BMJ Open Impact evaluation of the free maternal healthcare policy on the risk of neonatal and infant deaths in four sub-Saharan African countries: a quasi-experimental design with propensity score Kernel matching and difference in differences analysis

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ABSTRACT

To cite: Dwomoh D, Agyabeng K, Agbeshie K, *et al.* Impact evaluation of the free maternal healthcare policy on the risk of neonatal and infant deaths in four sub-Saharan African countries: a quasi-experimental design with propensity score Kernel matching and difference in differences analysis. *BMJ Open* 2020;**10**:e033356. doi:10.1136/ bmjopen-2019-033356

Prepublication history and additional material for this paper are available online. To view these files, please visit the journal online (http://dx.doi. org/10.1136/bmjopen-2019-033356).

Received 01 August 2019 Revised 29 January 2020 Accepted 19 March 2020

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Correspondence to Dr Duah Dwomoh; duahdwomoh@yahoo.com **Objective** Despite the huge financial investment in the free maternal healthcare policy (FMHCP) by the Governments of Ghana and Burkina Faso, no study has quantified the impact of FMHCP on the relative reduction in neonatal and infant mortality rates using a more rigorous matching procedure with the difference in differences (DID) analysis. This study used several rounds of publicly available population-based complex survey data to determine the impact of FMHCP on neonatal and infant mortality rates in these two countries.

Design A quasi-experimental study to evaluate the FMHCP implemented in Burkina Faso and Ghana between 2007 and 2014.

Setting Demographic and health surveys and maternal health surveys conducted between 2000 and 2014 in Ghana, Burkina Faso, Nigeria and Zambia.

Participants Children born 5 years preceding the survey in Ghana, Burkina Faso, Nigeria and Zambia.

Primary outcome measures Neonatal and infant mortality rates.

Results The Propensity Score Kernel Matching coupled with DID analysis with modified Poisson showed that the FMHCP was associated with a 45% reduction in the risk of neonatal mortality rate in Ghana and Burkina Faso compared with Nigeria and Zambia (adjusted relative risk (aRR)=0.55, 95% Cl: 0.40 to 0.76, p<0.001). In addition, infant mortality rate has reduced significantly in both Ghana and Burkina Faso by approximately 54% after full implementation of FMHCP compared with Nigeria and Zambia (aRR=0.46, 95% Cl: 0.36 to 0.59, p<0.001).

Conclusion The FMHCP had a significant impact and still remains relevant in achieving Sustainable Development Goal 3 and could provide lessons for other sub-Saharan countries in the design and implementation of a similar policy.

Strengths and limitations of this study

- The use of more rigorous statistical methods and data from repeated cross-sectional surveys improves the robustness of the impact estimate.
- This remains the first study that has quantified the impact of free maternal healthcare policy (FMHCP) on child survival.
- Evidence from this study can be used to inform policy decisions about the implementation of FMHCP in other sub-Saharan African countries.
- Unobserved factors could bias our study results if these factors affected interventions and comparison countries in different ways.
- We cannot interpret our results as causal since the data originate from a cross-sectional study design.

INTRODUCTION

Access to primary healthcare services remains low in many low-income and middle-income countries. According to the WHO report 2017, approximately half of the world population lack access to essential health services and it is estimated that over 100 million population are still pushed into extreme poverty because of out-of-pocket health expenditure. Peters and colleagues¹ as well as Jacobs and colleagues² have classified these factors into four main dimensions, namely geographical access, financial access, availability of healthcare and acceptability of healthcare service. Delay or lack of access to healthcare services due to financial constraints can affect child survival. Following the Abuja declaration for sub-Saharan African (SSA) countries to spend 15% of its public spending on healthcare at the turn of the century, Ghana in 2003 set up a National Health Insurance Scheme (NHIS) as a way of improving Universal Health Coverage.³ In September 2003, a policy exempting women in the four poorest regions of Ghana from delivery care fees was introduced by the Government of Ghana in an attempt to increase skilled birth attendance and reduce inequality in use of healthcare services.⁴ The policy was rolled out in all the 10 regions by the end of April 2005 but with serious challenges. Notable among them was the fact that the disbursement of funds to accredited health facilities was not forthcoming, and in October 2005 some health facilities started to charge clients again.⁴ In July 2008, the Government of Ghana through the NHIS implemented a national user free maternal care exemption policy to improve financial access to maternal health services and reduce maternal mortality rate, perinatal mortality rate, neonatal mortality rate (NMR) and infant mortality rate (IMR). The policy was popularly referred to as the free maternal healthcare policy (FMHCP). The main aim of the policy was to address financial barriers to demand healthcare services.

Burkina Faso is one of the countries in SSA which failed to achieve the target for Millennium Development Goal (MDG) goal number 5 (reduction of maternal mortality by 75% between 1990 and 2015).⁵ That notwithstanding, tremendous efforts have been made by Burkina Faso towards ensuring equitable access to maternal care services. For instance, maternal health financing and delivery reforms were developed and implemented, among which are the abolition of user fees for antenatal care (ANC) services in 2002, subsidisation of delivery costs for all women by 80% and by 100% for the poorest in 2007 and exemption of the poorest from payment of all user fees for all curative and preventive health services in 2009.⁶⁷ In this article, we refer to the policy implemented in Burkina Faso as FMHCP for easy reference to countries that have implemented the intervention.

Nigeria, for instance, did not have a clear federal policy on user fees in maternal and child health, and the regional variation at the primary and secondary levels is vast.⁸ Although Zambia removed user fees in 2006 in rural areas only,^{9 10} the policy had not been implemented properly and no impact had been seen in the following year or two.¹¹ That notwithstanding, fees are still payable (by cash) in urban areas and financial constraints still remain a significant barrier to institutional delivery.¹¹ The impact of these policies, particularly on access to health services and neonatal mortality, has not been evaluated using rigorous methods, and so the empirical basis for defending these policies is weak.¹² To determine the effectiveness of FMHCP in contributing to a reduction in the mortality rate relative to countries that do not have such policy, Propensity Score Kernel Matching with the difference in differences (DID) analysis was applied. Using a quasi-experimental design, the goal of this study is to determine whether the full implementation of FMHCP in Ghana and Burkina Faso contributed to the relative reduction in NMR and IMR between 2008 and

2014 compared with Nigeria and Zambia without such significant national health financing policy on maternal healthcare.

METHODS AND ANALYSES

Data sources

The data used in this study were obtained from 11 separate Demographic and Health Surveys (DHS) and 1 Malaria Indicator Survey (MIS). The DHS and MIS are nationally representative cross-sectional surveys which include common questions about the year of birth and survival status of all births to women of reproductive age (15-49 years). The DHS and MIS data sets are freely available and could be downloaded at the DHS website (http://dhsprogram.com) after completing the online data request registration form. With the exception of Burkina Faso that could not provide DHS but MIS data for 2014, each country contributed three different DHS data sets that were conducted between 2000 and 2014. That is, we used the pre-baseline data from 2001/2003to 2007/2008; baseline data 2007/2008 and end-line data 2013/2014. The unit of analysis in this study is the children of women born in 5 years (0-59 months) preceding the survey. Detailed distribution about number of live births in 5 years preceding the survey, number of women aged 15-49 interviewed, total number of women aged 15-49 in the country at the time of the survey, year of survey and survey response rate for eligible women, NMR and IMR per 1000 live births and cumulative incidence rate per 1000 person-years at risk can be found in online supplementary appendix table S1 A.

Patients and public involvement

Patients and the public were not involved.

Primary outcome measures

The primary outcomes of interest were IMR and the NMR. In this analysis, the IMR is defined as the probability of dying between birth and first birthday whereas NMR is defined as the probability of dying between birth and the first month of life.¹³ All deaths that were recorded within the first 28 days after birth were coded as 1 or otherwise 0 in defining a binary indicator variable for neonatal mortality. For infant mortality, deaths within 1 year after birth in the 5 years preceding each survey were coded as 1 otherwise 0 to define a binary indicator variable for infant mortality.

Exposure to FMHCP

Countries that have abolished at least 80% of user fees for institutional delivery in SSA between the periods of 2007 and 2014 and have DHS or MIS data readily available were included in this study as intervention countries. That notwithstanding, these countries should have conducted DHS between the periods of 2000 and 2008. This was necessary to test the parallel trend assumption which is a requirement for the validity of DID design
 Table 1
 Trend of neonatal and infant mortality between countries with and without FMHCP and description of the study participants: 2007/2008–2013/2014

		Intervention: FMHCP impl	ementation	
	Total %	No FMHCP %*	FMHCP %*	Rao–Scot χ ²
All-cause mortality in 5 years preceding the survey (95% Cl)	9.2 (8.9–9.5)	9.8 (9.5–10.1)	6.2 (5.9–6.6)	159.60***
All-cause neonatal deaths in 5 years preceding the survey (95% CI)	3.5 (3.3–3.6)	3.1 (2.9–3.3)	0.4 (0.3–0.4)	76.70***
All-cause infant deaths in 5 years preceding the survey (95% CI)	6.7 (6.5–7.0)	7.3 (7.1–7.6)	4.0 (3.6–4.3)	168.40***
Sex of household head				114.03***
Male	97 430 (88.4)	70 247 (83.4)	27 183 (16.6)	
Female	13 318 (11.6)	9740 (74.3)	3578 (25.7)	
Wealth quintile				2.00
Poorest	26 597 (23.3)	19 264 (82.9)	7333 (17.1)	
Poorer	25 526 (22.7)	18 862 (83.3)	6664 (16.7)	
Middle	22 913 (19.4)	16 412 (81.4)	6501 (18.6)	
Richer	20 303 (18.2)	14 198 (80.7)	6105 (19.3)	
Richest	15 409 (16.5)	11 251 (82.9)	4158 (17.1)	
Household size				20.26***
1–4	26 784 (25.8)	19 215 (79.8)	7569 (20.2)	
5–7	45 709 (41.5)	33 951 (82.9)	11 758 (17.1)	
8+	38 255 (32.8)	26 821 (83.5)	11 434 (16.5)	
Access to improved water				121.32***
Improved	89 000 (80.4)	61 284 (80.1)	28 049 (19.9)	
Unimproved	21 000 (19.6)	18 676 (91.4)	2711 (8.6)	
Missing	28 (0.01)	27 (98.4)	1 (1.6)	
Access to an improved toilet fac	cility			195.72***
Improved, not shared	26 000 (27.0)	22 493 (91.7)	3817 (8.3)	
Improved, shared	21 000 (22.5)	13 047 (71.0)	7762 (29.0)	
Unimproved	63 000 (50.1)	44 120 (82.4)	19 095 (17.6)	
	414 (0.4)	327 (81.2)	87 (18.8)	0.01
Place of residence	20,607 (20,0)	05 025 (02 0)	7500 (17 1)	0.61
Urban	32 627 (32.2)	25 035 (82.9)	7592 (17.1)	
Household ownership of bodnet	10 121 (01.0)	54 952 (62.0)	23 109 (16.0)	1012 50***
No bednet	43 000 (46 4)	36 880 (92 4)	6015 (7.6)	1013.32
Rednet	68 000 (53 6)	43 062 (73 6)	24 746 (26 5)	
Missing	45 (0.06)	45 (100 0)	0 (0 0)	
Mothers' current age	()		0 (0.0)	11.10***
<18 vears	3558 (3.3)	2803 (86.5)	755 (13.5)	
18–34 vears	80 000 (71.5)	58 000 (82.3)	22 000 (17.7)	
35+	27 000 (25.2)	20 000 (81.8)	7727 (18.2)	
Mothers' education				44.98***
None	53 000 (46.5)	32 000 (79.4)	21 000 (20.6)	
Primary	29 000 (23.2)	24 000 (86.6)	4475 (13.4)	
JHS	25 000 (25.5)	20 000 (81.7)	4686 (18.3)	
Secondary or higher	4241 (4.8)	3882 (92.4)	359 (7.6)	
Missing	16 (0.01)	11 (64.6)	5 (35.4)	

Continued

		Intervention: FMHCP imp	lementation	
	Total %	No FMHCP %*	FMHCP %*	Rao–Scot χ ²
Birth order				271.39***
1st birth	24 000 (21.1)	16 000 (75.8)	8167 (24.2)	
2nd birth	21 000 (18.9)	14 000 (76.1)	7351 (23.9)	
3rd birth	17 000 (15.2)	12 000 (83.1)	4386 (16.9)	
4th birth	49 000 (44.8)	38 000 (87.7)	11 000 (12.3)	
Multiple births				10.19**
Single	110 000 (96.4)	77 000 (82.4)	30 000 (17.6)	
Multiple	3994 (3.6)	2750 (79.1)	1244 (20.9)	
Child mortality estimate per co	untry			
Country	Year of survey	NMR per 1000 live births	IMR per 1000 live births	Cumulative incidence rate per 1000 person-years at risk
Burkina Faso	2003	31	81	67.9 (61.9–74.6)
	2010	28	65	44.3 (40.5–48.5)
	2014	27.3	61.4	23.9 (21.5–26.7)
Ghana	2003	43	64	30.0 (24.2–37.7)
	2008	30	50	28.5 (22.5–36.8)
	2014	29	41	15.1 (11.9–19.4)
Nigeria	2003	48	100	63.2 (55.6–72.1)
	2008	40	75	50.6 (47.7–53.7)
	2013	37	69	36.8 (34.3–39.6)
Zambia	2001–2002	37	95	70.5 (63.8–78.2)
	2007	34	70	44.7 (39.1–51.4)
	2014	24	45	26.5 (23.2–30.5)

P value notation: ***p<0.001, **p<0.01, *p<0.05.

Access to improved toilet facilities had a missing observation of 0.4%.

*% represents row percentages.

FMHCP, free maternal healthcare policy; IMR, infant mortality rate; NMR, neonatal mortality rate.

and its estimate. There were only two countries that implemented user fee reforms for maternal healthcare between 2007 and 2008. Ghana and Burkina Faso met these inclusion criteria and therefore were qualified as intervention countries. Although Zambia and Nigeria conducted DHS between 2000 and 2014, both countries did not have a universal exemption on user fees for institutional births during the study period and therefore were qualified to be used in the comparison groups. A similar study based on quasi-experimental design has provided a detailed explanation as to why Zambia, Cameroon and Nigeria could represent a valid comparison group compared with other countries in SSA in evaluating the impact of FMHCP on intermediate-term and long-term health outcomes.¹¹ Cameroon was excluded as a comparison country in this study because there was no survey conducted in 2007/2008 which represents the full policy implementation year.

Covariates assumed to be associated with child survival and included in the estimation of the propensity scores

The choice of the selected covariates in assessing risk factors of child survival was based on the analytical framework for the study of child survival in developing countries by Mosley and Chen.¹⁴ Specifically, we extracted data and performed the estimation of the propensity scores by using the following variables: household ownership of bednets, child's age and gender, mother's age at the time of the survey, mother's education level, household wealth, sex of the household head, urban or rural area of the household, birth order, multiple births and household size and household access to improved water and sanitation. We defined a household as having access to an improved water source if it has any of the following: piped water into the dwelling, yard or plot; public tap or standpipe, tube well or borehole; a protected dug well or protected spring; rainwater or bottled water. There is a direct correlation between access to an improved water source and infant survival.¹⁵ This analysis defines a household as having an improved sanitation if it has any of the following types of toilet facilities, and if this facility is not shared with another household: a flush or pour flush to piped sewer system, septic tank or pit latrine; a ventilated improved pit latrine; a pit latrine with a slab or a composting toilet. There is an inverse relationship between access to improved sanitation and infant mortality. Increasing access to improved sanitation is associated with lower levels of infant mortality.¹⁵ The estimation of the propensity scores was based on the binary logistic regression model that adjusted for the complex survey design structure of the data set (weighting, stratification and clustering).

Statistical analyses based on DHS and MIS data sets

Since the study pooled data from different surveys, the women's standard weights were denormalised. This was achieved by dividing the women's standard weight by the women survey sampling fraction, that is, the ratio of the total number of women aged 15-49 interviewed in the survey year over the total number of women aged 15-49 in the country at the time of the survey. The total number of women aged 15-49 interviewed in the survey year was obtained from the DHS data sets, while the total number of women aged 15–49 years in the country at the time of the survey was obtained from our world in data (https://ourworldindata.org/). Complex survey design characteristics (weighting, stratification and clustering) were adjusted in all the analyses. In particular, we used the sampling weights in the estimation of the propensity score model and also used the sampling weight times the Kernel weight obtained from the repeated cross-section as the weight variable in the final outcome analysis. This analytic technique has been shown to produce unbiased treatment effect estimates that are generalisable to the original survey target population.¹⁶ The Kernel function used in the weight estimation was Epanechnikov and the bandwidth selection was based on cross-validation of the means of covariates.¹⁷

To determine the impact of the policy on NMR and IMR, we performed a Propensity Score Kernel Matching with DID analysis using a modified Poisson regression model with robust standard errors. We estimated the average treatment effect (ATE) using propensity scores with Kernel matching adjustment and inverse probability of treatment weighting (IPTW). The data for this study originated from multistage complex surveys and to assess the impact of the intervention, there is a need to replicate random assignment. In experimental study design with random assignment, treatment groups (countries with FMHCP) and control groups (countries with no such policy) are similar on all background characteristics (observed and unobserved) as a consequence of the randomisation, allowing for straightforward comparison of outcomes. In contrast, in complex surveys, the intervention and comparison individuals may differ significantly on background characteristics. Thus, any difference in outcomes (NMR and IMR) between the two groups may be due to these background covariates or to the intervention itself. Matching procedures, followed by regression adjustment on the matched sample, can often be a stronger approach for estimating causal effects than regression on an unmatched sample.

The DID design is a known quasi-experimental method that is used frequently in policy evaluations to compare changes over time in a group unaffected by the policy intervention (comparison countries) with the changes over time in a group affected by the policy intervention (intervention countries) and attributes the 'DID' to the effect of the policy.¹⁹ Several sensitivity analyses were conducted to determine the robustness of our results. We tested whether the policy impact estimate is robust to the type of model specification using logit, probit and Cox proportional hazard models with robust standard errors. For the Cox model, the time-to-death with survival status as a censoring indicator was modelled. Finally, we tested whether the impact estimate is robust to different weighting procedures. First, we employed IPTW given by $w_i = \frac{T_i}{e_i} + \frac{1-T_i}{1-e_i}$ where e_k is the estimated propensity score for individual k and T_i is the treatment status indicator variable. The IPTW serves to weigh both the treated and control groups up to the full sample, in the same way, that surveys sampling weights weigh a sample up to a population.²⁰ We also applied weighting by the odds to estimate the ATE on the treated (ATT) given by $w_i = T_i + (1 - T_i) \frac{e_i}{1 - e_i}$. The DID design relies on the parallel trend assumption. This assumption stated that in the absence of the intervention (FMHCP), there would be no statistically significant difference in the trend of NMR and IMR between the intervention and the comparison countries. We relied on DHS data conducted between the years 2000 and 2008 to test this assumption. P values less than 0.05 were considered as statistically significant. Data cleaning and analysis were conducted using Stata V.15 (StataCorp).

RESULTS

Results using data from 2007 to 2014 showed that approximately 9.2% (95% CI: 8.9 to 9.5) of the 110 748 children in our sample died before reaching age 5. Within the same



Figure 1 KMSE at varying time points of FMHCP implementation. BF, Burkina Faso; FMHCP, free maternal healthcare policy; GHA, Ghana; KMSE, Kaplan–Meier survival estimate; NIG, Nigeria; ZAM, Zambia.

period, there was a statistically significant difference in the proportion of deaths between countries with FMHCP and those with no such policy (FMHCP=6.2% (95% CI: 5.9 to 6.6); no FMHCP=9.8% (95% CI: 9.5 to 10.1), Rao-Scot χ^2 test=159.6; p<0.001, table 1). The proportion of infant deaths was 6.7% (95% CI: 6.5 to 7.0). Among countries with FMHCP, the proportion of infant deaths was approximately 4.0% (95% CI: 3.6 to 4.3) compared with countries with no FMHCP where infant deaths were 7.3% (95% CI: 7.1 to 7.6) and the difference was statistically significant (Rao–Scot χ^2 test=168.4; p<0.001, table 1). The overall proportion of neonatal deaths was 3.5% (95% CI: 3.3 to 3.6). FMHCP countries recorded 0.4% (95% CI: 0.3 to 0.4) neonatal deaths compared with 3.1% (95% CI: 2.9 to 3.3) recorded by countries with no FMHCP (Rao-Scot χ^2 test=76.7, p<0.001).

NMR and IMR per 1000 live births decline between 2008 and 2014 in both FMHCP and non-FMHCP countries but the decline was steeper at all times in the FMHCP countries at various time points (figure 1).

Results on balancing and common support diagnostics of the Kernel-based matching

Balancing tests based on standardised mean difference and ratio of variances of the observed covariates between the two sets of countries (FMHCP and non-FMHCP) were conducted before and after Kernel-based matching. This was done to ascertain how the matching procedure has reduced biases in the means and variances of the observed covariates between FMHCP countries and non-FMHCP countries. The mean difference in the observed covariate between FMHCP and non-FMHCP countries reduced significantly after matching, making the two groups as similar as possible (online supplementary appendix table S1 B). The ratio of variances in the covariate between the two sets of countries was closer to 1 after matching than before matching (online supplementary appendix table S1 C). The results showed that the propensity score



Figure 2 Balancing the diagnostic test of the Kernel-based propensity score matching. FMHCP, free maternal healthcare policy.

with Kernel-based matching reduced covariate imbalance between countries with and without FMHCP. The results from the Kernel density, cumulative distribution and the box–whisker plots in figure 2 showed that matching has made FMHCP and non-FMHCP countries more similar in terms of the observed covariates, hence any change in the risk of neonatal and infant deaths could be attributed to FMHCP.

Results on the test of the parallel trend assumption

The fixed-effects model controls for all time-invariant differences between the individuals and the countrylevel factors such as differences in geographical location, so the estimated coefficients of the fixed-effects models cannot be biased because of omitted time-invariant characteristics.

The test of parallel trends showed that after controlling for baseline individual and country time-fixed effect characteristics, maternal, child and household characteristics including household ownership of bednet, both IMR and NMR did not differ between countries with FMHCP and those with no FMHCP before the implementation of FMHCP (NMR: aRR=0.91, 95% CI 0.71 to 1.16; p>0.05; table 2).

Mortality rates were declining in all of the study countries during this time period (NMR: aRR=0.88, 95% CI: 0.75 to 1.02; IMR: aRR=0.84, 95% CI: 0.76 to 0.94, p<0.05, table 2), but there was no evidence of trends being different between countries that have implemented FMHCP and comparison countries. In conclusion, the parallel trend assumption was not violated and therefore estimates from DID analyses were valid.

Impact of FMHCP on the risk of neonatal deaths

The results from the modified Poisson with DID using Propensity Score Kernel Matching showed that FMHCP is associated with 45% reduction in the risk of NMR in Ghana and Burkina Faso compared with Nigeria and Zambia (aRR=0.55, 95% CI: 0.40 to 0.76, p<0.001,

 Table 2
 Test of parallel trends assumption: risk of neonatal and infant mortality prior to free maternal healthcare policy implementation (2001–2008): modified Poisson model with robust SE on the unmatched sample

	Neonatal mortality: 20	00–2008	Infant mortality: 2000–2	2008
Covariates	uRR (95% CI)	aRR (95% CI)	uRR (95% CI)	aRR (95% CI)
Time baseline: 2008	ref	ref	ref	ref
End-line: 2014	0.86* (0.75 to 0.99)	0.88 (0.75 to 1.02)	0.83*** (0.76 to 0.91)	0.84** (0.76 to 0.94)
Intervention				
No FMHCP	ref	ref		ref
FMHCP-assumed it exited	0.85 (0.72 to 1.01)	0.89 (0.71 to 1.11)	0.94 (0.84 to 1.05)	0.92 (0.79 to 1.07)
Time*FMHCP	0.92 (0.74 to 1.15)	0.91 (0.71 to 1.16)	0.93 (0.81 to 1.08)	0.91 (0.78 to 1.08)
Sex of household head				
Male		ref		ref
Female		0.90 (0.73 to 1.11)		0.89 (0.76 to 1.03)
Mothers' current age				
<18 years		ref		ref
18–34 years		0.90 (0.78 to 1.03)		0.91 (0.82 to 1.00)
35+		2.33* (1.02 to 5.30)		2.40* (1.10 to 5.26)
Place of residence				
Urban		ref		ref
Rural		1.35*** (1.14 to 1.60)		1.29*** (1.14 to 1.45)
Household size				
1–4		ref		ref
5–7		0.48*** (0.40 to 0.57)		0.49*** (0.43 to 0.55)
8+		0.43*** (0.35 to 0.52)		0.43*** (0.37 to 0.49)
Access to improved water				
Improved		ref		ref
Unimproved		1.08 (0.92 to 1.27)		1.13* (1.01 to 1.26)
Access to an improved toilet fa	cility			
Improved, not shared		ref		ref
Improved, shared		0.81* (0.66 to 0.98)		0.76*** (0.66 to 0.87)
Unimproved		0.85 (0.72 to 1.01)		0.86* (0.77 to 0.97)
Mothers' education				
None		ref		ref
Primary		1.01 (0.84 to 1.2)		0.94 (0.83 to 1.06)
JHS		0.84 (0.69 to 1.02)		0.73*** (0.63 to 0.83)
Secondary or higher		0.93 (0.59 to 1.44)		0.55** (0.39 to 0.78)
Birth order				
1st birth		ref		ref
2nd birth		0.70** (0.57 to 0.86)		0.83* (0.71 to 0.98)
3rd birth		0.71** (0.55 to 0.9)		0.99 (0.84 to 1.17)
4th birth		1.03 (0.84 to 1.26)		1.20* (1.04 to 1.39)
Multiple births				
Single		ref		ref
Multiple		5.31*** (4.26 to 6.62)		3.70*** (3.11 to 4.40)
Household ownership of bedne	t			
No bednet		ref		ref
Bednet		0.91 (0.78 to 1.05)		0.95 (0.86 to 1.05)
Country fixed effect	Yes	Yes	Yes	Yes

P value notations: ***p<0.001, **p<0.01, *p<0.05.

With respect to Burkina Faso, 2010 demographic health survey data were used since they did not conduct any survey in 2008.

aRR, adjusted relative risk; FMHCP, free maternal healthcare policy; JHS, junior high school; uRR, unadjusted relative risk.

table 3). Sensitivity analyses based on different outcome model specification showed similar results (table 3).

Impact of FMHCP on the risk of infant deaths

IMR has reduced significantly in both Ghana and Burkina Faso by approximately 54% after full implementation of FMHCP compared with Nigeria and Zambia (aRR=0.46, 95% CI: 0.36 to 0.59, p<0.001; table 4). The series of sensitivity analysis that was conducted showed a similar impact on FMHCP(table 4). The analysis was adjusted for sex of the household head, mothers' current age, mothers' educational level, place of residence, wealth quintile, access to improved water and sanitation, birth order, multiple births and household ownership of bednet and country fixed effect.

DISCUSSION

This study quantified the contribution of FMHCP implementation in Ghana and Burkina Faso in the reduction of NMR and IMR. Child mortality within the implementation period in these two countries was compared with mortality in Nigeria and Zambia which do not have a significant major health financing reform in the period under consideration. It remains among the few studies to have compared the effectiveness of FMHCP in the four SSA countries using the more rigorous matching procedure with DID. Our impact evaluation found that the implementation of FMHCP led to a substantial reduction in both neonatal and infant mortality. This finding is consistent with what has been reported previously in the literature based on similar analytic technique.²¹ Although all the four countries studied did not attain the MDG 4, Ghana and Burkina Faso have seen a tremendous decline in the trend of NMR and IMR over the years. FMHCP was associated with substantial statistically significant reductions in IMR and NMR when these estimates were compared between Zambia and Nigeria.

It is estimated that the effective implementation of key maternal and child healthcare interventions could prevent up to 70% of neonatal deaths globally.^{22 23} The advantages of increasing access to facility delivery, prenatal and postnatal care through FMHCP are well documented in the literature.^{12 24} FMHCP contributes greatly to increased coverage of routine immunisation as women who visit and deliver in recommended health facilities were more likely to benefit from early immunisation. The policy also promotes early and accurate diagnosis of childhood illnesses after delivery and within the postpartum period. Education on malaria preventive measures after delivery and the administration of intermittent preventive treatment for pregnancy during antenatal are a few of the benefits women derived from the policy. The FMHCP is associated with high ANC attendance and institutional delivery by skilled attendants (midwives, nurses and doctors) at the time of delivery which consequently reduced neonatal deaths and to a larger extent infant mortality.^{25 26} Increasing access to the skilled birth attendant and emergency obstetric care is accepted as

the most crucial intervention for reducing maternal and newborn deaths. $^{\rm 27}$

Strengths and limitations

This study has several strengths and some limitations. The advantages of using DHS as our primary source data have been well documented.²⁸ Paramount among these several advantages include high response rates, national coverage, high-quality interviewer training, standardised data collection procedures across countries and consistent content over time, allowing comparability across populations cross-sectionally and over time. The use of DID models with Propensity Score Kernel Matching is seen as strong non-experimental study design options when randomisation is not feasible and provides more robust inference.¹⁹

The limitation of this study originates from the fact that the DID analytical technique is generally less robust than the randomised design even though the study established that the parallel trend assumption was not violated. Although Kernel matching maximises the chance of matching control to a treated individual, observations outside the range of common support are still discarded which could potentially reduce the sample size.

We highlighted the fact that our study could still suffer from the omission of important time-varying unobserved characteristics such as total annual health expenditure could bias our study results if the omitted variables affected Ghana, Burkina Faso and comparison countries in different ways. The reason is that DID attributes to the FMHCP policy intervention differences in mortality trends between the Ghana and comparison countries that occur from the time intervention begins (2008). If any other factor is present which affect the difference in trends between the two groups differently, then the estimate from DID could be biased. In particular, health funding sources like the US President Malaria Initiative (PMI), President's Emergency Plan for AIDS Relief (PEPFAR) and the Global Fund for HIV, tuberculosis and malaria are few of the foreign aid that could have an impact on child mortality.²⁹ For instance, Ghana and Zambia received funding support from PMI in 2008 but Burkina Faso has never benefited from PMI and Nigeria which only received funding from the PMI in 2011. Three out of the four countries studied continue to be benefited from PEPFAR but received the support at different times (Ghana; 2007, Burkina Faso; not at all, Zambia and Nigeria in 2004). The countries Ghana and Zambia still remain among the countries studied that have had the benefits of the US PMI since 2008 which also coincides with the year in which FMHCP policy became fully operational. The observed differentials among the four countries relative to foreign aid could impact on child mortality differently and bias the results.

With regards to Zambia and Nigeria, these two countries might not have a nationwide FMHCP but it is possible that there may be country-specific interventions put in place to curb the menace of child mortality.

	Sensitivity analysis based	on different model specific	cation				
	Modified Poisson model: clustering, weighti adjusted	ing and stratification were	Cox-proportional hazard model: clustering, weighting and stratification were adjusted	Logistic regression model: clustering, weighting and stratification were adjusted	Probit regression model: clustering, weighting and stratification were adjusted	Modified Poisson model: clustering stratification were	, weighting and e adjusted
	No Kernel matching aRR (95% Ci)	With Kernel matching based on PS	With Kernel matching based on PS aHR (05% CI)	With Kernel matching based on PS	With Kernel matching based on PS 8 (95% CI)	ATET weighting and PS-IPTW	ATE weighting and PS
Time							
Baseline: 2008	ref	ref	ref	ref	ref	ref	ref
End-line: 2014	0.92 (0.82 to 1.03)	0.63* (0.42 to 0.96)	0.62* (0.42 to 0.91)	0.59 (0.33 to 1.07)	-0.35* (-0.69 to 0.01)	0.61** (0.46 to 0.81)	0.67** (0.51 to 0.86)
Intervention							
No FMHCP	ref	ref	ref	ref	ref	ref	ref
FMHCP	0.66*** (0.53 to 0.83)	0.94 (0.74 to 1.19)	0.96 (0.77 to 1.19)	0.95 (0.62 to 1.47)	-0.04 (-0.28 to 0.2)	-0.81 (0.78 to 1.11)	0.93 (0.82 to 1.07)
Time*FMHCP	0.56*** (0.43 to 0.73)	0.55*** (0.40 to 0.76)	0.55*** (0.40 to 0.74)	0.44* (0.22 to 0.88)	-0.41* (-0.79 to 0.02)	0.57*** (0.42 to 0.77)	0.71 (0.46 to 1.08)
Sex of household head							
Male	ref	ref	ref	ref	ref	ref	ref
Female	0.85* (0.73 to 0.99)	0.73* (0.55 to 0.97)	0.81 (0.63 to 1.04)	1.08 (0.67 to 1.74)	0.02 (-0.24 to 0.29)	-1.99* (0.65 to 1)	0.78* (0.63, 0.95)
Mothers' current age							
<18 years	ref	ref	ref	ref	ref	ref	ref
18-34 years	0.43*** (0.35 to 0.54)	0.51** (0.35 to 0.75)	1.01 (0.71 to 1.43)	2.69** (1.40 to 5.18)	0.59** (0.21 to 0.98)	3.99*** (0.39 to 0.72)	0.53*** (0.39 to 0.71)
35+	0.46*** (0.36 to 0.59)	0.55** (0.35 to 0.86)	1.29 (0.85 to 1.95)	6.7*** (3.12 to 14.38)	1.11*** (0.67 to 1.56)	-3.17** (0.39 to 0.8)	0.5*** (0.35 to 0.72)
Place of residence							
Urban	ref	ref	ref	ref	ref	ref	ref
Rural	1.24** (1.08 to 1.42)	1.23 (0.87 to 1.74)	1.2 (0.88 to 1.64)	1.04 (0.65 to 1.65)	0.05 (-0.22 to 0.32)	1.94 (1.00 to 1.96)	1.21 (0.84 to 1.74)
Wealth quintile							
Poorest	ref	ref	ref	ref	ref	ref	ref
Poorer	1.01 (0.89 to 1.15)	1.04 (0.86 to 1.27)	1.03 (0.86 to 1.24)	0.81 (0.53 to 1.23)	-0.1 (-0.34 to 0.14)	0.72 (0.89 to 1.28)	0.98 (0.79 to 1.22)
Middle	0.86 (0.74 to 1.01)	0.93 (0.73 to 1.18)	0.96 (0.77 to 1.19)	0.93 (0.56 to 1.54)	-0.03 (-0.32 to 0.27)	0.2 (0.82 to 1.27)	0.89 (0.69 to 1.16)
Richer	0.91 (0.75 to 1.10)	0.83 (0.61 to 1.13)	0.87 (0.65 to 1.16)	0.88 (0.43 to 1.79)	-0.06 (-0.47 to 0.34)	-0.64 (0.70 to 1.20)	0.86 (0.61 to 1.2)
Richest	0.86 (0.68 to 1.11)	0.95 (0.56 to 1.61)	1.01 (0.62 to 1.62)	1.04 (0.46 to 2.34)	0.05 (-0.41 to 0.52)	-0.87 (0.60 to 1.22)	0.88 (0.55 to 1.42)
Household size							
							Continued

Open access

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Table 3 Continued	-						
	Sensitivity analysis based	on different model specific	ation				
	Modified Poisson model: clustering, weighti adjusted	ing and stratification were	Cox-proportional hazard model: clustering, weighting and stratification were adjusted	Logistic regression model: clustering, weighting and stratification were adjusted	Probit regression model: clustering, weighting and stratification were adjusted	Modified Poisson model: clustering, stratification were	weighting and adjusted
	No Kernel matching	With Kernel matching based on PS	With Kernel matching based on PS	With Kernel matching based on PS	With Kernel matching based on PS	ATET weighting and PS-IPTW	ATE weighting and PS
	aRR (95% CI)	aRR (95% CI)	aHR (95% CI)	aOR (95% CI)	β (95% CI)	aRR (95% CI)	aRR (95% CI)
1-4	ref	ref	ref	ref	ref	ref	ref
5–7	0.42*** (0.37 to 0.49)	0.46*** (0.38 to 0.56)	0.55*** (0.45 to 0.65)	0.49** (0.31 to 0.78)	-0.41** (-0.67 to 0.15)	8.18*** (0.38 to 0.55)	0.42*** (0.34 to 0.51)
8+	0.35*** (0.3 to 0.4)	0.41*** (0.33 to 0.52)	0.48*** (0.39 to 0.59)	0.33*** (0.22 to 0.49)	-0.64*** (-0.87 to 0.41)	8.67*** (0.32 to 0.49)	0.38*** (0.31 to 0.46)
Access to improved wate							
Improved	ref	ref	ref	ref	ref	ref	ref
Unimproved	1.06 (0.93 to 1.21)	1.06 (0.87 to 1.3)	1.06 (0.89 to 1.27)	1.28 (0.88 to 1.88)	0.15 (-0.07 to 0.36)	0.02 (0.82 to 1.22)	1.02 (0.82 to 1.28)
Access to an improved to	vilet facility						
Improved, not shared	ref	ref	ref	ref	ref	ref	ref
Improved, shared	0.84* (0.72 to 0.97)	0.97 (0.78 to 1.22)	0.97 (0.79 to 1.18)	1.08 (0.72 to 1.63)	0.04 (-0.20 to 0.28)	-0.43 (0.78 to 1.17)	0.90 (0.70 to 1.16)
Unimproved	0.84** 0.75 to 0.94)	0.92 (0.76 to 1.11)	0.92 (0.78 to 1.10)	1.10 (0.76 to 1.59)	0.05 (-0.15 to 0.26)	–1.00 (0.77 to 1.09)	0.86 (0.72 to 1.04)
Mothers' education							
None	ref	ref	ref	ref	ref	ref	ref
Primary	1.05 (0.93 to 1.20)	0.98 (0.78 to 1.23)	0.97 (0.78 to 1.20)	0.78 (0.52 to 1.18)	-0.14 (-0.38 to 0.10)	-0.80 (0.76 to 1.12)	0.99 (0.79 to 1.25)
SHL	0.94 (0.80 to 1.10)	0.93 (0.69 to 1.25)	0.86 (0.65 to 1.13)	0.62 (0.35 to 1.10)	-0.25 (-0.57 to 0.06)	-1.01 (0.72 to 1.11)	0.96 (0.80 to 1.16)
Secondary or higher	0.75 (0.55 to 1.03)	0.81 (0.46 to 1.41)	0.71 (0.42 to 1.21)	0.20*** (0.08 to 0.47)	-0.92*** (-1.44 to 0.41)	-1.37 (0.45 to 1.15)	0.79 (0.43 to 1.46)
Birth order							
1st birth	ref	ref	ref	ref	ref	ref	ref
2nd birth	0.94 (0.82 to 1.09)	0.69** (0.54 to 0.89)	0.62*** (0.49 to 0.77)	0.52* (0.30 to 0.89)	-0.38* (-0.67 to 0.08)	-2.89** (0.61 to 0.91)	0.74* (0.59 to 0.93)
3rd birth	0.98 (0.82 to 1.18)	0.89 (0.66 to 1.2)	0.73* (0.56 to 0.97)	0.60 (0.34 to 1.07)	-0.29 (-0.61 to 0.03)	-0.24 (0.75 to 1.25)	0.88 (0.67 to 1.15)
4th birth	1.32** (1.12 to 1.56)	1.18 (0.90 to 1.56)	0.87 (0.67 to 1.12)	0.46** (0.26 to 0.81)	-0.44** (-0.76 to 0.12)	2.21* (1.03 to 1.58)	1.29* (1.00 to 1.65)
Multiple births							
Single							
Multiple	5.84*** (4.97 to 6.86)	5.58*** (4.25 to 7.32)	4.73*** (3.78 to 5.92)	6.17*** (2.47 to 15.41)	0.96*** (0.52 to 1.4)	15.61*** (4.74 to 7.41)	5.58*** (4.37 to 7.12)
							Continued

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Table 3 Continue	pe						
	Sensitivity analysis based	on different model specific	ation				
	Modified Poisson model: clustering, weightir adjusted	ng and stratification were	Cox-proportional hazard model: clustering, weighting and stratification were adjusted	Logistic regression model: clustering, weighting and stratification were adjusted	Probit regression model: clustering, weighting and stratification were adjusted	Modified Poisson model: clustering stratification wer	, weighting and e adjusted
	No Kernel matching	With Kernel matching based on PS	With Kernel matching based on PS	With Kernel matching based on PS	With Kernel matching based on PS	ATET weighting and PS-IPTW	ATE weighting and PS
	aRR (95% CI)	aRR (95% CI)	aHR (95% CI)	aOR (95% CI)	β (95% CI)	aRR (95% CI)	aRR (95% CI)
Household ownership c	of bednet						
No bednet							
Bednet	0.98 (0.88 to 1.09)	0.96 (0.82 to 1.12)	0.96 (0.83 to 1.11)	1.01 (0.74 to 1.37)	0.01 (-0.16 to 0.19)	-1.04 (0.79 to 1.07)	0.93 (0.78 to 1.1)
Country fixed effect	Adjusted	Adjusted	Adjusted	Adjusted	Adjusted	Adjusted	Adjusted
P value notations: ***p<0.0 aHR, adjusted hazard ratio weighting; PS, propensity ;	01, **p<0.01, *p<0.05. ; aOR, adjusted odds ratio; aRR, adj scores; ref, reference category.	usted relative risk; ATE, average	treatment effect; ATET, average	e treatment effect on the treatec	;; FMHCP, free maternal healthcare pc	olicy; IPTW, inverse pr	obability of treatment

Even among the intervention countries, there may be other specific interventions that are tailored towards child mortality but were not controlled in the current study. For instance, the 'Rapid Scale-Up' programme in Burkina Faso has a component that focuses on integrated community case management and this policy has been found to reduce neonatal mortality by 6.2%.³⁰ Other interventions such as user fee exemption and mass radio campaigns have all been found to be associated with an increase in the healthcare utilisation among children under 5 in Burkina Faso which could have a direct positive impact in reducing NMR.^{31 32} In addition to the aforementioned interventions, it is worth emphasising that both Ghana and Burkina Faso receive support from the Global Fund in the fight against malaria, tuberculosis and HIV since 2003 and this might have contributed to why Burkina Faso and Ghana might be doing better in terms of reducing IMR and NMR. Despite the fact that our impact estimate of the policy may be imprecise and should be interpreted cautiously, we emphasised that the introduction of the FMHCP is associated with the reduction in both NMR and IMR which is an encouraging finding and an important contribution to the literature on the colossal benefits of FMHCP. DID still remains one of the robust quasi-experimental design to evaluate the impact of health intervention using cross-sectional timeseries data as it was the case in this study.

Policy implications

The findings from the study provide imperative evidence of an accelerated decline in child mortality rates after the introduction of FMHCP in the two West African countries. The additional investments in health tailored towards FMHCP implementation have yielded positive impacts. The implementation of the policy has reduced the financial burden associated with antenatal and postnatal care attendance and institutional delivery. Future studies should explore whether the investments made through FMHCP have spillover effects beyond the usual benefits associated with the policy, such as women empowerment, higher investment in the private sector, higher school attainment and increase in employment rate which might, in turn, lead to greater economic development. As the population of women keeps increasing geometrically in SSA, Governments should consider an alternative source of financing to sustain the policy.

CONCLUSION

The motivation of this study is to obtain more reliable evidence of how the implementation of the FMHCP in certain countries in the SSA has reduced child mortality compared with countries in the subregion with no such national policy. Our findings highlight the importance of FMHCP implementation in reducing the risk of neonatal and infant mortalities. We recommended that a similar policy should be implemented in other lower-income and

	Sensitivity analysis bé	ased on different model sp	secification				
	Modified Poisson model: clustering, we were adjusted	jghting and stratification	Cox-proportional hazard model: clustering, weighting and stratification were adjusted	Logistic regression model: clustering, weighting and stratification were adjusted	Probit regression model: clustering, weighting and stratification were adjusted	Modified Poisson model: clustering, weig were adjusted	phting and stratification
	No Kernel matching with PSM	With Kernel matching with PSM	With Kernel matching PSM	With Kernel matching with PSM	With Kernel matching with PSM	ATET weighting using PSM-IPTW	ATE weighting using PSM
	aRR (95% CI)	aRR (95% CI)	aHR (95% CI)	aOR (95% CI)	β (95% CI)	aRR (95% CI)	aRR (95% CI)
Time							
Baseline: 2008	ref	ref	ref	ref	ref	ref	ref
End-line: 2014	0.79*** (0.73 to 0.86)	0.62*** (0.48 to 0.8)	0.62*** (0.49 to 0.78)	0.45*** (0.35 to 0.58)	-0.45*** (-0.59 to 0.32)	0.64*** (0.53 to 0.78)	0.77** (0.64 to 0.91)
Intervention							
No FMHCP	ref	ref	ref	ref	ref	ref	ref
FMHCP	0.72*** (0.62 to 0.84)	0.87 (0.74 to 1.03)	0.89 (0.77 to 1.04)	0.86 (0.72 to 1.02)	-0.09 (-0.19 to 0.01)	0.84* (0.74 to 0.96)	0.83*** (0.76 to 0.92)
Time*FMHCP	0.49*** (0.39 to 0.61)	0.46*** (0.36 to 0.59)	0.45*** (0.35 to 0.57)	0.43*** (0.33 to 0.56)	-0.49*** (-0.64 to 0.34)	0.48*** (0.37 to 0.6)	0.55** (0.39 to 0.77)
Sex of household !	lead						
Male	ref	ref	ref	ref	ref	ref	ref
Female	0.88* (0.78 to 0.98)	0.75** (0.61 to 0.92)	0.81* (0.68 to 0.98)	0.96 (0.79 to 1.17)	-0.02 (-0.13 to 0.09)	0.87 (0.74 to 1.02)	0.88 (0.75 to 1.02)
Mothers' current a	ge						
<18 years	ref	ref	ref	ref	ref	ref	ref
18-34 years	0.49*** (0.41 to 0.58)	0.49*** (0.37 to 0.66)	0.86 (0.65 to 1.14)	3.1*** (2.2 to 4.38)	0.63*** (0.44 to 0.81)	0.49*** (0.38 to 0.63)	0.59*** (0.46 to 0.75)
35+	0.49*** (0.4 to 0.59)	0.47*** (0.33 to 0.66)	0.95 (0.69 to 1.31)	5.74*** (3.92 to 8.4)	0.96*** (0.75 to 1.17)	0.46*** (0.35 to 0.61)	0.52*** (0.39 to 0.68)
Place of residence							
Urban	ref	ref	ref	ref	ref	ref	ref
Rural	1.16** (1.05 to 1.29)	1.2 (0.96 to 1.5)	1.19 (0.97 to 1.46)	1.18 (0.96 to 1.45)	0.09 (-0.03 to 0.2)	1.14 (0.96 to 1.36)	1.13 (0.89 to 1.45)
Wealth quintile							
Poorest	ref	ref	ref	ref	ref	ref	ref
Poorer	1.02 (0.92 to 1.12)	1.01 (0.87 to 1.17)	1 (0.87 to 1.15)	0.97 (0.81 to 1.16)	-0.02 (-0.12 to 0.08)	1.04 (0.91 to 1.18)	0.99 (0.86 to 1.15)
Middle	0.88* (0.79 to 0.98)	0.92 (0.77 to 1.11)	0.95 (0.8 to 1.12)	0.99 (0.8 to 1.23)	-0.01 (-0.13 to 0.11)	0.92 (0.79 to 1.07)	0.93 (0.78 to 1.11)
Richer	0.86* (0.75 to 0.99)	0.91 (0.7 to 1.19)	0.95 (0.74 to 1.21)	0.93 (0.68 to 1.28)	-0.03 (-0.2 to 0.14)	0.89 (0.72 to 1.09)	0.8* (0.64 to 1)
Richest	0.69*** (0.57 to 0.82)	0.78 (0.54 to 1.12)	0.81 (0.58 to 1.13)	0.8 (0.57 to 1.11)	-0.14 (-0.32 to 0.05)	0.68** (0.52 to 0.89)	0.73 (0.5 to 1.06)
Household size							
1-4	ref	ref	ref	ref	ref	ref	ref
5-7	0.43*** (0.39 to 0.47)	0.45*** (0.38 to 0.52)	0.52*** (0.45 to 0.59)	0.54*** (0.45 to 0.65)	-0.34*** (-0.44 to 0.24)	0.46*** (0.4 to 0.52)	0.43*** (0.37 to 0.5)
8+	0.35*** (0.32 to 0.39)	0.4*** (0.34 to 0.48)	0.46*** (0.39 to 0.54)	0.43*** (0.35 to 0.52)	-0.45*** (-0.55 to 0.34)	0.42*** (0.36 to 0.49)	0.38*** (0.33 to 0.45)
Access to improve	d water						:
							Continued

Table 4 Contir	ned						
	Sensitivity analysis ba	sed on different model sp	becification				
	Modified Poisson model: clustering, wei were adjusted	ghting and stratification	Cox-proportional hazard model: clustering, weighting and stratification were adjusted	Logistic regression model: clustering, weighting and stratification were adjusted	Probit regression model: clustering, weighting and stratification were adjusted	Modified Poisson model: clustering, weig were adjusted	hting and stratification
	No Kernel matching with PSM	With Kernel matching with PSM	With Kernel matching PSM	With Kernel matching with PSM	With Kernel matching with PSM	ATET weighting using PSM-IPTW	ATE weighting using PSM
	aRR (95% CI)	aRR (95% CI)	aHR (95% CI)	aOR (95% CI)	β (95% CI)	aRR (95% CI)	aRR (95% CI)
Improved	ref	ref	ref	ref	ref	ref	ref
Unimproved	1.05 (0.96 to 1.16)	1.03 (0.89 to 1.19)	1.02 (0.89 to 1.17)	1.02 (0.88 to 1.17)	0 (-0.07 to 0.08)	0.99 (0.86 to 1.15)	1.05 (0.9 to 1.22)
Access to an impro	ved toilet facility						
Improved, not shared	ref	ref	ref	ref	ref	ref	ref
Improved, shared	1 0.87* (0.78 to 0.97)	0.94 (0.79 to 1.1)	0.94 (0.81 to 1.09)	0.82* (0.69 to 0.97)	-0.11* (-0.2 to 0.01)	0.97 (0.83 to 1.13)	0.94 (0.78 to 1.13)
Unimproved	0.88** (0.81 to 0.96)	0.92 (0.8 to 1.05)	0.93 (0.82 to 1.05)	0.88 (0.75 to 1.02)	-0.07 (-0.15 to 0.02)	0.93 (0.82 to 1.05)	0.92 (0.79 to 1.06)
Mothers' education							
None	ref	ref	ref	ref	ref	ref	ref
Primary	0.9* (0.82 to 0.99)	0.82* (0.7 to 0.97)	0.82** (0.71 to 0.95)	0.69*** (0.58 to 0.83)	-0.21*** (-0.31 to 0.11)	0.81** (0.71 to 0.93)	0.89 (0.76 to 1.03)
SHL	0.88* (0.78 to 0.98)	0.83 (0.67 to 1.02)	0.77** (0.64 to 0.94)	0.54*** (0.43 to 0.67)	-0.35*** (-0.47 to 0.24)	0.82* (0.69 to 0.97)	0.87 (0.73 to 1.03)
Secondary or higher	0.7** (0.56 to 0.89)	0.78 (0.53 to 1.17)	0.71 (0.48 to 1.04)	0.38*** (0.24 to 0.58)	-0.55*** (-0.78 to 0.31)	0.78 (0.53 to 1.14)	0.69 (0.43 to 1.11)
Birth order							
1st birth	ref	ref	ref	ref	ref	ref	ref
2nd birth	1.03 (0.93 to 1.15)	0.86 (0.72 to 1.03)	0.77** (0.65 to 0.91)	0.6*** (0.5 to 0.73)	-0.27*** (-0.38 to 0.17)	0.94 (0.82 to 1.08)	0.84* (0.71 to 1)
3rd birth	1.2** (1.05 to 1.37)	1.13 (0.91 to 1.4)	0.96 (0.79 to 1.17)	0.65*** (0.52 to 0.81)	-0.24*** (-0.36 to 0.12)	1.19 (1 to 1.43)	1.08 (0.88 to 1.34)
4th birth	1.59*** (1.42 to 1.79)	1.63*** (1.34 to 1.98)	1.25* (1.05 to 1.5)	0.7** (0.56 to 0.86)	-0.2** (-0.32 to 0.08)	1.66*** (1.42 to 1.93)	1.51*** (1.23 to 1.86)
Multiple births							
Single	ref	ref	ref	ref	ref	ref	ref
Multiple	4.37*** (3.86 to 4.95)	3.95*** (3.16 to 4.95)	3.57*** (2.96 to 4.3)	3.59*** (2.75 to 4.67)	0.72*** (0.57 to 0.87)	4.24*** (3.58 to 5.03)	4.43*** (3.71 to 5.29)
Household owners	nip of bednet						
No bednet	ref	ref	ref	ref	ref	ref	ref
Bednet	1.06 (0.98 to 1.14)	0.98 (0.88 to 1.09)	0.97 (0.88 to 1.08)	0.99 (0.88 to 1.12)	-0.01 (-0.08 to 0.06)	0.96 (0.86 to 1.06)	0.94 (0.85 to 1.05)
Country fixed effec	: Adjusted	Adjusted	Adjusted	Adjusted	Adjusted	Adjusted	Adjusted
P value notations: *** aHR, adjusted hazaπ probability of treatme	p<0.001, **p<0.01, *p<0.05. I ratic; aOR, adjusted odds int weighting; PS, propensit	ratio; aRR, adjusted relative r :y scores.	isk; ATE, average treatment	effect; ATET, average treatn	nent effect on the treated; FMHCP, f	ree maternal healthcare poli	cy; IPTW, inverse

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middle-income SSA countries to reduce the prevalence of neonatal and infant deaths.

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Acknowledgements The authors duly acknowledge the Demographic and Health Survey Program funded by the U.S. Agency for International Development (USAID), other donors and Ghana Statistical Service for making their data available to the public. We do acknowledge all mothers who were interviewed.

Contributors DD, KwA and PN conceived and designed the study. Data management and data cleaning were done by DD, GI and KoA. Statistical methods were drafted by DD, SB and GI. DD, PN, GI, KwA, KoA, SB and AY revised the draft critically. All authors have read and approved the final manuscript.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not required.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available in a public, open access repository. All data sets are public data and are freely available upon request.

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