

Supplementary appendix

Supplement to the article, “An analysis of clinical knowledge, absenteeism, and available of resources for maternal and child health: A cross-sectional quality of care study in 10 African countries”

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S1: STROBE Statement for cross-sectional, observational studies, with information for this study

	Item No	Recommendation
Title and abstract	1	<p>(a) Indicate the study's design with a commonly used term in the title or the abstract</p> <p><i>In the abstract, we include the following text indicating this is cross-sectional data analysis: "With nationally representative, cross-sectional data from ten countries in Sub-Saharan Africa, collected using clinical vignettes, unannounced visits, and visual inspections, we assess whether health providers are available and have sufficient knowledge and means to diagnose and treat patients suffering from conditions amenable to primary health care."</i></p> <p>(b) Provide in the abstract an informative and balanced summary of what was done and what was found</p> <p><i>In the abstract, we include data on methods and on results.</i></p>
Introduction		
Background/rationale	2	<p>Explain the scientific background and rationale for the investigation being reported</p> <p><i>We include the following text: "Measuring quality at various levels of the health system—especially where the system fails the people in need—remains rare in LMICs. Existing indicators of structural quality, such as availability of medicine, equipment, and qualified health attendants, are not sufficient to adequately capture the quality of care being offered.¹ The lack of quality measurement can help to explain both why there is limited research on how to effectively transform low-quality health systems into high-quality ones, and why there are few such successful reforms in LMICs. Actionable and credible data on quality, if properly disseminated, can play an important role in holding country health systems accountable for effective and safe care and can form the basis for quality improvement."</i></p>
Objectives	3	<p>State specific objectives, including any prespecified hypotheses</p> <p><i>We include the following text: "We use the data to measure dimensions we would argue are essential for providing quality care: What do providers know? Are providers available at health facilities? And what means do they have at their disposal?"</i></p>
Methods		
Study design	4	<p>Present key elements of study design early in the paper</p> <p><i>We lay out the design: "Our paper reports on the first large scale assessments of clinical knowledge, health provider absenteeism, and means to provide care in Sub-Saharan Africa. We use data collected through clinical vignettes, unannounced visits, and visual inspections of over 20,000 providers in over 8,000 facilities from ten Sub-Saharan African countries, which together represent over 40 percent of the region's total population."</i></p>

Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection <i>See the relevant text: “We draw upon data from the Service Delivery Indicators (SDI) program—an ongoing program with the aim of collecting informative and standardized measures of what providers know, what they do, and what they have to work with...To date, the SDI program has collected data from ten countries: Kenya (2018), Madagascar (2016), Mozambique (2014), Nigeria (2013), Niger (2017), Senegal (2010), Sierra Leone (2018), Tanzania (2016), Togo (2014), and Uganda (2013), and has since expanded to countries outside the Africa region. While every context is different, these ten countries serve as a reasonable proxy for much of Sub-Saharan Africa, with an average income per capita, life expectancy, and infant mortality similar to those of the continent as a whole (Appendix Table S2).”</i>
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of participants <i>See the relevant text: “All country datasets were representative at the national level except for Nigeria which, mainly owing to security concerns, covered 12 out of 37 states and was representative at the state level. The surveys used a multistage cluster-sampling design to ensure data were representative of rural and urban areas, private and public facilities, and of each facility type (e.g., front-line clinics versus hospitals).”</i>
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable <i>See the relevant text: “There are three essential conditions for quality care. First, providers have sufficient knowledge to diagnose and treat patients suffering from conditions that are amenable to care. Second, they are available to provide care when patients seek service. Third, providers have the means (e.g., equipment and drugs) to diagnose and treat patients.”</i> <i>“Combining data from the clinical vignettes (to measure knowledge) with data from the unannounced visit (to measure the extent to which providers are present in the facility at regular opening hours), and visual inspections (to quantify the availability of drugs and equipment), we estimate the probability that a facility is ready to provide care that meets minimum quality standards. A facility is defined as being ready to provide care that meets minimum quality standards for a given condition if there is at least one provider present in the clinic at a given day that can correctly diagnose and treat that condition, and the facility has a minimum set of drugs to treat it.”</i>
Data sources/ measurement	8*	For each variable of interest, give sources of data and details of methods of assessment (measurement). Describe comparability of assessment methods if there is more than one group <i>See the relevant text: “Diagnostic and treatment accuracy (knowledge) were measured using clinical vignettes focused on seven common conditions associated with the main causes of child and maternal mortality in Sub-Saharan Africa: malaria, tuberculosis, diarrhea, pneumonia, diabetes, neonatal asphyxia and post-partum hemorrhage. All seven of these are major global health challenges, with three of them – diarrhea,</i>

neonatal asphyxia, and malaria – together causing more than 180,000,000 disability-adjusted life years (Appendix Table S3). Whether providers are available to provide care at regular opening hours was assessed by verifying their presence through unannounced visits at the health facilities. Availability of drugs and functionality of equipment was assessed by visual inspections of storage facilities and consulting rooms at the health facilities.

Clinical vignettes have been used extensively in the medical literature to assess clinical knowledge of health workers. Data from vignettes studies are directly policy relevant as they are closely linked to the quality of medical training and quality improvement programs. Validation studies show that clinical vignettes, for relatively simple scenarios such as those assessed as part of the SDI program, perform similarly to standardized patients methods. Although a health worker's clinical vignette performance is not a measure of quality of care per se, it represents the highest level of quality care which s/he can deliver given his/her current knowledge, assuming that there are no constraints in drugs, equipment and motivations. (More details on the clinical vignettes are provided in Section S4 in the appendix.)

The clinical vignettes were adjusted to country-specific diagnostic and treatment protocols. All health providers assessed were asked for informed consent to participate. The enumerators presented the provider with the patient's symptoms and prompted the provider to take the patient's history, suggest physical examination and diagnostic tests that in principle could be conducted on site, make a diagnosis, and prescribe a course of treatment and follow-up management plan. For this study, we identified the lowest common denominators across the country-specific protocols to define comparable criteria across countries for diagnostic and treatment accuracy for each of the conditions. This methodological choice implies that knowledge is assessed at a lower standard than more stringent World Health Organization (WHO) protocols. Based on the responses to each vignette, we generated two variables: diagnostic accuracy and treatment accuracy, which are the percentage of respondents who provided an accurate diagnosis and – separately – an accurate treatment for a given condition. Both outcomes should be viewed as measuring the minimum required knowledge to prescribe correct diagnostic and treatment services. (See Table S7 for information on the coding of the two knowledge scores.) These outcomes do not represent the ideal level of knowledge; for example, they do not account for co-morbidities.

*To quantify whether providers are **available to provide medical service when patients seek care**, we followed a standard procedure in which enumerators visited each facility twice. During the first visit, information on all staff working at the facility, by professional cadre, was collected. (See Section S5 in the appendix for details on the selection of staff, **drugs, and equipment** for surveys.) A few days after the first visit, enumerators made a second unannounced visit to the facility and assessed the presence of a randomly pre-selected list of providers (or all providers, if less than 10 providers were available). Only staff on duty that day were included in the absenteeism measure. Country-level averages for each assessment were calculated using country-specific sampling weights.*

Combining data from the clinical vignettes (to measure knowledge) with data from the unannounced visit (to measure the extent to which providers are present in the facility at regular opening hours), and visual inspections (to quantify the availability of drugs and equipment), we estimate the probability that a facility is ready to provide care that meets minimum quality standards.”

Bias	9	Describe any efforts to address potential sources of bias
		<i>In order to minimize bias, the study includes nationally representative samples for all but one country in the sample (Nigeria). In Nigeria, the sample excluded some states due to security concerns but was representative at the state level for 12 of 37 states. In all countries, the team used a “multistage cluster-sampling design to ensure data were representative of rural and urban areas, private and public facilities, and of each facility type (e.g., front-line clinics versus hospitals).”</i>
Study size	10	Explain how the study size was arrived at
		<i>See the relevant text: “In total, the sample includes information from 89,826 vignettes from 13,754 providers and absenteeism observations from 22,747 providers across 8,061 facilities in 10 countries. In each country, the sample size was selected in order to provide representative data for the country and for sub-groups (e.g., rural versus urban areas).”</i>
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why
		<i>See the relevant text: “We present averages and 95% confidence intervals by first averaging all observations across facilities (or health workers) in each country (using country-specific facility weights to calculate a representative country average), and then averaging over the country means to calculate an overall average for the whole sample. (This is a suitable approach if we treat the country as the core unit of observation; i.e., we want to know how countries compare and how they perform as a group.)”</i>
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding
		<i>As above. In this analysis, we are comparing averages: “This procedure is equivalent to calculating a weighted average and its standard error across all observations in the sample, where the weight on observation i in country j is $\omega_{ij} = \prod_{k \neq j} F_k \times f_{ij}$, where f_{ij} denotes the (facility) weight of observation i in country j and $F_k = \sum_r f_{rk}$ is the sum of the facility weights in country k.”</i>
		(b) Describe any methods used to examine subgroups and interactions
		<i>See the relevant text: “Countries are divided according to performance. Figure S10 divides the countries into two groups based on performance. In the higher performance group, 49 percent of the health workers diagnose all or almost all of the conditions, compared to 11 percent in the lower performing group.”</i>
		<i>“Diagnosis and prescription of correct treatment is also examined according to the type of disease. For some conditions, the key constraint</i>

appears to be at the diagnostic stage, as with diarrhea, where 59% of the providers diagnosed correctly but 91% of those who did so prescribed the correct treatment, and neonatal asphyxia, where 59% of the providers diagnosed correctly but 93% of those who did so prescribed the correct treatment. For other conditions, the key constraint instead appears to be prescribing the correct treatment, as with tuberculosis, where 81% of the providers correctly diagnosed the condition but only 37% of these providers prescribed the correct treatment, and post-partum hemorrhage, where 71% of the providers correctly diagnosed the condition but only 45% of them prescribed the correct treatment.”

(c) Explain how missing data were addressed

The country-specific facility weights recreate a representative country average in the face of any missing facilities.

(d) If applicable, describe analytical methods taking account of sampling strategy

See the relevant text: We use “country-specific facility weights to calculate a representative country average.”

(e) Describe any sensitivity analyses

In addition to the average analysis, we provide country-by-country and condition-by-condition analysis to demonstrate that our results are not driven by a single country or condition.

Results

Participants	13*	<p>(a) Report numbers of individuals at each stage of study—eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed</p> <p><i>See the relevant text: “The analysis in the study is at the level of the provider. We used the full sample of providers available from the surveys. In total, the sample includes information from 89,826 vignettes from 13,754 providers and absenteeism observations from 22,747 providers across 8,061 facilities in 10 countries.”</i></p> <hr/> <p>(b) Give reasons for non-participation at each stage</p> <p><i>The surveys were carried out in partnership with national governments and so facilities participated.</i></p> <hr/> <p>(c) Consider use of a flow diagram</p> <p><i>Not applicable.</i></p>
Descriptive data	14*	<p>(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders</p> <p><i>Because the paper covers samples in ten different countries, and descriptive data on those samples are provided in country-specific studies, we provide overarching characteristics of the country samples (e.g., in Table S1) as compared to Africa as a whole.</i></p>

(b) Indicate number of participants with missing data for each variable of interest

Not applicable, given that facilities participated in the surveys.

Outcome data	15*	Report numbers of outcome events or summary measures <i>As discussed in the text, measures include (1) diagnostic accuracy, (2) rate at which the correct prescription is given to the patient, (3) rate of provider absence, (4) availability of drugs for treatment, (5) probability of care readiness, namely (i) whether there is at least one provider working in the clinic that can correctly diagnose and prescribe treatment for the condition(s); (ii) health worker attendance; and (iii) whether or not the facility has a minimum set of drugs to treat the condition(s).</i>
Main results	16	<p>(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and their precision (eg, 95% confidence interval). Make clear which confounders were adjusted for and why they were included</p> <p><i>The paper includes unadjusted estimates and confidence intervals: “(1) Across all conditions and countries, health care providers correctly diagnose 64% (95% CI 0.62-0.65) of the cases. For diabetes, 35% (95% CI 0.33-0.37) of the health workers provide a correct diagnosis (Figure 1A; Table S7). For the three main killer diseases for children in Sub-Saharan Africa, accounting for approximately one-third of all deaths among children under age 5 in the countries surveyed, average diagnostic accuracy is 57% (95% CI 0.54-0.59) for pneumonia, 59% (95% CI 0.56-0.61) for diarrhea and 85% (95% CI 0.82-0.89) for malaria.</i></p> <p><i>(2) Diagnostic knowledge is associated with correct treatment, but a correct diagnosis is not a guarantee for prescribing correct treatment. The providers prescribed the correct (minimum) treatment following a correct diagnosis in 72% of the cases. Across all conditions and countries, health care providers were able both to correctly diagnose and treat 45% (95% CI 0.43-0.46) of the cases. For diarrhea, malaria, and neonatal asphyxia a majority of the providers managed to do so (54% (95% CI 0.43-0.46); 70% (95% CI 0.66-0.73); and 55% respectively). For the other four conditions, the majority of providers either fail to diagnose the condition correctly or fail to provide the correct treatment (Figure 1B; Table 2).</i></p> <p><i>(3) The rate of provider absence is, on average, 30% (95% CI 0.29-0.32), with large variations across countries (see Table 2 and Table 4). For example, doctors in Togo and nurses in Uganda have absence rates of 50% (95% CI 0.26-0.74) and 47% (95% CI 0.43-0.50), respectively, while the average absence rate of doctors in Tanzania is 16% (95% CI 0.12-0.21).</i></p> <p><i>(4) On average, 42% (95% CI 0.40-0.45) of the facilities were stocked with all four (or all three for Kenya) types of drugs and 70% (95% CI 0.69-0.73) of the facilities had a minimum set of functioning medical equipment (see Table S13). ORS is available in 84% (95% CI 0.82-0.87) of the facilities, ACT in 77% (95% CI 0.75-0.80) of the facilities, antibiotics in 69% (95% CI 0.66-0.72) of the facilities and oxytocin in 62% (95% CI 0.60-0.65) of the facilities (see Table 2).</i></p> <p><i>(5) On average, only 14% (95% CI 0.12-0.15) of facilities are ready to provide selected child, neonatal, and maternity care that meets minimum</i></p>

		<p><i>quality standards, with (i) at least one provider available and able to correctly diagnose and treat diarrhea, pneumonia, post-partum hemorrhage and neonatal asphyxia, and (ii) minimum drugs as required (column (1), Table 5)."</i></p> <hr/> <p>(b) Report category boundaries when continuous variables were categorized</p> <p><i>Not applicable. The collected data includes only discrete variables. Any continuous variables are generated and are reported as continuous.</i></p> <hr/> <p>(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period</p> <p><i>Not relevant for this study.</i></p>
Other analyses	17	<p>Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses</p> <p><i>Countries are divided according to performance. Figure S11 divides the countries into two groups based on performance. In the higher performance group, 49 percent of the health workers diagnose all or almost all of the conditions, compared to 11 percent in the lower performing group.</i></p> <p><i>Diagnosis and prescription of correct treatment is also examined according to the type of disease. For some conditions, the key constraint appears to be at the diagnostic stage, as with diarrhea, where 59% of the providers diagnosed correctly but 91% of those who did so prescribed the correct treatment, and neonatal asphyxia, where 59% of the providers diagnosed correctly but 93% of those who did so prescribed the correct treatment. For other conditions, the key constraint instead appears to be prescribing the correct treatment, as with tuberculosis, where 81% of the providers correctly diagnosed the condition but only 37% of these providers prescribed the correct treatment, and post-partum hemorrhage, where 71% of the providers correctly diagnosed the condition but only 45% of them prescribed the correct treatment.</i></p>
Discussion		
Key results	18	<p>Summarise key results with reference to study objectives</p> <p><i>See the relevant text: "This study offers a single metric that quantifies the relative contribution of staff knowledge, staff availability, and other facility inputs to the probability of care readiness which meets minimum quality standards."</i></p> <p><i>"Our work shows severe gaps in care readiness in various countries in Sub-Saharan Africa, and health provider knowledge is a particularly severe constraint on the readiness of care across countries. In many facilities, providers are unable to diagnose accurately and propose appropriate treatment for conditions which account for major loss of life across the continent."</i></p>
Limitations	19	<p>Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss both direction and magnitude of any potential bias</p>

See the relevant text: “Not all misdiagnoses and mistreatments impose equal costs on patients and on health care systems. Some mistreatments – e.g., prescription of antibiotics for viral diarrhea – impose longer term costs on the patient population with greater resistance to antibiotics but limited short-term adverse impacts to the patient. In this analysis, we focus on the country-specific protocols, but one could imagine an analysis that weighs the direct and indirect cost to patients of different errors in diagnosis and treatment.

Furthermore, there are times that deviations from diagnostic and treatment protocols may be guided by information rather than ignorance. In the case of prescribing antibiotics for diarrhea, recent evidence suggests that antibiotics promote growth among young children, so medical providers could be incorporating that information into their treatment. That said, while a medical provider may overprescribe antibiotics for that reason in practice, there is less reason to expect that she would do so in a vignette as a response to a specific diarrheal condition. In contexts where providers seem to have limited skills, ensuring that guidelines adhere to the best and latest knowledge and practice, and encouraging providers to follow those guidelines, are likely to enhance the quality of care.”

Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence <i>See the relevant text: “Our findings highlight the need to adopt competency-based pre-service and in-service training for health care providers. As poor clinical knowledge is the key constraining factor at the facility level, improving drug availability or reducing health workers’ absenteeism would only modestly increase the average care readiness that meets minimum quality standards.”</i>
Generalisability	21	Discuss the generalisability (external validity) of the study results <i>See the relevant text: “This study uses the lowest common denominator across the country-specific protocols to define comparable care readiness criteria across countries. Moreover, there is the well-known gap between what health workers can do and what they actually do (the “can-do” gap).²² For these reasons, we expect that in most surveyed countries the probability of providing minimum quality care according to country-specific guidelines is lower than that reported in this study.”</i>
Other information		
Funding	22	Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based <i>See the relevant text: “The funding is from the World Bank Service Delivery Indicators Trust Fund (funded in large part by the Hewlett Foundation). The funding source did not have any role in the design, data collection, data analysis, data interpretation, or writing of the report.”</i>

*Give information separately for exposed and unexposed groups.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at <http://www.plosmedicine.org/>, Annals of Internal Medicine at <http://www.annals.org/>, and Epidemiology at <http://www.epidem.com/>). Information on the STROBE Initiative is available at www.strobe-statement.org.

Table S2: How the Study Countries Compare with Sub-Saharan Africa as a Whole

	GDP per capita, PPP (current international \$)	Life expectancy at birth (years)	Maternal mortality per 100,000 live births (modeled estimate)	Infant mortality rate per 1,000 live births	Incidence of malaria per 1,000 population at risk
Kenya	3,468	66	342	31	71
Madagascar	1,891	66	335	38	91
Mozambique	1,460	59	289	54	338
Niger	1,063	62	509	48	359
Nigeria	5,991	54	917	76	281
Senegal	3,783	67	315	32	65
Sierra Leone	1,602	54	1,120	79	380
Tanzania	3,227	64	524	38	113
Togo	1,774	60	396	47	371
Uganda	2,038	63	375	34	201
Average of ten study countries	3,999	59	632	55	221
Average of Sub- Saharan Africa	3,988	61	534	53	210

Source: World Development Indicators 2019.¹

Table S3. Disability adjusted life years lost to the diseases measured in the Service Delivery Indicators

	DALYs (Disability- Adjusted Life Years, thousands)	YLDs (Years Lived with Disability, thousands)	Prevalence (thousands)	Incidence (thousands)
Diarrhoeal diseases	81,000	10,465	93,473	6,292,937
Diabetes	67,900	38,575	475,996	22,936
Neonatal encephalopathy due to birth asphyxia and trauma <i>(closest in our study: neonatal asphyxia)</i>	56,500	9,704	538,238	2,753
Tuberculosis	45,000	3,120	1,929,209	8,966
Malaria	45,000	1,468	136,085	208,768
Pneumococcal meningitis <i>(closest in our study: pneumonia)</i>	3,080	325	3,557	445
Maternal haemorrhage	2,230	61	1,660	6,988

Source: Global Burden of Disease 2017.^{2,3}

S4. Vignettes: Protocol, administration, reason for use, and validation

A vignette is a hypothetical case in which the interviewer acts as a “patient” and provides a very brief description of symptoms. The provider is informed that the “patient” will comply with the provider’s instructions, medications, and tests. The provider is then invited to proceed exactly as he or she would under normal circumstances, asking questions about the history of the illness and performing necessary examinations and tests. The “patient” provides standardized predetermined answers to the questions and examination/test procedures. The questions, examinations, and tests that a provider uses are compared to protocol checklist. Data are recorded on each of the items on the checklist the provider provided; the diagnosis given; and the treatment suggested. Four of the seven vignettes used in this study were originally developed for use in Tanzania, with the items drawn from the national protocol.⁴ The list was subsequently expanded to include items required by the guidelines for the Integrated Management of Childhood Illnesses.^{5,6} Vignettes were administered to a random sample of doctors, medical assistants, nurses, and nurse-midwives (i.e., any staff that provide outpatient or maternity services) who were present in the facility on the day of the first, announced visit. Absenteeism on that first visit, announced in advance, was 17.2 percent (for nurses and midwives) and 25.4 percent (for doctors).

The assessments were carried out by a team of two enumerators who were trained in the administration of vignettes. One acted as the patient and the other recorded outcomes. Normally, the recorder should have medical training: for example, in Togo the enumerators were medical students and in Niger, enumerators were medical workers with at least a university degree.

We use data from seven vignettes capturing conditions associated with the main causes of child and maternal mortality in Sub-Saharan Africa: malaria, tuberculosis, diarrhea, pneumonia, diabetes, neonatal asphyxia and post-partum hemorrhage. All seven of these conditions are major global health challenges, with three of them only – diarrhea, neonatal asphyxia, and malaria – together causing an estimated 180,000 thousands disability-adjusted life years (see Table S3).

Table S5 below shows the starting scenario for the vignettes. Beyond the starting scenario, the vignettes proceed with additional information depending on the questions asked by the health provider.

Table S5. Initial scenarios for vignettes

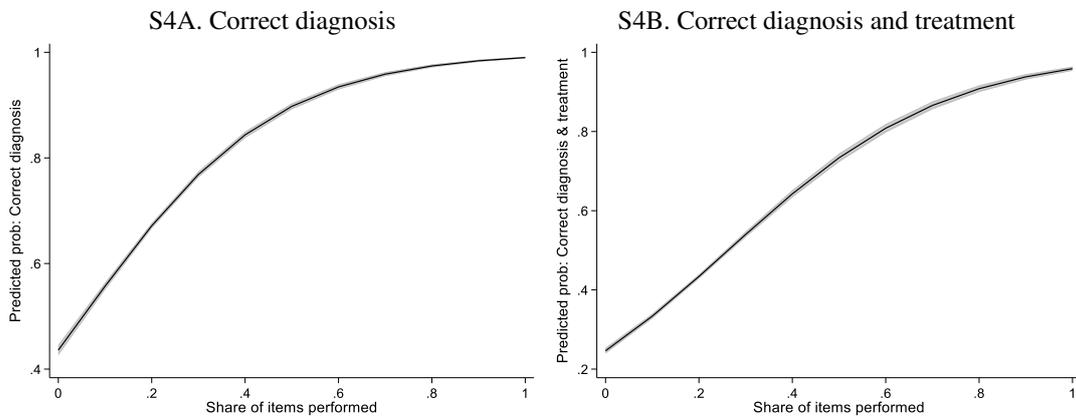
Vignette	Description
Acute diarrhea with severe dehydration	An adult presents a 1 year child with diarrhoea.
Pneumonia	An adult present a 5-year old child with cough.
Diabetes Mellitus	An adult presents at the facility feeling weak and without energy despite feeling hungry often and eating frequently. The man is 48yrs old and works as a clerk.
Pulmonary Tuberculosis	A pregnant woman presents at the facility after being told in the community that she should go for antenatal care.
Malaria + anemia	An adult presents a child with fever for some time who has gotten worse.
Post partum hemorrhage	A young woman presents at the facility with vaginal bleeding 24 hours after delivery in a health facility
Neonatal Asphyxia	A mother gives birth. The newborn is not crying. The newborn fails to establish regular breathing and appears pale and slightly blue.

There are a number of ways of scoring performance in patient case simulations, or vignettes. In this study we focus on diagnostic and treatment accuracy. We identified the lowest common denominators across the country-specific protocols to define comparable criteria across countries. This methodological choice implies that knowledge is assessed at a lower standard than more stringent WHO protocols.

The medical vignettes are used to measure clinic knowledge. That is, a health worker's clinical vignette performance is not a measure of quality of care per se. It does, however, represent a good measure of the quality care which the provider can deliver, assuming he/she has access to relevant medical infrastructure. There are alternative methods to test for clinical knowledge and performance, including direct observation of real patient consultations and the use of standardized patients. Observation of real patient consultations, like vignettes, may be subject to observer effects (also referred to as Hawthorne effects). A study in the U.S. finds that the quality score derived using data from vignettes was similar to the quality score with standardized patients (i.e., trained actors who make unannounced visits to the clinic), but a study in India shows large gaps between vignettes and actual practice with standardized patients.^{7,8} A study using data collected from 104 medical providers in northern Tanzania find that item by item, vignettes and direct observation produced identical results 63% of the time.⁸ Aggregate scores derived from vignettes tend to be higher than scores from direct clinical observation, suggesting that knowledge measured using vignettes should be interpreted as an upper-bound of what providers could do; i.e., the vignettes measure what the medical provider knows, but not necessarily whether the provider applies this knowledge in practice.^{9,10}

The main advantages of using medical vignettes compared to direct observation of real patient consultations (or re-examination of patients outside of the clinic by well-qualified personnel), are that they are far less time-consuming and they increase comparability by avoiding issues related to differences in case mix. Moreover they provide a direct measure of providers' clinical knowledge, rather than a combined measure of knowledge and effort. Data from medical vignettes is also directly policy relevant as the scores are closely linked to the quality of medical training.

Validation of vignettes typically involves comparing scores from vignettes with scores from direct observations.^{9,10} This is less relevant for this study as we focus on provider knowledge, realizing that in practice (some) providers may not always fully utilize their knowledge when diagnosing and treating patients. Still, the vignettes utilized in this study have been (indirectly) validated as a testing method for provider knowledge. One source of validation is the fact that providers with a high share of relevant items performed are much more likely to suggest the correct diagnosis and treatment.⁹ That is, better knowledge of which history taking question to ask and examinations to perform is highly correlated with both diagnostic and treatment accuracy. The two figures below, repeat that exercise using the full sample of vignettes and share of items performed (89,824 observations). We estimate a logit model with the binary response variable correct diagnosis [correct diagnosis and treatment] and with share of items performed as a continuous predictor and country and condition categorical predictors as control variables. The figures plot the adjusted probability of reaching the right diagnosis [right diagnosis and treatment]. As the share of items performed increases, the estimated probability of reaching a correct diagnosis and correct diagnosis followed by a correctly described treatment increase. The adjusted probability of correct diagnosis and treatment is estimated to be 33% for a provider performing 10% of the items as compared to 94% for a provider performing 90% of the items. For a provider who asks no history taking questions nor perform any examinations and tests, the adjusted probability of correct diagnosis and treatment is below 25%.



Note: Adjusted probability of reaching the right diagnosis (S4A) and right diagnosis and correctly described treatment (S4B) and 95% CI (shaded area). See text for details.

S6. Cross-country variation in treatment protocols for the vignettes

In each country, the Service Delivery Indicators team asked the government for the official treatment guidelines for each treatment, then validated those along with the vignette in a workshop with a government-identified in-country doctor. The result is that there is cross-country variation in treatment guidelines, but the variation is limited. To demonstrate, we provide the treatment guidelines listed in our data, resulting from the process above for post-partum hemorrhage, country by country. While there are minor variations across some countries, the general treatment protocols are very similar across countries.

Country	Treatment protocol
Kenya	Determine the cause, uterine massage, oxytocin, prostaglandins, surgery if other measures fail to stop bleeding, foley catheter, take blood grouping for cross matching, referral, advice on other signs
Madagascar	Determine the cause, uterine massage, IV line, oxytocin, IV infusion of plasmion, blood transfusion, prostaglandins, misoprostol, uterine revision, surgery if other measures fail to stop bleeding, foley catheter, referral
Mozambique	Uterine massage, Oxytocin, IV line, IV infusion of plasmion, blood transfusion, prostaglandins, misoprostol, uterine revision, surgery if other measures fail to stop bleeding, foley catheter, referral
Niger	Determine the cause, uterine massage, uterine revision, IV line, oxytocin, ergometrine, IV infusion of plasmion, blood transfusion, progstaglandins, misoprostol, surgery if other measures fail to stop bleeding, foley catheter, referral
Nigeria	IV line, take blood grouping for cross matching, foley catheter, uterine massage, oxytocin, prostaglandins, surgery if other measures fail to stop bleeding, referral
Sierra Leone	Determine the cause, IV line, oxytocin, ergometrine, IV infusion of plasmion, blood transfusion, prostaglandins, misoprostol, uterine revision, surgery if other measures fail to stop bleeding, foley catheter, referral
Tanzania	Determine the cause, uterine massage, IV line, oxytocin, IV infusion of plasmion, blood transfusion, prostaglandins, misoprostol, uterine revision, surgery if other measures fail to stop bleeding, foley catheter, referral
Togo	Determine the cause, IV line, uterine massage, IV infusion of plasmion, blood transfusion, take blood grouping for cross matching, foley catheter, uterine revision, prostaglandins, surgery if other measures fail to stop bleeding, referral
Uganda	Determine the cause, IV line, uterine massage, take blood grouping for cross matching, foley catheter, oxytocin, prostaglandins when available/misoprostol, surgery, referral
Minimum knowledge	Uterine massage and oxytocin

Table S7. Minimum knowledge

The minimum knowledge per vignette required for correct diagnose and correct treatment

<i>Vignette</i>	<i>Correct diagnosis</i>	<i>Correct treatment</i>
Pneumonia	Pneumonia	Antibiotics
Diarrhea with severe dehydration	Diarrhea with (severe) dehydration, or acute diarrhea, or diarrhea	ORS or rehydration therapy
Tuberculosis	Tuberculosis	Combination of anti-tuberculosis drugs
Malaria and anaemia	Malaria, or malaria and anemia, or severe malaria	Artemisinin-based combination therapy (ACT)
Neonatal asphyxia	Neonatal asphyxia or birth asphyxia	First aid assistance including establishing open airway and if need stimulate ventilation
Postpartum hemorrhage	Postpartum hemorrhage	Uterine massage and oxytocin
Diabetes	Diabetes	Refer patient to a specialist

S8. Selection of staff, drugs, and equipment for surveying

During the first survey visit, information on all staff working at the facility, by professional cadre, and whether or not they were conducting patient consultations, was collected. Then at most ten health workers were randomly sampled from the list of health workers present for the vignettes. Thus, the proportion of doctors, nurses, and midwives was not stratified and would roughly reflect the proportion of those professionals present at the clinic.

The World Health Organization Service Availability and Readiness Assessment (SARA) formed the basis for the list of drugs and equipment, intended to capture a list of essential or minimal items that should be available at every health facility.¹¹ Within each country, the SDI team collaborated with the government to adjust items on the list according to local guidelines and priorities. Availability of drugs and equipment was assessed during the first visit by visual inspections of storage facilities and consulting rooms at the health facilities.

A few days after the first visit, enumerators made a second unannounced visit to the facility and assessed the presence of a randomly pre-selected list of providers (or all providers, if less than 10 providers were available). Only staff on duty that day were included in the absenteeism measure. Staff who were away from the facility due to outreach or fieldwork were counted as present.

S9. Estimation of the probability of care readiness

The probability of care readiness that meets minimum quality standards for condition c in facility i (PCR_{ci}) was calculated as follows:

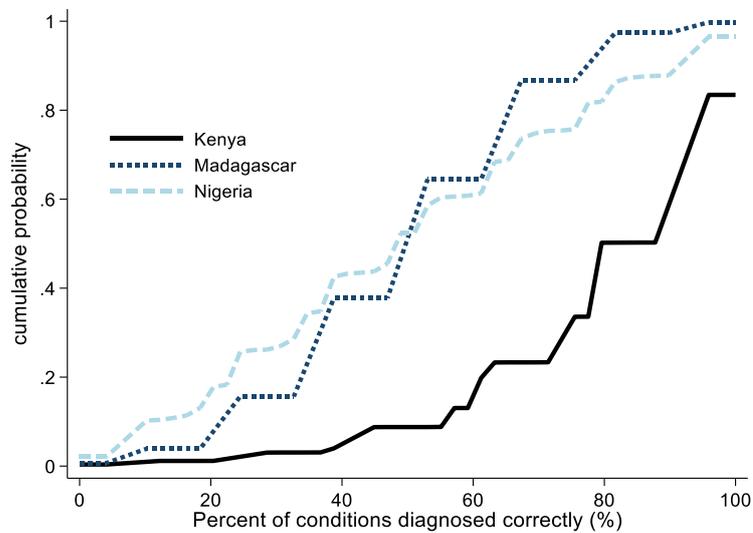
$$(A1) \quad PCR_{ci}(\theta_{ci}, p_{ij}, n_{cij}) = \theta_{ci} \left(1 - \prod_{j=1}^J (1 - p_{ij})^{n_{cij}} \right)$$

where j denotes cadre, or type (doctor, clinical officer, nurse, and community health workers), J is the number of types, n_{cij} denotes the number of providers in facility i of type j that are estimated to be able to describe the correct treatment for condition c , and p_{ij} the estimated probability that a type j provider will be available to provide services a given day.

Data from announced visits of a random sample of providers (up to 10 per facility) were used to estimate the average rate of presence of type j in facility i ; i.e. p_{ij} . In case no type j provider was assessed, we replace the mean presence rate for that type with the mean presence rate in the facility (across all other types).

The number of providers in facility i of type j that are able to describe the correct treatment for the three conditions (n_{ij}^c) is the sum of type j (in facility i) assessed providers that accurately describe the correct treatment, denoted n_{ij}^c , and the estimated number of non-assessed providers of type ij , denoted n_{ij}^e . n_{ij}^e is calculated as $(n_{ij}^c/n_{ij}^a) \times n_{ij}^{na}$, where n_{ij}^a is number of assessed type ij providers and n_{ij}^{na} is the number of non-assessed providers of type ij . In case no type j provider was assessed in facility i , n_{ij}^e is calculated $E(\text{accuracy}) \times n_{ij}^{na}$, where $E(\text{accuracy})$ is the country-specific treatment accuracy (share of providers of type j that correctly describe the correct treatment for the condition(s)).

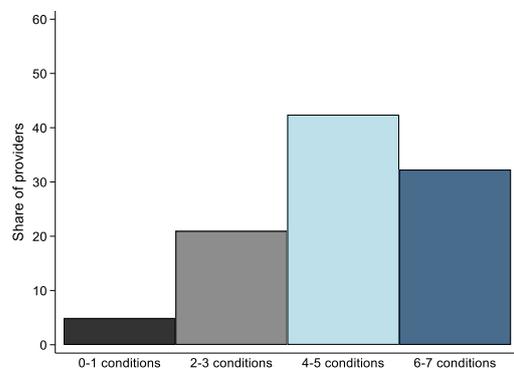
We report individual results for diarrhea, pneumonia, malaria, neonatal asphyxia, and postpartum hemorrhage in the main text and individual results for tuberculosis and diabetes in Table 3. Child care includes two conditions (diarrhea and pneumonia). Neonatal & maternity care includes two conditions (post-partum hemorrhage and neonatal asphyxia). Child, neonatal & maternity care includes four conditions (diarrhea, pneumonia, post-partum hemorrhage, and neonatal asphyxia). In Table 5 we add malaria (which leads to a large drop in sample size since the malaria vignette was not used in the Kenya survey).

Figure S10. Variation across the quality spectrum

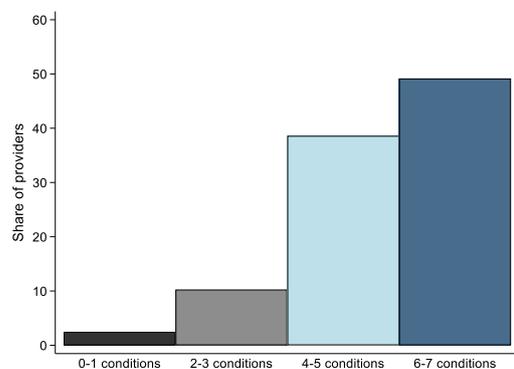
Notes: This figure shows the cumulative distributions of conditions diagnosed correctly for Kenya, Madagascar, and Nigeria. In Kenya, the curve is almost vertical at low scores, implying there are few providers that only manage to diagnose few conditions. In Nigeria and Madagascar, roughly 40 percent of the providers manage to diagnose 40 percent or less of the conditions, but Nigeria also has relative more providers that manage to diagnose most conditions; i.e., the slope of the curve is less steep in the case of Nigeria compared to Madagascar.

Figure S11. Clustering of Diagnostic and Treatment Accuracy

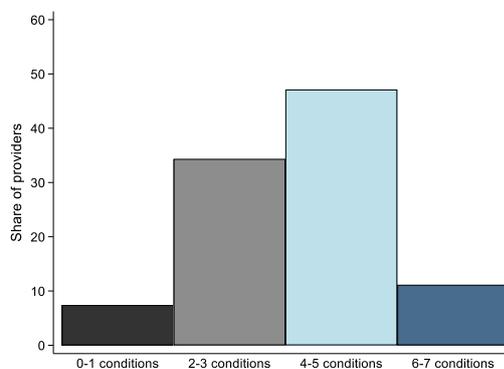
A. All countries



B. Higher performing countries



C. Lower performing countries



Notes: The figures show the share (%) of providers who correctly diagnose at each number of vignettes. Panel A illustrates the results for all countries, while Panel B illustrates the results for Kenya, Mozambique, Sierra Leone, Tanzania, and Uganda, and Panel C the results for Madagascar, Niger, Nigeria, and Togo. All estimates are (unweighted) mean outcomes across countries, with the country means calculated using country-specific sampling weights. All providers were tested for 7 conditions using vignettes in Madagascar, Mozambique, Nigeria, Niger, Sierra Leone, Tanzania, Togo, and Uganda, and for 6 conditions (not malaria) in Kenya.

Table S12. Over prescription of antibiotics

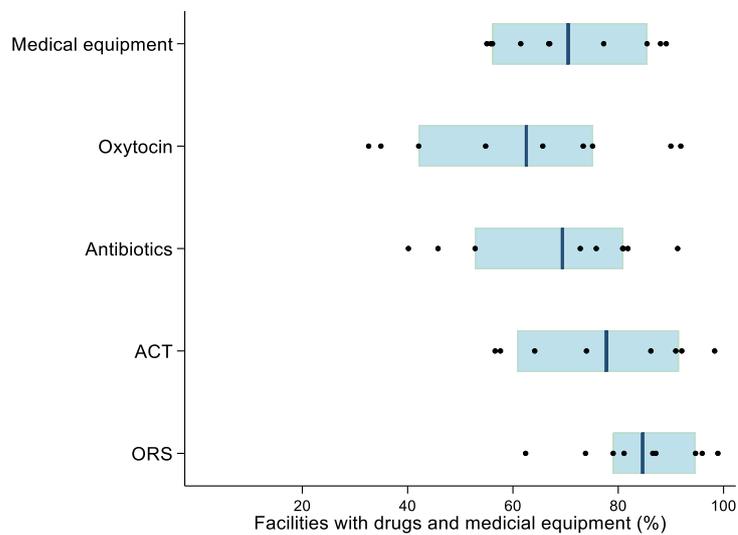
Prescribe antibiotics for diarrhea	(1) Independent of diagnosis	(2) Incorrect diagnosis	(3) Correct diagnosis
Kenya	0.40 (.38-.41) [4505]	0.49 (.45-.52) [1097]	0.37 (.35-.39) [3408]
Madagascar	0.35 (.29-.42) [642]	0.35 (.27-.43) [513]	0.36 (.22-.50) [129]
Mozambique	0.41 (.37-.46) [725]	0.44 (.36-.52) [189]	0.40 (.36-.46) [536]
Niger	0.17 (.12-.22) [519]	0.18 (.04-.31) [57]	0.17 (.11-.22) [462]
Nigeria	0.48 (.46-.50) [4549]	0.44 (.41-.46) [3313]	0.60 (.57-.64) [1236]
Senegal	0.20 (.10-.30) [152]	0.27 (.11-.42) [64]	0.16 (.05-.27) [88]
Sierra Leone	0.32 (.28-.35) [826]	0.47 (.30-.63) [44]	0.31 (.27-.35) [782]
Tanzania	0.30 (.24-.33) [542]	0.28 (.09-.48) [67]	0.30 (.24-.36) [475]
Togo	0.42 (.29-.54) [302]	0.47 (.32-.61) [233]	0.18 (.05-.31) [69]
Uganda	0.29 (.24-.33) [709]	0.27 (.22-.33) [422]	0.31 (.27-.35) [287]
All	0.33 (.31-.36) 13,374	0.36 (.32-.40) 5,915	0.32 (.29-.35) 7,458

Notes: The table reports the share of providers prescribe antibiotics for diarrhea, conditional on the diagnosis. All individual country statistics are calculated using country-specific sampling weights. 95% confidence intervals in parenthesis and number of observations in parenthesis.

Table S13. Access to drugs and medical equipment by country

	(1) Drugs	(2) Minimum medical equipment
Kenya	0.83 (.81-.85) [1646]	0.85 (.84-.87) [3064]
Madagascar	0.43 (.35-.52) [459]	0.67 (.59-.75) [459]
Mozambique	0.46 (.39-.53) [204]	0.77 (.71-.83) [204]
Niger	0.21 (.14-.27) [256]	0.55 (.46-.64) [256]
Nigeria	0.21 (.19-.23) [2358]	0.56 (.53-.58) [2366]
Senegal	NA	0.56 (.39-.74) [151]
Sierra Leone	0.55 (.51-.60) [544]	0.61 (.57-.66) [544]
Tanzania	0.56 (.48-.64) [399]	0.89 (.85-.93) [399]
Togo	0.42 (.27-.56) [180]	0.88 (.75-1.0) [180]
Uganda	0.14 (.09-.20) [390]	0.67 (.61-.73) [390]
All	0.42 (.40-.45) 6,435	0.70 (.67-.73) 8,013

Notes: The table reports the mean access to a set of essential drugs, and minimum set of medical equipment (by country), with all individual country statistics calculated using country-specific sampling weights. 95% confidence intervals in parenthesis and number of observations (facilities) in parenthesis. Drugs take the value 1 if the health facility is stocked with oral rehydration salts, antibiotics (amoxicillin or cotrimoxazole), artemisinin-based combination therapy, and oxytocin, 0 otherwise. Medical equipment takes the value 1 if the health facility is equipped with a minimum set of functional medical equipment (thermometer, stethoscope, and sphygmomanometer), 0 otherwise.

Figure S14. Drugs and medical equipment at the facility

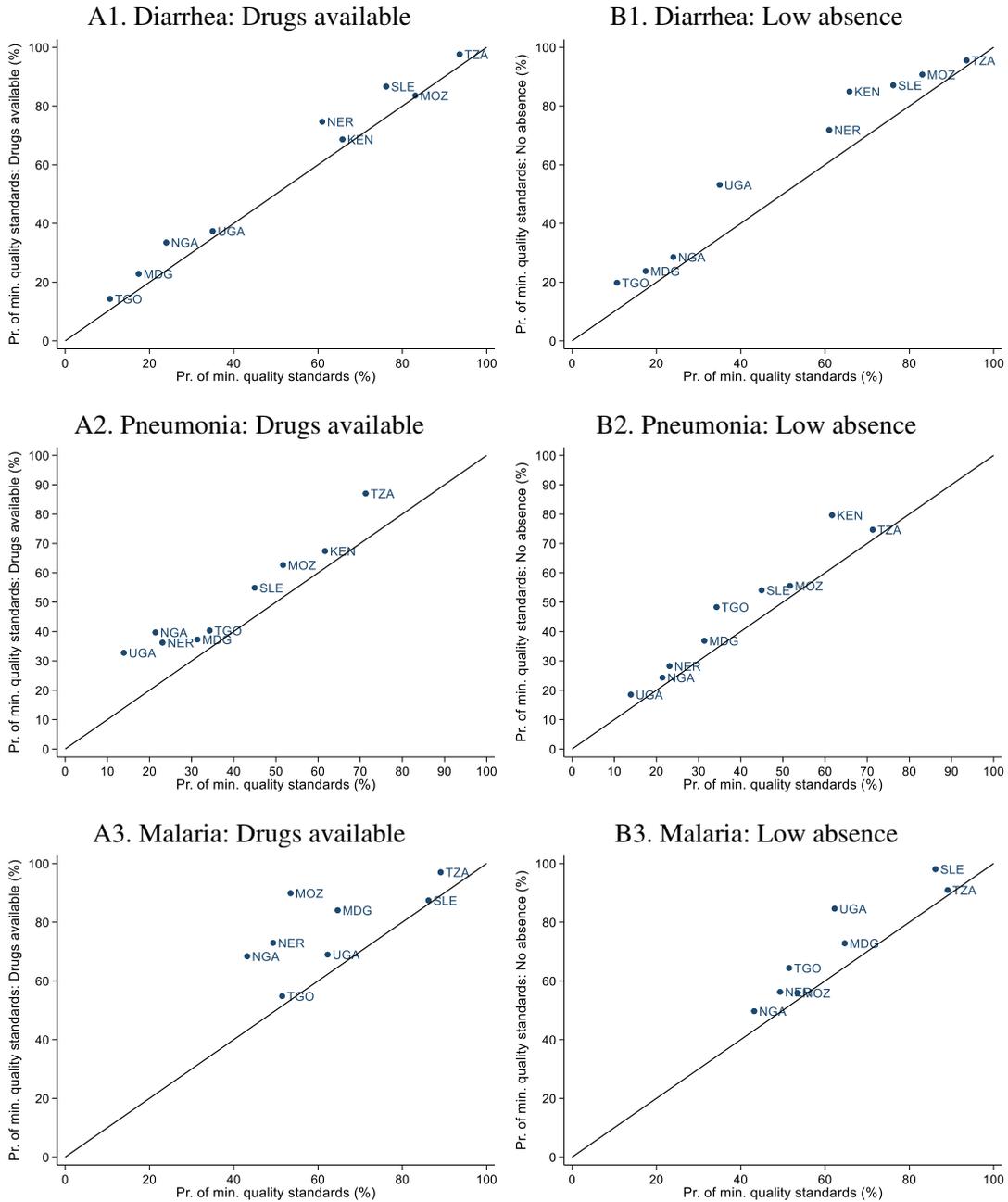
Notes: ACT stands for artemisinin-based combination therapy, a treatment for malaria. ORS stands for oral rehydration salts, a treatment for diarrhea. Dots represent country-specific means, calculated using country-specific sampling weights, vertical bars indicate median performance across countries, and boxes delineate the interquartile range (the middle 50 percent of values). Data are from visual inspections of storage facilities and consulting rooms at the health facilities from Kenya (2018), Madagascar (2016), Mozambique (2014), Nigeria (2013), Niger (2017), Senegal (2010), Sierra Leone (2018) Tanzania (2016), Togo (2014), and Uganda (2013). Medical equipment takes the value 1 if the health facility is equipped with a minimum set of functional medical equipment (thermometer, stethoscope, and sphygmomanometer), 0 otherwise. Drugs take the value 1 if the health facility is stocked with the drug, 0 otherwise. Data on availability of ACT was not collected for Kenya.

Table S15. Probability of care that meets minimum quality standards (%): by country

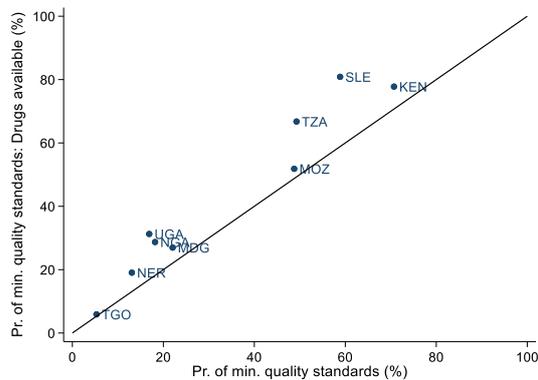
<i>Condition</i>	(1)	(2)	(3)	(4)
	Probability of Care Readiness			No. of facilities
	As observed	Assume available drugs	Assume provider presence	
Kenya	45.3 (43.0-47.6)	51.2 (49.0-53.5)	60.2 (57.7-62.7)	1,646
Madagascar	1.9 (-0.3-4.0)	2.0 (-0.2-4.1)	1.9 (-0.3-4.0)	459
Mozambique	16.4 (11.5-21.4)	19.6 (14.3-24.9)	17.2 (12.0-22.4)	204
Niger	4.3 (2.0-6.6)	8.2 (4.4-11.9)	5.8 (2.8-8.8)	256
Nigeria	3.8 (2.8-4.8)	6.4 (5.2-7.6)	4.3 (3.2-5.4)	2,358
Sierra Leone	25.5 (21.6-29.5)	40.6 (36.4-44.8)	31.0 (26.7-35.4)	399
Tanzania	25.4 (20.1-30.7)	39.1 (32.4-45.9)	27.4 (21.8-33.1)	180
Togo	0.0 -	0.0 -	0.0 -	390
Uganda	1.0 (-0.1-2.1)	4.7 (2.2-7.2)	1.3 (0.1-2.6)	544

Note: Column (1) reports the estimated probability of care readiness that meets minimum quality standards for child, neonatal & maternity care (diarrhea, pneumonia, post-partum hemorrhage, and neonatal asphyxia). Column 2 reports the estimated probability of care readiness that meets minimum quality standards, assuming essential drug(s) (oral rehydration salts or rehydration therapy for diarrhea; antibiotics for pneumonia; and oxytocin for postpartum hemorrhage) for treating the conditions are available. No essential drugs data were collected for neonatal asphyxia. Column 3 reports the estimated probability of care readiness that meets minimum quality standards, assuming no more than 5% absenteeism. All individual country statistics calculated using country-specific sampling weights. 95% confidence intervals in parenthesis. Number of facilities reported in column 4.

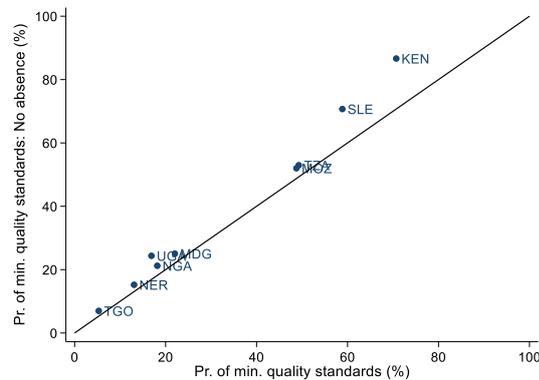
Figure S16. Comparing estimates of minimum quality care conditional on drugs availability and low absence.



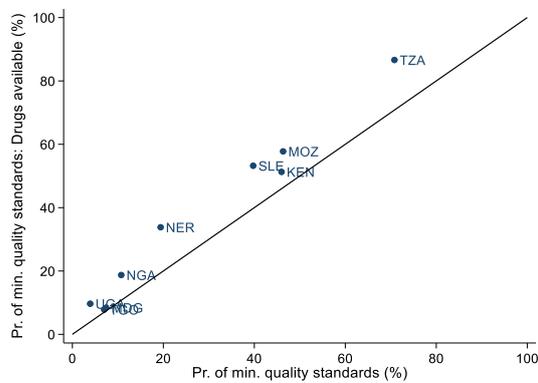
A4. Post-partum hemorrhage: Drugs available



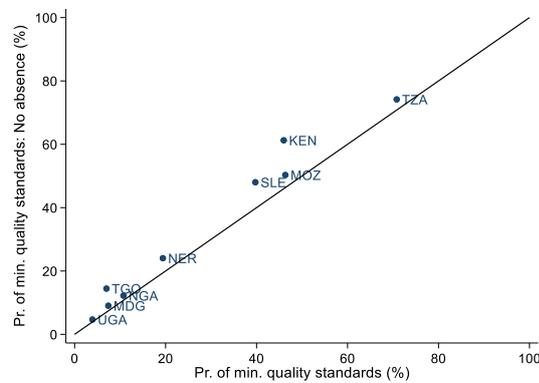
B4. Post-partum hemorrhage: Low absence



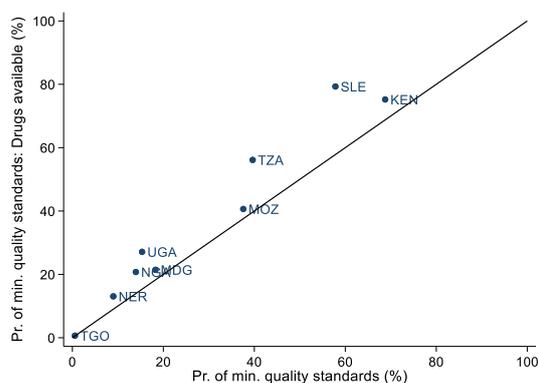
A5. Child care: Drugs available



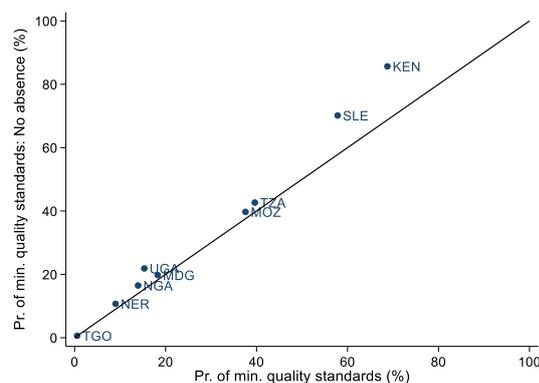
B5. Child: Low absence



A6. Neonatal and maternal care: Drugs available



B6. Neonatal and maternal care: Low absence

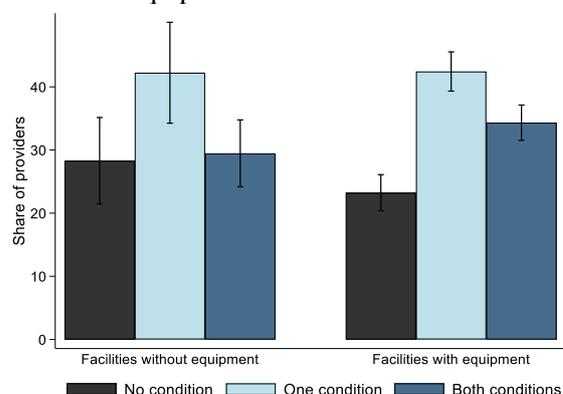


Note: Panel A plots the relationship between the probability of care readiness that meets minimum quality standards vs. the probability of care readiness that meets minimum quality standards assuming essential drugs treating the conditions are available. Panel B plots the relationship between the probability of care readiness that meets minimum quality standards, vs. the probability of care readiness that meets minimum quality standards assuming all providers are present at least 95% of the time. Countries that are higher than the 45 degree line are countries where our estimates suggest that having drugs available (Panel A) and reducing absenteeism (Panel B) would significantly improve the probability of care readiness that meets minimum quality standards. Essential drugs are ORS or rehydration therapy for diarrhea; antibiotics for pneumonia; ACT for malaria; and oxytocin for postpartum hemorrhage. Child care includes two conditions (diarrhea, pneumonia). Neonatal & maternal care includes two conditions (post-partum hemorrhage and neonatal asphyxia). Data are from clinical vignettes,

unannounced visits, and visual inspections from Kenya (2018), Madagascar (2016), Mozambique (2014), Nigeria (2013), Niger (2017), Sierra Leone (2018), Tanzania (2016), Togo (2014), and Uganda (2013). ISO 3-digit alphabetic codes are: KEN (Kenya), MDG (Madagascar), MOZ (Mozambique), NER (Niger), NGA (Nigeria), SLE (Sierra Leone), TZA (Tanzania), TGO (Togo), UGA (Uganda).

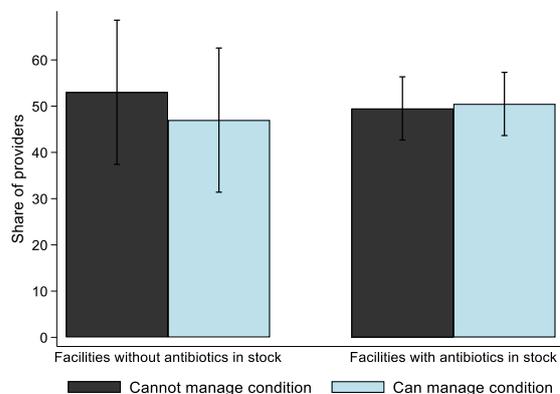
Figure S17. Share of providers who provide accurate diagnosis and treatment conditional on availability of equipment and drugs

A. Medical equipment



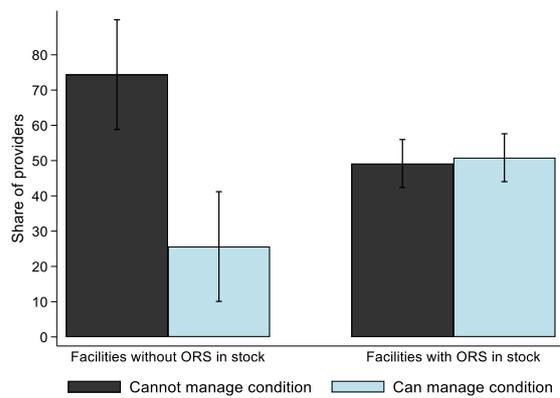
Note: Mean number of conditions (0,1,2) diagnosed and treated correctly by the most knowledgeable provider in the facility conditional on the facility having access or not to a minimum set of functional medical equipment (thermometer, stethoscope, and sphygmomanometer). Two conditions: diarrhea and pneumonia. The estimates are (unweighted) mean outcomes across countries, with the country means calculated using country-specific sampling weights. The error bars represent the 95-percent confidence interval. Data are from clinical vignettes and visual inspections of storage facilities and consulting rooms from Kenya (2018), Madagascar (2016), Mozambique (2014), Nigeria (2013), Niger (2017), Senegal (2010), Sierra Leone (2018) Tanzania (2016), Togo (2014), and Uganda (2013).

B. Antibiotics



Note: Share of the most knowledgeable provider in the facility that can provide correct diagnosis and treatment for pneumonia, conditional on the facility having access or not to antibiotics. The estimates are (unweighted) mean outcomes across countries, with the country means calculated using country-specific sampling weights. The error bars represent the 95-percent confidence interval. Data are from clinical vignettes and visual inspections of storage facilities and consulting rooms from Kenya (2018), Madagascar (2016), Mozambique (2014), Nigeria (2013), Niger (2017), Sierra Leone (2018) Tanzania (2016), Togo (2014), and Uganda (2013).

C. Oral rehydration salts (ORS)



Note: Share of the most knowledgeable provider in the facility that can provide correct diagnosis and treatment for pneumonia, conditional on the facility having access or not to antibiotics. The estimates are (unweighted) mean outcomes across countries, with the country means calculated using country-specific sampling weights. The error bars represent the 95-percent confidence interval. Data are from clinical vignettes and visual inspections of storage facilities and consulting rooms from Kenya (2018), Madagascar (2016), Mozambique (2014), Nigeria (2013), Niger (2017), Sierra Leone (2018) Tanzania (2016), Togo (2014), and Uganda (2013).

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