Geospatial evaluation of trade-offs between equity in physical access to healthcare and health systems efficiency

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ABSTRACT

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Correspondence to Dr Hari S lyer; hai161@mail.harvard.edu **Introduction** Decisions regarding the geographical placement of healthcare services require consideration of trade-offs between equity and efficiency, but few empirical assessments are available. We applied a novel geospatial framework to study these trade-offs in four African countries.

Methods Geolocation data on population density (a surrogate for efficiency), health centres and cancer referral centres in Kenya, Malawi, Tanzania and Rwanda were obtained from online databases. Travel time to the closest facility (a surrogate for equity) was estimated with 1 km resolution using the Access Mod 5 least cost distance algorithm. We studied associations between district-level average population density and travel time to closest facility for each country using Pearson's correlation, and spatial autocorrelation using the Global Moran's I statistic. Geographical clusters of districts with inefficient resource allocation were identified using the bivariate local indicator of spatial autocorrelation.

Results Population density was inversely associated with travel time for all countries and levels of the health system (Pearson's correlation range, health centres: -0.89 to -0.71; cancer referral centres: -0.92 to -0.43), favouring efficiency. For health centres, negative spatial autocorrelation (geographical clustering of dissimilar values of population density and travel time) was weaker in Rwanda (-0.310) and Tanzania (-0.292), countries with explicit policies supporting equitable access to rural healthcare, relative to Kenya (-0.579) and Malawi (-0.543). Stronger spatial autocorrelation was observed for cancer referral centres (Rwanda: -0.341; Tanzania: -0.259; Kenya: -0.595; Malawi: -0.666). Significant geographical clusters of sparsely populated districts with long travel times to care were identified across countries. Conclusion Negative spatial correlations suggested that the geographical distribution of health services favoured efficiency over equity, but spatial autocorrelation measures revealed more equitable geographical distribution of facilities in certain countries. These findings suggest that even when prioritising efficiency, thoughtful decisions regarding geographical allocation could increase equitable physical access to services.

INTRODUCTION

Achieving health equity through optimal allocation of healthcare resources is a priority for

Key questions

What is already known?

- Few prior evaluations of equity-efficiency trade-offs using travel time and population density have been conducted.
- Policy-makers would benefit from simple measures that can be readily estimated using existing data. These measures could help inform debates regarding the geographical placement of new facilities and health services, and enable monitoring of progress towards equity and/or efficiency goals.

What are the new findings?

- ► We studied associations between district-level population density (as a proxy for efficiency) and travel time (proxy for equitable physical access) using Pearson's correlations and the Global Moran's I test for spatial autocorrelation (geographical clustering of similar or dissimilar values of variables) in four African countries (Kenya, Malawi, Rwanda and Tanzania).
- While negative Pearson's correlations between population density and travel time were observed across countries and health systems levels, there was variability in negative spatial autocorrelation (geographical clustering of dissimilar values of population density and travel time). Weakest spatial autocorrelation was observed in Rwanda and Tanzania, countries with explicit policies supporting equitable access to rural healthcare.

What do the new findings imply?

- Applying this geospatial analytical approach could help policy-makers understand the trade-offs made between efficiency and equity in geographical access to health services across levels of the health system, and study how these decisions impact service delivery and population health.
- Geospatial data and analytical software are available for all countries around the world, facilitating consistent comparisons across countries.
- This approach can be adopted by health planners and policy-makers to monitor progress towards health systems goals, and identify geographical subregions with inefficient resource allocation to motivate an appropriate policy response.

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health systems worldwide.¹ Increasing equitable access to health services is one means of reducing health inequities.² Access can be defined using four dimensions^{3 4}: availability, geographical accessibility, affordability and acceptability. Geographical accessibility depends on decisions made by healthcare policy-makers about where to build healthcare facilities and how to distribute services throughout their coverage areas. Consequently, a better understanding of geographical accessibility to healthcare resources could inform policy decisions concerning the spatial distribution of services in ways that reduce health disparities. Decisions about where to place health services must balance trade-offs between equity of access versus efficiency of placement to leverage economies of scale.⁴⁻⁶

Urban and rural patients experience different barriers to accessing care. Geographical accessibility is often limited for rural patients, particularly in low-resource settings.⁷⁸ As the global burden of disease shifts towards non-communicable diseases that require more complex and specialised diagnosis and treatment (eg, pathology, chemotherapy and radiotherapy for cancer), urbanrural disparities may be exacerbated if services are preferentially deployed in urban areas.9-12 Policy-makers often decide to place diagnostic services in urban areas to benefit from economies of scale, because these specialised approaches may rely on efficiencies arising from concentrations of highly skilled workers, infrastructure and financial resources. However, placing services in urban areas could introduce geographical, financial and social barriers for rural patients that result in inequitable access to healthcare.¹³ Policy-makers may, therefore, choose to sacrifice some efficiency in favour of more equitable geographical access to healthcare services. Scientists have modelled the costs and benefits of these trade-offs when choosing testing options for HIV and tuberculosis. Their models account for the degree of economies of scale, availability of systems to transport specimens, timely return of results and costs incurred when patients are lost to follow-up.^{14 15} Policy-makers could benefit from knowing where their health systems lie along the equity versus efficiency spectrum as they decide where to place new healthcare services. Empirical assessment of the relationship between travel time as a proxy for equity and population density as a proxy for efficiency arising from economies of scale could provide useful information to policymakers hoping to evaluate the implications of these trade-offs.¹⁶

We propose the Geographic-Population Services Access (Geo-PSA) model, a novel conceptual framework and analytical approach to evaluate trade-offs between urban–rural equity in geographical access to health services and health systems efficiency. We illustrate how geospatial analytical approaches can be used within this framework to identify inefficient geographical allocation of health services and motivate appropriate public health responses. Finally, we demonstrate the flexibility of the Geo-PSA model to evaluate urban–rural disparities in access to (1) primary care services and (2) specialised oncology centres in four sub-Saharan African countries.

METHODS

The Geo-PSA model

Geo-PSA is a data-driven framework for evaluating tradeoffs between equitable geographical access and health systems efficiency when making decisions about placement of healthcare services. Geo-PSA links a conceptual framework for understanding and interpreting the relationship between equity and efficiency within the context of geographical distribution of health services with a geospatial analytical approach to empirically assess these trade-offs.

Conceptual framework

Travel time to the nearest health facility was chosen as a measure of equity because geographical access to health services is known to be a major barrier to accessing care, particularly for rural patients in low-resource settings.⁸ ¹³ ^{17–19} We chose high-resolution population density as a surrogate measure of efficiency arising from economies of scale. Population density reflects urbanicity, and thus, may provide an estimate of economies of scale and expected patient volume. In choosing population density as a proxy for efficiency and travel time as a proxy for equity, we selected metrics that could be readily acquired in diverse geographical contexts over a wide range of spatial scales. However, these measures do not account for other important dimensions of access, such as healthcare quality, which may lead patients to bypass nearby facilities for higher levels of care.^{19 20} Local knowledge regarding the relative importance of physical access compared with other barriers preventing patients from accessing care in a given geographical context is critical for interpreting and acting on Geo-PSA model results. The spatial scale chosen should be that which most closely aligns with the level of the health system where decisions regarding programmes and resources are made. In most countries, this will occur at district level.

We propose that plotting the relationship between travel time to nearest facility (vertical axis) and population density (horizontal axis) may reveal geographical distributions of facilities that reveal trade-offs between equity versus efficiency in healthcare systems, as well as identify settings that are inefficient (figure 1). Health systems that prioritise efficiency would be expected to have districts falling along a steeper line from the upper left quadrant (low-population density, long travel time) to the lower right quadrant (high-population density, short travel time). Health systems that favoured equity would be expected to have districts falling on a flatter line. Inefficient quadrants (upper right: high-population density, long travel time; lower left: low-population density, short travel time) can be used to understand how geographical allocation of resources might be improved. Online supplemental appendix table 1 provides examples of



(Efficiency)

Figure 1 The Geographic-Population services access model: a conceptual framework and analytical approach to inform policies for geographical allocation of health services that Optimise equity and efficiency. Note: the model provides a visual display of geographical data that captures proxies for efficiency (population density, x-axis) and equitable geographical access (travel time, y-axis). Efficient quadrants (upper left: low population density, long travel time; lower right: high population density, short travel time) and inefficient quadrants (lower left: low population density, short travel time; upper right: high population density, long travel time; upper right: high population density, long travel time) can be visualised. Significant outliers (A–D) can be detected using spatial statistical methods.

policy responses to respond to suboptimal geographical allocation of services relative to population.

Study setting and geospatial data sources

We chose to evaluate data from four sub-Saharan African countries: Kenya, Malawi, Rwanda and Tanzania. These countries were chosen because though they have majority rural populations and similar levels of life expectancy, they vary with respect to population density, economic development. The countries have also adopted different approaches to implementing government health policies addressed towards achieving equitable health service delivery. ²¹⁻²⁴ Finally, questions regarding tradeoffs between equity and efficiency are particularly pertinent as they expand access to care for cancer and other chronic diseases.^{25–27} In order to empirically evaluate the association between population density and travel time, we obtained multiple geospatial data inputs described in online supplemental file 1. These geospatial data are generated from administrative government databases and satellite imagery, and are available for most countries around the world for varying time periods. Governance of the public health system in Kenya, Malawi, Rwanda and Tanzania is decentralised, with decision making occurring at lower administrative levels. In order to reflect this in our analysis, we specifically chose to implement our method at district level in Malawi, Rwanda and Tanzania, and at county level in Kenya. Going forward, we refer to the geographical unit across countries as district level for clarity.

Descriptive characteristics for each of the four African countries are presented in table 1 using data from the World Bank Development Indicators²⁸ and the

International Agency for Research on Cancer obtained in 2018.²⁹ The total population ranged from 12.3 million (Rwanda) to 51.3 million (Kenya). Gross Domestic Product per capita in current US\$ ranged from US\$1311 (Malawi) to US\$3468 (Kenya). Life expectancy was similar for all countries, though Rwanda reported the highest (69 years). Estimated age-specific cancer mortality rates ranged from 94.6 (Tanzania) to 129.2 (Kenya) per 100 000. Population density was lowest in Tanzania (63.6 persons per km² and highest in Rwanda (498.7 persons per km². Over half of the population was reported to reside in rural settings for all countries.

Estimating travel time to closest health facility for patients in low-resource settings requires a different set of assumptions and geographical data than in highresource settings. This is because motorised transport via road networks, assumed to be the primary means of travel by patients in high-resource settings, may not always apply in more rural, low-resource settings. For this reason, geographical scientists proposed use of more detailed raster (gridded cell-based) methods that incorporate geographical datasets, including rivers and lakes, elevation and land cover classes. Incorporating these data enables more accurate consideration of the environment and modes of transport used patients in these contexts.³⁰

For raster (gridded) image files (land cover, elevation), we reprojected data using the Universal Transverse Mercator projection in metres corresponding to each country. We then applied a resampling algorithm using ArcMap V.10.2 to change the cell size of all rasters to 500 m to enable consistent spatial contributions of elevation and land cover to travel time estimates. Vector

referral centres				
Country	Kenya	Tanzania	Rwanda	Malawi
Total population (thousands)*	51 393	56318	12302	18143
GDP per capita (US\$ current)*	3468	3240	2252	1311
Life expectancy (years)*	66	65	69	64
Population density (persons per km ²)*	90.3	63.6	498.7	192.4
% Rural*	73	66	83	83
Districts (no)	47	171	30	27
Health centres (no)†	1038	675	486	457
Average population per health centre†	49512	83434	25313	39700
Median district-level travel time to health centre (min, median (IQR))†	44.8 (20.0–74.5)	84.9 (53.2–132.7)	19.5 (14.4–26.9)	55.7 (49.2–67.2)
Top five most frequent cancers‡	Breast, cervix, oesophagus, prostate, colorectum	Cervix, prostate, breast, colorectum, Kaposi sarcoma	Cervix, breast, colorectum, stomach, liver	Cervix, Kaposi sarcoma, oesophagus, non-Hodgkin's lymphoma, breast
Total cancer deaths‡	32987	28610	7662	13779
Age-standardised cancer mortality rate per 100 000‡	129.2	94.6	104.8	123.7
Cancer referral centres (no)§	12	10	6	6
Median district-level travel time to cancer research centre (min, median (IQR))†	127.7 (73.2–220.4)	356.0 (211.3–647.1)	59.9 (44.7–92.8)	296.2 (238.1–520.1)

*World Bank Development Indicators (2018).²⁸

†Authors' analysis of Maina et al. Geospatial Database of Health Facilities in sub-Saharan Africa (2019) (see text).³⁰

‡International Agency for Research on Cancer, Cancer Today Country Factsheets (2018).²

§Global Oncology Project Map (2020).32

GDP, gross domestic product.

datasets (roads, lakes, rivers, healthcare facilities) were acquired from public databases. We examined both geographical distribution of primary care facilities (ie, health centres) as well as tertiary care facilities (ie, cancer hospitals), hypothesising that countries might make different choices at different levels of care. Primary care health centre location data were acquired from a recently published database of government-run health facilities in sub-Saharan Africa.³¹ Lists of cancer hospitals and affiliated clinical research centres for each country were obtained from the Global Oncology online map of African cancer research and clinical centres.³² We obtained latitude and longitude for each cancer research centre by entering the name of the research centre for each country into Google Maps and extracting these coordinates. In sub-Saharan African settings, academic cancer centres often have greater availability of technical experts, training programmes and investments in supply chains, infrastructure and equipment needed to provide high-quality cancer care.^{33 34} We reviewed the list of Global Oncology cancer research and clinical centres and excluded those that corresponded to non-profit or national government agency headquarters, because these were not the locations where cancer care was being delivered. We assumed that the remaining cancer referral

centres would represent higher quality of care than other cancer care facilities and retained them for our analysis.

Estimating travel time to the nearest facility using geographical information systems

We estimated travel time to the nearest health facility using Access Mod 5, a WHO sponsored web-based tool.^{30 35} Briefly, the Access Mod 5 algorithm exploits high-resolution geospatial datasets and the spatial relationships between geographical features in neighbouring areas.³⁰ We assigned a travel speed and mode of transit to each land cover type (online supplemental appendix table 3). Access Mod 5 merges elevation, land cover, water bodies and road network databases into a single raster file, assigning user-defined speeds to each cell in a grid covering the entire geographical area. Next, we used the Access Mod 5 least cost distance algorithm to calculate the fastest route through a given area by minimising the time spent travelling between adjacent cells.³⁰ This analysis is augmented with information about elevation, accelerating or decelerating travel speeds depending on whether travel is uphill or downhill. This method produces a raster file that stores and displays estimated travel times for a specified geographical area as a grid, with each cell representing a 500 m by 500 m area. We ran

this analysis separately for each country and for each list of facilities (primary health centres and cancer referral centres.

Statistical analysis

In order to estimate district-level averages for population density and travel time, we calculated zonal statistics for population density and travel time, specifying the district as the geographical zone and taking the average population density and travel time over all cells in the district. District-level averages were log-transformed before conducting correlation analysis to correct the skewed distribution of these variables. We calculated Pearson correlation coefficients and 95% CIs between variables in each country, separately for travel time to closest health centre and travel time to closest cancer referral centre.

In order to evaluate geographical patterns in the relationship between our measures of equity and efficiency, we performed tests for spatial autocorrelation and cluster detection. Spatial autocorrelation arises when values of a variable in geographical space are positively or negatively correlated with values of another variable in neighbouring areas.³⁶ In our study, negative spatial autocorrelation would arise if densely populated districts were surrounded by districts with short travel time, or if sparsely populated districts were surrounded by districts with long travel time. This would produce clusters in the upper left to lower right quadrants of our conceptual model (figure 1), favouring efficiency. Conversely, positive spatial autocorrelation would arise if most districts clustered in the inefficient lower left to upper right quadrants. The magnitude of the spatial autocorrelation provides evidence regarding trade-offs between equity and efficiency-comparing spatial autocorrelation between countries and health system levels could reveal instances where equity was favoured.

We specified a queen's contiguity matrix to define neighbouring districts. The queen's contiguity matrix is used to classify districts based on their spatial relationships with one another. For a given district, using the queen's contiguity means that a bordering district in any direction would be considered a neighbour. The Global Moran's I³⁶ was calculated to determine whether positive or negative spatial autocorrelation between districtlevel population density and travel time was observed at national level. The bivariate local indicator of spatial autocorrelation (LISA)³⁷ was used to identify four types of geographical clusters of districts: (1) high population density, short travel time (high/short); (2) high population density, long travel time (high/long); (3) low population density, short travel time (low/short); (4) low population density, long travel time (low/long). Since the spatial weighting relies on neighbours, we excluded islands from our analysis. These four categories align with the efficient and inefficient quadrants in our conceptual framework (figure 1). Technical details about the geospatial analysis procedures are provided in the online supplemental methods appendix.

Permutation tests with 999 simulations were used for spatial statistical hypothesis tests. Global Moran's I and bivariate LISA analyses were conducted using GeoDa software.³⁸ We applied the Benjamini-Hochberg False Discovery Rate to our alpha level of 0.05 to correct for multiple testing across all of the districts.³⁹ As a sensitivity analysis, we also included results without correcting for multiple testing. In order to facilitate uptake of this analytical approach among government, non-profit or other healthcare organisations, we have included a data sharing appendix containing further information regarding the workflow and links to specific data elements and code used to generate the major figures in this paper.

Patient and public involvement

There was no patient or public involvement in the study.

RESULTS

Primary care at health centres

The number of health centres reported in the database used for this analysis ranged from 457 (Malawi) to 1038 (Kenya) (table 1). On average, each health centre served a population ranging from 25313 (Rwanda) to 83434 (Tanzania). The median district-level average travel time to the nearest health centre was shortest for Rwanda (19.5min, IQR: 14.4–26.9min) and longest for Tanzania (84.9min, IQR: 53.2–132.7min).

The bivariate LISA scatter plots may reflect differences in national priorities when balancing equitable geographical access and efficiency of health service delivery (figure 2). Average travel times to closest health centre were below 120min for most districts across countries. Correlations between population density and travel time were strongest in Kenya (r = -0.89, 95% CI -0.94 to -0.81) and weaker in Malawi (r=-0.80, 95% CI -0.91 to -0.61), Rwanda (r=-0.76, 95% CI -0.88 to -0.54) and Tanzania (r=-0.71, 95% CI -0.78 to -0.62). The Moran's I was statistically significant (permutation p<0.001), and negative for all countries, favouring efficiency (spatial clusters in the upper left and lower right quadrants of our framework). However, there was stronger evidence of negative spatial autocorrelation in Kenya (-0.579) and Malawi (-0.543) compared with Tanzania (-0.292) and Rwanda (-0.310), suggesting the placement of primary health centres may have favoured equity over efficiency in those countries. Maps revealed the geographical locations of clusters corresponding to the four quadrants in our conceptual framework (online supplemental appendix figure 1). Several significant clusters in the upper left quadrant (low population density, unusually long travel time), particularly near country borders were observed in all countries. Analyses without multiple testing correction revealed greater numbers of inefficient geographical clusters in Tanzania and Rwanda (online supplemental appendix figure 2).



Legend • NS • High/Long • Low/Long • High/Short

Figure 2 Pearson correlation between district-level travel time to the nearest primary care health centre and population per 10000 m² in Kenya (KEN), Tanzania (TZA), Rwanda (RWA) and Malawi (MWI). Colours correspond to districts with statistically significant bivariate local indicator of spatial autocorrelation clusters of (1): high population density/long travel time, (2): low population density/short travel time, (3): low population density/long travel time and (4): high population density/short travel time. Significance tests for district clusters were conducted using 999 permutation tests with an alpha=0.05. The Benjamini-Hochberg false discovery rate was applied to correct for multiple testing of clusters. Blue horizontal line denotes 120 min travel time. Dotted lines intersect at the median travel time and population density for each country. NS, non-significant.

Cancer referral centres

We applied the same model as for primary healthcare centres using locations of cancer referral and research centres in Kenya (n=12), Tanzania (n=10), Rwanda (n=6) and Malawi (n=6) (table 1). The median district-level average travel time to the nearest facility was shortest for Rwanda (59.9 min, IQR: 44.7–92.8 min) and longest for Tanzania (356.0 min, IQR: 211.3–647.1 min).

Figure 3 presents bivariate LISA plots for each country. The correlations between district-level travel time to cancer referral centres and population density were strongest for Kenya (r=-0.92, 95% CI -0.96 to -0.86) and Malawi (r=-0.87, 95% CI -0.94 to -0.74), and weaker for Rwanda (r=-0.78, 95% CI -0.89 to -0.59) and Tanzania (r=-0.43, 95% CI -0.55 to -0.30). The Moran's I statistic was significant and negative for all countries, favouring efficiency over equity (permutation p<0.001). Kenya (Moran's I: -0.595) and Malawi (-0.666) displayed stronger spatial autocorrelation than Rwanda (-0.341) and Tanzania (-0.259), again suggesting that Rwanda and Tanzania's health systems may have favoured equity in placement of their cancer services over efficiency.



Legend • NS • High/Long • Low/Short • Low/Long • High/Short

Figure 3 Pearson correlation between district-level travel time to nearest cancer centre and population per 10000 m² in Kenya (KEN), Tanzania (TZA), Rwanda (RWA) and Malawi (MWI). Colours correspond to districts with statistically significant bivariate local indicator of spatial autocorrelation clusters of (1): high population density/long travel time, (2): low population density/ short travel time, (3): low population density/long travel time and (4): high population density/short travel time. Significance tests for district clusters were conducted using 999 permutation tests with an alpha=0.05. The Benjamini-Hochberg false discovery rate was applied to correct for multiple testing of clusters. Blue horizontal line denotes 120 min travel time. Dotted lines intersect at the median travel time and population density for each country. NS, non-significant.

Significant district clusters were mostly identified in efficient quadrants (upper left and lower right), except for Tanzania which displayed some clusters in inefficient quadrants (online supplemental appendix figure 3). In analyses not correcting for multiple testing, three countries had at least one district with inefficient allocation (online supplemental appendix figure 4). Though Tanzania's correlation and spatial autocorrelation measures favoured equity, most districts had average travel times exceeding 120 min. Malawi also displayed travel times over 120 min for most districts, with stronger negative correlation and spatial autocorrelation measures than Tanzania. All countries but Rwanda displayed significant clusters of low population density and long travel times, which may require innovations in diagnostics or referral systems that reduce travel burden for rural patients.

DISCUSSION

Applying the Geo-PSA model in four African countries revealed trade-offs between urban-rural equity in geographical access and efficiency in health service delivery. Pearson correlation coefficients were uniformly negative, suggesting favouring of efficiency across all countries and health system levels. There were modest differences in the strength of the association between travel time and population density between countries, with Kenya and Malawi reporting stronger correlations than Rwanda and Tanzania at health centre level and cancer referral centre level. Tests for spatial autocorrelation within the Geo-PSA model allowed us to formally investigate geographical clustering of similar and dissimilar patterns of association between district-level population density and travel time. All tests for spatial autocorrelation were negative, implying that across countries, predominant geographical patterns were of regions exhibiting high population density with neighbours experiencing short travel time, and regions of low population density with neighbours experiencing long travel time. However, the magnitude of this spatial autocorrelation varied between countries, with Rwanda and Tanzania displaying weaker spatial autocorrelation than Kenya and Malawi. Given that all countries displayed strong negative aspatial correlations between population density and travel time that favour efficiency, spatial autocorrelation revealed how the geographical placement of facilities favoured equity more in some countries than others.

This study demonstrates how the Geo-PSA model can be used to monitor progress towards equitable urban–rural access across multiple countries. The governments of Rwanda and Tanzania have historically prioritised health equity through propoor healthcare financing and decentralised primary care services.^{21 24 40–42} Kenya's health system has focused on efficiency, and the placement of its facilities have been associated with rural–urban disparities in geographical access to primary care services.^{22 43} The health system in Malawi has pursued propoor policies, but its level of economic development has hampered

efforts to address urban-rural inequities in accessing primary care services for maternal and child health.² In the Geo-PSA model, strong Moran's I correlations between travel time to the closest facility and population density, like those observed in Kenya and Malawi, suggest that the placement of facilities favoured efficiency over equity compared with countries with weaker correlations (Rwanda and Tanzania). This relationship is independent of the overall population density, as Rwanda is the most densely populated of the four countries, while Tanzania is the least. These patterns may reflect differences in healthcare policy implementation in each country. Rwanda and Tanzania have taken steps towards equitable healthcare coverage in their respective countries, which have resulted in a greater rural orientation to cancer care delivery.^{25 41 44} Though national policy-making in Rwanda is centralised, implementation of healthcare delivery is decentralised and relies on evidence-based decision making to guide policies and planning.^{24 40 41} Since independence, Tanzania's government has pursued policies to ensure equitable access to healthcare and other public services, which prompted the country to build the necessary dispensaries, health centres and hospitals to reach rural populations.²¹ In contrast, the Government of Malawi faces resource constraints that limit its ability to apportion funds to increase the geographical extent of its health system.^{23 45} In Kenya, Barasa *et al* reported that although universal health coverage increased from 44% in 2003 to 52% in 2013, increases in service coverage and financial risk protection were largely concentrated among the wealthy, suggesting an urban bias in locations where health services are scaled up.⁴⁶

Policy implications

The framework presented here could support healthcare planning in a few important ways. First, the approach can be applied over time as countries increase coverage of healthcare services and introduce new facilities in a population. Countries can compare the Global Moran's I measure of negative spatial autocorrelation between population density and physical accessibility over time and across levels of the health system to monitor progress towards achieving equitable access to services.

Second, the proposed framework enables policymakers to identify locations with inefficient resource allocation (clusters in the lower left or upper right quadrants, reflecting low population density, short travel times or high population density, long travel times, respectively). Prior research in this area focused on modelling the impact of shifting hospital locations on socioeconomic inequities in geographical access to health services to find an optimal balance between efficiency and equity. Our approach offers a simple analytical model for evaluating trade-offs between equity and efficiency. Policy-makers can decide how best to respond to these inefficiencies based on the level of the health system and their resource constraints⁶ (box 1).

Box 1 Applications of Geographic-Population Services Access (Geo-PSA) model in other health services research settings

- A high-income country seeking to reduce health disparities has resources to deploy a limited number of decentralised clinical breast examination screening centres. Using the Geo-PSA model, the national task force is able to identify clusters of low population density, high travel time clusters where such a test would increase equitable access to testing.
- An international task force is assembled to study the geographical distribution of laboratory diagnostics around the world. Using the Geo-PSA model, countries can be compared based on their tradeoffs between efficiency and urban–rural equity of access to specific laboratory services.
- The national government in a low-income country is rapidly scaling up testing in the wake of a global pandemic, and seeks to determine how best to leverage existing laboratories as well as new tools to identify the disease. Using the Geo-PSA model allows the government to evaluate the current geographical accessibility of its laboratories, areas where new laboratories may need to be built (high population, long travel time), and areas where point-of-care diagnostics or specimen transport should be deployed (low population density, long travel time). Armed with this information, the government can work with other partners to develop a plan for increasing coverage of its testing services.

For primary care services, policy-makers may be willing to sacrifice efficiency to increase access to infectious disease tests, under-5 care and maternal care. For cancer diagnostics, innovations that leverage technology or improved referral systems may be needed to reduce geographical barriers for rural patients. Based on this analysis, programmes in Kenya and Malawi may favour strategies to help rural patients access urban cancer centres, through transport subsidies or patient waiting homes. Tanzania and Rwanda, which already have more geographically accessible health services, may instead introduce systems improvements and technology to lower levels of the health system. Several novel technologies have already been piloted to overcome geographical access barriers for rural populations. For example, GeneXpert devices, which have already been deployed for diagnosis of tuberculosis and other infectious diseases, are being adapted to diagnose breast cancer.47 Some global oncology programmes have begun experimenting with telepathology services.^{48 49} As more researchers test novel approaches to deliver cancer care in sparsely populated rural areas, the solutions will benefit rural cancer patients everywhere.

Some limitations should be considered when applying the Geo-PSA model. First, the geographical analysis used to estimate travel time relies heavily on assumptions regarding the mode of transportation and travel speeds and may underestimate the travel times for individual patients.⁵⁰ Analysts should take care to consider the appropriate spatial scale for their analysis, and use high-quality road network data and to the extent they are

able, verify the speeds of transit and mode of transit of the population under study. A strength of geographical analytical estimates of travel time is objective measurement, incorporation of elevation and land cover, and ability to conduct cross-country, macrolevel analyses. Even if exact times for individual patients are estimated with error, the method still enables ranking of places with longer or shorter travel times to reach facilities. Caution may be needed when interpreting results from geographical cluster analysis for regions that do not have neighbours or occur at national borders. The Geo-PSA model does not account for availability of specific equipment and tests, quality of services or health outcomes. However, these limitations are offset by several strengths. Growing government and non-profit databases contain harmonised geographical datasets to capture the various geographical inputs required to estimate travel time for countries around the world.^{51–53} Furthermore, the availability of data at multiple administrative levels⁵⁴ and remote sensing data provide far more flexibility in terms of defining spatial scales and characterising the specific geographical area of interest.^{30 52 53} The ability to rapidly estimate travel times for multiple countries, apply spatial statistical analysis to evaluate efficiency and equity in varied contexts, and application of geospatial analysis to locate clusters of inefficient allocation of services to motivate an appropriate policy response. Urban-rural disparities in physical access to healthcare services have been observed within both high-income⁵⁵⁻⁵⁸ and low-income and middle-income⁵⁹⁻⁶² countries around the world, suggesting that questions regarding trade-offs between physical accessibility and efficiency may apply to other settings outside of sub-Saharan Africa.

CONCLUSION

In summary, we introduce a novel geospatial analytical model, Geo-PSA, to evaluate trade-offs between equity in geographical access to healthcare and efficiency in four African countries. This flexible model can be applied by health policy and services researchers across different disease areas and geographical contexts. The model can be used to monitor progress towards equitable access to healthcare services at national level, compare countries' trade-offs between equity and efficiency, and inform local policies to respond to inefficient geographical allocation of health services.

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Contributors HSI, JF and TRR conceived the study and design. JF and TRR facilitated acquisition of the data. HSI and NGW analysed the data. HSI, NGW, MCC, SH, LFS, JF and TRR interpreted the data. HSI drafted the manuscript. HSI, JF, NGW, SH, LFS, MCC and TRR provided critical review and final approval of the manuscript.

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Data availability statement Data for preparing the scatter plots presented in this study are available in a public, open access repository. Additional data are available upon request to the corresponding author. Data used in this study include tables with health centre and cancer referral centre locations and gridded image files with travel time data (500m resolution) and population density data (100m resolution). These data were generated using publicly available datasets and so no conditions apply constraining their use. We have made selected data and code used to generate the major figures in the paper available at the following link: https://github.com/hiyer09/geopsa_paper.

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REFERENCES

- Inter-Agency and Expert Group on SDG Indicators. Report of the Inter-Agency and expert group on sustainable development goal indicators (E/CN.3/2017/2), Annex III, 2017. Available: https://wwwsciencedirect-com.ezp-prod1.hul.harvard.edu/science/article/pii/ S2214109X17304722 [Accessed 20 Apr 2020].
- 2 Marmot M, Commission on Social Determinants of Health. Achieving health equity: from root causes to fair outcomes. *Lancet* 2007;370:1153–63.
- 3 Oliver A, Mossialos E. Equity of access to health care: outlining the foundations for action. *J Epidemiol Community Health* 2004;58:655–8.
- 4 Peters DH, Garg A, Bloom G, et al. Poverty and access to health care in developing countries. Ann N Y Acad Sci 2008;1136:161–71.

- 5 Anand S. The concern for equity in health. in: public health, ethics, and equity. New York: Oxford University Press, 2004: 15–20.
- 6 Farmer P, Basilico M, Kerry V, et al. Global health priorities for the early twenty-first century. in: Reimagining global health: an introduction. Berkeley and Los Angeles, California: University of California Press, 2013: 320–9.
- 7 Norheim OF. Ethical perspective: five unacceptable trade-offs on the path to universal health coverage. Int J Health Policy Manag 2015;4:711–4.
- 8 Hartley D. Rural health disparities, population health, and rural culture. *Am J Public Health* 2004;94:1675–8.
- 9 Kyei-Nimakoh M, Carolan-Olah M, McCann TV. Access barriers to obstetric care at health facilities in sub-Saharan Africa-a systematic review. Syst Rev 2017;6:110.
- 10 Atun R, Jaffray DA, Barton MB, *et al*. Expanding global access to radiotherapy. *Lancet Oncol* 2015;16:1153–86.
- 11 Juran S, Broer PN, Klug SJ, et al. Geospatial mapping of access to timely essential surgery in sub-Saharan Africa. BMJ Glob Health 2018;3:e000875. Aug.
- 12 Kingham TP, Alatise OI, Vanderpuye V, et al. Treatment of cancer in sub-Saharan Africa. Lancet Oncol 2013;14:e158–67.
- 13 Wilson ML, Fleming KA, Kuti MA, et al. Access to pathology and laboratory medicine services: a crucial gap. Lancet 2018;391:1927–38.
- 14 Strasser R. Rural health around the world: challenges and solutions. *Fam Pract* 2003;20:457–63.
- 15 Cassim N, Coetzee LM, Schnippel K, et al. Estimating implementation and operational costs of an integrated tiered CD4 service including laboratory and point of care testing in a remote health district in South Africa. *PLoS One* 2014;9:e115420.
- 16 Sohn H, Kasaie P, Kendall E, et al. Informing decision-making for universal access to quality tuberculosis diagnosis in India: an economic-epidemiological model. *BMC Med* 2019;17:155.
- 17 Lwasa S. Geospatial analysis and decision support for health services planning in Uganda. *Geospat Health* 2007;2:29–40.
- 18 Grimes CE, Bowman KG, Dodgion CM, *et al.* Systematic review of barriers to surgical care in low-income and middle-income countries. *World J Surg* 2011;35:941–50.
- 19 Kelly C, Hulme C, Farragher T, et al. Are differences in travel time or distance to healthcare for adults in global North countries associated with an impact on health outcomes? A systematic review. BMJ Open 2016;6:e013059.
- 20 Brownlee S, Chalkidou K, Doust J, et al. Evidence for overuse of medical services around the world. *Lancet* 2017;390:156–68.
- 21 Jonsson U. Ideological framework and health development in Tanzania 1961-2000. Soc Sci Med 1986;22:745–53.
- 22 Makau-Barasa LK, Greene S, Othieno-Abinya NA, *et al.* A review of Kenya's cancer policies to improve access to cancer testing and treatment in the country. *Health Res Policy Syst* 2020;18:2.
- 23 Abiiro GA, Mbera GB, De Allegri M. Gaps in universal health coverage in Malawi: a qualitative study in rural communities. *BMC Health Serv Res* 2014;14:234.
- 24 Iyer HS, Chukwuma A, Mugunga JC, et al. A comparison of health achievements in Rwanda and Burundi. *Health Hum Rights* 2018;20:199–211.
- 25 Tapela NM, Mpunga T, Hedt-Gauthier B, et al. Pursuing equity in cancer care: implementation, challenges and preliminary findings of a public cancer referral center in rural Rwanda. *BMC Cancer* 2016;16:237.
- 26 Gopal S, Krysiak R, Liomba NG, et al. Early experience after developing a pathology laboratory in Malawi, with emphasis on cancer diagnoses. PLoS One 2013;8:e70361.
- 27 Rositch AF, Unger-Saldaña K, DeBoer RJ, et al. The role of dissemination and implementation science in global breast cancer control programs: frameworks, methods, and examples. *Cancer* 2020;126 Suppl 10:2394–404.
- 28 World Bank. World development indicators, 2017.
- 29 Ferlay J, Ervik M, Lam F, et al. Global cancer Observatory: cancer today. Lyon, France: International Agency for Research on Cancer, 2018. https://gco.iarc.fr/today
- 30 Ray N, Ebener S. AccessMod 3.0: computing geographic coverage and accessibility to health care services using anisotropic movement of patients. *Int J Health Geogr* 2008;7:63.
- 31 Maina J, Ouma PO, Macharia PM, *et al*. A spatial database of health facilities managed by the public health sector in sub Saharan Africa. *Scientific Data* 2019;6:1–8.
- 32 Global Oncology. The go MAP, 2020. Available: http://www. thegomap.org/?data=projects [Accessed 20 Apr 2020].
- 33 Shulman LN, Mpunga T, Tapela N, et al. Bringing cancer care to the poor: experiences from Rwanda. Nat Rev Cancer 2014;14:815–21.

- 34 Stulac S, Binagwaho A, Tapela NM, et al. Capacity building for oncology programmes in sub-Saharan Africa: the Rwanda experience. Lancet Oncol 2015;16:e405–13.
- 35 AccessMod 5. Modelling physical accessibility to health care. Available: https://www.accessmod.org [Accessed 4 Mar 2020].
- 36 Cliff A, Ord J. Spatial processes: models and applications. London: Pion, 1981: 266.
- 37 Anselin L. Local indicators of spatial Association-LISA. Geogr Anal 1995;27:93–115.
- 38 Anselin L. Exploring spatial data with GeoDa: a workbook. University of Illinois, Urbana-Champaign: Center for Spatially Integrated Social Science, 2005.
- 39 Castro MC, Singer BH. Controlling the false discovery rate: a new application to account for multiple and dependent tests in local statistics of spatial association. *Geographical Analysis* 2006;38:180–208.
- 40 Basinga P, Gertler PJ, Binagwaho A, *et al.* Effect on maternal and child health services in Rwanda of payment to primary healthcare providers for performance: an impact evaluation. *The Lancet* 2011;377:1421–8.
- 41 Binagwaho A, Wagner CM, Gatera M, et al. Achieving high coverage in Rwanda's national human papillomavirus vaccination programme. Bull World Health Organ 2012;90:623–8.
- 42 van EGM. Rural health development in Tanzania: a case-study of medical sociology in a developing country. BRILL, 1976: 206.
- 43 Noor AM, Amin AA, Gething PW, et al. Modelling distances travelled to government health services in Kenya. Trop Med Int Health 2006;11:188–96.
- 44 McCree R, Giattas MR, Sahasrabuddhe VV, et al. Expanding cervical cancer screening and treatment in Tanzania: stakeholders' perceptions of structural influences on scale-up. Oncologist 2015;20:621–6.
- 45 Masamba L. The state of oncology in Malawi in 2015. *Malawi Med J* 2015;27:77–8.
- 46 Barasa E, Nguhiu P, McIntyre D. Measuring progress towards sustainable development goal 3.8 on universal health coverage in Kenya. *BMJ Glob Health* 2018;3:e000904.
- 47 Kimambo A, Ng D. Validation of the GeneXpert breast cancer STRAT4 assay for rapid analysis of breast cancer biomarker status from fine-needle aspiration biopsies in Tanzania (GX-BCB): preliminary results. *Am J Clin Pathol* 2018;150:S138.
- 48 Mpunga T, Hedt-Gauthier BL, Tapela N, et al. Implementation and validation of Telepathology triage at cancer referral center in rural Rwanda. J Glob Oncol 2016;2:76–82.
- 49 Mpunga T, Tapela N, Hedt-Gauthier BL, et al. Diagnosis of cancer in rural Rwanda: early outcomes of a phased approach to implement

anatomic pathology services in resource-limited settings. *Am J Clin Pathol* 2014;142:541–5.

- 50 Rudolfson N, Gruendl M, Nkurunziza T, *et al.* Validating the global surgery geographical accessibility indicator: differences in modeled versus patient-reported travel times. *World J Surg* 2020;44:2123–30.
- 51 Thomson DR, Linard C, Vanhuysse S, et al. Extending data for urban health decision-making: a menu of new and potential Neighborhood-Level health determinants datasets in LMICs. J Urban Health 2019;96:514–36.
- 52 Tonne C, Basagaña X, Chaix B, *et al*. New frontiers for environmental epidemiology in a changing world. *Environ Int* 2017;104:155–62.
- 53 Frumkin H, Haines A. Global environmental change and noncommunicable disease risks. *Annu Rev Public Health* 2019;40:261–82.
- 54 DIVA-GIS free, simple & effective. Available: https://www.diva-gis. org/ [Accessed 3 Sep 2020].
- 55 Shi X, Alford-Teaster J, Onega T, et al. Spatial access and local demand for major cancer care facilities in the United States. Ann Assoc Am Geogr 2012;102:1125–34.
- 56 Rosenthal MB, Zaslavsky A, Newhouse JP. The geographic distribution of physicians revisited. *Health Serv Res* 2005;40:1931–52.
- 57 Kelly C, Hulme C, Farragher T, et al. Are differences in travel time or distance to healthcare for adults in global North countries associated with an impact on health outcomes? A systematic review. BMJ Open 2016;6:e013059.
- 58 Wang F, Luo W. Assessing spatial and nonspatial factors for healthcare access: towards an integrated approach to defining health professional shortage areas. *Health Place* 2005;11:131–46.
- 59 Rocha TAH, da Silva NC, Amaral PV, et al. Access to emergency care services: a transversal ecological study about Brazilian emergency health care network. *Public Health* 2017;153:9–15.
- 60 Nwakeze NM, Kandala N-B. The spatial distribution of health establishmentsin Nigeria. *African Population Studies* 2011;25.
- 61 Attaei MW, Khatib R, McKee M, et al. Availability and affordability of blood pressure-lowering medicines and the effect on blood pressure control in high-income, middle-income, and low-income countries: an analysis of the pure study data. *Lancet Public Health* 2017;2:e411–9.
- 62 Say L, Raine R. A systematic review of inequalities in the use of maternal health care in developing countries: examining the scale of the problem and the importance of context. *Bull World Health Organ* 2007;85:812–9.