



BMJ Open Peer-based Retention Of people who Use Drugs in Rural Research (PROUD-R²): a multisite, randomised, 12-month trial to compare efficacy of standard versus peer-based approaches to retain rural people who use drugs in research

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ABSTRACT

Introduction Rural communities bear a disproportionate share of the opioid and methamphetamine use disorder epidemics. Yet, rural people who use drugs (PWUD) are rarely included in trials testing new drug use prevention and treatment strategies. Numerous barriers impede rural PWUD trial engagement and advancing research methods to better retain rural PWUD in clinical trials is needed. This paper describes the Peer-based Retention Of people who Use Drugs in Rural Research (PROUD-R²) study protocol to test the effectiveness of a peer-driven intervention to improve study retention among rural PWUD.

Methods and analysis The PROUD-R² study is being implemented in 21 rural counties in three states (Kentucky, Ohio and Oregon). People who are 18 years or older, reside in the study area and either used opioids or injected any drug to get high in the past 30 days are eligible for study inclusion. Participants are allocated in a 1:1 ratio to two arms, stratified by site to assure balance at each geographical location. The trial compares the effectiveness of two retention strategies. Participants randomised to the control arm provide detailed contact information and receive standard retention outreach by study staff (ie, contacts for locator information updates, appointment reminders). Participants randomised to the intervention arm are asked to recruit a 'study buddy' in addition to receiving standard retention outreach. Study buddies are invited to participate in a video training and instructed to remind their intervention participant of follow-up appointments and encourage retention. Assessments are completed by intervention, control and study buddy participants at 6 and 12 months after enrolment.

Ethics and dissemination The protocol was approved by a central Institutional Review Board (University of Utah). Results of the study will be disseminated in academic conferences and peer-reviewed journals, online and print media, and in meetings with community stakeholders.

Trial registration number NCT03885024

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ The intervention design uses a combination of elements of successful peer navigator models for treatment retention and peer-driven models of recruitment to improve study retention.
- ⇒ The multisite nature of the study representing geographically diverse rural areas of the US strengthens potential generalisability.
- ⇒ Because the intervention is embedded in the participant retention protocol, participants and site staff cannot be blinded to arm assignment.
- ⇒ A limitation of the protocol is that participants may refer study buddies with whom they have weak social connections making it difficult to leverage the relationships for retention.
- ⇒ Relationships between intervention participants and study buddies may dissolve over time and present challenges for leveraging those ties for continued retention in the intervention arm.

INTRODUCTION

Opioid use disorder (OUD) is a national crisis in the USA.¹ Emergence of illicit fentanyl, which is 50–100 times more potent than morphine, as an adulterant of heroin, methamphetamine and other drugs has precipitously increased overdose. Rural communities bear a disproportionate share of the opioid and methamphetamine use disorder epidemics.^{2–3} Clinical trials testing new drug use prevention and treatment strategies are urgently needed. Yet, rural communities are rarely included in such trials and numerous barriers may impede retention for rural participants.^{4–5} Advancing research

methods to better retain rural people who use drugs (PWUD) in clinical trials addresses a major gap in development of new interventions to address drug use and other conditions in rural America and improves generalisability of inferences from clinical trials.

Poor retention is a common threat to validity in substance use disorder (SUD) research,^{6–8} particularly in rural communities.⁹ Barriers include perceived stigma, scheduling difficulties, transportation difficulties, community-level distrust of the research process and limited investigator experience working in non-urban communities.⁹ The National Drug Abuse Treatment Clinical Trials Network (CTN) identified individual-level and site-level facilitators of retention.^{10–15} A review of 24 completed CTN trials including 11 000 participants seeking treatment for SUD found that gender and race/ethnicity did not affect retention. Younger participants, however, had lower retention.¹² Length of trial and participant burden were not strong predictors of retention, suggesting a need to identify key variables that influence trial participation.¹⁵ Few trials, however, enrolled rural participants.

Peer navigators may be an untapped resource for improving clinical trial retention. Peers share characteristics with participants, such as demographics and drug use history, which may facilitate rapport with PWUD.¹⁶ Peers are effective at engaging and retaining hard-to-reach urban populations in SUD treatment and other clinical care.^{17 18} A systematic review of nine studies in mainly urban settings reported that peer-delivered recovery support services in SUD treatment improved outcomes,¹⁷ but their potential in rural settings is less clear.

Respondent-driven sampling (RDS)^{19 20} is a common peer-based strategy that has been used to recruit participants in biological and behavioural studies in over 80 countries,^{21 22} including SUD studies.^{23 24} RDS is a network-based sampling technique that forms chains of respondents, where purposively sampled initial participants, or ‘seeds’, are identified and given a limited number of coupons to recruit drug-using peers, who then recruit their peers. The value of peers in facilitating recruitment of PWUD into research is well established, but its effectiveness in improving study retention needs more rigorous evaluation.

The Peer-based Retention Of people who Use Drugs in Rural Research (PROUD-R²) study tests the effectiveness of a peer-driven intervention to improve study retention among PWUD in rural communities. PROUD-R² overcomes two historical roadblocks in clinical trials research: (1) widely dispersed rural populations that complicate clinical trials implementation and (2) inclusion of individuals with SUD whose social circumstances impede clinical trial participation. PROUD-R² began with a formative survey on factors, motivations and barriers influencing research participation and retention among rural PWUD,⁵ followed by the launch of the trial described in this manuscript. The PROUD-R² intervention focuses on optimising study retention in this special population and

prepares a cohort of rural PWUD for participation in clinical trials. We hypothesise that participants who receive a peer-based retention strategy will be more likely to be retained in the study at 6 and 12 months compared with participants who receive standard retention approaches.

METHODS

Study Setting

PROUD-R² leverages the National Rural Opioids Initiative (NROI),²⁵ a multistate consortium assessing interventions to increase access to care and reduce opioid overdose deaths and comorbidities in rural America. The three sites involved in PROUD-R² include 21 rural counties in Ohio (n=6), Oregon (n=3) and Kentucky (n=12).

The Oregon counties lie in the southwest coastal and mountainous areas of the state, areas with high overdose rates. These counties have a rural population from 38% to 45%.²⁶ The Ohio and Kentucky counties fall within Appalachia. These counties have been designated as ‘Distressed’ or ‘At-Risk’ counties based on several economic indicators, including unemployment, per capita income and poverty rates.²⁷ All six Ohio counties and 10 of Kentucky’s twelve counties are within the top 5% of counties in the USA most vulnerable to continued high HCV transmission and the potential for an HIV outbreak among people who inject drugs.²⁸ One Ohio county and 10 Kentucky counties are 100% rural with the remaining counties ranging from 54% to 81%.²⁶

Eligibility criteria

People who are 18 years or older, reside in the study area, and either used opioids or injected any drug to get high in the past 30 days are eligible for study inclusion. There are no exclusion criteria. Investigators may remove a participant from the study if worsening health precludes participation; participant poses a safety risk to study staff; participation is determined to be due to external pressure; or the study is terminated by the institutional review board (IRB), data safety monitoring board (DSMB) or funder. Participants are not prohibited from concurrent research or care as a condition of PROUD-R² participation.

Randomisation

Participants are allocated in a 1:1 ratio to two arms, stratified by site to assure balance at each geographical location. Research staff at each site log into a REDcap randomisation module hosted at Oregon Health and Science University. Randomisation is performed following completion of baseline data collection. Study buddies (described below) are not randomised or enrolled in the trial as intervention or control participants.

Trial arms

The trial compares the effectiveness of two retention strategies (table 1). Participants randomised to the control arm receive standard retention outreach by study staff. Participants randomised to the intervention arm are

Table 1 Description of staff tasks in the PROUD-R² participant retention protocol by study arm

	Intervention participants	Control participants
Within 3 weeks*†	Remind participant weekly to refer study buddy for a US\$10 incentive	Not Applicable
1 month*†	Remind participant weekly to refer study buddy for a US\$10 incentive	Contact participant to verify or update locator information (US\$10 incentive)
2 months	Remind participant to refer study buddy if they have not done so already	Not applicable
3 months*	Contact participant to verify/update locator information (US\$10 incentive if completed before 5 months)	
5 months	Contact participant to remind them of their appointment	
	Contact study buddy to remind them that their peer is due for follow-up on (date)	Not applicable
1 week*	For those who are unreachable, contact participant to remind them of their appointment	
2 weeks*	Follow-up PROUD-R survey window opens. For those who are unreachable, contact participant to remind them of their appointment and contact participants' contacts	
3 weeks*	For those who are unreachable, contact participant to remind them of their appointment and staff conduct home visit‡	
24 hours prior to appointment	Contact participant to remind them of their appointment	
6 months	Follow-up PROUD-R survey	
15 min after appointment time*	For those who miss appointment, contact participant to remind them of their appointment	
1 week*	For those who miss appointment, contact to ask them to reschedule and mail letter to the participant	
2 weeks*	For those who miss appointment, contact for appointment reminder and conduct home visit‡ for those who are unreachable Intervention only: Contact study buddy to remind them that their peer is due for follow-up by (date)	
3 weeks*	For those who miss appointment, contact participant's contacts to remind them of participant's appointment and mail a letter to participant's contacts	
24 hours before 7 months*	For those who miss appointment, contact participant to remind them that survey window is closing	
7 months	Follow-up PROUD-R survey window closes	
9 months - 13 months	3-month to 7-month process described above repeats	

*Jail logs are searched to identify if participant is in jail.

†Oregon and Ohio also encourage participants coenrolled in their Rural Opioid Initiative (ROI) studies and PROUD-R2 (intervention, control and study buddy participants) to refer peers for respondent-driven sampling.

‡Suspended due to COVID-19 restrictions.

PROUD-R2, Peer-based Retention Of people who Use Drugs in Rural Research.

asked to recruit a 'study buddy' in addition to receiving standard retention outreach. Study buddies remind their intervention participant of follow-up appointments and encourage attendance. Intervention participants and study buddies are advised against interacting if pressure/coercion to participate arises, they do not feel safe, or they feel the interaction could trigger unwanted drug use. Assessments are completed at 6 and 12 months after enrolment. [Table 2](#) summarises the activities planned for participant contact in the intervention and control arms.

Control condition

Following standard procedures used in longitudinal research with PWUD, participants provide detailed locator form information (online supplemental

appendix A) to assist with retention and/or contact for future research including names, pseudonyms, phone numbers, addresses, email addresses, social media contact information, employer contact information, probation/parole officer information if applicable and contact information for up to three people who should know how to reach the participant if contact information changes. Participants are contacted at the mid-point of each follow-up interval (ie, at 3-month postenrolment and 9 months postenrolment) to update locator information and remind them about their follow-up appointment date. Participants receive US\$10 for updating or verifying locator information at each interval.

Table 2 Participant timeline

Time point	Study period					
	Enrolment	Allocation	Postallocation			Closeout
	-t ₁	0 months	0 months	6 months	12 months	13 months
Enrolment						
Eligibility screen	X					
Informed consent	X					
Randomisation		X				
Interventions						
Standard retention (control)			X	X	X	
Standard retention+peer retention (intervention)			X	X	X	
Assessments						
Baseline survey	X					
Follow-up survey				X	X	
Analysis						X

Study staff contact the participant using provided information and, if unsuccessful, attempt to reach one of their contacts from the locator form. Staff send another reminder by phone, text or private message on social media at predefined intervals (table 2). If participants cannot be reached via phone or text messaging, staff attempt contact by email and mail. Sites use the same phone/text message script for appointment reminders and record information on the number of attempted contacts, number of successful contacts, intervals between contacts and type of contact with each participant. In Kentucky and Oregon, staff may offer transportation or accompany participants to the field office if COVID-19 precautions allow. Offering transportation is infeasible in Ohio. In Kentucky, participants who are incarcerated and consented to be contacted for follow-up while incarcerated may complete follow-up surveys from jail (with permission from jail administrative staff). In Oregon, staff are permitted to maintain contact while participants are incarcerated, and schedule postrelease follow-up visits. In Ohio, study staff routinely check jail and prison records to identify participants who may be unable to attend follow-up appointments.

Intervention condition

In addition to receiving the same retention strategies given to control participants, intervention participants are also provided a 'golden ticket' at the time of randomisation to refer a study buddy. They receive US\$10 for referral of an eligible study buddy who enrolls in the study. The ticket includes a unique number to link the study buddy and intervention participant; if tickets are lost, names and demographic data are used to link individuals.

Study buddies

Study buddies who enrol in the study receive a 6 min peer retention training video suggesting ways to encourage peer participation in the study without coercion; the

messages build on those used in CTN trials^{12 14 29} and previous research.^{30–35} During the iterative development process, we sought feedback on scripts and preliminary video clips from Community Advisory Boards of PWUD and/or peer support specialists at each site. The final video, available online,³⁶ describes the study, discusses the importance of retention and shows a mock peer-to-peer interaction demonstrating a study buddy reminding a peer about their follow-up appointment, inquiring about best contact methods and briefly discussing barriers. When possible, staff show videos to study buddies in person following completion of their baseline survey. Due to social distancing measures during COVID-19, staff administer some surveys by phone and send a url for the video via text message, email or social media. The video is book-ended by a short series of questions that allow staff to determine whether a study buddy watched the video and their duration of viewing. Following the video, staff ask participants about their understanding of the video and clarify content for those who did not fully understand.

Staff contact the study buddy 1 month prior to their peer's follow-up appointment date and 2 weeks afterward if the appointment is missed. In these contacts, the staff encourage the study buddy to inform the peer (ie, the intervention arm enrollee) that they are due for a follow-up but do not provide the peer's contact information to the study buddy. Study buddies are not members of the control or intervention arms but do complete the same data collection procedures and receive the same retention protocol as the control participants. Study buddies are required to meet the same eligibility criteria as other participants except for the drug use criterion, which was removed for study buddies in February 2022.

Outcomes

The primary and secondary outcomes are to compare the proportion of participants retained in intervention

and control arms at 12 and 6 months postbaseline, respectively. Participants are considered 'retained' if they at least partially complete the follow-up survey during the 164–210 days postbaseline assessment window for 6-month retention and 344–390 days window for 12-month retention.

Data collection

NROI sites use a harmonised RDS strategy to recruit PWUD.²⁵ Issues of stigma, mobility, legality and absence of adequate sampling frames for high-risk populations³⁷ create challenges for recruitment in these populations. RDS helps to address these challenges.^{20 23 38} Each PROUD-R² site recruited rural PWUD for the NROI using RDS. In Kentucky and Oregon, RDS for NROI commenced in February and March 2018, respectively, and in Ohio, began in November 2019. Participants who enrol in the NROI studies are invited to participate in the proposed PROUD-R² study. In addition, staff conduct outreach at syringe service programmes and other community venues to recruit. The total proposed sample size for PROUD-R² is 700 participants.

Community-based field staff conduct surveys that are programmed in REDCap web-based data collection system.³⁹ Staff administer surveys in-person or by phone based on participant preference. Each site uses the same survey, which collects detailed information on demographic characteristics, sexual and drug-related risk behaviour, criminal justice involvement, SUD treatment, medical care access and attitudes toward clinical trials. Staff administer follow-up surveys at 6 months and 12 months postbaseline. In Ohio, participants receive US\$40 for completing each survey, and in Kentucky and Oregon, participants receive US\$15 for each survey. The amount is higher in Ohio to improve comparability given that Kentucky and Oregon participants also have a chance to enrol in other ongoing projects wherein they receive additional incentives.

Follow-up surveys ask participants whether participants received encouragement from peers to attend their follow-up assessment; if so, from whom (name and demographic data) and whether they have a coupon referral code (see paragraph above). Information provided about who encouraged them to follow-up determines whether an incentive is owed to the study buddy. Study buddies receive US\$10 for successfully encouraging their intervention participant's retention at each follow-up.

Relationships between intervention participants and study buddies may evolve over time as people become unreachable due to incarceration, hospitalisation, movement outside of the study area or achieve different stages of recovery. Therefore, at the end of each follow-up survey, participants in the intervention arm are asked about the nature of their relationship with their study buddy. Items assess frequency of contact and where (city, state) the study buddy currently lives and whether they are homeless, incarcerated, hospitalised or in inpatient treatment. Participants are also asked whether they think this person

is in treatment and/or recovery from substance use and how that affected their interactions. Similar dyadic questions were used for network research among rural PWUD.⁴⁰ The data on the relationship between participant/study buddy will be investigated as a possible factor associated with retention in exploratory analyses.

Blinding, fidelity and contamination

Investigators remain blinded through recruitment and follow-up until completion of primary and secondary analyses, using uninformative participant labels. Study biostatisticians will also use these labels during analyses. Due to the nature of the interventions, participants and site staff administering the intervention are not blinded. These staff are instructed to use uninformative labels when discussing participants with blinded investigators.

We will use descriptive analyses to evaluate fidelity. Among individuals in the intervention arm, we will report the number and percentage of participants who did not recruit a study buddy, whose study buddy did not watch the training video, and who were not encouraged or actively discouraged by their study buddy to remain in the study. Fidelity assessment will be used to help inform the Per-Protocol Population. We will also use data from a retention task tracker programmed in REDCap that field staff use to record whether they successfully completed each task described in table 1.

Contamination is possible given participants in different study arms may be connected socially. Participants are asked whether they communicated with any peers about their follow-up visit and, if so, which peers(s) (eg, first name, nickname, first initial of last name, approximate age, gender, race and referral code (if known)) and the nature of the communication (ie, appointment reminder, received or gave encouragement to stay enrolled, messages about the importance of the research). With this detail, analysts will cross-reference information, identify potential contamination across study conditions, and conduct sensitivity analyses as appropriate.⁴¹

Data management

Data are stored in a secure REDCap database, separate from the randomisation database behind a firewall. The data manager assesses transferred data for completeness, queries sites regarding any inconsistencies, and code merged data files for analysis. The list linking participants to their unique identifier is password protected and maintained at research sites. To protect participant confidentiality, only de-identified data are shared for analysis. Data transmitted between sites contain only a unique identifier and no protected health information.

Statistical methods

The intention-to-treat (ITT) population will contain all randomised participants according to their assigned study arm. The per-protocol population will include participants who complete the trial as originally allocated. For instance, participants from the intervention arm who

do not recruit a study buddy or lose contact with their study buddy during follow-up as well as those from the control arm who are determined to have received support from an enrolled study buddy will be excluded from this population. We hypothesise that the proportion of participants retained in the intervention arm will be greater than the control arm at 12 (primary outcome) and 6 (secondary outcome) month follow-up in the ITT population. Participants who do not attend their 6-month or 12-month visit will be considered 'not retained'. There should be no missingness in the outcome in the way it has been defined and we will not conduct imputation for covariates with missing data. Type 1 error (α) will be set to 0.05 in primary and secondary outcome analyses, while $\alpha=0.1$ will be used for subgroup and exploratory analyses except where specified. All tests will be two tailed.

We will use a generalised linear mixed model (GLMM) with a logit link (ie, mixed effects logistic regression) with a fixed effect for study arm and random effects for site and RDS chain nested within site. We will include covariates: gender; age at enrolment; race/ethnicity; education; drug of choice and unstable housing status. Covariates will be measured at baseline except for drug of choice and unstable housing status, which may be updated from follow-up surveys. If a covariate causes collinearity or convergence problems when modelling, it will be excluded.

We will report exponentiated coefficients (ORs) for the study arm variable from models with and without covariates along with corresponding 95% CIs based on bootstrapped standard errors. For reporting and

dissemination, we will transform ORs to relative risk,^{42 43} which are recommended for prospective studies. Relative risk estimated from the model including covariates will represent intervention effectiveness. Sensitivity analyses will be performed repeating this analytic strategy with (1) a measure of the intervention intensity in lieu of study arm in the ITT population and (2) the per-protocol population. Supplementary preplanned analyses will explore intervention effects separately for each site as well as known or hypothesised associations with retention within groups, for example, specific drug use (opioids vs methamphetamine, opioids alone vs polysubstance use) and those experiencing unstable housing.

Another exploratory assessment will be performed using similar methods to other analyses but following a purposeful risk factor model building strategy. Among participants in the intervention arm, we will explore factors associated with retention at 12 months. Participant characteristics, study buddy characteristics, relationship between participant and study buddy, and study buddy contact characteristics will be assessed as fixed effects in univariable GLMM; the random effects used in primary analyses will be included in all models. Variables associated with retention at 12 months at a $p<0.25$ threshold in univariable models will be considered for inclusion in a multivariable model. Factors significantly associated with the outcome at $p<0.10$ as well as confounding variables will be retained in the final model. A confounder will be defined as a covariate from the list above resulting in a 10% change or greater in the estimated ORs when included in the model. Adjusted

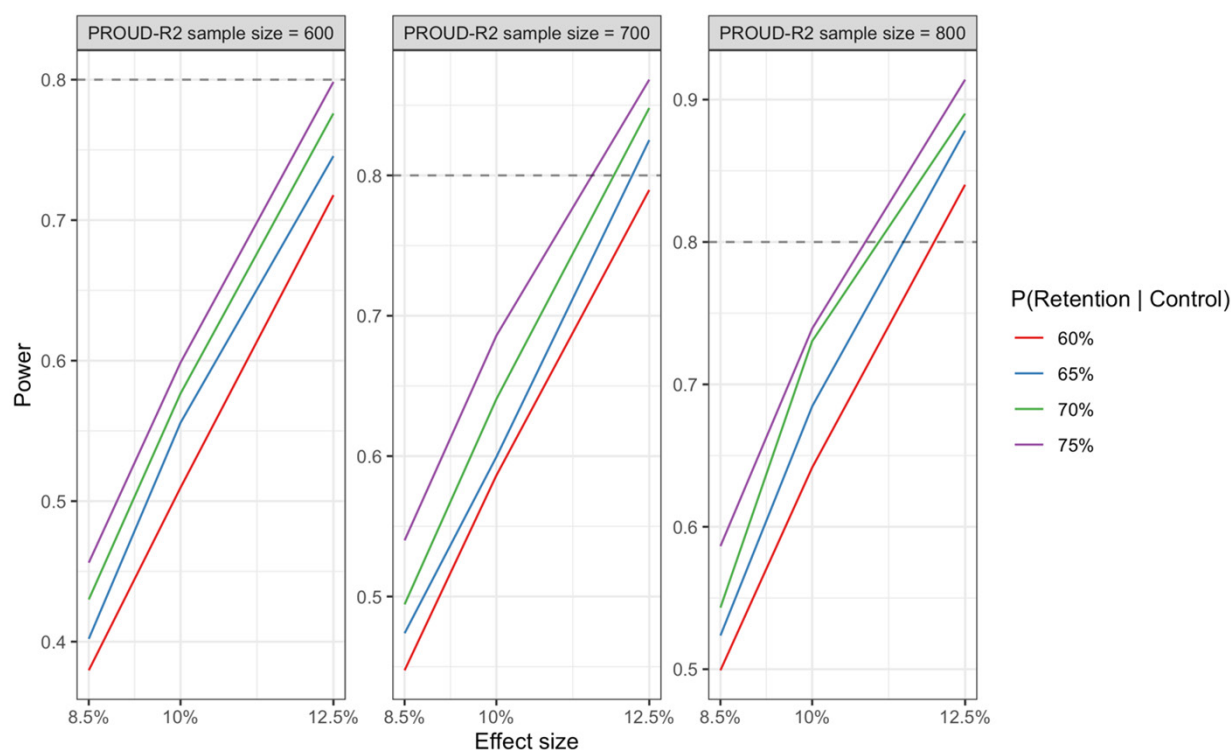


Figure 1 PROUD-R² study power. PROUD-R², Peer-based Retention Of people who Use Drugs in Rural Research.

ORs and corresponding 95% CIs will be calculated from the final model.

Power calculation

Assuming a target sample size of 700 PROUD-R² participants, we estimate that we will have at least 80% power to detect an approximately 11%–12.5% increase in retention attributable to the intervention (figure 1). Power estimates assume >65% retention in the control arm, between-RDS chain variance of 0.05, and two-sided hypothesis test at $\alpha=0.05$. We simulated RDS chain structures based on NROI recruitment data collected prior to the start of PROUD-R². Power calculations are based on simulations using GLMM with a logit link; simulated models included a fixed effect term for study group and random intercept for RDS chain. Calculations were performed using R.

Data monitoring

A DSMB comprising an addiction medicine physician, a statistician and two behavioural scientists with expertise in research among PWUD oversees the study. The DSMB is independent of the sponsor and competing interests. The DSMB will meet at least annually to review emerging data, and make recommendations about the trial's conduct, including stopping the trial. At each meeting, the DSMB will determine whether a change in the protocol is warranted, there are safety concerns and formally vote to allow the study to continue. A report on DSMB meetings and activities will be sent to the funding agency within a month after each meeting. No formal interim analyses are planned.

Social harms

Social harms related to participation will be actively assessed and documented. For this study, social harms are defined as any intended or unintended cause of physical; emotional; or psychosocial injury or hurt from one participant to another, a participant to themselves, or an institution to a participant, occurring as a result of study participation.⁴⁴ Participants will complete a social harms questionnaire at each study visit. Study staff are trained to provide appropriate care, counselling and referral as needed. Any identified social harms are reported to study investigators who determine severity and provide details to the IRB as required.

Auditing

A PROUD-R² staff person assesses REDCap data for missingness and data quality and provides feedback to site coordinators regarding any issues that need to be addressed. Investigators in Kentucky, Ohio and Oregon review study consent materials to assure appropriate documentation of consent at least semiannually.

Patient and public involvement

Participants nor the public were directly involved in the development of the research question, outcome measures or conduct of the trial. However, elements of

PROUD-R²'s design were informed by a formative survey of rural PWUD on factors influencing study retention.⁵ Also, during the iterative process of developing the intervention training video for study buddies, we sought feedback from Community Advisory Boards of PWUD and/or peer support specialists at each site about burden (ie, length) and content. Longitudinal follow-up of participants will end so that analysis can begin, making it impossible to disseminate results directly to each cohort member. However, results will be distributed via study social media pages, websites and to community partners who serve PWUD.

ETHICS AND DISSEMINATION

Sites participating in PROUD-R² rely on a Single IRB (University of Utah IRB) for human subjects oversight; local IRBs cede responsibility. Modifications to the protocol (eg, changes to eligibility criteria, outcomes or analyses, changes in study procedures) and revisions to consent forms and other participant-facing documents are submitted to the Single IRB for approval. Study staff complete human subjects training and are approved as personnel by the Single IRB. Protocol modifications are submitted to the Single IRB prior to implementation and reflected in ClinicalTrials.gov. Approval from the funding agency will be sought for major protocol modifications such as changes in inclusion criteria or aims prior to submitting those changes to the Single IRB.

All participants enrolling in PROUD-R² complete an informed consent process. Potential participants' understanding of study procedures are assessed using a comprehension tool included within the informed consent document. In addition to describing the protocol, risks and benefits, this consent form states that the participant, if randomised to the intervention arm, will provide permission for staff to reveal their name to the study buddy whom they referred so that person can help encourage their retention in the study. Participants are advised that they can refuse to engage with their study buddy, which may be important should different stages of recovery or interpersonal conflicts arise. Consent procedures are completed over the phone or in a private area with only the participant and study staff present.

Findings will be disseminated to the public and health-care professionals in peer-reviewed journals, professional conferences and community forums. Authorship eligibility guidelines follow ICMJE criteria. We will submit manuscripts to NIHMS to be made publicly available no later than 12 months after the official date of publication in compliance with the funder's open access policy. Deidentified data will be made available to interested parties on submission and approval of a written request describing data security protocols and intended use.

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Competing interests None declared.

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Appendix A. PROUD-R² Participant Locator Form

☐ Entered in locator database ☐ Baseline form, **or** ☐ Updates to earlier form Date (mm/dd/yy): _____

Staff: Do not label this paper with an ID number. File separately & alphabetically by last name.

Thank you for agreeing to let us get in touch with you in the future. We are going to ask you some questions to help us contact you. The information that you share will be kept in a separate place from your consent form and from what we learn from you in the interview. We will only use this information to locate you in the future, and it will not be given to anyone else. One thing we will ask is whether we can contact someone you know (like your mother) to reach you. If we do that, we will not tell that person *anything* except that you have been asked to take part in a health study. All questions are optional – please provide only the information you feel comfortable providing.

For people who screened ineligible:

☐ I consent to be contacted in the future by the research team regarding my willingness to participate in future research studies about rural health, substance use and related harms.

Full name:	<input type="text"/>	<input type="text"/>	<input type="text"/>
	First name	Middle name	Last name
	<input type="text"/>		<input type="text"/>
	Nicknames, aliases, other names		Maiden name(s)
Date of birth:	<input type="text"/>	Social security number (optional):	<input type="text"/>
	mm/dd/yy	Full number or <u>first</u> 5 digits	xxx-xx-xxxx

Are you homeless? No ☐ Yes ☐ → If yes, answers below should be where you spend most of your time.

Physical address:
(where you sleep and spend time)

* Street Apartment or unit number

* City State Zipcode

Whose address is this? *

Full Name (first and last) or Agency Name

Mailing address: (where you get important mail like checks)

* Street or PO Box Apartment or unit number

* City State Zipcode

Whose address is this? *

Full Name (first and last) or Agency Name

Phone numbers:			
* ()	()	()	
Cell phone	Home phone	Other phone	
*			
Whose phone is this?	Whose phone is this?	Whose phone is this?	

Email address: * @ .

Other email address: _____@_____. _____.

SOCIAL MEDIA:

May we contact you by Facebook messenger? We would only use Facebook messenger to set up appointments with you. We would not disclose any private information. ☐ No ☐ Yes → If yes, would you be willing to complete interviews by phone or video through Facebook Messenger? ☐ No ☐ Yes

***Note:** Messages would come from our study Facebook community page. We will send you an invitation to like or follow our page, however your Facebook friends may be able to see this activity. If you decide to not like or follow our page, please check your messenger junk folder for messages from our study team. Additionally, we advise you not to disclose any private information in Facebook messages that you send to us.

What is your name on Facebook?

Privacy disclaimer: We will not share your private information, however all communication using Facebook messenger is subject to Facebook's privacy and data policies. Facebook collects content and other information messaged or communicated with others. This can include information in or about the content you provide, such as the location of a photo or the date a file was created. More information can be found at www.facebook.com/about/privacy/

Instagram: Snapchat:

Other:

WORK INFORMATION:

Are you employed? ☐ No (skip section) ☐ Yes → If yes, can we contact you at work? ☐ No (skip section) ☐ Yes

Company/agency name: Supervisor name:

Company/agency address:
Street or PO Box Office or unit number

City State Zipcode

Company/agency phone:
Work phone Extension

SOCIAL SERVICES INFORMATION:

Do you have a social worker, case manager, counselor, parole officer, etc that you are comfortable with us contacting if we need to reach you? ☐ No (skip section) ☐ Yes

Name: Agency name:

Address:
Street or PO Box Apartment or unit number

City State Zipcode

Phone: () Extension:

Email:

WHO WILL ALWAYS KNOW HOW TO REACH YOU (E.G., MOTHER, FATHER, ETC)?

Full name: * * *

First name Last name Relationship to you

Address: * *

Street or PO Box Apartment or unit number

* * *

City State Zipcode

Phone: *() Email:

Please give me the TWO best people to contact who will always know where to locate you. Interviewer note: try to get at least 2 addresses/phone numbers and encourage them to name people beyond people who are also in the study. Use the space provided at the end to collect additional information if the participant provides.

PERSON 1

Full name: * * *

First name Last name Relationship to you

Address: *

Street or PO Box Apartment or unit number

* * *

City State Zipcode

Phone: *() Email:

PERSON 2

Full name: * * *

First name Last name Relationship to you

Address: *

Street or PO Box Apartment or unit number

City

State

Zipcode

Phone: () Email:

What is the best way to reach you (check all that apply)?☐ Text message ☐ Phone call ☐ Email ☐ Facebook**If you do not want us to leave a voicemail or send a text message, please indicate that here:**☐ No text messages ☐ No voicemails**What are the best days to reach you?** ☐Sun. ☐Mon. ☐Tues. ☐Wed. ☐Thurs. ☐Fri. ☐Sat.**What are the best times to reach you?** ☐Morning ☐Afternoon ☐Evening**Do you have a preference for the gender of the staff person who contacts you?**☐ No preference ☐ Female ☐ MaleIf we have trouble contacting you, may we visit you at the address you provided? ☐ Yes ☐ No***Note:** We will make every attempt to contact you in advance.If yes, what days are the best days to reach you? ☐Sun. ☐Mon. ☐Tues. ☐Wed. ☐Thurs. ☐Fri. ☐Sat.What are the best times to reach you? ☐Morning ☐Afternoon ☐Evening

Where do you hang out?

If one year from now, someone owed you \$1000 and your phone number and address had changed, how would that person find you to give it to you?

Additional notes or locator information:
