

Automated phone call and text reminders for childhood immunisations (PRIMM): a randomised controlled trial in Nigeria

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ABSTRACT

Background Sub-Saharan Africa has high under-5 mortality and low childhood immunisation rates. Vaccine-preventable diseases cause one-third of under-5 deaths. Text messaging reminders improve immunisation completion in urban but not rural settings in sub-Saharan Africa. Low adult literacy may account for this difference. The feasibility and impact of combined automated voice and text reminders on immunisation completion in rural sub-Saharan Africa is unknown.

Methods We randomised parturient women at the Mother and Child Hospitals Ondo State, Nigeria, owning a mobile phone and planning for child immunisation at these study sites to receive automated call and text immunisation reminders or standard care. We assessed the completion of the third pentavalent vaccine (Penta-3) at 18 weeks of age, immunisation completion at 12 months and within 1 week of recommended dates. We assessed selected demographic characteristics associated with completing immunisations at 12 months using a generalised binomial linear model with 'log' link function. Feasibility was assessed as proportion of reminders received.

Results Each group had 300 mother–baby dyads with similar demographic characteristics. At 18 weeks, 257 (86%) and 244 (81%) (risk ratio (RR) 1.05, 95% CI 0.98 to 1.13; $p=0.15$) in the intervention and control groups received Penta-3 vaccine. At 12 months, 220 (74%) and 196 (66%) (RR 1.12, 95% CI 1.01 to 1.25; $p=0.04$) in the intervention and control groups received the measles vaccine. Infants in the intervention group were more likely to receive Penta-3 (84% vs 78%, RR 1.09, 95% CI 1.01 to 1.17; $p=0.04$), measles (73% vs 65%, RR 1.13, 95% CI 1.02 to 1.26; $p=0.02$) and all scheduled immunisations collectively (57% vs 47%, RR 1.13, 95% CI 1.02 to 1.26; $p=0.01$) within 1 week of the recommended date. No demographic character predicted immunisation completion. In the intervention group, 92% and 86% reported receiving a verification reminder and at least one reminder during the study period, respectively.

Conclusion Paired automated call and text reminders significantly improved immunisation completion and timeliness.

Trial registration number NCT02819895.

Key questions

What is already known?

- In sub-Saharan Africa, text message reminders alone improve childhood immunisation uptake in urban areas but not in rural areas where literacy levels are lower.
- Low adult literacy rates in sub-Saharan Africa may limit the use of stand-alone text messaging reminder systems.

What are the new findings?

- In a semi-rural setting in Nigeria, paired automated voice call and text immunisation reminders significantly improved the completion and timely receipt of immunisations.
- Cost of transportation, distance to and length of time spent at the immunisation clinics were additional significant barriers to immunisation reported in this setting.

What do the new findings imply?

- Our results add to the body of literature on the impact of phone reminders on immunisation completion and inform research on innovative solutions capable of improving access to and efficiency of immunisation clinics.

INTRODUCTION

Vaccine-preventable diseases are a leading cause of under-5 mortality.¹ Sub-Saharan Africa (SSA) and Southeast Asia—regions with poor immunisation uptake and completion rates—carry 80% of the global under-5 mortality burden.^{1,2} These regions contribute significantly to the stalled global immunisation coverage.³ Nigeria has the highest number of underimmunised and unimmunised children worldwide.^{3,4} A 2017 Nigerian National Immunization Coverage Survey estimated that 37% and 40% of children in Nigeria aged 12–24 months were underimmunised and unimmunised, respectively.⁵ A

fully immunised child is defined as one who received one Bacille Calmette–Guerin (BCG), three pentavalent (ie, diphtheria, tetanus, pertussis, *Haemophilus influenzae* type B and hepatitis B), one yellow fever and measles vaccine within the first year of life. The third dose of the pentavalent vaccine (Penta-3) is the global benchmark for vaccine reporting. To meet the 2020 Global Vaccine Action Plan targets, interventions that are innovative, cost-effective, region and country specific are needed.

In high-income countries, immunisation reminders through automated or real-time text and or calls improve immunisation uptake and completion.⁶ According to the Nigerian National Immunization Coverage Survey report, lack of awareness of recommendations for a child's immunisation is the biggest reason for not completing the immunisation series—accounting for 42% of the reported barriers.⁵ With high mobile phone penetration in Nigeria and other SSA countries, phone reminders have the potential to help overcome this barrier.

Evidence from randomised controlled trials (RCTs) conducted in urban areas of Zimbabwe, Nigeria and Burkina Faso suggest text reminders alone improve immunisation completion and timeliness.^{7–9} However, a recent RCT from rural Kenya contradicts these findings.¹⁰ In this study, text message alone did not improve immunisation completion, but improved timeliness of vaccine receipt. Only when text reminders were paired with a higher monetary incentive did completion rates improve.¹⁰ Text reminders assume the recipient is literate enough to read and understand the message. However, 38% of African adults (some 153 million) are illiterate—two-thirds of these being women.¹¹ Adult literacy is often lower in rural compared with urban areas.^{12 13} This may account for the difference in effectiveness—an indication that text reminders alone are not appropriate in this setting. In a study from a rural community in Nigeria, the use of manual phone call immunisation reminders significantly improved immunisation completion—a 41% increase from controls.¹⁴ However, this method is time and human resource dependent and unlikely to garner national application. With literacy being a limitation in some areas, automated audible reminders in the native languages may provide added benefit to text reminders.

We conducted the Phone Reminder for Immunisation (PRIMM) trial to test the feasibility and effect of pairing automated text and call reminders on immunisation completion in a semi-rural setting in Nigeria. We hypothesised that the Penta-3 vaccine and the completion of the childhood immunisation series assessed at 12 months of age would significantly improve with this intervention.

METHODS

Study design

A two-arm parallel RCT was conducted at the Mother and Child Hospital Ondo Town (MCH-Ondo) and Akure (MCH-Akure), in Ondo State, Nigeria. The conduct, analysis and reporting of results are in accordance with

the Consolidated Standards of Reporting Trials guidelines for reporting parallel group randomised trials.¹⁵

The Nigerian government provides routine childhood immunisation at no cost to recipients. Government-run immunisation clinics are locations where vaccines are routinely received. Dictated by the national programme on immunisation,¹⁶ the routine immunisation schedule in Ondo State, Nigeria is as follows:

- Birth: BCG, hepatitis B virus and oral polio vaccine (OPV-0).
- Six weeks: Penta-1, OPV-1, pneumococcal conjugate vaccine (PCV-1) and rotavirus vaccine (Rota-1).
- Ten weeks: Penta-2, OPV-2, Rota-2 and PCV-2.
- Fourteen weeks: Penta-3, OPV-3, PCV-3 and inactivated polio vaccine (IPV).
- Six months: vitamin A.
- Nine months: measles and yellow fever vaccine.

Immunisation clinic record-book audits were planned to assess vaccination uptake. However, the study was impacted by a hospital workers' strike lasting 6 weeks (2 February to 15 March 2017). The strike interrupted enrolment, immunisation receipt and monitoring. During this period, parents sought other government and private clinics for immunisations. We therefore amended our study protocol to include phone audits for reporting of immunisation uptake between weeks 18 and 30 of each participant's study enrolment period. This was to limit misclassification of the immunisation endpoint. We incorporated the phone call immunisation audit into the planned mid-study survey. The applicable ethical review bodies approved the amendment.

Study participants and setting

Parturient women and their healthy newborn infants delivered at MCH-Ondo and Akure were eligible for enrolment. We included mothers of healthy newborn babies, who owned a mobile phone and planned to attend the MCH immunisation clinics. We excluded mothers of ill newborns, multiple births and those without mobile phones.

The MCHs are state-run facilities. They provide free healthcare services to pregnant women and children under age 5, with most patients being middle-income and low-income families. Each hospital runs its own immunisation clinic. Ondo State is in the southwest region of Nigeria. The main local language is Yoruba. The projected 2016 population size from the 2006 national census for Ondo Town and Akure are 389 900 and 486 300, respectively.¹⁷ The primary occupations of citizens include farming, artisanship, trading and public service.¹⁷

Study intervention

We developed a customised Windows software application (app) designed to send automated voice call text and email immunisation reminders. We integrated a secure cloud communications platform, called Twilio, into the app. Messaging and voice were sent by Twilio through the app. Date of birth of the newborn and the

phone number of the mother and father, when provided, were inputted into the app. The immunisation reminders were autocalculated from date of birth of the child and tailored to the local immunisation schedule. At enrolment, the registered phone number(s) received a verification message. Thereafter, reminders were sent 2 days and the day before the scheduled date of the Penta-1, 2, 3 and measles immunisations at 08:00. Eight sets of reminders were sent to each participant. The delivery of text and call occurred at the same time. Voicemail service was not available during the study period.

The automated text message reminder was in English. The text reminder read, *'Reminder from MCH—Your baby's next immunisation visit is in 2 days (or 1 day as appropriate). Immunisation protects your child against killer diseases. Please bring your baby for this visit.'* The automated call reminder was in English and Yoruba. The duration of the call was 50 s, had a 5 s delay before starting and expressed the same message as the text. It cost US\$0.0075 to send a text and US\$0.015/min for an automated call. There was no cost to the recipient. Whether study participants received or read the text message and whether participants listened to audio messages in its entirety could not accurately be determined from our telecommunications platform.

Study procedures

A research assistant at each site assessed mothers daily in the postpartum ward for eligibility. We systematically recorded the number of screened women and the reason for exclusion; however, due to a clerical error at MCH-Ondo, the exact numbers and reasons are unavailable.

In developing the study protocol, we took into consideration that the shortest interval between the pentavalent vaccines is 3 weeks. In Nigeria, the immunisation schedule allows for only 4 weeks between Penta-1 and 2, and between Penta-2 and 3.¹⁸ We anticipated a potential stacking of reminders, and immunisation ineligibility, if there was any delay in receiving a scheduled immunisation greater than 1 week. Hence, a priori, we determined the reminders for Penta-2 and 3 would be recalculated from the date the Penta-1 and 2 were administered, respectively. For those who did not receive either Penta-1 and or 2, the Penta-2 and 3 reminders were sent 2 weeks after the Penta-1 and 2 were past due, respectively. We performed daily audits of the immunisation clinic record book. When vaccines were received later than expected, the vaccine receipt date was used to calculate the next scheduled vaccine accordingly.

A child-health immunisation card, which listed the ages when a child was to receive his/her immunisation, comprised standard care. The intervention group received the automated text and call reminder plus standard care, while the control group received only standard care.

We obtained data for the primary outcome from immunisation clinic record books maintained at immunisation clinics and during the mid-study phone survey

of all participants. Study research assistants called each study participant on the telephone and obtained verbal reports—name and date—of when Penta-1, 2 and 3 were received. We assessed the receipt of measles vaccine solely from the immunisation clinic record book. This was because the health workers' strike did not affect the receipt or monitoring of the measles immunisation. We did not physically audit the child-health immunisation card given to parents and caregivers.

Study outcome

The primary outcome was the proportion of infants who received the Penta-1, 2 and 3 immunisations (henceforth referred to as Penta-3) at 18 weeks of age. The administration of BCG vaccination occurs at hospital discharge. Our intervention did not influence BCG receipt, and so it was not included in our primary outcome. We defined the secondary outcomes as completing Penta-3 and the measles immunisation by 12 months of age and receiving each within 1 week of the recommended time.

We assessed feasibility by the proportion of participants who received the verification text and call at enrolment, as well as those who reported receiving the reminders during the mid-study survey. Additionally, using a socio-ecological framework, we designed and administered a pre-study survey to assess sociodemographic characteristics of the mothers. We categorised the survey questions into maternal demographics, knowledge about and attitude towards immunisations, mothers' health and health-seeking behaviour, household demographic construct and access to health facility and health information related to immunisation. The mid-study survey was to assess acceptability of the intervention, perception of phone reminders by both groups and perceived barriers to completing immunisations.

Randomisation and blinding

Mother–infant dyad assignments to study groups in a 1:1 ratio was by a permuted randomisation scheme,¹⁹ using balanced random blocks of 6, 8 or 10. We stratified the randomisation by study site to account for centre population differences.

Randomisation was done in May 2016 at The Children's Hospital of Philadelphia. Allocation assignments were stored in sealed opaque envelopes and mailed via courier to the local study principal investigators. Only after obtaining written informed consent did the local study teams know the allocation assignment. Neither study participants nor research team were blinded. However, the immunisation clinic staff—those who administered and recorded immunisation—were blinded to study group allocations.

Sample size and statistical methods

Based on audits of the 2015 MCH-Ondo immunisation records and statewide reports, the baseline Penta-3 completion rate estimate was 75%. To account for a 10% loss to follow-up, we needed 300 mother–infant dyads in

each study arm to have a statistical power of 80% and an alpha level of 0.05, to detect a 10% difference in the primary outcome. We deemed this 10% difference to be of public health importance.

With the protocol amendment, we defined two study populations for the primary outcome analysis. First, a modified intention-to-treat (mITT) population—defined as all randomised subjects regardless of where immunisation was received or audited—immunisation record books or phone calls. The second was the per-protocol (PP) population—defined as all subjects who received immunisations only at MCH-Ondo or Akure and had immunisation receipt audited solely from the immunisation clinic record books.

We compared demographic characteristics and post-study survey variables between study groups using standard descriptive statistics. We used two-sample t-test or Wilcoxon rank-sum test for continuous variables and χ^2 test or Fisher's exact test for categorical variables. Risk ratios and risk difference were calculated for the primary and secondary outcomes. In a post hoc analysis, we used a generalised linear model for binomial distributions with 'log' link function to examine the association of selected demographic characteristics by study group on immunisation completion at 12 months. The selected variables were based on demographic factors reported in the literature to influence immunisation completion.^{5 10 20–22} We tested interaction effects of the demographic factors with the study group and report the p values for the interaction effects. Results are expressed as risk ratio along with their corresponding 95% CIs. Data were analysed using Stata V.15.1 (StataCorp, College Station, Texas, USA) with a two-sided significance level of 0.05.

RESULTS

We enrolled 250 mother–infant dyads from MCH-Ondo and 350 from Akure between August 2016 and June 2018. Five (0.8%) infants died during the follow-up period, 48 (8%) were lost to follow-up for the primary outcome and 184 (31%) for the 12-month outcomes (figure), while 155 (26%) received the Penta-3 immunisation at a different clinic.

Characteristics of study subjects

Infant and maternal demographic information were similar in both groups (table 1). Newborns were 48% female; the mean (SD) birth weight was 3040 g (485) and median (IQR) gestational age at birth was 38 (37–38) weeks, respectively (table 1). The majority of mothers were aged 18–35 years (88%), married (98%), and had a university (58%) or secondary education (39%). There were no differences in maternal knowledge about and attitude towards immunisation, mothers' health and health-seeking behaviour, household demographics or access to health facility and health information related to immunisation (table 1).

Primary and secondary analyses

For the Penta-3 completion, there was no significant difference between the intervention and control groups in either the mITT population assessed at 18 weeks (86% vs 81%, risk ratio (RR) 1.05, 95% CI 0.98 to 1.13; p=0.15) (table 2), 12 months (online supplementary table 1 and figure) or in the PP population (91% vs 88%, RR 1.03, 95% CI 0.97 to 1.10; p=0.37) (online supplementary table 2). The proportion of infants who completed the 12-month immunisation series was significantly higher in the intervention compared with the control group (74% vs 66%, RR 1.12, 95% CI 1.01 to 1.25; p=0.03) (table 2). To achieve completion of the immunisation series for one infant, 12.5 parents would need to receive the call and text message reminders. The proportion of infants who received the Penta-3 and measles immunisation within 1 week of the expected date was also significantly higher in the intervention compared with the control groups (84% vs 78%, RR 1.09, 95% CI 1.01 to 1.17, p=0.04 and 73% vs 65%, RR 1.33, 95% CI 1.02 to 1.26, p=0.02, respectively) (table 3). Those who collectively received all immunisations within a week of the expected date were also significantly higher in the intervention compared with the control group (57% vs 47%, RR 1.22, 95% CI 1.04 to 1.43; p=0.01) (table 3). In the post hoc subgroup analysis, there was no difference in the immunisation rates by the selected demographic variables assessed (table 4).

Feasibility, acceptability of intervention and barriers to completing immunisation assessed during the mid-study survey

At enrolment, 277 (92%) of those in the intervention group received a confirmatory text and call. Failed delivery resulted from poor local telecommunication service. It affected either internet or mobile phone connectivity. The mid-study survey was completed by 276 (92%) and 281 (94%) subjects in the control and intervention groups, respectively. In the intervention group, 86% reported they received reminders at least once during the study period. Twenty-five mothers (13%) reported receiving a partial or an unclear text reminder. Sixty-six per cent thought the text and calls were overall useful in reminding them of their child's appointment and 78% reported it reminded them all or some of the time of appointment dates.

Based on the survey data, the biggest reported barriers to completing immunisations were long wait times in 308 (55%) and transportation cost in 187 (34%). Forgetfulness (12; 2.2%) was one of the least reported barriers to completing the routine immunisation series (online supplementary tables 3 and 4).

DISCUSSION

Paired automated calls with text immunisation reminders significantly improved the proportion of infants who completed all routine immunisations by 12 months of age and the timeliness of vaccines administered late in

Table 1 Infant and maternal demographics by study group

	Control (n=300)	Intervention (n=300)	Total (N=600)
Infant demographics			
Female, n (%)	137 (47.4)	152 (52.6)	289 (48.2)
Birth weight, g, mean (SD)	3081.0 (505.5)	3118.7 (463.1)	3099.8 (484.7)
Gestational age, weeks, median (IQR)	38 (37–38)	38 (37–38)	38 (37–38)
Maternal demographics			
Age, years, mean (SD)	30 (5)	30 (5)	30 (5)
Married, n (%)	292 (97)	297 (99)	589 (98)
Maternal education†			
► University	163 (54.3)	183 (61.0)	346 (57.7)
► Secondary	123 (41.0)	109 (36.3)	232 (38.7)
► Primary	14 (4.7)	7 (2.3)	21 (3.5)
► No education	0 (0.0)	1 (0.3)	1 (0.2)
Maternal profession, n (%)*			
► Professional, top civil servant, politician or businesswoman	116 (38.7)	110 (37.0)	226 (37.9)
► Middle bureaucrats, technicians, skilled artisans, well-to-do trader	70 (23.3)	75 (25.3)	145 (24.3)
► Unskilled worker	60 (20.0)	56 (18.9)	116 (19.4)
► Housewife, unemployed	54 (18.0)	56 (18.9)	110 (18.4)
Christian, n (%)	279 (93.0)	280 (93.3)	559 (93.2)
No of children age ≤5, n (%)			
► 1	149 (50.5)	164 (54.8)	313 (52.7)
► 2	108 (36.6)	111 (37.1)	219 (36.9)
► 3	38 (12.9)	24 (8.0)	62 (10.4)
Maternal access to health facility and health information n (%)			
Mother can drive	40 (13)	48 (16)	88 (15)
Mother owns car	25 (8)	37 (12)	62 (10)
Transportation to hospital n (%)			
► Walk	13 (4.3)	8 (2.7)	21 (3.5)
► Public transportation	239 (79.7)	237 (79.0)	476 (79.3)
► Personal car	48 (16.0)	55 (18.3)	103 (17.2)
Average cost of transportation to immunisation clinic (US\$), median (IQR)	0.6 (0.4–0.9)	0.6 (0.4–0.9)	0.6 (0.4–0.9)
Average time to the hospital (min), median (IQR)	20 (12.5–30)	20 (10–25)	20 (10–25)
Report viewing or hearing advertisements about immunisations, n (%)			
Most times (at least twice a week)	152 (50.8)	165 (55.0)	317 (52.9)
Sometimes (at least twice a month/<twice a week)	118 (39.5)	111 (37.0)	229 (38.2)
Never	29 (9.7)	24 (8.0)	53 (8.8)
Mother's knowledge about and attitude towards immunisations, n (%)			
Agree, immunisations prevent disease (agree)	295 (98.3)	296 (98.7)	591 (98.5)
Agree, immunisation prevents diseases that can lead to death	245 (81.7)	247 (82.3)	492 (82.0)
Disagree, immunisations harmful to child	292 (97.3)	290 (97.0)	582 (97.2)
Knew immunisations are completed within 1 year of life	245 (82.2)	251 (83.7)	496 (82.9)
Able to name two diseases immunisation prevents	170 (56.7)	184 (61.3)	354 (59.0)
Mother's health and health-seeking behaviours, n (%)			
Rates health as good	295 (98.3)	290 (96.7)	585 (97.5)

Continued

Table 1 Continued

	Control (n=300)	Intervention (n=300)	Total (N=600)
Attended antenatal care	298 (99.7)	296 (98.7)	594 (99.2)
Received antimalaria prophylaxis	262 (87.3)	259 (86.3)	521 (86.8)
Received prenatal vitamin	237 (80.1)	227 (76.9)	464 (78.5)
Household sociodemographic construct			
Father's age, years, mean (SD)	36.3 (5.79)	36.1 (5.67)	36.2 (5.73)
Father's education† n (%)			
► University	194 (64.9)	192 (64.0)	386 (64.4)
► Secondary	100 (33.4)	102 (34.0)	202 (33.7)
► Primary	4 (1.3)	5 (1.7)	9 (1.5)
► No school	1 (0.3)	1 (0.3)	2 (0.3)
Domestic help present, n (%)	45 (15.1)	45 (15.1)	90 (15.1)
Family member at home to help, n (%)	216 (72.0)	220 (73.8)	436 (72.9)
Father's perception of immunisation positive, n (%)	297 (99.7)	299 (99.7)	596 (99.7)
Extended family perception of immunisation positive, n (%)	273 (91.3)	267 (89.0)	540 (90.2)
Has someone to bring child for immunisation if mother cannot, n (%)	92 (30.8)	98 (32.9)	190 (31.8)
Has someone to watch over other child while at immunisation visit, n (%)	158 (68.4)	163 (70.9)	321 (69.6)

*A director or manager of a government entity is an example of a top civil servant, while a clerk, typist or cashier is an example of a middle bureaucrat.

†Educational level indicates having some or completed the level of education.

the immunisation schedule. For the primary outcome, however, Penta-3 completion rates were higher than anticipated in both groups and not significantly different between groups.

For the Penta-3 completion, our findings differ from trials where text messages reminders alone were used in urban SSA settings,⁷⁻⁹ but are similar to a trial in rural Kenya.¹⁰ This may indicate that low literacy may not account for the lack of effectiveness of text reminders. An inappropriate study population may also explain this difference. Although we did not assess literacy level directly, 58% and 39% of our study participants had at least some university or secondary education, respectively.

Lack of 'awareness' ranks highest among barriers to immunisation completion reported by the Nigerian National Immunization Survey.⁵ The relatively high

awareness among the standard of care group in our study may reflect above-average site-specific counselling. Cost of transportation, distance to and overall 'conduciveness' at the clinics were the most frequent barriers to immunisation in our study population. These barriers have also been reported in trials conducted in Kenya and Nigeria.⁸⁻²² Furthermore, they have attracted research interest from organisations like the Bill and Melinda Gates foundation in recent years.²³

Findings from a study in rural Nigeria where real-time phone call immunisation reminders significantly improved immunisation completion premised our intent to use automated calls.¹⁴ The Nigeria study found a 41% increase in immunisation completion.¹⁴ Our intervention also resulted in improvements in immunisation completion at 12 months, with a smaller effect. Methodological

Table 2 Effect of Intervention on primary and secondary outcomes by study group*

Immunisation	Control (n=300)	Intervention (n=300)	RR (95% CI)	RD (95% CI)	P value
Penta-1† n (%)	289 (97)	285 (95)	0.98 (0.95 to 1.02)	-0.02 (-0.05 to 0.015)	0.31
Penta-2† n (%)	278 (93)	276 (92)	0.99 (0.95 to 1.04)	-0.01 (-0.04 to 0.05)	0.76
Penta-3† n (%)	244 (81)	257 (86)	1.05 (0.98 to 1.13)	0.04 (-0.02 to 0.10)	0.15
Measles‡ n (%)	196 (66)	220 (74)	1.12 (1.01 to 1.25)	0.08 (0.01 to 0.15)	0.03

*Modified intention-to-treat analysis; includes all subjects regardless of where immunisation was received or how it was audited.

†Assessed at 18 weeks.

‡Assessed at 12 months.

RD, risk difference; RR, risk ratio.

Table 3 Timely receipt of immunisation—individually and collectively by study group*†

Immunisation n (%)	Control (n=300)	Intervention (n=300)	RR (95% CI)	P value
Penta-1 at 6 weeks	250 (83)	257 (86)	1.03 (0.96 to 1.10)	0.43
Penta-2 at 10 weeks	252 (84)	256 (85)	1.02 (0.95 to 1.09)	0.65
Penta-3 at 14 weeks	233 (78)	253 (84)	1.09 (1.01 to 1.17)	0.04
Measles at 9 months	194 (65)	220 (73)	1.13 (1.02 to 1.26)	0.02
All immunisations	140 (47)	171 (57)	1.22 (1.04 to 1.43)	0.01

*Modified intention-to-treat analysis. Includes all subjects regardless of where immunisation was received or how it was audited.

†Timeliness defined as receiving vaccinations within 1 week of their due date.

RR, risk ratio.

differences may explain effect size differences. While our study sent out two reminder calls 2 days and the day before the immunisation due date, the prior published study used two real-time reminder phone calls plus four recalls if immunisations were missed as the primary intervention.¹⁴ Furthermore, and likely of importance, is the absence of the human element in automated calls. The ability to empathise, counsel and reassure mothers during a real-time phone call is lost with automation.

However, with current resource and organisational limitations, real-time phone calls plus recalls is time and human resource dependent and likely to result in a high implementation cost. Taking it to scale and sustenance will also be a challenge. Since most studies where text was used yielded between 60% and 80% completion rates, automated text and/or calls could serve as a first-pass reminder to be followed by real-time phone calls when immunisations are delayed. This may prove to be a more

Table 4 Subgroup analysis of completing immunisation by 12 months of age by study group

Characteristic	Control (n=300) (%)	Intervention (n=300) (%)	Stratum-specific RR (95% CI)	P value
Sex	112/163 (69)	116/148 (78)	1.14 (1.00 to 1.30)	0.74
▶ Male	86/137 (63)	105/152 (69)	1.10 (0.93 to 1.30)	
▶ Female				
Maternal education	124/163 (76)	144/183 (79)	1.22 (1.00 to 1.49)	0.17
▶ Completed university	74/137 (54)	77/117 (66)	1.03 (0.92 to 1.16)	
▶ Did not complete university				
Mothers age (years)	28/40 (70)	22/30 (73)	1.05 (0.78 to 1.41)	0.21
▶ >35	169/259 (65)	199/270 (74)	1.13 (1.01 to 1.27)	
▶ ≤35				
No of children under age 5	94/146 (64)	104/135 (77)	1.20 (1.03 to 1.39)	0.23
▶ >1	101/149 (68%)	117/164 (71%)	1.05 (0.91 to 1.22)	
▶ ≤1				
Mother owns a car	180/275 (65)	192/263 (73)	1.12 (1.00 to 1.25)	0.88
▶ No	18/25 (72)	29/37 (78)	1.09 (0.81 to 4.47)	
▶ Yes				
Mother's mode of transportation	171/252 (68)	183/245 (75)	1.10 (0.98 to 1.23)	0.51
▶ Public transportation or walk	27/48 (56)	38/55 (69)	1.23 (0.91 to 1.67)	
▶ Personal car				
Able to name two diseases immunisation prevents	64/92 (70)	57/82 (70)	1.00 (0.98 to 1.23)	0.51
▶ No	134/208 (64)	164/218 (75)	1.23 (0.91 to 1.67)	
▶ Yes				
Someone to bring child for immunisation if mother cannot	145/207 (70%)	151/200 (76%)	1.08 (0.96 to 1.21)	0.27
▶ No	53/92 (58%)	70/98 (71%)	1.24 (1.00 to 1.15)	
▶ Yes				
Study site location	59/125 (47%)	73/125 (58%)	1.24 (0.98 to 1.57)	0.25
▶ Ondo Town	139/175 (79%)	148/175 (85%)	1.06 (0.96 to 1.18)	
▶ Akure				

Data are n/N (%) and RR (95% CI). P values obtained from an interaction term between intervention.

RR, risk ratio.

cost-effective strategy. Digital immunisation registries can potentially improve the efficiency of this process by simplifying identification of defaulters.

In our study, we demonstrated modest feasibility instituting a customised automated text and call immunisation reminder system. However, poor phone and internet connectivity resulted in 8% delivery failure at enrolment. Furthermore, only 86% reported receiving at least one set of reminders during the study period. Phones that were switched off or had poor reception when messages were sent were the most common reported reasons for not receiving phone reminders. This is a limitation in web-based text and call systems which lack the ability to queue messages until the switched off phone is turned back on or a phone in poor reception areas moves to an area with better reception. In addition, voicemail is uncommon in these settings. We speculate that a system that uses the available local telecommunication network, which is usually more redundant, and has the capacity to provide more accurate log data, may yield better results.

Our study had some limitations. A 2015 institutional audit and state reported Penta-3 completion rates informed our sample size estimation. This likely underestimated the true institutional completion for those committed to receiving their immunisations at only MCH clinic. Furthermore, we introduced selection bias by recruiting only parents who planned for their child to receive immunisations only at MCH. Ondo state government during the study period provided free maternal care. Women in lower socioeconomic classes may have been those who lived further away and only used MCH for its free services. However, this was necessary to allow us to ascertain our intended primary outcome (immunisations confirmed by clinic records). The inclusion of women who own phones limits the generalisation of our results. Another limitation was the workers' strike. Unfortunately, this reflects a common reality in SSA and a need to strengthen the healthcare system. We were, however, able to retain 92% of the enrolled population for the primary outcome. Due to a clerical error, the number of subjects who refused or were ineligible for the study at MCH-Ondo could not be determined (figure). Another limitation is our inability to access reminder messaging log from our web-based platform. This is a limitation in most similar trials from SSA with only one trial reporting log data.¹⁰ Finally, although we intended to overcome low literacy levels by adding voice call reminders, 97% of study participants had at least secondary education.

The strengths of our study include a randomisation strategy that resulted in balanced demographic characteristics between study groups. We also used a more practical approach to study participant's message receipt as a proxy to assess feasibility.

CONCLUSIONS

In a semi-rural region in Nigeria, the implementation of an automated text and call immunisation system was

modestly feasible. The phone reminders significantly improved immunisation completion and timeliness. This study adds to the growing evidence of the effectiveness of phone reminders in SSA. Factors such as transportation cost, distance to and overall 'immunisation clinic conduciveness' were the most frequently reported barriers to immunisation. Combining reminder systems with innovative solutions capable of improving access to and efficiency of immunisation clinics are likely to improve immunisation uptake and completion in Nigeria and other SSA countries.

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Data sharing statement All available data can be obtained by contacting the corresponding author. All data requests should be submitted to the corresponding author for consideration. Access to anonymised data may be granted following review.

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