factor to show the complete picture of the study subjects. This calls for proper documentation of patient data. It would also be useful to know the percentage of malaria infected people and who were not presenting at the health center with suspected malaria. However, the report of our study will be an important source of data that will indirectly provide information concerning the implementation of malaria prevention and control measures in the study area where there is a high burden of the disease.

In conclusion, despite the many efforts that

are in place to control it, malaria remains one of the most important causes of morbidity in our study area. The predominant identified species was *P. falciparum*. Thus, concerned stakeholders should strengthen sustainable malaria prevention and control measures to restrain the problem.

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REFERENCES

- World Health Organization. World Malaria Report 2005. 20 Avenue Appia, 1211 Geneva 27, Switzerland. Available at: <http://apps.who.int/iris/bitstream/10665/43213/1/9241593199_eng.pdf>. Accessed on 27 Dec 2016.
- Rowe K, Rowe Y, Snow W, *et al.* The burden of malaria mortality among African children in the year 2000. *Int Journal of Epidemiol* 2006; 35:691–704.
- Ghebreyesus I, Deresa W, Written KH, *et al.* The epidemiology and Ecology of health and death in Ethiopia. Addis Ababa *Ethiopia J-Health Deve* 2007; 21(2).
- Grant A, Roussillon C, Paul R, Sakuntabhai A. The genetic control of immunity to Plasmodium infection. *BMC Immunology* 2015; 16:14.
- World Health Organization. Eliminating malaria, WHO/HTM/GMP/2016.3; 20 Avenue Appia, 1211 Geneva 27, Switzerland. Available at: <http://www.who.int/about/licensing/copyright_for_m/en/index.html>. Accessed on 1 Jan 2017.
- World Health Organization. Global technical strategy for malaria 2016-2030; 20 Avenue Appia, 1211 Geneva 27, Switzerland. Available at: <www.who.int/about/licensing/copyright_form/en/index.html>. Accessed on 1 Jan 2017.
- World Health Organization; WHO global malaria program, World malaria report 2014. Available at <www.who.int/malaria>. Accessed on 15 Dec 2016.
- Lengrar C. Insecticide treated nets for malaria control real gains. Bulletin of WHO, Geneva Switzerland, World Health Organization 2004; 82-84.
- Epidemiology and distribution of malaria in Ethiopia. Available at: <<http://moodle.digital-campus.org/mod/page/view.php?id=14803>>. Accessed on 1 Jan 2017.
- FDRE, MOH. Malaria prevention and control programs. Available at: <http://www.moh.gov.et/malaria?p_p_auth=Ws9d2X5o&p_p_id=77&p_p_lifecycle=0&p_p_state=maximized&p_p_mode=view&_77_struts_action=%2Fjournal_content_search%2Fsearch>. Accessed on; 4 Jan 2017
- World Health Organization. World Malaria Report 2016. Regional Profile. Available at: <<http://www.who.int/malaria/publications/world-malaria-report-2016/WMR-2016-regional-profiles.pdf?ua=1>>. Accessed on 4 Jan 2017.
- FDRE, MoH. Malaria Epidemiological profile. Available at: <<http://www.moh.gov.et/malaria>>. Accessed; on 4 Jan 2017.
- President's malaria initiative. Ethiopia-malaria operational plan FY 2016.
- African Union update on malaria control in Africa. Special summit of Africa Union on HIV/AIDs, TB and malaria (ATM): Abuja, Nigeria 2006.
- The Ethiopian Health and Nutrition Research Institute & partners. Ethiopia National Malaria Indicator Survey; Addis Ababa Ethiopia 2011.
- WHO Global malaria programme. World Malaria Report 2014; 20 Avenue Appia, 1211 Geneva 27, Switzerland.
- Cheesbrough M. District laboratory practice in tropical countries. Part one, Second Edition. Cambridge University 2009; p244-245.
- Tangpukdee N, Duangdee C, Wilairatana P, Krudsood S. Malaria Diagnosis: A Brief Review. *Korean J Parasitol* 2009; 47(2): 93.
- Ghansah A, Amenga-Etego L, Amambua-Ngwa A, *et al.* Monitoring parasite diversity for malaria elimination in sub-Saharan Africa. *Science*. 2014; 345(6202):1297-8.
- Yihenew G, Adamu H, Petros B. The impact of cooperative social organization on reducing the prevalence of malaria and intestinal parasite infections in Awramba, a rural community in South Gondar, Ethiopia. *Interdiscip Perspect Infect Dis* 2014;378780.
- Deressa W, Ali A, Enqusellassie F. "Self-treatment of malaria in rural communities, Butajira, southern Ethiopia," Bulletin of the World Health Organization, 2003; 81(4): 261.
- Yewhalaw D Getachew Y, Tushune K, *et al.* The effect of dams and seasons on malaria incidence and anopheles abundance in Ethiopia. *BMC Infect Di* 2013; 13: 161.