Behaviour change interventions to reduce risky substance use and improve mental health in children in care: the SOLID three-arm feasibility RCT

Hayley Alderson,¹ Eileen Kaner,^{1*} Rebecca Brown,¹ Denise Howel,¹ Elaine McColl,¹ Deborah Smart,¹ Alex Copello,² Tony Fouweather,¹ Ruth McGovern,¹ Heather Brown,¹ Paul McArdle³ and Raghu Lingam^{1,4}

Declared competing interests of authors: Eileen Kaner sat on the Public Health Research Research Funding Board (2010–16) and reports National Institute for Health Research (NIHR) Public Health Research grants during the conduct of this study. Denise Howel was a member of NIHR Health Services and Delivery Research Commissioning Board (2012–15) and is a member of NIHR Programme Grants for Applied Research Subpanel (2017–20). Elaine McColl was a member of the NIHR Journals Library Editorial Group; she was an editor for the NIHR Programme Grants for Applied Research programme (2013–16) and was a member of the NIHR Clinical Trials Unit Standing Advisory Committee until 2016. She reports grants from NIHR Public Health Research programme during the conduct of this study and other NIHR Journals Library-funded grants outside the submitted work.

Disclaimer: This report contains transcripts of interviews conducted in the course of the research and contains language that may offend some readers.

Published September 2020

DOI: 10.3310/phr08130

Scientific summary

The SOLID three-arm feasibility RCT Public Health Research 2020; Vol. 8: No. 13

DOI: 10.3310/phr08130

NIHR Journals Library www.journalslibrary.nihr.ac.uk

¹Institute of Health and Society, Newcastle University, Newcastle upon Tyne, UK

²School of Psychology, University of Birmingham, Birmingham, UK

³Child and Adolescent Mental Health Services, Tyne and Wear NHS Foundation Trust, St Nicholas Hospital, Newcastle upon Tyne, UK

⁴Faculty of Medicine, University of New South Wales, Randwick, NSW, Australia

^{*}Corresponding author eileen.kaner@newcastle.ac.uk

Scientific summary

Background

Looked-after children and care leavers (henceforth referred to as children in care) are young people who have been placed under the care of the local authority, in many instances as a result of a history of abuse and or neglect. Compared with their peers, these young people have a fourfold increased risk of drug and alcohol use and significantly increased risk of mental health disorders. To date, there is a lack of robust evidence on the most effective interventions to decrease this risk of substance use in this high-risk group of young people.

Aim

The Supporting Looked After Children and Care Leavers In Decreasing Drugs, and alcohol (SOLID) pilot trial aimed to assess the feasibility and acceptability of a definitive three-arm multicentre randomised controlled trial (two behaviour change interventions and care as usual) to reduce risky substance use (illicit drugs and alcohol), and improve mental health in children in care (aged 12–20 years).

The study had two linked phases: (1) formative qualitative work, followed by (2) an external pilot randomised controlled trial.

Objectives

Phase 1: formative study

- To adapt two behaviour change interventions for children in care to help reduce risky substance use: (1) motivational enhancement therapy and (2) social behaviour and network therapy. The adaptations to the interventions were made with involvement from children in care and drug and alcohol treatment-seeking young people, their carers, drug and alcohol workers, and social workers working with children in care to ensure acceptability and feasibility of the intervention packages.
- To refine and produce manuals for the intervention packages for integration into care pathways for children in care.
- To conduct a survey with drug and alcohol service leads across England to characterise usual care.

Phase 2: external pilot randomised controlled trial

Primary objective

• To conduct a three-arm pilot randomised controlled trial [comparing motivational enhancement therapy, social behaviour and network therapy and control (usual care)], to determine if rates of eligibility, recruitment and retention of children in care, and acceptability of the interventions, are sufficient to recommend a definitive multicentre randomised controlled trial.

Secondary objectives

- To establish data yield, data quality and acceptability of the proposed outcomes measures for self-reported alcohol and drug use, mental health and well-being, sexual behaviour and placement stability 12 months post recruitment in order to inform a sample size calculation for a definitive multicentre randomised controlled trial.
- To assess acceptability and engagement with the adapted motivational enhancement therapy and social behaviour and network therapy interventions by children in care, their carers and front-line drug and alcohol workers.
- To carry out a process evaluation to include fidelity of intervention delivery and qualitative assessment of the barriers to successful implementation.
- To develop a core intervention delivery package, potentially of a single optimised intervention, linked to a theory of change model to use in the definitive trial.
- To develop and assess tools to collect data on costs and health benefits, and carry out a value of information analysis to inform the appropriate sample size in a definitive study.
- To apply prespecified STOP/GO criteria and determine if a definitive multicentre randomised controlled trial is feasible and, if so, develop a full trial protocol. Criteria for progression to a definitive trial were recruitment of 60% of children in care identified as eligible, 80% of participants attending 60% of offered sessions and retention of 70% of participants after 12 months' follow-up.

Methods

Public and patient involvement

Public and patient involvement was sought at multiple time points throughout the study. Public and patient involvement representatives included children in care, local authority employees, drug and alcohol practitioners and non-looked-after children. Their contributions have influenced the study design and they have co-designed study documentation and the adapted motivational enhancement therapy and social behaviour and network therapy manuals.

Formative phase

The formative research consisted of five separate, but interconnected, stages: (1) the selection of two evidence-based interventions suitable for adaptation to be used with children in care; (2) development of a theory of change model; (3) conducting qualitative interviews and focus groups with key stakeholders to examine the principles behind the motivational enhancement therapy and social behaviour and network therapy approaches; (4) analysis of the qualitative data; and (5) the co-production of the final interventional manuals.

Participants (children in care, foster carers, residential workers, social workers and drug and alcohol workers) were recruited purposively to ensure maximum diversity.

In-depth one-to-one semistructured interviews, dyad interviews and focus groups were used to explore the assumptions inherent within our logic models, the principles behind the adapted motivational enhancement therapy and social behaviour and network therapy approaches, their relevance to children in care and the broader therapeutic approaches, inclusive of the key behavioural and motivational domains that the interventions should address when working with the population of children in care.

Interviews and focus groups were audio-recorded and transcribed verbatim. Data were collected until data saturation. Transcripts were anonymised and identifiable participant details removed. Pseudonyms were allocated to each transcript and have been used in all reports and publications to maintain participants' anonymity.

Survey

A telephone/online survey of all drug and alcohol providers was carried out to describe the drug and alcohol treatment services offered in each local authority in England as of 2018. The survey aimed to define usual care for a definitive future study.

Randomised controlled trial

The three-arm randomised controlled trial compared the adapted motivational enhancement therapy and adapted social behaviour and network therapy interventions developed in the formative phase with usual care. The trial involved children in care across six local authorities in north-east England. Trial participants were screened for drug and alcohol use with the validated six-question CRAFFT [car, relax, alone, forget, friends, trouble(s)] screening tool administered by their social worker. Children in care, aged 12–20 years, who reported drug or alcohol use within the previous 12 months and who were able to provide assent or informed consent in English, depending on their age, were eligible to participate in the trial. Assent was taken from children aged < 16 years and informed consent for those aged \geq 16 years. Those who were currently receiving treatment from drug and alcohol services, were due to move out of the area, or unable to give informed consent (due to acute or severe mental health difficulties, mental capacity or language barriers) were ineligible to participate.

Data were collected at baseline and 12 months post baseline, using participant-completed questionnaires on a tablet computer. The baseline and follow-up questionnaires measured self-reported drug and alcohol use (Alcohol Use Disorder Identification Test and Alcohol, Smoking and Substance Involvement Screening Tool), mental health and well-being (Strengths and Difficulties Questionnaire and Warwick–Edinburgh Mental Wellbeing Scale), and health-related quality of life (EuroQol-5 Dimensions, five-level version). The follow-up questionnaire also collected data on placement stability, use of health and social service, self-reported sexual behaviour, and antisocial and criminal behaviour. Data using timeline follow-back substance use and self-reported occasions of 'drunkenness' in the last 30 days were collected.

Individual randomisation was stratified by placement type (residential/non-residential), site and age band (12–14 years/> 14 years) to reflect risk profile for substance use. Interventions were delivered by experienced young people's drug and alcohol practitioners who received 2 full days' training in the adapted allocated intervention; either social behaviour and network therapy or motivational enhancement therapy. Participants allocated to the control group received usual care, which involved their social worker making a referral along the usual drug and alcohol service pathway, as required.

Process evaluation

A detailed process evaluation, using both qualitative and quantitative methods was conducted, and involved children in care, their foster carers, residential workers, social workers, and drug and alcohol practitioners. In-depth one-to-one interviews, dyad interviews and focus groups were used to explore the key lessons learned from implementing the Supporting Looked After Children and Care Leavers In Decreasing Drugs, and alcohol (SOLID) trial (both the interventions and the trial processes). Quantitative methods (practitioner intervention log, audio files) were used to assess the quality of intervention delivery (treatment fidelity) by applying a validated process rating scale developed in the United Kingdom Alcohol Treatment Trial.

Economic analysis

The study conducted an exploratory return on investment analysis, which aimed to assess the feasibility of a within-trial economic analysis in the context of a definitive trial.

Ethics and consent

A favourable ethics opinion was granted by Newcastle and North Tyneside 1 National Research Ethics Service Committee (16/NE/0123). Informed consent (assent for those aged < 16 years) was taken from all participants. For children in care aged < 16 years, informed assent was requested with an accompanying adult (parent, carer, social worker, children's home lead) present. If the accompanying adult did not have parental responsibility then the research team contacted the adult with parental responsibility to obtain informed consent prior to the young person taking part in the research.

An adverse events procedure was implemented; however, no adverse events were reported throughout the study.

Analysis

Qualitative data

The transcripts from the qualitative interviews (formative and process evaluation) were subject to thematic analysis, an iterative process, using the constant comparative method. Qualitative software (NVivo 10, QSR International, Warrington, UK) aided in the organisation of thematic codes and categories. The analysis of drug and alcohol practitioner data within the process evaluation was informed by normalisation process theory.

Quantitative trial data

The main outcomes were feasibility outcomes to inform the design of a future definitive study. Descriptive statistics were used and no formal comparisons were drawn as the sample size was not powered to detect differences (*Table a*).

Results

Formative

The manual development incorporated adaptations suggested by key stakeholders (n = 65: 24 children in care, eight non-care young people and 33 professionals). Key adaptations made were the need to focus on overcoming mistrust; insecure attachments that children in care experience due to their history

TABLE a The STOP/GO criteria

	Criteria				
STOP/GO criteria	Green	Amber	Red	Achieved	
Percentage of eligible participants consenting to trial	≥ 60	40-60	< 40	53	
Percentage of children attending 60% of offered sessions	≥80	20-80	< 20	9	
Percentage of participants retained to 12 months' follow-up	≥ 70	50-70	< 50	54	
Were interventions delivered with fidelity?	Yes	Unclear	No	Unclear	
Were interventions perceived acceptable by children in care and workers?	Yes	Unclear	No	Low uptake of intervention by children, but acceptable to workers	
Does the value of information analysis show future research is worthwhile?	Worthwhile	Unclear	Not worthwhile	No available data	

[©] Queen's Printer and Controller of HMSO 2020. This work was produced by Alderson et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

of abuse and/or neglect; and their interactions with the care system. Owing to the fragmented nature of support networks available to children in care, more flexibility is required regarding social network members. There is a need to use creative methods to engage children in care. Finally, children in care are known to have higher rates of comorbid mental health problems and higher levels of risk-taking behaviour; therefore, treatment goals need to be wider than substance misuse alone to accommodate the diverse needs of this population group. The manuals underpinned the training delivered to the drug and alcohol practitioners in preparation for the intervention delivery.

Survey

In total, 122 (82%) of 149 national drug and alcohol services completed the survey. The survey highlighted the high levels of variation in drug and alcohol service across England, with regard to screening and intervention delivery. None of the services reported delivering manualised evidence-based interventions. Just over half of the local authorities (n = 67, 55%) said that they offered a 'bespoke service'; however, this was often not specified in many cases. Thirty-five (29%) local authorities reported offering structured work.

Randomised controlled trial

Of the 1450 eligible children in care aged 12–20 years, 860 (59%) were screened for drug and alcohol use by social workers over a period of 1 year. A total of 211 (24.5%) eligible children met the inclusion criteria of the trial, and 112 young people (7.7% of the original eligible cohort) consented and were randomised into the trial arms.

In total, 15 out of the 76 (20%) participants allocated to an intervention arm attended any of the offered motivational enhancement therapy or social behaviour and network therapy sessions. Sixty participants (54%) completed the 12-month follow-up questionnaires. The pilot randomised controlled trial did not meet the prespecified 'GO' criteria, demonstrating that a definitive randomised controlled trial is not feasible. As the primary outcomes were recruitment and retention to 12 months' follow-up, no formal comparisons are drawn between groups, as the sample size had not been powered to detect group differences.

Process evaluation

There were 109 stakeholders involved in the process evaluation (37 children in care and 72 professionals). Findings illustrated that the principles of the adapted interventions were acceptable to the different professional groups and the method of collecting data on a tablet computer worked well. However, the combination of multiple steps in the study process and the time lost between screening and first appointment set up within the current referral pathways, meant that the process was not swift enough to engage participants in the trial and the interventions. In addition, even though many participants were engaged in risky drug and alcohol use, acknowledgement of risk and the need to reduce their use was rare. The combination of these factors meant that the adapted interventions could not be delivered to a sufficient extent to fully test acceptability in practice.

Cost of intervention delivery

A log was completed by drug and alcohol practitioners to help the research team calculate the cost of delivering the interventions. The practitioners seemed able and willing to fill out the log. Unfortunately, the sample size was too small to conclude whether or not the tool is acceptable to use in a definitive study.

Fidelity of intervention delivery

Practitioners uploaded nine out of a potential 26 audio-recordings. The quality of the recordings was variable and due to the limited number we are unable to assess the internal validity of the interventions being delivered.

Economics

Our exploratory return on investment analysis concluded that a medium to large health effect would need to be demonstrated before the intervention would be considered cost-effective.

Synthesis of findings

The criteria to determine the feasibility of progressing to a full definitive trial were not met. Major challenges were found; social care staff were overstretched, resulting in screening and recruitment being problematic. This, compounded with children in care experiencing complex care arrangements and placement instability, resulted in intervention delivery being challenging, and the fidelity of the intervention could not be assessed owing to low uptake. The current format of the intervention pathway was not feasible to deliver.

This is the first UK-based pilot feasibility trial that assessed the feasibility of delivering behaviour change interventions to decrease drug and alcohol use and support mental health of children in care. Several key lessons have been learnt to inform future service delivery and research for children in care.

Model of care

The screen and treat model used in SOLID has been shown to be problematic. Any future trial needs to think about how best to engage children in care. The current model of referring participants on to another service does not work, interventions need to be delivered opportunistically within enhanced social care pathways.

Delivery agent and support

The SOLID trial tried to use the existing drug and alcohol services and standard referral system to deliver novel interventions. Children's services departments are often less research mature and very stretched. Without additional, dedicated 'in-service' academic support, potentially in the form of embedded researchers/academic social workers, research with children in care will prove difficult. An embedded researcher has the potential to be jointly managed by local authorities and universities, facilitating clearance to engage clients; this could significantly change the research culture within units. A new way of working, in which drug and alcohol workers could be allocated to social care services and residential units, would decrease referral times and could be a new way of delivering these interventions. The National Institute for Health Research Clinical Research Network has now extended support into research taking place in non-NHS settings, such as health and social care and public health. These amendments could facilitate the necessary change as it has within NHS research trials.

Conclusion

This study found that many children in care do not identify themselves as needing a drug and alcohol intervention, despite reporting use of substances and linked risky behaviours. This mismatch between the views of professionals and those of children in care justifies further attention. Future ecologically relevant models of care are needed for children in care to improve the outcomes of these potentially vulnerable young people across their life course.

Study and trial registration

This study is registered as PROSPERO CRD42018098974 and ISRCTN80786829.

Funding

This project was funded by the National Institute for Health Research (NIHR) Public Health Research programme and will be published in full in *Public Health Research*; Vol. 8, No. 13. See the NIHR Journals Library website for further project information.

Public Health Research

ISSN 2050-4381 (Print)

ISSN 2050-439X (Online)

This journal is a member of and subscribes to the principles of the Committee on Publication Ethics (COPE) (www.publicationethics.org/).

Editorial contact: journals.library@nihr.ac.uk

The full PHR archive is freely available to view online at www.journalslibrary.nihr.ac.uk/phr. Print-on-demand copies can be purchased from the report pages of the NIHR Journals Library website: www.journalslibrary.nihr.ac.uk

Criteria for inclusion in the Public Health Research journal

Reports are published in *Public Health Research* (PHR) if (1) they have resulted from work for the PHR programme, and (2) they are of a sufficiently high scientific quality as assessed by the reviewers and editors.

Reviews in *Public Health Research* are termed 'systematic' when the account of the search appraisal and synthesis methods (to minimise biases and random errors) would, in theory, permit the replication of the review by others.

PHR programme

The Public Health Research (PHR) programme, part of the National Institute for Health Research (NIHR), is the leading UK funder of public health research, evaluating public health interventions, providing new knowledge on the benefits, costs, acceptability and wider impacts of non-NHS interventions intended to improve the health of the public and reduce inequalities in health. The scope of the programme is multi-disciplinary and broad, covering a range of interventions that improve public health.

For more information about the PHR programme please visit the website: https://www.nihr.ac.uk/explore-nihr/funding-programmes/public-health-research.htm

This report

The research reported in this issue of the journal was funded by the PHR programme as project number 14/183/08. The contractual start date was in March 2016. The final report began editorial review in May 2019 and was accepted for publication in October 2019. The authors have been wholly responsible for all data collection, analysis and interpretation, and for writing up their work. The PHR editors and production house have tried to ensure the accuracy of the authors' report and would like to thank the reviewers for their constructive comments on the final report document. However, they do not accept liability for damages or losses arising from material published in this report.

This report presents independent research funded by the National Institute for Health Research (NIHR). The views and opinions expressed by authors in this publication are those of the authors and do not necessarily reflect those of the NHS, the NIHR, NETSCC, the PHR programme or the Department of Health and Social Care. If there are verbatim quotations included in this publication the views and opinions expressed by the interviewees are those of the interviewees and do not necessarily reflect those of the authors, those of the NHS, the NIHR, NETSCC, the PHR programme or the Department of Health and Social Care.

© Queen's Printer and Controller of HMSO 2020. This work was produced by Alderson *et al.* under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

Published by the NIHR Journals Library (www.journalslibrary.nihr.ac.uk), produced by Prepress Projects Ltd, Perth, Scotland (www.prepress-projects.co.uk).

Editor-in-Chief of **Public Health Research** and NIHR Journals Library

Professor Ken Stein Professor of Public Health, University of Exeter Medical School, UK

NIHR Journals Library Editors

Professor John Powell Chair of HTA and EME Editorial Board and Editor-in-Chief of HTA and EME journals. Consultant Clinical Adviser, National Institute for Health and Care Excellence (NICE), UK, and Senior Clinical Researcher, Nuffield Department of Primary Care Health Sciences, University of Oxford, UK

Professor Andrée Le May Chair of NIHR Journals Library Editorial Group (HS&DR, PGFAR, PHR journals) and Editor-in-Chief of HS&DR, PGFAR, PHR journals

Professor Matthias Beck Professor of Management, Cork University Business School, Department of Management and Marketing, University College Cork, Ireland

Dr Tessa Crilly Director, Crystal Blue Consulting Ltd, UK

Dr Eugenia Cronin Senior Scientific Advisor, Wessex Institute, UK

Dr Peter Davidson Consultant Advisor, Wessex Institute, University of Southampton, UK

Ms Tara Lamont Senior Scientific Adviser (Evidence Use), Wessex Institute, University of Southampton, UK

Dr Catriona McDaid Senior Research Fellow, York Trials Unit, Department of Health Sciences, University of York, UK

Professor William McGuire Professor of Child Health, Hull York Medical School, University of York, UK

Professor Geoffrey Meads Professor of Wellbeing Research, University of Winchester, UK

Professor John Norrie Chair in Medical Statistics, University of Edinburgh, UK

Professor James Raftery Professor of Health Technology Assessment, Wessex Institute, Faculty of Medicine, University of Southampton, UK

Dr Rob Riemsma Reviews Manager, Kleijnen Systematic Reviews Ltd, UK

Professor Helen Roberts Professor of Child Health Research, UCL Great Ormond Street Institute of Child Health, UK

Professor Jonathan Ross Professor of Sexual Health and HIV, University Hospital Birmingham, UK

Professor Helen Snooks Professor of Health Services Research, Institute of Life Science, College of Medicine, Swansea University, UK

Professor Ken Stein Professor of Public Health, University of Exeter Medical School, UK

Professor Jim Thornton Professor of Obstetrics and Gynaecology, Faculty of Medicine and Health Sciences, University of Nottingham, UK

Professor Martin Underwood Warwick Clinical Trials Unit, Warwick Medical School, University of Warwick, UK

Please visit the website for a list of editors: www.journalslibrary.nihr.ac.uk/about/editors

Editorial contact: journals.library@nihr.ac.uk