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RESEARCH ARTICLE

REVISED A qualitative exploration of advantages and

disadvantages to using technology in the process of

randomised controlled trial recruitment [version 2; peer

review: 2 approved, 1 approved with reservations]

Lauren A. Muldowney¹, Sinéad M. Hynes², Megan Oglesby¹, Christopher P. Dwyer³

¹School of Medicine, University of Galway, Galway, County Galway, Ireland

²Occupational Therapy, School of Health Sciences, University of Galway, Galway, County Galway, Ireland
³Teacher Education, Technological University of the Shannon Midlands Midwest, Athlone, Westmeath, Ireland

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Abstract

Background

Despite its importance, recruiting a sufficient sample size for randomised controlled trials (RCTs) can pose a significant challenge, which has real-world impact on reliability of evidence, trial completion and ultimately, patient care. Technology has potential to enhance the recruitment process, but there is a lack of evidence regarding its current use and effectiveness. Consistent with findings from the PRioRiTy I study, the current research aims to explore the advantages and disadvantages to using technology during the recruitment process for RCTs.

Methods

Semi-structured interviews (n=7) were conducted with researchers involved in RCT recruitment in Ireland. Subsequently, a Public & Patient Involvement (PPI) panel focus group (n=3) was conducted to add further depth to these findings. The data were qualitatively analysed through 'Reflexive Thematic Analysis' to extract prominent themes.

Results

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1. Marie Falahee ^(D), University of Birmingham, Birmingham, UK

Open Peer Review

- 2. **Upasana Baruah**, National Ageing Research Institute, Victoria, Australia
- 3. **Milan Bimali** D, University of Arkansas for Medical Sciences, Little Rock, USA

Any reports and responses or comments on the article can be found at the end of the article.

A superordinate theme arose: *Tech is just a medium so that you can reach more people'*, along with two themes, which were corroborated by the PPI focus group: *Technology is used if and when the benefits outweigh the costs'* and *Success of recruitment through technology depends on the nature of the study.'*

Conclusions

This study provided a deeper understanding of the factors which influence researchers to employ technology in recruitment for RCTs. Implications suggest that future researchers should aim to adapt their recruitment approaches to meet digital tool preferences of their target cohort; and engage with patient groups in the community to allow networking opportunities for future studies. This research may contribute towards maximising efficiency in RCT recruitment.

Keywords

trial recruitment, thematic analysis, trials methodology, patient and public involvement, technology in RCT recruitment

Corresponding author: Christopher P. Dwyer (christopher.dwyer@tus.ie)

Author roles: Muldowney LA: Data Curation, Formal Analysis, Investigation, Methodology, Project Administration, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Hynes SM**: Conceptualization, Funding Acquisition, Methodology, Project Administration, Resources, Supervision, Visualization, Writing – Review & Editing; **Oglesby M**: Formal Analysis, Methodology, Writing – Review & Editing; **Dwyer CP**: Conceptualization, Formal Analysis, Funding Acquisition, Methodology, Project Administration, Resources, Supervision, Visualization, Formal Analysis, Funding Acquisition, Methodology, Project Administration, Resources, Supervision, Visualization, Formal Analysis, Funding Acquisition, Methodology, Project Administration, Resources, Supervision, Visualization, Formal Analysis, Funding Acquisition, Methodology, Project Administration, Resources, Supervision, Visualization, Writing – Review & Editing

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REVISED Amendments from Version 1

Updates have been made to the discussion, specifically the limitations section with respect to further detailing implications of the study's, arguably, small sample size. Additions have also been made to the methodology, specifically details regarding the recruitment strategy; transcription and accuracy checking; explicit statement of COREQ checklist completion (which has also been added as an appendix); explicit statement of NVivo 12's use to help facilitate coding. Details regarding ethics have been elaborated upon for further clarity.

Any further responses from the reviewers can be found at the end of the article

Introduction

Randomised controlled trials (RCTs) are widely considered to provide the highest level of evidence for an intervention's efficacy¹⁻³, with implications for participants and policymakers alike, making the optimisation of their efficiency and reliability a research priority^{4.5}. A vital factor for ensuring the RCT success is achievement of an adequate sample size, as failure to do so increases the likelihood of type-II errors, trial extension and uncertainty surrounding what might be a potentially beneficial therapy⁶. However, many trials do not achieve their desired sample size – just 56% of UK RCTs conducted between 2004 to 2016 achieved such targets⁷. While this was an improvement on 1994–2002 rates⁸, there remains a lack of understanding of optimal recruitment methods for RCTs.

RCT-participation is influenced by many variables, including perceived benefit, peer encouragement, the opportunity to help others and effective trial communication^{8–10}. With respect to the latter, despite technological advances in communication methodologies and an ever-growing internet engagement (e.g. through social media), there remains a lack of research on the advantages and disadvantages of utilising technology in RCT recruitment – an issue which has been identified as an important research priority⁵, given its potential to transcend more traditional trial communication methods and allow for a wider range of participants in RCTs.

Extant literature suggests that a considerable proportion of researchers may be using technology in some capacity in their clinical studies and along with its development, more opportunities to create and maintain effective contact with potential participants arise¹¹. Further to more traditional methods of recruitment (e.g., post, advertisement, and posters), a vast range of digital resources are available for use in RCT recruitment, from social media and direct-messaging to automated screening of health records and data mining. However, a disproportionate number of studies assess some technological methods (e.g. social media) as opposed to others¹². Moreover, as there is great variety in the types of technology researchers may use in recruitment, assessing the effectiveness of each may be challenging.

Though there is a limited body of research in this area, Rosa $et \ al.^{13}$ found that trial efficiency, lowered costs, enhanced stakeholder-involvement and reaching a more diverse range of participants are all advantages of using technology in RCT

recruitment. Conversely, disadvantages may include privacy issues, inadequate infrastructure, exclusivity to particular populations and a lack of human interaction¹⁴. In practice, however, there is little evidence of the true barriers and facilitators to using technology in RCT recruitment¹³. Thus, the aim of the current study is to further explore the advantages and disadvantages of using technology in the RCT recruitment process and to facilitate deeper understanding of the pertinent issues. This may have future implications for the development of recruitment strategies for the benefit of both researchers and members of the public who are eligible to participate in such RCTs.

Methods

Ethical statement

This research was reviewed by and granted approval by the NUI Galway Research Ethics committee (Ref: 2021.05.007) on May 7th, 2021. Formal written consent was obtained by all research participants in this study. All participants were made aware that their data would be pseudonymised for report in the study and that the data would be made available as an open resource (see *Data Availability*). Participants were also made aware that they could terminate participation at any time.

Study design

A series of one-to-one semi-structured interviews were conducted with consenting researchers who had previously been involved in the recruitment process for a RCT, with the aim of exploring perceived advantages and disadvantages of using technology during trial recruitment. An inductive, interpretive qualitative approach was used to explore participants' perspectives, identify clear and relevant themes¹⁵ and gain insight into participants' experiences of topics lacking deep understanding¹⁶. Following analysis of the interview data (Phase 1), a focus group was conducted with a Public & Patient Involvement (PPI) panel (i.e. people with lived experience of a particular condition as consultants throughout the research process^{17,18}) in order to review the findings and further elaborate on concepts and themes, consistent with their experiences of being recruited for RCTs (Phase 2). As focus groups facilitate added depth of shared ideas through interactive discussion^{19,20}, a PPI focus group was implemented in this manner to both add depth and richness to the analysis and interpretation of findings and provide a means of ensuring trustworthiness of the Phase 1 data9. All data were collected and analysed via reflexive thematic analysis; thus, facilitating the inductive approach through an iterative, recursive process of identifying, analysing, forming and revising themes from the collected data²¹.

Materials

Zoom²², a cloud-based videoconferencing application, was used to conduct and record the interviews and focus group. The semi-structured interview guide was developed in light of findings from the PRioRiTy I study⁵ by a group of researchers experienced in the recruitment process (see Table 1). Questions within the guide were designed to both facilitate elaboration on experience(s) of using technology in recruitment, as well as to seek recommendations for its use in the future. Following its development, the interview guide was pilot-tested with the research team. In an effort to reduce bias, PPI members

Table 1. Semi-Structured Interview Template.	
1. What kind of recruitment strategies have you used where technology has been involved?	
(b) Is there one technological method(s) you'd recommend over another?	
2. Why was it decided to use technology to recruit?	
3. Was there anything about using this technology that you found advantageous or facilitated	recruitment?
4. Was there anything about using this technology that you found impeded recruitment?	
5. Were you able to achieve the sample required?(a) If not, have you ever worked on study without technological support in recruitment whe achieved? (a.1) Why do you think that is?(b) Do you think using a recruitment strategy without this technology would have yielded a	re the required sample was similarly sized sample?
6. What was your retention rate like for this RCT?(a) Do you think your recruitment approach impacted retention?	
7. Can you describe the cohort you were seeking to recruit with regard to their age group, sex socioeconomic status?(a) Do you think that the type of cohort you were recruiting (e.g. with respect to age, gender status had an impact on your recruitment process)?	and educational or , ethnicity, socioeconomic
8. Did the use of technology cost anything above and beyond that of not using it with respect (a) Do you think it was worth these costs?	to resources?
9. Has COVID impacted the way in which you think about recruiting for RCTs?	
 How might technology be used to improve recruitment for randomised controlled trials in (a) What do you think of the real-world implications of such recommendations (e.g. with reaccessibility and appropriateness). 	the future? spect to feasibility,
11. Do you think there are any other potential barriers to using technology for recruitment the discussed?	at have not yet been
12. Do you think there are any other potential factors that can enable recruitment that have ne (a) Any final thoughts/comments	ot yet been discussed?

involved with the Phase 2 focus group were not included in the development or pilot-testing of the interview guide. A completed COREQ checklist is presented in Appendix A.

Procedure

Previously funded RCTs, conducted in Ireland since 2006, were identified by database searches of Ireland's four major funding bodies. Direct contact was then made to researchers on 50 identified trials, inviting participation via email. Specifically, in each of the 50 trials, both the lead author and corresponding author (if different individuals) were identified for contact, based on the commonality of these individuals having the most input into a study's administration. If these individuals were not involved in the recruitment process, they were asked in the email to advise who best to contact in this context (given that not all published articles include a breakdown of author involvement). In the event of non-response (i.e. after two weeks, a second email was sent.

In addition, a descriptive flyer was circulated on social media using Twitter, Facebook and LinkedIn, with a contact email made available for interested parties. Potential participants were provided with information about the study in the email, as well as a participant information sheet and consent form. The information sheet was also forwarded to eight clinical research facilities, based in Ireland, for further circulation. Eight individuals expressed interest in participating in an interview, one of whom was unable to take part due to scheduling issues. All of the remaining seven (N = 7) individuals were eligible according to the criteria that the individual was a consenting (1) researcher (2) previously involved in RCT recruitment, (3) based in Ireland; and participated in the virtual audio-recorded interviews (conducted between July-August 2021; mean duration 28 mins). Data were coded, analysed and themes identified (Phase 1)⁴.

Consistent with COREQ checklist guidelines (see Appendix A), it is notable that author LAM, a female medical student, who was awarded this particular research position as part of the HRB-TMRN Summer Student Scholarship, conducted the interviews and focus group. LAM had previously conducted qualitative research, as part of a project funded by the NUI Galway School of Medicine and received further training in qualitative methodology for this particular role, including interviewing skills and thematic analysis by the senior researchers involved in the current study (i.e. SMH and the PI, CPD). Notably, data were co-coded and co-analysed along with MO and CPD, both of whom are well-experienced in qualitative research. Participants recruited for the study were not known to the researcher conducting the interviews; and, aside from email correspondence to arrange suitable interview times, first meeting between researcher and participants was at the time of interviews. Upon meeting, LAM engaged with participants to build rapport. In addition to information provided in the participant information sheets, a brief rationale for the research was also relayed prior to the interview proper. Interviewer characteristics (e.g. biases, assumptions, interest, etc.) were not engaged prior to the interviews, so as to not bias participants' accounts.

In Phase 2, a PPI focus group (N= 3; duration 52 mins) was conducted in light of findings from the previous phase, in order to further explore and elaborate on concepts and themes. PPI panel members were first presented the semi-structured interview questions asked of the interviewees as a means of introducing the concepts of interest and to further explore consistency between perceptions of the panel (i.e. former/potential RCT participants) and then actual Phase 1 responses, particularly with respect to advantages and disadvantages of using technology in the recruitment process. The PPI panel was asked the questions prior to being advised of Phase 1 responses and themes, in order to limit any potential bias.

Data analysis

Transcribed data were checked by two researchers for context and errors through reading of the transcripts along with audio, followed by a subsequent re-reading (LAM and CPD). NVivo 11 was used to help facilitate coding. Consistent with Braun and Clarke²³, (1) data familiarisation began during data collection and involved the reading and re-reading of the interview transcripts, accompanied by observational note-taking; (2) researchers then systematically generated concise, meaningful codes; (3) identification of patterns from/within the data, both of which were discussed by the researchers; and (4) themes identification; prior to completion of the iterative review and refinement of these themes against the transcripts, for the purpose of ensuring their credibility24. Thus, trustworthiness of the data and credibility of the findings were ensured in multiple ways, such as triangulation (e.g. multiple observers/observations and analysts/analyses), researcher immersion in the data for ensuring rich descriptions, as well as consultation of the PPI panel.

Results

Table 2 presents relevant demographic information for each interview participant $(N=7; 6f, 1m)^{25}$. PPI panel members (N=3; 1f, 2m) consisted of individuals living with a chronic illness who have either taken part or have been previously eligible to participate in a RCT relevant to their illness. Pseudonyms were used for all participants in this study.

Phase 1

Consistent with the aim of this study, a number of advantages and disadvantages of using technology in the RCT recruitment process were identified, as presented in Table 3. With respect to the study's other aim – that is, to facilitate deeper understanding of the issues pertinent to using technology in the recruitment process, findings from the reflexive thematic analysis identified one over-arching, super-ordinate theme; two themes and five sub-themes (see Figure 1).

Super-ordinate theme: "Tech is just a medium so that you can reach more people"

The overarching theme – or message – of the findings was that the use of technology in recruitment is a means of reaching as many people as possible to ensure the sample necessary for the study. If a certain technology could facilitate that and was feasible to utilise, then it would be engaged. This superordinate theme is further explored throughout the analysis regarding the themes and sub-themes.

Pseudonym	Research Level	Trial Design	Field of Study	Participant Age Group	Technology Used
Anne	Post-Doc	Pilot RCT	Endocrinology	Young Adults (18-25)	 Phone Calls Postal service
Beatrice	Senior Lecturer	Pilot RCT	General Practice	Not specified (18+)	EmailsMailing ListsSocial Media (Twitter)
Craig	Post-Doc	RCT	Mental Health	School-age children	EmailsPhone callsParticipant Recruitment Websites
Deirdre	PI	RCT	Mental Health	Older adults (65+)	• Social Media (Facebook, Twitter)
Eleanor	PhD	RCT	Neurology	Not specified (18+)	Videoconferencing (Zoom)Phone Calls
Frances	Lecturer	RCT	Breast Cancer	Not specified (18+)	 Phone Calls Videoconferencing Emails
Grace	Clinical Research Coordinator	RCT	Intensive Care	Not specified (18+)	Phone Calls

Table 2. Demographic information.

Medium	Advantages	Disadvantages
Overall Use of Technology	 Can be less resource-intensive Allows more targeted recruitment on a larger scale Preferable to face-to-face contact in certain contexts 	 Participants may still want to meet in person Requires IT skills of both participant and researcher Data security considerations Issue of representativeness
Phone Calls	 Quick, direct way to contact potential participants and check their status in the 'sign-up process' More personal 	• Requires initial contact by other means be it clinical or otherwise
Emails	 Can attach relevant documents (e.g. information leaflets) Cheap, fast Allows reminders to be sent Recipient has more agency about when to engage with the information 	 Less personal and thus easier to ignore Cold-emailing tends to be less successful than when a relationship has been established/a gatekeeper is used Requires more IT skills than traditional phone calls or post
Post	• May be preferable to participants who like to physically read information on paper	 Less frequently used nowadays, especially for younger people. Many unfamiliar with going to post office Relatively cheap Slower
Videoconferencing	Most effectively simulates a face-to-face meeting	Platforms are of variable qualitySome IT skills required
Social Media	 Quick and cheap Allows widespread communication to large numbers of people Specific social media sites can be utilised according to demographics desired 	 Needs to be shared by those with larger following to reach required participants Can be too widespread, difficult to reach specific populations May exclude certain cohorts e.g., older people who may be less engaged with social media Less room for information - character limits etc. Can be considered less credible in some cases
Mailing Lists	 Allows researcher to target a very specific desired population Those signed up are more likely to be interested in research 	• Require trust/relationship with a relevant body to allow access to mailing list
Participant Recruitment Websites	 Monetary incentives are provided for participants Can recruit large numbers in short space of time 	• Expensive for researchers and create disadvantage among researchers who cannot afford to use them

Table 3. Advantages and disadvantages of using technology in the RCT recruitment process.

Theme 1: Technology is used if and when the benefits outweigh the costs

The first theme addresses how technology is utilised to facilitate recruitment when its use is perceived as convenient and inexpensive with respect to financial cost, time and other resources to both researchers and patients. Data from the interviews suggest that technology was not used just for the sake of using it, but with the intention of meeting logistical needs of a particular study's recruitment process.

"I'm a little bit wary when people say that they're going to use technology as a way around things,

whereas technology, I think, is a facilitator for things that are going to work or not work in the first place" - Craig

This is a key finding of the study, which emphasises the utility of technology as a tool that has potential to enhance the recruitment process but cannot necessarily eliminate all existing barriers. Use of technology must be relevant in the context of meeting an unmet need in research recruitment and should never make this process more difficult.

Within this theme, two further subthemes were identified (i.e. 'Avoidance of resource-intensive methods', 'Proclivity



Figure 1. Outline of themes identified.

for easy and fast'). The tendency towards avoiding 'resourceintensive methods' was noted throughout the interviews. Cost and budget considerations tended to influence what was feasible in the recruitment process. The cost of purchasing a new technology is cited as a barrier to its use, as recruitment may not be the first priority in the allocation of funds. Likewise, cost to the potential participant and its impact on the decision to take part was also an important consideration. For example:

"You wouldn't often think about allocating a lot of resources to the recruitment strategies. ...We'd be relying on existing networks... If it's something that takes a lot more on their part, I'd be less inclined to use it... it comes down to the time and cost for the participants and how likely it is that they're going to engage with us." - Beatrice

Aside from monetary cost, time was a resource impacted by the choice of recruitment method. The use of technology to increase reach, with respect to advertising recruitment, saves time in the recruitment process. Where a large sample of participants is required in a timely manner, technological methods can provide a quick means to access the patient cohort needed:

"Time-wise, I think, casting a wider net - which I could do with tech – it saved me time compared to... recruiting much more locally, where you're dependent on a small few eligible samples, where the hit rate is much lower." - Craig

'*Proclivity for easy and fast*' was the second subtheme, which captured the essence of technology, with respect to its general purpose and strength of meeting the needs of the research study. Thus, the identification of this theme is somewhat unsurprising. The use of existing technological resources for

the purpose of convenience became evident throughout the interviews – essentially, 'use what's there':

"Just think about the budget you will need. If you're going to do things virtually... we just used the resources that we have. Your sponsor can just have... already... some platforms and nowadays with video calls, it's like all the companies have some sort of platform already arranged." - Frances

Despite budgetary concerns surrounding use of technology, less resource-intensive 'technological' methods were utilised by researchers (e.g. phone calls and emails). These were described as a cheap and convenient means for contacting interested parties:

"Because it was email, I suppose, it's not something that we needed to train ourselves. Skills-wise, we could do it; time-wise, we could do it; and it didn't cost anything, financially, or in terms of time. So, I don't think it was a bigger drain on... it's very low resource way for me." - Beatrice

Use of technology may also facilitate ease with respect to reducing the time that prospective participants may have to spend in face-to-face environments. Technology can be used to minimise such contact, while simultaneously reducing travel costs. For example, use of videoconferencing allowed some researchers to meet with potential participants and engage in the informed consent process virtually:

"It's more comfortable for the patient – just doing this step online; they are going to come one more time to the clinic for the baseline. But, at least, they don't need to come for the informed consent and they can just be comfortable at home... not having to drive here." – Frances This concept also has consequences for the 'scalability' of technology, which potentially allows for a larger pool of participants to be reached, thus making technological methods more convenient in comparison to traditional methods in large-scale studies. Where a specific target population is required, technology may just be 'easier':

"Whether we would have got the spread [without technology]... I don't think we would have. We would have been relying more on a convenience sample. So, I definitely think that this gave us the scale to reach more practices." – Beatrice

Theme 2: Success of recruitment through technology depends on the nature of the study

The second theme describes the variability of recruitment methods required, depending on the type of study being conducted. It acknowledges the vast differences among individuals and their preferences for engagement with a research study. Three subthemes were also identified: *recruitment strategy should be dictated by the cohort; gatekeeping and referrals; and ethical considerations.*

It was clearly recognised throughout the interviews that the '*recruitment strategy should be dictated by the cohort.*' This subtheme recognises the diversity among potential participants and, consequently, their variable engagement with technology. Age featured as a point of comparison regarding levels of engagement with various means of communication. For example, it was suggested that more mature cohorts might prefer less technologically advanced methods communication (e.g. mail/post), whereas younger cohorts more technologically advanced methods (e.g. email, text and video-calls). According to Grace:

"It's the older generation versus the younger generation; and the older generation is what we still see a lot of in our cohort of patients. So, even though we have emails to say it to some people... they like the written printed-out version"

The preferred method of communication, be it through advertising the study, providing an information sheet or even obtaining informed consent, is an important point of consideration when targeting a specific group of participants. Similarly, the variability of IT skills among potential participants should also be considered. For example, when technology is a feature, there are often some participants who require more support in accessing or using the technology:

"One of the initial problems was the technology itself and technical issues about setting up, but then usually their children or grandchildren will be there to set them up with the video call – but then if they're alone, we wouldn't get a lot of inputs out of them." – Eleanor

This reflects an important consideration regarding the first theme – while it is preferable to choose an efficient method with respect to resources, ease and speed, the recruitment method also needs to be accessible to the target cohort or it will, inevitably, fail to recruit the desired cohort.

Even though technology can be a barrier for some potential participants, if there is a will, there is often a way with respect to finding the supports needed to engage the technology (such as in Grace's example of having aid from family members). However, it must also be acknowledged that support is not always feasible. Thus, technology is a hurdle to some.

Whereas thinking and discussion often revolved around the level or type of technology used in the recruitment process, the manner in which recruitment was *targeted* was also an important factor. Largely consistent with the super-ordinate theme that '*tech is just a medium so that you can reach more people*', the concept of targeted advertisement is vital for considering how best to reach a *particular* audience, especially if the required cohort is 'specific' or 'niche'. In this context, the interviews suggested that regardless of whether or not 'technology' is utilised as a means for recruitment, the method is only useful if it reaches the cohort suitable for the study:

"Social media feeds and stuff are all based on how much somebody's using them – if if someone's sharing it and they've only got 20 people on it, that's a disadvantage. It's more advantageous if it's picked up by somebody with a high [number] of followers ... Then the advantage is there; especially... if any of the [health service] national groups pick it up and forward it on, you get much better reach. So, it's really dependent on who's actually posting it, whose news feed and how it then gets marketed; and that's the thing as clinicians, we aren't taught marketing... and that's what you're basically asked to do." - Deirdre

Notably, just as the cohort *should* dictate the recruitment method, the recruitment method *could* potentially dictate the cohort. As alluded to above, the way recruitment is advertised and conducted could impact the diversity or representativeness of the sample, which might be otherwise unaccounted for within the recruitment design.

"Depending on how [technology's]used, you may only get the people who are very engaged; so, if it's something like Twitter or social media, thinking about how representative those people are and how equal is it for everyone to be able to engage with that technology." - Beatrice

The second subtheme, '*Gatekeeping & Referrals*', addressed the concept of a 'gatekeeper' and their utility in research recruitment. The term was aptly defined by one of the researchers interviewed:

"It's even more integral now with GDPR because you can't access people's information... so, a gatekeeper is a person who has access to that information and can share your study information with people, but they're not... abusing their access to the information – they're not pressuring the person." - Beatrice This concept emerged across the interviews and illustrated a sense of trust that aided study participation. Providing information about a study via a trusted individual or organisation (e.g. study advertisement or a participant information sheet) was perceived as a successful strategy:

"I think having a credible source be the messenger for your study is really valuable. Whether that also involves technology, that's totally possible; but, I think having other GPs endorse my study or having them kind of promote the study or suggest it to their colleagues carries more weight than me trying to do so."- Beatrice

This sub-theme also suggests that relationships and networking similarly provided a solid foundation for researchers to gain access to recruit from particular groups, irrespective of using technology:

"It was a little bit of a nightmare...And then the only reason that I was able to recruit... after that is because I knew somebody who knew somebody." - Craig

The final subtheme elicited was *ethical considerations*, which encompassed the systemic structures (e.g. committees) underpinning ethical research as well as the increasing concern for data security within society. Research ethics committees, while crucial for maintaining the safety of participants and integrity of research, emerged as a source of frustration for researchers:

"The recruitment strategies were... it's slightly different in each hospital, because the ethics committees required different strategies. So that was a bit frustrating, but we had to do it." - Anne

Likewise, ethical approval requirements were perceived as a barrier to recruitment:

"The more interesting the sample, the more difficult it was to access them and I suspect that's the same in other areas too. It certainly complicated recruitment further. So, I honestly think that the ethics committees, generally speaking, make it unnecessarily harder. I know that there are certain laws that they have to abide by and certain principles, but I find that they're not very pragmatic and that can hinder recruitment." - Craig

Notably, it can be argued that this quote also relates back to the concept of 'recruitment being dictated by the cohort, in the sense that 'the more interesting the sample, the more difficult it was to access them' suggests that specific populations that may be the target of a study may require a different strategy for recruitment – less so an issue of whether technology would or would not be advantageous, rather an issue of ethical consideration – particularly, if the 'interesting sample' was deemed vulnerable.

Notwithstanding the obstacles encountered in gaining ethical approval, data security was recognised as a vital aspect of research and recruitment. Respect for participant data and acknowledgement of the trust participants have in researchers to protect data responsibly was evident among the interview data:

"There should always be a backup, so that you can protect the integrity of the study and the integrity of the data that you're collecting... for the patients' privacy... We need to protect their data." – Grace

Data protection and security is a matter of public concern that has gained considerable attention in recent years. Trust is required in the recruitment process for patients to consent to sharing personal and often sensitive data for the purposes of research. Clarity should be provided when enrolling participants to a trial with regard to how their information will be collected, stored and used. This is of utmost significance in the context of technological methods of recruitment, both in the sense of how data are collected securely at this early stage of a trial and in setting the precedent for further trial communication. Robust processes need to be demonstrated to avoid distrust, which could influence retention and completion of trial enrolment.

Phase 2

Following presentation of the interview guide to the PPI panel and subsequent discussion of interview results, the PPI focus group findings were largely consistent with those of Phase 1 and provided further perspective and depth to the identified themes. The first theme, *Technology is used if and when the benefits outweigh the costs*, was addressed by the panel, who agreed that technology should be used when it is convenient and when the cost is acceptable. In this respect, the panel paid particular focus to emphasising that *Avoidance of resourceintensive methods* should, as recommended in Phase 1, extend to consideration of potential participants' resources (be it time, financial or otherwise), as recruitment strategies can either limit cost or be costly. For example: –

"And like I live over in [county] and I was driving over 45 miles over to Galway If I could have done it over the web, like, it'd be great." – Calum

Likewise, according to Ben:

"You'll have a certain cohort of people that, say they've lost time from work or they've lost time from their family... If you have somebody that has chronic fatigue, every hour that you have to do something, that's precious. So, you're going to focus on your family or you're going to focus on your kids, rather than on research. So, how do you compensate for that?"

Thus, in order for recruitment to be successful, methods of accommodating participants to engage with research should be understood and engaged by the research team – ultimately influencing both recruitment and study design.

The concept of '*Proclivity for easy and fast*', was understood by the panel to mean that researchers choose the method which is easiest and fastest for the research team. For example, by choosing tools that are already accessible via their institution rather than exploring other options:

"I'm at the stage now where if I was told by my consultant that I was going to be called on [specific videoconferencing platform] I would say, 'Do you know what? I'm fine. I'll wait until I can come and see you in person' – I can't be doing this. It's exhausting – especially if you have any kind of extra ability issues going on at the same time. It's just – no." – Arlene

Like researchers, patients exhibit a preference for recruitment methods that are convenient and least disruptive to their lives. However, these methods may not always be congruent between researchers and their desired sample, which is consistent with discussion of the second theme, 'Success of recruitment through technology depends on the nature of the study' – particularly, the sub-theme of 'recruitment strategy should be dictated by the cohort'. For example:

"I think that whoever is designing whatever trial – they need to know what their patient cohort uses, not what they use." – Ben

In a cumulative sense, it must also be acknowledged that, through presentation of the subtheme regarding ease and speed, there may be the assumption by the PPI that researchers only consider these issues from their own perspective, where it could well be the case that the researcher also thinks that such a method would facilitate potential participants. Though the suggestion of uniformity may not necessarily facilitate a 'happy medium' between researchers and potential participants, the concept of 'uniformity' introduced here may more so refer to the need for researchers and potential participants to 'get on the same page' - in terms of congruency, with respect to what methods work best and for who, as addressed above. Thus, it may be the case that a mixed-approach for utilising technology, via recruitment through multiple avenues, might work best in this context. Such a mixed-approach for designing the recruitment strategies was favourable among the panel, for example, in recognition of the natural heterogeneity of individual patients and patient groups:

"I think you just need to have a balance. I think you can't go one way or the other... if you want to include everybody or try and hit as many people as you can." – Arlene

Consistent with this diversity of patient preference, the panel described methods which they themselves preferred, with some having a tendency towards the written word (sometimes facilitated through technology), while others appreciated the growing opportunities technology provides. According to Calum:

"I suppose like a few paragraphs... text-based stuff... so you can read it – something physical, tangible even if it is a PDF document – and you can kind of mull over it and you know [researcher]'s email address was in it. So that's the way I'm quite happy working with."

According to Ben:

"If I have to go somewhere, it's harder and harder to leave these four walls. It's incredible, what's happened now is that I can access stuff that's in Switzerland and the United States... I'm not geographically bound anymore, which is incredibly liberating."

This issue of diversity is additionally interesting to consider because even when putting participants before the ease, speed cost or even preferences of the researcher(s), there remains no guarantee that all individuals within a targeted cohort will prefer the same means of communication as others. Levels of engagement with each medium will vary, as will access to technology and infrastructure, regardless of the cohort or, in the context of the PPI focus group, patient group one belongs to. This point may also have some implications for the likelihood of finding uniformity (as addressed above) between researchers and potential participants.

Moving forward, relationships and networking were also emphasised as being integral to accessing a desired patient group. Similar to the 'gatekeeping and referrals' subtheme, value was placed on having a trusted person or body to connect the patient with the researcher:

"We have a forum that does a whole lot of wonderful things locally and they have very tight networks, so, if you have access...to one person – then they can spread it throughout their network very quickly, and that will aid your recruitment." – Ben

Likewise, according to Calum:

"So long as you've someone trustworthy... It's not just some kind of random email you open up that could be God knows what." - Calum

Upon delving further into the feasibility of networking, the contrast between the worlds of researchers and participants emerged as a potential barrier to accessing local groups. The PPI panel felt that both entities seem to occupy different spaces, which can make the process of forging connections between the two more difficult:

"Researchers live in their bubble and everyone else isn't in their bubble... which is quite unusual because we need each other. We need the researchers, just as much as they need us; and as patients, we want to hear from them. We want to be talking to them. We would like to be influencing the studies that they're doing; yet, they don't seem to have access... researchers need to get out of their offices and... they need to be part of that community."" – Ben Nonetheless, the panel was also understanding of researchers' perspectives, acknowledging that ethical considerations from Phase 1 findings may also play a role in their decision-making – arising as another potential barrier to networking. This is an aspect of research culture the panel seemed to feel was old-fashioned, which could be improved upon to aid recruitment. Likewise, consistent with the dynamic nature of technology, recruitment strategies evolve over time; and so, there may be a need to shift the status quo within research to keep up with this evolution. This panel viewed such a shift as a significant challenge:

"I think there's a fear amongst researchers that they will be doing something that's unethical... There's a reluctance amongst some people to change the way they've done it historically, 'this is the way we've always done it, and this is the way we've done our trial' and to change that mindset is a real challenge." - Ben

Discussion

Interpretation of results

Upon analysis of Phase 1 data, one overarching theme was identified: 'Tech is just a medium so that you can reach more people'. This represents the concept that the utility of technology lies predominantly in its ability to scale up the recruitment process and may not require any great consideration beyond this utility, except for issues relating to cost and the nature of the study itself. Notably, the term 'technology' was presented to participants in a purposefully vague manner, so as to not bias their views. It was hoped that some would endeavour to provide their definition to better conceptualise technology in this context. Though no such definition was provided, a series of different technologies were discussed mostly in terms of web-based applications (e.g. email, social media, text-messaging and video-calls). Phone calls and post/mail were also discussed, but not necessarily in terms of 'technology'.

Herein lies an interesting consideration – technology evolves over time. In research 60 years ago, the telephone might be considered among peak technologies. Before that, post/mail might be considered as such as well. Indeed, something as simple as the printed word was once considered technology. Just because time has elapsed and technology evolved does not mean that these means of communication have ceased to be 'technology'; for example, given that people still use post/mail and the telephone to communicate. In the results it was found that some people prefer these older technologies. Thus, if the aim of recruitment is to reach as many people as possible to achieve one's sample size, the age or format of a technology does not matter, just if it remains relevant and useful for achieving the necessary sample.

Two central themes were addressed within the superordinate. The first, 'Technology is used if and when the benefits outweigh the costs', presents the idea that availability of resources impact how technology is implemented to recruit for RCTs and also how researchers take preference for the method which is easiest and fastest for both themselves and the potential participants. The second, 'Success of recruitment through technology depends on the nature of the study', addressed the contrast between different study designs and procedures, as well as how this influences the recruitment process. Within this theme, the variation in participants' engagement with technology, the pivotal role gatekeepers and referrals play in successful recruitment and the impact of ethical considerations on recruitment were also discussed. In summary, choosing a successful method(s) for recruitment requires consideration of the specific characteristics of the target cohort; and establishing networks that link participants and researchers is often vital, regardless of 'technology'.

Phase 2 data corroborated these findings, while also shedding light on the results from a patient perspective. The panel found value in the themes' message and re-emphasised that resource considerations should extend to the cost a potential participant undertakes as a research participant. The necessity for a culture shift in research was also highlighted to bridge the communication gap between researchers and patients. Moreover, the heterogeneity of individual patients within a group may influence their engagement with a specific technology or method of communication. Willingness to incorporate multiple methods to reach a larger subset of desired patient groups may be a key step in combating this issue.

The findings highlight a number of similarities and contrasts with extant, though limited research. Of note, the current research was particularly consistent with the qualitative portion of a study by Blatch-Jones *et al.*¹⁴ which argues that the availability and acceptability of strategies across different patient groups requires consideration as does the trade-off between utilising digital strategies and more personal interaction.

Regarding the common use of specific technology, participants mentioned phone calls, emails, social media, participant recruitment websites, post, videoconferencing and mailing lists, which were consistent with a recent review by Frampton et al.12 of 105 RCTs. While studies have cited use of an electronic health record to allow automated identification of potential participants¹⁴, this was not an approach that arose from our study - perhaps due to incompatibility of current electronic health records for this type of use in the Irish context. With that, extant literature has not been able to comprehensively compare the efficacy of individual methods of digital recruitment; instead, focusing on particular methods in isolation, such as social media²⁶. While two participants in the current research noted the utility of social media in reaching a larger number of widely distributed potential participants, the need for a link or 'gatekeeper' to successfully aid this method was also emphasised.

Limitations

Though a number of interesting findings were yielded, some specific limitations must be addressed. First, the sample size can be argued to be small. Overall, eight individuals expressed interest in participating, in which the irony of the situation is acknowledged - wherein there was difficulty in recruiting for a study on recruitment. It is perhaps the case that the adequacy of the recruitment strategy used in this study's approach requires consideration with respect to the thoroughness of efforts to make contact with potential participants. However, efforts made were as thorough as possible within the time-frame of the summer studentship (see again, Footnote 1). Though it can be argued that the data yielded from this 'small' sample was qualitatively rich, it can also be argued that, despite such perceived 'richness', this remains inadequate for data saturation. On the other hand, there is indeed another argument to be made that saturation (or, at the very least, consistency) was reached with respect to the observable repetition of themes. For example, in the cases of the five sub-themes identified (which were thematically organised into two themes and one superordinate theme), one was discussed by 5 of 7 participants (i.e. Proclivity for easy and fast), three by 6 of 7 participants (i.e. Avoidance of resource-intensive methods; Gatekeeping & Referral; and Ethical Considerations) and one by 7 of 7 participants (Recruitment strategy should be dictated by the cohort). Moreover, the sample attained is consistent with recent research⁹ which utilised a similar dual-phase approach with one-to-one interviews conducted prior to a corroboratory focus group. Furthermore, it can also be argued that sample size in qualitative research is a subjective consideration - the priority should be achieving a sample that is not too large that it obstructs the process of deep analysis, consistent with the notion that the more relevant information available within the sample, the lower the number of participants needed^{27,28}. Thus, in the current research, it is reasonably argued that the sample size is justified on account of the richness and extensive analysis of the available qualitative data.

Furthermore, as addressed above, a standardised definition of technology was not presented to interviewees, which allowed for a wider interpretation of 'technology' in this study. It is possible that this limited some level of comparability between interviewees' responses. However, consistent with discussion regarding the super-ordinate theme identified, it can also be appreciated how this added depth in allowing each researcher to individually contemplate what technology means, in general, as well as in the context of their own trial.

Conclusion

The results of this study contribute a deeper understanding of the pertinent issues for using technology in recruiting potential participants for RCTs. Key factors that influence the choice to use technology were highlighted (e.g. cost, ease, speed/time and the nature of the study), with particular emphasis on technology being no more than a medium to achieve research aims. The need to adapt recruitment design to suit target cohorts is of significance. The current research arguably succeeded in answering Healy *et al.*'s⁵ PrioRiTy question 10: '*what are the advantages and disadvantages to using technology during the recruitment process?*', through providing a thorough list (again, Table 3) and elaborating on technology as, simply, a medium for communication and both advantages and disadvantages to its use as contextual –likely to be utilised when easy, fast, lacking resource-intensiveness and consistent with the nature of the study.

In keeping with the qualitative nature of this study, the findings cannot be generalised to larger research populations; however, they have facilitated deeper understanding of the advantages and disadvantages of using technology in the RCT recruitment and other pertinent issues. Such understanding provides a useful starting point for further research to investigate the themes identified - for example, future research should aim to: adapt their recruitment approach to acknowledge the preferences and costs to potential participants (regardless of how 'technologically advanced' it may be) and engage with potential participants to facilitate networking opportunities for future studies. Further research into the use of technology in RCT recruitment is warranted, particularly surrounding its effectiveness when used in multiple ways or in combination with more traditional methods. In light of these findings, the identified advantages and disadvantages of using technology for RCT recruitment can be appreciated, while also acknowledging that the use of technology is a multifaceted decision in this process, integrating resource considerations, study design issues, as well as potential participant preferences.

Data availability

Open Science Framework: Advantages and disadvantages to using technology in the process of randomised controlled trial recruitment. https://doi.org/10.17605/OSF.IO/67GM5²⁵.

This project contains the following underlying data:

- Focus Group anonymised final.docx
- Interviews anonymised.docx

Data are available under the terms of the Creative Commons Attribution 4.0 International license (CC-BY 4.0).

No	Item	Guide questions/description	Response	
Dom	ain 1: Research team and			
Perso	onal Characteristics			
1.	Interviewer/facilitator	Which author/s conducted the interview or focus group?	LAM	
2.	Credentials	What were the researcher's credentials? E.g. PhD, MD	N/A (Med Student)	
3.	Occupation	What was their occupation at the time of the study?	Med Student	
4.	Gender	Was the researcher male or female?	Female	
5.	Experience and training	What experience or training did the researcher have?	LAM had previously conducted qualitative research, as part of a project funded by the NUI Galway School of Medicine and received further training in qualitative methodology for this particular role, including interviewing skills and thematic analysis by the senior researchers involved in the current study (i.e. SMH and the PI, CPD).	
Relati	Relationship with participants			
6.	Relationship established	Was a relationship established prior to study commencement?	None.	
7.	Participant knowledge of the interviewer	What did the participants know about the researcher? e.g. personal goals, reasons for doing the research	In addition to information provided in the participant information sheets, a brief rationale for the research was also relayed prior to the interview proper.	
8.	Interviewer characteristics	What characteristics were reported about the interviewer/facilitator? e.g. <i>Bias, assumptions, reasons and interests in the research topic</i>	Interviewer characteristics (e.g. biases, assumptions, interest, etc.) were not engaged prior to the interviews, so as to not bias participants' accounts.	
Dom	ain 2: study design			
Theoretical framework				
9.	Methodological orientation and Theory	What methodological orientation was stated to underpin the study? e.g. grounded theory, discourse analysis, ethnography, phenomenology, content analysis	An inductive, interpretive qualitative approach via reflexive thematic analysis was used.	
Participant selection				
10.	Sampling	How were participants selected? <i>e.g. purposive, convenience, consecutive, snowball</i>	Purposive	
11.	Method of approach	How were participants approached? <i>e.g. face-to-face, telephone, mail, email</i>	Zoom teleconferencing	
12.	Sample size	How many participants were in the study?	10 completed	
13.	Non-participation	How many people refused to participate or dropped out? Reasons?	1 dropped out due to scheduling.	
Setting				
14.	Setting of data collection	Where was the data collected? <i>e.g. home, clinic, workplace</i>	Workplace	
15.	Presence of non- participants	Was anyone else present besides the participants and researchers?	Not during data collection/interviews	
16.	Description of sample	What are the important characteristics of the sample? <i>e.g. demographic data, date</i>	That all participants were involved in the recruitment of participants fro RCTs	
Data collection				
17.	Interview guide	Were questions, prompts, guides provided by the authors? Was it pilot tested?	Questions, prompts, guides were provided by the authors and pilot-tested.	

Appendix A: COREQ Checklist

No	Item	Guide questions/description	Response	
18.	Repeat interviews	Were repeat interviews carried out? If yes, how many?	No	
19.	Audio/visual recording	Did the research use audio or visual recording to collect the data?	Yes	
20.	Field notes	Were field notes made during and/or after the interview or focus group?	Yes	
21.	Duration	What was the duration of the interviews or focus group?	28 mins (mean)	
22.	Data saturation	Was data saturation discussed?	Yes	
23.	Transcripts returned	Were transcripts returned to participants for comment and/or correction?	No	
Domain 3: analysis and findings				
Data analysis				
24.	Number of data coders	How many data coders coded the data?	3	
25.	Description of the coding tree	Did authors provide a description of the coding tree?	Yes.	
26.	Derivation of themes	Were themes identified in advance or derived from the data?	Derived from data	
27.	Software	What software, if applicable, was used to manage the data?	Microsoft Office	
28.	Participant checking	Did participants provide feedback on the findings?	Yes, in Phase 2	
Reporting				
29.	Quotations presented	Were participant quotations presented to illustrate the themes / findings? Was each quotation identified? e.g. participant number	Yes	
30.	Data and findings consistent	Was there consistency between the data presented and the findings?	Yes	
31.	Clarity of major themes	Were major themes clearly presented in the findings?	Yes	
32.	Clarity of minor themes	Is there a description of diverse cases or discussion of minor themes?	Yes	

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Current Peer Review Status: 💙 👘

Version 2

Reviewer Report 28 August 2024

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Milan Bimali 匝

University of Arkansas for Medical Sciences, Little Rock, USA

Randomized controlled trials are considered "gold standard" in biomedical research. One of the reasons for failure of RCTs is their inability to accrue study participants. This manuscript covers an important aspect related to recruitment. It focuses on the use of technology in the context of participant recruitment in RCTs. This is important because the use of technology is becomingly increasingly inevitable and its application in recruitment in RCTs can be an important topic for scientific community. The manuscript is based on a qualitative approach and makes use of structured interviews with researchers involved in RCT while drawing on their perspective and opinions related to the use of technology in RCT recruitment. The analysis was done using the "Reflexive Thematic" method to identify key themes. The authors identified two themes – technology is used if and when benefits outweigh the costs; recruitment success through technology depends on the nature of the study.

I have included below my review comments:

- 1. The authors has consistently the term "adequate sample size" throughout the manuscript. The term "adequate" appears very subjective and vague. Often RCTs are powered and the sample size is statistically justified. In some studies, the sample size is justified based on precision estimates. The authors must reconsider the use of term "adequate".
- 2. The authors have used the "reflexive thematic analysis". I would suggest adding a brief description about it, so readers have a basic understanding of it.
- 3. The "Introduction" section must recognize that as of now, not all RCTs have the option of incorporating technology in recruitment. This may be especially true for studies in resource limited setting where technology is not available.
- 4. The limited sample size in this study has been acknowledged by the author. This has greatly limited the generalizability of study findings. It is also unclear whether there was a self-selection bias at play here. Was there a systematic difference in the fundamental characteristics of studies whose investigator agreed to participate in the study?
- 5. It would also be helpful if the authors delved a bit more into either the introduction or discussion on the ethical issues related to use of technology in recruitment.

Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound? Partly

Are sufficient details of methods and analysis provided to allow replication by others? $\ensuremath{\mathsf{Yes}}$

If applicable, is the statistical analysis and its interpretation appropriate? Not applicable

Are all the source data underlying the results available to ensure full reproducibility? $\ensuremath{\mathsf{Yes}}$

Are the conclusions drawn adequately supported by the results? $\ensuremath{\mathsf{Yes}}$

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Biostatistics

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Reviewer Report 08 June 2024

https://doi.org/10.21956/hrbopenres.15257.r40525

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Marie Falahee ២

University of Birmingham, Birmingham, England, UK

Thank you for addressing my comments.

Is the work clearly and accurately presented and does it cite the current literature?

Yes

Is the study design appropriate and is the work technically sound?

Yes

Are sufficient details of methods and analysis provided to allow replication by others? $\ensuremath{\mathsf{Yes}}$

If applicable, is the statistical analysis and its interpretation appropriate?

Yes

Are all the source data underlying the results available to ensure full reproducibility? $\ensuremath{\mathsf{Yes}}$

Are the conclusions drawn adequately supported by the results? $\ensuremath{\mathsf{Yes}}$

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Psychology, Rheumatology, Patient and Public Involvement in Research

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Version 1

Reviewer Report 30 April 2024

https://doi.org/10.21956/hrbopenres.15069.r36770

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Upasana Baruah

National Ageing Research Institute, Victoria, Australia

This research paper explores the role of technology in the recruitment process for Randomized Controlled Trials (RCTs) by conducting a dual-phase qualitative study involving interviews and a patient and public involvement (PPI) focus group. The study reveals an overarching theme that perceives technology as a medium to broaden participant reach. The findings emphasize the importance of considering resource costs, study characteristics, and participant preferences when incorporating technology into recruitment strategies. The discussion delves into the evolving nature of technology, addresses central themes related to cost-benefit analysis and study-specific recruitment success, and effectively integrates patient perspectives from the PPI focus group. The conclusion summarizes key findings, highlights the study's contribution to understanding technology's role in RCT recruitment, acknowledges limitations, and suggests avenues for future research, providing a comprehensive overview of the nuanced considerations surrounding the utilization of technology in clinical trial recruitment.

While the paper adeptly explores the role of technology in RCT recruitment through a wellexecuted dual-phase qualitative study, a noteworthy limitation lies in the small sample size, particularly in the PPI focus group where data saturation becomes challenging with only three participants. The authors acknowledge this limitation, but clarity on the rationale for this specific number is lacking. Furthermore, while the recruitment process is adequately detailed, inclusion of participant characteristics, such as gender, experience, or roles in recruitment, would enhance transparency and generalizability.

The paper mentions using Microsoft Word for analysis without specifying the process, necessitating clarification on whether any qualitative software was employed for data management and analysis. The data management and safety plan require elaboration for a comprehensive understanding.

The results section might benefit from a more succinct presentation of main themes and subthemes, avoiding exhaustive detail.

Additionally, the initial paragraphs of the discussion seem repetitive of the results and could be condensed to emphasize key discussion points more concisely.

Is the work clearly and accurately presented and does it cite the current literature? $\ensuremath{\mathsf{Yes}}$

Is the study design appropriate and is the work technically sound? Yes

Are sufficient details of methods and analysis provided to allow replication by others? Partly

If applicable, is the statistical analysis and its interpretation appropriate? $\ensuremath{\mathsf{Yes}}$

Are all the source data underlying the results available to ensure full reproducibility? $\ensuremath{\mathsf{Yes}}$

Are the conclusions drawn adequately supported by the results?

Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: dementia, carer support interventions, psycho-social interventions delivered remotely, mental health

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Author Response 29 May 2024 Christopher Dwyer Many thanks for your comments. We considered each in turn and have amended the manuscript in light of these considerations. Other comments and questions we answer here.

1. While the paper adeptly explores the role of technology in RCT recruitment through a wellexecuted dual-phase qualitative study, a noteworthy limitation lies in the small sample size, particularly in the PPI focus group where data saturation becomes challenging with only three participants. The authors acknowledge this limitation, but clarity on the rationale for this specific number is lacking.

Response 1: Many thanks for this comment. Please see Response 1 to Reviewer 1.

2. Furthermore, while the recruitment process is adequately detailed, inclusion of participant

characteristics, such as gender, experience, or roles in recruitment, would enhance transparency and generalizability.

Response 2: These are presented in Table 2.

3. The paper mentions using Microsoft Word for analysis without specifying the process, necessitating clarification on whether any qualitative software was employed for data management and analysis.

Response 3: The line "Microsoft Office was used" is a typo – perhaps a partial remnant of a previous version. This line has been removed from the revised version of the manuscript. NVivo 12 was used to help facilitate coding. This has been added to the manuscript.

4. The data management and safety plan require elaboration for a comprehensive understanding.

Response 4: Data collection, analysis and availability are all reported. Details regarding ethics have been elaborated upon for further clarity.

5. The results section might benefit from a more succinct presentation of main themes and sub-themes, avoiding exhaustive detail.

Response 5: We would argue that the great detail presented is necessary to fully encapsulate participants' perspectives and facilitate interpretation. We hope that Figure 1 succeeds in achieving a more succinct presentation for those who prefer such an approach and that the greater detail will cater to those who prefer a more comprehensive approach.

6. Additionally, the initial paragraphs of the discussion seem repetitive of the results and could be condensed to emphasize key discussion points more concisely.

Response 6: Though we recognise that there is some level of repetition, this is

purposeful for two reasons: first, given that the results section is on the lengthy side, it's useful to begin the discussion with a brief summary, as a reminder for readers; second, the summary provides an outline for the interpretation of results. Yes, findings were interpreted (to an extent) within the results section, but added nuance and further implications are presented in this discussion.

Competing Interests: N/A

Reviewer Report 16 January 2024

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? Marie Falahee 匝

University of Birmingham, Birmingham, England, UK

Thank you for the opportunity to review this interesting manuscript describing a qualitative study of researchers' perspectives on the use of technology to facilitate recruitment to randomized clinical trials (RCTs). This is an important area of investigation given the crucial importance of successful trial recruitment for evidence-based medicine and well-documented challenges towards achieving this.

The manuscript is extremely well written and clearly presented, and the introduction develops a strong rationale for why this research is needed. The major weakness of this study is the small sample size, though this is appropriately noted by the authors in the discussion section in relation to the adequacy of the data. However, it could be argued that saturation is unlikely to have been achieved (for example, richer insight into the relationships between equality, diversity and inclusivity issues and sample bias might be expected) and reflection on the adequacy of the recruitment approach used in this study, perhaps balanced against what is feasible within a summer studentship research project, is needed in the discussion.

Further detail about recruitment procedures in the method section would also be helpful. The authors aimed to recruit "researchers who had previously been involved in the recruitment process for a RCT." Most trials involve large, multidisciplinary research teams. Were all members of the identified trials approached, or just the principal investigator? Furthermore, recruitment is often delivered by clinical research staff who may not be identifiable as part of the investigator team. Were such staff invited to participate? More details about the characteristics of those approached to take part would be helpful for the reader, particularly in relation to their professional roles (e.g. clinical, academic, methodologist, etc.). Were those contacted also invited to identify/invite others who had been involved in recruitment?

Of those researchers who agreed to participate, it would be interesting to know (especially given

the topic of this manuscript) the recruitment approach that led to their participation, if this information is available. For example, which had been invited by the research team directly to take part in the study and which contacted the research team as a result of seeing an advert on social media / via a colleague?

Minor comments:

The results section suggests that audio recorded interviews were transcribed and checked for accuracy. This should be noted in the method section, with details of the approach taken (e.g. by whom).

It is commendable that the authors appear to have applied the COREQ checklist. It would be helpful to include an explicit statement to this effect in the main text of the methods, rather than in a footnote, and to provide a completed COREQ checklist as supplementary material.

Patient and public partners could have been involved in other aspects of the research, such as the development of the qualitative interview schedule and the development of the recruitment procedures, though this is not described.

Whilst reading the manuscript I noticed some typographical errors. For example, in Table 3: 'frquently' should be 'frequently'. There is erroneous repetition of 'With' in the second sentence of the second paragraph in the introduction.

Is the work clearly and accurately presented and does it cite the current literature? $\ensuremath{\mathsf{Yes}}$

Is the study design appropriate and is the work technically sound? Partly

Are sufficient details of methods and analysis provided to allow replication by others? Partly

If applicable, is the statistical analysis and its interpretation appropriate? Not applicable

Are all the source data underlying the results available to ensure full reproducibility? $\ensuremath{\mathsf{Yes}}$

Are the conclusions drawn adequately supported by the results?

Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Psychology, Rheumatology, Patient and Public Involvement in Research

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have

significant reservations, as outlined above.

Author Response 29 May 2024

Christopher Dwyer

Many thanks for your comments. We considered each in turn and have amended the manuscript in light of these considerations. Other comments and questions we answer here.

1. The major weakness of this study is the small sample size, though this is appropriately noted by the authors in the discussion section in relation to the adequacy of the data. However, it could be argued that saturation is unlikely to have been achieved (for example, richer insight into the relationships between equality, diversity and inclusivity issues and sample bias might be expected) and reflection on the adequacy of the recruitment approach used in this study, perhaps balanced against what is feasible within a summer studentship research project, is needed in the discussion.

Response 1: As indicated, we certainly acknowledge the small sample size perspective – indeed, a larger sample was desired. We also acknowledge, that saturation may not have been reached. However, if one is to subscribe to the perspective that saturation operationally refers to the observation of the same themes emerging, there is an argument that this study achieved that, at least to some extent; for example, in the cases of the five sub-themes identified (which were thematically organised into two themes and one superordinate theme) one was discussed by 5 of 7 participants (i.e. *Proclivity for easy and fast*), three by 6 of 7 participants (i.e. *Avoidance of resource-intensive methods; Gatekeeping & Referral;* and *Ethical Considerations*) and one by 7 of 7 participants (*Recruitment strategy should be dictated by the cohort*). With that, the limitations section has been amended to reflect such further considerations as well as the points made in the reviewer's comment.

2. Further detail about recruitment procedures in the method section would also be helpful. The authors aimed to recruit "researchers who had previously been involved in the recruitment process for a RCT." Most trials involve large, multidisciplinary research teams. Were all members of the identified trials approached, or just the principal investigator? Furthermore, recruitment is often delivered by clinical research staff who may not be identifiable as part of the investigator team. Were such staff invited to participate? More details about the characteristics of those approached to take part would be helpful for the reader, particularly in relation to their professional roles (e.g. clinical, academic, methodologist, etc.). Were those contacted also invited to identify/invite others who had been involved in recruitment?

Response 2: The following has been added to the methodology: "Specifically, in each of the 50 trials, both the lead author and corresponding author (if different individuals) were identified for contact, based on the commonality of these individuals having the most input into a study's administration. If these individuals were not involved in the recruitment process, they were asked in the email to advise who best to contact in this context (given that not all published articles include a break-down of author

involvement). In the event of non-response (i.e. after two weeks, a second email was sent." With that, we recognise that in cases where no response was received, contact details could have been found for other researchers from the study and an email sent querying who most appropriate to contact in light of the null response. However, such activity would have been time costly and as the reviewer correctly identified with respect to what is feasible within a summer studentship research project, such activity was not engaged.

3. Of those researchers who agreed to participate, it would be interesting to know (especially given the topic of this manuscript) the recruitment approach that led to their participation, if this information is available. For example, which had been invited by the research team directly to take part in the study and which contacted the research team as a result of seeing an advert on social media / via a colleague?

Response 3: Many thanks for this comment – it is certainly an interesting and very relevant point. Unfortunately, we do not have data on this. Moreover, though it would certainly be interesting, we're not entirely sure how much we could take from it, interpretation-wise, given it being based on only 8 occurrences (including the participant lost to scheduling). However, should future research further investigate this topic, for example, via survey – where a much larger sample size would be required, inclusion of such data collection is certainly recommended.

4. The results section suggests that audio recorded interviews were transcribed and checked for accuracy. This should be noted in the method section, with details of the approach taken (e.g. by whom).

Response 4: Such discussion appears in the data analysis section of the methodology. The approach is now described and the researchers who checked for accuracy identified.

5. It is commendable that the authors appear to have applied the COREQ checklist. It would be helpful to include an explicit statement to this effect in the main text of the methods, rather than in a footnote, and to provide a completed COREQ checklist as supplementary material.

Response 5: Though we have maintained the footnote, given that it facilitates sufficient detail (and would perhaps divert the logic elsewhere placed, we have included a brief note about this in the methodology as well, with the completed COREQ checklist added as an appendix.

6. Patient and public partners could have been involved in other aspects of the research, such as the development of the qualitative interview schedule and the development of the recruitment procedures, though this is not described.

Response 6: Although we certainly see the value of engaging PPI for the development of interview schedules such as this, the focus of our rationale for the interview guide was development in light of both findings from the PRioRiTy I study⁵ and, specifically,

a group of researchers experienced in the recruitment process (e.g. with respect to theory, administration and other background activities). Moreover, PPI (especially those involved in Phase 2) was excluded from interview guide development in an effort to reduce bias. With that, PPI views and perspectives certainly come through in light of their Phase 2 engagement. Likewise, we certainly see the value of engaging PPI for the development of recruitment procedures, but given the nature of the primary cohort (i.e. researchers) as well as time constraints associated with the study, a researcher developed recruitment protocol was utilised.

7. Whilst reading the manuscript I noticed some typographical errors. For example, in Table 3: 'frquently' should be 'frequently'. There is erroneous repetition of 'With' in the second sentence of the second paragraph in the introduction.

Response 7: Many thanks for spotting these – they have been amended.

Competing Interests: N/A