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Cost effectiveness of medical devices to diagnose pre-eclampsia in low-resource settings

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Abstract

Background—Maternal mortality remains a major health challenge facing developing countries, with pre-eclampsia accounting for up to 17 percent of maternal deaths. Diagnosis requires skilled health providers and devices that are appropriate for low-resource settings. This study presents the first cost-effectiveness analysis of multiple medical devices used to diagnose pre-eclampsia in low- and middle-income countries (LMICs).

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Authors' contributions

Supervised the development of the decision tree framework and generation of the analysis parameters, designed the cost-effectiveness analyses: ZMM. Identified and selected medical devices for inclusion in the study and obtained parameter values for use in the analysis: JPH ASS JA. Contributed to the development of the compendia used in the study: ASS. Performed the cost-effectiveness analyses in TreeAge: ZMM EP AS. Contributed to the development of the decision tree and analysis parameters: TRBJ. Conceived of the study, conceived of and supervised the creation of one of the compendia used in the study, supervised data collection: KS. Drafted the manuscript: ZMM ASS AS TRBJ KHS. Read and approved the final manuscript: ZMM JPH ASS EP AS JA TRBJ KHS.

Conflicts of interest

None declared.

Ethical approval

Not required.

Methods—Blood pressure and proteinuria measurement devices, identified from compendia for LMICs, were included. We developed a decision tree framework to assess the cost-effectiveness of each device using parameter values that reflect the general standard of care based on a survey of relevant literature and expert opinion. We examined the sensitivity of our results using one-way and second-order probabilistic multivariate analyses.

Results—Because the disability-adjusted life years (DALYs) averted for each device were very similar, the results were influenced by the per-use cost ranking. The most cost-effective device combination was a semi-automatic blood pressure measurement device and visually read urine strip test with the lowest combined per-use cost of \$0.2004 and an incremental cost effectiveness ratio of \$93.6 per DALY gained relative to a baseline with no access to diagnostic devices. When access to treatment is limited, it is more cost-effective to improve access to treatment than to increase testing rates or diagnostic device sensitivity.

Conclusions—Our findings were not sensitive to changes in device sensitivity, however they were sensitive to changes in the testing rate and treatment rate. Furthermore, our results suggest that simple devices are more cost-effective than complex devices. The results underscore the desirability of two design features for LMICs: ease of use and accuracy without calibration. Our findings have important implications for policy makers, health economists, health care providers and engineers.

Keywords

cost-effectiveness analysis; medical devices; maternal mortality; pre-eclampsia; low-income and middle-income countries; global health; low resource settings

1. Introduction

Maternal mortality remains a major health challenge facing developing countries. Each year more than 280,000 women die due to complications related to childbirth, with the vast majority of these deaths occurring in low-income countries [1]. Very few countries achieved the Millennium Development Goal 5 of reducing maternal mortality by three-quarters by 2015 [2]. In low-income countries pre-eclampsia accounts for 11 to 17 percent of maternal mortality [3]. Pre-eclampsia is characterized by high blood pressure and elevated levels of protein in the urine. A diagnosis of pre-eclampsia is made if blood pressure is above 140/90 mmHg and there is more than 30 mg/dL of protein in the urine after 20 weeks of gestation [4]. When not properly managed, pre-eclampsia may progress to eclampsia, which is characterized by the onset of seizures and can lead to dangerous complications such as stroke, heart failure, thrombosis, systemic endothelial dysfunction, HELLP syndrome, placental abruption, or even death [4]. The majority of deaths due to pre-eclampsia are preventable if the symptoms can be identified and if treatment can be administered in a timely manner. Treatment for pre-eclampsia consists of medical and surgical options, including the administration of magnesium sulfate and delivery through induction of labor or cesarean section [5].

The diagnosis of pre-eclampsia requires not only skilled health providers, but accurate medical devices that are appropriate for target settings, enabling users to identify pre-

eclamptic women when the condition can be managed. In low- and middle-income countries (LMICs) access to the types of medical devices used to diagnose pre-eclampsia in high-income countries is hindered by high cost, limited trained users and inadequate distribution channels. A recent Lancet Commission called for collective approaches by academia, private sectors, non-governmental and international organizations, and ministries of health to prioritize pressing needs, co-identify appropriate solutions, and devise sustainable implementation plans [6]. Cost-effectiveness analysis provides a unifying framework to guide the allocation of scarce resources to reduce maternal mortality and morbidity. It has important implications for health policy, health care provision, and clinical and biomedical engineering.

Few studies have examined the cost-effectiveness of devices or procedures to reduce maternal mortality in LMICs. Tsu et al. examined the cost-effectiveness of the active management of third-stage labor [7]. Shmueli et al. performed an economic assessment of screening for pre-eclampsia using uterine artery Doppler and serum biomarkers relative to no screening, in an Israeli healthcare system [8]. Simon et al. assessed the cost-effectiveness of using magnesium sulfate for pre-eclampsia in low GNI and high GNI countries, and the incremental cost of preventing one case of eclampsia [9]. Hadker et al. found a novel diagnostic test to be more cost effective in managing a typical pregnancy than are blood tests, urine tests, and uterine artery Doppler ultrasounds from the perspective of the UK health system [10]. Meads et al. compared 27 screening tests in a UK setting and found a no-test, treat-all strategy to be the most cost effective [11]. The current study is the first to generate and compare cost-effectiveness ratios for multiple medical screening devices specifically designed for use in LMICs.

The purpose of this study is three-fold: (1) to tabulate cost and effectiveness data for medical devices appropriate for diagnosing pre-eclampsia in LMICs, (2) to develop a decision-tree framework for evaluating interventions that improve diagnosis and treatment of pre-eclampsia based on the costs and benefits to society as a whole, and (3) to generate cost-effectiveness estimates to guide decision-making in clinical practice and health policy to ultimately reduce maternal mortality. This study presents findings from a decision analysis model of medical devices used to diagnose pre-eclampsia in the population of pregnant women residing in LMICs from a societal perspective.

2. Methods

To identify the set of medical devices to include in this study we conducted several literature searches spanning 2000–2015 using PubMed, SciVerse Scopus, WHO Compendium of New and Emerging Health Technologies, Appropedia's Medical Devices Compendium and Medline. We contacted both health care professionals in the field of maternal health and product developers involved in the design and testing of maternal health related medical devices designed for use in LMICs to inquire about prototypes in the pipeline. Of the initial list of 15 devices identified, only eight devices had sufficient cost and effectiveness data to be included in the study (Table 1).

Pre-eclampsia is diagnosed by blood pressure measurement with subsequent confirmation of proteinuria by urinalysis. The mercury sphygmomanometer is the widely accepted “gold standard” manual blood pressure measurement device. The other blood pressure measurement devices in the study are either automatic or semi-automatic and designed for use in LMICs. Both auscultatory and oscillometric blood pressure measurement devices constrict blood flow with an inflatable arm cuff. Oscillometric devices use volumetric changes in the cuff to calculate blood pressure, while auscultatory devices require a stethoscope to listen to Kortokoff sounds in the downstream artery to determine pressure. Single-use dipsticks are the standard proteinuria testing method in LMICs. Multistix Pro ILS and Bayer Diagnostics India’s Uristik include a color chart to determine the amount of protein in the urine. The Siemens Clinitek 50 device with microalbumin strip test and Siemens/Bayer DCA 2000+ Analyzer accept a urine sample and display the protein content numerically.

The appropriate sensitivity and specificity measurements for each device were drawn from the literature (Table 1). We calculate sensitivity values for the blood pressure devices using the bias-corrected modeling methodology of Wan et al., which predicts sensitivity rates based on information on the mean and standard deviation using predictions from regression modeling of studies included in the systematic review that measured all three parameters[12]. Costs were obtained from the literature where available, and otherwise directly from the manufacturer or distributor.

We developed a decision-tree framework for a study population of pregnant women in LMICs (Figure 1). The decision tree models the probability that a pregnant woman is tested, stepwise, first for pre-eclampsia (i.e. blood pressure measurement), and then followed by confirmatory proteinuria analysis for women with blood pressure above 140/90 mmHg, correctly diagnosed, and treated successfully. Each terminal node reveals the disability-adjusted life years (DALYs) associated with a particular sequence of patient outcomes. We used standard cost-effectiveness methodology to assess costs, DALYs gained relative to the baseline and incremental cost per DALY gained [13, 14]. We identify undominated, absolute dominated and extended dominated interventions. The parameter values in the model reflect the generally available standard of access and care based on our review of the literature and expert opinion (Table 2). We also performed a comprehensive set of sensitivity analyses to evaluate the robustness of our findings to changes in parameter values.

No official measures are recorded for pre-eclampsia testing rates in LMICs, so we assumed that women are tested for pre-eclampsia if they have attended at least four antenatal care visits (51% of pregnancies). This assumption is conservative; many women will be tested for pre-eclampsia during one of the first three prenatal visits, especially if these visits occur late in the pregnancy. Our estimate of the survival rate of untreated pre-eclampsia comes from a population-based survey of a cohort of over 20,000 pregnant women in six West African countries [15]. Nevertheless, the sensitivity analysis includes results from the range of parameter values found in the literature (78.8% – 85.6%) [15, 16]. We modeled magnesium sulfate injection as the standard management technique for pre-eclampsia because it has been accepted as the “drug of choice” for treating pre-eclampsia and is included on the WHO’s list of priority medicines for mothers and children, the package of commodities

needed to achieve the Millennium Development Goal 5 [17]. Magnesium sulfate is easy to administer, even in LMICs, and demonstrably extremely effective, with a success rate of 99.2% in a large international trial [5]. False positives receiving magnesium sulfate should not experience a negative impact on mortality or morbidity. Side effects of the treatment are minimal and rare. The availability of magnesium sulfate is poor in LMICs despite its recent addition to essential drug lists. With little data available about the availability of magnesium sulfate, we used 16% as our best base case estimate based on author's (Johnson) experience, but also examined a wide range of possible values so that our results are generalizable to many settings. If magnesium sulfate is unavailable, clinicians may provide valium or a lytic cocktail, however these alternatives will have little to no effect on mortality rates. Cost of treatment with magnesium sulfate was estimated as \$13 and \$18 per treatment in low- and middle-GNI countries, respectively [9]. The cost of severe morbidity due to eclampsia is \$80 and \$722 in low- and middle-income countries, respectively [9]. To calculate DALYs associated with the 3.8% of cases resulting in permanent disability due to severe complications of pre-eclampsia, we used a health-related quality of life (HRQL) weight of 0.25 [18–20]. The only parameters in the model that vary by device are sensitivity and specificity. The time period of analysis capture outcomes for the entire lifespan of the pregnant women in our simulation.

All costs are expressed in 2015 U.S. dollars and, in the absence of available data on training or maintenance costs, reflect only the incremental cost of purchasing the device. The devices are designed to be easy to use and require minimal training; their use can be built into existing provider care and no additional equipment is needed. We therefore do not include provider training, time, or additional equipment costs.

To identify the impact of uncertainty about parameter values on our results we performed one-way sensitivity analyses as well as second-order probabilistic multivariate sensitivity analyses. We performed a standard set of 5,000 independent Monte Carlo simulations where the outcome at each node of the decision tree was determined by draws from independent binomial distributions. For parameters that the one-way sensitivity analyses identified as being particularly influential, we examined a wide range of plausible values in the Monte Carlo analysis. Because survival rates with and without access to pre-eclampsia treatment are correlated, we examined two additional scenarios: a high-resource/high-skill environment with higher than base-case survival rates at all terminal nodes and a low-resource/low-skill environment with relatively lower survival rates at all terminal nodes. Analyses were performed with TreeAge Pro 2015.

3. Results

Table 3 shows the main results on the cost-effectiveness of each intervention. Because the DALYs averted for each device were very similar, the results primarily depended on the per-use cost ranking. With the lowest combined per-use cost of 0.204 cents and incremental cost-effectiveness ratio of \$93.6 per DALY gained relative to a baseline scenario with no access to testing or treatment, the Microlife semi-automatic blood pressure monitor with the Uristik single-use proteinuria strips was the most cost-effective combination of medical devices to improve the diagnosis of pre-eclampsia. The device combinations on the cost-

effectiveness frontier (undominated) were, in order of increasing incremental cost-effectiveness ratio, Nissei DM3000/CLINITEK strip test, the sphygmomanometer/CLINITEK strip test, and the sphygmomanometer/DCA 2000+. The SpotVital blood pressure device and the Multistix proteinuria test were strongly dominated in all combinations. To be on the cost-effectiveness frontier, the SpotVital device would need to have a per-use cost that implies well over 10,000 uses at the quoted price, which is implausible in LMICs. Extended (weak) dominance ruled out two combinations: Nissei DM3000/Uristik and sphygmomanometer/ Uristik.

We performed one-way sensitivity analyses of the results to changes in our benchmark model parameters (see tornado diagram in Figure 2). Our results were most sensitive to the specificity rate for the blood pressure device. Our results were insensitive to parameter values that were very high (treatment success rate, survival rate for untreated pre-eclampsia) or very low (rate of pre-eclampsia, rate of severe morbidity due to untreated or unsuccessfully treated pre-eclampsia).

To address the issue that minimal clinical trial data exists to identify some of our benchmark decision tree parameters, we present the results of a comprehensive sensitivity analysis (Table 4). The first column shows results from a set of simulations based on the base case parameters for the most cost-effective device combination of the Microlife blood pressure measurement device with the Uristik proteinuria strips. The largest improvements in the ICER come from increases in the specificity of the blood pressure measurement (Microlife) and the treatment access rate for pre-eclampsia. It is worth noting that in countries with a life expectancy greater than 65 years, this intervention pays for itself. In no case does the ICER estimate for the Microlife, the most cost-effective pre-eclampsia diagnosis device, exceed the threshold to be considered cost-effective in low-income countries [21].

Survival rates for untreated pre-eclampsia and successfully treated pre-eclampsia are likely to be positively correlated based on overall health status, access to health care, available health resources and the skill level of health care providers. Columns 2 and 3 of Table 4 show the results from a similar sensitivity analysis to as Column 1, but using the high-resource and low-resource scenario values. These scenarios typify comparisons of countries (e.g. South Africa vs. Nigeria or India), regions (e.g. urban vs. rural), or facility types (e.g. tertiary hospital vs. local clinic). The Microlife device had an ICER of \$78.88 in the low resource environment relative to the benchmark scenario with no access to testing and \$115.05 in the high resource environment.

4. Discussion

This study is designed to help translate scientific advances into policy and practice in low-income countries. We provide a clear framework for decision-making and assess the areas that are most sensitive to uncertainty. In the era of implementation science and implementation engineering, cost-effectiveness should guide not only the development of new devices and procedures but also the implementation and evaluation [22].

Our findings show that among the devices analyzed, the most cost-effective devices for diagnosing pre-eclampsia are simple rather than complex medical devices. For example, the Microlife blood pressure device is manually inflated but provides automated read-outs, and the Uristik strip test is read visually, in comparison with devices such as the SpotVital Signs or the DCA 2000+ which are fully automated. These devices are all low-cost and feasible to implement even in very low-income countries. These devices are likely to be at least as cost-effective as screening for pre-eclampsia using uterine artery Doppler and serum biomarkers [8], prophylactic magnesium sulfate for pre-eclampsia [9], community-based distribution of misoprostol for treatment or prevention of postpartum hemorrhage [23], screening for bacteriuria or syphilis, skilled birth attendance, management of obstructed labor, postpartum hemorrhage and maternal sepsis [24]. Furthermore, these devices are well within the range considered cost-effective in low-income countries based on the World Health Organization definition.

Our cost-effectiveness estimates were not particularly sensitive to changes in testing rate, however they were sensitive to changes in device specificity and the treatment access rate (Table 4). Increasing the testing rate by 20 percentage points (from 40% to 60%) has almost no effect on the ICER whereas increasing the blood pressure device specificity by 20 percentage points (from 70% to 90%) reduces the ICER by \$180. Intuitively, there is little benefit to identifying pre-eclamptic women when treatment access is low. This has important implications for device design: when treatment rates are low, a reasonably sensitive device that is low in cost is preferred to a highly-sensitive but high-cost device. When it comes to design for global health, the perfect may be the enemy of good.

Our results highlight the importance of two desirable features for LMIC settings: reusability and accuracy in the absence of calibration. High recurring costs, such as single-use components and maintenance costs, limit the cost-effectiveness of these devices in low-income countries. Even though Microlife and Nissei DM3000 require batteries (or AC power) to operate, their batteries are rechargeable rather than disposable. The most cost-effective devices for blood pressure measurement (Microlife, Nissei DM3000 and Mercury sphygmomanometer) are highly accurate and require limited to no calibration.

Other major design features common to the most cost-effective devices are the ability to withstand variable environmental conditions (temperature and humidity), portability and usability, which are essential characteristics of devices to be used by health care workers in rural areas. Microlife and Nissei DM3000 have digital screens for displaying blood pressure, which minimizes the likelihood of human error. Uristik and Multistix test strips can be easily interpreted by the immediate change in color in response to the urine droplet.

Our assumptions are all conservative and our results are robust to variance on the dimensions of uncertainty as demonstrated in Tables 4. Our estimates understate the DALYs gained from each device in that they don't include the health benefits to the unborn child from improved diagnosis of pre-eclampsia, which are likely quite substantial. While these results do not provide accurate absolute cost effectiveness of testing diagnostic in scenarios where diagnostic equipment are not already accessible and must be purchased; however, the

analysis does provide informative data on the relative cost effectiveness of testing women for preeclampsia using these common technologies.

One limitation of our study is the limited data available on several input parameters of our model and on medical device effectiveness and maternal mortality in general. More precise measures of these input parameters will improve our cost-effectiveness estimates to better guide implementation and health policy. Values from standardized trials similar to the MAGPIE trial that compare effectiveness across devices and across settings would be ideal. We have addressed this limitation with a comprehensive set of probabilistic sensitivity analyses, however, additional effectiveness and survival data would enable us to widen the range of devices examined and draw more specific policy implications from the study. We are not able to model the full cost structure due to limited data on labor costs for training and delivering the test, device maintenance costs, time patients spend traveling and waiting for care, costs of follow-on-care following serious complications due to pre-eclampsia and offsetting health care cost savings due to prevented cases of eclampsia.

Cost-effectiveness analysis considers every stage of implementation in the local environment from when the patient presents at a health facility through diagnosis, treatment and patient outcome. By taking into account the big picture, it makes implicit tradeoffs explicit such as diagnosis versus treatment or cost versus effectiveness. In settings where treatment is rationed or unavailable, the most effective diagnostic device is not necessarily the most desirable, but rather the one that achieves a certain threshold of effectiveness at the lowest cost. Our results provide additional evidence that "hand me down" technology from high-income countries is unlikely to be appropriate in LMICs because developed countries have focused on the most effective solution, regardless of cost [6, 25, 26].

Our results have important implications for routine healthcare provision in both clinical and policy contexts. While other studies have evaluated the cost effectiveness of screening and treatment strategies in high income countries, to our knowledge no other study has evaluated the cost effectiveness of medical screening technologies designed for use in LMIC. While other studies have evaluated the cost effectiveness of screening and treatment strategies in high income countries, to our knowledge no other study has evaluated the cost effectiveness of medical screening technologies designed for use in LMIC. These are precisely the type of results needed to inform evidence-based medicine, which allows policy-makers to standardize procedures in LMICs and thereby reduce maternal mortality. Cost-effectiveness analysis is an important tool that incorporates information about the local environment to produce tailored policies, devices, and procedures.

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Appendix

Complete decision tree representing the pathways of diagnosis and treatment for a pre-eclampsic pregnant woman presenting at an antenatal care facility in a developing country.

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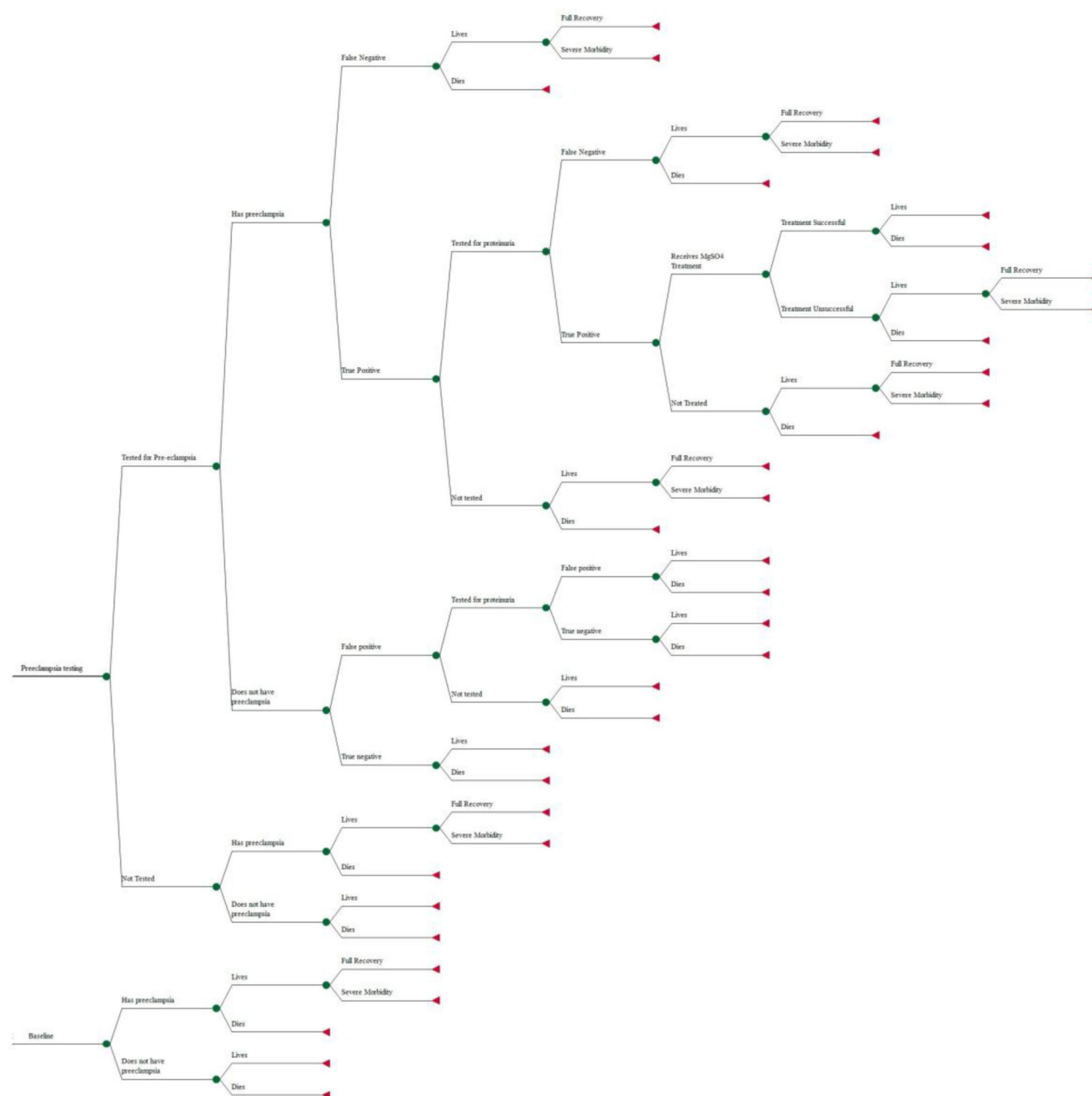


Figure 1. A standard decision tree representing the pathways of diagnosis and treatment for a pre-eclamptic pregnant woman presenting at an antenatal care facility in a developing country

Notes: Full analysis includes analogous branches of decision tree for non-pre-eclamptic pregnant woman. Triangles represent terminal nodes. Parameter values are found in Table 2.

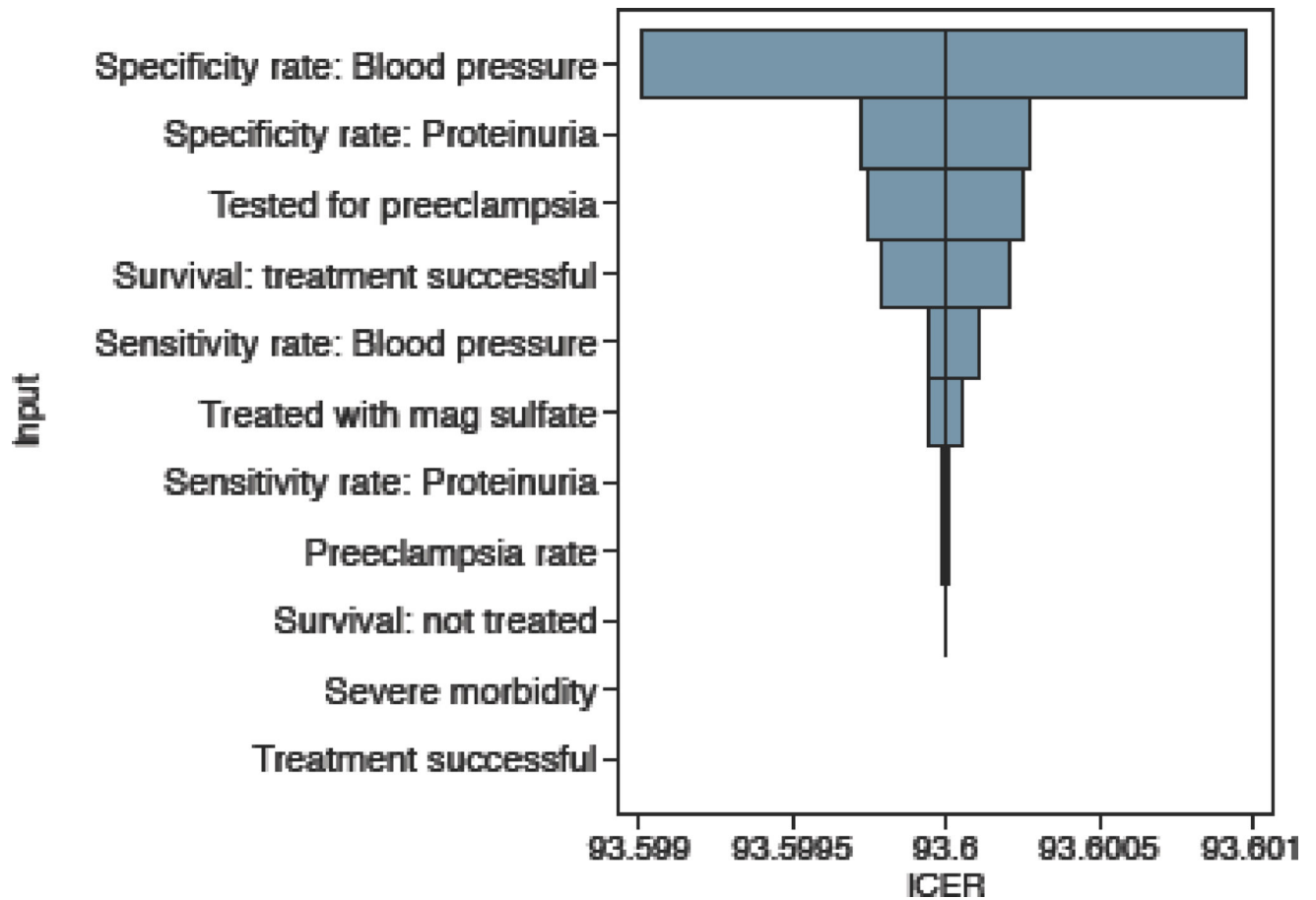


Figure 2. Tornado diagram: One-way sensitivity analysis of cost-effectiveness of pre-eclampsia intervention to variations in decision tree parameters

Notes: Base case = ICER of \$93.6/ DALY gained using Microlife/Uristik device combination. Standard tornado diagram calculated from base case parameter values, varying one parameter at a time by .1 percentage point. Inputs ranked by size of effect on output mean.

Table 1

Proteinuria and blood-pressure devices

Device Category	Device Type	Device Name	Description	Per-use Cost (\$)	Device sensitivity	Device specificity
Diagnostic: Proteinuria	Dipstick Test	CLINITEK Status microalbumin strip test [27]	Clinitek Status automated analyzer with single use plastic strips that detect albumin and creatinine in urine.	5.70–5.85	0.955	0.533
		Multistix Pro 10LS [28]	Single use plastic strips that detect proteinuria (visually read).	3.29	0.263	0.729
		Uristik [29]	Single use plastic strips that detect glucose and protein in urine (visually read).	0.2	0.680	0.680
		DCA 2000+ [27]	Automated, battery-powered device that displays the proteinuria content of a urine sample on an LCD screen.	12.90	0.994	0.840
Diagnostic: Blood-pressure	Auscultatory	Mercury Sphygmomanometer [30]	Manual device that uses a mercury column to display the pressure of a cuff inflated around the upper arm. Requires a trained operator to listen to blood sounds with a stethoscope.	0.009–0.039	0.966	0.877
	Hybrid	Nissei DM-3000 [31]	Hybrid device with manual auscultatory and automatic oscillometric settings, with digital display. This study uses values from the auscultatory setting only.	0.076–0.11	0.918	0.914
	Oscillometric	Spot Vital Signs [32]	Rechargeable battery-powered device that automatically inflates cuff, obtains blood pressure through a built-in algorithm, and displays result on a digital readout.	0.15 – 0.29	0.931	0.858
		MicroLife [33]	Manually inflated device that automatically determines blood pressure with a built-in algorithm, and displays result digitally.	0.004	0.910	0.905

Notes: Costs are in US 2015 dollars.

Table 2

Parameters, values, and sources used in pre-eclampsia decision tree analysis

Parameter Description	Value	Source
% of pregnancies that are pre-eclamptic	2.80%	WHO Global Survey on maternal and perinatal health 2009 [34]
% of pregnancies tested for pre-eclampsia	51.00%	WHO Global Health Observatory [35]
Survival rate for undiagnosed or untreated pre-eclampsia	94.40%	Pruhal, 2000 [15]
% of diagnosed pre-eclampsia treated	16.00%	Varies worldwide; Dr. Johnson
Success rate of treatment among pre-eclamptic pregnancies	99.20%	The Magpie Trial Collaborative Group [5]
Survival rate for successfully treated pre-eclamptic pregnancies	99.43%	World Health Statistics 2011 [36]; Dr. Martin
Survival rate for unsuccessfully treated pre-eclamptic pregnancies	94.40%	Pruhal, 2000 [15]; Dr. Martin
Rate of severe complications for pre-eclampsia survivors	3.80%	Sibai, 2005 [4], Steeger, 2010 [37]
HRQL for lifetime following severe complications	0.25	Lee et al., 2010, Aberg et al. (2011), WHO (2004) [18–20]
Life expectancy	55 yrs	
Discount rate	3%	

Table 3

Costs, DALYs averted and incremental cost effectiveness ratios (ICER) for devices used to detect pre-eclampsia.

Device	Total cost of intervention	Incremental cost	DALYs	Incremental effectiveness (DALYs averted)	ICER (cost per DALY gained)
Baseline (no screening)	0.080		0.05137		
Microlife, Uristik	0.305	0.2243	0.04898	0.00240	94
Nissei (low), Uristik	0.322	0.0174	0.04896	0.00002	829
Nissei (high), Uristik	0.339	0.0173	0.04896	0.00000	*
Sphygmo (low), Uristik	0.369	0.0467	0.04883	0.00013	368
Sphygmo (high), Uristik	0.384	0.0153	0.04883	0.00000	*
Microlife, Multistix	0.452	0.0828	0.05045	-0.00162	*
Sphygmo (low), Multistix	0.457	0.0883	0.05039	-0.00156	*
Nissei (low), Multistix	0.458	0.0895	0.05044	-0.00161	*
Nissei (high), Multistix	0.476	0.1068	0.05044	-0.00161	*
Spot Vital Signs (low), Uristik	0.481	0.1123	0.04892	-0.00009	*
Spot Vital Signs (high), Uristik	0.553	0.1837	0.04892	-0.00009	*
Sphygmo (high), Multistix	0.567	0.1981	0.05039	-0.00156	*
Spot Vital Signs (low), Multistix	0.686	0.3171	0.05042	-0.00160	*
Nissei (low), CLINITEK(low)	0.716	0.3472	0.04798	0.00085	408
Nissei (low), CLINITEK(high)	0.724	0.0084	0.04798	0.00000	*
Microlife, CLINITEK(low)	0.731	0.0150	0.04801	-0.00003	*
Nissei (high), CLINITEK(low)	0.733	0.0173	0.04798	0.00000	*
Nissei (high), CLINITEK(high)	0.742	0.0257	0.04798	0.00000	*
Spot Vital Signs (high), Multistix	0.757	0.0413	0.05042	-0.00245	*
Microlife, CLINITEK	0.758	0.0418	0.04801	-0.00003	*
Sphygmo (low), CLINITEK(low)	0.903	0.1867	0.04780	0.00018	1055
Sphygmo (low), CLINITEK(high)	0.914	0.0112	0.04780	0.00000	*
Sphygmo (high), CLINITEK(low)	0.918	0.0153	0.04780	0.00000	*
Sphygmo (high), CLINITEK(high)	0.929	0.0265	0.04780	0.00000	*
Nissei (low), DCA 2000+	0.948	0.0453	0.04784	-0.00004	*
Nissei (high), DCA 2000+	0.965	0.0627	0.04784	-0.00004	*

Device	Total cost of intervention	Incremental cost	DALYs	Incremental effectiveness (DALYs averted)	ICER (cost per DALY gained)
Microlife, DCA 2000+	0.976	0.0738	0.04787	-0.00007	*
Spot Vital Signs (low), CLINITEK(low)	1.082	0.1792	0.04793	-0.00013	*
Spot Vital Signs (low), CLINITEK(high)	1.094	0.1917	0.04793	-0.00013	*
Spot Vital Signs (high), CLINITEK(low)	1.153	0.2506	0.04793	-0.00013	*
Spot Vital Signs (high), CLINITEK(high)	1.166	0.2631	0.04793	-0.00013	*
Sphygmo (low), DCA 2000+	1.199	0.2959	0.04765	0.00015	2026
Sphygmo (high), DCA 2000+	1.214	0.0153	0.04765	0.00000	*
Spot Vital Signs (low), DCA 2000+	1.404	0.2057	0.04779	-0.00013	*
Spot Vital Signs (high), DCA 2000+	1.476	0.2771	0.04779	-0.00013	*

Notes: Costs in US 2015 dollars. ICERs from decision tree simulation where testing rate is not 100%. Devices ordered by cost of intervention. Effectiveness calculated relative to scenario with no pre-eclampsia. Baseline scenario has no access to diagnostic testing or clinical management (see text).

Asterisk (*) indicates that ICER is not applicable due to dominance by prior strategy. These rows are skipped in calculating incremental costs and effectiveness in subsequent rows.

Italics indicates extended dominance; see text.

Table 4

Sensitivity analysis for base case, low-resource and high-resource country scenarios

	ICER	ICER	ICER
	Base case	Low resource	High resource
Tested rate			
20	93.58	78.85	114.95
40	93.63	78.89	115.02
51 (base case)	93.62	78.88	115.05
60	93.62	78.90	115.05
80	93.61	78.89	115.06
Treated rate			
16 (base case)	93.62	78.88	115.05
20	76.10	64.12	93.52
40	41.04	34.60	50.42
60	29.36	24.76	36.06
80	23.52	19.84	28.87
Device sensitivity, Microlife			
70	119.59	100.77	146.99
80	105.49	88.90	129.71
90	94.57	79.71	116.21
91 (base case)	93.62	78.88	115.05
95	89.95	75.81	110.55
99	86.62	72.99	106.47
99.5	86.22	72.67	105.96
Device sensitivity, Uristik			
68 (base case)	93.62	78.88	115.05
70	91.10	76.79	111.99
80	80.47	67.83	98.89
90	72.18	60.84	88.74
95	68.70	57.91	84.43
99	66.17	55.78	81.33
99.5	65.88	55.52	80.94
Device specificity, Microlife			
70	278.55	234.62	342.39
80	188.34	158.65	231.49
90	98.13	82.68	120.60
90.5 (base case)	93.62	78.88	115.05
95	53.03	44.69	65.15
99	16.95	14.30	20.63
99.5	12.44	10.50	15.25
Device specificity, Uristik			
68 (base case)	93.62	78.88	115.05

	ICER	ICER	ICER
	Base case	Low resource	High resource
70	88.51	74.57	108.77
80	62.96	53.05	77.36
90	37.41	31.54	45.95
95	24.63	20.78	30.24
99	14.41	12.17	17.68
99.5	13.14	11.09	16.10
Discount rate			
0%	0.02	0.02	0.02
3% (base case)	93.62	78.88	115.05
6%	-0.03	-0.03	-0.03
Life expectancy			
55 (base case)	93.62	78.88	115.05
65	0.07	0.07	0.07
HRQL			
0.2	167.40	123.91	257.45
0.25 (base case)	93.62	78.88	115.05
0.3	64.98	57.87	74.08

Notes: Table reports simulation results of ICERs from varying the mean of each parameter as indicated for Microlife/Uristik device combination. Each cell represents a separate simulation. Low-resource country scenario (e.g., Nigeria) has high general mortality: untreated survival 92.4%; treated survival 98.93%. High-resource country scenario (e.g., South Africa) has low general mortality: untreated survival 96.4%; treated survival 99.93%.