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Waning success: a 2013–2022 spatial and temporal trend analysis of malaria in Ethiopia

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Abstract

Background Despite consecutive decades of success in reducing malaria transmission, Ethiopia went off track towards its goal of malaria elimination by 2030, as outlined in the NMCP malaria strategy. Recent malaria outbreaks in Ethiopia are attributed to the emergence and spread of diagnostic and drug-resistant *Plasmodium falcipa-rum*, increased insecticide resistance in major vectors and the spread of invasive *Anopheles stephensi*. The effects of the COVID-19 pandemic, environmental anomalies and internal conflicts have also potentially played a role in increasing malaria transmission. This study aimed to evaluate the contribution of environmental factors and *An. stephensi* to the spatiotemporal trends of recent malaria cases in Ethiopia.

Methods Clinical malaria case data reported weekly between January 2013 and January 2023 were obtained from the Ethiopian Public Health Institute (EPHI), Addis Ababa. A negative binomial regression model was used to explain the variability and potential overdispersion in the weekly number of malaria cases reported across Ethiopian administrative zones. This model incorporated fixed effects for selected environmental factors and random effects to capture temporal trends, zone specific seasonal patterns, spatial trends at the zone level, and the presence of *An. stephensi* and its impact.

Results Our negative binomial regression model highlighted 56% variability in the data and slightly more than half (55%) was due to environmental factors, while the remainder was captured by random effects. A significant nationwide decline in malaria risk was observed between 2013 and 2018, followed by a sharp increase in early 2022. Malaria risk was higher in western and northwestern zones of Ethiopia compared to other zones. Zone-specific seasonal patterns, not explained by environmental factors, were grouped into four clusters of seasonal behaviours. The presence of *An. stephensi* was not shown to have any significant impact on malaria risk.

Conclusions Understanding the spatial and temporal drivers of malaria transmission and therefore identifying more appropriate malaria control strategies are key to the success of any malaria elimination and eradication programmes in Ethiopia. Our study found that approximately 50% of malaria risk variability could be explained by environmental,

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temporal, and spatial factors included in the analysis, while the remaining variation was unexplained and may stem from other factors not considered in this study. This highlights the need for a better understanding of underlying factors driving local malaria transmission and outbreaks, to better tailor regional programmatic responses.

Keywords Anopheles stephensi, Environmental factors, Malaria risk, Spatiotemporal, Ethiopia

Background

Global efforts towards malaria control and elimination were heightened following the launch of Roll Back Malaria programme in 1998 and the declaration of Abuja in 2000, with the target of reducing malaria-related deaths by 50% before the end of 2010 [1]. Since then, investments have been made to promote universal coverage of insecticide-treated mosquito nets (ITNs), diagnostic testing, and updated malaria treatment guidelines [2]. These efforts saw a drop in malaria cases and deaths from 238 million and 736,000 in 2000 to 229 million to 409,000 in 2019, respectively [3]. Between 2000 and 2015, eight countries eliminated malaria and many others reduced transmission to low levels [4]. Motivated by these achievements, in 2015 WHO developed a global technical strategy (GTS) for 2016-2030 [2] to reduce malaria related mortality and incidence by at least 90% by 2030 from the 2015 baseline [4].

However, between 2015 and 2019, the global rate of decline was decidedly lower, at around 2% [5], highlighting that the programme might fall short of the targets for 2030 [6]. Importantly, the 2023 WHO malaria report showed that the global case incidence was off course by 55% [7]. Factors contributing to this discrepancy include the emergence of drug and diagnostic-resistant parasites [8, 9], mosquito resistance to insecticides [7, 10], the COVID-19 pandemic [10], the global rise in temperatures [7], expansion of *Anopheles stephensi* [8, 11], and weakening of control programmes due to conflict and internal population displacement [12].

In line with global momentum, in 2004, Ethiopia began scaling up interventions to prevent and control malaria, including the distribution of insecticide treated nets (ITNs), indoor residual spraying (IRS), and the introduction of rapid diagnostic tests and artemisinin-based combination therapies [13, 14]. Moreover, the 'test-and-treat' policy was implemented in 2010 [13]. This achieved a reduction in malaria-related deaths of 54% between 2000 and 2016 [15], and Ethiopia was one of four countries that were on course to meet the GTS target by 2020 [16].

With these gains, Ethiopia set a goal to achieve zero indigenous malaria cases in 565 and 1046 districts by 2025 and 2030, respectively [17]. To support these goals, the risk of malaria at the district level was stratified into five districts based on annual parasite incidence (API,

cases/1000 people per year), altitude and expert opinions: high risk (API \geq 50), medium risk (API \geq 10 & <50), low risk (API > 5 & <10), very low risk (API > 0 & \leq 5), and malaria-free areas (API=0) [14]. However, efforts to prioritize elimination were complicated by a rapid growth in malaria in 2022 with an estimated 1,732,562 malaria cases [7], nearly a two-fold increase from the 2018 baseline of 962,087 [16]. Factors attributed to the nationwide upsurge included climatic anomalies [18], biological threats including *An. stephensi* [11, 19] and emergence of drug and diagnostic-resistant *Plasmodium falciparum* (*P. falciparum*) [8, 9], service interruptions due to the COVID-19 pandemic [14], and widespread internal conflicts [7].

To evaluate risk factors associated with the spatial and temporal patterns of malaria risk in Ethiopia, we therefore utilised a geospatial modelling technique [20]. Our approach can support decision-makers in tailoring interventions at a local scale and to inform technologies, strategies, and target populations [21].

Methods

Background to the study

Ethiopia is characterized by Kola or hot lowlands, Weyna Dega, and Dega or cool highlands with altitudes of \leq 1500 meters above sea level (masl), 1500–2400 masl, and>2400 masl, respectively. Rainfall is strongly correlated with altitude and therefore varies significantly across the country [22]. This has resulted in heterogeneous malaria transmission in a bimodal pattern [14]. The bimodal seasonal pattern of malaria transmission is mainly associated with rainfall. In most parts of Ethiopia the major transmission peaks occur from September to December following the main rainy season (June to August) and small peaks from April to June following the minor rainy season (March to May) [22, 23]. This seasonal pattern includes a lag phase of approximately 1 month following the end of the rainy season [24]. Altitude is also a major driver of malaria transmission and areas that lie below 2000 masl, where approximately 60% of Ethiopians reside, are considered malarious [17], [25]. Overall, Plasmodium falciparum and Plasmodium vivax are the co-endemic species in different proportions with

Variable	Code	Unit	Source	Resolution	
Horizontal easterly wind speed, at a height of ten metres above the Earth's surface	ital easterly wind speed, u10 Metres per second ght of ten metres he Earth's surface		ERA5-Land hourly reanalysis data, Copernicus's Climate Data Store, by European	Temporal: hourly 1950–present Spatial: 1100 m	
Horizontal northernly wind speed, at a height of ten metres above the Earth's surface	v10	Metres per second	Centre for Medium-Range Weather Forecasts. These data are processed retrospectively		
Leaf area index, high vegetation (evergreen trees, deciduous trees, mixed forest/woodland, and interrupted forest)	lai_hv	Square meter per square meter	and ground data assimilation methods		
Leaf area index, low vegeta- tion (crops and mixed farming, irrigated crops, short grass, tall grass, tundra, semidesert, bogs and marshes, evergreen shrubs, deciduous shrubs, and water and land mixtures)	lai_lv	Square meter per square meter			
Skin (Earth surface) temperature	skt	Kelvin			
Total precipitation	tp	Meter			
Volume of water in soil layer 1 (0–7 cm)	swvl1	cubic meter per cubic meter			
Gridded population density	pop_density	Number of people per square kilometer	Socioeconomic Data and Appli- cations Center, NASA, Version 4	Temporal: Year 2020 Spatial: 1000 m	
Land cover	land_cover	-	CCI Land Cover (LC) team, Euro- pean Space Agency	Temporal: Year 2016 Spatial: 20 m	
Elevation	Elevation	Meter	U.S. Geological Survey	Temporal: Year 2007 Spatial: 450 m	
Presence of invasive species	Invasive	Binary (presence/absence)	WHO Global Malaria Programme	Temporal: yearly Spatial: zone-level	

Table 1 Variables used for contributing factors to spatial and temporal variation in weekly malaria cases across Ethiopian zones

a national average of 65% and 35% of all cases respectively [14].

Malaria case data

The Ethiopia Public Health Institute (EPHI) [26] is responsible for collecting data related to public health emergency management (PHEM) [26]. The public health institute analyses weekly PHEM data on diseases that are classified as public health threats in Ethiopia, including malaria, to identify areas of concern and respond with appropriate measures. PHEM data collected at the health facility level is aggregated by catchment, district, zone, and region.

Clinical malaria cases PHEM data between January 2013 to January 2023 were obtained from EPHI. Data was stratified to the finest spatial resolution available at woreda level, the third administrative division in Ethiopia. In the Ethiopian health system, a primary hospital is expected to serve a woreda with an average population size of 60,000 to 100,000. However, significant administrative border changes were implemented between 2013 and 2022. Therefore, to reduce spatial uncertainties as a result of these changes, data were aggregated to zone

level, the second administrative division which was less affected. Data completeness improved over time; the annual national rate of missing records ranged from 17.6% in 2015 to 5% in 2021. Ten zones had no missing records and the average rate of missing records was approximately 5% in all other zones. West Gondar zones had the highest missing rates (40% to 79% of records) potentially due to internal conflict.

To facilitate georeferencing of weekly malaria surveillance data we utilised the 2021 Ethiopia administrative division shapefile and its associated population size provided by the United Nations Office for the Coordination of Humanitarian Affairs [27, 28]. This shapefile includes 13 regions and 92 zones. However, to align with the recording structure of the malaria surveillance data, East Bale zone and Bale zone, Dire Dawa rural zone and Dire Dawa urban zone were all merged within the retrieved shapefile. Therefore, all analyses in this study are conducted exclusively on 90 zones in 13 regions of Ethiopia (Supplementary Table 1).

Table 2	Random effect te	erms included in t	the log-linear	model for the week	ly risk of malar	ia in Ethiopian zones

Term	Notation	Representing	Probability distribution	Parameters (variances)
Temporal trend	$ au_t$	Overall long-term and large-scale temporal variation in the entire Ethiopia (not by zone)	First-order random walk	σ_{τ}^2
Zone-specific seasonality	s _{i,t}	Recurring seasonal variations in each zone not explained by environmental factors	Cyclic second-order random walk tem- poral pattern and spatially exchange- able	σ_s^2
Zone-level spatial trend	ξi	Spatial variations between zones arise from dependencies or heterogeneity that are not explained by environmental factors	Besag-York-Mollié (BYM)	$\sigma^2_{m{\xi}, Besag}$ and $\sigma^2_{m{k}, iid}$
Invasive species	$v_{i,t}$	Presence of An. stephensi in a zone from time i	Independent and identically distributed	σ_v^2

Risk factors

Remote sensing data for Ethiopia across the whole study period were retrieved from open repositories (Table 1). Spatial and temporal resolutions of environmental factors varied across some variables. This can introduce bias due to a mismatch between the scale of change in each variable and the scale of measurement. To address this, we aggregated the environmental factors to match the spatial and temporal scale of the case counts. Specifically, for original spatial and temporal scales, we applied spatial aggregation to all environmental factors at zone level and temporal aggregation at weekly level. This aggregation process involved the computation of summary statistics, including mean, minimum, maximum and standard deviation for each variable per zone and per week. The only exception was land cover, where we calculated percentage per zone for the ten categories of land cover.

To assess the impact of *An. stephensi*, we combined data on 114 mosquito sampling attempts across Ethiopia between 2016 and 2023 from the WHO malaria threats map dataset [29] and other recent fieldwork [11] (Table 1).

Modelling of spatiotemporal variations

Although the Poisson model is commonly used for spatial and spatiotemporal count data, it assumes that the mean and variance are equal, which can be restrictive. In contrast, the negative binomial model allows the variance to be greater than the mean, a condition called overdispersion. When data shows more variability than expected under a Poisson model, the negative binomial model is a more suitable choice [30]. To address variability and potential overdispersion in the weekly number of malaria cases across Ethiopian zones, denoted by $y_{i,t}$ where *i* is the zone and *t* the week, a negative binomial regression model was chosen over a Poisson model to examine the spatiotemporal relationships of malaria with the risk factors, denoted by $x_{i,t} = (x_{i,t,1}, \dots, x_{i,t,M})$, where *M* is the total number of risk factors. The mean of the distribution of $y_{i,t}$ is:

$$\mu_{i,t} = E[y_{i,t}] = P_{i,t} \times \theta_{i,t}$$

and variance is:

$$\sigma_{i,t}^2 = Var(y_{i,t}) = \mu_{i,t} \times \left(1 + \frac{\mu_{i,t}}{\varphi}\right)$$

here $P_{i,t}$ and $\theta_{i,t}$ denote the known exposed population size and expected relative risk of malaria infection in the zone *i* and week *t*, respectively, and $\varphi > 0$ is the dispersion parameter. As a common approach in disease mapping with aggregated areal count data, including the population size $P_{i,t}$ as a multiplicative offset term to model the mean, allows adjustment for population variations. Thus, the risk $\theta_{i,t}$ represents the impact of all other factors besides population fluctuations [31] and is represented by a log-linear model:

$$\theta_{i,t} = \exp(\alpha + f_{i,t} + r_{i,t})$$

where:

- Intercept: α
 - expresses the unmodelled overall country-wide average of expected risk of malaria;
- Fixed effect of environmental factors: $f_{i,t} = \sum_{j=1}^{M} \beta_j x_{i,t,j}$
 - o each environmental factor $x_{i,t,j}$ has its own coefficient β_j ;
 - Random effect of other sources of variations: $r_{i,t} = \tau_t + s_{i,t} + \xi_i + \nu_{i,t}$
 - Represents the additive combination of temporal trend, zone-specific seasonality, zone-level spatial trend and effect of invasive species, as summarised in Table 2.



Fig. 1 Weekly clinical malaria cases in Ethiopia from January 2013 to January 2023, EPHI

As a common model in the context of disease mapping and epidemiology, a Besag-York-Mollié (BYM) model was considered for possible spatial heterogeneity and dependence between zones. This model includes two components, enabling it to capture both spatial autocorrelation among zones and heterogeneity across different zones, even after accounting for environmental factors and existing spatial autocorrelation.

A Bayesian approach with integrated nested Laplace approximation method [32] was used for parameter estimation and statistical inference of the proposed model. The missing weekly counts for some zones were considered to be missing completely at random. This means that the likelihood of missing data is the same for all observations, regardless of any specific characteristics or values. Based on this assumption, any analysis performed on the available data remains unbiased, as the missingness does not systematically influence the results. All the computations for this approach were implemented through the R package INLA [33, 34]. Full computational details are provided in Gómez-Rubio (2020) [35].

To evaluate spatial patterns in the residuals of the models (which can support investigation of additional factors driving malaria) a hierarchical cluster analysis was employed to group similar zones based on their estimated random effects. Among various methods, we used the complete linkage method to calculate the distance between clusters [36]. In this approach, the distance between two clusters is defined as the maximum distance between any pair of spatial random effects from the zones within each cluster.

Results

Progress in malaria control and elimination efforts has stalled since 2021

There were temporal variations in the weekly count of clinical malaria cases in the country during the study period, January 2013 to January 2023 (Fig. 1). Most years exhibited intra-annual bimodal seasonal patterns and a long-term temporal trend that declined between 2013 to 2018, and then increased from 2021 onwards. Seasonal peaks tended to occur in the second half of the year and notable variability among different regions as evident in the summary statistics of the weekly counts of clinical malaria cases by year and by region, respectively, provided in Supplementary Tables 1 and 2.

Binomial model estimates of the malaria risk across different weeks and zones (in 90 zones); mean (lines) and associated 95% credible intervals (grey shading) for the aggregated country-wide risk of malaria cases, $\theta_t = \sum_{i=1}^{90} \theta_{i,t}$, per 10,000 population (i.e. 10,000 × θ_t), over the study period revealed declining risk from 2013 to 2018 followed by a sharp increased risk from early 2021 (Fig. 2).

The spatial pattern of total clinical malaria cases per 10,000 population across zones of Ethiopia reveals a higher concentration of clinical malaria cases in western and northwestern zones (Fig. 3).

The estimated risk of malaria cases per 10,000 population, $10,000 \times \frac{1}{525} \sum_{t=1}^{525} \theta_{i,t}$, showed higher risk in west and northwest zones, compared to other zones. at the zone level for the 525 weeks, accounted for the overall effect of environmental factors as well as spatial and temporal random effects (Fig. 4, right panel). The means of the spatial random effect (Fig. 4 left panel), represent spatial fluctuations not accounted for by other components in the model. The sign and magnitude of these means indicate a general trend: negative spatial effects



Fig. 2 Mean (solid line) and 95% credible interval (grey shading) of the aggregated country-wide risk of malaria per 10,000 population in Ethiopia between January 2013 and January 2023, EPHI



Fig. 3 Rates of clinical malaria cases per 10,000 population across zones of Ethiopia between 2013 and 2023. Grey zones indicate no recorded cases. Lakes are represented in white. For 2022 and 2023, records were missing for North Ethiopia (West Gondar) was due to internal conflict



Fig. 4 Means of the spatial random effect (left) and the estimated mean risk of malaria per 10,000 population [37] across zones of Ethiopia

in the eastern zones and positive spatial effects in the western and northwestern zones. This suggests that predictions based solely on the fixed effects of environmental factors may underestimate or overestimate the risk of malaria. Specifically, underprediction occurs in the West, while overprediction occurs in the East.

Approximately 50% of variability in malaria trends over the last decade, as captured by the model, can be attributed to environmental factors

A goodness-of-fit measure (Bayesian R-squared) indicated that approximately 56% of variation in the weekly number of clinical malaria cases in Ethiopia is explained by the model. Of this explained variation, 55% is attributable to environmental factors.

In terms of the importance of the different residual model components (therefore excluding fixed effects), the zone-specific seasonal random effect $s_{i,t}$ and spatially structured part of the spatial random effect ξ_i are components with larger variances and therefore contribute more to describing unexplained variations (Fig. 6A). Subsequent analyses (Bayesian R-squared) [38] highlighted that approximately 56% of variation in the weekly number of clinical malaria cases in Ethiopia is explained by the model with all components. Within the explained component, 55% is due to fixed effects.

From the fixed effects variables described in Table 1, seven were found significantly associated with weekly malaria variation over the zones (Fig. 6B). In particular the (1) horizontal eastly wind speed at a height of up to 10 m (u10): Higher average wind speed within a zone

is associated with a slight (5%) increase in malaria risk. Surface soil moisture (swvl1), (2) the maximum amount of water held in the top soil layer (0–7cm) across each zone significantly increases the risk of malaria by 7%. Conversely, both the standard deviation and minimum of swvl1 within each zone showed a decrease of around 4% and 7% in malaria risk respectively, with each unit increase, (3) Leaf area index for low vegetation: The mean leaf area index for low vegetation decreases malaria risk by 17% and (4) Elevation: higher minimum elevation is associated with 69% increase in malaria risk while mean elevation is associated with 75% decrease (Fig. 5).

Observed clustering and seasonality is unexplained by environmental factors

The estimated zone-specific seasonal random effects indicated that there is residual seasonal variation (the part of seasonality not explained by environmental factors) across both space and time (Supplementary Fig. 1). Some zones, like Kilbati, do not show a clear seasonal pattern beyond what environmental factors can explain, while others exhibit distinct cycles, with winter declines and summer increases. A hierarchical cluster analysis using a complete linkage method [36] identified four geographical clusters (Fig. 6, top panel) with local seasonality not explained by environmental variables (Supplementary Fig. 2). Zones in Cluster 4 demonstrate consistent seasonal patterns, with significant declines in March and April and significant peaks in October and November. In contrast, zones in Cluster 2 and, to some extent, those in Cluster 1 showed no consistent seasonality. Zones in



Fig. 5 Estimated means and their corresponding 95% credible intervals of logarithms of variance for all random effect terms within the considered model: spatial random effect ξ_{i} , overall temporal random effect τ_{t} , zone-specific seasonal random effect $s_{(i,t)}$ and the presence or absence of invasive species $v_{(i,t)}$ (**A**). Forest plot of the estimated means and their corresponding 95% credible intervals of statistically significant coefficients for environmental factors in the considered model for clinical malaria cases in Ethiopia (**B**)

Cluster 3, however, consistently peak in June. In terms of malaria risk representation, zones in Cluster 3 have a significantly higher malaria risk [95% Confidence interval (CI): 10.8-15.9] than Cluster 2 (95% *CI:* 4.96–7.37). Zones in clusters 1 and 4 tend to have an overall medium risk (Supplementary Table 3). The "[(95% Confidence interval (CI): 10.8-15.9]" should change to: "95% confidence interval (*CI*): 10.8-15.9]".

Discussion

Progress in malaria control and elimination efforts in Ethiopia has stalled since 2021

This study revealed that malaria transmission in Ethiopia exhibited a cyclical seasonal pattern and a declining longterm temporal trend between 2013 and 2018, followed by increases in transmission up to 2021 similar to those observed in the WHO African Region [7]. The models identified a higher concentration of clinical malaria cases in the western and northwestern zones bordering countries.

Approximately 50% of variability in malaria trends over the last decade in Ethiopia could be attributed to environmental factors

The observed lower variance in overall temporal effects implies that large-scale temporal variations play a lesser role. Fixed effects (represented by the selected seven environmental variables), account for approximately 55% of explained variance, and the larger variances in zone-specific seasonal and spatially structured random effects highlight the importance of local variations in understanding the remaining unexplained variations in malaria risk (approximately 45%). While environmental variables like easterly horizontal wind speed at a height of 10 m, surface soil moisture, and higher minimum elevation were associated with increased malaria risk, lower mean leaf area index of low vegetation and mean elevation were associated with decreased risk. However, a substantial amount of unexplained seasonal variation across space and time remained, as captured by zone-specific seasonal random effects, and tended to cluster in regions that are not always geographically contiguous. The significantly higher risk observed in the northwest and lower risk in the southwestern part of Ethiopia (Figs. 3 and 4) is potentially due to a complex interplay of factors. The model outputs are consistent with Ethiopian malaria transmission patterns, which are seasonal, inter-annual and spatially heterogeneous, except in the low-lying southwestern border areas experiencing perennial transmission [13, 14]. Factors influencing mosquito breeding, parasite survival, and consequently malaria transmission, including environmental factors such as temperature and rainfall [39], might play important roles in the observed variation. Variation in malaria seasonality across zones, as well as clustering following similar patterns (Figs. 4 and 6), is consistent with previous studies that documented significant geographical heterogeneity in malaria transmission. Regional and subregional spatial clustering and the presence of hotspots were attributed to environmental and socioeconomic factors. In the northwest, from 2009 to 2010 focal upsurges in malaria cases



Fig. 6 Hierarchical clustering of Ethiopia's zones based on the similarity of their estimated zone-specific seasonal random effects in 4 clusters (top panel) with the posterior mean and 95% confidence intervals of seasonal effects displayed for each cluster (bottom panels)

with spatiotemporal patterns were reported [40]. Low malaria risk in eastern zones compared to the north-western zones might be due to variations in rainfall patterns and seasonality as observed elsewhere [39]. The overall trend in our risk map (Fig. 4) aligns with the Ethiopian national malaria risk stratification map [14], showing a relatively higher burden in the western and

northwestern parts of the country which is also consistent with non-epidemic years [13]. Factors that could have a bearing on the nationwide upsurge in addition to the environmental and socioeconomic variation might be the consequences of long and widespread internal conflict and population displacement [12] as well as the COVID-19 pandemic [10] which disrupted the health system. Therefore, the adverse effects of the interplay between climatic anomalies and conflictassociated displacement in populations at high risk of neglected tropical diseases is highlighted as a strategic research focus [12]. The other potential risk factor which could have contributed to the changes in clinical malaria trends is *An. stephensi*, reported mainly in east and southeastern Ethiopia [8, 41] but not in others [42]. We need to be cautious but our model identified low malaria in eastern Ethiopia where *An. stephensi* is wellestablished [11, 19].

In our model, a horizontal easterly wind speed at a height of ten meters above the Earth's surface was associated with a 5% increase in malaria risk. This might be explained by the fact that wind can assist mosquitoes in host-seeking [43] and wind-borne migration of mosquitoes [44]. Importantly, the direction of the wind from villages to the breeding habitats has been linked with an increased population size of mosquitoes compared to the opposite wind direction [45]. However, wind can also generate waves on the water surface, which can be fatal to the aquatic life stage of mosquitoes in large reservoirs compared to small rain-made breeding habitats [46].

Surface soil moisture (the volume of water in the first layer (0–7 cm) of surface soil) is a good predictor of malaria risk in this study, as indicated elsewhere [47]. Previous studies from Nigeria [48] and Uganda [49] have shown that the vegetation index was an important contributor to malaria risk. Conversely, the mean leaf area index for low vegetation decreased malaria risk by 17% in our study. Importantly, a decrease in elevation (altitude) was significantly associated with an increased relative risk of malaria in our study, while the average higher elevation was not. However, warmer years could promote malaria transmission at higher altitudes in Ethiopia and Colombia [50].

Observed local clustering and seasonality is unexplained by environmental factors

Our study also suggests the existence of local (smallscale) spatial and temporal variations more important than large-scale variations that could shape Ethiopia's malaria risk distribution (Fig. 6A and B). This is evident from the substantial variance in zone-specific seasonal effects, highlighting how seasonal fluctuations within zones influence overall risk. Potentially local human behaviours and mosquito ecology [51] as well as socioeconomic conditions [52] might vary across zones and seasons, impacting transmission dynamics. Importantly, significant variation in the spatially structured effect suggests that local spatial patterns matter. Shared patterns between neighbouring zones could contribute to this spatial clustering [53]. Importantly, within each cluster, it is likely that zones share common factors that were not considered in the model such as mosquito vector population, human behavioral factors, differences in healthcare structure, and malaria interventions. biological threats such as drug and diagnostic-resistant *P. falciparum* reported in different parts of Ethiopia [8, 9].

Our study revealed the spatial and temporal trends of malaria across Ethiopian zones. The local variation in transmission emphasizes the need for tailored interventions across zones but also their potential as sources of epidemics [54]. We acknowledge that the findings should be cautiously interpreted. The inherent incompleteness of the PHEM data and the use of different time scales with environmental data might have resulted in an underestimation of risks. Considering more risk factors, socioeconomic and near-surface temperature, and considering a time lag with nonlinear modelling might have improved the outcome.

Conclusions

Our findings underline the importance of environmental factors, accounting for 55% of the variability explained by our model, as drivers of the spatiotemporal distribution of malaria risk in Ethiopia. The local clustering and seasonality unexplained by environmental factors call for further exploration into tailored focal responses. The presence of a sizable proportion (45%) of risk factors accounting for changes in the trends of clinical malaria remain unexplained by our model; information about biological threats (vector dynamics; change in insecticide resistance and/or biting and resting behavior, occurrence of drug and diagnostic resistance in Plasmodium populations) and years in the grip of internal conflict could have played important roles. This study showed that reduced effectiveness of malaria control efforts in a country after years of progress towards elimination highlights the need for multi-sectoral coordination, and a localised and coordinated response beyond the health system.

Supplementary Information

The online version contains supplementary material available at https://doi. org/10.1186/s40249-024-01259-4.

Additional file 1

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Author contributions

EG, LS, YA, DW, ALW and MJD conceptualized and designed the study. GMA, TA, AJ, NN, AK, MA, NT, DT and BK curated the data. AJ, GMA and LS conducted

the statistical analysis. EH, EM, TA, LS, AJ and EE drafted the manuscript. EG, MJD, AMR, DW, ALW and LS critically reviewed the manuscript. All authors reviewed, edited and approved the manuscript.

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Availability of data and materials

The data underlying this article will be shared upon reasonable request to the corresponding author.

Declarations

Ethics approval and consent to participate

The study is part of the CEASE (Controlling emergent An. stephensi in Sudan and Ethiopia), the umbrella Project received approval from the National (Ref: 02/256/630/14) and the ALERT/AHRI (Ref: P0-08-22) research ethics committees, Addis Ababa, Ethiopia. This manuscript used case data made readily available by the Ethiopian Federal Ministry of Health malaria programme (PHEM), thus consent was not necessary.

Consent for publication

Not applicable.

Competing interests

No competing interests were declared by the authors.

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References

- WHO. The Abuja Declaration and the plan of action: an extract from the African Summit on Roll Back Marlaria, Abuja, 25 April 2000 (WHO/CDS/ RBM/2000.17). Geneva: World Health Organization; 2003. https://iris.who. int/handle/10665/67816.
- WHO. A framework for malaria elimination. Geneva: World Health Organization; 2017.
- WHO. World malaria report 2020. Geneva: World Health Organization; 2020. Report No.: 978-92-4-001579-1. https://www.who.int/publicatio ns/i/item/9789240015791.
- 4. WHO. Global technical strategy for malaria 2016–2030. Geneva: World Health Organization; 2015.
- Ansah EK, Moucheraud C, Arogundade L, Rangel GW. Rethinking integrated service delivery for malaria. PLoS Glob Public Health. 2022;2(6): e0000462.
- WHO. Global technical strategy for malaria 2016–2030, 2021 update. Geneva: World Health Organization; 2021. https://www.who.int/publi cations/i/item/9789240031357.
- WHO. World Malaria Report 2023. Geneva: World Health Organization; 2023.
- Emiru T, Getachew D, Murphy M, Sedda L, Ejigu LA, Bulto MG, et al. Evidence for a role of *Anopheles stephensi* in the spread of drug and diagnosis-resistant malaria in Africa. Nat Med. 2023;29(12):3203–11.

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- Fola AA, Feleke SM, Mohammed H, Brhane BG, Hennelly CM, Assefa A, et al. *Plasmodium falciparum* resistant to artemisinin and diagnostics have emerged in Ethiopia. Nat Microbiol. 2023;8(10):1911–9.
- 10. WHO. Global technical strategy and targets for malaria 2016–2030. Geneva: World Health Organization; 2024. https://apps.who.int/gb/ ebwha/pdf_files/WHA77/A77_6-en.pdf.
- Ashine T, Eyasu A, Asmamaw Y, Simma E, Zemene E, Epstein A, et al. Spatiotemporal distribution and bionomics of *Anopheles stephensi* in different eco-epidemiological settings in Ethiopia. Parasit Vectors. 2024;17:166.
- Kelly-Hope LA, Harding-Esch EM, Willems J, Ahmed F, Sanders AM. Conflict-climate-displacement: a cross-sectional ecological study determining the burden, risk and need for strategies for neglected tropical disease programmes in Africa. BMJ Open. 2023;13(5): e071557.
- Taffese HS, Hemming-Schroeder E, Koepfli C, Tesfaye G, Lee M-C, Kazura J, et al. Malaria epidemiology and interventions in Ethiopia from 2001 to 2016. Infect Dis Poverty. 2018;7:103.
- NMEP. Ethiopia Malaria Elimination Strategic Plan: 2021–2025. Addis Ababa: Ministry of Health; 2020. http://repository.iifphc.org/xmlui/handle/123456789/1526.
- Girum T, Shumbej T, Shewangizaw M. Burden of malaria in Ethiopia, 2000–2016: findings from the Global Health Estimates 2016. Trop Dis Travel Med Vaccines. 2019;5:11.
- WHO. World Malaria Report 2019. Geneva: World Health Organization; 2019. https://iris.who.int/bitstream/handle/10665/330011/9789241565 721-eng.pdf?sequence=1.
- NMER. National Malaria Elimination Roadmap: 2017–2030. Addis Ababa: Ministry of Health; 2021. https://www.scribd.com/document/587696971/ Malaria-Elimination-Roadmap-Ethiopia-final-H.
- Woyessa A, Siebert A, Owusu A, Cousin R, Dinku T, Thomson MC. El Niño and other climatic drivers of epidemic malaria in Ethiopia: new tools for national health adaptation plans. Malar J. 2023;22:195.
- Balkew M, Mumba P, Yohannes G, Abiy E, Getachew D, Yared S, et al. An update on the distribution, bionomics, and insecticide susceptibility of *Anopheles stephensi* in Ethiopia, 2018–2020. Malar J. 2021;20:263.
- Collins O, Duffy K. A mathematical model for the dynamics and control of malaria in Nigeria. Infect Dis Model. 2022;7(4):728–41.
- Smith TA, Chitnis N, Penny M, Tanner M. Malaria modeling in the era of eradication. Cold Spring Harb Perspect Med. 2017;7(4): a025460.
- 22. FMoH. An epidemiological profile of malaria in Ethiopia. Addis Ababa: Ministry of Health; 2014.https://webarchive.lshtm.ac.uk/www.linkmalaria. org/sites/link/files/content/country/profiles/Ethiopia%20Epi%20Report% 20(240314).pdf.
- FMoH. Ethiopia Health and Health Related Indicators 2010–2011. Addis Ababa: Ministry of Health; 2012. https://ghdx.healthdata.org/record/ethio pia-health-and-health-related-indicators-2010-2011.
- 24. Haileselassie W, Parker DM, Taye B, David RE, Zemene E, Lee MC, et al. Burden of malaria, impact of interventions and climate variability in Western Ethiopia: an area with large irrigation based farming. BMC Public Health. 2022;22:196.
- FMoH. Ethiopia: National Malaria Guidelines, Fourth edition. Addis Ababa: Ministry of Health; 2017. https://reliefweb.int/report/ethiopia/ethiopianational-malaria-guidelines-fourth-edition-november-2017.
- EPHI. Ethiopia National Malaria Indicator Survey 2015. Addis Ababa: Ethiopian Public Health Institute; 2016. https://www.malariasurveys.org/ documents/Ethiopia_MIS_2015.pdf.
- Ethiopia: Subnational Administrative Boundaries [Internet]. United Nation. 2021. Available from: https://data.humdata.org/dataset/cb58fa1f-687d-4cac-81a7-655ab1efb2d0. Accessed 24 April 2024.
- Ethiopia: Subnational Population Statistics [Internet]. United Nation. 2022. Available from: https://data.humdata.org/dataset/cod-ps-eth. Accessed 24 April 2024.
- Malaria Threats Map: Invasive vector species [Internet]. World Health Organization. 2024. Available from: https://apps.who.int/malaria/maps/ threats/. Accessed 24 April 2024.
- Hilbe JM. Modeling count data. 32 Avenue of the Americas, New York, NY 10013–2473, USA: Cambridge University Press; 2014.
- Lawson AB. Bayesian disease mapping: hierarchical modeling in spatial epidemiology. 3rd edn: Chapman and Hall/CRC; 2018.
- Rue H, Martino S, Chopin N. Approximate Bayesian inference for latent Gaussian models by using integrated nested Laplace approximations. J R Stat Soc Series B Stat Methodol. 2009;71(2):319–92.

- 33. Martins TG, Simpson D, Lindgren F, Rue H. Bayesian computing with INLA: new features. Comput Stat Data Anal. 2013;67:68–83.
- 34. Wang X, Yue YR, Faraway JJ. Bayesian regression modeling with INLA. New York: Chapman and Hall; 2018. p. 324.
- 35. Gómez-Rubio V. Bayesian inference with INLA. Boca Raton, FL: Chapman & Hall/CRC Press; 2020.
- Everitt BS, Landau S, Leese M, Stahl D. Cluster analysis. New York: John Wiley & Sons, Ltd; 2011.
- Osland A, Beyene D, Vefring HK, De Wit TR, Wright MS. Identification and characterization of human B-cell epitopes in recombinant antigens of *Leishmania aethiopica*. Parasite Immunol. 1996;18(5):265–9.
- Gelman A, Goodrich B, Gabry J, Vehtari A. R-squared for Bayesian regression models. Am Stat. 2019;73:3.
- Thomson MC, Ukawuba I, Hershey CL, Bennett A, Ceccato P, Lyon B, et al. Using rainfall and temperature data in the evaluation of national malaria control programs in Africa. Am J Trop Med Hyg. 2017;97(Suppl 3):32.
- Alemu K, Worku A, Berhane Y. Malaria infection has spatial, temporal, and spatiotemporal heterogeneity in unstable malaria transmission areas in northwest Ethiopia. PLoS ONE. 2013;8(11): e79966.
- Zhou G, Taffese HS, Zhong D, Wang X, Lee M-C, Degefa T, et al. Resurgence of clinical malaria in Ethiopia and its link to *Anopheles stephensi* invasion. Pathogens. 2024;13:9.
- Teka H, Golassa L, Medhin G, Balkew M, Sisay C, Gadisa E, et al. Trend analysis of malaria in urban settings in Ethiopia from 2014 to 2019. Malar J. 2023;22:235.
- Cummins B, Cortez R, Foppa IM, Walbeck J, Hyman JM. A spatial model of mosquito host-seeking behavior. PLoS Comput Biol. 2012;8(5): e1002500.
- Huestis DL, Dao A, Diallo M, Sanogo ZL, Samake D, Yaro AS, et al. Windborne long-distance migration of malaria mosquitoes in the Sahel. Nature. 2019;574(7778):404–8.
- 45. Endo N, Eltahir EA. Prevention of malaria transmission around reservoirs: an observational and modelling study on the effect of wind direction and village location. Lancet Planet Health. 2018;2(9):e406–13.
- 46. Endo N, Eltahir EA. Modelling and observing the role of wind in *Anopheles* population dynamics around a reservoir. Malar J. 2018;17:48.
- Kalthof MWML, Gravey M, Wijnands F, Karssenberg D. Predicting continental scale malaria with land surface water predictors based on malaria dispersal mechanisms and high-resolution earth observation data. GeoHealth. 2023;7(10):e2023000811.
- Okunlola O, Oloja S, Ebiwonjumi A, Oyeyemi O. Vegetation index and livestock practices as predictors of malaria transmission in Nigeria. Sci Rep. 2024;14:9565.
- 49. Okiring J, Routledge I, Epstein A, Namuganga JF, Kamya EV, Obeng-Amoako GO, et al. Associations between environmental covariates and temporal changes in malaria incidence in high transmission settings of Uganda: a distributed lag nonlinear analysis. BMC Public Health. 2021;21:1962.
- Siraj AS, Santos-Vega M, Bouma MJ, Yadeta D, Carrascal DR, Pascual M. Altitudinal changes in malaria incidence in highlands of Ethiopia and Colombia. Science. 2014;343(6175):1154–8.
- Gryseels C, Durnez L, Gerrets R, Uk S, Suon S, Set S, et al. Re-imagining malaria: heterogeneity of human and mosquito behaviour in relation to residual malaria transmission in Cambodia. Malar J. 2015;14:165.
- Rouamba T, Nakanabo-Diallo S, Derra K, Rouamba E, Kazienga A, Inoue Y, et al. Socioeconomic and environmental factors associated with malaria hotspots in the Nanoro demographic surveillance area, Burkina Faso. BMC Public Health. 2019;19:249.
- Alemu A, Abebe G, Tsegaye W, Golassa L. Climatic variables and malaria transmission dynamics in Jimma town, South West Ethiopia. Parasit Vectors. 2011;4:11.
- Kouame RMA, Edi AVC, Cain RJ, Weetman D, Donnelly MJ, Sedda L. Joint spatial modelling of malaria incidence and vector's abundance shows heterogeneity in malaria-vector geographical relationships. J Appl Ecol. 2024;61(2):365–78.