The effects of text reminders on the use of family planning services: evidence from a randomised controlled trial in urban Mozambique

Jessica Leight, Catherine Hensly, Marcos Chissano, Elana Safran, Liza Ali, Domingos Dustin, Julian Jamison

ABSTRACT

Introduction Reduction of unmet need for contraception is associated with enhanced health outcomes. We conducted a randomised controlled trial in Mozambique analysing the effects of text messages encouraging use of family planning services.

Methods This trial was conducted within a sample of women served by the Integrated Family Planning Program implemented by Population Services International, in which community health workers provide clinic referrals for family planning services. The evaluation enrolled 5370 women between 20 January and 18 December 2020 who received a referral, reported access to a mobile phone and provided consent. Women were randomly assigned to a treatment group that received a series of text message reminders encouraging them to visit a clinic or to a control arm. An intention-to-treat analysis was conducted to analyse the effect of reminders on the probability of a clinic visit and contraceptive uptake. The final analysis includes 3623 women; 1747 women were lost to follow-up.

Results Women assigned to receive the text reminders are weakly more likely to visit a clinic (risk difference 2.3 percentage points, p=0.081) and to receive a contraceptive method at a clinic (2.2 percentage points, p=0.091), relative to a base rate of 48.0% and 46.9%, respectively. The effect on clinic visits is larger and statistically significant in the prespecified subsample of women enrolled prior to the COVID-19-related state of emergency (3.2 percentage points, p=0.042).

Conclusion Evidence from this trial suggests that text message reminders are a promising nudge that increases the probability that women receive contraception.

Trial registration number AEARCTR-0005383.

INTRODUCTION

Worldwide, an estimated 10% of women of reproductive age are characterised by an unmet need for family planning—defined as women who want to stop or delay childbearing but are not using any method of contraception—including 17% of women in sub-Saharan Africa.1 Access to and use of modern contraception has substantial health benefits, enhancing birth spacing and thus reducing maternal and neonatal mortality and morbidity.2-4 There is also evidence of economic benefits in terms of higher earnings for women and enhanced educational outcomes for children.5 Accordingly, high levels of unmet need in low- and middle-income countries (LMICs) can have meaningful consequences for welfare. While access to family planning has been increasing, progress has been slow in some regions, particularly in sub-Saharan Africa.6

Ensuring universal access to sexual and reproductive healthcare is identified as one of the Sustainable Development Goals,7 and in Mozambique, the site of this evaluation, a commitment to the Family Planning 2020
global partnership was made in 2012 with the objective of accelerating progress towards family planning goals.\textsuperscript{8} Despite this commitment, unmet need remains high: national unmet need for modern family planning method in Mozambique was estimated as 23\% in 2015.\textsuperscript{3} Among adolescent girls aged 15–19, a total of 46\% are pregnant for the first time or already have one child, and in this group, only 14\% are using any contraception method.\textsuperscript{9}

Given these persistent challenges around access to contraception, researchers and policymakers have identified potential barriers informed by behavioural science for women who have unmet need for family planning or have not accessed their preferred method. For example, limited attention and present bias may prevent women from taking short-term steps to use a contraception method despite their long-term family planning goals. ‘Nudges’ designed to address these barriers have potential to be integrated into family planning policies and programming. The evidence around the effects of nudges on health-related outcomes is substantial\textsuperscript{10,11} but derived primarily from high-income countries, and there is relatively limited literature from LMICs.

More specifically, one popular nudge is reminders or other information delivered by text message or mobile phone applications. Existing reviews of mobile health (mHealth) interventions in LMICS have noted there is no consistent evidence that these interventions lead to behaviour change,\textsuperscript{12} and highlighted that many published papers do not provide any evidence on health outcomes.\textsuperscript{13–15} In family planning specifically, a number of published studies evaluate the effects of short message service (SMS) or rapid message interventions, but report only effects on variables such as contraceptive knowledge or attitudes. These evaluations do not measure or do not find any evidence of shifts in behaviours such as contraceptive use.\textsuperscript{16–20} Accordingly, a broader evidence base is needed.\textsuperscript{21}

This paper reports on a randomised controlled trial conducted in the context of a large community health worker programme in urban and periurban Mozambique led by Population Services International (PSI). A series of text message reminders was designed to encourage women who had received a referral from a community health worker to visit a health clinic for a family planning consultation. The objective of the trial was to evaluate the effect of these text reminders on the probability of a clinic visit as well as the probability of contraceptive uptake.

**METHODS**

**Trial design and participants**

This two-arm, parallel randomised controlled trial was conducted between January 2020 and January 2021 in urban and periurban areas of two provinces in Mozambique, Nampula (including Nampula city, Angoche, Ilha de Mocambique, Murrupula and Nacala Porto) and Sofala (including Beira and Dondo). The trial was jointly conducted by PSI and researchers based at the Office of Evaluation Sciences in the US General Services Administration and the International Food Policy Research Institute. The trial protocol and Consolidated Standards of Reporting Trials checklist are available as supporting information (online supplemental material S1, S2).

The target sample for the study was women of reproductive age served by community health workers in the Integrated Family Planning Program (IFPP). PSI delivered IFPP services in the provinces of interest, deploying community health workers (known as promoters) who offered women information about family planning in visits to homes or neighbourhoods in their service area. Visits focused on providing information about family planning and addressing common myths, and did not entail the direct provision of any family planning methods; rather, promoters provided women who voluntarily expressed demand for family planning with referrals to public health facilities, where family planning counselling from a nurse and contraceptive methods are available at no cost. Promoters also conducted up to three follow-up visits with women as needed, to provide more information or address concerns about side effects of a method obtained. PSI recruited around 100—120 promoters in the target regions, organised into teams of around five promoters each working with a single supervisor. Promoters conducted around 13—15 visits per day on average and were compensated per visit conducted and for each woman referred who visited a clinic.

In addition, promoters recorded information about each visit and the woman who participated in PSI’s mobile application, Connecting with Sarah (CwS). PSI and public nurses in local health clinics used the same mobile application to record when women presented a referral for family planning counselling, and to record if a contraceptive method was provided.

The following eligibility criteria were specified for enrolment in the trial. Women 18 or older were eligible if they were visited by a promoter and received a referral to a health facility for further services, provided a phone number, and provided written consent for inclusion in the evaluation. Enrolment commenced on 20 January 2020 and was suspended on 4 April 2020 due to the state of emergency (SOE) declared in Mozambique as a result of the COVID-19 pandemic. Enrolment then resumed on 1 October 2020, concluding on 18 December 2020. The target sample size was specified to be 5000 women enrolled; given this sample size, the trial could detect an increase in the probability of a clinic visit of 4 percentage points. The trial ultimately closed following a shorter enrolment period due to COVID-19 disruptions, and the timing of enrolment closure was dictated by the timeline for broader programme conclusion.

The two provinces included in this evaluation, Nampula and Sofala, are highlighted in figure 1. Demographic data from the 2015 AIDS Indicator Survey conducted by the Demographic and Health Surveys programme can be used to characterise the sample. In Sofala, 26\% of urban

Adults surveyed report no education and 43% report primary education, while in Nampula, the corresponding figures are 36% and 41%. Nationwide in Mozambique, 35% of urban adults report no education, and 48% report primary education. Socioeconomic indicators are generally higher in Sofala, where 82% of urban adult residents surveyed report access to electricity, and around 70% report their residences have cement walls and cement floors; in Nampula, 64% of urban adults report access to electricity at home; 46% have cement floors; and 35% have cement walls. Nationwide in Mozambique, 38% report access to electricity; 40% have cement floors; and 30% have cement walls.9

Randomisation and masking

Eligible women enrolled in the evaluation had their consent forms verified by trial staff, and the sample was then aggregated across the two study provinces. Women were randomly assigned to the treatment or control arm using the phone number provided on their consent form. Randomisation was conducted at least twice weekly by PSI staff in Maputo in Stata V.14 using a reproducible seed, employing stratification at the level of the promoter supervisor. On average, randomisation was conducted 5 days following the provision of consent. (Given that only two provinces are included in the evaluation, randomisation at the province level was infeasible.)

The intended allocation ratio of participants to the treatment and control arm was 1:1. When the study was launched, the randomisation code had an unintentional error that slightly increased the probability of assigning women to treatment: if the number of women in a particular strata (defined by supervisor identity on a given randomisation day) was odd, the final observation was uniformly assigned to treatment. (The identity of the final observation was, however, determined randomly via assignment of random numbers.) This error was corrected as of 28 February 2020.

Given the nature of the intervention, it was not possible to mask participants assigned to the treatment arm to their assignment. However, participants assigned to the control arm may have been blind to their assignment. Promoters and health staff at the health facilities were blind to study group assignment.

Intervention design

The intervention of interest was a series of short text message reminders designed to encourage take-up of health facility visits for family planning counselling by women who had been provided referrals by promoters. The messages were developed by the research and implementation teams with the objective of providing women with targeted, brief reminders about the importance of family planning; the opportunity to visit the health facility; and the opportunity to follow-up with their promoter, as needed. The text of the reminders can be found in table 1 in English; in the trial, all messages were in Portuguese. To maintain an appropriate level of confidentiality, the messages refer to family planning but also use more general terms such as health, hospital and promoter.

Table 1 also describes message timing as follows: on the same day as randomisation, women assigned to the treatment arm were registered by the vendor (SISLOG) who was responsible for delivering the text messages. Message timing is defined relative to this registration date. (On average, the time elapsed between the date of promoter visit and the randomisation and the registration date was 5 days, the time required to verify and aggregate the sample of women providing consent across all promoters.) Messages were sent on the first and fourth days following registration, and then 1, 2, 3 and 4 weeks following the first message.

Women who received the messages were of course free to share information received with members of their social networks, and this may have included women who were enrolled in the evaluation and assigned to the control arm. Any such communication between treatment and control beneficiaries would serve to reduce the magnitude of the estimated treatment effect.
Data and outcomes

The data employed in this analysis were collected via a mobile application, CwS, used by promoters and health facility nurses. Following each promoter visit, promoters used a PSI-provided smartphone to access the CwS platform and recorded information about the beneficiary (age; current contraceptive use, if any; and access to a mobile phone), and whether a referral was issued. The data were uploaded at the conclusion of the visit and were automatically stamped with the date, time and global positioning system (GPS) coordinates. Similarly, nurses or health staff at referring facilities used CwS to record visits from women who visit for family planning services with a referral, and the two records were linked by a referral number. These facility records were also time-stamped and include information about the type of contraceptive method provided (counselling only, contraceptive pill, Depo-Provera, Sayana Press, contraceptive implant, intrauterine implant or other).

The analysis includes data from 20 January 2020, the day on which enrolment was launched, until 31 January 2021, 6 weeks following the last enrolment. Between 5 April and 1 October 2020, enrolment in the evaluation was paused, but we still observe data from clinics in this period and are thus able to identify if women who were already enrolled in the evaluation visited a clinic. Our data do not include any records of women’s receipt of contraceptive methods in locations other than public clinics (ie, pharmacies or private clinics). Also, should a woman visit a public clinic and fail to provide her referral information, she could still receive family planning services at no cost, but this visit would not be recorded in the CwS application and thus would not be visible in our data. Data from the 2003 Demographic and Health Survey (not reported in more recent survey rounds) suggest that among women reporting use of modern contraceptives, less than 5% report receipt from a private clinic, nurse or pharmacy, suggesting that our data presumably include the majority of contraceptive receipt for this population.25

The primary outcome of interest is a binary variable for a clinic visit following promoter referral. Secondary outcomes of interest include a binary variable for contraceptive uptake at the clinic and a continuous variable capturing the number of days between the promoter visit and the woman’s visit to the clinic, conditional on the observation of a clinic visit. There were no changes to the primary outcomes following the commencement of the trial.

For the variables corresponding with clinic visit and contraceptive uptake, these variables are coded as one if any facility activity is recorded on any day following the woman’s promoter visit. Within a given randomisation stratum, all women included were randomised into the sample on the same day and visited by promoters within the same approximately 3-day period; accordingly, within a randomisation strata, all are observed in the sample for approximately the same period of time. There is, however, variation across strata in how long women were observed in the sample: women enrolled in the evaluation in the first month were observed for roughly 10 months, while women enrolled in the final month were observed for 1 month. The inclusion of binary variables for strata in the primary specification will adjust for this variation across strata in the observable period. In addition, as a robustness check, we define an additional variable equal to clinic visit within 30 days, coded as one for all observations in which the respondent is observed visiting a clinic within this time period.

The analysis also draws on administrative data from SISLOG, a local text message vendor, allowing us to report whether text messages were recorded as delivered or undelivered because the number was out of service. (However, one of the three cell service companies in Mozambique did not provide this information to SISLOG, and messages sent to customers of this company would never be recorded as ‘delivered’; accordingly, the estimated delivery rate can be considered to be a lower bound.) We are not able to access any data reporting whether a message was read.

Statistical analysis

The methods used in the statistical analysis were prespecified in a registered analysis plan.25 We estimate risk
differences between the treatment and the control arms using ordinary least squares in an intent-to-treat framework. We use a linear model for ease of interpretation and cluster the SEs with respect to the day of randomisation to account for potential correlation in the error term for women who were visited by promoters and referred to clinics in the same period.25 We prespecified the estimation of a model that adjusted for randomisation strata (the interaction of randomisation day and a binary variable for supervisor), age, province and distance to the nearest health facility. Distance is estimated as the linear distance employing the GPS coordinates associated with the promoter visit conducted at home and the coordinate of the closest health facility, and included as a series of binary variables capturing deciles of distance to allow for non-linear effects.

The analysis plan also prespecified analysis of heterogeneity along a number of dimensions: beneficiary age, contraceptive use at first meeting with the promoter, sole ownership of phone, time elapsed between promoter visit and randomisation, and distance to clinic. Following the suspension of enrolment due to the COVID-19-related SOE, the analysis plan was also updated to specify the separate analysis of treatment effects in the pre-SOE and post-SOE period.

In addition, we estimate a treatment on the treated effect in a two-stage least squares specification. In this model, the independent variable is a binary variable equal to 1 if the phone number reports receipt of all eight reminder messages and 0 otherwise (and zero for all phone numbers assigned to the control arm), and we instrument for this variable with treatment assignment. The model adjusts for the same covariates described in the ITT model, and standard errors are again clustered with respect to the day of the randomisation. This analysis was also prespecified.

In some cases, the same phone number was enrolled in the evaluation more than once, or the same number was enrolled and randomised once but was observed multiple times in promoter data. This duplication could be accurate (multiple women use the same phone) or could be recorded in error. There are 201 cases in which a phone number was enrolled more than once, and 531 cases in which the phone number was observed more than once in outcome data; there are 598 phone numbers characterised by at least one form of duplication. Given this pattern, all the variables in the primary specification are converted to phone number-level means: the dependent variable (a binary variable for a clinic visit), the independent variable capturing treatment assignment and all covariates. The same procedure was used when a phone number that was uniquely enrolled via the consent forms matched to multiple observations of that number in the CwS database, indicating the number was recorded for separate visits to different women. Due to this procedure, there is some continuous variation in both the treatment and the dependent variables between 0 and 1, corresponding to phone numbers that were partially treated: some women who enrolled using this number were assigned to treatment, and some women were assigned to control. For example, if a phone number was enrolled twice (listed on two separate consent forms) and was randomised once to treatment and one to control, treatment status for this phone number would be coded as 0.5.

Intervention costs were also tracked and analysed as part of this trial using a provider perspective. This cost estimate includes the full cost of the contract with the vendor who provided text message services, and an estimate of staff and administrative costs required to design and implement the intervention.

Role of the funding source
The funders of the study (United States Agency for International Development) had no role in study design, data collection, analysis, interpretation or writing of the results. The corresponding author (JL) had full access to all data in the study and the final responsibility for the decision to submit for publication.

Patient and public involvement
During the design phase of this trial, interviews of women served by IFPP as well as focus groups of promoters, supervisors and PSI staff were convened in order to provide feedback about the challenges faced by women in accessing contraception, and by promoters in serving them. In addition, PSI staff shared sample messages with programme beneficiaries in focus groups to ensure clarity and relevance of the text messages and made adjustments based on participant feedback. An author reflexivity statement is attached (online supplemental material S5).

RESULTS
Figure 2 depicts the trial profile. During the period of enrolment, 110 200 women were visited by PSI promoters and assessed for eligibility; 41 642 women received referrals to health facilities. This sample receiving referrals was screened for inclusion in the trial, and 13 586 women met the primary criteria (provided a mobile phone number). A total of 5370 women provided written consent and were enrolled in the evaluation. There was some duplication of phone numbers across enrolled women; some women were registered by the text message vendor using the same phone number, either accurately (because two women in the sample share a number) or in error. Accordingly, 5169 unique phone numbers were enrolled: there were 170 numbers that were registered twice, 14 numbers that were registered three times and 1 number that was registered four times. Within the sample, 2728 enrolled women were randomly assigned to treatment, and 2642 enrolled women were randomised to control.

The phone numbers recorded on consent forms were matched to the phone numbers available in the CwS database. Only 3623 enrolled women could be matched with an administrative record of their promoter and (if...
applicable) clinic visits, corresponding to 3468 unique phone numbers. This results in 1747 women with written consent who were lost to follow-up or 32.5% of the sample: 906 or 33.2% assigned to the treatment arm and 841 or 31.8% assigned to the control arm, a difference that is not statistically significant (p=0.281). Women who were lost to follow-up are missing outcome data and summary data on covariates, as all CwS data are linked to the phone number.

We present characteristics of the study sample in Table 2 for the full sample and the sample enrolled both pre-SOE and post-SOE. We also report summary statistics for the sample observed in treatment and control arms. The average age of women enrolled in the sample is 25.7 years. Distance to facility is low on average (less than four kilometres), though this is higher pre-SOE (5.02 kilometers) than post-SOE (2.63 kilometers). Within the sample, 69.6% report they are the sole owner of the phone number they have provided. Among the women, 30.9% report they are currently using contraception on their first interaction with the promoter, and this probability is dramatically higher pre-SOE (44.0%) compared with post-SOE (20.9%). We also report the p value corresponding to a $\chi^2$ test of the joint hypothesis that these covariates are balanced across treatment and control arms; this hypothesis cannot be rejected (p=0.903).

Considering the outcome variables of interest, the probability of a clinic visit following a referral is around 48% on average, slightly lower pre-SOE (43.7%) compared with post-SOE (51.3%). The probability of a clinic visit within a month of the promoter’s visit is 44.7%.
Table 2 Characteristics of sample respondents

<table>
<thead>
<tr>
<th></th>
<th>All</th>
<th>Pre-SOE</th>
<th>Post-SOE</th>
<th>Assigned to control</th>
<th>Assigned to treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (cont.)</td>
<td>25.652 (3468)</td>
<td>25.338 (1502)</td>
<td>25.891 (1966)</td>
<td>25.671 (1679)</td>
<td>25.634 (1789)</td>
</tr>
<tr>
<td>Distance to facility (cont.)</td>
<td>3.661 (3468)</td>
<td>5.018 (1502)</td>
<td>2.625 (1966)</td>
<td>3.580 (1679)</td>
<td>3.738 (1789)</td>
</tr>
<tr>
<td>Sole phone owner (binary)</td>
<td>69.6% (2289/3468)</td>
<td>67.2% (965/1502)</td>
<td>71.4% (1324/1966)</td>
<td>70.5% (965/1502)</td>
<td>68.7% (1324/1966)</td>
</tr>
<tr>
<td>Current user of contraception (binary)</td>
<td>30.9% (843/3468)</td>
<td>44.0% (583/1502)</td>
<td>20.9% (260/1966)</td>
<td>29.7% (583/1502)</td>
<td>32.0% (260/1966)</td>
</tr>
<tr>
<td>Joint χ² test of balance across covariates</td>
<td>p=0.903</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Probability of clinic visit (binary)</td>
<td>48.0% (1547/3468)</td>
<td>43.7% (607/1502)</td>
<td>51.3% (940/1966)</td>
<td>47.4% (607/1502)</td>
<td>48.6% (940/1966)</td>
</tr>
<tr>
<td>Probability of clinic visit within 1 month (binary)</td>
<td>44.7% (1441/3468)</td>
<td>41.1% (573/1502)</td>
<td>47.5% (868/1966)</td>
<td>43.6% (573/1502)</td>
<td>45.8% (868/1966)</td>
</tr>
<tr>
<td>Probability of contraceptive receipt (binary)</td>
<td>46.9% (1505/3460)</td>
<td>42.3% (585/1499)</td>
<td>50.4% (920/1961)</td>
<td>46.3% (585/1499)</td>
<td>47.4% (920/1961)</td>
</tr>
<tr>
<td>Days elapsed: promoter to clinic visit (cont.)</td>
<td>11.459 (1779)</td>
<td>11.435 (704)</td>
<td>11.475 (1075)</td>
<td>11.661 (845)</td>
<td>11.276 (934)</td>
</tr>
<tr>
<td>Probability of receiving any text reminder (treatment arm only)</td>
<td>79.4%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Probability of receiving all text reminders (treatment arm only)</td>
<td>62.1%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

This table reports summary statistics for the full sample for the sample enrolled in the pre-COVID and post-COVID state of emergency period; and for the sample assigned to the treatment and control arms. The labels indicate whether the variable of interest is continuous (for which the mean and the number of observations is reported) or binary (for which the mean and the number of observations is reported). For the purposes of the table, any phone number that is enrolled multiple times and partially treated is included as part of the treatment arm.

(without any time limit) is 46.9%. Around 11 days elapse on average between the promoter visit and the woman’s visit to a clinic (median days elapsed is eight). In analysing summary statistics for numbers assigned to treatment and control, there is little evidence of any meaningful differences across the two arms.

Data from the text message vendor on intervention implementation are also reported. On average, phone numbers assigned to the treatment arm reported receipt of 79.4% of text messages, though as noted earlier this may reflect a lower bound due to missing information from one cell service company. Of the enrolled phone numbers, 62.1% reported receipt of all eight text messages.

Table 3 presents the primary results analysing the effect of the intervention on clinic visits and contraceptive take-up. In the intent-to-treat analysis, the estimated effect of the text reminders on the probability of a clinic visit is positive and marginally statistically significant at the 10% level: women who received text reminders were 2.3 percentage points (95% CI –0.003% to 0.048%, p=0.081) more likely to visit a clinic for family planning services, relative to a probability of 47.4% in the control arm. This is a proportional effect of 4.9%. The effect is larger (3.2 percentage points, or a proportional effect of 6.8%) and statistically significant at the 5% level in the pre-SOE period (95% CI 0.001% to 0.063%, p=0.042) and the same magnitude but noisily estimated in the post-SOE period (95% CI –0.017% to 0.064%, p=0.254). As a robustness check, we also estimate the effect of the reminder on a clinic visit within 1 month and here observe an effect that is larger vis-à-vis the main treatment estimate (3.7 percentage points) and statistically significant at the 1% level in the full sample (95% CI 0.012% to 0.062%, p=0.004).

We also report the estimated treatment effects for the two additional secondary variables of interest. For the probability of receiving a contraceptive method at a clinic, the estimated effect of the text reminders is positive (2.3 percentage points) and statistically significant at the 10% level (95% CI –0.004% to 0.048%, p=0.091) though again significant at the 5% level when restricted to the pre-SOE period. For the time elapsed between promoter visit and clinic visit (conditional on observing a clinic visit), we observe a negative coefficient of –1.219 that is statistically significant at the 1% level (95% CI –2.133% to –0.306%, p=0.009).

Table 4 presents the two-stage least squares analysis. The estimated effect of receiving all reminder messages is now larger (3.6 percentage points, 95% CI –0.005% to 0.076%, p=0.083) or a proportional effect of 7.6%. There is a 5.8 percentage point increase in the probability of a clinic visit within a month (95% CI 0.019% to 0.098%, p=0.004) and a 3.5 percentage point increase in the probability of contraceptive receipt (95% CI –0.006% to 0.077%, p=0.092). Redemption time is around 2 days shorter on average (95% CI –3.478 to –0.507, p=0.009).
Table 5 presents the results by subsamples. The positive treatment effect is observed to be larger (4.1 percentage points) for women under the age of 25 (95% CI −0.001 to 0.083, p=0.055). By contrast, the estimated effect for women over the age of 25 is in fact negative, though statistically insignificant. There is no evidence of any meaningful heterogeneity in response with respect to whether the woman reports she is currently using contraception on meeting the promoter, whether she reports she is the sole owner of the phone registered, the time elapsed between the promoter visit and the date on which the woman enters the randomisation sample, or distance to the health facility.

Cost data for the intervention suggests a total cost of $7593.87 for the implementation of the text reminders, excluding the costs of the evaluation itself. Given the number of women targeted by the intervention (enrolled in the treatment arm and successfully reached by at least one text reminder), the cost per woman targeted is $2.74.

DISCUSSION

The evidence presented here from a randomised trial of text reminders suggests that this intervention can be an effective strategy to increase facility visits by women referred for family planning services in urban and periurban Mozambique. The effect of the text reminders is positive and statistically significant: women who received text reminders are more likely to visit a clinic, are more likely to report receiving a contraceptive method at a clinic and visit the clinic more rapidly (conditional on ever reporting a visit). Moreover, the effects are particularly large for younger women and prior to the COVID-19-related SOE. In addition, the fact that the effect on contraceptive use is also positive indicates that the marginal woman encouraged to visit the clinic is more likely to report receiving a contraceptive method at a clinic and visit the clinic more rapidly (conditional on ever reporting a visit). Moreover, the effects are particularly large for younger women and prior to the COVID-19-related SOE.

Table 3  Effect of the text message reminders on primary and secondary outcomes: intent-to-treat analysis

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Full sample</th>
<th>Pre-SOE</th>
<th>Post-SOE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Clinic visit (primary)</td>
<td>0.023 (−0.003 to 0.048)</td>
<td>0.032 (0.001 to 0.063)</td>
<td>0.023 (−0.017 to 0.064)</td>
</tr>
<tr>
<td>N</td>
<td>3468</td>
<td>1502</td>
<td>1966</td>
</tr>
<tr>
<td>Clinic visit within 1 month (secondary)</td>
<td>0.037 (0.012 to 0.062)</td>
<td>0.042 (0.008 to 0.075)</td>
<td>0.040 (0.000 to 0.080)</td>
</tr>
<tr>
<td>N</td>
<td>3468</td>
<td>1502</td>
<td>1966</td>
</tr>
<tr>
<td>Received contraceptive method (secondary)</td>
<td>0.022 (−0.004 to 0.048)</td>
<td>0.034 (0.003 to 0.065)</td>
<td>0.021 (−0.019 to 0.062)</td>
</tr>
<tr>
<td>N</td>
<td>3460</td>
<td>1499</td>
<td>1961</td>
</tr>
<tr>
<td>Days between promoter visit and clinic visit</td>
<td>−1.219 (−2.13 to −0.306)</td>
<td>−0.643 (−2.170 to 0.884)</td>
<td>−1.302 (−2.541 to −0.062)</td>
</tr>
<tr>
<td>N</td>
<td>1779</td>
<td>704</td>
<td>1075</td>
</tr>
</tbody>
</table>

This table reports estimated risk differences corresponding to the effect of assignment to the text message treatment for the full sample, the sample enrolled pre-SOE and the sample enrolled post-SOE. For phone numbers that were enrolled more than once, we identify the phone number as pre-SOE if at least one enrolment was recorded pre-SOE. Eight phone numbers are missing data for method choice that would allow us to identify contraceptive receipt. Days between promoter visit and clinic visit are coded as missing for women who never reported a clinic visit. Each cell reports the coefficient and 95% CI, and p value and the number of observations.

Table 4  Effect of the text message reminders on primary and secondary outcomes: treatment-on-the-treated analysis

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Coefficient</th>
<th>95% CI</th>
<th>p value</th>
<th>N</th>
</tr>
</thead>
<tbody>
<tr>
<td>Clinic visit (primary)</td>
<td>0.036 (−0.005 to 0.076)</td>
<td>p=0.083</td>
<td>3468</td>
<td></td>
</tr>
<tr>
<td>Clinic visit within 1 month (secondary)</td>
<td>0.058 (0.019 to 0.098)</td>
<td>p=0.004</td>
<td>3468</td>
<td></td>
</tr>
<tr>
<td>Received contraceptive method (secondary)</td>
<td>0.035 (−0.006 to 0.077)</td>
<td>p=0.092</td>
<td>3460</td>
<td></td>
</tr>
<tr>
<td>Days between promoter visit and clinic visit</td>
<td>−1.993 (−3.478 to −0.507)</td>
<td>p=0.009</td>
<td>1779</td>
<td></td>
</tr>
</tbody>
</table>

This table reports estimated risk differences corresponding to the effect of receipt of a full set of eight text messages for the full sample, using a two-stage least squares analysis in which receipt of text messages is instrumented by treatment assignment. Eight phone numbers are missing data for method choice that would allow us to identify contraceptive receipt. Days between promoter visit and clinic visit are coded as missing for women who never reported a clinic visit. Each cell reports the coefficient and 95% CI, and p value and the number of observations.
### Table 5  Effect of the text message reminders on primary and secondary outcomes: intent-to-treat analysis for prespecified subsamples

<table>
<thead>
<tr>
<th>Subsample of interest</th>
<th>Current non-user of contraception</th>
<th>Current user of contraception</th>
<th>Reports shared ownership of phone</th>
<th>Reports sole ownership of phone</th>
<th>Age ≤25</th>
<th>Age &gt;25</th>
</tr>
</thead>
<tbody>
<tr>
<td>Treatment effect (risk difference)</td>
<td>0.030 (−0.014 to 0.074)</td>
<td>0.022 (−0.025 to 0.070)</td>
<td>−0.004 (−0.064 to 0.055)</td>
<td>0.032 (−0.003 to 0.068)</td>
<td>0.041 (−0.001 to 0.083)</td>
<td>−0.016 (−0.064 to 0.033)</td>
</tr>
<tr>
<td><strong>p</strong></td>
<td>0.184</td>
<td>0.356</td>
<td>0.883</td>
<td>0.074</td>
<td>0.055</td>
<td>0.53</td>
</tr>
<tr>
<td><strong>N</strong></td>
<td>2206</td>
<td>1262</td>
<td>917</td>
<td>2551</td>
<td>1918</td>
<td>1550</td>
</tr>
</tbody>
</table>

This table reports estimated risk differences corresponding to the effect of assignment to the text message treatment on the probability of a clinic visit for the specified subsamples. For the analysis of heterogeneity, phone numbers that were repeated in the analysis and thus have a constructed (continuous) value for the covariate are included in the subsample reporting non-zero values; that is, current user of contraception includes phone numbers with multiple observations in which at least one is identified as a current user of contraception; current non-user of contraception includes those phone numbers for which no recorded observation reports any contraceptive use. 28 observations are missing estimated distance to the closest facility. Each cell reports the coefficient and 95% CI, p values and the number of observations.

In general, the intervention. This paper joins a limited literature to date analysing the effects of mHealth interventions and nudges informed by behavioural science on primary healthcare use, including use of sexual and reproductive healthcare, in LMICs. Only two previous papers to our knowledge have analysed the effects of a nudge on family planning uptake in LMICs. Only two previous papers to our knowledge have analysed the effects of a nudge on family planning uptake. One evaluation conducted in urban Kenya found that there was no incremental effect of text message reminders for women offered vouchers to receive family planning at no cost at a private clinic. A second evaluation of vouchers and SMS reminders in Kenya targeting family planning. In particular, the previous intervention—designed to enhance contraceptive counselling and communication with a health provider—had a large positive effect on adoption of long-acting reversible contraception. A third recent paper applied an mHealth intervention designed to enhance contraceptive counselling by reminding women who recently delivered to use any contraceptive method. This paper joins a limited literature to date analysing the effects of mHealth interventions and nudges informed by behavioural science on primary healthcare use, including use of sexual and reproductive healthcare, in LMICs. Only two previous papers to our knowledge have analysed the effects of a nudge on family planning uptake. One evaluation conducted in urban Kenya found that there was no incremental effect of text message reminders for women offered vouchers to receive family planning at no cost at a private clinic. A second evaluation of vouchers and SMS reminders in Kenya targeting family planning. In particular, the previous intervention—designed to enhance contraceptive counselling and communication with a health provider—had a large positive effect on adoption of long-acting reversible contraception. A third recent paper applied an mHealth intervention designed to enhance contraceptive counselling by reminding women who recently delivered to attend antenatal care facilities in Zanzibar led to a 13 percentage point increase in the use of antenatal care. In general, the magnitude of the effect observed here (ranging from 2 to 6 percentage points for the sample under 25, versus 13 percentage points for the full sample) is consistent with the literature on the effects of mHealth interventions on knowledge or attitudes around contraception, rather than safety and ultimately choosing not to use any contraceptive method.
In addition to encouraging take-up of contraception, the text reminders were particularly effective in encouraging women to visit the clinic promptly: the effect on clinic visits within 30 days was proportionally around 60% larger than the effect on any clinic visit, and among women who did visit clinics, they visited on average 1 day sooner relative to a median of 8 days. The timing of visits is potentially of relevance, given that use of long-acting contraceptives is relatively unusual in this sample: among women who visit a clinic, less than a quarter received a long-acting method (3% received an intrauterine device, and 19% received a contraceptive implant). (These summary statistics were not presented in the Results section, as they are simple means rather than treatment effects.) These patterns are broadly similar to those observed in the most recent Demographic and Health Survey conducted in Mozambique in 2011, as only 10% of women reporting use of contraception in that survey reported use of a long-acting method.32

In a context in which use of short-acting contraceptives is dominant, interventions that target more timely clinic visits may have significant health implications if they effectively reduce contraceptive discontinuation, given evidence from the literature around high rates of discontinuation in LMICs.33 34 Discontinuation is especially high for short-acting methods,35 though it is also observed for long-acting methods,36 and is particularly common for adolescents.37 Women who have discontinued previous use without switching to a new method thus constitute a substantial share of the population of women with unmet need.38 39 While we do not have any systematic data about discontinuation of contraceptive methods in this evaluation, text messages encouraging more rapid follow-up at clinics or with promoters may be a promising strategy to reduce discontinuation (allowing for method refill or counselling about side effects or options for switching), particularly for the adolescent population that is shown to be more responsive to the reminders.

It is important to note that this evaluation was conducted in 2020 during a year of acute uncertainty and disruption linked to the COVID-19 pandemic and in a general environment of health-related fears. In Mozambique, an SOE linked to the pandemic was in effect from 31 March 2020 to 7 September 2020 (thereafter transitioning to a state of public calamity); however, promoters continued to operate throughout this period, and health clinics remained open. (Enrolment in the study was paused during the SOE, as described in more detail earlier.) Previous work by this research team analysing the short-term effects of the SOE on the behaviour of PSI promoters and women found a modest short-term drop in both service provision and use through June 2020, followed by a relatively rapid rebound.40 However, the fact that the effect of the text reminders is generally reduced and more noisily estimated in the post-SOE period constitutes suggestive evidence that in this period, other barriers (eg, anxiety about visiting clinics or increased economic stress) may have become more salient, and our intervention did not effectively target these barriers.

This analysis has a number of strengths and weaknesses. It is only the third evaluation to analyse the effectiveness of text reminders on contraceptive uptake in LMICs and the first to do so including a sample of women who are not exclusively postpartum. We analyse a substantial sample using administrative data (ie, not relying on self-reported use measures or additional data collection) and successfully completed the evaluation in an environment of substantial COVID-related disruptions. Key limitations of the evaluation include a high attrition rate (due to an inability to match phone numbers recorded manually on consent forms to phone numbers recorded in administrative data by promoters) and the reliance on a relatively narrow administrative dataset. Accordingly, we are unable to analyse intervention effects on use of contraception in other contexts (outside of public health clinics) and are unable to analyse longer-term use or other health outcomes. We also do not have access to any data on women’s literacy and their comprehension or recall of the messages. Finally, our findings are informative about the effectiveness of the reminders on a population of women that has already received referrals from health promoters and may not be informative about the effects of this intervention in a broader population.

CONCLUSION

This paper presents evidence from a randomised controlled trial evaluating a series of text message reminders designed to increase facility visits for family planning services in urban and peri-urban Mozambique. The results suggest that the reminders may be a promising strategy to increase clinic visits and associated uptake of contraception following referrals by community-based promoters, and these effects may be largest for younger women. Future programming and research work can explore the potential of incorporating mobile reminders to encourage consistent uptake of family planning services.

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Contributors JL, CH, MC, ES and JJ designed the evaluation. MC, LA and DD led the field implementation and administration of the evaluation in the field. JL conducted the analysis and drafted the manuscript. All authors provided the
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