

Using Routine Health Information Data for Research in Low- and Middle-Income Countries: A Systematic Review

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Abstract

Background : Routine health information systems (RHISs) support resource allocation and management decisions at all levels of the health system, as well as strategy development and policy-making in many low- and middle-income countries (LMICs). Although RHIS data represent a rich source of information, such data are currently underused for research purposes, largely due to concerns over data quality. Given that substantial investments have been made in strengthening RHISs in LMICs in recent years and that there is a growing demand for more real-time data from researchers, this systematic review builds upon the existing literature to summarize the extent to which RHIS data have been used in peer-reviewed research publications.

Methods : Using terms 'routine health information system', 'health information system', or 'health management information system' and a list of LMICs, four electronic peer-review literature databases were searched from inception to February 20 2019: PubMed, Scopus, EMBASE, and EconLit. Articles were assessed for inclusion based on pre-determined eligibility criteria and study characteristics were extracted from included articles using a piloted data extraction form.

Results : We identified 132 studies that met our inclusion criteria, originating in 37 different countries. Overall, the majority of the studies identified were from Sub-Saharan Africa and were published within the last five years. Malaria and maternal health were the most commonly studied health conditions, although a number of other health conditions and health services were also explored.

Conclusions : Our study identified an increasing use of RHIS data for research purposes, with many studies applying rigorous study designs and analytic methods to advance program evaluation, monitoring and assessing services, and epidemiological studies in LMICs. RHIS data represent an underused source of data and should be made more available and further embraced by the research community in LMIC health systems.

Background

Routine health information systems (RHISs) collect and provide information at regular intervals on services and activities delivered in health facilities¹. RHISs have been implemented in many low and middle-income country (LMIC) health systems to support resource allocation and day-to-day management decisions at facility, district, provincial, and national levels, as well as to facilitate strategy development and policy-making^{2,3}. Despite the fact that RHISs are being implemented at scale in many LMICs, and that they have been widely recognized as an important component of health systems strengthening^{4,5}, prior studies have suggested that researchers continue to prefer using intermittent cross-sectional population-based surveys rather than RHISs data to conduct studies, including the monitoring of health programs and policy evaluations⁶⁻⁸.

In order to improve health system performance, reliable, timely, and transparent data on health services are crucial^{9,10}. RHISs collect such data and thus could provide important insights into health system performance⁴. Substantial investments have been made in the development and strengthening of RHISs in many LMICs over the past two decades^{5,11}, and interventions targeting data collection, processing, analysis, and dissemination have increased the accessibility of RHIS data^{5,12}. While early RHISs were established using paper-based health facility reports, newer web-based systems have been adopted in many LMICs over the last decade^{13,14}. The most common of these is the District Health Information System 2 (DHIS 2) platform, which is used as the foundation for the national health management information systems (HMIS) in 46 countries and piloted in another 21 countries¹⁵. Studies have shown that the implementation of newer information and communication technology systems, along with supportive feedback

mechanisms to encourage their use in routine practice, can lead to substantial improvements in RHIS reporting and data quality ^{5,13,16,17}.

Despite the documented improvements in data quality, recent studies have shown a persistent underuse of RHIS data for research purposes in LMICs ^{8,18}. A number of factors may contribute to the underuse of RHIS data. Numerous studies and commentators have questioned the usefulness of RHIS-sourced data to monitor and evaluate health services due to data quality concerns, such as incompleteness and inaccuracy ^{19–23}. Additionally, RHIS data are often not publicly available for secondary analyses, which further limits their use ²⁴. Due to these concerns, the research community has shown a persistent preference to use data sourced from intermittent cross-sectional population-based surveys rather than data sourced from RHIS to conduct research on health service utilization and policy evaluation in LMICs ^{8,18,25,26}. However, population-based surveys also have drawbacks, including the fact that they may be costly ²⁶ and are often unable to generate sufficient data at the district or other subnational-levels ²⁷. In addition, reliance on such data may encourage the use of potentially weak evaluation designs ⁸ and may make establishing an appropriate baseline challenging when trying to evaluate specific programs ²⁸. Intermittent cross-sectional population-based surveys themselves also suffer from a number of quality concerns and thus should not be considered the gold standard for estimating service coverage or other population-based estimates ²⁹.

Given the potential of RHISs to play a greater role in the evaluation of health policy and programs and to monitor the performance of health systems, it is important to better understand the extent to which such data are currently being used in research studies. To date, there have been no systematic reviews of the use of RHIS data for research purposes beyond studies that were specific to malaria control¹⁸, a gap this paper seeks to address. Specifically, we systematically reviewed the published literature to identify and describe the different ways in which RHIS data have been used in peer-reviewed research, including the types of health conditions studied. We also summarized the different methodologies that have been used to analyze RHIS data in research and the types of strategies that were applied to circumvent common RHIS data issues, such as incomplete or inaccurate data. It is our goal to provide guidance to other researchers who may be interested in using such data for research purposes by helping them to gain a better understanding on how such data have been successfully used in other contexts.

Methods

This systematic literature review followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines. Peer-reviewed published studies that used data from RHISs in LMICs were included in this study, where RHISs were defined as data systems designed to collect and generate information on services provided by health facilities at regular intervals of a year or less ¹. This included data systems that collect information on clinical service delivery, pharmaceuticals, or diagnostic service provision, as well as routine administrative management. Conversely, systems that collect individual-level data for clinical decision-making purposes and pilot systems to test the implementation of a new data collection component or method were not considered to be RHISs.

Search strategy

Relevant studies were identified through an electronic search of four databases of peer-reviewed literature: PubMed, Scopus, EMBASE, and EconLit – from inception through February 20, 2019, the date we launched the search. For each database, we identified studies that contained any of the following free text terms in their titles or abstracts: ‘routine health information system’, ‘health information system’, or ‘health management information system’, and any LMIC, as defined by the World Bank’s 2019 classifications (appendix 1). Articles were included in the study if they met the following criteria: a) full-text article available in English, b) original research, and c) used data from a RHIS in at least

one LMIC for research purposes. In order to be considered as having used data from a RHIS for research purposes, studies had to involve an analysis, either descriptive or analytical, of RHIS data, or applied RHIS data to inform their study design. We excluded studies that: a) only described RHISs, b) only described the administrative decision-making process, c) only focused on RHIS data collection issues, or d) only assessed RHIS data quality.

Selection of studies

Figure 1 shows the number of articles identified and retained at each stage of the systematic review process. After removing duplicates from the various database searches, we identified 1459 potential articles. Two reviewers independently screened the search results by title and abstract for inclusion eligibility. When there was insufficient information to determine eligibility at the title and abstract screening stage, the article was included for full-text screening. Full texts of the potentially eligible articles were then obtained and further screened for inclusion eligibility. At both stages, the reasons for excluding individual articles were recorded. The full-texts for all but one article were found. Disagreements at each stage were resolved through discussion. Where an agreement could not be reached, a third reviewer made the final determination.

Data extraction and analysis

Two authors extracted data from all included studies using a piloted data extraction form. For each included article, data were extracted on study design, study objective, disease or health condition categories, study sample, description of RHIS data used, use of other data sources, analytic methods of RHIS data, strategies applied to circumvent data quality issues, and study findings. Due to the heterogeneity of the studies in terms of study design, study purpose, health conditions, and analysis methods, we thematically analyzed the studies according to research purpose, types of diseases studied, analytic methods applied, impact factor of journals in which the articles were published, and types of strategies used to circumvent RHIS data quality issues.

Results

Of the 1459 unique articles retrieved from the database search, 132 studies met the inclusion criteria after full-text screening and were thus included in the review. The characteristics of these studies are presented in Table 1. Our review identified studies from 37 different countries. Three quarters of the studies were from Sub-Saharan African countries (74%), followed by South Asia (11%). The vast majority of the studies were published in the last decade, and more than half were published after 2014 (55%), suggesting an increase in the use of RHIS data for research purposes over time. Most of the studies included an analysis of RHIS data (97%), and a few used RHIS data to inform the study but did not describe analysis of RHIS data. One study, for example, used information from RHIS to justify for the selection of the indicators to be used at the individual-level in their study. Among the studies that analyzed RHIS data, most utilized an ecological study design (79%). Of those, more than half included statistical inferences (61%), while the remaining studies only used RHIS data for descriptive purposes (39%). Nearly a fifth of the studies were mixed methods or case studies (18%), a third of which included statistical analyses of RHIS data (33%). A quarter of articles included a description of how they managed missing data (25%), while only a small number of studies described how they detected and dealt with extreme values (14%).

[Table 1 to be placed here.]

Types of disease and research purpose

Figure 2 shows the different research purposes for which RHIS data were used, along with the health topics investigated. The most common purpose of the studies was program evaluation (51%). RHIS data have been used to

evaluate a wide range of interventions, ranging from programs that targeted specific diseases to interventions or policies that affected multiple types of diseases or health services. These included: the effect of malaria control strategies³⁰⁻³⁶, user fee exemption policies³⁷⁻⁴⁰, health financing schemes⁴¹⁻⁴⁴, interventions on health governance⁴⁵⁻⁵³, the administration of new vaccines and vaccination campaigns⁵⁴⁻⁵⁶, as well as community-level interventions such as approaches to enhance community participation and improve referrals from traditional birth attendants in increasing the demand for maternal and child care⁵⁷⁻⁵⁹.

Additionally, RHIS data were used to monitor or assess service provision (23%) and to describe disease epidemiology (17%). Similar to the program evaluation studies, these studies also investigated a diverse set of health services and the allocation of healthcare resources. Some of these studies found large discrepancies between RHIS data and an estimated disease burden in populations or highlighted the lack of service provision. A few studies also used RHIS data to describe specific programs⁶⁰⁻⁶⁴, conduct impact evaluations (non-programmatic)⁶⁵⁻⁶⁸, and estimate costs^{69,70}. Most of the studies investigated a communicable disease (95%), of which malaria was most studied health condition (24%). A few studies focused on mental health (2%), diabetes mellitus (1%), and permanent tooth extraction (1%). Only two studies used RHIS data to research the health workforce or the equity of funding allocations^{71,72}.

Analytic methods using RHIS data

Among articles that conducted statistical analyses using RHIS data (n=68), time series analyses to test or account for trends were most commonly performed (25%), followed by geostatistical analyses (16%), pre-post comparisons (15%), interrupted time series (ITS) (10%), and difference-in-difference analyses (7%). Other longitudinal analyses (13%), other cross-sectional analyses (12%), and scenario analysis on cost effectiveness (2%) were also conducted. Table 2 presents the range of methodologies identified across studies using RHIS data, as well as the corresponding articles.

[Table 2 to be placed here.]

Time series analysis

Time series analysis using RHIS data was most often applied to evaluate programs and identify disease epidemiology, with one study assessing the impact of an infectious disease outbreak on primary health service utilization⁷³. Studies analyzed indicators using large quantities of monthly or yearly data to estimate change (range of time units: 5 – 168). For instance, two-thirds of the studies analyzed three or more years of monthly data. Many of the studies utilized the highly disaggregated nature of the data by using either facility or district level data, with the exception of two studies which modelled national trends^{33,74}. Studies commonly applied strategies to account for temporal autocorrelation and the correlation between geographical units, including generalized linear models⁵⁸, multi-level analysis^{75,76}, and ordinary least-squares regression with adjustment for seasonality and lag^{34,37,77}. Among studies that modelled multiple facilities or administrative regions, random effects were commonly applied to account for heterogeneity.

In addition to RHIS data, a number of included studies incorporated data from external sources in their models based on geographical location such as district or region. Studies of malaria, for example, commonly included climate data from satellites in their models to control for important temporal factors, for example precipitation, humidity, and temperature^{77,78}. Other studies incorporated information from other national community surveys, health facility surveys, and program data as covariates^{34,75}. While most studies controlled for potential confounders by including covariates in analytic models, one study on maternal health service applied propensity score matching to further remove biases from differences in covariate distribution³⁷.

Geostatistical analysis

Geostatistical analyses using RHIS data were predominantly conducted for epidemiological purposes and the monitoring and assessment of service provision by exploiting geospatial information included in the RHIS at the facility or district level. Three of the studies that applied geostatistical analysis were cross-sectional, while the remainder were spatial-temporal. About half of the studies focused on malaria, of which three compared and illustrated various kriging methods to provide a reliable estimate of malaria burden amid missing reporting^{79–81}, and one study applied geostatistical modeling to select the most relevant health facility indicators for severe malaria outcomes⁸². Studies on other topics investigated the spatial or spatial-temporal dynamics of malaria in pregnancy⁸³, childhood diarrhea⁸⁴, clustering of malaria and HIV⁸⁵, and meningitis⁸⁶. About half of the studies did not include data from external sources, and others triangulated data sourced from satellite data, Demographic and Health Surveys, national Malaria Indicator Surveys, and Service Delivery Indicator Surveys in their analyses. Studies that included covariates in the geostatistical analysis applied Bayesian hierarchical Poisson model or Bayesian geostatistical negative binomial models^{82,87,88}.

Pre-post comparison analysis

Pre-post comparison was commonly applied among studies that used RHIS data for program evaluation, and several studies used simple descriptive statistics to compare the periods before and after interventions. As pre-post comparison is subject to the limitation of temporal confounders and secular trends, two of the studies included contextual factors in regression modelling^{35,89}.

Interrupted time series analysis

Most of the studies that conducted ITS analysis used it to evaluate interventions, and one assessed the impact of an infectious disease outbreak on maternal and child health service use⁶⁸. The studies used large quantities of monthly data to model trend and level change (range of time unit: 44 – 132). RHIS data were minimally aggregated in these studies, which mostly analyzed facility or district level data, and similar to studies using time series analysis, accounted for autocorrelation through incorporating autoregressive structures or clustered standard errors in their modelling.

As ITS analyses are generally unaffected by confounding variables that do not change over time by design⁹⁰, baseline characteristics were typically not included in these models. Nonetheless, ITS analyses can be affected by time-varying confounders that rapidly change and some models included contextual factors from other data sources, such as climate and program data. To strengthen the quasi-experimental design, two studies also included a contrast group of time series to control for contextual changes that occurred at the same time as the interventions^{38,45}.

Difference-in-difference analysis

Five studies applied difference-in-difference techniques using a wide range of time periods (range of time units: 4 – 48) and level of geographical units (facility, district, provincial). Only one study included contextual characteristics from other data sources in its analysis. Analytic methods varied from descriptive comparison between and within intervention and control groups^{41,59,91,92}, to ordinary least square regression with propensity score matching⁴².

Impact of research using RHIS data

Most of the studies that conducted statistical analyses using RHIS data were published in journals with impact factors (88%, figure 3), two-thirds of which were two or higher, and more than a fifth of which were greater than three. Among those studies published in journals with the highest impact factors, most of them focused on program evaluation (53%), followed by monitoring and assessment of service provision (20%), epidemiology (20%) and impact evaluation (7%). These studies encompassed a range of health topics commonly studied using RHIS data.

Strategies to circumvent RHIS data quality issues

Data quality is commonly cited as a barrier to using RHIS data in research, and slightly more than a quarter of the included studies described the strategies that they used to handle missing data and/or identify extreme values (table 3). These strategies consisted of exclusion, imputation, interpolation, verification, and accounting for missing data in modeling. Exclusion of missing data was the most common practice, and among studies that used this technique, they excluded facilities from the analytic samples^{38,41,45,52,65,91,93-98}, restricted the study period based on explicit criteria^{54,99}, or applied sensitivity analysis to compare various exclusion criteria^{41,100,101}. Imputation methods varied from assigning specific values to the missing observation^{42,86,91,102-104}, to various modeling strategies such as conditional autoregressive model⁸⁷, generalized linear regression¹⁰³, and iterative singular value decomposition¹⁰³. A sensitivity analysis was also conducted to select a specific imputation strategy¹⁰³. Interpolation involves predicting values at unsampled locations. Methods described included the use of space-time kriging⁷⁹⁻⁸¹, and the adjustment of results by calibrating with other relevant information^{52,53,55}. Some studies assumed data were missing at random, which was accounted for in specific modeling methods such as mixed-effect models^{65,103}. When the source of data could be reached, some studies also described verifying the missing information using registries where the original data were recorded^{39,78,99,105,106}.

[Table 3 to be placed here.]

Slightly fewer articles described methods to identify and handle extreme values in the RHIS data, of which three types of strategies emerged: setting specific thresholds, visual inspection, and analytic assessment. Thresholds were set based on the distribution of the data, such as proportions or standard deviations from univariate regression. Several studies used visual inspection of outliers^{38,81}, while the use of jackknifing analysis and the identification of influential points through Cook's distance statistics were also applied^{107,108}. Upon identification of extreme values, several strategies were utilized: exclusion, replacement with the average value, replacement with the missing value, verification with a data source, or discounting the observation in statistical estimation. However, studies that replaced the extreme value with an explicit value potentially introduced bias into their estimates. A few studies also described the strategies applied to assess the reliability of the RHIS data, some of which were routine processes administered in the health systems^{39,106}.

Discussion

In recent years, there have been increased investments made to improve the quality of RHIS data in many LMICs. Over the same time period, we found an increase in published research using RHIS sourced data, especially over the past five years, likely due to the increased availability, accessibility, and quality of RHIS data¹⁸. While these studies have made contributions to the literature, we also found that the total number of studies conducted (n=132) remains a small part of the overall literature base on health system evaluation and performance in LMICs.

Malaria and maternal health conditions were the most commonly studied health conditions, despite the fact that RHISs collect data on a wide range of other diseases and conditions. In particular, the use of RHIS data for non-communicable diseases (NCDs) research was very limited. As LMICs are undergoing an epidemiologic transition and the importance of NCDs is increasing¹⁰⁹, LMIC health systems face the increasing challenges of addressing the dual burden of communicable and non-communicable diseases^{110,111}. In spite of the limited implementation of non-communicable diseases interventions¹¹¹, the few studies that used RHIS data for non-communicable disease research mainly analyzed the gap in service provision and estimated disease burden, highlighting the large unmet need for health care in affected populations. A couple of the studies described how their research was limited by data availability and quality,

such as the lack of diagnostic categories of the investigated health conditions in the RHIS. Future research should investigate how RHIS data on non-communicable diseases could better help to provide insights on its epidemiology and service provision to address these health conditions.

Our systematic review found that many of the studies took advantage of some of the features of RHIS data, in particular by exploiting the high frequency nature of these data at the level of health facilities, as well as combining external information to enhance estimations and enable assessing new research questions. The triangulation of populational health characteristics, environmental factors, and service coverage strengthens the analysis and the understanding of their influence¹¹². In addition, the overlay of different information in analyses of RHIS data allows for the advancement research methods. For instance, a recent study demonstrated how to assess the effects of facility readiness on severe malaria outcomes through constructing a composite facility readiness index based on health facility characteristics and spatial data, and using RHIS data as the outcome variable⁸². The detailed routine nature of RHIS data and the ability to link with other geographically based information, including data on population, environmental, health behavior, and facility characteristics, can generate high impact research and advance our understanding of disease epidemiology and health improvement efforts in LMICs.

Despite the increasing use of RHIS data for research purposes, the quality of these data remains imperfect and such issues should be identified and addressed in order to limit estimation error and bias. RHIS data quality issues remain a particular concern in some settings^{113–115}, however, other studies have shown that strategies that have been implemented to improve RHIS data across different international contexts can be successful^{5,116}. Multiple strategies were discussed in the articles we reviewed in our paper, including strategies to address common data quality issues such as missingness and data validity, for example the simple exclusion of missing data and various imputation and interpolation methods. However, the majority of the studies that used RHIS data did not describe the extent of the quality issues or the steps they took to overcome them. The use of sensitivity analysis in assessing the effect of specific cut-offs or methods was scarce. Explicit descriptions of the extent of the data quality issues and the reasons for selecting a particular approach should be encouraged in future research.

While our review used major databases and systematic methods, it nevertheless has some limitations that are worth noting. First, we included only peer-reviewed studies that were published in English, and therefore may have overlooked potentially relevant studies published in the grey literature or written in other languages. Additionally, given our focus on original research, we did not search the broader body of literature for books, reports, or grey literature. Our literature search also identified phrases that described health information systems in title and abstracts only, possibly resulting in the exclusion of studies that only mentioned RHIS data use in the full text. Finally, additional variants on these search terms may have generated more articles or a slightly different set of articles.

Conclusions

In this systematic review we summarized the use of data collected from RHISs in LMICs. Overall, we found that researchers are increasingly using data sourced from RHISs to conduct health system planning and evaluation studies in LMIC health systems, however these data likely remain underutilized by the broader research community. As many of the studies included in this review were published in prominent journals and were able to use strong quasi-experimental or geo-spatial methods, we believe this makes the case for greater use of these data for research purposes in the future, which will likely happen as RHIS data become more openly available to the research community. However, there is a need to help build the case to use these data for a broader range of health conditions and to develop more of a consensus on methods to deal with data imperfections, given that our findings underlined the limited use and comparison of these methods. That said, our review clearly demonstrates the feasibility of use RHIS data in conjunction

with rigorous study designs and analytic methods in LMICs. We suggest that future program evaluations should consider their use more broadly, to assess an increased variety of health conditions in conjunction with, or as a replacement for, household or facility survey methods.

Abbreviations

DHIS 2	District health information system 2
HMIS	Health management information system
ITS	Interrupted time series
LMIC	Low- and middle-income country
NCD	Non-communicable disease
PRISMA	Preferred reporting items for systematic reviews and meta-analyses
RHIS	Routine health information system

Declarations

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Authors' contributions: YWH, KH, and KAG identified the search strategy. YWH and KH conducted the literature screening, and KAG was consulted when required. YWH, KH, and BRI extracted data from the included studies. YWH wrote the manuscript. KAG and MRL provided feedback and edits on the manuscript. All authors read and approved the final manuscript.

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References

1. Hotchkiss DR, Diana ML, Foreit KGF. How can routine health information systems improve health systems functioning in low and middle-income countries? assessing the evidence base. *Adv Health Care Manag.* 2012. doi:10.1108/S1474-8231(2012)0000012006
2. AbouZahr C, Boerma T. Health information systems: the foundations of public health. *Bull World Health Organ.* 2005;83:578-583.
3. World Health Organization. Everybody's business—strengthening health systems to improve health outcomes: WHO's framework for action. 2007.

4. Lippeveld T, Sauerborn R, Bodart C, Organization WH. Design and implementation of health information systems. 2000.
5. Mutale W, Chintu N, Amoroso C, et al. Improving health information systems for decision making across five sub-Saharan African countries: Implementation strategies from the African Health Initiative. *BMC Health Serv Res*. 2013;13(Suppl 2):1-12.
6. Hogan DR, Stevens GA, Hosseinpoor AR, Boerma T. Monitoring universal health coverage within the Sustainable Development Goals: development and baseline data for an index of essential health services. *Lancet Glob Heal*. 2018;6(2):e152-e168. doi:[https://doi.org/10.1016/S2214-109X\(17\)30472-2](https://doi.org/10.1016/S2214-109X(17)30472-2)
7. Cutts FT, Claquin P, Danovaro-Holliday MC, Rhoda DA. Monitoring vaccination coverage: Defining the role of surveys. *Vaccine*. 2016;34(35):4103-4109. doi:<https://doi.org/10.1016/j.vaccine.2016.06.053>
8. Wagenaar BH, Sherr K, Fernandes Q, Wagenaar AC. Using routine health information systems for well-designed health evaluations in low- and middle-income countries. *Health Policy Plan*. 2016;31(1):129-135. doi:10.1093/heapol/czv029
9. World Health Organization. *Monitoring the Building Blocks of Health Systems: A Handbook of Indicators and Their Measurement Strategies*. Geneva, Switzerland; 2010.
10. World Health Organization. *Framework and Standards for Country Health Information Systems (Second Edition)*. Geneva, Switzerland; 2012.
11. Warren AE, Wyss K, Shakarishvili G, Atun R, de Savigny D. Global health initiative investments and health systems strengthening: a content analysis of global fund investments. *Global Health*. 2013;9(1):30. doi:10.1186/1744-8603-9-30
12. Gimbel S, Mwanza M, Nisingizwe MP, et al. Improving data quality across 3 sub-Saharan African countries using the Consolidated Framework for Implementation Research (CFIR): Results from the African Health Initiative. *BMC Health Serv Res*. 2017. doi:10.1186/s12913-017-2660-y
13. Kiberu VM, Matovu JKB, Makumbi F, Kyoziira C, Mukooyo E, Wanyenze RK. Strengthening district-based health reporting through the district health management information software system: the Ugandan experience. *BMC Med Inform Decis Mak*. 2014;14:40. doi:10.1186/1472-6947-14-40
14. Hazel E, Wilson E, Anifalaje A, Sawadogo-Lewis T, Heidkamp R. Building integrated data systems for health and nutrition program evaluations: lessons learned from a multi-country implementation of a DHIS 2-based system. *J Glob Health*. 2018;8(2):20307. doi:10.7189/jogh.08.020307
15. Dehnavieh R, Haghdoost A, Khosravi A, et al. The District Health Information System (DHIS2): A literature review and meta-synthesis of its strengths and operational challenges based on the experiences of 11 countries. *Heal Inf Manag J*. June 2018:1833358318777713. doi:10.1177/1833358318777713
16. Githinji S, Oyando R, Malinga J, et al. Completeness of malaria indicator data reporting via the District Health Information Software 2 in Kenya, 2011–2015. *Malar J*. 2017;16:344. doi:10.1186/s12936-017-1973-y
17. MEASURE Evaluation. *Lessons in Health Information System Strengthening: What Worked in the Democratic Republic of the Congo*. Chapel Hill, USA; 2019. <https://www.measureevaluation.org/resources/publications/wp-19-229>.
18. Ashton RA, Bennett A, Yukich J, Bhattarai A, Keating J, Eisele TP. Methodological considerations for use of routine health information system data to evaluate malaria program impact in an era of declining malaria transmission. *Am J Trop Med Hyg*. 2017. doi:10.4269/ajtmh.16-0734
19. Rowe AK, Kachur SP, Yoon SS, Lynch M, Slutsker L, Steketee RW. Caution is required when using health facility-based data to evaluate the health impact of malaria control efforts in Africa. *Malar J*. 2009;8(1):209.

20. Hazel E, Requejo J, David J, Bryce J. Measuring Coverage in MNCH: Evaluation of Community-Based Treatment of Childhood Illnesses through Household Surveys. *PLoS Med*. 2013. doi:10.1371/journal.pmed.1001384
21. Lim SS, Stein DB, Charrow A, Murray CJL. Tracking progress towards universal childhood immunisation and the impact of global initiatives: a systematic analysis of three-dose diphtheria, tetanus, and pertussis immunisation coverage. *Lancet*. 2008;372(9655):2031-2046. doi:https://doi.org/10.1016/S0140-6736(08)61869-3
22. Cutts FT, Izurieta HS, Rhoda DA. Measuring Coverage in MNCH: Design, Implementation, and Interpretation Challenges Associated with Tracking Vaccination Coverage Using Household Surveys. *PLOS Med*. 2013;10(5):e1001404. https://doi.org/10.1371/journal.pmed.1001404.
23. Sychareun V, Hansana V, Phengsavanh A, Chaleunvong K, Eunyoung K, Durham J. Data verification at health centers and district health offices in Xiengkhouang and Houaphanh Provinces, Lao PDR. *BMC Health Serv Res*. 2014;14(1):255. doi:10.1186/1472-6963-14-255
24. Gloyd S, Wagenaar BH, Woelk GB, Kalibala S. Opportunities and challenges in conducting secondary analysis of HIV programmes using data from routine health information systems and personal health information. *J Int AIDS Soc*. 2016. doi:10.7448/IAS.19.5.20847
25. Aqil A, Lippeveld T, Hozumi D. PRISM framework: a paradigm shift for designing, strengthening and evaluating routine health information systems. *Health Policy Plan*. 2009;24(3):217-228.
26. Short Fabic M, Choi Y, Bird S. A systematic review of Demographic and Health Surveys: data availability and utilization for research. *Bull World Health Organ*. 2012. doi:10.2471/BLT.11.095513
27. Fernandes QF, Wagenaar BH, Anselmi L, Pfeiffer J, Gloyd S, Sherr K. Effects of health-system strengthening on under-5, infant, and neonatal mortality: 11-year provincial-level time-series analyses in Mozambique. *Lancet Glob Heal*. 2014. doi:10.1016/S2214-109X(14)70276-1
28. Ratnayake R, Ratto J, Hardy C, et al. The effects of an integrated community case management strategy on the appropriate treatment of children and child mortality in Kono District, Sierra Leone: A program evaluation. *Am J Trop Med Hyg*. 2017. doi:10.4269/ajtmh.17-0040
29. Bishai DM. Global initiatives in universal childhood immunisation. *Lancet*. 2008;372(9655):2004-2005. doi:https://doi.org/10.1016/S0140-6736(08)61853-X
30. Seitio-Kgokgwe O, Mashalla Y, Seloilwe E, et al. Developing block wise composite health index in Yavatmal district, Maharashtra state, India: An analysis of available data. *Malar J*. 2017;16(1 PG-):35. doi:10.4103/2008-7802.160056
31. Pagnoni F, Convelbo N, Tiendrebeogo J, Cousens S, Esposito F. A community-based programme to provide prompt and adequate treatment of presumptive malaria in children. *Trans R Soc Trop Med Hyg*. 1997;91(5 PG-512-7):512-517. NS -.
32. Rehman A, Mann A, Schwabe C, et al. Five years of malaria control in the continental region, Equatorial Guinea. *Malar J*. 2013;12(PG-154):154. doi:10.1186/1475-2875-12-154
33. Nyarango P, Gebremeskel T, Mebrahtu G, et al. A steep decline of malaria morbidity and mortality trends in Eritrea between 2000 and 2004: the effect of combination of control methods. *Malar J*. 2006;5(PG-33):33. doi:10.1186/1475-2875-5-33
34. Over M, Bakotéé B, Velayudhan R, Wilikai P, Graves PM. Impregnated Nets Cannot Fully Substitute for DDT: The Field Effectiveness of Alternative Methods of Malaria Prevention in Solomon Islands, 1993-99. *World Bank, Policy Res Work Pap Ser 3044, 2003*. 2003;(PG-). https://libproxy.wlu.ca/login?url=https://search.proquest.com/docview/56217804?accountid=15090 NS -.
35. Okullo AE, Matovu JKB, Ario AR, et al. Malaria incidence among children less than 5 years during and after cessation of indoor residual spraying in Northern Uganda. *Malar J*. 2017;16(1 PG-319):319. doi:10.1186/s12936-

36. Sande S, Zimba M, Mberikunashe J, Tangwena A, Chimusoro A. Progress towards malaria elimination in Zimbabwe with special reference to the period 2003-2015. *Malar J.* 2017. doi:10.1186/s12936-017-1939-0
37. Manthalu G. User fee exemption and maternal health care utilisation at mission health facilities in Malawi: An application of disequilibrium theory of demand and supply. *Health Econ.* 2019;(PG-). doi:10.1002/hec.3856
38. Nguyen H, Zombré D, Ridde V, De Allegri M. The impact of reducing and eliminating user fees on facility-based delivery: a controlled interrupted time series in Burkina Faso. *Health Policy Plan.* 2018;33(8 PG-948-956):948-956. doi:10.1093/heapol/czy077
39. Njuguna J, Kamau N, Muruka C. Impact of free delivery policy on utilization of maternal health services in county referral hospitals in Kenya. *BMC Health Serv Res.* 2017. doi:10.1186/s12913-017-2376-z
40. Ridde V, Richard F, Bicaba A, Queuille L, Conombo G. The national subsidy for deliveries and emergency obstetric care in Burkina Faso. *Health Policy Plan.* 2011. doi:10.1093/heapol/czr060
41. Steenland M, Robyn P, Compaore P, et al. Performance-based financing to increase utilization of maternal health services: Evidence from Burkina Faso. *SSM - Popul Heal.* 2017;3(PG-179-184):179-184. doi:10.1016/j.ssmph.2017.01.001
42. Falisse JB, Ndayishimiye J, Kamenyero V, Bossuyt M. Performance-based financing in the context of selective free health-care: An evaluation of its effects on the use of primary health-care services in Burundi using routine data. *Health Policy Plan.* 2015. doi:10.1093/heapol/czu132
43. Witter S, Zulfiquar T, Javeed S, Khan A, Bari A. Paying health workers for performance in Battagram district, Pakistan. *Hum Resour Health.* 2011. doi:10.1186/1478-4491-9-23
44. Ir P, Horemans D, Souk N, Van Damme W. Using targeted vouchers and health equity funds to improve access to skilled birth attendants for poor women: A case study in three rural health districts in Cambodia. *BMC Pregnancy Childbirth.* 2010. doi:10.1186/1471-2393-10-1
45. Iyer HS, Hirschhorn LR, Nisingizwe MP, et al. Impact of a district-wide health center strengthening intervention on healthcare utilization in rural Rwanda: Use of interrupted time series analysis. *PLoS One.* 2017;12(8 PG-e0182418):e0182418. doi:http://dx.doi.org/10.1371/journal.pone.0182418
46. Lima RT de S, Fernandes TG, Balieiro AA da S, et al. Primary Health Care in Brazil and the Mais Médicos (More Doctors) Program: an analysis of production indicators. *Cien Saude Colet.* 2016. doi:10.1590/1413-81232015219.15412016
47. Mikkelsen-Lopez I, Cowley P, Kasale H, Mbuya C, Reid G, de Savigny D. Essential medicines in Tanzania: does the new delivery system improve supply and accountability? *Heal Syst.* 2014. doi:10.1057/hs.2013.14
48. Mphatswe W, Mate K, Bennett B, et al. Improving public health information: a data quality intervention in KwaZulu-Natal, South Africa. *Bull World Health Organ.* 2012. doi:10.2471/blt.11.092759
49. Newbrander W, Ickx P, Feroz F, Stanekzai H. Afghanistan's Basic Package of Health Services: Its development and effects on rebuilding the health system. *Glob Public Health.* 2014;9(SUPPL.1 PG-S6-S28):S6-S28. doi:10.1080/17441692.2014.916735
50. Santos LMP, Oliveira A, Trindade JS, et al. Implementation research: towards universal health coverage with more doctors in Brazil. *Bull World Health Organ.* 2017. doi:10.2471/BLT.16.178236
51. Scott VE, Sanders D. Evaluation of how integrated HIV and TB programs are implemented in South Africa and the implications for rural-urban equity. *Rural Remote Health.* 2013.
52. Vindigni SM, Riley PL, Kimani F, et al. Kenya's emergency-hire nursing programme: A pilot evaluation of health service delivery in two districts. *Hum Resour Health.* 2014;12(1 PG-). doi:10.1186/1478-4491-12-16

53. Field E, Usurup J, Nathan S, Rosewell A. Contextual factors and health service performance from the perspective of the provincial health administrators in Papua New Guinea. *Rural Remote Health*. 2018. doi:10.22605/RRH4484
54. Inchauste L, Patzi M, Halvorsen K, Solano S, Montesano R, Iñiguez V. Impact of rotavirus vaccination on child mortality, morbidity, and rotavirus-related hospitalizations in Bolivia. *Int J Infect Dis*. 2017. doi:10.1016/j.ijid.2017.06.006
55. Nanyunja M, Lewis RF, Makumbi I, et al. Impact of Mass Measles Campaigns among Children Less Than 5 Years Old in Uganda. *J Infect Dis*. 2003. doi:10.1086/368026
56. Tafesse B, Tekle E, Wondwossen L, et al. Effects of polio eradication activities on routine immunization: lessons from the 2013 outbreak response in Somali region of Ethiopia. *Pan Afr Med J*. 2017;27(Suppl 2 PG-4):4. doi:10.11604/pamj.supp.2017.27.2.10569
57. Mochache V, Irungu E, El-Busaidy H, Temmerman M, Gichangi P. "Our voices matter": A before-after assessment of the effect of a community-participatory intervention to promote uptake of maternal and child health services in Kwale, Kenya 11 Medical and Health Sciences 1117 Public Health and Health Services. *BMC Health Serv Res*. 2018;18(1 PG-). doi:10.1186/s12913-018-3739-9
58. Kitui J, Dutton V, Bester D, Ndirangu R, Wangai S, Ngugi S. Traditional Birth Attendant reorientation and Motherpacks incentive's effect on health facility delivery uptake in Narok County, Kenya: An impact analysis. *BMC Pregnancy Childbirth*. 2017;17(1 PG-125):125. doi:10.1186/s12884-017-1307-7
59. Paudel D, Shrestha I, Siebeck M, Rehfuess E. Impact of the community-based newborn care package in Nepal: a quasi-experimental evaluation. *BMJ Open*. 2017;7(10 PG-e015285):e015285. doi:10.1136/bmjopen-2016-015285
60. Khurmi M, Sayinzoga F, Berhe A, et al. Newborn Survival Case Study in Rwanda - Bottleneck Analysis and Projections in Key Maternal and Child Mortality Rates Using Lives Saved Tool (LiST). *Int J MCH AIDS*. 2017;6(2 PG-93-108):93-108. doi:10.21106/ijma.214
61. Lindgren TG, Deutsch K, Schell E, et al. Using mobile clinics to deliver HIV testing and other basic health services in rural Malawi. *Rural Remote Health*. 2011.
62. Mapira P, Morgan C. The contribution of church health services to maternal health care provision in Papua New Guinea. *P N G Med J*. 2011.
63. Robertson LJ, Szabo CP. Community mental health services in southern Gauteng: An audit using Gauteng district health information systems data. *South African J Psychiatry*. 2017;23(1 PG-a1055):a1055. doi:http://dx.doi.org/10.4102/sajpsychiatry.v23i0.1055
64. Yellapurkar M V. Health management information system in leprosy control programme. *Indian J Lepr*. 1984;56(1 PG-86-97):86-97. <https://www.scopus.com/inward/record.uri?eid=2-s2.0-0021634802&partnerID=40&md5=9c9d755790bfa4154b2372c12bdeeb37> NS -.
65. Wagenaar BH, Augusto O, Beste J, et al. The 2014–2015 Ebola virus disease outbreak and primary healthcare delivery in Liberia: Time-series analyses for 2010–2016. *PLoS Med*. 2018;15(2):e1002508. doi:10.1371/journal.pmed.1002508
66. Quaglio G, Pizzol D, Bome D, et al. Maintaining Maternal and Child Health Services During the Ebola Outbreak: Experience from Pujehun, Sierra Leone. *PLoS Curr*. 2016;8. doi:10.1371/CURRENTS.OUTBREAKS.D67AEA257F572201F835772D7F188BA5
67. Sesay T, Denisiuk O, Shringarpure KK, et al. Paediatric care in relation to the 2014–2015 Ebola outbreak and general reporting of deaths in Sierra Leone. *Public Heal Action*. 2017. doi:10.5588/pha.16.0088
68. Sochas L, Channon AA, Nam S. Counting indirect crisis-related deaths in the context of a low-resilience health system: The case of maternal and neonatal health during the Ebola epidemic in Sierra Leone. *Health Policy Plan*. 2017;32(suppl_3):iii32-iii39. doi:10.1093/heapol/czx108

69. Cassim N, Coetzee LM, Schnippel K, Glencross DK. Estimating the cost-per-result of a national reflexed Cryptococcal antigenaemia screening program: Forecasting the impact of potential HIV guideline changes and treatment goals. *PLoS One*. 2017;12(8 PG-e0182154):e0182154. doi:http://dx.doi.org/10.1371/journal.pone.0182154
70. Novaes H, Itria A, Silva G, Sartori A, Rama C, Soárez P. Annual national direct and indirect cost estimates of the prevention and treatment of cervical cancer in Brazil. *Clinics (Sao Paulo)*. 2015;70(4 PG-289-95):289-295. doi:10.6061/clinics/2015(04)12
71. Namaganda G, Oketcho V, Maniple E, Viadro C. Making the transition to workload-based staffing: using the Workload Indicators of Staffing Need method in Uganda. *Hum Resour Health*. 2015;13(PG-89):89. doi:10.1186/s12960-015-0066-7
72. Kusemererwa D, Alban A, Obua OT, Trap B. An exploratory study on equity in funding allocation for essential medicines and health supplies in Uganda's public sector. *BMC Health Serv Res*. 2016;16((Obua) Ministry of Health, Pharmacy Division, Lourdel Road, Wandegaya, Kampala, Uganda PG-453):453. doi:http://dx.doi.org/10.1186/s12913-016-1698-6
73. Wagenaar BH, Augusto O, Beste J, et al. The 2014-2015 Ebola virus disease outbreak and primary healthcare delivery in Liberia: Time-series analyses for 2010-2016. *PLoS Med*. 2018;15(2 PG-e1002508):e1002508. doi:10.1371/journal.pmed.1002508
74. Nanyunja M, Lewis RF, Makumbi I, et al. Impact of mass measles campaigns among children less than 5 years old in Uganda. *J Infect Dis*. 2003;187(SUPPL. 1 PG-S63-S68):S63-S68. doi:10.1086/368026
75. Luboga SA, Stover B, Lim T, et al. Did PEPFAR investments result in health system strengthening? A retrospective longitudinal study measuring non-HIV health service utilization at the district level. *Health Policy Plan*. 2016;31(7 PG-897-909):897-909. doi:10.1093/heapol/czw009
76. Wagenaar B, Gimbel S, Hoek R, et al. Effects of a health information system data quality intervention on concordance in Mozambique: time-series analyses from 2009-2012. *Popul Health Metr*. 2015;13(PG-9):9. doi:10.1186/s12963-015-0043-3
77. Gunda R, Chimbari M, Mukaratirwa S. Assessment of Burden of Malaria in Gwanda District, Zimbabwe, Using the Disability Adjusted Life Years. *Int J Environ Res Public Health*. 2016;13(2 PG-244):244. doi:10.3390/ijerph13020244
78. Hajison PL, Mwakikunga BW, Mathanga DP, Feresu SA. Seasonal variation of malaria cases in children aged less than 5 years old following weather change in Zomba district, Malawi. *Malar J*. 2017. doi:10.1186/s12936-017-1913-x
79. Gething PW, Noor AM, Goodman CA, et al. Information for decision making from imperfect national data: Tracking major changes in health care use in Kenya using geostatistics. *BMC Med*. 2007. doi:10.1186/1741-7015-5-37
80. Gething PW, Noor AM, Gikandi PW, et al. Improving imperfect data from health management information systems in Africa using space-time geostatistics. *PLoS Med*. 2006;3(6 PG-0825-0831):825-831. doi:10.1371/journal.pmed.0030271
81. Gething P, Atkinson P, Noor A, Gikandi P, Hay S, Nixon M. A local space-time kriging approach applied to a national outpatient malaria dataset. *Comput Geosci*. 2007;33(10 PG-1337-1350):1337-1350. doi:10.1016/j.cageo.2007.05.006
82. Ssempiira J, Kasirye I, Kissa J, et al. Measuring health facility readiness and its effects on severe malaria outcomes in Uganda. *Sci Rep*. 2018;8(1 PG-17928):17928. doi:10.1038/s41598-018-36249-8
83. Kamuliwo M, Kirk K, Chanda E, et al. Spatial patterns and determinants of malaria infection during pregnancy in Zambia. *Trans R Soc Trop Med Hyg*. 2015;109(8 PG-514-21):514-521. doi:10.1093/trstmh/trv049

84. Beyene H, Deressa W, Kumie A, Grace D. Spatial, temporal, and spatiotemporal analysis of under-five diarrhea in Southern Ethiopia. *Trop Med Health*. 2018;46(PG-18):18. doi:10.1186/s41182-018-0101-1
85. Gwitira I, Murwira A, Mberikunashé J, Masocha M. Spatial overlaps in the distribution of HIV/AIDS and malaria in Zimbabwe. *BMC Infect Dis*. 2018;18(1 PG-598):598. doi:10.1186/s12879-018-3513-y
86. Philippon S, Broutin H, Constantin de Magny G, et al. Meningococcal meningitis in Mali: a long-term study of persistence and spread. *Int J Infect Dis*. 2009;13(1 PG-103-9):103-109. doi:10.1016/j.ijid.2008.05.1223
87. Bennett A, Yukich J, Miller JM, et al. A methodological framework for the improved use of routine health system data to evaluate national malaria control programs: Evidence from Zambia. *Popul Health Metr*. 2014. doi:10.1186/s12963-014-0030-0
88. Sartorius B, Cohen C, Chirwa T, Ntshoe G, Puren A, Hofman K. Identifying high-risk areas for sporadic measles outbreaks: lessons from South Africa. *Bull World Health Organ*. 2013. doi:10.2471/BLT.12.110726
89. Njuguna J. Effect of eliminating open defecation on diarrhoeal morbidity: An ecological study of Nyando and Nambale sub-counties, Kenya. *BMC Public Health*. 2016;16(1 PG-). doi:10.1186/s12889-016-3421-2
90. Bernal JL, Cummins S, Gasparrini A. Interrupted time series regression for the evaluation of public health interventions: A tutorial. *Int J Epidemiol*. 2017. doi:10.1093/ije/dyw098
91. Assegaai T, Reagon G, Schneider H. Evaluating the effect of ward-based outreach teams on primary healthcare performance in North West Province, South Africa: A plausibility design using routine data. *S Afr Med J*. 2018;108(4 PG-329-335):329-335. doi:10.7196/SAMJ.2017.v108i4.12755
92. Anwari Z, Shukla M, Maseed B, et al. Implementing people-centred health systems governance in 3 provinces and 11 districts of Afghanistan: a case study. *Confl Health*. 2015;9(PG-2):2. doi:10.1186/1752-1505-9-2
93. Ruton H, Musabyimana A, Gaju E, et al. The impact of an mHealth monitoring system on health care utilization by mothers and children: An evaluation using routine health information in Rwanda. *Health Policy Plan*. 2018. doi:10.1093/heapol/czy066
94. Sibomana H, Rugambwa C, Mwenda JM, et al. Impact of routine rotavirus vaccination on all-cause and rotavirus hospitalizations during the first four years following vaccine introduction in Rwanda. *Vaccine*. 2018. doi:10.1016/j.vaccine.2018.01.072
95. Gebrehiwot TG, Sebastian MS, Edin K, Goicolea I. The Health Extension Program and its association with change in utilization of selected maternal health services in tigray region, Ethiopia: A segmented linear regression analysis. *PLoS One*. 2015. doi:10.1371/journal.pone.0131195
96. Buchmann EJ, Mnyani CN, Frank KA, Chersich MF, McIntyre JA. Declining maternal mortality in the face of persistently high HIV prevalence in a middle-income country. *BJOG An Int J Obstet Gynaecol*. 2015. doi:10.1111/1471-0528.13064
97. Walsh A, Ndubani P, Simbaya J, Dicker P, Brugha R. Task sharing in Zambia: HIV service scale-up compounds the human resource crisis. *BMC Health Serv Res*. 2010. doi:10.1186/1472-6963-10-272
98. Heymann DL, Chen L, Takemi K, et al. Global health security: The wider lessons from the west African Ebola virus disease epidemic. *Lancet*. 2015. doi:10.1016/S0140-6736(15)60858-3
99. Luboga SA, Stover B, Lim TW, et al. Did PEPFAR investments result in health system strengthening? A retrospective longitudinal study measuring non-HIV health service utilization at the district level. *Health Policy Plan*. 2016. doi:10.1093/heapol/czw009
100. Mugeni C, Levine AC, Munyaneza RM, et al. Nationwide implementation of integrated community case management of childhood illness in Rwanda. *Glob Heal Sci Pract*. 2014. doi:10.9745/ghsp-d-14-00080
101. Verguet S phane, Jassat W, Bertram MY, et al. Supplementary immunization activities (SIAs) in South Africa: Comprehensive economic evaluation of an integrated child health delivery platform. *Glob Health Action*. 2013.

doi:10.3402/gha.v6i0.20056

102. Amouzou A, Kachaka W, Banda B, Chimzimu M, Hill K, Bryce J. Monitoring child survival in “real time” using routine health facility records: Results from Malawi. *Trop Med Int Heal*. 2013. doi:10.1111/tmi.12167
103. Inguane CA, Gloyd S, Manuel JL, et al. Assessment of linkages from HIV testing to enrolment and retention in HIV care in Central Mozambique. *J Int AIDS Soc*. 2016. doi:10.7448/IAS.19.5.20846
104. Wagenaar BH, Gimbel S, Hoek R, et al. Effects of a health information system data quality intervention on concordance in Mozambique: Time-series analyses from 2009-2012. *Popul Health Metr*. 2015. doi:10.1186/s12963-015-0043-3
105. Pires G, Rosa M, Zangarote M, Chicumbe S. Determinants of stillbirths occurred in health facilities in Zambezia province, Mozambique (2013-2014). *Rev Bras Saude Matern Infant*. 2016;16(4 PG-415-420):415-420. doi:10.1590/1806-93042016000400004
106. Njuguna J. Effect of eliminating open defecation on diarrhoeal morbidity: An ecological study of Nyando and Nambale sub-counties, Kenya. *BMC Public Health*. 2016. doi:10.1186/s12889-016-3421-2
107. Sebastian MS, Lemma H. Efficiency of the health extension programme in Tigray, Ethiopia: A data envelopment analysis. *BMC Int Health Hum Rights*. 2010. doi:10.1186/1472-698X-10-16
108. Jat TR, Sebastian MS. Technical efficiency of public district hospitals in Madhya Pradesh, India: a data envelopment analysis. *Glob Health Action*. 2013. doi:10.3402/gha.v6i0.21742
109. Stanaway JD, Afshin A, Gakidou E, et al. Global, regional, and national comparative risk assessment of 84 behavioural, environmental and occupational, and metabolic risks or clusters of risks for 195 countries and territories, 1990-2017: A systematic analysis for the Global Burden of Disease Stu. *Lancet*. 2018. doi:10.1016/S0140-6736(18)32225-6
110. Remais J V, Zeng G, Li G, Tian L, Engelgau MM. Convergence of non-communicable and infectious diseases in low- and middle-income countries. *Int J Epidemiol*. 2013. doi:10.1093/ije/dys135
111. Heller O, Somerville C, Suggs LS, et al. The process of prioritization of non-communicable diseases in the global health policy arena. *Health Policy Plan*. June 2019. doi:10.1093/heapol/czz043
112. Victora CG, Black RE, Boerma JT, Bryce J. Measuring impact in the Millennium development Goal era and beyond: A new approach to large-scale effectiveness evaluations. *Lancet*. 2011. doi:10.1016/S0140-6736(10)60810-0
113. Endriyas M, Alano A, Mekonnen E, et al. Understanding performance data: Health management information system data accuracy in Southern Nations Nationalities and People’s Region, Ethiopia. *BMC Health Serv Res*. 2019. doi:10.1186/s12913-019-3991-7
114. Ward K, Mugenyi K, Benke A, et al. Enhancing workforce capacity to improve vaccination data quality, Uganda. *Emerg Infect Dis*. 2017. doi:10.3201/eid2313.170627
115. Maïga A, Jiwani SS, Mutua MK, et al. Generating statistics from health facility data: the state of routine health information systems in Eastern and Southern Africa. *BMJ Glob Heal*. 2019;4(5):e001849. doi:10.1136/bmjgh-2019-001849
116. Necochea E, Tripathi V, Kim YM, et al. Implementation of the Standards-Based Management and Recognition approach to quality improvement in maternal, newborn, and child health programs in low-resource countries. *Int J Gynecol Obstet*. 2015. doi:10.1016/j.ijgo.2015.04.003
117. Wright J, Vazé P, Russell G, et al. Seasonal aspects of weight-for-age in young children in Zimbabwe. *Public Health Nutr*. 2001. doi:10.1079/phn2000100
118. Gunda R, Chimbari M, Shamu S, Sartorius B, Mukaratirwa S. Malaria incidence trends and their association with climatic variables in rural Gwanda, Zimbabwe, 2005-2015. *Malar J*. 2017;16(1 PG-393):393. doi:10.1186/s12936-017-2036-0

119. De Melo Cunha MAG, Lino PA, Dos Santos TR, Vasconcelos M, Lucas SD, De Abreu MHNG. A 15-year time-series study of tooth extraction in Brazil. *Med (United States)*. 2015;94(47 PG-e1924):e1924. doi:<http://dx.doi.org/10.1097/MD.0000000000001924>
120. Larsen DA, Winters A, Cheelo S, et al. Shifting the burden or expanding access to care? Assessing malaria trends following scale-up of community health worker malaria case management and reactive case detection. *Malar J*. 2017;16(1 PG-441):441. doi:10.1186/s12936-017-2088-1
121. Ngabo F, Tate J, Gatera M, et al. Effect of pentavalent rotavirus vaccine introduction on hospital admissions for diarrhoea and rotavirus in children in Rwanda: a time-series analysis. *Lancet Glob Heal*. 2016;4(2 PG-e129-36):e129-36. doi:10.1016/S2214-109X(15)00270-3
122. Ir P, Korachais C, Chheng K, Horemans D, Van Damme W, Meessen B. Boosting facility deliveries with results-based financing: A mixed-methods evaluation of the government midwifery incentive scheme in Cambodia. *BMC Pregnancy Childbirth*. 2015. doi:10.1186/s12884-015-0589-x
123. Landoh E, Tchamdja P, Saka B, et al. Morbidity and mortality due to malaria in Est Mono district, Togo, from 2005 to 2010: a times series analysis. *Malar J*. 2012;11(PG-389):389. doi:10.1186/1475-2875-11-389
124. Björkman A, Shakely D, Ali AS, et al. From high to low malaria transmission in Zanzibar - Challenges and opportunities to achieve elimination. *BMC Med*. 2019. doi:10.1186/s12916-018-1243-z
125. Oguttu DW, Matovu JKB, Okumu DC, et al. Rapid reduction of malaria following introduction of vector control interventions in Tororo District, Uganda: a descriptive study. *Malar J*. 2017. doi:10.1186/s12936-017-1871-3
126. Schoon MG. Impact of inter-facility transport on maternal mortality in the free state province. *South African Med J*. 2013;103(8 PG-534-537):534-537. doi:10.7196/SAMJ.6828
127. Kakchapati S, Ardkaew J. Modeling of malaria incidence in nepal. *J Res Health Sci*. 2011;1(1 PG-7-13):7-13. <https://www.scopus.com/inward/record.uri?eid=2-s2.0-79959717379&partnerID=40&md5=259481ca13eda8f1ef59d22356034ffd> NS -.
128. Mufunda J, Nyarango P, Usman A, et al. Roll back malaria - An African success story in Eritrea. *South African Med J*. 2007.
129. Kamuliwo M, Chanda E, Haque U, et al. The changing burden of malaria and association with vector control interventions in Zambia using district-level surveillance data, 2006-2011. *Malar J*. 2013. doi:10.1186/1475-2875-12-437
130. Philippon S, Broutin H, Constantin de Magny G, et al. Meningococcal meningitis in Mali: a long-term study of persistence and spread. *Int J Infect Dis*. 2009. doi:10.1016/j.ijid.2008.05.1223
131. Bosomprah S, Dotse-Gborgbortsi W, Aboagye P, Matthews Z. Use of a spatial scan statistic to identify clusters of births occurring outside Ghanaian health facilities for targeted intervention. *Int J Gynecol Obstet*. 2016. doi:10.1016/j.ijgo.2016.04.016
132. Himanshu M, Källestål C. Regional inequity in complete antenatal services and public emergency obstetric care is associated with greater burden of maternal deaths: Analysis from consecutive district level facility survey of Karnataka, India. *Int J Equity Health*. 2017. doi:10.1186/s12939-017-0573-3
133. Souza ML. Registro de distúrbios mentais no Sistema de Informação da Atenção Básica do Brasil, 2014. *Epidemiol e Serv saude Rev do Sist Unico Saude do Bras*. 2016;25(2 PG-405-410):405-410. doi:10.5123/S1679-49742016000200018
134. Wanzira H, Muyinda R, Lochoro P, et al. Quality of care for children with acute malnutrition at health center level in Uganda: a cross sectional study in West Nile region during the refugee crisis. *BMC Health Serv Res*. 2018. doi:10.1186/s12913-018-3366-5

Tables

Table 1. Characteristics of research studies that used RHIS data.

	n	Percent
Geographical region		
East Asia and Pacific	8	6.1
Latin America and the Caribbean	9	6.8
Middle East and North Africa	2	1.5
South Asia	15	11.4
Sub-Saharan Africa	98	74.2
Year of publication		
<2000	3	2.3
2000-2004	7	5.3
2005-2009	10	7.6
2010-2014	40	30.3
2015-2019	72	54.5
RHIS data as source or to inform study		
Data source	128	97.0
Inform study	4	3.0
Types of study design		
Ecological study - cross-sectional	13	9.8
Ecological study - longitudinal	51	38.6
Ecological study - descriptive	41	31.1
Case study	11	8.3
Mixed methods study	13	9.8
Cross-sectional study	1	0.8
Pre- and post-intervention study	1	0.8
Nested clustered randomized controlled trial	1	0.8
Data use purpose		
Program evaluation	67	50.8
Epidemiology	23	17.4
Monitoring and assessment of service provisions	30	22.7
Program description	6	4.5
Impact evaluation	4	3.0
Cost estimation	2	1.5
Health conditions/service type		
General (multiple aspects)	21	15.9
Secondary health utilization	2	1.5
General causes of death	1	0.8
Maternal and Child health/healthcare	12	9.1
Maternal health/healthcare	24	18.2
Child health/healthcare	11	8.3
Vaccine prevented childhood illnesses	10	7.6
Malaria	30	22.7
Malaria & HIV/AIDS	1	0.8
Malaria & other parasitic diseases	1	0.8
HIV and related diseases	8	6.1
Mental health/healthcare	3	2.3
Other diseases	5	3.8
Healthcare workforce and other resources	2	1.5
Data issue of RHIS: missingness		
Described how missing data was managed	33	25.0

No description of how missing data was managed	99	75.0
Data issue of RHIS: outlier		
Described how outlier was detected	19	14.4
No description of how outlier was detected	113	85.6

Table 2. Types of analytic methods applied among studies that analyzed RHIS data.

Data use purpose	Type of disease/service studied	Range of data (unit)	Level of aggregation	Analytic methods	Other information sources included	Reference
<i>Time series analysis</i>						
Epidemiology	Child health, malaria, tooth extraction	15 (year) - 120 (month)	Ward, municipal, district	Time series correlograms; ordinary least-squares regressions adjusted for seasonality and lag; non-linear time series correlation and regressions	GPS coordinates, Climate Hazards Group Infrared Precipitation with Station Data, satellite data, meteorological department data, program data	78,117-119
Program evaluation	General, maternal and child health, maternal health, vaccine prevented childhood illnesses, malaria	5 (year) - 168 (month)	Facility, district, region, nation	Ordinary least squares regression; negative binomial generalized linear model; random effects negative binomial regressions; switching regression methods weighted by propensity scores	Program data, program reports, data from Bureau of Statistics and Ministry of Health, Malaria Indicator Survey, Demographic Health Survey, Health Facility Survey, community survey, satellite data, sentinel site case-investigations/surveillance, abstraction from hospital registries	33,34,37,40,54,55,58,75,76,94,120,121
Impact evaluation (non-program)	General	84 months (month)	Facility	Linear mixed-effect time-series analysis with a segmented regression parameterization	None	73
<i>Interrupted time series analysis</i>						
Program evaluation	General, maternal and child health, maternal health, malaria	53 (month) - 132 (month)	Facility, intervention vs. control groups, district	Generalized least square model with autoregressive structure; generalized least square model with controls, with autoregressive process and moving average process; segmented linear regression	Meteorology Department data, program data, facility survey	38,45,93,95,122,123
Impact evaluation (non-program)	Maternal and child health	44 (month)	District	Segmented linear regression with district fixed effect and clustered standard error at district level	Demographic Health Survey	68
<i>Difference-in-difference analysis</i>						
Program evaluation	General, child health, maternal health	4 (year) - 48 (month)	Facility, district, province	Ordinary least squares regression with and without	Verified data from Performance-Based Financing system	41,42,59,91,92

propensity score matching; Wilcoxon rank-sum test on median difference-in-differences between facilities; descriptive comparison of means

<i>Pre-post comparison analysis</i>						
Program evaluation	Child health, maternal and child health, vaccine prevented childhood illnesses, malaria, HIV or related diseases	2 (year) - 48 (month)	Facility, district	Chi-square test; Pearson correlation; Wilcoxon signed-rank test; paired sample t-test; linear regressions; Poisson regression; negative binomial regression; logistic regression	Bureau of Statistics data, program reports, Meteorological Department data, entomological sentinel surveys, Demographic and Health Survey, UN Interagency Group for Childhood Mortality Estimation (CME Info) database, abstraction from facility registers, community surveys, vital registry, provincial maternal death notification register	35,39,48,57,100,101,124-126
Impact evaluation (non-program)	Child health	26 (month)	District	Pearson chi-square test	District hospital registers, Safe and dignified burials for all deaths database	67
<i>Other longitudinal analysis</i>						
Epidemiology	Maternal health, malaria	12 (year) - 16 (year)	District	Chi-square test; negative binomial regression	Review of hospital death records	96,127
Monitoring and assessment of service provision	HIV or related diseases	3 (year)	District	Descriptive comparison over time	Surveys with health facility managers	97
Program evaluation	General, child health, malaria, malaria and other parasitic diseases	3 (year) - 24 (month)	Facility, district, nation	Poisson regression to explore association between intervention coverage and disease burden; Mann-Whitney U Test to compare prevalence in intervention and non-intervention area; linear regression model; student t-test	Sentinel surveillance data, program reports, national facility and community survey, Bureau of Statistics data, program data	47,52,66,106,128,129
<i>Geostatistical analysis</i>						
Epidemiology	Child health, malaria, malaria and HIV/AIDS, meningococcal meningitis	1 (year) - 520 (week)	District	Cluster analysis; cross-correlations of different spatial scales between time series of cases; Bayesian	Malaria Indicator Survey, Demographic Health Survey, program data	83-85,88,130

hierarchical
Poisson model
and smoothed
model estimates
plotted on
district maps

Monitoring and assessment of service provision	Malaria, maternal health	1 (year) - 57 (month)	Facility, district	Kriging (ordinary kriging, space-time ordinary kriging, local space-time ordinary kriging); Bayesian geostatistical negative binomial model	Service Delivery Indicator Survey	79-82,131
Program evaluation	Malaria	36 (month)	District	Bayesian geostatistical models and Bayesian generalized linear models	Malaria Indicator Survey, malaria control program data, satellite data, Demographic Health Survey, ACTWatch household surveys	87
<i>Other cross-sectional analysis</i>						
Epidemiology	Maternal health	Median of 24 months	Province	Linear regression model	None	105
Monitoring and assessment of service provision	General, child health, maternal health, mental health	1 (year)	Facility, district, municipality, state	Descriptive statistics, Tobit regression model, bivariate and multivariate linear regression models,	Nutrition Service Delivery Assessment, abstraction from Integrated Nutrition Register, structured questionnaire with district health officers, District-level household and facility surveys, National Register of Health Service Providers, data from Institute of Geography and Statistics	107,132-134
Program evaluation	HIV and related diseases	1 (year)	District	Mixed-methods	Register reviews and a series of patient folder (health record) reviews	51

Table 3. Strategies applied in research articles to counter issues of RHIS data.

Type of strategy	Description of strategy
Missing data	
Exclusion	Exclude facility data if a certain threshold was reached (e.g. more than two-thirds of months in a year; more than a sixth of baseline data; facilities with any missing data)
	Restrict analysis to a period with a low level of missing data
	Sensitivity analysis to compare analysis of restricted period and full period
Imputation	Assign missing observations with mean-value for the year
	Assign missing observations with the average of precedent and subsequent data
	Imputation using conditional autoregressive model
	Missing value was replaced as positive (binary form) to prevent exaggeration of the fade-out effect
	Sensitivity analysis of imputation strategies: 1) single imputation using means, trimmed means, and median, 2) Poisson generalized linear modeling, 3) iterative singular value decomposition method
Interpolation	Interpolation using space-time kriging
	Adjust results by dividing each indicator by the percentage of reports submitted
	Adjust the data by calibrating to the total population using proportion reported in a household survey to have occurred in health facilities
Verification	Manual verification of the missing data with register at the health facility
Account in the modeling method	Missing data was assumed missing at random and accounted for in the mixed-effect models using standard maximum likelihood estimation
Identifying extreme values	
Specific threshold	Establishing a lower and upper limit based on proportion of the annual average or feasible value
	Univariate regression on individual facility-level to identify deviation from the mean time trend (e.g. if exceed 8 standard deviations)
Visual	Visual inspection of outliers
Analytic assessment	Jackknifing analysis to assess influence
	Student residual higher than an absolute value of 2 and influence on the estimated coefficients determined by high Cook's distance statistics
Handling of extreme values	
Exclusion	Extreme values were excluded from analyses
Replacing extreme value with average	Extreme values were assigned the average value of the year; with exceptions of low average values
Replacing extreme value with missing	Outliers set to missing
Verification with data source	Any drastic change in monthly data reported electronically were manually verified with register at the health facility. Discrepancies were replaced with data in the register
Discount observation in estimation	Outliers were allocated a dummy coding to discount the observation in the calculation of coefficients
Assess reliability	
Data validation process	Randomly selected 10% of the total sample to check accuracy and reliability of data with reports and registers
	Verify data with another source (e.g. payroll)
	Established routine data validation process by health information and records officer (e.g. monthly data review)

Figures

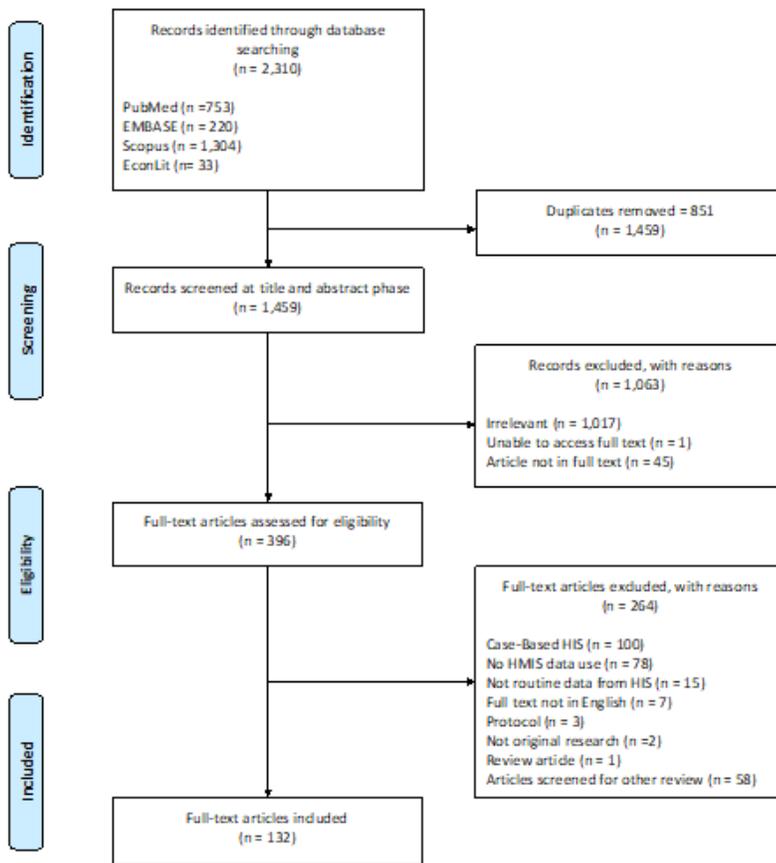


Figure 1

PRISMA flowchart of study identification and screening process of publications use RHIS data.

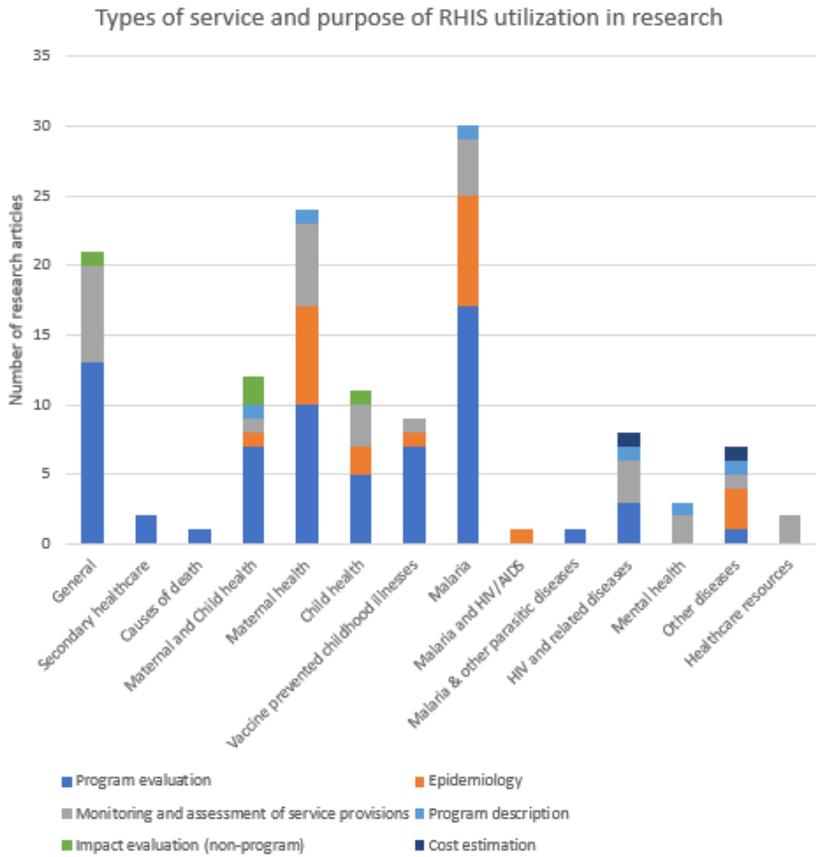


Figure 2

Types of service and research purpose of RHIS data use (n=132).

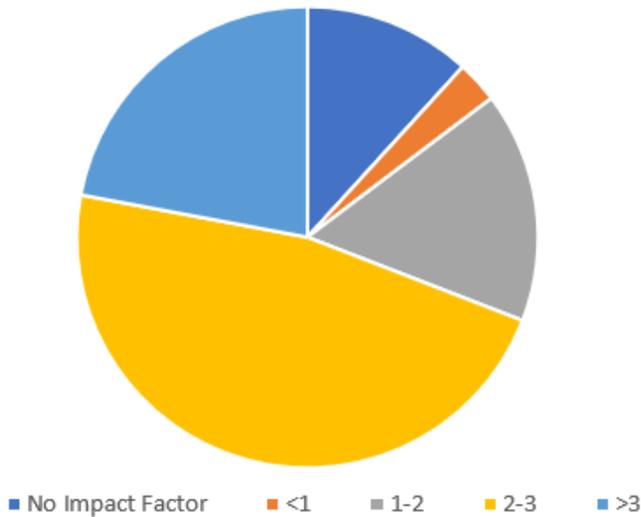


Figure 3

Distribution of impact factor of journals that published research studies that conducted statistical analysis of RHIS data (n=68).

Supplementary Files

This is a list of supplementary files associated with this preprint. Click to download.

- [Appendix1.docx](#)
- [PRISMA2009checklist.doc](#)