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Abstract

A self-efficacy enhancement alcohol reduction intervention for men on-remand in prison: the APPRAISE feasibility pilot RCT

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Background: As many as 70% of remand prisoners have admitted to being under the influence of alcohol when committing the crime leading to their imprisonment. Providing support and advice regarding alcohol consumption can be effective in some groups of people. There is little evidence regarding this for men on remand in prison.

Objective: To pilot the study measures and evaluation methods to assess the feasibility of conducting a future definitive multicentre, pragmatic, parallel group, randomised controlled trial.

Design: A two-arm, parallel group, individually randomised pilot study of a self-efficacy-enhancing psychosocial alcohol intervention to reduce levels of alcohol consumption for males on remand in prison and on liberation.

Setting: Two purposively selected prisons in Scotland and England.

Participants: Adult men on remand in prison with an Alcohol Use Disorders Identification Test score of ≥ 8 .

Intervention: The APPRAISE intervention delivery comprised four steps: Step 1: 1 \times 40-minute face-to-face session, delivered by a trained practitioner from Change Grow Live in prison. Steps 2, 3 and 4: 20-minute sessions conducted by phone, on or as close as possible to days 3, 7 and 21 post liberation. Control: assessment, screening and referral onto further alcohol support options.

Main outcome measures: Recruitment and retention rates, completion of follow-ups, outcome measures at 12 months and interventions delivered. The primary outcome for the pilot study was alcohol consumed in the 28 days prior to Time Point 2, assessed using the extended Alcohol Use Disorders Identification Test-C.

Results: Of 182 men on remand approached across two study sites, 132 were randomised (90 in England; 42 in Scotland) with 46 randomised to intervention and 44 to care as usual in England and 22 randomised to intervention and 20 to care as usual in Scotland. A total of 53 in-prison interventions were delivered. One day-3 post-liberation intervention was delivered, no day-7 and one day-21. At 12 months, of 132 randomised, 18 (13%) were followed up, 53 (40%) were not liberated; 47 (36%) were uncontactable and 14 (11%) had been released but could not be located. Data completeness was 96% at baseline and 8% at 12 months. The process evaluation reported good acceptability of the intervention with investment in time, capacity and space to support implementation identified. The economic study produced guidance on how to assess costs associated with implementing the APPRAISE intervention which could be applied more broadly.

Harms: No adverse events or side effects were noted.

Conclusions: A future definitive trial would be possible, but only if follow-up mechanisms can be addressed as well as full access to recidivism and health data. Collaboration with the probation service in future could offer the opportunity to develop a robust process and system to optimise follow-up post liberation. Dedicated resources to support the intervention delivery both in and out of the prison setting are recommended.

Limitations: Coronavirus disease discovered in 2019 impacted recruitment and follow-up, with access to prisons restricted. We were unable to deliver the post-liberation element of the intervention. We did not include probation services or other agencies in the trial.

Trial registration: This trial is registered as Current Controlled Trials ISRCTN36066.

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List of supplementary material

Report Supplementary Material 1 Secondary outcome results stratified by site and Summary statistics for mechanisms of change

Supplementary material can be found on the NIHR Journals Library report page (<https://doi.org/10.3310/KNWT4781>).

Supplementary material has been provided by the authors to support the report and any files provided at submission will have been seen by peer reviewers, but not extensively reviewed. Any supplementary material provided at a later stage in the process may not have been peer reviewed.

List of abbreviations

AA	Alcoholics Anonymous sessions	GDPR	General Data Protection Regulation
ABC	activity-based costing	GP	general practitioner
ABI	alcohol brief intervention	HMPPS	His Majesty's Prison and Probation Service
APPRAISE	A two-arm parallel group individually randomised Prison Pilot study of a male Remand Alcohol Intervention for Self-efficacy Enhancement	HMPS	His Majesty's Prison Service
AUDIT	Alcohol Use Disorders Identification Test	HRA	Health Research Authority
AUDIT-C	Alcohol Use Disorders Identification Test – Consumption	IRAS	Integrated Research Application System
BECCI	Behaviour Change Counselling Index	ITT	intention-to-treat
CEA	cost-effectiveness analysis	MAR	missing-at-random
CGL	Change Grow Live	MCAR	missing-completely-at-random
CHEERS	Consolidated Health Economic Evaluation Reporting Standards	mNCA	model non-commercial agreement
CI	chief investigator	MoJ	Ministry of Justice
CJM	community justice mentor	MRC	Medical Research Council
CJS	criminal justice system	MRC-PHIND	Medical Research Council Public Health Intervention Development
CJSW	criminal justice social worker	NAEQ	Negative Alcohol Expectancy Questionnaire
CONSORT	Consolidated Standards of Reporting Trials	NHS	National Health Service
COVID-19	coronavirus disease discovered in 2019	NICE	National Institute for Health and Care Excellence
CRF	case report form	NIHR	National Institute for Health and Care Research
DART	Drug and Alcohol Recovery Team	NPT	normalisation process theory
DMEC	Data, Monitoring and Ethics Committee	PHE	Public Health England
DRSEQ-R	Drinking Refusal Self-efficacy Questionnaire – revised	PIS	Participant Information Sheet
ECTU	Edinburgh Clinical Trials Unit	PMG	project management group
EQ-5D-5L	EuroQol-5 Dimensions, five-level version	PNC	Police National Computer
FRAMES	Feedback, Responsibility, Advice, Menu, Empathy, Self-efficacy	PPE	personal protective equipment
GDP	gross domestic product	PPI	patient and public involvement
		PRISM-A	Alcohol Brief Interventions for Male Remand Prisoners
		PSS	personal social services
		RA	research assistant

LIST OF ABBREVIATIONS

RAEC	Research and Access Ethics Committee	TIDieR	Template for Intervention Description and Replication
RCT	randomised controlled trial	TLFB-28	timeline followback-28
SAS	statistical analysis system	ToR	terms of reference
SD	standard deviation	TP	time point
SIMD	Scottish Index of Multiple Deprivation	TSC	Trial Steering Committee
SPS	Scottish Prison Service	UK	United Kingdom
SPSS	statistical package for social science	WEMWBS	Warwick-Edinburgh Mental Well-being Scale
		WHO	World Health Organization

Plain language summary

What was the question?

We know many men on remand report being under the influence of alcohol at the time of their arrest. Having a short conversation providing alcohol support and advice (known as an 'intervention') can be effective. However, we do not know if an alcohol intervention for men on remand in prison is possible to deliver or if we can 'test' the benefit of it. The aim of the study was to 'test' an intervention named APPRAISE using a small study, for men on remand in one Scottish and one English prison.

What did we do?

We aimed to get 180 men in this study. Men taking part were asked to tell us about their alcohol drinking. Men reporting 'risky' drinking were split into two groups by chance. One group were to be given the APPRAISE intervention while in prison and once released. The other group did not receive the intervention. We then set out to measure their drinking levels after 12 months. We interviewed different people involved in the study to find out what they thought.

What did we find?

We successfully recruited 132 men but had to stop due to coronavirus disease discovered in 2019, as we were unable to go into the prisons. We delivered 53 out of 68 interventions in prison but not once men were released; we were only able to contact 18 out of 132 men at 12 months. People we interviewed stated that having an intervention to reduce risky drinking would be acceptable; however, this would require investment, time, space, capacity and trust.

What does this mean?

Before we can plan a larger study we need to identify the best way to locate men once released from prison, to deliver the whole intervention and measure its effect on drinking levels.

Scientific summary

The prevalence of at-risk drinking, which includes drinking at levels that harm a person's health, is far higher amongst those in contact with the criminal justice system (73%). For those on remand in prison, the prevalence is between 62% and 68%. This compares to 35% in the general population.

Alcohol brief interventions (ABIs) are a secondary prevention activity, aimed at those individuals who are drinking in a pattern that is likely to be harmful to health and/or well-being. Similarly, the theoretical validity of a self-efficacy-enhancing alcohol intervention in other settings has shown evidence of potential effect.

Study aims and objectives

Objective 1: to pilot the study measures and evaluation methods to assess the feasibility of conducting a future definitive multicentre, pragmatic, parallel-group randomised controlled trial.

- 1a. Is it feasible to conduct a future multicentre RCT of a self-efficacy-enhancing psychosocial alcohol intervention for men on remand?
- 1b. Can we obtain reasonable estimates of the parameters necessary to inform the design and sample size calculation for a future definitive multicentre RCT? This includes standard deviations of potential continuous primary outcomes and estimates of recruitment met across trial arms and study, retention and follow-up rates.
- 1c. How well do participants complete the questionnaires necessary for a future definitive RCT?
- 1d. Can we collect economic data needed for a future definitive RCT?
- 1e. Can we access recidivism data from the Police National Computer (PNC) databases for trial participants?
- 1f. Can we access health data from routine National Health Service (NHS) data sources for trial participants?

Objective 2: to assess intervention fidelity.

- 2a. What proportion of the interventions are delivered as per protocol?
- 2b. Is there any evidence of contamination between the two conditions and/or between those workers delivering the intervention?
- 2c. To what extent was the intervention changing process variables consistent with the underpinning theory?

Objective 3: to qualitatively explore the feasibility and acceptability of a self-efficacy-enhancing psychosocial alcohol intervention and study measures to staff and for men on remand and on liberation.

- 3a. How acceptable are the trial and intervention procedures (including context and any barriers and facilitators) to the following key stakeholders: men on remand in prison and on liberation; prison staff (including healthcare staff); commissioners; policy-makers and third-sector partners?

Objective 4: to assess whether operational progression criteria for conducting a future definitive randomised controlled trial are met across trial arms and study sites and, if so, develop a protocol for a future definitive trial. (Operational progression criteria are based on previous research results.)

- 4a. Do the two prisons invited to the study agree to take part?
- 4b. Based on knowledge from previous data, do at least 90 eligible participants consent to take part and be randomised across the trial arms?
- 4c. Do at least 70% of participants who consent to the trial receive the intervention?
- 4d. Are at least 60% of those who received the intervention followed up at 12 months across trial arms and study sites?

Objective 5: to ascertain what alcohol services are available in male remand prisons and how coronavirus disease discovered in 2019 has affected services.

- 5a. To ascertain what alcohol services are currently provided within male remand prisons.
- 5b. To explore the prison governors' understanding of brief interventions.
- 5c. To understand how the coronavirus disease discovered in 2019 pandemic has impacted the services available in male remand prisons.
- 5d. To identify whether the impact of the COVID-19 pandemic upon services could be avoided in the future.

Methods

Phase 1 pilot trial

Phase I was a two-arm, parallel-group, individually randomised pilot study of a self-efficacy-enhancing psychosocial alcohol intervention for men on remand in prison to provide data, including economic, recidivism and health data, on feasibility and an assessment of the likely impact of the APPRAISE intervention to inform the feasibility of a future definitive multicentre RCT.

The Phase I pilot trial was undertaken in two prison settings, one in Scotland and one in England. Those eligible to participate were adult men detained on remand in either the Scottish study site or the England study site who had been in prison for 3 months or less and had an Alcohol Use Disorders Identification Test (AUDIT) screening score of 8 or more. The original recruitment target was 180 participants, 90 at each study site.

The target was reached at the England study site; however, restricted access to the prison estates due to COVID-19 meant we were unable to recruit any further participants at the Scottish study site. As a result, we recruited 132 participants (90 in England and 42 in Scotland), who were randomised. In England, 46 participants were randomised to the intervention and 44 to care as usual. For participants in Scotland, 22 were randomised to the intervention and 20 to care as usual. Allocation was conducted at the level of the participants, randomised to the active or control intervention using stratified block randomisation by site, via sealed envelopes, based on a predetermined random number allocation carried out by the study Trials Unit.

The APPRAISE intervention focused on enhancing self-efficacy and comprised four steps: Step 1 comprised a 1 × 40-minute face-to-face session in which the nine elements were delivered by an interventionist in the prison setting. Steps 2, 3 and 4 were 20-minute booster sessions to be delivered by phone, on or as close as possible to days 3, 7 and 21 post liberation. Interventions were delivered by Change Grow Live practitioners who had received prior training in intervention delivery. Care as usual across both sites comprised an alcohol assessment and referral onto further alcohol support options if requested. Due to COVID-19 restrictions, access to both study sites for the research assistants (RAs) and research team was halted, resulting in liberation data not being accessible. There was limited capacity of the interventionists to follow up on liberation data of those in the intervention group or deliver post-liberation intervention.

Data were collected at TP0 before randomisation (baseline), TP1 (6 months) and TP2 (12 months). Follow-up assessments were attempted where participants had been (a) not liberated, (b) liberated and in the community or (c) liberated and then re-incarcerated, and were conducted by phone; hard copies of the follow-up questionnaire were sent by post with an accompanying letter to be completed and returned to the study team via a pre-paid envelope, or via hard copy in prison. As a result, modifications were made to the follow-up method to also include contact via text message, WhatsApp, Facebook and an electronic Qualtrics link to the survey sent by phone or e-mail. The case report forms (CRFs) were adapted to facilitate self-completion.

The primary outcome was alcohol consumed in the 28 days prior to TP2 (12-month follow-up), assessed using the extended AUDIT-C. The following secondary outcome measures were used across the three time points: Warwick-Edinburgh Mental Well-being Scale (WEMWBS); Drinking Refusal Self-efficacy Questionnaire – revised (DRSEQ-R); Negative Alcohol Expectancy Questionnaire (NAEQ); EuroQol-5 Dimensions, five-level version; Readiness to Change Ruler; Economic Form 90.

Results

Of 182 men on remand approached across two study sites, 132 were randomised (90 in England; 42 in Scotland) with 46 randomised to intervention and 44 to care as usual in England, and 22 randomised to intervention and 20 to care as usual in Scotland.

A total of 53 in-prison interventions were delivered. One day-3 post-liberation intervention was delivered, no day-7 post-liberation intervention was delivered and one day-21 post-liberation intervention was delivered. At 12 months out of all 132 randomised, 18 were followed up, 53 (40%) were not liberated, 47 (36%) were uncontactable and 14 (11%) had been released but could not be located. Data completeness was 96% at baseline and 8% at 12 months.

Mindful of the very small sample sizes and that as such caution should be applied, provisional indications suggest that self-efficacy may be a determinant of alcohol consumption and further exploration of interventions targeting self-efficacy should be considered. We were able to develop a micro-costing methodology protocol and collect provisional scoping of routine data sources to support future cost-effectiveness analyses. We were unable to access both PNC and NHS data for participants in either study prison.

Phase II process evaluation and survey

In Phase II, the aim of the process evaluation was threefold: first, to assess how the intervention was implemented, second, to undertake some preliminary exploration of change mechanisms underpinning the intervention, and third, to assess the acceptability and context within which the intervention was delivered through interviews with study participants and key stakeholders involved in supporting the trial. A survey of prisons in Scotland and England was conducted to ascertain what alcohol services were available for men on remand in prisons and how COVID-19 had affected these services.

Study records and semistructured interviews provided the data for the process evaluation. Purposive sampling was used for interviews. Due to COVID-19, the target of 40 interviews was not attainable. Thematic analyses using a NPT lens to support thematic identification. All interviews were digitally recorded and transcribed verbatim ahead of analysis.

High levels of practitioner behaviour change skills were identified from the Behaviour Change Counselling Index (BECCI) scores of the four intervention delivery sessions recorded at the Scotland site. Differences in median dose delivered between study sites were noted. CGL intervention training evaluation was positive. From the data available evidence of contamination was limited.

Fifteen semistructured interviews were conducted with three participant groups [remand participants, Change Grow Live (CGL) team and wider stakeholders] from across the two study sites. The themes generated suggested a strong acceptability of the intervention with investment in time, capacity and space identified to support implementation, as well as the buy-in from all stakeholders, development of trust and relationships as key facilitators to supporting behaviour change.

From 59 prisons in Scotland and England, successful e-mails were sent to 55 prison governors. Seventeen (31%) were completed. The findings confirmed that the COVID-19 pandemic had undoubtedly impacted upon alcohol and wider prison services available to men on remand.

Strengths of the study

- The APPRAISE study is, to the best of our knowledge, the first pilot trial of an alcohol-focused self-efficacy-enhancing intervention for men on remand.
- The study provides significant insight into the feasibility and acceptability of pilot trials for this particular population.
- The ethical approval process for the prison setting across devolved countries provided useful insights into how to navigate the submission and approvals process across a range of ethics committees.
- The study provided good evidence of the feasibility of recruitment, training interventionists and subsequent in-prison delivery of the APPRAISE intervention to 73% of men on remand, randomised to the intervention, within a highly complex prison setting.
- The economic findings provide valuable insights, as we believe this to be the first of its kind for this type of intervention in this setting.
- Useful insight was gained regarding the access to PNC and NHS Service Use Data.
- The process evaluation provided insights into the perspectives of the remand participants, the intervention delivery team and wider stakeholders on the acceptability and feasibility of intervention implementation.
- Our patient and public involvement (PPI) colleague, a co-applicant on the study, provided unique insights and guidance throughout the study.

Limitations of the study

- Protracted multiple ethical approvals across devolved countries and processes meant a significant delay in recruitment commencing, particularly for the Scotland site.
- We were unable to deliver the post-liberation elements of the intervention.
- The onset of the COVID-19 pandemic resulted in prison restrictions and no access to the prison site, meaning we were unable to identify if participants had been released or not at 6 months, with no access at the England site at 12 months and very limited access at the Scotland site at 12 months.
- Although the COVID-19 pandemic was undoubtedly a factor in the low percentage followed up, only 13% (18/132) of those who received the intervention were followed up at 12 months.
- Changes to the study protocol to include self-completion of follow-up data may have made it difficult for participants to complete the survey, resulting in larger proportions of missing data as compared to RA survey completion at baseline, where missing data were minimal.
- We did not include probation services in the trial, which on reflection would have strengthened our post-liberation follow-up process.

Recommendations regarding a definitive randomised trial

From the study progression criteria for a full RCT, we have identified the following recommendations:

1. Buy-in for a research trial of this nature in prison requires significant pre-study trust and relationship development, buy-in from the prison estate, governor, prison officers, peer mentors and embedded third-sector services.
2. Recruitment and randomisation of men on remand to a future APPRAISE RCT are possible, with trust in the research team an important factor.
3. Training team members of existing alcohol services to deliver APPRAISE as per protocol is possible.
4. Delivery of the in-prison APPRAISE intervention is possible and would require appropriate space, time and team member capacity.
5. Economic evaluation is possible.
6. Post-liberation intervention delivery and follow-up would only be possible if there was a robust follow-up process identified and in place.
7. Further exploration of the inclusion of and collaboration with the probation service in the service delivery and implementation of the APPRAISE intervention, and that of voluntary groups and agencies at local community levels engaged specifically in post-discharge support services, may

offer higher success rates of post-liberation intervention delivery and a more robust follow-up process.

8. PPI membership should be strengthened to reflect the complexity of the prison setting and the range of stakeholders within the criminal justice system.

Conclusion

Addressing alcohol harm in prisons, at what can be considered a 'teachable moment', can provide men with an opportunity for reflection on their risky drinking and their current position.

Many men on remand do not have the opportunity to access mainstream prison health or public health services. An intervention such as APPRAISE offers an opportunity to provide an extended behavioural-based alcohol intervention to men on remand. The APPRAISE study has identified that despite the complexities of ethical approval and the time taken to build relationships and trust, it is possible to undertake key elements of a future RCT, but not all, namely follow-up. Further focused research needs to be undertaken to explore, identify and develop a robust process and system to optimise follow-up post liberation.

The evidence base to meet the needs of men on remand in relation to risky drinking remains weak. However, there are opportunities to build on the work of APPRAISE to ensure equal access to interventions that have the potential to positively impact their relationship with and use of alcohol.

Trial registration

This trial is registered as Current Controlled Trials ISRCTN36066.

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Chapter 1 Introduction and background

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For individuals in contact with the criminal justice system, the prevalence of at-risk drinking is far higher than in the general population. There is very little evidence of efficacy or effectiveness of alcohol interventions in reducing risky drinking amongst those in the criminal justice system and in particular men on remand in prison. This is compounded by the limited evidence for the optimum timing of delivery, recommended length, content, implementation and economic benefit of an extended alcohol intervention in the prison setting. APPRAISE (A two-arm parallel group individually randomised Prison Pilot study of a male Remand Alcohol Intervention for Self-efficacy Enhancement) will therefore provide vital evidence to inform a future definitive randomised controlled trial (RCT) of an extended alcohol intervention for men on remand in prison. The remainder of this chapter provides the background research and rationale for APPRAISE, presents the study aims and objectives, and concludes by describing the structure of the remaining chapters of this report.

Prevalence of at-risk drinking in the United Kingdom criminal justice system

Hazardous drinking is defined as a repeated pattern of drinking that increases the risk of psychological or physical problems,² whereas harmful drinking is defined by the presence of these problems.³ Drinking at hazardous or harmful levels is categorised as risky or at-risk drinking.

The prevalence of at-risk drinking, which includes drinking at levels that harm a person's health, is far higher amongst those in contact with the criminal justice system (73%).⁴⁻⁹ In the UK, between 51% and 83% of incarcerated people are classified as risky drinkers.¹⁰ For those on remand in prison, the prevalence is between 62% and 68%.⁸ This compares to 35% in the general population.¹¹ Furthermore, rates of alcohol dependence among those who are incarcerated (43%) are 10 times higher than for the general population.⁸

Interventions

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Alcohol brief interventions (ABIs) are a secondary prevention activity, which is aimed at people who are drinking in a pattern that is likely to be harmful to health and/or well-being. They have been frequently shown to be effective in primary health care,^{12,13} but they are typically delivered by practitioners who are not addiction specialists, to non-treatment, opportunistic populations.¹⁴

ABIs largely consist of two different approaches: simple structured advice, which after screening raises awareness through provision of personalised feedback and advice on steps to reduce drinking behaviour and its adverse consequences; and an extended brief intervention, which generally involves behaviour-change counselling. Extended brief intervention introduces and evokes change by giving the participant

the opportunity to explore their alcohol use, as well as their motivations and strategies for change. Both forms share the common aim of helping people to change drinking behaviour to promote health.¹⁴

There is a wide variation in the duration and frequency of ABIs. However, typically they consist of between one and four sessions and are very short in nature (between 5 and 60 minutes).¹⁵ They generally include personalised feedback on alcohol intake in relation to what the recommended limits are and discussion of both health and social risks, and may include setting personal targets which can include psychological and motivational interviewing.¹⁵ One example of this is using the FRAMES (Feedback, Responsibility, Advice, Menus, Empathy, Self-efficacy) approach.¹⁴

Due to the established links between risky drinking and crime and the costs to society, in both health and social care, it is important to find interventions that are effective. It has been shown that interventions that capitalise on the 'teachable moment' are conducive to behaviour change, for example where individuals consider their alcohol use within the context of their offending behaviour and its punitive consequences.^{16,17}

There is robust evidence from systematic reviews and meta-analyses to indicate that short alcohol interventions are effective in reducing alcohol consumption amongst risky drinkers in healthcare settings.^{18,19} There is very little evidence of efficacy or effectiveness of alcohol interventions in reducing risky drinking amongst those in the criminal justice system, including the prison system,^{20,21} and in particular remand prisoners. However, there has recently been evidence in the UK that alcohol (and drug) interventions can have an effect in reducing recidivism.²² Furthermore, short alcohol interventions have been shown to reduce recidivism in the probation setting.⁷ Nevertheless, there is limited evidence for the optimum timing of delivery, recommended length, content, implementation and economic benefit of an extended alcohol intervention in the prison setting. Likewise, there are weaknesses within the current evidence base with regard to the theoretical underpinnings of such interventions. This risks an intervention with a weak theoretical base and poorly specified 'active' ingredients, and which is less likely to deliver the desired outcomes.

Current evidence of alcohol interventions for those in the criminal justice system

A recent systematic review of the efficacy of psychosocial alcohol interventions for incarcerated people found that interventions in the prison setting have the potential to positively impact their relationship with and use of alcohol;²¹ however, because of small numbers and the use of different outcome measures it was not possible to conduct a meta-analysis or generalise findings. Notably, none of the studies focused on men on remand, as compared with other subgroups within prisons. Remand refers to those who are either unconvicted or convicted and unsentenced, held in custody awaiting trial and/or sentencing.

Rationale for APPRAISE

In 2021, 16% of the prison population in England and Wales, and 19% in Scotland, were remand prisoners.^{23,24} This equated to around 12,750 in England and Wales and 1700 in Scotland. Approximately 25% of individuals will not receive a custodial sentence.²⁵ People spend on average 10 weeks in custody while on remand.²⁶ During this time, many do not have the opportunity to access mainstream prison health or public health services.²⁷

It has been shown that there is a complex interplay between individual and contextual factors and risky drinking behaviours and alcohol-related crime.²⁸ Addressing alcohol harm in prisons, at what can be considered a 'teachable moment', where there is an opportunity for people to reflect on their risky drinking and their current position, could potentially reduce the risk of re-offending and costs to society, and help to address health inequalities.²⁹ While there is an inevitable uncertainty in estimating

the costs of alcohol-related crime and disorder, most estimates suggest they represent a considerable economic burden.

A recent Cabinet Office estimate reported that alcohol-related crime in England cost society £21.5 billion.³⁰ However, this estimate is outdated and there are concerns regarding the assumptions and methodological judgements used in deriving this estimate.³¹ Better-quality estimates from four high-income countries placed the total costs of alcohol at 2.6% of gross domestic product (GDP) in 2007, of which 3.5% was made up of law-enforcement costs.^{32,33} It has been shown that intervening to reduce alcohol use is cost-effective, generating both long- and short-term savings. Public Health England (PHE) estimated that every £1 invested in effective alcohol treatment brings a social return of £5. Evidence estimates that health savings of £4.3 m and crime savings of £100 m per year can be made as a result of appropriate alcohol interventions.³⁴ It has also been suggested that providing effective treatment is likely to significantly reduce the costs relating to alcohol as well as increase individual social welfare.³⁵

In 2017, the Scottish Parliament Health and Sports Committee held an inquiry into prisoner health care to consider how health and social care and medicines in prison are accessed and the effectiveness of health and social care in prison. The Inquiry report set out recommendations for the Scottish government to prepare a strategic plan to address prison social and health care.³⁶ Furthermore, National Institute for Health and Care Excellence (NICE) Guidelines [NG57]³⁷ for people in prison acknowledge that adequate healthcare provision for prisoners would reduce pressure on community services later.

Theoretical background

The intervention in this study (the APPRAISE study) builds on previous research which has explored the theoretical validity of a self-efficacy-enhancing alcohol intervention,^{38,39} and was originally tested as part of a pilot cluster RCT in a general hospital setting with evidence of a potential effect.³⁸ In our recently completed Medical Research Council Public Health Intervention Development (MRC-PHIND) funded PRISM-A (Alcohol Brief Interventions for Male Remand Prisoners) study, we undertook development and refinement of this self-efficacy-enhancing alcohol intervention within the prison setting, working with men on remand to include a synthesis of their views, with reviews of the evidence base and theoretical underpinnings.⁴⁰

This current study worked solely with men on remand. We acknowledge that there are also women prisoners who have alcohol-use disorders. There is evidence that women face different issues when imprisoned, and different barriers to services, which are important to note.^{41,42}

The PRISM-A study with men on remand, carried out in two prisons in the UK, showed a high prevalence of risky drinking [82% scored more than 8 on the Alcohol Use Disorders Test (AUDIT)]. These numbers are comparable to other findings in the prison system in the UK,⁴⁻⁸ but more than three times as high as in primary care settings.¹¹ In the PRISM-A study, we were able to access, recruit, consent and identify those who were risky drinkers. We also found high levels of willingness of staff and participants to engage with the self-efficacy-enhancing intervention we tested for acceptability, and a willingness to participate in a trial that would involve follow-up at 6 and 12 months. Analysis of 24 in-depth interviews with male remand participants demonstrated strong substantiation that the intervention could help men on remand to develop skills and strategies that would be particularly useful on liberation. We also identified through the interviews a stronger preference amongst the participants for interventions of longer duration and that such interventions should incorporate a post-liberation component.

This is an under-researched area with large gaps in knowledge. A systematic review of alcohol interventions for offenders in the criminal justice setting identified a lack of evidence-based intervention strategies and highlighted the paucity of knowledge in this area, and in particular the lack of UK-based studies and the absence of rigorous studies focusing on men on remand.^{8,21}

Given the limited evidence on the effectiveness of psychosocial alcohol interventions for men on remand, we aimed to address this gap by carrying out a two-armed individually randomised pilot study to assess the feasibility and acceptability of a self-efficacy-enhancing intervention that had been developed during the PRISM-A study.⁴⁰ The results from this study will enable us to undertake a future definitive RCT evaluating its effectiveness and cost-effectiveness.

The study aligns to the MRC framework,⁴³ using mixed methods with two linked phases conducted in two sites in the UK: Scotland and England.

Phase I involves a two-arm, parallel-group, individually randomised pilot study. The pilot evaluation provides data on feasibility and an assessment of the likely impact of the APPRAISE intervention to inform the feasibility of a future definitive multicentre RCT.

Phase II comprises a process evaluation. We have drawn on the MRC framework for process evaluation to guide the planning, design and proposed conduct of the process evaluation.⁴⁴ The aim of the process evaluation is threefold: first, to assess how the intervention was implemented, second, to undertake some preliminary exploration of change mechanisms underpinning the intervention and third, to assess the context within which the intervention is delivered.

An additional element was added to the study, as part of the study extension, during and in response to the COVID-19 pandemic. Following approval from National Institute for Health and Care Research (NIHR), we carried out a survey of prisons in Scotland, England and Wales to ascertain what alcohol services are available in male remand prisons and how coronavirus disease discovered in 2019 (COVID-19) has affected these services.

Study aims and objectives

APPRAISE comprised a two-arm, parallel-group, individually randomised, pilot study of a self-efficacy-enhancing psychosocial alcohol intervention for men on remand in prison. It aimed to provide the evidence to support the design of a future definitive multicentre RCT for which new funding would be sought. APPRAISE had the following objectives and questions.

Objective 1: to pilot the study measures and evaluation methods to assess the feasibility of conducting a future definitive multicentre, pragmatic, parallel-group RCT.

- Is it feasible to conduct a future multicentre RCT of a self-efficacy-enhancing psychosocial alcohol intervention for men on remand?
- Can we obtain reasonable estimates of the parameters necessary to inform the design and sample size calculation for a future definitive multicentre RCT? This includes standard deviations of potential continuous primary outcomes and estimates of recruitment, retention and follow-up rates.
- How well do participants complete the questionnaires necessary for a future definitive RCT?
- Can we collect economic data needed for a future definitive RCT?
- Can we access recidivism data from the Police National Computer (PNC) databases for trial participants?
- Can we access health data from routine NHS data sources for trial participants?

Objective 2: to assess intervention fidelity.

- 2a. What proportion of the interventions are delivered as per protocol?
- 2b. Is there any evidence of contamination between the two conditions and/or between those workers delivering the intervention.
- 2c. To what extent was the intervention changing process variables consistent with the underpinning theory?

Objective 3: to qualitatively explore the feasibility and acceptability of a self-efficacy-enhancing psychosocial alcohol intervention and study measures to staff and for men on remand and on liberation.

3a. How acceptable are the trial and intervention procedures (including context and any barriers and facilitators) to the following key stakeholders: men on remand in prison and on liberation; prison staff (including healthcare staff); commissioners; policy-makers and third-sector partners?

Objective 4: to assess whether operational progression criteria for conducting a future definitive RCT are met across trial arms and study sites and, if so, develop a protocol for a future definitive trial. Operational progression criteria are based on previous research results.⁷

4a. Do the two prisons invited to the study agree to take part?

4b. Based on knowledge from previous data, do at least 90 eligible participants consent to take part and be randomised across the trial arms?

4c. Do at least 70% of participants who consent to the trial receive the intervention?

4d. Are at least 60% of those who received the intervention followed up at 12 months across trial arms and study sites?

Objective 5: to carry out a survey to ascertain what alcohol services are available in male remand prisons and how COVID-19 has affected services.

Structure of the report

The report comprises nine chapters, providing details of the background to the study, the rationale for the study, and an account of the research design, methods and findings. Reporting of public patient involvement (PPI) involvement and on equality diversity and inclusion is included. The report concludes by answering whether the study met its aims and objectives, learning from the study, strengths and limitations, while reflecting on the impact of COVID-19 and details of the recommendations for a future definitive trial.

This first chapter has provided the background research and rationale for the study and presented the study aims and objectives. [Chapter 2](#) details the development of the intervention, presents the various elements of the intervention and theoretical underpinnings, and aligns itself to the Template for Intervention Description and Replication (TIDieR) framework.⁴⁵ delivery mechanisms as well as the logic model. Intervention training is also described. [Chapter 3](#) reports the pilot trial methods including the management of the study, governance as well as ethics and research clearance. [Chapter 4](#) presents the pilot trial findings, addressing Objectives 1 and 4. [Chapter 5](#) reports the qualitative process evaluation methods. [Chapter 6](#) presents the qualitative process evaluation findings, addressing Objectives 2 and 3. [Chapter 7](#) details the results of the collection and access of economic data and recidivism data for trial participants, addressing Objective 1. [Chapter 8](#) presents the APPRAISE COVID-19 survey methods and results. This was an additional piece of work we agreed with NIHR to carry out during COVID-19 and as part of the study extension period. [Chapter 9](#) presents the discussion, PPI overview, strengths and limitations, recommendations for future RCT and conclusion. The aim of PPI was to provide meaningful engagement with stakeholders who had direct experience of the prison system, ranging from those who have experience of incarceration to those who run third-sector organisations supporting people on release from prison. Our PPI co-applicant was present to contribute in all PMG meetings. The Guidance for Reporting Instrument of Patients and the Public – short form (GRIPP2) form⁴⁶ identifies the PPI study engagement (see [Appendix 1](#)). Finally, recommendations for a future definitive trial are presented. Due to COVID-19, we faced unavoidable modifications to the APPRAISE study. These are reported across chapters at the relevant points. To support completeness of reporting these unavoidable and important modifications to the trial due to COVID-19 (extenuating circumstances), we also completed the CONSERVE checklist⁹ (see [Appendix 2](#)).

Chapter 2 Intervention development

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This chapter reports on the origins of the APPRAISE intervention and its theoretical underpinnings, and provides a detailed description of the study intervention itself and the nature of the training for those delivering it. For completeness of the APPRAISE intervention reporting and replicability, the TIDieR checklist is used to support the structure of the chapter (see [Appendix 3](#)).⁴⁵

Origins and development of the APPRAISE intervention

The APPRAISE intervention is an extended self-efficacy-enhancing alcohol intervention developed for men on remand with self-reported drinking that is harmful or hazardous. The APPRAISE intervention is based on previous ABI research focusing on self-efficacy as an appropriate basis for a single-session ABI intervention to enhance drinking refusal self-efficacy in a general hospital population.^{38,39}

From this original work, additional intervention modification was undertaken as part of the PRISM-A study on the basis of our learning from interviews with men on remand ($n = 24$).⁴⁰ The aim of these interviews was to explore the acceptability of a self-efficacy-enhancing ABI,⁴⁰ including listening to feedback regarding the nine elements of the intervention: (1) preliminary discussion, (2) acquiring and providing information, (3) self-monitoring, (4) increasing awareness, (5) situation appraisal and appropriate coping strategies, (6) goal-setting, (7) relapse, (8) self-evaluation/self-reinforcement and (9) culmination.^{39,40}

The nine intervention elements of the single-session delivery were found to be acceptable to men on remand who were interviewed as part of the PRISM-A study. Feedback regarding frequency and intensity of contact identified a preference for more than one session, with a desire for additional sessions being delivered in the community following liberation. The participants reported that this would provide the opportunity to put the skills gained as a result of the single-session intervention into practice, once liberated, in real-world situations in which alcohol is widely available to them.

This evidence subsequently informed the development and refinement of the single-session ABI to an extended four-session ABI, incorporating a single session to be delivered in the prison setting with the remaining three additional booster sessions to be delivered following liberation.

APPRAISE intervention: theory, elements and logic model

The APPRAISE intervention is based on Social Cognitive Theory and designed to increase self-efficacy and other self-regulatory skills to reduce alcohol consumption. [Figure 1](#) illustrates this process in the APPRAISE logic model. According to the theory, self-efficacy is a central determinant of health behaviour change.⁴⁷ Primary sources of self-efficacy that can be targeted through interventions are mastery experience, verbal persuasion, vicarious experience and physiological state.^{39,40,47} All these sources of self-efficacy were addressed in the intervention (see [Table 1](#) for all elements of the APPRAISE intervention). Mastery experiences should become more likely due to intervention elements addressing feedback, goal-setting, planning for release, including plans for coping responses, and self-monitoring of the intervention. Verbal persuasion was delivered by the interventionist, as were ways to achieve a

INTERVENTION DEVELOPMENT

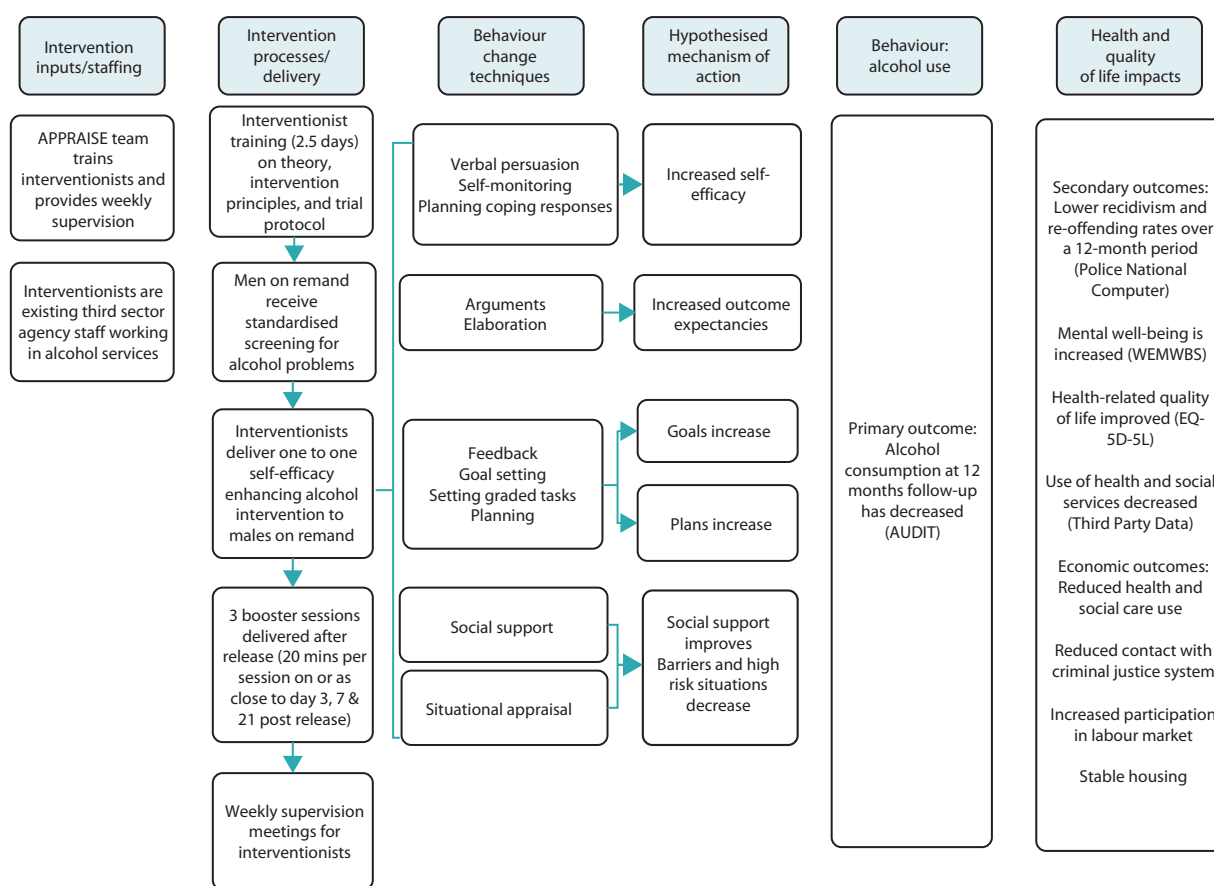


FIGURE 1 APPRAISE logic model.

calmer physiological state during high-risk situations. The intervention was designed to also increase outcome expectations by delivering arguments and elaborations and seeking social support. The intervention aimed to develop better self-regulation skills, thus enabling individuals to address their alcohol consumption behaviour. Liberation then offered participants the opportunity to implement and act on the targeted behaviours learned in the intervention in their 'real life' social context. The aim was for participants to have the ability to develop and build upon these skills and self-belief through success and mastery, with their efforts leading to the adoption and maintenance of reduced alcohol consumption. Reducing alcohol consumption can provide a sense of achievement and success, which should further increase self-efficacy.³⁹

The six proposed mechanisms of change in the logic model are: self-efficacy, outcome expectations, goals, plans, social support and high-risk situations.

1. Self-efficacy will be measured using: (i) an item on the Readiness to Change Ruler: 'How confident are you that you could reduce (or stop) drinking when you are released?' and separately (ii) the Drinking Refusal Self-efficacy Questionnaire.
2. Alcohol expectancy will be measured using the Negative Alcohol Expectancy Questionnaire.
3. Goals will be measured using an item on the Readiness to Change Ruler: 'How much do you want to reduce (or stop) drinking when you are released?'
4. Plans will be measured using: (i) an item on the Readiness to Change Ruler: 'How ready are you to reduce (or stop) drinking when you are released?' and (ii) 'I have made a detailed plan regarding when, where and how to reduce my drinking'.
5. Social support will be measured using the following items: (i) 'The people who are important to me support me in reducing my drinking' and (ii) 'I have someone I can talk to about reducing my drinking' in the Readiness to Change Ruler.

TABLE 1 APPRAISE elements

	Opening strategies
Element 1: preliminary discussion	Introduction to APPRAISE study Introduction to APPRAISE Consent, confidentiality, engagement rules, trust
Element 2: acquiring and providing information	Feedback on AUDIT score Establish perception of impact of alcohol on health and life Standard units of alcohol Recommended drinking levels Alcohol-related health problems Legal drink/drive limit Tips on reducing consumption Where to obtain information/support (prison and liberation)
Element 3: self-monitoring	Diary card – when, where, who, type of drink, why
Element 4: increasing awareness	Balance sheet – pros and cons of drinking Physiological sensations identified Alternative appraisal of somatic sensations identified Strategies to reduce
Element 5: situation appraisal and appropriate coping strategies	High-risk situations and antecedents of over-drinking identified Alternative coping strategies identified Coping strategies verbalised by participant Praise provided Strategies developed further through co-production Strategies modelled by interventionist Participant verbalises strategies and visualises them Plan for exposure to low-risk situations and avoidance of high-risk situations General control strategies: reduction in rate of drinking, sipping, drinking low-alcohol drinks and alternating between soft or low-alcohol drinks and higher-alcohol drinks
Element 6: goal-setting	Setting realistic sub-goals (short-term) Facilitating success and increasing motivation
Element 7: relapse	What happens if you relapse What caused the relapse? How do I understand relapse?
Element 8: self-evaluation and self-reinforcement	Using my alcohol diary as a means of self-evaluation and self-reinforcement Self-congratulations and rewarding my success What do I attribute my success to?
Element 9: culmination	Reflections and conclusions. Plans and goals reiterated and confirmed

6. Barriers and high-risk situations will be measured using the following items: (i) 'I know in which situations it will be difficult for me to reduce my drinking' and separately (ii) the question 'to what extent do you consider this to be a high-risk situation for you?' in the Drinking Refusal Self-efficacy Questionnaire.

APPRAISE intervention materials

The APPRAISE intervention consisted of three documents:

1. APPRAISE Intervention Tool [V3_20October] (see [Appendix 4](#))
2. Intervention Tool Postcard (Unit card) [V2_25Sept19] (see [Appendix 5](#)) and
3. Follow-up Manual [V1_25 Sept19] (see [Appendix 6](#)).

These documents together supported the delivery of the behaviour-change methods, linked directly to the determinants/mechanisms of change that focused on the targeted behaviour, in order to achieve the desired outcomes.

Document 1: the APPRAISE Intervention Tool was an eight-page, double-sided A5 booklet containing all nine Intervention Elements ([Table 1](#): APPRAISE elements) and was the material used for the first intervention session.

Document 2: the Intervention Tool Postcard (Unit card) was postcard-sized and provided brief information about recommended drinking levels and units that was given to individuals to take away with them during Element 2 of the first intervention session.

Document 3: the Follow-up Manual was a six-page double-sided A5 booklet for use during all three post-liberation follow-up intervention sessions (Day 3, Day 7 and Day 21). The manual contained Element 1 as a way of re-establishing dialogue following on from the first intervention session, followed by Elements 5–9 of the APPRAISE intervention.

APPRAISE intervention procedures

The APPRAISE intervention was delivered in a safe, non-judgemental environment where engagement rules and trust were affirmed. This was with the aim of enabling and supporting the intervention activities. The APPRAISE Intervention Tool booklet contained '*Things to remember*' – notes for the interventionist to support the delivery of the intervention elements. This reminded the interventionist of the key areas within each element to be emphasised during intervention delivery. It also provided a mechanism of supporting intervention fidelity in relation to delivery of the intervention changing process variables consistent with the underpinning theory.

While the APPRAISE intervention has specific core elements, the intervention itself was person-centred in that it became individualised during the course of the participant-interventionist conversation and subsequent intervention delivery.

The starting point for this approach was assessing participants' readiness to change by way of completion of the Readiness to Change Ruler.⁴⁸ The scoring then provided information and understanding as to where the participant was in relation to their readiness to change. Similarly high-risk drinking situations were identified during Element 1 to provide further information that would build upon all the elements moving forward. This initial discussion was key to ensuring the interventionist was able to understand, recognise and respond effectively in relation to gauging how participants perceived their drinking and their motivation to change and how they used alcohol in their lives. Discussing answers to both the Readiness to Change Ruler and high-risk situations supported the delivery of an

individual realistic starting point for both participant and interventionist. Depending on the results of these elements, each element could then be individualised to participants' particular needs.

The participants AUDIT score, obtained following screening, was shared with them during Element 2 to feedback and provide information on recommended drinking levels, units and the impact alcohol may have been having on themselves, others and their life on the whole. Giving the Intervention Tool Postcard (Unit card) to the participant provided an easy-to-carry postcard-size information resource which could be studied following intervention sessions and post liberation.

In addition, the booklet had a perforated page that was removed at the end of Element 8 and was given to and kept by the individual receiving the intervention. The page was a takeaway 'personal planner' with reinforcement triggers of the key behaviour-change methods and mechanisms.

APPRAISE intervention training

Intervention training was provided at both study sites to members of the Change Grow Live (CGL) team who were supporting the delivery and which included those practitioners who would be delivering the APPRAISE intervention itself.

The aims of the training for interventionists were to:

- have an awareness and understanding of the APPRAISE intervention
- understand AUDIT Scores and what they mean
- have an awareness of the Readiness to Change Ruler and Risky Situation Assessments and why we use them
- be confident working with the APPRAISE booklet; and
- be able to deliver the APPRAISE intervention.

The training material '(APPRAISE training plan1_VG) (see [Appendix 7](#)) for the APPRAISE intervention provides interventionists with:

- a written brief on the theoretical underpinnings of the intervention
- paper copies of the three APPRAISE intervention documents
- paper copy of the APPRAISE elements
- online narrated review of the three APPRAISE intervention documents
- intervention delivery video
- paper copy of the AUDIT instrument; and
- opportunity to engage in intervention delivery role-play exercises.

On completion of the training, a six-question Likert scale survey (see [Appendix 8](#)) as a form of evaluation of the training undertaken was self-completed by interventionists and CGL team members from both study sites (see [Intervention implementation](#)).

APPRAISE intervention delivery mode and location

Following completion of baseline assessments and randomisation, those allocated to the intervention arm were invited to attend their first of three intervention sessions. This was delivered face to face by the identified interventionist at each study site. It was delivered in a quiet room either on the residential wing, a landing or in one of the CGL designated rooms and was delivered at both sites by CGL interventionists. Four members of the CGL team from the England study site and two members from the Scotland study site attended the intervention delivery training.

At the beginning of data collection, both study sites had allocated two interventionists. Due to resource capacity and sick leave during the data-collection period, Scotland had one interventionist and England had two interventionists able to deliver the interventions.

We asked the three interventionists for details of their education and previous training experience. This request was made during COVID-19 when we were only able to communicate via e-mail. While there was a willingness to provide the information, due to the incapacity of interventionists to find the time we were unable to obtain that information.

The APPRAISE intervention comprises nine elements (*Table 2*) to be delivered in four steps: Step 1 comprises a 1 × 40-minute face-to-face session in which the nine elements are covered and delivered by a trained practitioner from CGL in the prison setting. Steps 2, 3 and 4 are 20-minute sessions conducted by phone, on or close to day 3, 7 and 21 post liberation. The post-liberation sessions include elements 1 (preliminary discussion), 5 (situation appraisal), 6 (goal-setting), 7 (relapse), 8 (self-evaluation/self-reinforcement) and 9 (culmination). *Table 3* provides details of the intervention sessions, duration, intensity and dose.

TABLE 2 APPRAISE intervention element descriptions

Element	Description
Element 1: preliminary discussion	Participants re-introduced to study and APPRAISE, made aware of the purpose of the engagement, emphasising confidentiality, consent and trust. Mirrored aspects of 'Opening strategies'. ⁴⁸ Readiness to Change Ruler completed and list of high-risk drinking situations identified on a scale of 0–5.
Element 2: acquiring and providing information	Focus on participants reported level of alcohol consumption as per screening AUDIT score and their perception of the impact of alcohol on their health and life. Six key pieces of information provided: Standard units of alcohol; Recommended drinking levels; Alcohol-related health problems; Legal drink/drive limit; Tips on reducing consumption; Where to obtain information/support. Intervention Tool Postcard (Unit Card) given to keep.
Element 3: self-monitoring	Participant introduced to the idea of keeping a diary post liberation: they are guided through how they would complete the diary with the support of the interventionist. They are informed that the diary will encourage them to reflect and record when, where and with whom, if anyone, they have consumed alcohol, why and what type of drink to enable self-monitoring on liberation, acting as a source of self-evaluation and record of progress. ⁴⁹
Element 4: increasing awareness	Pros and cons of drinking recorded on a balance sheet of drinking. Physiological drinking sensations experienced discussed with strategies to identify and reduce with alternative appraisal of somatic sensations identified, for example alternative stress-relieving strategies, such as relaxation techniques rather than alcoholic drink.
Element 5: situation appraisal and appropriate coping strategies	High-risk situations and antecedents of over-drinking identified and alternative coping strategies considered. Situations are identified in order that appropriate coping and response strategies are practised prior to the situation next occurring. Coping strategies verbalised by participant, praise provided and strategies developed further and modelled by interventionist. Participants talk through the strategy and visualise themselves carrying it out (covert modelling). The better ingrained and more automated the response is, the higher one's self-efficacy and the lower the probability of relapse. ⁵⁰ Avoidance of high-risk situations may be necessary in the first instance (although can be unrealistic) until mastery with low-risk situations is achieved (progressive mastery). Core set of general control strategies also discussed: reduction in rate of drinking, sipping drinks, low-alcohol-content drinks and alternating between soft or low-alcohol and higher-alcohol drinks.
Element 6: goal-setting	Realistic sub-goals set, short-term so that unachievable goals initially avoided. Setting proximate goals in relation to self-efficacy development can facilitate success promptly with increase in motivation towards accession to distal goals. This approach likened to a stepladder approach of focusing on the few rungs in front of you. For example, goal may be to reduce quantity of alcohol consumption on certain days of the week.

TABLE 2 APPRAISE intervention element descriptions (continued)

Element	Description
Element 7: relapse	Likelihood of relapse discussed and importance of identifying the events that resulted in the relapse, not attributing relapse to stable causes, such as ability or uncontrollable causes as this can result in lowering of self-efficacy and expectation of future success. Attribution to relapse discussed and focus on appraisal due to unstable causes and not to a personal failure/ability. Majority of relapses occur as a result of emotional distress, social pressure and interpersonal conflicts, therefore appropriate relapse strategies are discussed and practised.
Element 8: self-evaluation and self-reinforcement	Reward for success through affective self-reaction, for example, self-congratulations. The alcohol diary is a means of self-evaluation and self-reinforcement, success is attributed to their ability and skill, resulting in feelings of pride.
Element 9: culmination	Reflection of preceding discussion, conclusions drawn, praise given on progress and decisions made. Plans and goals reiterated and confirmed. Agreement regarding mobile phone follow-up on liberation.

TABLE 3 Outline of APPRAISE intervention – sessions, duration, intensity, dose

Element	Elements of intervention ^a	Enhancing self-efficacy	Delivery mode and location	Number of sessions and location
1	Preliminary discussion	Verbal persuasion	Face-to-face (P) Mobile phone (L)	1 (P) 3 (L)
2	Acquiring and providing information	Verbal persuasion	Face-to-face (P)	1 (P)
3	Self-monitoring	Verbal persuasion	Face-to-face (P)	1 (P)
4	Increasing awareness	Physiological state	Face-to-face (P)	1 (P)
5	Situation appraisal and appropriate coping strategies ^a	Vicarious experience	Face-to-face (P) Mobile phone (L)	1 (P) 3 (L)
6	Goal-setting ^a	Verbal persuasion	Face-to-face (P) Mobile phone (L)	1 (P) 3 (L)
7	Relapse ^a	Performance attainment	Face-to-face (P) Mobile phone (L)	1 (P) 3 (L)
8	Self-evaluation/self-reinforcement ^a	Performance attainment	Face-to-face (P) Mobile phone (L)	1 (P) 3 (L)
9	Culmination	Performance attainment	Face-to-face (P) Mobile phone (L)	1 (P) 3 (L)

P, prison; L, liberation.

^a Elements 5–8 highly rated by participants in the intervention and feasibility study (PRISM-A) form key focus of intervention delivery by mobile phone on liberation.

APPRAISE post-liberation intervention modifications due to coronavirus disease discovered in 2019

The outbreak of the COVID-19 pandemic had a significant impact on the project. Due to COVID-19 restrictions, access to both study sites for the RAs and research team was halted. It became clear quite quickly that the impact of the pandemic resulted in there being limited or no capacity for the interventionists to follow up on liberation data of those in the intervention group or deliver

post-liberation intervention. Prison staff were also unable to provide us with liberation data, as their focus was rightly on the COVID-19 response in prisons. Similarly, RAs were unable to go into the prisons and access liberation data.

Modifications on timing of the post-liberation intervention delivery were made in an attempt to provide flexibility, if there was an opportunity to deliver the post-release interventions. This meant that post-release intervention sessions could still be attempted where possible with the originally intended interval between them maintained to the extent possible (4 days between sessions 2 and 3, and 14 days between sessions 3 and 4).

Chapter 3 Pilot trial methods

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APPRAISE was a two-arm, parallel-group, individually randomised pilot study of a self-efficacy-enhancing psychosocial alcohol intervention for men on remand in prison.¹ It was intended to provide the evidence to support the design of a future definitive multicentre RCT, for which new funding would be sought.

The study aligned to the MRC framework,⁴³ using mixed methods with two linked phases.

- Phase I involved a two-arm, parallel-group, individually randomised pilot study. The pilot evaluation provides data to inform the feasibility of a future definitive multicentre RCT.
- Phase II comprised an embedded process evaluation. The MRC framework for process evaluation guided the planning, design and proposed conduct of the process evaluation.⁴⁴

This chapter presents the methods of the Phase I pilot trial with the pilot trial findings presented in [Chapter 4](#), and relates to research Objectives 1 and 4 in our protocol. Objectives 2 and 3 are discussed in later chapters.

The primary research objectives of the Phase I pilot study were:

Objective 1: to pilot the study measures and evaluation methods to assess the feasibility of conducting a future definitive multicentre, pragmatic, parallel-group RCT.

- 1a. Is it feasible to conduct a future multicentre RCT of a self-efficacy-enhancing psychosocial alcohol intervention for men on remand?
- 1b. Can we obtain reasonable estimates of the parameters necessary to inform the design and sample size calculation for a future definitive multicentre RCT? This includes standard deviations of potential continuous primary outcomes and estimates of recruitment, retention and follow-up rates.
- 1c. How well do participants complete the questionnaires necessary for a future definitive RCT?
- 1d. Can we collect economic data needed for a future definitive RCT?
- 1e. Can we access recidivism data from the PNC databases for trial participants?
- 1f. Can we access health data from routine NHS data sources for trial participants?

Objective 4: to assess whether operational progression criteria for conducting a future definitive RCT are met across trial arms and study sites and, if so, develop a protocol for a future definitive trial. Operational progression criteria are based on previous research results.⁷

- 4a. Do the two prisons invited to the study agree to take part?
- 4b. Based on knowledge from previous data, do at least 90 eligible participants consent to take part and be randomised across the trial arms?
- 4c. Do at least 70% of participants who consent to the trial receive the intervention?
- 4d. Do at least 60% of those who received the intervention get followed up at 12 months across trial arms and study sites?

Setting and site selection

The study occurred in two study sites hosting men on remand within the prison estate of the Scottish Prison Service (SPS) and His Majesty's Prison and Probation Service (HMPPS), one in Scotland and one in England respectively.

The complexity of the prison estate cannot be overestimated. Our learning from previous work had highlighted how essential it was to engage with our colleagues within the prison system with regard to the study. Prison governors and associated personnel at both prison sites were involved in discussions regarding the study while we were developing the initial funding application. We had built good operational knowledge as well as relationships with key personnel over the course of securing funding.

The prison governor at each site was the primary gatekeeper and contact for the study team. This proved vital in previous prison research work and again for this study. In the months leading up to data collection commencing, while waiting for ethics approval, we had several meetings with the governors at both study sites and also with appointed research study contacts for day-to-day operations. Visits were made to both prison sites to share details of the study with prison officers, health professionals and CGL over several months. The strong relationships formed were essential to our understanding of the complexities of the prison estates and the day-to-day work carried out there, providing insight into how we could undertake the research study with minimal disruption to the estate operations.

Eligibility criteria

Men detained on remand had to meet the following criteria:

- Inclusion criteria
 - detained in either the SPS Scottish study site or His Majesty's Prison Service (HMPS) North East England study site
 - have been in the prison setting for 3 months or less
 - aged 18 years and over
 - informed consent given
 - ≥ 8 on the AUDIT screening tool⁵¹
- Exclusion criteria
 - previously recruited to the study
 - unable to give informed consent or deemed incompetent/unable to make an informed decision regarding consent
 - identified as a risk to self and/or others by prison staff
 - judged to be under the influence of an illicit substance by prison or research staff
 - currently taking Antabuse
 - on a segregative rule under the prison rules
 - not able to understand the documents, which are in the English language, or agree to the research assistant (RA) working with them to understand them.

Prison and/or research staff were responsible for making a subjective judgement as to whether or not a remand person was under the influence of an illicit substance and whether the level of intoxication was likely to have an influence on risk or capacity to understand/consent.

Participant selection and recruitment

The prison staff were key to facilitating the selection and recruitment of study participants at both sites. At the England study site, peer induction workers were also key to this process. The RAs spent time developing good working relationships with prison staff and peer induction workers on the wing.

Information regarding males on remand entering the estate was made available to the RA with the selection and recruitment process across both study sites, as outlined in [Table 4](#).

Consent

The RA at each site reviewed the Participant Information Sheet (PIS) with men on remand who met the eligibility criteria, provided a written and verbal description of the study, answered any questions or queries and then invited them to consider participating. The RA obtained informed consent from those willing to participate. Those who did not wish to participate at this point were thanked for their time and received no further interaction with regard to participating in the study.

We were aware that freedom of consent can easily be undermined for prisoners and those in the criminal justice system, which means these individuals may be more vulnerable to exploitation or abuse by researchers: learning disabilities, illiteracy and language barriers are prevalent within these populations. The prevalence of these characteristics alongside the power differential between researcher and potential participant meant that particular care was needed to ensure that valid, freely given and fully informed consent was achieved. For the purposes of this research, we considered valid

TABLE 4 Participant selection and recruitment at each study site

Scotland	England
Men on remand were given PIS by reception staff as part of their admittance.	Men on remand were held on the Induction wing. During the first 3 weeks, men on remand who were eligible were given PIS by the RA where possible. Thereafter, the RA was able to join the daily Induction session and provide the PIS and talk about the research with all new men on remand.
Review new arrival list sheet daily. Highlight initial eligibility of potential participants (< 3 months, high risk, Rule 95 ^a).	Drug and Alcohol Recovery Team (DART) provided list of new arrivals daily. RA reviewed the list. Highlighted initial eligibility of potential participants (< 3 months, high risk).
Arrive on wing with list and checked with prison officers any on the list eligible who may be at high risk/excluded/ Rule 95. Prison officers would also identify to RA anyone who had expressed an interest to them about participating in the study. RA informed prison officer whom they wished to approach.	Arrive on wing with list and checked with prison officers any on the list eligible who may be at high risk/excluded. Prison officers and Peer Induction would also identify to RA anyone who has expressed an interest to them about participating in the study. RA informed prison officer whom they wished to approach.
If approval given, cell number identified and potential participant approached via door hatch. If they agreed to speak with RA, prison officer would open pad door and escort potential participant and RA to the interview room.	If approval given, cell number identified and potential participant approached via door hatch. If they agreed to speak with RA, prison officer would open pad door and escort potential participant and RA to the interview room.
Interview room on landing with window. RA would sit on seat nearest the door and potential participant on the other side of the table. RA reaffirmed study information, confirms exclusion and inclusion criteria and consents as appropriate. Following completion prison officer would escort participant back to the cell. If potential participant did not wish to participate, prison officer would escort them back to the cell.	Interview room glass room in middle of wing. RA reaffirmed study information, confirms exclusion and inclusion criteria and consents as appropriate. Following completion prison officer would escort participant back to the cell. If potential participant did not wish to participate, prison officer would escort them back to the cell.
<p>^a Rule 95 – the Governor may order in writing that a prisoner must be removed from association with other prisoners, either generally or to prevent participation in a prescribed activity or activities.⁵²</p>	

consent as underpinned by adequate information being provided to the potential study participants and that they had the capacity to decide for themselves. A capable person would:

- understand the purpose and nature of the research
- understand what the research involves, its benefits (or lack of benefits), risks and burdens
- understand the alternatives to taking part
- be able to retain the information long enough to make an effective decision
- be able to make a free choice; and
- be capable of making *this particular decision* at the time it needs to be made.

Participants were asked to consent to their data being used for the present study as well as consenting to long-term follow-up (beyond this study) and for appropriate linkage to be conducted.

Stopping rule/discontinuation/withdrawal criteria

Participants were reminded during the trial that they were free to withdraw at any time without having to give a reason and without it affecting their care in the prison setting or on liberation. Where a participant wished to withdraw, we honoured their wish. We also made attempts to ascertain and document the reason for withdrawal, recording this within the case report form (CRF). Participants who requested withdrawal from the study intervention were asked if they would be willing to remain in the study for the purposes of follow-up data collection. Data provided at the point of withdrawal were retained and used in the study analysis, where the patient consented to this.

Potential reasons for participants being withdrawn from the study were: participant withdrawal of consent; RA discretion in the best interest of the participant to withdraw; the RA was informed by Prison, NHS or service staff that it was in the best interests of the participant to withdraw; an adverse event required discontinuation of participation in the trial; or termination of the trial by the sponsor.

Sample size

The aim was to recruit at least 180 participants in total comprising 90 participants per study arm across the two study sites. From prison data obtained from correspondence with SPS and the Ministry of Justice (MoJ), we had estimated that approximately 50% of participants would be liberated while the rest would remain incarcerated, leaving 45 participants per study arm (90 in total) across the two study sites.

The target sample size

1. would enable us to calculate two-sided 95% confidence intervals around proportions recruited, liberated, and dropping out in each study arm with half-widths of < 0.15
2. exceeds the 30 per group recommendation of Lancaster *et al.*,⁵³ and the 35 participants per group recommendation of Teare *et al.*,⁵⁴ for estimating key unknown design parameters, for example standard deviation with sufficient precision when the primary outcome is continuous; and
3. would ensure that within each study arm within each site we were satisfying the minimum 12 per group rule of thumb of Julious for pilot trials.⁵⁵

Participant screening

The AUDIT tool⁵¹ is considered the 'gold standard' alcohol screening instrument and was developed by the World Health Organization (WHO). It is a 10-item screening tool designed to assess alcohol

consumption, drinking behaviours and alcohol-related harms. The score is the total sum derived from the 10 questions. Scores range from 0 to 40. A score of 8–15 indicates hazardous/increasing-risk alcohol use, 16–19 indicates harmful/high-risk alcohol use and 20–40 indicates likely dependence. The AUDIT has been used previously in the criminal justice system,⁷ and was found to be feasible and acceptable with men on remand in the PRISM-A study.⁴⁰

After obtaining informed consent, the RAs screened all study participants using the AUDIT. Those with a score of ≥ 8 were considered eligible for the study. Those scoring < 8 on AUDIT were thanked for their time and took no further part in the study. Those with an AUDIT score of ≥ 8 completed the baseline assessment and measures and were then randomised.

Randomisation and blinding

The randomisation process was supported by the Edinburgh Clinical Trials Unit (ECTU). Due to the restriction of the use of electronic equipment and no access to mobile phones by the research team and RAs while in the prison setting, we were unable to use a randomisation system, such as an interactive voice response telephone system or one accessed over the web. Allocation was conducted at the level of the participants, randomised to the active or control intervention using stratified block randomisation by site, via opaque sealed envelopes, based on a predetermined random number allocation carried out by the ECTU. Allocation was conducted by researchers opening the next sequential sealed opaque envelope after consent had been provided and the baseline assessment completed. Therefore, RAs were blind to group allocation at baseline.

The RAs were not involved in the delivery of the active and control interventions but were aware of study allocation of participants as the trial progressed. Allocation concealment was used whereby neither the person delivering the interventions nor the participant was aware of the study allocation until they were irrevocably entered into the trial. Both the trial statistician (RP) and health economists (AS, JB) were blind to group allocation prior to the final analysis and only had details of study participants by study number.

The trial statistician and health economists were only to be unblinded if requested to do so by the Data Monitoring and Ethics Committee Project Steering Group (DMEC), due to safety concerns. This was not necessary during the study and unblinding took place at the final analysis stage.

Self-report primary outcome measure

The proposed primary outcome measure for a future definitive study was total alcohol consumed (in units) in a 28-day period. This can be ascertained using the 28-day timeline followback questionnaire (TLFB-28).⁴³ Three other variables can be derived from the data: per cent days abstinent, drinks per drinking day (secondary outcome measures) and total number of days where alcoholic drinks are consumed. The TLFB-28 was to be conducted by the RA at time points (TP) 1 and 2.

Self-report secondary outcome measures

Secondary outcomes measures were to be completed at TPO (baseline), TP1 (6 months) and TP2 (12 months).

Alcohol use frequency, quantity (on a typical occasion) and binge drinking were assessed using the AUDIT.³⁴ In order to assess the utility of using Average Drinks per Day derived from the AUDIT at 12 months as a primary outcome measure we planned to randomise the order of presentation of

TLFB-28 and AUDIT and conduct a levels-of-agreement analysis, using TLFB-28 as a gold standard, to explore whether AUDIT is an acceptable proxy for TLFB-28. The order of presentation of TLFB and AUDIT was to be randomised in advance by a secure remote randomisation service. (See section '[Follow-up plans post-COVID 19](#)' for changes to the use of AUDIT and TLFB-28 at TP1 and TP2.)

The following study measures were to be used across the three time points.

Warwick-Edinburgh Mental Well-being Scale

Warwick-Edinburgh Mental Well-being Scale⁵⁶ comprises 14 items with five response categories, ranging from 'None of the time' to 'All of the time'. The items are all worded positively and cover both feeling and functioning aspects of mental well-being. Items are scored on a range from 1 to 5, providing a total score between 14 and 70.

Drinking Refusal Self-efficacy Questionnaire – revised

This revised version of the DRSEQ⁵⁷ assesses a person's belief in his/her ability to resist alcohol. There are 19 items, each scored on a five-point Likert scale. Responses range from 1 (I am very sure I could NOT resist) to 6 (I am very sure I could resist), providing a total score between 19 and 114. A lower score indicates less self-efficacy.

Three subscales can also be calculated: Social Pressure (SP) (score range 5–30), Emotional Relief (ER) (score range 7–42) and Opportunistic (OP) (score range 7–42).

Negative Alcohol Expectancy Questionnaire

The NAEQ⁵⁸ assesses the extent to which people expect negative consequences to occur if they drink. There are 60 items in the questionnaire. Responses are measured on a five-point Likert scale ranging from 1, which is 'highly unlikely', to 5, which is 'highly likely', providing a total score between 60 and 300. A lower score indicates more negative expectancies.

The items, themselves, are grouped onto three dimensions (same-day, next-day and continued-drinking expectations). The next-day and continued-drinking dimensions are combined to form the distal expectancy subscale (score range 39–195) and the same-day, the proximal subscale (score range 21–105).

EuroQol-5 Dimensions, five-level version

The EQ-5D-5L⁵⁹ comprises two parts. The first is a visual analogue scale and the second is a descriptive system. There are five dimensions within the descriptive system: mobility, self-care, usual activities, pain/discomfort, anxiety/depression. Each has five levels: no problems, slight problems, moderate problems, severe problems, and extreme problems. Participants convey their health state by identifying the most appropriate statement in each of the five dimensions. Each dimension score is combined to describe health state. An X is used for the visual analogue score to identify the health of the participant on that particular day.

Readiness to Change Ruler

The Readiness to Change Ruler is used to assess a person's readiness to change their alcohol use.¹⁰ There are six readiness to change rulers on the assessment form. Each ruler represents a linear scale from 1 to 10, and the participant marks on the ruler their current position in the change process.

The Readiness to Change Ruler was completed immediately prior to and immediately following the initial intervention session. This was carried out by the interventionist as part of the initial intervention delivery.

Economic Form 90

We adapted the terminology of the Economic Form 90⁶⁰ (the Economic Form 90 was originally designed for American context) (see [Chapter 7, Methods](#)) to produce a service user questionnaire to determine

social costs including in the domains of health and social care use, criminal justice involvement, unemployment, and welfare (see [Appendix 9](#)).

Baseline data collection and assessment of outcomes

Baseline data (TPO) collection was conducted prior to randomisation to prevent observer or respondent bias to questions asked or answered at baseline by knowledge of the randomisation group, with follow-up data collected at TP1 (6 months) and TP2 (12 months).

The baseline assessment data were collected via researcher-led completion of hard-copy questionnaires embedded within the Baseline CRF (TPO). Due to restrictions on the use of electronic equipment in the prison estate, we were unable to use electronic devices to record study data on site. The RA was able to answer any questions and offered clarification as needed. The baseline assessment took approximately 35 to 60 minutes. Participants also completed the Participant Locator Form (see [Appendix 10](#)). The Locator Form was used to collect information from participants about a range of methods, modes and formats of contacting them at follow-up if they were liberated. Immediately after, participants were randomised, informed of their allocation,⁶¹ and thanked for their time.

The baseline CRF recorded demographic data on age, known court dates, ethnicity, postcode [Scottish Index of Multiple Deprivation (SIMD)], relationship status, and educational achievement. In addition, baseline data on proposed secondary outcome measures were recorded via the Readiness to Change Ruler,¹⁰ the EQ-5D-5L,⁵⁹ the Warwick-Edinburgh Mental Well-being Scale,⁵⁶ the Drinking Refusal Self-efficacy Questionnaire – revised,⁵⁷ the Negative Alcohol Expectancy Questionnaire,⁵⁸ and a revised version of the Economic Form 90,⁶⁰ which recorded data relating to hospital and or A/E visits, general practitioner (GP) registration and visits, engagement with Community Psychiatric Nurse, engagement with any addiction services, emergency ambulance use, engagement with social worker, current living arrangements, ever been homeless, employment status, number of arrests and number of court visits.

Control condition (care as usual)

The control condition (care as usual) consisted of similar components at both study sites.

England control condition

All men held on remand are screened by a reception nurse on reception. If a substance misuse need is identified in the screening, they are referred to the Drug and Alcohol Recovery Team (DART) and the team then sees them the following day to complete the triage and comprehensive assessment. Those men who decline the offer of the DART service are then revisited to be given harm-reduction information and a point of contact if they need to self-refer further into their sentence. Posters are displayed around all of the establishment and anyone or any service can refer an individual to DART.

They refer to the relevant community substance misuse treatment service for an individual's release and ensure the service user understands when and where their appointment is. They also offer additional support through the gate with their Reconnect Navigators, who will support those individuals who are more complex and struggle to engage to their first appointment.

Scotland control condition

For those service users who expressed that they are looking for alcohol support, the support is person-centred and based on the person's needs. Following assessment, individuals would agree a recovery plan with their worker, the recovery coordinator, and support Scotland Mental Well-Being Scale may be alcohol awareness, harm reduction, motivational support and relapse prevention. The frequency of this would be dependent on the individuals' needs and would not be time-limited. People with alcohol issues could be referred to the relevant region's alcohol service for counselling support if they wanted this. If

the individual was returning to the region where the prison was, they would be offered through-care support from a recovery support service. This would include practical support as well as substance misuse support to help them reintegrate back into the community.

Intervention condition

Following completion of baseline assessments and randomisation, those allocated to the intervention arm were invited to attend their first of three intervention sessions. This was delivered face to face by the identified interventionist at each study site. The APPRAISE intervention was delivered at both sites by CGL.

The in-prison and post-liberation elements of the APPRAISE intervention are described in full detail in [Chapter 2](#).

Follow-up data collection

We worked with our PPI co-applicant and former community justice mentor, who mentored men in prison and on liberation, to advise on the strategies to maximise retention and follow-up. These strategies were based on her own experiences of following up her mentees on liberation and were also strategies used by CGL community justice workers. Each study participant provided their preferred method of follow-up by completing the Locator Form at baseline.

Follow-up assessments were to be attempted at 6 months (TP1) and 12 months (TP2) post-randomisation where the participant had been (a) not liberated, (b) liberated and in the community, or (c) liberated and then re-incarcerated. An 8-week window was in place for the follow-up to be completed for both of the time points. If a participant was not contacted at 6 months they were still approached again at 12 months. Follow-up was to be conducted by the RA at each study site. However, due to changes in personnel at both study sites during the study, all follow-ups for both study sites were not conducted by the RA who undertook recruitment and baseline measurements. Feedback from all RAs noted that participants responded to follow-up more positively when it was the RA they knew and remembered from recruitment and baseline. There was a sense of trust that had been built which was less evident when the new RAs were undertaking the follow-ups.

Follow-up plans pre-coronavirus disease discovered in 2019

Participants were initially followed up using one of the following methods, determined by the information they provided at baseline interview as part of the Locator Form completion:

- by phone call (at varying times and days of the week)
- via hard copies of the follow-up questionnaire sent by post with an accompanying letter to be completed and returned to the study team via a pre-paid envelope; two reminders were sent; or
- via hard copy in prison for completion with the RA.

Follow-up plans post-coronavirus disease discovered in 2019

As a result of the impact of COVID-19 on the planned follow-up strategy, modifications were made to the follow-up method, procedures, data collection format, measures, and CRFs. These were discussed by the project management group (PMG) and then at the Trial Steering Committee (TSC). Ethical approval for these amendments was sought and gained.

Additional follow-up approaches were added: text message, WhatsApp, Facebook and an electronic Qualtrics link to the survey sent by phone or e-mail. Follow-ups were to be conducted over the phone by the RA. Where study participants were identified as being incarcerated at follow-up, we would

attempt to make appropriate arrangements with the relevant prison to establish if it was possible to visit and undertake the follow-up assessment. Due to the duration of the COVID-19 pandemic, access to the England study site was still not possible and for the Scotland site, although access had started to return, it was restricted and was only possible in a few instances, by which time the study had concluded.

Participants who had given permission to use social media were also sought on Facebook and one attempt to contact them was carried out (we were limited to one attempt as per ethics committee stipulation).

COVID-19 restrictions meant that we were unable to undertake face-to-face interviews. We therefore developed alternative formats of CRFs for participants to complete at 6- and 12-month follow-ups. Our amendment allowed us to send electronic copies via Qualtrics link to participants (if they had consented to us contacting them in this way) and would allow for self-completion using hard copies. We simplified the CRFs to accommodate this and removed the TLFB-28, which required in-person administration. We therefore replaced the AUDIT and TLFB-28 with the extended-item AUDIT-C questionnaire. Previous studies have shown AUDIT-C and the TLFB-28 demonstrating excellent levels of agreement on alcohol consumption.⁴ An overall total score of ≥ 5 is deemed as AUDIT-C positive, and indicates increasing or higher-risk drinking.⁶²

The NAEQ includes three sub-sections. The final sub-section was removed as it was considered as having potential to cause distress when self-completed. This was discussed at length by the PMG and with particular input from our PPI co-applicant. The layout of the Health Econ Form was also simplified to support self-completion.

Statistical methods/analysis plan

The progress of participants through the APPRAISE pilot trial is presented in accordance with CONSORT (Consolidated Standards of Reporting Trials) guidelines⁴⁰ to allow descriptions of key parameters required for a future RCT: eligibility rates, consent, adherence, retention at follow-up and data completeness of outcome measures.

Overall statistical principles

Analyses were based on a modified intention-to-treat (ITT) population unless otherwise stated. The modified ITT population included all participants who were randomised into the APPRAISE study, if they had provided valid non-missing data for each outcome at the relevant time points. Participants were analysed according to the trial arm they were randomised to, irrespective of the treatment they received, continuing study eligibility, or compliance post randomisation. Participants who discontinued with the intervention, withdrew, or were non-compliant with the protocol were included in the ITT population, provided that valid outcome data were recorded for them.

It was assumed that any missing data were missing-completely-at-random (MCAR). Although a multiple imputation analysis was specified in the protocol paper, this approach is only valid when the proportion of missing data lies between 5% and 40%.⁶³ Very high amounts of missing data at follow-up meant that no multiple imputation was performed and all missing data were left as missing.

Lancaster *et al.*⁵³ recommend that the analysis of a pilot study should be mainly descriptive and focus on confidence intervals. Therefore, the analyses focused on calculating descriptive statistics and two-sided 95% confidence intervals rather than on hypothesis tests or *p*-values in general. Outliers were identified by viewing boxplots and/or histograms of the outcome variables of interest and were queried at the data-checking stage where an error had been suspected. All analyses included outliers as standard unless otherwise stated.

Primary outcome measure

The primary outcome was to be collected at TP0, TP1 and TP2. The primary outcome for the pilot study was alcohol consumed in the 28 days prior to TP2, assessed using the extended Alcohol Use Disorders Identification Test – Consumption (AUDIT-C). This was calculated using questions 1 and 2 of the AUDIT-C, which address frequency and quantity of alcohol consumed.^{64,65} Responses to question 1 were recoded; Never = 0, Monthly or less = 0.25, 2–4 times per month = 0.75, 2–3 times per week = 2.5, 4–5 times per week = 4.5, 6 or more times per week = 7. Question 2 was recoded; 1–2 drinks = 1.5, 3–4 = 3.5, 5–6 = 5.5, 7–9 = 8, 10–12 = 11, 12–14 = 13 and more than 14 = 16. The weekly consumption was derived by multiplying the recoded values for question 1 and question 2 together and then multiplying the product by 4 to derive 28-day consumption.

The original AUDIT-C measure is known for right censoring quantity of consumption and as a consequence being less responsive to change in heavy drinkers. Extending the AUDIT-C response categories we aimed to have a trade-off between minimal completion burden and granular data. We initially tested our assumptions about categorisation of consumption in data sets that contained the extended AUDIT-C and TLFB and found minimal benefit of allocating a daily consumption value over 16 units per drinking day, something that has been further confirmed in other studies using the extended AUDIT-C to assess weekly consumption.⁶⁶

Secondary outcome measures

The secondary outcome measures are the Readiness to Change Ruler,¹⁰ the EQ-5D-5L,⁵⁹ the Warwick-Edinburgh Mental Well-being Scale,⁶⁷ the Drinking Refusal Self-efficacy Questionnaire – revised,⁵⁷ the Negative Alcohol Expectancy Questionnaire,⁵⁸ and a revised version of the Economic Form 90.⁶⁰ Measurements were recorded at baseline (TP0) using a combination of self-completion and researcher-led completion of a hard-copy questionnaire. These measures were to be collected again at 6 months (TP1) and 12 months (TP2) post randomisation where participants were (a) not liberated, (b) liberated and in the community or (c) liberated but back in prison.

Recruitment, adherence and follow-up retention analyses

Trial management data, the Intervention Delivery Logs, and follow-up questionnaires were used to assess recruitment, eligibility, adherence, and retention at follow-up. For recruitment and eligibility, the following trial management data (recorded from the initiation of the trial until full recruitment is reached) were considered by each study site where possible:

- number entering on remand
- number given information sheet
- number approached
- number consented
- number randomised and
- number followed up at each time point.

These numbers were used to calculate appropriate proportions, which are reported for the entire study and by site. Exact 95% confidence intervals were calculated for key overall proportions relating to recruitment, adherence and retention.

Descriptive summaries

Descriptive statistics were calculated for the following variables:

- baseline characteristics
- numbers receiving the in-prison and post-liberation intervention
- duration of the in-prison intervention sessions
- reasons why the in-prison intervention was not delivered
- numbers followed-up

- reasons for being unable to follow-up
- information on data completeness at baseline and follow-up time points
- the primary outcome (28-day alcohol consumption)
- secondary outcomes.

In general, categorical variables were presented as counts and percentages. Discrete and continuous variables were presented using the mean, median, standard deviation (SD), minimum, maximum, and number of responses, split by randomisation group for the entire study and by study site.

A side-by-side boxplot was produced to show the age distribution of participants randomised to each site.

Primary outcome analyses

Means and 95% confidence intervals were calculated for 28-day alcohol consumption in each randomised group and site.

For the primary analysis, the mean difference in alcohol consumption at 12 months post randomisation (TP2) between the two randomisation groups was calculated (with 95% confidence intervals), both overall and stratified by site. Linear regression was used, adjusting for baseline alcohol consumption (TPO).

The mean difference was calculated assuming that any missing outcome data were missing-at-random (MAR) and only patients with AUDIT-C data collected at TPO and TP2 were included (i.e. we performed a complete-cases analysis).

Secondary outcome analyses

The psychometric measurements were treated as secondary outcomes, and descriptive statistics were calculated based on these, stratified by site and randomised group.

[Table 5](#) contains information on the number and type of measurements for each psychometric tool.

Information on the derivation of the psychometric outcome measures is provided below, along with details of how missing individual item scores were accounted for in the calculation of total scores.

1. AUDIT-C: the primary outcome is alcohol consumed in the 28 days prior to TP2, assessed using the extended AUDIT-C. This is calculated using questions 1 and 2 of the AUDIT-C, which address frequency and quantity of alcohol consumed. Responses to question 1 were recoded; Never = 0, Monthly or less = 0.25, 2–4 times per month = 0.75, 2–3 times per week = 2.5, 4–5 times per week = 4.5, 6 or more times per week = 7. Question 2 was recoded; 1–2 drinks = 1.5, 3–4 = 3.5,

TABLE 5 Overview of the psychometric outcome measurements recorded in the baseline and follow-up questionnaires

Tool	Abbreviation	Number and type of measurements
Alcohol Use Disorders Identification Test for consumption	AUDIT-C	1 score (0–12) treated as discrete and categorical
Readiness to Change Ruler	–	6 scores (1–10) treated as discrete
EQ-5D-5L	–	5 categorical variables and 1 score (0–100) treated as discrete
Warwick-Edinburgh Mental Well-being Scale	WEMWBS	1 score (14–70) treated as discrete
Drinking Refusal Self-efficacy Questionnaire – revised	DRSEQ-R	1 score (19–114) treated as discrete
Negative Alcohol Expectancy Questionnaire	NAEQ	2 separate scores (21–105; 18–90) treated as discrete

- 5–6 = 5.5, 7–9 = 8, 10–12 = 11, 12–14 = 13 and more than 14 = 16. The weekly consumption was derived by multiplying the recoded values for question 1 and question 2 together and then multiplying the product by 4 to derive 28-day consumption. The AUDIT-C was also used to calculate a total score, which was calculated as the sum of the first three questions of the AUDIT questionnaire. The first question was scored as 0 (never), 1 (monthly or less), 2 (2–4 times a month), 3 (2–3 times a week), 4 (4–5 times a week, or 6 or more times a week); the second question scored as 0 (1 to 2), 1 (3 to 4), 2 (5 to 6), 3 (7 to 9), 4 (10 to 12, 12 to 14, or more than 14); and the third question scored as 0 (never), 1 (less than monthly), 2 (monthly), 3 (weekly), 4 (daily or almost daily). If any participants answered 'Never' to AUDIT-C Q1 then their overall score was 0. Otherwise, if any of the first three questions were missing then the AUDIT-C was also missing.
2. The Readiness to Change Ruler is used to assess a person's readiness to change their alcohol use. There are six readiness to change rulers on the assessment form. Each ruler represents a linear scale from 1 to 10, and the participant marks on the ruler their current position in the change process. The score for each question was regarded as the integer value closest to the mark made by the participant on the ruler. No missing-value imputation was performed.
 3. EQ-5D-5L consists of five descriptive dimensions (mobility, self-care, usual activities, pain/discomfort, anxiety/depression) and a question on how the respondent would rate their health. Each descriptive dimension has a score of 1–5 with levels: no problems (1), slight problems (2), moderate problems (3), severe problems (4), and unable to or extreme problems (5). The visual analogue score consists of a scale numbered from 0 ('best health you can imagine') to 100 ('worst health you can imagine'). No missing-value imputation was performed.
 4. The WEMWBS consists of 14 items with five response categories. Each item was scored on a range from 1 to 5, and a total score was calculated with a possible range of 14 to 70. A higher WEMWBS score indicates a higher level of mental well-being. The total score was treated as missing if 50% or more of the individual items were missing. If fewer than 50% of the items were missing, then each of the missing individual item scores was imputed using the mean of the remaining non-missing questions before calculation of the total score.
 5. The DRSEQ-R consists of 19 items with five response categories following a Likert scale. Each item is scored on a range from 1 to 6, providing a total score between 19 and 114. A lower score indicates less self-efficacy to resist alcohol. The total score was treated as missing if 50% or more of the questions were missing. If fewer than 50% of the items are missing, then each of the missing individual item scores was imputed using the mean of the remaining non-missing questions before calculation of the total score.
 6. The NAEQ at baseline consists of 60 items with five response categories following a Likert scale. Each item is scored on a range from 1 to 5, with 1 indicating 'highly unlikely' and 5 indicating 'highly likely'. The NAEQ was used to generate two expectancy scores: same day (sum of items 1–21) and next day (sum of items 22–39).⁶⁸ Each expectancy score was treated as missing if 50% or more of the questions were missing. If fewer than 50% of the items were missing, then each of the missing individual item scores was imputed using the mean of the remaining non-missing questions before calculation of the expectancy score. A higher score indicates more negative expectancies of drinking alcohol.

Logic model analyses

An exploratory analysis was performed to explore key behavioural markers (mediators) of change and how these may influence the effect of the delivered intervention. The six proposed mechanisms of change were: self-efficacy, outcome expectations, goals, plans, social support and high-risk situations.

In exploratory analysis, Kendall's tau correlation coefficients were calculated to assess the correlation between each of the mechanisms of change and the primary outcome (AUDIT-C) at 12 months.

Software

Statistical Product and Service Solutions (SPSS) version 27 (IBM Corp. Released 2020. IBM SPSS Statistics for Windows, Version 27.0. Armonk, NY: IBM Corp) was used to produce the side-by-side

boxplot of age. RStudio version 1.4.1103 [RStudio Team (2021). RStudio: Integrated Development Environment for R. RStudio, PBC, Boston, MA. www.rstudio.com/] was used to analyse the screening and recruitment data and produce the recruitment graph. All other tables and analyses were produced using statistical analysis system (SAS) software version 9.4 (SAS Institute Inc., Cary, NC, USA).

Ethics approval

Ethics approval for the study was sought and secured from five different organisations:

1. Integrated Research Application System (IRAS) (East of Scotland Research Ethics Service) REC reference 19/ES/0068, IRAS project ID 261003
2. HMPPS Ministry of Justice, National Research Committee reference 2019240
3. SPS Research Access and Ethics Committee (RAEC) approval letter dated 13 June 2019
4. the University of Scotland School of Health in Social Science Nursing Studies Research Panel approval letter dated 26 September 2019
5. NHS R&D in Scotland (R&D No. 2019/0268).

The University of Edinburgh acted as trial sponsor (CAHSS 18/2/02). A model non-commercial agreement (mNCA) was in place for the England study site but not for the Scotland study site following discussions with Health Research Authority (HRA). The mNCA is strongly recommended as the agreement between sites for clinical trials/investigations; it is not mandatory for a non-commercial study. For the Scotland study site it was not considered the appropriate mechanism for agreement and therefore a site agreement between the Sponsor and the SPS site was drawn up and put in place. The agreement for this took many months of discussion with stakeholders and was a long protracted process causing delays to the start of recruitment at the Scotland study site.

Management of the study

Two groups were in place to support the research management of the study. The PMG was convened and comprised the chief investigator (CI), co-applicants, our PPI member who was also a co-applicant, the Project Manager and researchers from both study sites working on the project. Professor Jeremy Bray chaired the PMG. The frequency of the PMG meetings was determined by the needs of the project. The PMG's responsibility was to ensure appropriate and timely study implementation. The constitution and composition of the PMG and TSC were in line with NIHR research governance guidelines.

The TSC was appointed to provide independent assessment of the study progress, adherence to the protocol, patient safety and well-being of study participants, consideration of new information of relevance to the progress of the study and support of decision-making on the merits of a future definitive trial. A subcommittee DMEC was also convened following approval from NIHR-PH as this was a pilot feasibility study. It was agreed that because the study was a feasibility study rather than a definitive trial, it was sufficient to have the group as a subcommittee rather than having it sealed off as a separate entity. The terms of reference (ToR) for the TSC with the ToR for the subcommittee added were agreed (see [Appendix 11](#)).

Research governance

The study was managed and conducted in accordance with the Medical Research Council's Guidelines on Good Clinical Practice in Clinical Trials (www.mrc.ac.uk), which included compliance with national and international regulations on the ethical involvement of patients in clinical research (including the Declaration of Helsinki). Before the study commenced, all required approvals were obtained and all conditions of approvals were met. All staff assisting with the study were adequately informed about the protocol and their trial-related duties.

The study was conducted at each site in compliance with the study protocol and all protocol amendments. All forms, reports, and other records were designed to maintain participant confidentiality. All data were held in a secure environment with participants' information identified by a unique participant identification code. All records were kept in a secure storage area in compliance with the Data Protection Act (2018) and the General Data Protection Regulation (GDPR) 2019 regarding collection, storage, processing and disclosure of personal information. Trial management support was provided by the Clinical Trials Unit we were working with.

Changes to the original study protocol

During the course of the study, a number of amendments were made to the study protocol, the majority as a result of the impact of the COVID-19 pandemic. Several of these changes have been discussed above, but this section provides a comprehensive catalogue of substantial changes. All proposed amendments to the study were discussed by the CI, PMG and TSC. Once agreed, the process involved discussion with the study sponsor, whose responsibility it was to determine if an amendment was substantial or not.

Substantial amendments were submitted to the East of Scotland Research Ethics Service Committee by the CI, on behalf of the sponsor. Once approval was obtained, changes to the study protocol were then implemented. ([Table 6](#) provides details of the substantial protocol changes approved and made.)

TABLE 6 Substantial APPRAISE protocol changes

Original aim	Changes to protocol
Changes to recruitment	<p>Our original aim was to recruit 180 participants across two prison sites, Scotland (N = 90) and England (N = 90). Recruitment began in England on 2 December 2019 and was completed on 28 February 2020 (N = 90). Recruitment began in Scotland on 15 January 2020 and was halted due to COVID-19 restrictions on 13 March 2020 (N = 44).</p>
Changes to intervention delivery	<p>As a result of COVID-19, recruitment at HMP Scotland was halted on 13 March and a decision to stop any further recruitment was made on 12 June. Total consented: 134. Total included: 132 (90 in England, 42 in Scotland). This change was approved on 16 July 2020.</p>
Changes to 6- and 12-month follow up timing	<p>Unfortunately, due to COVID-19 restrictions, outstanding interventions in Scotland (N = 5) were not possible and post-release intervention sessions at both sites were halted. Post-release intervention sessions had to be halted because we were unable to access release data in the prisons. This meant that intervention sessions could not be delivered on or as close as possible to days 3 (session 2), 7 (session 3) and 21 (session 4) post-release. It was agreed that post-release intervention sessions should still be attempted where possible and that the originally intended interval between them should be maintained (4 days between sessions 2 and 3, and 14 days between sessions 3 and 4).</p>
	<p>Due to the impact of COVID-19, we extended the possible follow-up window from 8 weeks to 12 weeks. This was to allow for both post-release intervention delivery and 6- and 12-month data collection. This change was approved on 21 September 2020.</p>

TABLE 6 Substantial APPRAISE protocol changes (continued)

Original aim	Changes to protocol
Changes to 6- and 12-month follow-up procedures	<p>Due to the impact of COVID-19, face-to-face follow-ups were no longer appropriate. Therefore, an amendment was sought to allow hard copies and electronic copies of the survey to be sent to participants.</p> <p>This change was approved on 21 September 2020.</p>
Changes to 6- and 12-month CRFs	<p>To allow for the self-completion of questionnaires, a number of changes were required.</p> <p>These changes were approved on 21 September 2020.</p> <ul style="list-style-type: none"> • The timeline followback-28 (TLFB-28) was removed from the follow-up CRFs. This is because this is not an appropriate self-completion tool. • The Negative Alcohol Expectancy Questionnaire includes three sub-sections. The final sub-section was removed as the project management group identified it as having the potential to cause distress. • The layout of the Service Use Questionnaire was simplified to support self-completion.
Changes to interview schedules	<p>To explore the unique impact of COVID-19 on the feasibility of this and any future project, additional questions have been added to our interview guides. The questions will allow us to identify whether changes to service utilisation and delivery have occurred as a result of COVID-19 and establish whether there are likely to be long-term impacts and changes as a result. This will help inform the feasibility of any future trial.</p> <p>This change was approved on 4 September 2020.</p>
Inclusion of a survey	<p>We have proposed designing and distributing a short survey to all UK prisons with a male remand population to ask what their current ABI delivery process is, how this has been impacted by COVID-19, and whether they would be willing to work with us on a potential future RCT to explore the efficacy of a self-efficacy-enhancing psychosocial alcohol intervention. This information would inform the feasibility of a future RCT.</p>
Changes to proposed end date	<p>We requested a 9-month extension which changed the end date from 31 March to 31 December 2020.</p> <p>This change was approved on 4 September 2020.</p>

Chapter 4 Pilot trial results

Data regarding the number of participants consented, recruited and randomised to the study are presented. In addition, we present data on baseline characteristics and primary and secondary outcome measures. Information on data completeness across the three time points is provided. Numbers receiving the intervention and numbers followed up are also shown.

Overall study recruitment

Both study sites agreed to take part with the necessary organisation approvals and ethics approvals were secured. Recruitment opened at the England site on 2 December 2019 and closed on 28 February 2020, as the required sample size of 90 participants had been met. The Scotland site opened to recruitment on 15 January 2020 and was halted due to COVID-19 restrictions on 13 March 2020, with 42 individuals recruited to the study. Following discussions with the site governor and the head of research at the Scottish Prison Service, HMP Scotland was closed to any further recruitment on 12 June 2020. Differences in start date between study sites were due to difficulties caused by the model non-commercial agreement (mNCA) requested by the Health Research Authority (see [Ethics approval](#)).

[Table 7](#) provides an overview of the screening and recruitment data at the two sites.

Recruitment methods varied between the two sites as the prison profiles, regimes and access were different. As such, the screening and recruitment data for the two sites are considered in more detail separately.

England study site recruitment

[Table 8](#) summarises the screening and recruitment data for England by week number. Unlike the Scotland site, where all individuals on remand were given an information sheet about the trial, not all of the individuals entering HMP England received an information sheet.

The number of prisoners on remand at the start of recruitment was not provided by the England site. Reasons why individuals were excluded in England are summarised in [Table 9](#).

Scotland study site recruitment

[Table 10](#) contains information on the number of individuals entering HMP Scotland on remand by week of recruitment. At the start of recruitment, there were 162 prisoners on remand, and an additional

TABLE 7 Summary of screening and recruitment by site

	Entering on remand (N = 656)	Given information sheet (N = 705) ^a	Number approached (N = 182)	Number randomised (N = 132)
England	497 (75.8)	384 (54.5)	134 (73.6)	90 (68.2)
Scotland	159 (24.2)	321 (45.5)	48 (26.4)	42 (31.8)

^a The Scotland figure includes 162 prisoners who were on remand when recruitment began.

TABLE 8 Summary of screening and recruitment at the England site by week

Week number	Date at start of week	No. (%) entering on remand (N = 497)	No. (%) given info sheet (N = 384)	No. (%) approached (N = 134)	No. (%) randomised (N = 90)
1	2 December 2019	44 (8.9)	31 (8.1)	3 (2.2)	3 (3.3)
2	9 December 2019	35 (7)	23 (6)	3 (2.2)	1 (1.1)
3	16 December 2019	28 (5.6)	19 (4.9)	10 (7.5)	2 (2.2)
4	6 January 2020	47 (9.5)	43 (11.2)	19 (14.2)	6 (6.7)
5	13 January 2020	45 (9.1)	38 (9.9)	15 (11.2)	9 (10)
6	20 January 2020	68 (13.7)	59 (15.4)	15 (11.2)	8 (8.9)
7	27 January 2020	52 (10.5)	36 (9.4)	13 (9.7)	9 (10)
8	3 February 2020	44 (8.9)	30 (7.8)	16 (11.9)	7 (7.8)
9	10 February 2020	62 (12.5)	56 (14.6)	19 (14.2)	15 (16.7)
10	17 February 2020	51 (10.3)	30 (7.8)	2 (1.5)	13 (14.4)
11	24 February 2020	21 (4.2)	19 (4.9)	19 (14.2)	17 (18.9)

No., number; info, information.

TABLE 9 Reasons for exclusions at the England site

	Number (%) of individuals (N = 88)
Self-identified non-drinker	10 (11.4)
Not in pad (cell)	9 (10.2)
Refused	4 (4.5)
Not feeling well	12 (13.6)
Isolation/segregation	2 (2.3)
Risk to others	6 (6.8)
High risk	8 (9.1)
Operational	5 (5.7)
Alcohol-dependant	11 (12.5)
Not able to understand the documents	10 (11.4)
Health care	10 (11.4)
Already participated	1 (1.1)

159 individuals entered the prison on remand during the data-collection period. During recruitment, 195 individuals were excluded, and 86 were not approached due to lack of resources and time.

[Table 11](#) provides information on the number of screened individuals with 0, 1, 2, 3 or 4 reasons for being excluded from the study.

Of those who had at least one reason for being excluded at the Scotland site, there was a total of 210 exclusion reasons, and these have been summarised in [Table 12](#). Of note is the largest exclusion reason,

TABLE 10 Summary of screening and recruitment at the Scotland site by week

Week number	Date at start of week	No. (%) entering on remand (N = 159)	No. (%) randomised (N = 42)
1	13 January 2020	7 (4.4)	4 (9.5)
2	20 January 2020	23 (14.5)	6 (14.3)
3	27 January 2020	16 (10.1)	5 (11.9)
4	3 February 2020	22 (13.8)	4 (9.5)
5	10 February 2020	13 (8.2)	2 (4.8)
6	17 February 2020	17 (10.7)	5 (11.9)
7	24 February 2020	15 (9.4)	5 (11.9)
8	2 March 2020	25 (15.7)	5 (11.9)
9	9 March 2020	21 (13.2)	6 (14.3)

No., number.

TABLE 11 Information on the number of screened individuals with 0, 1, 2, 3 or 4 reasons for being excluded from the study at the Scotland site

Number of reasons	Number (%) of individuals excluded (N = 195)
0	9 (4.6)
1	165 (84.6)
2	19 (9.7)
3	1 (0.5)
4	1 (0.5)

TABLE 12 Information on the number of exclusion reasons at the Scotland site

	Number (%) of exclusion reasons (N = 210)
More than 3 months on remand	74 (35.2)
Operational reasons (hall)	44 (21)
Transferred	5 (2.4)
Liberated	49 (23.3)
Unable to give informed consent	1 (0.5)
Risk to self	7 (3.3)
Risk to others	4 (1.9)
Segregative rule	9 (4.3)
Not able to understand the documents	3 (1.4)
Sentenced	5 (2.4)
Self-identified non-drinker	3 (1.4)
Awaiting extradition	3 (1.4)
Audit < 8	2 (1)
Already active participant	1 (0.5)

which was being on remand for more than 3 months. Being liberated ($n = 49$) and operational reasons ($n = 44$) were the second and third most prevalent reasons for exclusion respectively.

We had not created a standard proforma of consistent categories for exclusion, which resulted in different terminology being used for the exclusion criteria at the two study sites. We also noted that some criteria were much more frequent at one study site compared to the other (there was not much consistency between sites). Following several discussions by the PMG we agreed, for the purposes of the pilot trial and to support our understanding of these differences, that for a future trial we would continue the site-specific reasons and as such the exclusions are reported separately.

Consented and randomised

There were 134 participants who gave informed consent (90 from the England site and 44 from Scotland). A total number of 132 participants were randomised. Two participants from the Scotland site who were consented did not proceed to randomisation. One participant was considered to have lacked capacity and the other scored less than 8 on the AUDIT. [Table 13](#) provides data on the numbers randomised to the intervention and control arms at each study site.

Randomisation recruitment rates

It was anticipated that recruitment would start on the 1 September 2019 and would last for 4–6 months, with recruitment being quicker in England compared to Scotland. We were predicting this based on our learning from the PRISM-A study.⁴⁰ Variations in the speed of recruitment can partially be explained by access opportunities. Due to the different regimes within the prisons, the amount of time on the halls was less in Scotland than in England. The regimes also impacted on the amount of time the researchers were able to be on site. For example, in England the researcher was on site 5 days a week. In Scotland the researcher was on site 3 days a week.

[Table 14](#) contains information on the number of participants randomised by month and by site, with [Figure 2](#) providing a graphical overview of the recruitment rate by site and as a total. The planned rate of recruitment starts from 2 December in [Figure 3](#) to correspond with when the England site was opened to recruitment.

Baseline characteristics

The demographic characteristics of the randomised participants can be found in [Tables 15](#) and [16](#). The average age of the participants was 34 years (median 31, SD 10), the majority of participants were

TABLE 13 Number (%) randomised and allocation

	Site					
	England		Scotland		Both sites	
	N	%	N	%	N	%
Intervention	46	51	22	52	68	52
Control	44	49	20	48	64	48
Total	90	100	42	100	132	100

TABLE 14 Number of randomised participants by month-year and site

Month year	England (N = 90)	Scotland (N = 42)	Total (N = 132)
December 2019	6 (6.7)	0 (0)	6 (4.5)
January 2020	32 (35.6)	15 (35.7)	47 (35.6)
February 2020	52 (57.8)	16 (38.1)	68 (51.5)
March 2020	0 (0)	11 (26.2)	11 (8.3)

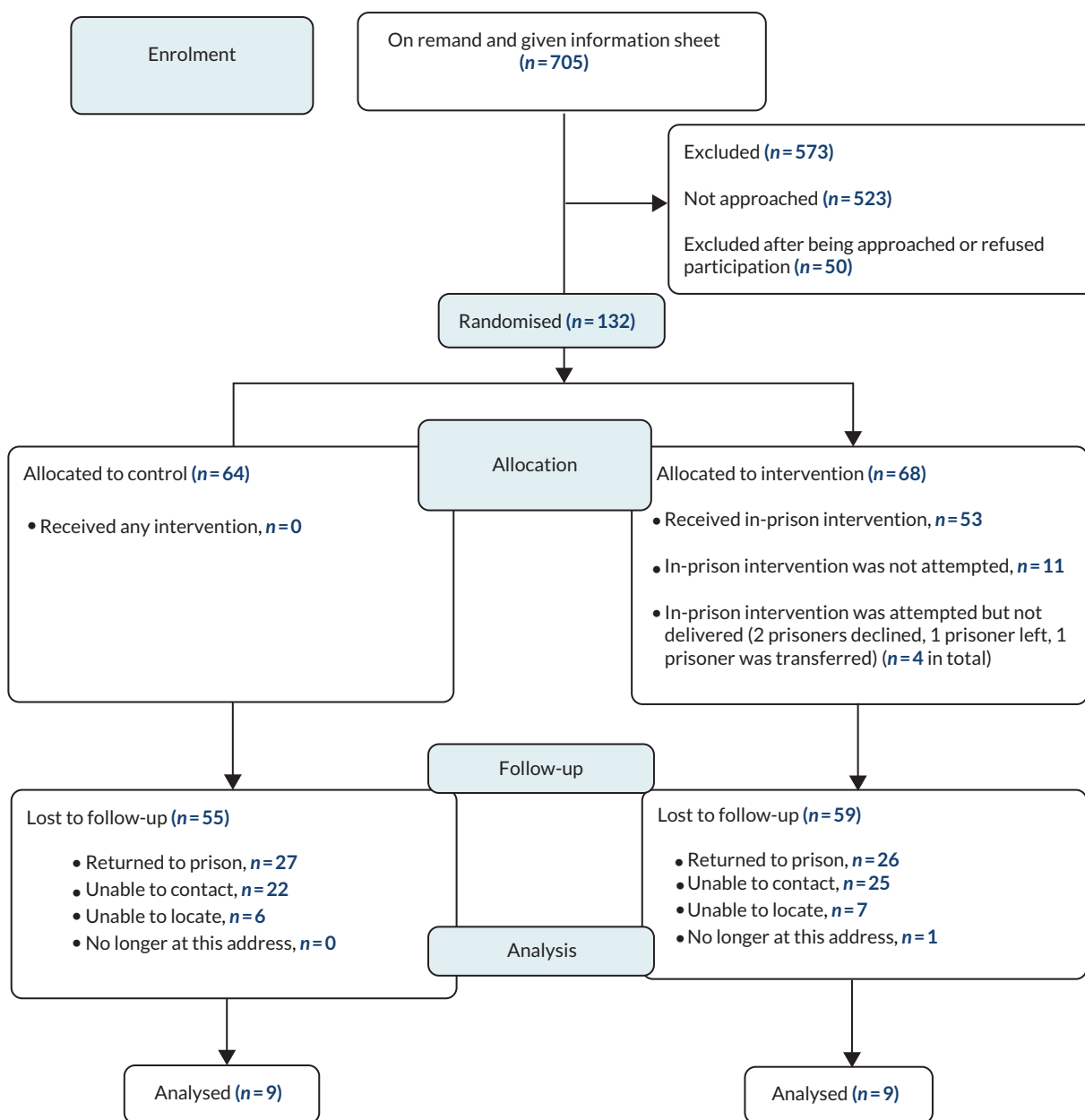


FIGURE 2 Consolidated Standards of Reporting Trials 2010 flow diagram.

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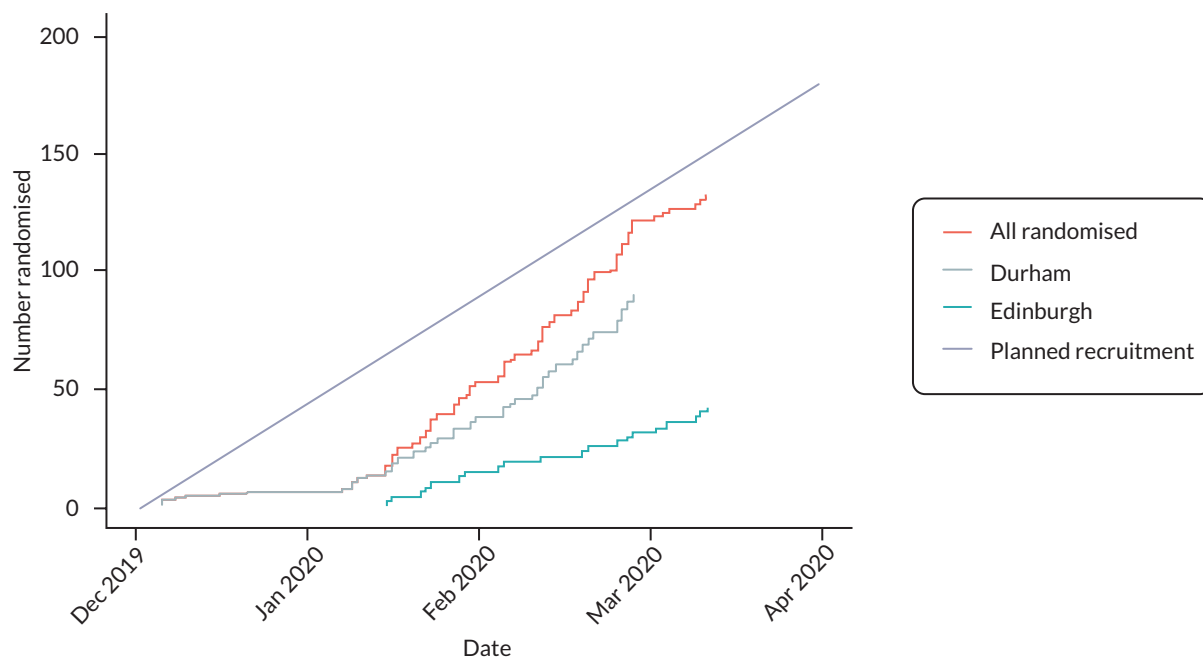


FIGURE 3 Recruitment graph of planned recruitment and overall recruitment by study site.

TABLE 15 Frequency table of the categorical baseline characteristics of the randomised participants, stratified by site and randomised group

		Randomisation allocation					
		Intervention		Control		Overall	
		N	%	N	%	N	%
Site							
England	Total number of prisoners	46	67.6	44	68.8	90	68.2
	Ethnicity						
	White Scottish	0	0.0	0	0.0	0	0.0
	White British	45	66.2	38	59.4	83	62.9
	White Irish	0	0.0	2	3.1	2	1.5
	White other	0	0.0	1	1.6	1	0.8
	Asian British	0	0.0	1	1.6	1	0.8
	Mixed race	1	1.5	1	1.6	2	1.5
	Other	0	0.0	1	1.6	1	0.8
	Relationship status						
	Single	31	45.6	33	51.6	64	48.5
	In a relationship (same sex or opposite)	14	20.6	8	12.5	22	16.7
	Married/civil partnership	1	1.5	2	3.1	3	2.3
	Divorced	0	0.0	1	1.6	1	0.8
	Highest educational achievement						
	University degree	1	1.5	0	0.0	1	0.8

TABLE 15 Frequency table of the categorical baseline characteristics of the randomised participants, stratified by site and randomised group (*continued*)

		Randomisation allocation					
		Intervention		Control		Overall	
		N	%	N	%	N	%
Scotland	College degree	0	0.0	0	0.0	0	0.0
	A-level/highers/advanced highers	1	1.5	0	0.0	1	0.8
	GCSEs/standard grades/National Certificate	14	20.6	18	28.1	32	24.2
	Recognised trade apprenticeship	10	14.7	8	12.5	18	13.6
	No formal qualifications	20	29.4	18	28.1	38	28.8
	Total number of prisoners	22	32.4	20	31.3	42	31.8
	Ethnicity						
	White Scottish	17	25.0	16	25.0	33	25.0
	White British	2	2.9	4	6.3	6	4.5
	White Irish	0	0.0	0	0.0	0	0.0
	White other	1	1.5	0	0.0	1	0.8
	Asian British	0	0.0	0	0.0	0	0.0
	Mixed race	0	0.0	0	0.0	0	0.0
	Other	2	2.9	0	0.0	2	1.5
	Relationship status						
	Single	13	19.1	10	15.6	23	17.4
	In a relationship (same sex or opposite)	8	11.8	8	12.5	16	12.1
	Married/civil partnership	1	1.5	2	3.1	3	2.3
	Divorced	0	0.0	0	0.0	0	0.0
	Highest educational achievement						
University degree	2	2.9	1	1.6	3	2.3	
College degree	2	2.9	2	3.1	4	3.0	
A-level/highers/advanced highers	0	0.0	0	0.0	0	0.0	
GCSEs/standard grades/National Certificate	10	14.7	9	14.1	19	14.4	
Recognised trade apprenticeship	3	4.4	4	6.3	7	5.3	
No formal qualifications	5	7.4	4	6.3	9	6.8	
Both sites	Total number of prisoners	68	100.0	64	100.0	132	100.0
	Ethnicity						
	White Scottish	17	25.0	16	25.0	33	25.0
	White British	47	69.1	42	65.6	89	67.4
	White Irish	0	0.0	2	3.1	2	1.5
	White other	1	1.5	1	1.6	2	1.5
	Asian British	0	0.0	1	1.6	1	0.8

continued

TABLE 15 Frequency table of the categorical baseline characteristics of the randomised participants, stratified by site and randomised group (*continued*)

	Randomisation allocation					
	Intervention		Control		Overall	
	N	%	N	%	N	%
Mixed race	1	1.5	1	1.6	2	1.5
Other	2	2.9	1	1.6	3	2.3
Relationship status						
Single	44	64.7	43	67.2	87	65.9
In a relationship (same sex or opposite)	22	32.4	16	25.0	38	28.8
Married/civil partnership	2	2.9	4	6.3	6	4.5
Divorced	0	0.0	1	1.6	1	0.8
Highest educational achievement						
University degree	3	4.4	1	1.6	4	3.0
College degree	2	2.9	2	3.1	4	3.0
A-level/highers/advanced highers	1	1.5	0	0.0	1	0.8
GCSEs/standard grades/National Certificate	24	35.3	27	42.2	51	38.6
Recognised trade apprenticeship	13	19.1	12	18.8	25	18.9
No formal qualifications	25	36.8	22	34.4	47	35.6

TABLE 16 Summary statistics of baseline characteristics of the randomised participants, stratified by site and randomised group

			N	Mean	SD	Median	Min	Max
Age	Site	Randomisation allocation						
		England	Intervention	46	31.0	7.7	30.0	19.0
		Control	44	34.4	12.6	30.0	19.0	86.0
		Overall	90	32.7	10.5	30.0	19.0	86.0
	Scotland	Randomisation allocation						
		Intervention	22	37.3	10.1	37.5	23.0	60.0
		Control	20	34.4	9.3	33.0	22.0	55.0
		Overall	42	35.9	9.7	34.5	22.0	60.0
	Both sites	Randomisation allocation						
		Intervention	68	33.1	8.9	31.0	19.0	62.0
		Control	64	34.4	11.6	31.0	19.0	86.0
		Overall	132	33.7	10.3	31.0	19.0	86.0
Alcohol consumption: number of drinks consumed in previous 28 days according to AUDIT-C	Site	Randomisation allocation						

TABLE 16 Summary statistics of baseline characteristics of the randomised participants, stratified by site and randomised group (*continued*)

			N	Mean	SD	Median	Min	Max	
AUDIT-C total score	England	Intervention	46	75.5	66.4	33.0	8.0	198.0	
		Control	44	102.1	72.3	110.0	10.5	198.0	
		Overall	90	88.5	70.2	80.0	8.0	198.0	
	Scotland	Randomisation allocation							
		Intervention	22	113.8	78.4	104.5	11.0	198.0	
		Control	20	98.0	78.4	81.0	5.5	198.0	
	Both sites	Randomisation allocation							
		Intervention	68	87.9	72.2	71.5	8.0	198.0	
		Control	64	100.8	73.6	110.0	5.5	198.0	
	Readiness to Change Ruler: how much do you really want to stop (or cut down) drinking at the present time?	England	Intervention	46	8.7	2.1	9.0	3.0	12.0
			Control	44	9.4	2.0	9.0	4.0	12.0
			Overall	90	9.0	2.1	9.0	3.0	12.0
		Scotland	Randomisation allocation						
			Intervention	22	9.7	2.3	9.5	5.0	12.0
			Control	20	9.3	2.4	9.0	5.0	12.0
Both sites		Randomisation allocation							
		Intervention	68	9.0	2.2	9.0	3.0	12.0	
		Control	64	9.4	2.2	9.0	4.0	12.0	
Readiness to Change Ruler: how much do you really want to stop (or cut down) drinking at the present time?		England	Intervention	46	7.2	3.5	9.0	1.0	10.0
			Control	44	7.3	3.2	9.0	1.0	10.0
			Overall	90	7.2	3.4	9.0	1.0	10.0
		Scotland	Randomisation allocation	22	7.8	2.8	9.0	1.0	10.0
			Intervention						
			Control	20	7.6	3.0	9.5	1.0	10.0
	Both sites	Randomisation allocation	42	7.7	2.9	9.0	1.0	10.0	
		Intervention	68	7.4	3.3	9.0	1.0	10.0	

continued

TABLE 16 Summary statistics of baseline characteristics of the randomised participants, stratified by site and randomised group (*continued*)

			N	Mean	SD	Median	Min	Max
		Control	64	7.4	3.1	9.0	1.0	10.0
		Overall	132	7.4	3.2	9.0	1.0	10.0
Readiness to Change Ruler: if you decided to stop or cut down drinking right now, how confident do you feel about succeeding with this?	Site	Randomisation allocation						
	England	Intervention	46	7.6	2.8	9.0	1.0	10.0
		Control	44	8.2	2.1	9.5	3.0	10.0
		Overall	90	7.9	2.5	9.0	1.0	10.0
	Scotland	Randomisation allocation						
		Intervention	22	7.0	2.7	7.5	1.0	10.0
		Control	20	7.3	3.0	9.0	1.0	10.0
	Overall	42	7.1	2.8	8.0	1.0	10.0	
	Both sites	Randomisation allocation						
		Intervention	68	7.4	2.8	8.0	1.0	10.0
		Control	64	7.9	2.4	9.0	1.0	10.0
	Overall	132	7.6	2.6	8.5	1.0	10.0	
Readiness to Change Ruler: how ready are you to stop (or cut down) drinking right now?	Site	Randomisation allocation						
	England	Intervention	46	8.2	2.6	10.0	1.0	10.0
		Control	44	8.3	2.6	10.0	1.0	10.0
		Overall	90	8.2	2.6	10.0	1.0	10.0
	Scotland	Randomisation allocation						
		Intervention	22	7.8	3.2	9.0	1.0	10.0
		Control	20	9.0	1.9	10.0	3.0	10.0
	Overall	42	8.4	2.7	10.0	1.0	10.0	
	Both sites	Randomisation allocation						
		Intervention	68	8.0	2.7	9.0	1.0	10.0
		Control	64	8.5	2.4	10.0	1.0	10.0
	Overall	132	8.3	2.6	10.0	1.0	10.0	
Readiness to Change Ruler: I have someone I can talk to about reducing my drinking	Site	Randomisation allocation						
	England	Intervention	46	7.3	3.6	10.0	1.0	10.0
		Control	44	7.9	3.0	10.0	1.0	10.0
		Overall	90	7.6	3.3	10.0	1.0	10.0

TABLE 16 Summary statistics of baseline characteristics of the randomised participants, stratified by site and randomised group (*continued*)

			N	Mean	SD	Median	Min	Max	
Readiness to Change Ruler: the people who are important to me support me in reducing my drinking	Scotland	Randomisation allocation							
		Intervention	22	5.6	3.7	5.0	1.0	10.0	
		Control	20	6.5	3.9	7.5	1.0	10.0	
		Overall	42	6.0	3.8	6.5	1.0	10.0	
	Both sites	Randomisation allocation							
		Intervention	68	6.7	3.7	9.0	1.0	10.0	
		Control	64	7.5	3.3	10.0	1.0	10.0	
		Overall	132	7.1	3.5	9.5	1.0	10.0	
	Readiness to Change Ruler: I have made a detailed plan regarding when, where and how to reduce my drinking	England	Randomisation allocation						
			Intervention	46	7.7	3.5	10.0	1.0	10.0
			Control	44	8.5	2.6	10.0	1.0	10.0
			Overall	90	8.1	3.1	10.0	1.0	10.0
Scotland		Randomisation allocation							
		Intervention	21	6.7	3.8	8.0	1.0	10.0	
		Control	18	6.2	3.8	6.5	1.0	10.0	
		Overall	39	6.5	3.8	8.0	1.0	10.0	
Both sites		Randomisation allocation							
		Intervention	67	7.4	3.6	10.0	1.0	10.0	
		Control	62	7.8	3.2	10.0	1.0	10.0	
		Overall	129	7.6	3.4	10.0	1.0	10.0	
Readiness to Change Ruler: I have made a detailed plan regarding when, where and how to reduce my drinking	England	Randomisation allocation							
		Intervention	46	3.2	3.2	1.0	1.0	10.0	
		Control	44	5.0	4.1	3.0	1.0	10.0	
		Overall	90	4.0	3.8	1.5	1.0	10.0	
	Scotland	Randomisation allocation							
		Intervention	22	3.7	3.5	1.5	1.0	10.0	
		Control	20	3.8	2.8	3.0	1.0	10.0	
		Overall	42	3.7	3.1	2.0	1.0	10.0	
	Both sites	Randomisation allocation							
		Intervention	68	3.3	3.3	1.0	1.0	10.0	
		Control	64	4.6	3.8	3.0	1.0	10.0	
		Overall	132	3.9	3.6	2.0	1.0	10.0	

continued

TABLE 16 Summary statistics of baseline characteristics of the randomised participants, stratified by site and randomised group (*continued*)

			N	Mean	SD	Median	Min	Max	
EQ-5D: mobility	Site	Randomisation allocation							
	England	Intervention	46	1.4	0.8	1.0	1.0	4.0	
		Control	44	1.1	0.5	1.0	1.0	3.0	
		Overall	90	1.3	0.7	1.0	1.0	4.0	
	Scotland	Randomisation allocation							
		Intervention	22	1.1	0.4	1.0	1.0	2.0	
		Control	20	1.8	1.0	1.0	1.0	4.0	
	Overall	Randomisation allocation							
		Intervention	68	1.3	0.7	1.0	1.0	4.0	
		Control	64	1.3	0.7	1.0	1.0	4.0	
	EQ-5D: self-care	Site	Randomisation allocation						
		England	Intervention	46	1.2	0.6	1.0	1.0	4.0
Control			44	1.2	0.5	1.0	1.0	3.0	
Overall			90	1.2	0.6	1.0	1.0	4.0	
Scotland		Randomisation allocation							
		Intervention	22	1.4	1.0	1.0	1.0	4.0	
		Control	20	1.2	0.5	1.0	1.0	3.0	
Overall		Randomisation allocation							
		Intervention	68	1.2	0.8	1.0	1.0	4.0	
		Control	64	1.2	0.5	1.0	1.0	3.0	
EQ-5D: usual activities (e.g. work, study, housework, family or leisure activities)		Site	Randomisation allocation						
		England	Intervention	46	1.4	0.7	1.0	1.0	3.0
	Control		44	1.3	0.8	1.0	1.0	4.0	
	Overall		90	1.4	0.7	1.0	1.0	4.0	
	Scotland	Randomisation allocation							
		Intervention	22	2.1	1.4	1.5	1.0	5.0	
		Control	20	2.1	1.3	2.0	1.0	5.0	
	Overall	Randomisation allocation							
		Intervention	68	1.6	1.0	1.0	1.0	5.0	

TABLE 16 Summary statistics of baseline characteristics of the randomised participants, stratified by site and randomised group (*continued*)

			N	Mean	SD	Median	Min	Max
EQ-5D: pain/discomfort	Site	Control	64	1.6	1.0	1.0	1.0	5.0
		Overall	132	1.6	1.0	1.0	1.0	5.0
	England	Intervention	46	2.3	1.3	2.0	1.0	5.0
		Control	44	1.8	1.1	1.0	1.0	5.0
	Scotland	Intervention	22	1.9	1.3	1.0	1.0	5.0
		Control	20	2.0	1.3	1.0	1.0	4.0
	Both sites	Intervention	68	2.1	1.3	2.0	1.0	5.0
		Control	64	1.9	1.2	1.0	1.0	5.0
		Overall	132	2.0	1.2	1.0	1.0	5.0
		Overall	132	2.0	1.2	1.0	1.0	5.0
EQ-5D: anxiety/depression	Site	Control	64	1.9	1.2	1.0	1.0	5.0
		Overall	132	2.0	1.2	1.0	1.0	5.0
	England	Intervention	46	2.9	1.2	3.0	1.0	5.0
		Control	44	2.5	1.2	2.5	1.0	5.0
	Scotland	Intervention	22	3.1	1.2	3.0	1.0	5.0
		Control	20	3.6	1.4	4.0	1.0	5.0
	Both sites	Intervention	68	3.0	1.2	3.0	1.0	5.0
		Control	64	2.9	1.3	3.0	1.0	5.0
		Overall	132	2.9	1.3	3.0	1.0	5.0
		Overall	132	2.9	1.3	3.0	1.0	5.0
EQ-5D: visual analogue scale	Site	Control	64	1.9	1.2	1.0	1.0	5.0
		Overall	132	2.0	1.2	1.0	1.0	5.0
	England	Intervention	46	49.7	23.0	50.0	0.0	100.0
		Control	44	61.0	22.0	60.0	10.0	100.0
	Scotland	Intervention	22	61.4	21.1	62.5	5.0	90.0
		Control	20	61.1	21.3	62.5	20.0	100.0
	Both sites	Intervention	68	55.5	22.0	55.0	0.0	100.0
		Control	64	61.0	22.0	60.0	10.0	100.0

continued

TABLE 16 Summary statistics of baseline characteristics of the randomised participants, stratified by site and randomised group (*continued*)

			N	Mean	SD	Median	Min	Max
Warwick-Edinburgh Mental Well-being Scale: total score	Both sites	Randomisation allocation						
		Intervention	68	53.5	22.9	50.0	0.0	100.0
		Control	64	61.0	21.6	60.0	10.0	100.0
		Overall	132	57.1	22.5	60.0	0.0	100.0
	Site	Randomisation allocation						
	England	Intervention	46	34.4	13.7	32.0	14.0	66.0
		Control	44	40.2	10.6	39.0	19.0	63.0
		Overall	90	37.2	12.6	36.0	14.0	66.0
	Scotland	Randomisation allocation						
		Intervention	21	41.6	13.1	41.0	24.0	70.0
Drinking Refusal Self-efficacy Questionnaire: total score	Both sites	Randomisation allocation						
		Intervention	67	36.7	13.8	35.6	14.0	70.0
		Control	64	40.5	10.5	40.0	19.0	63.0
		Overall	131	38.5	12.4	38.0	14.0	70.0
	Site	Randomisation allocation						
	England	Intervention	45	72.8	24.8	73.5	22.2	114.0
		Control	44	70.8	22.2	69.6	33.8	109.8
		Overall	89	71.8	23.4	71.0	22.2	114.0
	Scotland	Randomisation allocation						
		Intervention	21	58.5	20.1	55.8	27.1	99.0
Negative Alcohol Expectancy Questionnaire: same-day expectancy score	Both sites	Randomisation allocation						
		Intervention	66	68.2	24.2	67.6	22.2	114.0
		Control	64	68.0	23.0	68.6	19.0	109.8
		Overall	130	68.1	23.5	68.1	19.0	114.0
	Site	Randomisation allocation						
	England	Intervention	45	52.3	16.6	54.2	21.0	87.3
		Control	44	55.3	18.4	54.4	23.0	105.0
		Overall	89	53.7	17.5	54.2	21.0	105.0
	Scotland	Randomisation allocation						
		Intervention	21	61.2	12.2	62.0	28.7	86.1

TABLE 16 Summary statistics of baseline characteristics of the randomised participants, stratified by site and randomised group (*continued*)

			N	Mean	SD	Median	Min	Max
Control			19	64.8	9.7	67.2	46.2	82.0
Overall			40	62.9	11.1	64.6	28.7	86.1
Negative Alcohol Expectancy Questionnaire: next-day expectancy score	Both sites	Randomisation allocation						
		Intervention	66	55.1	15.8	56.0	21.0	87.3
	Control		63	58.1	16.8	58.0	23.0	105.0
		Overall	129	56.6	16.3	56.0	21.0	105.0
	England	Randomisation allocation						
		Intervention	45	41.4	16.2	39.0	18.0	71.0
	Control		44	47.9	20.1	43.9	18.0	90.0
		Overall	89	44.6	18.5	41.3	18.0	90.0
	Scotland	Randomisation allocation						
		Intervention	21	54.8	16.9	54.0	20.0	87.0
	Control		19	60.7	12.7	62.0	38.0	88.0
		Overall	40	57.6	15.2	58.0	20.0	88.0
Both sites	Randomisation allocation							
	Intervention	66	45.7	17.5	45.6	18.0	87.0	
Control		63	51.8	19.1	51.0	18.0	90.0	
	Overall	129	48.6	18.5	49.0	18.0	90.0	

single [87 (66%)] and considered themselves to be White British or White Scottish. Regarding highest educational achievement, 36% had no formal qualifications, 19% had a recognised trade apprenticeship, 39% had GCSEs or equivalent, 1% had A-levels or equivalent, while 6% were educated to university or college degree level. There was substantial variability in terms of the number of drinks consumed per month at baseline: the SD was 73 (range 5–198 drinks per month). The median number of drinks consumed per month was 80 (mean 94). The mean total score of the AUDIT-C was 9.2 (SD 2.2, range 3 to 12). A total score of < 5 on the AUDIT-C is regarded as low risk, while a score of 5 or more indicates hazardous drinking. Overall, 98% of participants were in the 'at-risk' category for drinking based on the AUDIT-C (≥ 5) at baseline, including 23% who recorded the maximum value of 12 on the AUDIT-C (indicating extremely high levels of drinking). Only two participants were considered to be at low risk (2%). These two participants had an AUDIT-C of 9 and were therefore eligible for the study.

On average, high scores were recorded on the readiness to change ruler at baseline except for the item 'I have made a detailed plan regarding when, where and how to reduce my drinking', indicating that most participants had not made a detailed plan regarding when, where and how to reduce their drinking. The EQ-5D item scores suggested that participants had moderate problems with anxiety/depression on average; all other EQ-5D item scores were low on average. The WEMWBS had a mean of 38.5 (SD 12.4), indicating probable clinical depression in most participants.

Figure 4 shows side-by-side boxplots of the ages of participants in each site.

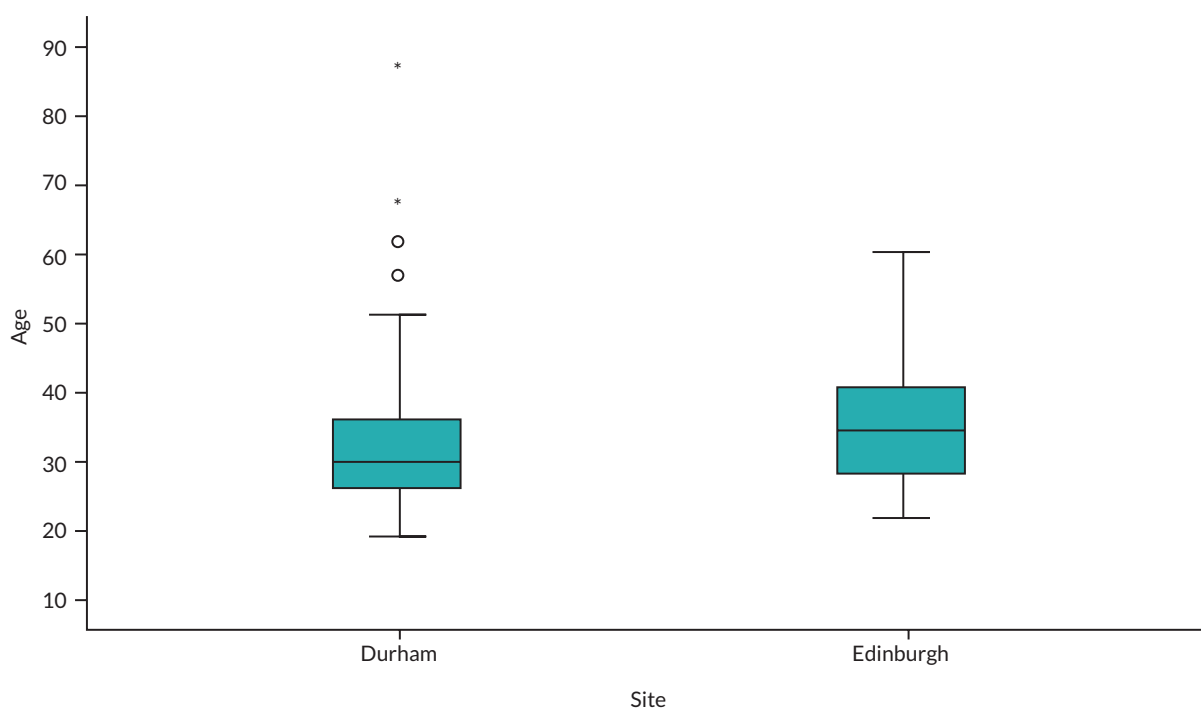


FIGURE 4 Ages of participants at HMP England and HMP Scotland.

In-prison and post-liberation intervention delivery

[Table 17](#) contains information on whether those randomised to the intervention arm received the in-prison and post-liberation intervention sessions. Out of 68 participants randomised to the intervention condition, 53 received the in-prison intervention session (77.9%, 95% CI 66.2% to 87.1%). This percentage is greater than the 70% threshold specified as one of the criteria necessary for progression to a future larger randomised controlled trial (Objective 4c).

Reasons for why the in-prison intervention was not delivered are shown in [Table 18](#).

Only one 'day-3 post-prison intervention' session was attempted and was delivered (duration 30 minutes) at the Scotland site. No 'day-7 post-prison intervention' sessions were attempted. Only one 'day-21 post-prison intervention' session was attempted and was delivered to one participant (duration 30 minutes) at the England site.

The main reasons why participants did not receive the post-prison interventions were (1) they could not be contacted and (2) CGL staff were unable to contact participants due to the impact of the COVID-19 pandemic.

Data for the duration (minutes) of the intervention sessions across both study sites are presented for all but one of the 53 interventions delivered in [Table 19](#). There was a marked difference in the median time taken to deliver the intervention between the two sites: the intervention duration was half an hour in England compared to 1 hour in Scotland. The shortest time for an intervention was only 5 minutes in England, whereas in Scotland it was 30 minutes. The maximum time for delivery was more aligned across sites, being 70 minutes and 80 minutes in England and Scotland, respectively.

TABLE 17 Number (%) of participants randomised to the intervention arm who received the in-prison and post-prison intervention delivery

	England (N = 46)	Scotland (N = 22)	Total (N = 68)
In prison			
Yes	40 (87)	13 (59.1)	53 (77.9)
No	6 (13)	9 (40.9)	15 (22.1)
First (3 days) post prison			
Yes	0 (0)	1 (4.5)	1 (1.5)
No	40 (87)	12 (54.5)	52 (76.5)
N/A ^a	6 (13)	9 (40.9)	15 (22.1)
Second (7 days) post prison			
Yes	0 (0)	0 (0)	0 (0)
No	40 (87)	13 (59.1)	53 (77.9)
N/A ^a	6 (13)	9 (40.9)	15 (22.1)
Third (21 days) post prison			
Yes	1 (2.2)	0 (0)	1 (1.5)
No	39 (84.8)	13 (59.1)	52 (76.5)
N/A ^a	6 (13)	9 (40.9)	15 (22.1)

a Not applicable (N/A) as the in-prison session was not received.

TABLE 18 Reasons why in-prison intervention was not delivered

	Site		
	England	Scotland	Both sites
Session attempted but participant declined	1	1	2
Prisoner was sentenced	0	2	2
Prisoner was released	1	3	4
Prisoner was transferred	1	0	1
Session not attempted because the prisoner had a crisis	0	1	1
Session not attempted because prisoner had COVID-19	0	2	2
Session not attempted for unknown reason	3	0	3
Total	6	9	15

TABLE 19 Intervention duration times (minutes)

	N	Miss	Mean	SD	Median	Min	Max
Duration of session (minutes)	40	0	31.8	11.2	30.0	5.0	70.0
Site							
England							
Scotland	12	1	53.8	14.9	60.0	30.0	80.0
Both sites	52	1	36.8	15.2	35.0	5.0	80.0

Follow-up and withdrawal

There were no formal withdrawals from the study. However, only 3 out of 132 randomised participants recorded any follow-up data at 6 months (2.3%, 95% CI 0.5% to 6.5%), and only 18 out of 132 randomised participants recorded any follow-up data at 12 months (13.6%, 95% CI 8.3% to 20.7%) (Table 20). The low numbers of follow-ups were due to the impact of COVID-19.

Tables 21 and 22 show the numbers of follow-ups at 6 and 12 months respectively within each site and randomised allocation group.

Out of the 44 participants receiving the intervention, only 13% were followed up to 12 months. Note that this percentage is much lower than our progression criteria (Objective 4d) specifying that at least 60% of those who received the intervention should be followed up at 12 months across trial arms and study sites. This progression criterion was set prior to the COVID-19 pandemic.

TABLE 20 Overall numbers (%) followed up by site and overall

	England (N = 90)	Scotland (N = 42)	Total (N = 132)
6-month follow-up			
Yes	0 (0)	3 (7.1)	3 (2.3)
No	90 (100)	39 (92.9)	129 (97.7)
12-month follow-up			
Yes	4 (4.4)	14 (33.3)	18 (13.6)
No	86 (95.6)	28 (66.7)	114 (86.4)

TABLE 21 Number (%) followed up at 6 months split by site and randomised allocation group

		Randomisation allocation					
		Intervention		Control		Overall	
		N	%	N	%	N	%
Site	Recording data at TP1						
England	No	46	67.6	44	68.8	90	68.2
	Yes	0	0.0	0	0.0	0	0.0
	Total	46	67.6	44	68.8	90	68.2
Scotland	Recording data at TP1						
	No	20	29.4	19	29.7	39	29.5
	Yes	2	2.9	1	1.6	3	2.3
	Total	22	32.4	20	31.3	42	31.8
Both sites	Recording data at TP1						
	No	66	97.1	63	98.4	129	97.7
	Yes	2	2.9	1	1.6	3	2.3
	Total	68	100.0	64	100.0	132	100.0

TABLE 22 Number (%) followed up at 12 months by site and overall

		Randomisation allocation					
		Intervention		Control		Overall	
		N	%	N	%	N	%
Site	Recording data at TP2						
England	No	44	64.7	42	65.6	86	65.2
	Yes	2	2.9	2	3.1	4	3.0
	Total	46	67.6	44	68.8	90	68.2
Scotland	Recording data at TP2						
	No	15	22.1	13	20.3	28	21.2
	Yes	7	10.3	7	10.9	14	10.6
	Total	22	32.4	20	31.3	42	31.8
Both sites	Recording data at TP2						
	No	59	86.8	55	85.9	114	86.4
	Yes	9	13.2	9	14.1	18	13.6
	Total	68	100.0	64	100.0	132	100.0

The greatest increase in numbers of those followed up at 12 months was seen at the Scotland site, where we were able to have some access to the prison site and identify where participants were located; however, this was minimal. In England where there was no access to the estate this meant at the time of intervention delivery and follow-ups we did not know who had been liberated and who had not liberated. However, this access remained very limited. Reasons for being unable to follow up at 12 months are shown in [Table 23](#). Out of all 132 randomised participants, 53 (40%) were not liberated or released at 12 months follow-up, 47 (36%) were uncontactable and 14 (11%) could not be located even though they had been released.

Data completeness

[Table 24](#) contains information on data completeness at baseline and follow-up for all 132 randomised participants. Complete data collection refers to when a participant has recorded data for all relevant questionnaires and outcomes. Incomplete data collection refers to when the participants have recorded data on the primary outcome, but have not recorded data for one or more secondary outcomes.

The information we have collected on data completeness helps us to address Objective 1c ('How well do participants complete the questionnaires necessary for a future definitive RCT?'). The majority of participants did not complete questionnaires, which was at least partly due to the difficulties in following up participants because of the COVID-19 pandemic and lockdown restrictions. However, considering the small numbers of participants who did complete the questionnaires, nearly half of them (9/21) could not (or would not) complete all the questionnaires or record data in response to every question.

TABLE 23 Table showing reasons for being unable to follow up at 12 months split by site and randomised allocation group

	Liberation/release status	Randomisation allocation					
		Intervention		Control		Overall	
		N	%	N	%	N	%
Site	Not liberated/released	24	54.5	26	61.9	50	58.1
England	Liberated and in community but unable to contact	16	36.4	11	26.2	27	31.4
	Liberated and in community but unable to locate	4	9.1	5	11.9	9	10.5
	Total	44	100	42	100	86	100
Scotland	Not liberated/released	2	13.3	1	7.7	3	10.7
	Liberated and in community but unable to contact	9	60.0	11	84.6	20	71.4
	Liberated and in community but unable to locate	4	26.7	1	7.7	5	17.9
	Total	15	100	13	100	28	100
Both sites	Not liberated/released	26	44.1	27	49.1	53	46.5
	Liberated and in community but unable to contact	25	42.4	22	40.0	47	41.2
	Liberated and in community but unable to locate	8	13.6	6	10.9	14	12.3
	Total	59	100	55	100	114	100

TABLE 24 Overall data completeness at baseline and follow-up

	Baseline	6 months	12 months
Complete data collection	127 (96.2)	1 (0.8)	11 (8.3)
Incomplete data collection	5 (3.8)	2 (1.5)	7 (5.3)
No data recorded	0	129 (97.7)	114 (86.4)

Outcomes

Primary outcome

The intended primary outcome measure for the pilot RCT was the TLFB-28. As outlined in [Chapter 3](#) this was amended to AUDIT-C. [Table 25](#) shows descriptive statistics for estimated alcohol consumption in the previous 28 days at each time point, restricted to those participants with any follow-up information. Please note that alcohol information recorded at baseline may be an underestimate because a different AUDIT questionnaire was used at this time point and the formats of the first few question options were different (e.g. '4–5 times a week' drinking was not distinguished from '6 or more times a week' drinking).

[Table 26](#) shows means and 95% confidence intervals for alcohol consumption (number of drinks consumed in the past 28 days) within each site and randomised group. Note that the amount of missing data is very substantial, so it is difficult to interpret these results.

TABLE 25 Descriptive statistics for primary outcome: alcohol consumption in previous 28 days according to the AUDIT-C, split by site and randomised group

			N	Mean	SD	Median	Min	Max	
Both sites	Time point	Randomisation allocation							
		Baseline							
		Intervention	9	95.2	83.8	80.0	10.5	198.0	
			Control	9	107.8	70.3	110.0	11.0	198.0
		Overall	18	101.5	75.3	104.5	10.5	198.0	
	6 months follow-up	Randomisation allocation							
		Intervention	2	10.8	0.4	10.8	10.5	11.0	
		Control	1	11.0	.	11.0	11.0	11.0	
		Overall	3	10.8	0.3	11.0	10.5	11.0	
	12 months follow-up	Randomisation allocation							
		Intervention	9	106.1	159.8	16.5	0.0	448.0	
		Control	9	41.4	58.2	8.0	0.0	154.0	
	Overall	18	73.8	121.3	14.5	0.0	448.0		
Site	Time point	Randomisation allocation							
England	Baseline	Intervention	2	60.3	70.4	60.3	10.5	110.0	
		Control	2	121.5	31.8	121.5	99.0	144.0	
		Overall	4	90.9	56.9	104.5	10.5	144.0	
	12 months follow-up	Randomisation allocation							
		Intervention	2	13.5	4.2	13.5	10.5	16.5	
		Control	2	117.0	52.3	117.0	80.0	154.0	
	Overall	4	65.3	67.0	48.3	10.5	154.0		
Scotland	Baseline	Randomisation allocation							
		Intervention	7	105.2	89.6	80.0	11.0	198.0	
		Control	7	103.9	79.6	110.0	11.0	198.0	
		Overall	14	104.5	81.4	95.0	11.0	198.0	
	6 months follow-up	Randomisation allocation							
		Intervention	2	10.8	0.4	10.8	10.5	11.0	
		Control	1	11.0	.	11.0	11.0	11.0	
		Overall	3	10.8	0.3	11.0	10.5	11.0	
	12 months follow-up	Randomisation allocation							
		Intervention	7	132.6	174.3	24.0	0.0	448.0	
		Control	7	19.8	40.2	4.5	0.0	110.0	
		Overall	14	76.2	134.9	12.0	0.0	448.0	

TABLE 26 Mean and 95% confidence intervals for alcohol consumption

Randomisation allocation	Site	Baseline	6 months	12 months
Intervention	England	60.3 (0 to 692.4)	-	13.5 (0 to 51.6)
	Scotland	105.2 (22.4 to 188.1)	10.8 (7.6 to 13.9)	132.6 (0 to 293.7)
	Overall	95.2 (30.8 to 159.7)	10.8 (7.6 to 13.9)	106.1 (0 to 228.9)
Control	England	121.5 (0 to 407.4)	-	117.0 (0 to 587.1)
	Scotland	103.9 (30.2 to 177.5)	-	19.8 (0 to 57.0)
	Overall	107.8 (53.7 to 161.8)	-	41.4 (0 to 86.2)

For the primary analysis, the mean difference with 95% confidence interval in alcohol consumption at 12 months post randomisation (TP2) between the two randomisation groups was calculated (Table 27), both overall and stratified by site. Linear regression was used, adjusting for baseline alcohol consumption (TP0). Note that the linear regression model for England was based on only four data points and the 95% confidence intervals are extremely wide. Due to the substantial amount of missing data, extreme caution is required when interpreting these results.

Secondary outcomes

Summary statistics of the secondary outcomes are presented in Table 28 for both sites combined. Please see supplementary material for the same tables stratified by site (Tables S1.1 and S1.2 in Report Supplementary Material 1). Although sample sizes are small, the standard deviations reported at 12 months follow-up could be used to inform sample size calculations for future studies in conjunction with the standard deviations reported in the baseline characteristics (see Table 16). This addresses our Objective 1b ('Can we obtain reasonable estimates of the parameters necessary to inform the design and sample size calculation for a future definitive multicentre RCT?'). This also gives us some insight into Objective 2c ('To what extent was the intervention changing process variables consistent with the underpinning theory').

The supplementary material also contains detailed tables of individual questionnaire item scores, overall and split by site (Tables S3.1 to S3.7 in Report Supplementary Material 1).

The secondary outcome results, change from baseline, show the average scores of participants in the intervention and control group at baseline, 6 months and 12 months. Due to missing data and small sample sizes, we are not able to interpret change from baseline to 6 months, nor group differences in change for 6 to 12 months. For change from baseline to 12 months, we can look at patterns of change in the data in mechanisms of change, albeit with caution. Readiness to change does not show increasing or at least maintained values; rather it seems to slightly decrease across items. One exception is the item 'I have made a detailed plan regarding when, where and how to reduce my drinking', which increases from baseline to 12 months in both groups (at baseline intervention: $M = 4.17$, control: 3.0; at 12 months intervention: $M = 6.1$, control 5.1).

Logic model results

An exploratory analysis was performed to explore key behavioural markers (mediators) of change and how these may influence the effect of the delivered intervention. The six proposed mechanisms of change were: self-efficacy, outcome expectations, goals, plans, social support and high-risk situations.

TABLE 27 Mean and 95% confidence intervals for the difference in alcohol consumption between intervention/control groups

Site	Mean difference intervention – control (95% CI)
England	-117.5 (-919.1 to 684.0)
Scotland	112.6 (-42.4 to 267.6)
Overall	67.2 (-57.0 to 191.5)

TABLE 28 Secondary outcome results

			N	Miss	Mean	SD	Median	Min	Max		
AUDIT-C total score	Time point	Randomisation allocation									
		TP0	Intervention	9	0	8.6	3.0	9.0	5.0	12.0	
		Control	9	0	9.9	1.8	10.0	7.0	12.0		
		Overall	18	0	9.2	2.5	9.0	5.0	12.0		
	TP1	Randomisation allocation									
		Intervention	2	0	4.5	2.1	4.5	3.0	6.0		
		Control	1	0	6.0	.	6.0	6.0	6.0		
		Overall	3	0	5.0	1.7	6.0	3.0	6.0		
	TP2	Randomisation allocation									
		Intervention	9	0	6.9	4.0	6.0	0.0	12.0		
		Control	8	1	4.5	4.4	4.0	0.0	10.0		
		Overall	17	1	5.8	4.2	6.0	0.0	12.0		
	Readiness to Change Ruler: how much do you really want to reduce (or stop) drinking?	Time point	Randomisation allocation								
			TP0	Intervention	9	0	8.3	2.8	10.0	3.0	10.0
			Control	9	0	7.6	3.6	9.0	1.0	10.0	
		Overall	18	0	7.9	3.1	9.5	1.0	10.0		
TP1		Randomisation allocation									
		Intervention	2	0	4.5	6.4	4.5	0.0	9.0		
		Control	1	0	10.0	.	10.0	10.0	10.0		
		Overall	3	0	6.3	5.5	9.0	0.0	10.0		
TP2		Randomisation allocation									
		Intervention	8	1	8.1	3.1	9.5	1.0	10.0		
		Control	8	1	7.0	3.9	8.5	1.0	10.0		
		Overall	16	2	7.6	3.4	9.0	1.0	10.0		

continued

TABLE 28 Secondary outcome results (continued)

			N	Miss	Mean	SD	Median	Min	Max		
Readiness to Change Ruler: if you decided to stop or cut down drinking right now, how confident do you feel about succeeding with this?	Time point	Randomisation allocation									
		TP0	Intervention	9	0	8.4	1.5	9.0	6.0	10.0	
		Control	9	0	8.7	1.3	9.0	6.0	10.0		
		Overall	18	0	8.6	1.4	9.0	6.0	10.0		
	TP1	Randomisation allocation									
		Intervention	2	0	9.0	1.4	9.0	8.0	10.0		
		Control	1	0	7.0	.	7.0	7.0	7.0		
		Overall	3	0	8.3	1.5	8.0	7.0	10.0		
	TP2	Randomisation allocation									
		Intervention	8	1	6.8	3.6	8.0	1.0	10.0		
		Control	7	2	7.6	1.8	7.0	6.0	10.0		
		Overall	15	3	7.1	2.9	8.0	1.0	10.0		
	Readiness to Change Ruler: how ready are you to stop (or cut down) drinking right now?	Time point	Randomisation allocation								
			TP0	Intervention	9	0	9.4	1.0	10.0	7.0	10.0
			Control	9	0	9.7	0.5	10.0	9.0	10.0	
		Overall	18	0	9.6	0.8	10.0	7.0	10.0		
TP1		Randomisation allocation									
		Intervention	2	0	5.0	7.1	5.0	0.0	10.0		
		Control	1	0	10.0	.	10.0	10.0	10.0		
		Overall	3	0	6.7	5.8	10.0	0.0	10.0		
TP2		Randomisation allocation									
		Intervention	8	1	8.1	3.1	9.5	1.0	10.0		
		Control	7	2	6.3	3.2	6.0	1.0	10.0		
		Overall	15	3	7.3	3.2	8.0	1.0	10.0		
Readiness to Change Ruler: I have someone I can talk to about reducing my drinking		Time point	Randomisation allocation								
			TP0	Intervention	9	0	7.8	3.3	10.0	1.0	10.0
			Control	9	0	7.1	3.8	9.0	1.0	10.0	
		Overall	18	0	7.4	3.5	9.5	1.0	10.0		
	TP1	Randomisation allocation									
		Intervention	2	0	7.5	3.5	7.5	5.0	10.0		
		Control	1	0	0.0	.	0.0	0.0	0.0		
		Overall	3	0	5.0	5.0	5.0	0.0	10.0		
	TP2	Randomisation allocation									
		Intervention	8	1	6.8	3.7	7.5	1.0	10.0		

TABLE 28 Secondary outcome results (continued)

			N	Miss	Mean	SD	Median	Min	Max
Readiness to Change Ruler: the people who are important to me support me in reducing my drinking		Control	7	2	7.0	3.2	7.0	2.0	10.0
		Overall	15	3	6.9	3.3	7.0	1.0	10.0
	Time point	Randomisation allocation							
	TP0	Intervention	8	1	8.5	1.9	9.0	5.0	10.0
		Control	9	0	6.4	4.0	8.0	1.0	10.0
		Overall	17	1	7.4	3.2	8.0	1.0	10.0
	TP1	Randomisation allocation							
		Intervention	2	0	7.5	3.5	7.5	5.0	10.0
		Control	1	0	5.0	.	5.0	5.0	5.0
		Overall	3	0	6.7	2.9	5.0	5.0	10.0
	TP2	Randomisation allocation							
	Intervention	8	1	6.9	3.8	8.0	1.0	10.0	
	Control	7	2	6.9	3.1	6.0	3.0	10.0	
	Overall	15	3	6.9	3.4	8.0	1.0	10.0	
Readiness to Change Ruler: I have made a detailed plan regarding when, where and how to reduce my drinking	Time point	Randomisation allocation							
	TP0	Intervention	9	0	4.7	4.2	2.0	1.0	10.0
		Control	9	0	3.0	2.2	3.0	1.0	7.0
		Overall	18	0	3.8	3.4	2.5	1.0	10.0
	TP1	Randomisation allocation							
		Intervention	2	0	5.0	4.2	5.0	2.0	8.0
		Control	1	0	7.0	.	7.0	7.0	7.0
		Overall	3	0	5.7	3.2	7.0	2.0	8.0
		TP2	Randomisation allocation						
		Intervention	8	1	6.1	3.4	6.5	1.0	10.0
	Control	7	2	5.1	2.8	5.0	1.0	10.0	
	Overall	15	3	5.7	3.0	6.0	1.0	10.0	
EQ-5D: mobility	Time point	Randomisation allocation							
	TP0	Intervention	9	0	1.1	0.3	1.0	1.0	2.0
		Control	9	0	1.6	0.9	1.0	1.0	3.0
		Overall	18	0	1.3	0.7	1.0	1.0	3.0
	TP1	Randomisation allocation							
		Intervention	2	0	2.0	1.4	2.0	1.0	3.0
		Control	1	0	1.0	.	1.0	1.0	1.0
	Overall	3	0	1.7	1.2	1.0	1.0	3.0	

continued

TABLE 28 Secondary outcome results (continued)

			N	Miss	Mean	SD	Median	Min	Max	
EQ-5D: self-care	TP2	Randomisation allocation								
		Intervention	8	1	1.5	0.8	1.0	1.0	3.0	
		Control	9	0	1.2	0.4	1.0	1.0	2.0	
		Overall	17	1	1.4	0.6	1.0	1.0	3.0	
	Time point	Randomisation allocation								
		TP0	Intervention	9	0	1.0	0.0	1.0	1.0	1.0
		Control	9	0	1.2	0.7	1.0	1.0	3.0	
	Overall	18	0	1.1	0.5	1.0	1.0	3.0		
	TP1	Randomisation allocation								
		Intervention	2	0	1.0	0.0	1.0	1.0	1.0	
		Control	1	0	1.0	.	1.0	1.0	1.0	
		Overall	3	0	1.0	0.0	1.0	1.0	1.0	
TP2	Randomisation allocation									
	Intervention	8	1	1.1	0.4	1.0	1.0	2.0		
	Control	9	0	1.3	0.7	1.0	1.0	3.0		
	Overall	17	1	1.2	0.6	1.0	1.0	3.0		
EQ-5D: usual activities (e.g. work, study, housework, family or leisure activities)	Time point	Randomisation allocation								
		TP0	Intervention	9	0	1.9	1.4	1.0	1.0	5.0
		Control	9	0	1.8	1.3	1.0	1.0	5.0	
	Overall	18	0	1.8	1.3	1.0	1.0	5.0		
	TP1	Randomisation allocation								
		Intervention	2	0	2.0	1.4	2.0	1.0	3.0	
		Control	1	0	1.0	.	1.0	1.0	1.0	
		Overall	3	0	1.7	1.2	1.0	1.0	3.0	
	TP2	Randomisation allocation								
		Intervention	8	1	1.9	1.4	1.0	1.0	4.0	
		Control	9	0	1.3	0.7	1.0	1.0	3.0	
		Overall	17	1	1.6	1.1	1.0	1.0	4.0	
EQ-5D: pain/discomfort	Time point	Randomisation allocation								
		TP0	Intervention	9	0	2.1	1.4	1.0	1.0	4.0
		Control	9	0	1.9	1.4	1.0	1.0	4.0	
	Overall	18	0	2.0	1.3	1.0	1.0	4.0		
	TP1	Randomisation allocation								
		Intervention	2	0	2.5	2.1	2.5	1.0	4.0	
		Control	1	0	1.0	.	1.0	1.0	1.0	

TABLE 28 Secondary outcome results (continued)

			N	Miss	Mean	SD	Median	Min	Max
		Overall	3	0	2.0	1.7	1.0	1.0	4.0
	TP2	Randomisation allocation							
		Intervention	8	1	2.8	1.3	3.0	1.0	4.0
		Control	9	0	1.7	1.1	1.0	1.0	4.0
		Overall	17	1	2.2	1.3	2.0	1.0	4.0
EQ-5D: anxiety/ depression	Time point	Randomisation allocation							
	TP0	Intervention	9	0	2.7	1.1	3.0	1.0	4.0
		Control	9	0	3.1	1.8	4.0	1.0	5.0
		Overall	18	0	2.9	1.5	3.0	1.0	5.0
	TP1	Randomisation allocation							
		Intervention	2	0	1.0	0.0	1.0	1.0	1.0
		Control	1	0	2.0	.	2.0	2.0	2.0
		Overall	3	0	1.3	0.6	1.0	1.0	2.0
	TP2	Randomisation allocation							
		Intervention	8	1	2.9	1.7	3.0	1.0	5.0
		Control	9	0	2.8	1.3	2.0	1.0	5.0
		Overall	17	1	2.8	1.5	3.0	1.0	5.0
EQ-5D: visual analogue scale	Time point	Randomisation allocation							
	TP0	Intervention	9	0	60.0	24.7	65.0	25.0	90.0
		Control	9	0	71.2	15.9	70.0	50.0	90.0
		Overall	18	0	65.6	21.0	68.0	25.0	90.0
	TP1	Randomisation allocation							
		Intervention	2	0	86.0	15.6	86.0	75.0	97.0
		Control	1	0	95.0	.	95.0	95.0	95.0
		Overall	3	0	89.0	12.2	95.0	75.0	97.0
	TP2	Randomisation allocation							
		Intervention	9	0	57.2	29.9	70.0	0.0	95.0
		Control	9	0	57.2	18.9	60.0	20.0	80.0
		Overall	18	0	57.2	24.3	60.0	0.0	95.0
Warwick-Edinburgh Mental Well-being Scale: total score	Time point	Randomisation allocation							
	TP0	Intervention	9	0	46.0	12.3	42.0	28.0	70.0
		Control	9	0	42.2	9.8	41.0	29.0	56.0
		Overall	18	0	44.1	10.9	42.0	28.0	70.0

continued

TABLE 28 Secondary outcome results (continued)

			N	Miss	Mean	SD	Median	Min	Max	
Drinking Refusal Self-efficacy Questionnaire: total score	TP1	Randomisation allocation								
		Intervention	2	0	55.0	1.4	55.0	54.0	56.0	
		Control	1	0	52.8	.	52.8	52.8	52.8	
		Overall	3	0	54.3	1.6	54.0	52.8	56.0	
		TP2	Randomisation allocation							
		Intervention	8	1	44.3	13.1	46.5	23.0	61.0	
		Control	9	0	47.7	9.0	49.5	32.0	57.0	
		Overall	17	1	46.1	10.9	49.0	23.0	61.0	
		Time point	Randomisation allocation							
		TP0	Intervention	9	0	73.4	19.1	79.2	39.1	96.0
			Control	9	0	63.9	30.7	70.0	19.0	102.8
			Overall	18	0	68.6	25.3	76.5	19.0	102.8
Negative Alcohol Expectancy Questionnaire: same-day expectancy score	TP1	Randomisation allocation								
		Intervention	2	0	93.4	9.7	93.4	86.6	100.3	
		Control	1	0	86.1	.	86.1	86.1	86.1	
		Overall	3	0	91.0	8.1	86.6	86.1	100.3	
		TP2	Randomisation allocation							
		Intervention	8	1	64.0	33.2	63.1	19.0	109.0	
		Control	8	1	81.3	23.2	85.8	40.0	114.0	
		Overall	16	2	72.7	29.1	82.3	19.0	114.0	
		Time point	Randomisation allocation							
		TP0	Intervention	9	0	52.7	16.6	52.5	28.7	86.1
			Control	9	0	57.8	14.6	63.0	35.0	75.6
			Overall	18	0	55.2	15.4	53.3	28.7	86.1
		TP1	Randomisation allocation							
			Intervention	1	1	29.0	.	29.0	29.0	29.0
			Control	0	1
			Overall	1	2	29.0	.	29.0	29.0	29.0
		TP2	Randomisation allocation							
			intervention	8	1	53.9	19.7	51.0	31.0	79.3
		Control	8	1	50.3	16.0	48.7	28.0	77.0	
		Overall	16	2	52.1	17.4	48.7	28.0	79.3	
Negative Alcohol Expectancy Questionnaire: next-day expectancy score	Time point	Randomisation allocation								
	TP0	Intervention	9	0	47.1	18.7	47.0	20.0	82.0	
		Control	9	0	53.8	19.3	55.0	18.0	88.0	
		Overall	18	0	50.5	18.8	49.5	18.0	88.0	

TABLE 28 Secondary outcome results (continued)

		N	Miss	Mean	SD	Median	Min	Max
TP1	Randomisation allocation							
	Intervention	1	1	33.0	.	33.0	33.0	33.0
	Control	0	1
	Overall	1	2	33.0	.	33.0	33.0	33.0
TP2	Randomisation allocation							
	Intervention	8	1	50.1	22.6	45.5	23.0	87.2
	Control	8	1	48.4	14.5	45.0	30.0	71.0
	Overall	16	2	49.3	18.4	45.5	23.0	87.2

Summary statistics for the mechanisms of change at each time point and within each randomised group are presented in [Report Supplementary Material 1, Tables S2.1 to S2.6](#).

In exploratory analysis, Kendall's tau correlation coefficients were calculated to assess the correlation between each of the mechanisms of change and the primary outcome (AUDIT-C) at 12 months. Correlation coefficients and *p*-values are shown in [Table 29](#), along with valid sample size *N* that each correlation analysis is based on.

TABLE 29 Kendall's tau correlation coefficients of each of the mechanisms of change against the primary outcome

Mechanism of change	N	Kendall's tau correlation coefficient	P-value
Self-efficacy: Readiness to Change Ruler: 'How confident are you that you could reduce (or stop) drinking when you are released?'	15	-0.31	0.13
Self-efficacy: Drinking Refusal Self-efficacy Questionnaire Total	16	-0.43	0.023
Alcohol expectancy: Negative alcohol expectancy questionnaire: Same-day expectancy score	16	0.02	0.93
Alcohol expectancy: Negative alcohol expectancy questionnaire: Next-day expectancy score	16	0.11	0.55
Goals: Readiness to Change Ruler: 'How much do you want to reduce (or stop) drinking when you are released?'	16	0.27	0.18
Plans: Readiness to Change Ruler: 'How ready are you to reduce (or stop) drinking when you are released?'	15	0.17	0.41
Plans: Readiness to Change Ruler: 'I have made a detailed plan regarding when, where and how to reduce my drinking'	15	-0.19	0.34
Social support: 'The people who are important to me support me in reducing my drinking'	15	-0.26	0.22
Social support: 'I have someone I can talk to about reducing my drinking'	15	-0.22	0.28
High-risk situations: How much could I resist drinking when I am out for dinner?	16	-0.46	0.026

continued

TABLE 29 Kendall's tau correlation coefficients of each of the mechanisms of change against the primary outcome (*continued*)

Mechanism of change	N	Kendall's tau correlation coefficient	P-value
High-risk situations: How much could I resist drinking when someone offers me a drink?	16	-0.47	0.019
High-risk situations: How much could I resist drinking when my spouse or partner is drinking?	13	-0.54	0.021
High-risk situations: How much could I resist drinking when my friends are drinking?	16	-0.35	0.084
High-risk situations: How much could I resist drinking when I am at a pub or club?	16	-0.26	0.21
High-risk situations: How much could I resist drinking when I am angry?	16	-0.34	0.088
High-risk situations: How much could I resist drinking when I feel frustrated?	16	-0.33	0.10
High-risk situations: How much could I resist drinking when I am worried?	16	-0.24	0.24
High-risk situations: How much could I resist drinking when I feel upset?	16	-0.36	0.077
High-risk situations: How much could I resist drinking when I feel nervous?	15	-0.39	0.062
High-risk situations: How much could I resist drinking when I feel sad?	16	-0.52	0.009
High-risk situations: How much could I resist drinking when I am watching TV?	15	-0.47	0.031
High-risk situations: How much could I resist drinking when I am at lunch	16	-0.48	0.019
High-risk situations: How much could I resist drinking when I am on the way home from work?	15	-0.53	0.012
High-risk situations: How much could I resist drinking when I am listening to music or reading?	16	-0.46	0.026
High-risk situations: How much could I resist drinking when I am by myself?	16	-0.57	0.006
High-risk situations: How much could I resist drinking when I have just finished playing sport?	14	-0.45	0.046
High-risk situations: How much could I resist drinking when I first arrive home?	16	-0.57	0.006
High-risk situations: How much could I resist drinking when I feel like celebrating?	16	-0.13	0.52
High-risk situations: Drinking Refusal Self-efficacy Questionnaire Total	16	-0.43	0.023

Mindful of the very small sample sizes and that as such caution should be applied, provisional indications suggest that self-efficacy may be a determinant of alcohol consumption and further exploration of interventions targeting self-efficacy should be considered. We observed moderate correlations between markers of self-efficacy and high-risk situations and alcohol consumption, whereas the correlations between the other mechanisms for change and alcohol consumption were weak.

Serious adverse events

There were no serious adverse events reported during the pilot RCT.

Chapter 5 Process evaluation methods

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In this chapter, we present the methods for Phase II of the study which comprised the process evaluation (addressing Objectives 2 and 3). Elements of the process evaluation, where quantitative data were also collected, were integrated within the pilot trial (Objectives 2a and 2c) with cross-referencing to related results tables for these specific objectives provided.⁴⁴

Process evaluation aims and objectives

The new MRC framework for process evaluation guided the planning, design and proposed conduct of the process evaluation.³² The process evaluation set out to assess intervention fidelity, to consider the extent to which the behaviour change markers were appropriate to the underpinning intervention and to explore the feasibility and acceptability of the APPRAISE intervention and study measures to staff and for men on remand and following liberation. Objectives 2 and 3 set out the key questions that were to be answered.

Objective 2: to assess intervention fidelity

- 2a. What proportion of the interventions are delivered as per protocol?
- 2b. Is there any evidence of contamination between the two conditions and/or between those workers delivering the intervention?
- 2c. To what extent was the intervention changing process variables consistent with the underpinning theory?

Objective 3: to qualitatively explore the feasibility and acceptability of a self-efficacy-enhancing psychosocial alcohol intervention and study measures to staff and for men on remand and on liberation

- 3a. How acceptable are the trial and intervention procedures (including context and any barriers and facilitators) to the following key stakeholders: men on remand in prison and following liberation; prison staff (including healthcare staff); commissioners; policy-makers and third-sector partners?

Process evaluation methods and design

Table 30 summarises the different components of the process evaluation and how they relate to the objectives of the APPRAISE pilot study. It also identifies the main research question for each process evaluation component and the main sources of data that were employed to answer them.

Both quantitative and qualitative methods were employed to provide the data required to answer the process evaluation questions. The quantitative data providing evidence related to the implementation dose and quality as well as the mechanisms of change associated with the intervention were integrated in the pilot trial. The study participants and wider stakeholder semistructured interviews were conducted separately from the pilot trial. This was due to the impact on access and availability during the COVID-19 pandemic.

TABLE 30 Process evaluation process: data source and results

Process evaluation component	Objective	Main research questions	Data source	Results
Implementation (i) Dose: delivered	2a	What proportion of the interventions were offered and subsequently successfully delivered?	Quantitative data from participant study records, intervention booklets of successful (includes duration) and unsuccessful delivery (includes reasons for latter).	Chapter 4, In-prison and post-liberation intervention delivery, Tables 17–19.
Implementation (ii) Dose: quality (fidelity)	2b	To what extent was the intervention delivered as intended?	Three cases were audio-recorded (with participant consent) of intervention delivery in the 1st APPRAISE session. Data were coded to assess delivery of essential theoretical elements and the use of the Behaviour Change Counselling Index (BECCI) ⁷² to assess intervention fidelity.	Chapter 6, Intervention implementation results
Contamination	2b	Is there any evidence of contamination between: those who receive the intervention(s) and those who are in the control condition and/or between those workers delivering the intervention?	Qualitative interviews with men on remand from intervention and control arm. Qualitative interviews with intervention workers and non-interventionists at each study site.	Chapter 6, Contamination results
Mechanism of change	2c	To what extent was the intervention delivery consistent with the underpinning theory? Are the key behavioural markers appropriate for the mechanism of behaviour change?	Quantitative data from self-reported alcohol self-efficacy and alcohol expectancies.	Chapter 4, Outcomes Table 28; Supplementary Material 1 (Tables S3.1–S3.7) Chapter 4, Logic model results Table 29; Supplementary Material 1 (Tables S2.1–S2.6)
	3a	How acceptable is the trial intervention and procedures to men on remand and on liberation? What are the barriers and facilitators?	Qualitative interviews with men on remand in both conditions.	Chapter 6, Remand participants
	3a	How acceptable is the trial intervention and procedures to key stakeholders? What are the barriers and facilitators?	Qualitative interviews with prison staff (including health-care staff); commissioners; policy-makers and third-sector partners.	Chapter 6, CGL team participants and Stakeholder participants
Context	3a	What factors will enable the delivery and or impact of the intervention at scale?	Qualitative interviews with stakeholders including prison staff (including healthcare staff); commissioners; policy-makers and third-sector partners.	Chapter 6, Context, Remand participants, CGL team participants and Stakeholder participants

In particular, the qualitative interview data analysis sought to provide evidence to verify the feasibility and acceptability of the APPRAISE study intervention to better understand and explain the social processes associated with the intervention through the lens of normalisation process theory (NPT).⁶⁹⁻⁷¹

Intervention implementation

To confirm intervention implementation, data collection was planned in two ways: intervention booklets (dose delivered) and audio recordings (dose quality).

Dose delivered

The interventionists were asked to use intervention booklets to record contact attempts, sessions offered, sessions delivered, reasons for unsuccessful delivery, session dates, and duration of sessions to assess whether the intervention was delivered as per protocol (see [In-prison and post-liberation intervention delivery](#), Tables 17–19).

Dose quality (fidelity)

For the intervention condition, we had planned to audio record 20% of sessions delivered at each time point (4 sessions in total × 9 participants = 36) and for the control condition, we had planned to audio record 20% of the care-as-usual interventions (4 sessions in total × 9 participants = 36). However, we were unable to secure approval from the England study site to take in audio recording equipment, which meant that the in-prison intervention and control conditions could not be recorded there. Following discussions with the Scotland study site intervention team, recording control intervention delivery was not deemed feasible or practical.

Interventionists at the Scotland study site were aware that we were recording the intervention delivery sessions for the purposes of undertaking intervention fidelity assessment.

The Audio Recordings and Behaviour Change Counselling Index (BECCI)⁷² was used to provide insight into the Behaviour Change Counselling Skills of the interventionists from the audio recording of the APPRAISE intervention delivery sessions. To calculate the overall BECCI score, the mean across the item responses is calculated to provide the Practitioner BECCI score. The BECCI mean score will correspond to the points given on the Likert scales on the checklist.

The interventionists at both study sites received training (see [Chapter 2, APPRAISE intervention training](#)) and self-completed a six-question Likert scale evaluation (see [Appendix 8](#)) on completion of the training.

Contamination

During the qualitative interviews with the participants and interventionists and those delivering care as usual, we explored the possibility of contamination. This was with the purpose of identifying any evidence of contamination between those in the intervention and control arm and/or the intervention and non-intervention (care as usual) workers. We had hoped to note the wing/area in the prison where the participant was located and which other participants in the study they were likely to have most contact with. However, this was not feasible in terms of the practicalities within the running of the estates at each study site and the impact of COVID-19.

Mechanism of change

Quantitative and qualitative data were obtained to explore the mechanisms of change and the extent to which the intervention changing process variables were consistent with the underpinning theory.

The six proposed mechanisms of change in the logic model are: self-efficacy, outcome expectations, goals, plans, social support. These quantitative measures are discussed in [APPRAISE intervention: theory, elements and logic model](#). In the pilot study the causal mechanisms were measured to provide increased understanding of how the delivered intervention influenced change through the quantitative assessment of key behavioural markers (mediators) of change. These were to be assessed at TP0, TP1 and TP2 and an exploratory prognostic analysis undertaken to explore the nature of change within these domains and their relationship to observed outcome.

In addition, we also conducted a qualitative exploration of study participant and stakeholder responses to, and interactions with, the intervention, providing the opportunity to identify acceptability, barriers and facilitators.

Context

We undertook qualitative interviews with study participants and wider stakeholders to contribute to our understanding of the context within which APPRAISE was delivered, to enable the delivery and or impact of the intervention at scale. We also recorded any differences in delivery in and across study sites where local organisational and devolved country contexts may have had implications for future intervention design and delivery.

Eligibility and recruitment to interviews

Those eligible to be interviewed were drawn from those registered in the pilot trial and from a range of stakeholders who had been involved in different aspects of the study or where, if this intervention were to be implemented at scale, they would be involved in some capacity.

Recruitment to all interviews was significantly impacted by the ongoing COVID-19 pandemic. The challenges of being able to locate men on remand hampered our ability to recruit them to take part in the process evaluation interviews. Similarly, the availability and capacity of some stakeholders (commissioners, policy-makers and prison healthcare professionals), as a result of the ongoing COVID-19 pandemic, meant it was not possible to secure interviews. Interviews were conducted at a time convenient to the participant and were conducted by JF and VG.

Sampling

Participants for all the interviews were selected via purposive sampling within three main groups.⁷³ The proposed number of participants to recruit was 40 interviews in total across the two study sites: men on remand from intervention ($n = 8$ at each site) [$n = 16$] and control arm ($n = 8$ at each site) [$n = 16$], $N = 32$; stakeholders (interventionists and those delivering usual care, prison governor, prison officers, commissioners, policy-makers, third-sector partners and prison healthcare professionals) ($N = 8$). However, due to the impact of COVID-19 we were unable to reach this target.

Participant group 1: men on remand who had been enrolled onto the APPRAISE pilot study and had been randomised.

Participant group 2: interventionists, who were invited to take part if they had delivered the APPRAISE intervention to study participants.

Participant group 3: stakeholders, who were invited to take part if they had been involved in different aspects of the study or where, if this intervention were to be implemented at scale, they would be involved in some capacity.

Informed consent

Each participant was required to provide their informed consent before taking part in an interview. We reinforced to participants that taking part in the interviews was voluntary, even if they had been involved in the main study or had delivered the interventions. It was also imperative to ensure that a participant's decision to take part was based on a clear understanding of what was involved. All participants, therefore, received a verbal explanation of the study, at the time of the qualitative data collection, even in the case of male remand prisoner participants or APPRAISE interventionists. The verbal explanation included the provision of a PIS and then the interviewer went through all the elements in the PIS. When this had been completed, those in Stakeholder 1 group did not have to provide further written consent at this stage as they had already provided written consent on entering the study, which included taking part in future interviews. For those in Stakeholder 2 and 3 groups, each had to sign a consent form.

All participants were given the opportunity to ask questions or to clarify anything that was not clear, or they did not understand. In addition, all participants were reassured that they did not have to answer any questions that they did not understand or that made them feel uncomfortable and that they could withdraw their consent to taking part, at any time during the interview, and even following data collection.

Data management

During the timeline of the study, the COVID-19 pandemic came into play and therefore many of the interviews had to be conducted virtually using videoconferencing software or via telephone with Dictaphone facility as all interviews were to be transcribed verbatim. In Scotland as COVID-19 restrictions eased in the prison three of the four remand participant interviews were able to take place face to face and were digitally recorded, with the fourth taking place by phone.

Audio files were uploaded securely to a password-protected site by the researchers, from which the approved transcription company was able to access the files. On return, a quality check was conducted to ensure the accuracy of the transcribed files and then all audio recordings were securely destroyed, and the resulting transcripts were stored on password-protected drives at the University of Edinburgh and Teesside University. Upon storage, all interview transcripts were anonymised, with any personal or identifiable participant information being removed, and each transcript was assigned a unique participant number for analysis purposes.

Any pieces of hard-copy data from the Scotland site, such as the completed consent forms with a participant's name on, were sent by secure courier to the England study team.

Qualitative data analysis

Following the transcription of the interviews, Braun and Clarke's approach to thematic analysis was used, supporting a structured yet reflexive analytic process.⁷⁴ Because of the atheoretical nature of Braun and Clarke's approach, we chose to use NPT as a lens to support thematic identification.⁷⁴

Thematic data analysis is often underpinned by theoretical concepts. In this study, we employed the NPT, an implementation theory, to better understand what the qualitative data could tell us around

implementing the APPRAISE intervention for future work. NPT was developed to explore and develop the understanding around the factors affecting whether an intervention can be incorporated into practice, and the context in which the work of the new intervention happens.⁶⁹⁻⁷¹ NPT aims to facilitate a better understanding of the challenges arising during the embedding of a new practice by considering the components of four different constructs; Coherence ('sense-making'), Cognitive Participation ('relational' work), Collective Action ('operational' work) and Reflexive Monitoring ('appraisal' work).⁶⁹⁻⁷¹ The theory lends itself well when considering qualitative data around implementing a new intervention in different contexts and in this work was used as a tool to structure the thinking around the different factors arising that would affect the future implementation of the APPRAISE intervention.

The interview data in the APPRAISE study were analysed within each of the three main groups; remand participants, CGL team and wider stakeholders. The first stage consisted of data familiarisation, essentially reading through the transcripts to elicit any initial thoughts and feelings emerging from the data. The next stage involved forming early codes from the data, the process of which is referred to as 'coding'. A code refers to 'the most basic segment, or element, of the raw data or information that can be assessed in a meaningful way regarding the phenomenon'.⁷⁵ Each transcript was read and coded by JF and was then double coded by GW at data analysis meetings to ensure consistency and refined further or developed further as necessary. The next stages involved the codes being assimilated and emergent data themes were mapped against the four main NPT constructs. Areas of divergence and discrepancy were explored and the data checked for fit.

Chapter 6 Process evaluation results

I think COVID and the impact of COVID, we'll be dealing with this for a number of years going forward. And I think the longer term effects of COVID, and some of the almost the PTSD from COVID is something that's going to play out over the next few years

D400S

As described in [Chapter 5](#), quantitative and qualitative methods were employed to provide the data required to answer the process evaluation questions. The results of the implementation dose and quality as well as the mechanisms of change associated with the intervention are presented first. This is then followed by the results from the qualitative interview data, which focused on the feasibility and acceptability of the APPRAISE study intervention as well as any reflections on potential contamination and the context within which the intervention was delivered.

Firstly, it is important to acknowledge that the impact of the COVID-19 pandemic was felt by all participants. The pandemic resulted in prison lockdowns, social distancing and increased social isolation; pandemic measures reduced the prison staff capacity and the space available within the prison.

All participants reported on the impact of the COVID-19 pandemic on mental health. The lockdowns were identified as heightening mental health issues, by facilitating social isolation and reducing available support to the prisoners even further. The stakeholders and the interventionists recalled struggling with the different and ever-changing demands within the prison, while dealing with the unprecedented lack of social contact and restrictions outside of the prison.

Intervention implementation results

To confirm intervention implementation, data collection was planned in two ways: intervention booklets (dose delivered) and audio recordings (dose quality).

Dose delivered results

The full results of the in-prison and post-liberation intervention doses delivered are presented in [In-prison and post-liberation intervention delivery, Tables 17–19](#). To recap, of the 68 participants randomised to the intervention condition, 53 received the in-prison intervention session (77.9%, 95% CI 66.2% to 87.1%). This percentage is greater than the 70% threshold specified as one of the criteria necessary for progression to a future larger randomised controlled trial (Objective 4c). Reasons why the in-prison intervention was not delivered are shown in [Table 18](#). They varied from participant declining, being sentenced, released, transferred, in crises, had COVID-19 and unknown.

Only one 'day-3 post-prison intervention' session was attempted and was delivered (duration 30 minutes) at the Scotland site. No 'day-7 post-prison intervention' sessions were attempted. Only one 'day-21 post-prison intervention' session was attempted and was delivered to one participant (duration 30 minutes) at the England site. The main reasons why participants did not receive the post-prison interventions were (1) they could not be contacted and (2) CGL staff were unable to contact participants due to the impact of the COVID-19 pandemic.

Data for the duration (minutes) of the intervention sessions across both study sites are presented for all but one of the 53 interventions delivered in [Table 19](#). There was a marked difference in the median time taken to deliver the intervention between the two sites: the intervention duration was half an hour in England compared to 1 hour in Scotland. The shortest time for an intervention was only 5 minutes in England whereas in Scotland it was 30 minutes. The maximum time for delivery was more aligned across sites, being 70 minutes and 80 minutes in England and Scotland, respectively.

Dose quality (fidelity) results

Interventionist training evaluation scores

Six practitioners from CGL took part in and participated in the training. Two interventionists and two team leaders at the England site and two interventionists at the Scotland study site (see the section [Intervention implementation](#)) completed a five-item 'agree' to 'disagree' Likert scale evaluation asking the trainees to score the Intervention Training sessions (see [Appendix 8](#)). Ratings for the six questions were all 4 and 5, with the exception of three practitioners from England who rated Q5 'The videos used during the training were helpful' a 3. One England practitioner also rated Q3 'The training was what I expected' a 3.

England interventionists and team leaders provided some written evaluation feedback. There was no written evaluation feedback from the Scotland interventionists who completed the training.

1. 'More understanding now'
2. 'Much more info now – trainer knew her stuff and explained really well – video clip delivery not scripted'
3. 'Thank you, nice relaxed training'
4. 'The intervention video – too long and should just be an overview, not full session'

The Audio Recordings and Behaviour Change Counselling Index scores

During the course of the pilot trial, a total of three intervention sessions were successfully recorded at the Scotland site. Three recordings were reviewed and the BECCI instrument was completed. The mean Practitioner BECCI score for each of the recorded sessions was 4, which indicates the interventionist has been practising BCC to a great extent for all three of the recorded APPRAISE intervention sessions.

Contamination results

During the qualitative interviews with the participants and interventionists and those delivering care as usual, we explored the possibility of contamination. This was with the purpose of identifying any evidence of contamination between those in the intervention and control arm and/or the intervention and non-intervention (care as usual) workers. There was mention of discussion of the study between one of the remand participants and another (see sub-section 'Coherence: acceptability of the APPRAISE intervention' in the section 'Remand participants' below) but the participant stated that he did not know if the other participant got the intervention or not. No other evidence was obtained of contamination; however, it would be prudent to explore this again in any future work in the absence of a global pandemic, lockdown and restrictions across the prison estate. We had hoped to note the wing/area in the prison where the participant was located and which other participants in the study they were likely to have most contact with. However, this was not feasible in terms of the practicalities within the running of the estates at each study site.

Mechanism of change results

Quantitative and qualitative data were obtained to explore the mechanisms of change. The results of the quantitative data are presented in [Chapter 4, Outcomes, Table 28](#) and [Supplementary Material 1, Tables S3.1 to S3.7](#), and [Table 29](#) and [Report Supplementary Material 1, Tables S2.1 to S2.6](#).

In addition, we also conducted a qualitative exploration of men on remand, and CGL team and stakeholder responses to, and interactions with, the intervention, providing the opportunity to identify mechanisms of change and influencing factors (see section 'Interview results' below).

Context

Interviews with the three participant groups contributed to our understanding of the context within which APPRAISE was delivered, to enable the delivery and or impact of the intervention at scale. We also explored any differences in delivery in and across study sites where local organisational and devolved country contexts may have had implications for future intervention design and delivery. The next section provides the interview findings.

Interview results

Fifteen semistructured interviews were conducted with three participant groups from across the two study sites (Scotland and England) and are presented by participant group in [Table 31](#).

All remand participants were from the Scotland study site only. For the CGL team and wider stakeholder groups, participants from each of the two study sites were distinguished by their participant code, by using an E for England and an S for Scotland.

Remand participants

A total of four male remand prisoners were interviewed. Due to the COVID-19 regulations, following the completion of the recruitment of participants to the study, England project staff were unable to be granted further face-to-face access to prisoners. Therefore, as previously mentioned, all prison interview participants were from Scotland. The key themes emerging from the remand participants' reported data aligned to each of the NPT constructs respectively were: Acceptability of the APPRAISE intervention; Utility and Value to ensure 'buy-in' in the APPRAISE intervention; Gaining trust and 'opening up'; and

TABLE 31 Breakdown of the interview sample by group

Participant group	Participants	Number of interviews	Participant codes
Participant group 1: remand participants	Male remand prisoner participating in APPRAISE pilot study	4	Participant A
			Participant B
			Participant C
			Participant D
Participant group 2: CGL team	APPRAISE Interventionists	6	E200S
			E300S
			S100S
			S200S
			S300S
			S600S
Participant Group 3: wider stakeholders	Prison staff RAs	5	E100S
			E400S
			E500S
			S400S
			S500S

finally Provision of ongoing support to affect change (Table 32). Within each theme there were sub-themes drawn from the data that collectively provided evidence to support our understanding of the experiences of the remand participants interviewed who had received the intervention.

Coherence: acceptability of the APPRAISE intervention

When exploring the responses around the initial screening process, using the baseline questionnaire to assess alcohol use, in the main, participants were happy to answer all of the questions in the questionnaire and, in general, the questions appeared to be acceptable to the population group, with only one participant saying they found some questions too personal.

I did feel comfortable about the questions and I understood what they meant, most of them nay bother eh I felt fine answering them

Participant A

Some of them are a bit personal like, but I suppose I felt alright about them, aye

Participant B

The APPRAISE intervention was reported by the remand participants as being useful and having a purpose.

Good because like, like I say, I didn't ever talk to anybody about, so it's good to have somebody that you can talk to, and say I'm feeling like this and tempted to like, use drugs or drink

Participant B

It seemed useful, do you know what I mean, to help me out going forward, you know what I mean? Get a better understanding, and not going back to old habits basically

Participant D

In addition, some participants were already able to reflect upon being able to see the benefit of engaging with such an intervention. The data for Participant B also provide evidence of some form of acknowledgement between remand participants about the intervention.

I don't even know if he got the intervention, but he said he had spoke to yous. And they, it seemed, like he'd already stopped drinking

Participant B

The data suggest that remand participants were cognisant of the APPRAISE intervention and could reflect upon being participants in the study, the purpose of it and how it could be useful.

TABLE 32 Remand participant summary of NPT informed interview themes and subthemes

NPT construct	Theme	Subthemes
Coherence	Acceptability of the APPRAISE intervention	<ul style="list-style-type: none"> Acceptability influenced participation in the new intervention. Remand participants viewed the intervention as having a useful purpose.
Cognitive participation	Utility and Value to ensure APPRAISE intervention 'buy-in'	<ul style="list-style-type: none"> Perceived value influences motivation and willingness to change.
Collective action	Gaining trust and 'opening up'	<ul style="list-style-type: none"> Therapeutic relationship shapes the implementation. Those involved in delivery key in facilitating trust.
Reflexive monitoring	Provision of ongoing support to affect change	<ul style="list-style-type: none"> Practicalities of providing continued support perceived as being required.

Cognitive participation: utility and value to ensure APPRAISE intervention 'buy-in'

When considering the implementation of the APPRAISE intervention on a wider scale, a key contributor to this, which is incorporated within the Cognitive Participation construct of the NPT, is the notion of participants recognising the value of an intervention and hence facilitate their 'buy-in'. Participants were seen to value the idea that by receiving support to reduce their drinking, it could have a positive impact on other elements of their life, including reducing re-offending and maintaining their relationships. A specific example of this was a participant who talked about being able to see the benefit of the intervention in allowing individuals to reduce their offending rates as a positive consequence of reducing their drinking.

I think it's important, I believe that most people who have committed offences have been on alcohol or drugs eh? So, I think it is maybe important to hear people like come in prisons and say they can do stuff. And speak about drink, because there's big problems there, it is. If they didn't drink, they probably wouldn't be half the people in prison. If they never had the problems with a drink. Yeah. So, if the good guys get a bit of help, then maybe it would make them not commit their offences and they wouldn't come back.

Participant A

The notion of possible utility and value of engaging in such an intervention appeared to link to the acceptability to engage and the motivation and willingness to change their current drinking behaviour. Remand participants recognised that they needed help with reducing their alcohol consumption outside of prison, from completing the initial baseline question and intervention session, and hence appeared to welcome receiving support and having a new way of approaching it.

I wanted that support, I wanted that definitely, and into, I wanted that into it as well, you know, how you can change and how you can do better with your life, you know what I mean? It's just that wee bit knowledge, that what I never had and that goes a long way when you've got it and then you're off doing it, you know what I mean?

Participant C

Within the data, remand participants were seen to talk about not knowing where to go to access or receive the support or help they needed when early in the process of thinking about reducing their alcohol consumption.

If you've got that from inside going out, then you've got more of a chance I would say of you actually doing it.

Participant C

Therefore, as the first intervention session was delivered within the prison setting, it was seen as beneficial to participants to allow them to consider new habits once liberated.

Collective action: gaining trust and 'opening up'

The idea of gaining trust and building a rapport during the first intervention session was reported recurrently, and is one which is highly indicative of the Collective Action construct of the NPT. Building and maintaining trust and having staff members who possess the relevant characteristics and qualities to be able to do so was seen to significantly increase the acceptability of the APPRAISE intervention.

some people just cannae, I don't know they just cannae talk to people, do you know what I mean, for whatever reason, so I, that might be like your only barrier would be like getting through to people and saying, listen were just here to talk about this, if you want to talk but just many people understand why they don't want to talk to people about things but, but whatever went on in their life basically they're not willing to share.

Participant B

The issue of being unable or not wanting to talk about or vocalise problems to staff was discussed by the participants. However, one participant talked about feeling more comfortable to talk to a peer and it feeling more acceptable and relatable to engage with someone who had lived experience or who could empathise with an individual's situation and offer a positive pathway for change.

I mean, not to say you're intimidating or anything like that. You're not, but it's like, men don't like to talk generally, especially on issues. They kind of close down. But yeah, you need to get some people inside, I guess. You know, so if you get someone here that goes out and reaches out to people in the hall there, that would help I think. It's easier than them coming to you. If there's somebody like me going to them, and going like, listen, man, I think that'd be good for you, I've done it and I can tell you this, this is what's helped me, this is what it's like, you don't have to be fearful, if you don't want to, you know what it is, you don't have to do it. You don't have to say this. You don't have to say that. You know but give it a chance.

Participant C

It was clear that being able to open up and to speak to the individual delivering the intervention was important to the prison participants. The same participant talked about the importance of building a rapport with the support worker and gaining a relationship of trust to facilitate the delivery of the intervention.

I've got a lot of trust, I've got a lot of trust in this lassie. And that's only because she was helping me, she was helping me with things I didn't know nothing about, do you know what I mean?

Participant C

The sharing of information with the intervention delivery staff emerged as key. A challenge around the acceptability of the APPRAISE intervention was observed when the prison participants talked about being unable to 'open up' and talk to a new interventionist or support worker about their personal experiences or alcohol consumption.

Reflexive monitoring: provision of ongoing support to affect change

When asked about what would improve the intervention session or add to its usefulness or making it worthwhile, participants considered the offer of a continuous and ongoing package of support to be most useful.

Continuous, continuous support, you know what I mean. And letting people know, you're definitely having the possibility of contacting people, you know, you're just not being left to your own devices, to your own thoughts.

Participant D

All four participants welcomed the idea of having continuous and ongoing support, that they could access easily and build on the content of the intervention, in times of need for additional support.

The discussion on the post-release intervention sessions identified them as being less acceptable and feasible to remand participants. The remand participants talked about the importance of the post-release intervention support being obtained quickly, before an individual was able to return to their old habits.

I think if that, work, that help was still there right at the gate, waiting for the guy to go out to you'd have many a chance of like benefiting.

Participant C

However, many of the participants reflected upon the challenges of a participant being released from prison and establishing any reduction in their alcohol consumption, regardless of receiving an ABI

in prison. They talked about an individual being released from prison and then finding themselves in situations where alcohol was freely accessible again or being in the company of other individuals that they used to drink with or who were drinking around them and falling back into previous risky drinking habits.

I dunno like, some people might work like, aye you might work with people in the jail, and like outside the jail, but it would be a lot harder outside the jail, because obviously other people could come in here with all good intentions of going to interventions, or whatever it may be, but when it gets out there it's all going to be different, eh?

Participant B

On a more practical point, participants spoke about the difficulties that they would face in contacting an interventionist once they were released and vice versa. One participant talked about lacking a phone, and hence being unable to receive calls to arrange a follow-up session with a support worker or to contact them quickly if he needed any additional support.

You see now, outside everybody expects people to have phones. I was like, I've not got a phone, you know what I mean, so, they'd look at me and go, well some of my workers would go, 'what you've not got a phone?' How, now, wait a minute and they might now think different and go, 'we need to write you a letter'. You know, and that, a letter can take days to try and get to you, whereas a quick phone call is like there and then isn't it.

Participant C

Therefore, different methods of keeping in contact with men post-release appeared to be a key area warranting further consideration in future work, to facilitate successful implementation and success of the intervention over time.

Change Grow Live team participants

A total of six members of the CGL team participated in the semistructured interviews. The team members were all responsible for delivering alcohol and substance use interventions, within the prison setting, prior to the study and were all employed by the respective commissioned service provider at the time of the study taking place. This included those who delivered the APPRAISE intervention at both study sites and had received the intervention training. From the CGL team participant data, four key themes emerged: making sense of new ways of intervention practice; engaging in the APPRAISE intervention content; investing in alcohol prevention interventions; and scaling up the APPRAISE intervention ([Table 33](#)). Subthemes for each were identified, providing an insight into the context and system within which the intervention was delivered and providing key areas for recommendations if the APPRAISE intervention were to be scaled up and implemented. The CGL team participants reflected upon several factors related to the prison setting and alcohol services that they reported were relevant to the acceptability, feasibility and implementation of the APPRAISE intervention.

Coherence: making sense of new ways of intervention practice

In order to deliver the APPRAISE intervention, the CGL team who were to be involved in delivering the intervention and supporting the study were required to undertake APPRAISE training. Even though they were likely to have been delivering/managing/supporting a form of alcohol brief intervention within the prison setting previously, they were asked to complete training. Interview participants were asked about the training they received to determine whether it was fit for purpose and whether it was acceptable and feasible to them. Training is a key part of the NPT's Coherence construct as it relates to the participants' understanding of what was expected of them and being able to see the value of a new intervention or practice.

TABLE 33 Change Grow Live team participant summary of NPT informed interview themes and subthemes

NPT construct	Theme	Sub-themes
Coherence	Making sense of new ways of intervention practice	<ul style="list-style-type: none"> Acceptable and appropriate intervention training instruments 'fit for purpose'. Currency in understanding the differences between current and new practice.
Cognitive participation	Engaging in the APPRAISE intervention content	<ul style="list-style-type: none"> Encouragement and motivation viewed as important part of engaging in a new practice.
Collective action	Investing in alcohol prevention interventions	<ul style="list-style-type: none"> The perceived conflict and reality of alcohol being left behind. Time and space required for delivery of APPRAISE intervention.
Reflexive monitoring	Scaling up the APPRAISE intervention	<ul style="list-style-type: none"> Navigating the terrain to understand the impact of the APPRAISE intervention.

Generally, the training was well received by the interventionists. It was seen to be pitched at an appropriate level for them to understand the components of the APPRAISE ABI and to feel confident with the delivery following the session.

When we came to doing the APPRAISE books we knew exactly, yeah. It went well.

E300S

Even participants who were not feeling enthusiastic about undertaking the training felt that it was a valuable experience and allowed them to gain the skills and knowledge they needed to deliver the intervention.

I was quite keen not to go through it. But I'm glad we did. Because otherwise I wouldn't have known that tool so well.

S200S

The less acceptable elements of the training appeared to be the role-play components, the amount of paperwork and the overall length of the session. One participant said they did not usually enjoy role-playing in training sessions, but they agreed that it added value to the training, in developing their understanding around the best ways to deliver an intervention to a prisoner.

I'm always apprehensive about role playing. It's not anybody's favourite thing to do. But actually, it was, it was quite good.

S200S

The feedback about the length of the training was that the session was too long, and that the content could be covered effectively in a shorter time frame.

I'm thinking maybe even just the one half-day probably would have been enough. Erm, cos yeah, I sort of felt like I had it after the first one, kind of thing.

S100S

In addition, the same participant felt that the amount of materials provided was overwhelming, and that they didn't have time to look at them post training.

I remember getting a lot of paperwork though, supposed to read through and didn't. I had every good intention of reading the paperwork. Yes. Just never found the time.

S100S

When talking about the intervention itself and the content, the interventionists were largely positive and indicated that it was well received by both the prisoners and the delivery staff.

everyone we talked to thought it went very well.

E3002

They seemed like the, you know, it was interesting for them, it was a little bit different. Yeah, no, it was, it was good. Yeah, there was a little bit of resistance with some, but not with many, I would say.

S2005

The fact that the ABI was different from the usual treatment and was 'more structured' and 'repetitive' in content appeared to be a facilitator for wider implementation.

I think there was actually a bit of intrigue around it, just because it was something new and something different. And you tend to find that does sort of pique people's interest? That actually the idea of this something new, something different. That might be what works for them? Was something worth sort of buying into?

S1005

Being able to see the difference from current ways of working or usual care is part of the Coherence construct of NPT. Being able to identify how the APPRAISE intervention is different and valuing those differences can be a key facilitator of early implementation.

Cognitive participation: engaging in the APPRAISE intervention content

Similarly, to the remand participants, the CGL team participants stressed the importance of staff motivation and encouragement. Building a rapport between the prisoner and the interventionist delivering the intervention meant the prisoner was more likely to share their alcohol consumption habits and engage with the intervention content.

You know, just conversation, building some kind of rapport and relationship.

S2005

That's the most important thing. Motivation and encouragement that's the most important thing.

E3002

CGL team participants reported their belief that if the staff were motivated and encouraging when delivering the ABI, the prisoners were more likely to engage and take more from it. This is indicative of the Cognitive Participation construct of the NPT, which considers how key individuals can drive the intervention forward and support the implementation of an intervention. If the APPRAISE interventionists are motivated and can see the value of such an intervention, this could facilitate the implementation further.

The need for staff buy-in is also reflective of the Cognitive Participation construct of NPT, where implementation is supported by the relevant individuals buying into an intervention. The issue of staff capacity and space is encompassed within the Collective Action construct, which relates to the need for those involved in delivery of the intervention to have the motivation, capacity and ability to deliver it.

Collective action: investing in alcohol prevention interventions

From the interviews it was clear that the services and treatment options that were currently available to the men on remand were limited. The interventionists valued an ABI that was targeted at men on remand, as they generally had fewer services available to them, and it provided a window of opportunity to offer support post release.

There are a lot of services in here that are only available for people who have been convicted.

200S

The lack of service and treatment options available to the remand population at the time the interviews were undertaken was accompanied by reports that alcohol consumption was being viewed as less important or the 'lesser of evils' within the remand prison setting. One participant labelled alcohol as 'being continuously left behind', as the primary focus tended to be on substances that were more likely to cause significant damage in the short term, by overdosing or causing immediate harm.

I think alcohol is left behind a bit. A lot of concentration is on heroin, and methadone. Because the powers that be are so worried about people overdosing, which, obviously we all are.

E300S

CGL team participants reported how it had been eye-opening for the remand prisoners involved in the study to better understand the risks associated with alcohol consumption as opposed to drug use.

On a more practical level and building upon the prison setting as a suitable or feasible location for intervention delivery, a commonly reported issue was the lack of space. All of the participants reported room availability being very limited and often interventionists had struggled to find acceptable rooms to deliver the APPRAISE intervention.

Issues with rooms is kind of permanently a thing. And there, there's never enough space.

S100S

As the interventionists were delivering the APPRAISE intervention over and above their usual service provision and need for rooms, room availability was even more stretched than usual and available space was very limited.

I couldn't, because there weren't enough rooms to facilitate the APPRAISE project, whether it was you in with interviewing somebody or [name removed] over to do the intervention with somebody. And it was just like for goodness sake, but you know. But, again, that's life. And that's how it goes. And that is just one of the ties with having such little room in this place.

S100S

more dedicated time, more dedicated space. And at the end of the day, you need, you need that buy in.

S100S

This is reflective of the Collective Action construct of NPT as it highlights the importance of how the host organisation, that is, the prison estate, is able to adequately support the implementation of the new practice; the pressures on suitable room availability could hinder the feasibility of an intervention future roll-out.

Reflexive monitoring: scaling up the APPRAISE intervention

A specific element of the intervention content that the interventionists from the Scotland study site reported they had found was a challenge for men on remand during intervention delivery was in navigating the variable numbering system of different scales. Two CGL team participants talked about the need to standardise the numbering so as to not exclude or frustrate the interventionists during delivery.

But the one thing that confused me and confused a few of the guys that I did the brief intervention with, was the change in the numberings. So, I think one of the, some of them were between one and five, and some of them were between one and 10. And that did throw people off, myself included.

S100S

One participant talked about the difficulties of maintaining implementation fidelity in an ever-changing, dynamic environment, such as the remand prison setting.

different workers have different approaches and different service users have different appointments.

S300S

Similarly, to the remand participants, the CGL team participants found the post-release follow-up sessions to be less acceptable and feasible to conduct in practice. Follow-ups were made challenging by inability to make contact with prisoners post release and this was further hampered by the COVID-19 pandemic, which made it impossible to meet face-to-face. The interventionists reflected upon the different methods of establishing contact with the prisoners. Letters were often seen as helpful, as a tool for prisoners to refer back to, and also sending text messages, prior to making a phone call, to introduce who was calling them.

sending a letter can be quite helpful, because sometimes it's just that kind of post release panic of trying to get everything sorted out. And then the dust settles, and then they go, 'Oh, yeah, I was offered that support'.

S100S

I would send that text explaining who I am because I know these guys and they won't answer numbers they don't know.

S600S

After being a part of the APPRAISE pilot study, the interventionists appeared to be positive about it being scaled up and extended as a larger study to determine the effectiveness of the intervention and could see the potential of the APPRAISE intervention in the setting.

yeah, I think it's, a bigger study would be good. Very definitely.

E300S

I quite liked the project. Would be nice to see it done like on a larger scale because I do think it has, it has potential.

S100S

When asked to reflect upon the factors that would enable the delivery and facilitate future scale-up, a key facilitator of the pilot study in the setting was the researcher being able to understand the context and the management of the prison and hence being flexible and accommodating to any unexpected situations that arose.

I would say that the research team was flexible, which was good, and because they appreciated the difficulties and the pressures that we are often under as a team, due to various factors such as staffing, for example.

S300S

Being flexible aided the acceptability of conducting a research study within a prison and the researchers, within the APPRAISE study, were able to build a rapport with staff and support them with any issues as required, which would also be important within a future trial or considering future roll-out.

A commonly talked about barrier to research within the prison was capacity, in addition to the issue of space. Thinking specifically about staff capacity, interventionists talked about already having a high case load of prisoners, prior to working on the APPRAISE study, and therefore they often felt apprehensive about the thought of a research study adding to their burden.

I think concern for anything that we end up doing that's an addition to somebody's core duties that they do, there's always just like, concerns about how that's fitting in. But then, I think they had a bit more reassurance once it's like, 'okay, we're only going to ask you for this amount of commitment'

S300S

Therefore, exploring ways to ameliorate these issues would contribute to future feasibility of roll-out. On a positive note, the interventionists were in agreement that the APPRAISE study did not add significantly to their workload in practice. However, it has to be remembered that the majority of the post-liberation intervention delivery did not take place due to COVID-19. And participants did talk about how they thought a larger-scale trial in the future may be more difficult in terms of capacity.

would just be capacity for workers so I guess if it was a larger study, it's just how?

S300S

Stakeholder participants

The final group of interview participants consisted of stakeholders who were involved with the delivery or the management of the APPRAISE pilot study. This group consisted of five participants ranging from the researcher who was managing one of the study sites and going into the prison to recruit the participants, to prison staff who were involved with the undertaking of the study and recruitment. The four main themes were: understanding the diversity of the prison population for APPRAISE intervention delivery; acceptability and utility of APPRAISE intervention champion; the value of networking; and awareness of the individuality of the prison environment and vulnerability of the remand population. Sub themes emerging from the stakeholder interviews are presented ([Table 34](#)).

The themes and subthemes emerging from the stakeholder interview data were very similar to those of the interventionist data. Similar findings were reported around the prison's logistical constraints affecting the APPRAISE intervention feasibility, such as space and staff capacity, the importance of engagement with prisoners and the lack of currently available support for prisoners on remand. Overlapping findings are not discussed further in this section to avoid repetition. The novel themes and subthemes that came out of the stakeholder data were around the intricacies of the prison environment and how it affected the study and intervention delivery, the individual differences across the setting and across the participants, and further barriers and facilitators to delivery of the APPRAISE intervention.

TABLE 34 Wider stakeholder summary of NPT informed interview themes and subthemes

NPT construct	Theme	Subthemes
Coherence	Understanding the diversity of the prison population for APPRAISE intervention delivery	<ul style="list-style-type: none"> • Cognisance of different drinking patterns of the prison population. • Overly complex language as a barrier where literacy and comprehension are a factor.
Cognitive participation	Acceptability and utility of the APPRAISE intervention champion	<ul style="list-style-type: none"> • Relevance of lived experience as an influencer in the delivery of alcohol intervention.
Collective action	The value of networking	<ul style="list-style-type: none"> • Joined-up different treatment services can shape APPRAISE delivery.
Reflexive monitoring	Awareness of the individuality of the prison environment and vulnerability of the remand population	<ul style="list-style-type: none"> • Modifying relational work in the appraisal of a new practice or intervention.

Coherence: understanding the diversity of the prison population for APPRAISE intervention delivery

The need for joined-up care and signposting to different services links to another key finding from the stakeholder data, around understanding the remand prison population. A key facilitator in ensuring that the APPRAISE intervention was acceptable and feasible in practice was the delivery staff being able to understand the population. As the stakeholder group included mainly prison staff, it included people who understood this population well.

we need to understand the cohorts we're dealing with as well as the remand group, particularly younger drinkers and particularly the kind of thinking around it.

E500S

This included being able to understand that younger drinkers were likely to have different drinking patterns, in comparison to an older population group. Therefore, having an understanding of the diverse population group and how to work with them would be advantageous when considering the future delivery and wider implementation.

A barrier which was reflected upon briefly in the prisoner findings, and directly linked to understanding the population involved, was considering the language used in both the intervention and questionnaire. Several stakeholders who had significant experience working with remand prisoners talked about the need to simplify the language for it to be acceptable for the population. The findings highlighted that the population group often had literacy or comprehension difficulties.

I get that I think some of it is come from, you're already, you know, evidence based and use trials. You know questionnaires, but you're working with a different clientele?

E100S

Another stakeholder participant talked about the fact that the language that was acceptable to academics isn't always acceptable for this population group. This led to the discussion around the need to 'demystify' overly complex language.

I think the other things is finally around brief interventions is to demystify an awful lot of language that people use in treatment eh I think academics are the worst for it.

E500S

Cognitive participation: acceptability and utility of the APPRAISE champion

Thinking about some of the findings which might increase the acceptability and utility of the APPRAISE intervention in practice, it was reported by one stakeholder that the use of 'intervention champions' could be advantageous. The stakeholder talked about the importance of having an individual, within the prison setting, who clearly saw the purpose of the intervention and who was motivated to engage others to deliver and implement it. The stakeholder participant was able to see the effectiveness of having an intervention champion in other settings, therefore they believed it would be useful to employ this strategy within the remand prison setting.

they become your champions, they do it everywhere else why not do it around this particular intervention, because when they are gone, they trained the next leader.

E500S

This finding is concurrent with the Cognitive Participation construct of the NPT, as it includes the concept that key individuals are responsible for driving an intervention forward. Similarly to the prisoner participants, the stakeholder participants could also see the benefit and relevance of employing people with lived experience of alcohol use or the prison setting, in the intervention delivery process, as it appeared more acceptable to the prisoners.

that's where lived experiences are really important, peer support, peer mentors, mutual aid groups, recovery networks, visible recovery.

E500S

The discussion around prisoners and their individual needs is linked to another finding that prison staff were often apprehensive about allowing research to be undertaken in the remand setting. One participant talked about how they felt protective of prisoners taking part in research studies and that they had to be able to see the benefit to the prisoner in taking part in the research, before allowing a study to take place.

Some staff are very protective of their prisoners as well and they don't like to see tours coming in or high school kids coming in, or university guys just coming in, because we don't want people in our care to be treated like, you know, it's like a fishbowl.

S400S

This is again reflective of the NPT Cognitive Participation construct, as the stakeholders needed to be able to see the value of the new ABI to the prisoners, before agreeing to take part in the research study.

Collective action: the value of networking

Another perceived facilitator to the APPRAISE intervention delivery was being able to work with other services to create a network and to offer signposting, and stakeholder participants frequently talked about the importance of adopting a 'joined-up' approach. One participant talked about promoting a 'social network', to support the different treatment services.

If your cat is sick and your car breaks down who do you ring? You don't ring the same person, so from that perspective that's why networks are really important, your social networks, so what you need to understand is what social networks and what other services do they buy into, where do they go.

E500S

Working with other services and the sub-theme of trust came across in the findings. This is reflective of the Collective Action construct of NPT, as it includes the concept of participants being able to maintain their trust in each other's work and expertise throughout the work of an intervention, such as the APPRAISE ABI.

Reflexive monitoring: awareness of the individuality of the prison environment and vulnerability of the remand population

As expected, the remand prison setting was found to be dynamic and largely unpredictable. This meant that there was a constant pressure felt by the prison staff to be able to adapt to the changing environment and situations.

my main interest is in this place is keeping the place safe, keeping it decent, but keeping people alive when they come in, because I manage an awful lot of chaos in a very confined custodial setting.

E400S

They've also usually got quite a lot of chaotic things happen in the background.

S400S

Specific challenges that were discussed across the two study sites, and at times which affected the study and recruitment, included prisoners' illicit drug and alcohol use, physical violence and the prison going into lockdown. Poor mental health was also a commonly reflected-upon issue that added to the unpredictable nature of the population. The reported stigma of being on remand was felt to contribute to this, with one participant reflecting upon the challenges associated with a prisoner adapting to their environment.

Just a week on remand can mean family breakdown, loss of job, loss of status, stigma, self-esteem, down the toilet, you know, so that's you, you know, you've got it branded, is right in the middle of your forehead.

S500S

The trauma experienced by the remand prisoners as reported by one stakeholder participant identified that this also extended to their release, and the APPRAISE study findings indicated that this was a contributing factor to why the follow-ups were difficult in practice (aside from COVID-19), as the prisoners struggled to 're-establish their lives' post release.

And you know that that traumatic, that trauma they've just went through, probably might not be the one trauma they have. They're now leaving custody to try and re-establish their life under a great deal of stress and pressure, and probably are here at least in part due to alcoholism.

S400S

The individual nature of both the prisoners and the prison setting was highlighted. Stakeholder participants commonly reflected that what worked in one prison may not necessarily work in another, due to the individuality of the settings. However, there was a consensus that although there were differences experienced across prisons, there would also be a certain degree of consistency.

obviously quite different in general quite different prisons, but you would expect some consistency.

S400S

The idea of individuality also extended to the prisoners and acknowledging their differences and individual needs. Stakeholders talked about the importance of the services, offered to remand prisoners, not just acting as a tick-box exercise, but being able to adapt and encompass prisoners' individual differences.

I think, trying to just keep away from a map that's got tick boxes on it, is better. We need to talk to that individual about their individual needs and how it's affecting their individual well-being.

S400S

One participant talked about how prisoners were often seen to be excluded and labelled as 'hard to reach', due to them not engaging in a particular activity or service. However, this was often more complex than that because prisoners were seen to be likely to be struggling with various issues, or it was found that the service did not meet their needs or was not acceptable to them etc.

services generically are quite often designed where people don't feel they're included.

E500S

Therefore, it appeared to be important to consistently appraise the ABI to ensure it is not a 'one size fits all' approach and that it meets a range of different prisoners' needs. This is reflective of the Reflexive Monitoring construct of NPT, which highlights the importance of modifying relational work in the appraisal of a new practice or intervention. Clearly clarifying which components of the ABI can be subject to modifications and which need to exist as 'core components' will aid implementation in a future trial.

Summary of findings

Analysis of the 16 interviews highlighted key findings from the qualitative work, in regard to the acceptability and the feasibility of the APPRAISE intervention. The in-prison APPRAISE intervention session appeared to be acceptable and feasible to receive and deliver according to all of the participants we asked. The intervention sessions delivered post liberation appeared to be more challenging in practice and less feasible to complete because of a plethora of different factors, such as being unable

to re-establish contact with males upon their release and individuals relapsing to their previous risky alcohol consumption when alcohol was more freely available. Therefore, it would be imperative to explore different ways to make contact with the men post release and to provide more emphasis to strategies of avoiding relapse in future work, if pursuing an intervention which included post-release sessions. Picking up on the value of networking, it may be prudent to explore how the Probation Service could play a role in facilitating making contact with the men post liberation.

The prisoner participants appeared to welcome receiving support for making changes to their alcohol consumption and talked about the utility and value of the intervention in order to ensure buy-in from them. Importantly, they also mentioned the need for the provision of ongoing support and the practicalities of that. Fundamental to the men was the importance of fostering an environment that promoted an individual to feel comfortable and trusting of the interventionist, in order for the men to open up and share their personal experiences and discuss their consumption patterns and potentially other areas of their lives that were relevant to drinking.

The interventionist findings also included the importance of building a rapport with prisoners when undertaking the intervention sessions. In addition, they talked about how the prison setting and culture affected the acceptability and the feasibility of implementing an intervention due to the lack of currently available options to the remand population, the lack of available space in the prison and limited staff capacity. Investing in alcohol prevention interventions were identified as important as other behaviours that impact on health and well-being. The intervention training and content appeared to be largely acceptable and were key to identifying the purpose of the intervention, while understanding the impact of the intervention would facilitate scaling up in the longer term, although it was thought that the time taken and the materials used could be reduced to ensure the feasibility in future roll-out. The delivery significantly benefited from staff buying into the intervention and being motivated and engaged with the content.

The final stakeholder group expressed the need for implementation to fully understand and respond to the diversity across the prison population and the different drinking patterns that may exist. They also believed that the language used could be modified and simplified to be more acceptable for the remand population. In addition, the need to create safe spaces, fostering inclusivity and adaptation to meet the individual needs of the remand population, was identified. An awareness of the vulnerability within the prison environment for those on remand and being cognisant of this in working to implement new interventions within the estate was identified as being required. The relevance of the lived experience and that those with it can have a role to play alongside the wider network of services could shape the implementation of the APPRAISE intervention.

Finally, all participants interviewed appeared to be enthusiastic about facilitating future work in the area and building upon the delivery of the ABI in the APPRAISE study. Researchers were able to introduce the study effectively, develop rapport with the prison staff and maintain flexibility to ensure minimal disruption to the setting, even during the unprecedented COVID-19 pandemic. The fact that participants relayed that the services and treatment options currently available to remand prisoners were extremely limited further highlights the need to broaden the existing services in this area and work towards a future trial of the APPRAISE intervention to determine the APPRAISE intervention effectiveness.

This, therefore, further confirmed the importance of the remand prison setting as a suitable location for delivering the APPRAISE intervention and the importance of the wider network that could support implementation.

Although this did not appear to have a negative impact on delivery, in a future roll-out it may be prudent to condense training sessions and the accompanying materials to ensure that it remains acceptable and feasible to a larger cohort of interventionists.

Chapter 7 Economic study

Were the APPRAISE intervention to demonstrate effectiveness, the next barrier to widespread adoption would be cost efficiency. While stated policy criteria exist in the UK health sector to guide the design of health research,⁷⁶ the APPRAISE intervention encompasses a wider stakeholder group, including the prison and wider criminal justice system (CJS) stakeholders, and the prisoners themselves. These groups additionally require convincing of value from their own perspective to help alleviate implementation barriers. Hence, a standard health and social care perspective alone may not provide sufficient evidence needed by all parties in their independent decision-making, and a broader taxonomy of outcomes is necessary. Conducting any analysis in a prison setting adds challenges to data collection and analysis, hence the feasibility of collecting data to inform data analysis requires exploring and refining prior to any significant investment in a larger definitive trial. To that end, we present the findings of the APPRAISE economic substudy, including a micro-costing methodology protocol, and a revision of the Economic Form 90 for a UK context,⁶⁰ and some provisional scoping of routine data sources to support future cost-effectiveness analyses (CEA) in this area. We anticipate this work to be valuable as we believe this to be the first of its kind for an extended ABI in this setting and that the methods we have discussed will likely be relevant and transferable across other ABI settings.

Full details of the wider trial methodology are given elsewhere in this report and in our published study protocol.¹ This chapter focuses exclusively on the economic evaluation pilot study. Specifically, we discuss efforts undertaken to address the study Objectives 1d, 1e and 1f.

- 1d. Can we collect economic data needed for a future definitive RCT?
- 1e. Can we access recidivism data from the PNC databases for trial participants?
- 1f. Can we access health data from routine NHS data sources for trial participants?

To aid UK policy relevance, the base-case costing perspective was based on the NICE recommended NHS and personal social services (PSS) perspective,⁷⁶ with the addition of criminal justice and prison service costs to future support prison sector buy-in. The economic study focuses on the opportunity cost of intervention implementation from these perspectives. Further planned analyses aimed to widen this to a whole societal perspective with the further inclusion of employment, welfare, and third-sector addiction services, as well as including future cost savings from the NHS and PSS perspectives. For this pilot feasibility study, the base year for all costs was set to the 2020–1 financial year, selected at the latest financial year from which any prisoner provided data on balance internal and external validity. Discounting and inflation adjustments were not necessary as the time horizon for analysis was under 1 year, and all price estimates related directly to the base year. A Consolidated Health Economic Evaluation Reporting Standards (CHEERS) checklist indicating the location of key economic principles and methods can be found in [Appendix 12](#).

Methods

The economic evaluation pilot study aimed to evaluate the feasibility of collecting economic data to inform design of a potential future RCT of the APPRAISE intervention in a UK prison setting. This had three principal components:

1. development and pilot testing of an activity-based costing (ABC) exercise adapting recommendations from Drummond *et al.*,⁷⁷ which aimed to estimate the resources used to deliver the ABI programme, and equivalent usual care, and associated financial cost of these
2. pilot testing of the collection of self-report data on prisoners' health care, social care, criminal justice, welfare payments, employment, and third-sector addiction-service utilisation at baseline and at 6- and 12-month follow-ups using an adapted Economic Form 90⁶⁰

3. an examination of the feasibility of accessing recidivism data from the PNC databases, and health data from routine NHS data sources for research purposes within a prisoner population.

Of note are overlaps between potential data items obtainable via components 2 and 3. We sought to compare data quality to both validate our survey data instrument and identify the preferred data-collection approach to provide an informed balance of data quality and participant self-report burden. We initially planned to publish a health economic analysis plan prior to analysis; however, disruptions due to the pandemic necessitated a number of ad hoc adaptations.

The activity-based costing exercise

The proposed ABC exercise was based on the recommendations by Drummond *et al.*,⁷⁷ and consisted of four principal steps:

1. Step 1 – a taxonomy of intervention activities was created. A 1-day consultation exercise was held, with both the staff delivering the intervention and the research staff involved in the APPRAISE pilot study, to identify all of the resources that would be required in the implementation of the APPRAISE intervention, at scale, in a future hypothetical roll-out, as opposed to those only required for research purposes. A detailed narrative was constructed, for each study site, which documented the key components of the intervention delivery and the process. The intervention activities were listed in the sequence performed, in relation to each participant, and the resources that were used for each activity.
2. Step 2 – resources used for each activity were measured. The staff delivering the APPRAISE interventions in each study site were asked to note the duration of each intervention activity and the necessary materials and commodities that were required before, during, and after each stage of intervention. Examples of these resources measured included: the time taken to locate participants using their preferred method of contact; the preparation and printing of the intervention materials; and the actual time taken for the delivery of the intervention sessions. Security protocols and procedures prohibited the use of use electronic equipment for recording this information. In order to comply, handwritten research diaries were instead compiled. These were either transcribed verbatim or were scanned and shared retrospectively within the economic research team.
3. Step 3 – identification of unit cost. Where possible, unit costs for each resource use item were identified based on the study invoices and targeted (non-systematic) literature searches or sourced from expert knowledge within the team. Priority was given to the estimates considered to be best fit to a hypothetical future roll out (at scale) where multiple estimates were located.
4. Step 4 – calculation of costs. The resources identified in Step 2 were combined with the unit costs from Step 3 to generate estimations of the costs of each activity.

These calculations are not intended as an accurate estimation of costs of a hypothetical future roll-out, but rather as a demonstration of how these might be calculated in a future trial. Of particular note is that differences in prison procedures would be expected to yield differences in durations and costs of each type of activity, and that a larger study would need to account for this. To this end, separate outputs from Steps 1 and 2 were generated and compared for each site.

Adaptation of the Economic Form 90

The items of the Economic Form 90 were systematically examined in consultation with a UK economist and experts in the prison and addiction-services fields to identify UK equivalents for each entry and anything missing, and remove anything not relevant in a UK context. Questions were additionally rephrased for clarity to a UK reader and PPI input was solicited.

Unit costs for each item were identified by a non-systematic literature search, alongside those from the ABC exercise, with the same preference given to national-level unit costs to maximise relevance to a hypothetical future roll out (at scale).

Examination of Police National Computer and National Health Service data sources

Permission to access PNC and PHS data sets was requested from prison authorities. Upon receipt these were to be examined for data quality and compared to results of the Form 90 survey to cross validate. Results from this comparison would inform the optimal balance of data items to be obtained from each source in a future definitive trial. Consideration was to be given to any missing items, and if these were estimable via other sources, such as literature or modelling.

Results

Disruption due to coronavirus disease discovered in 2019 pandemic

Follow-up data collection for multiple parts of the study became disrupted as researchers were unable to access prison services due to revised safety protocols. This had the following notable impacts on the economic substudy:

- It was not possible collect the anticipated number of Economic Form 90 surveys, preventing analysis and validation efforts. However, the adaptation process was completed prior to onset of the pandemic, and scoping of potential sources for price weights was possible remotely.
- Prison staff experienced sharp increases in workloads which they needed to prioritise over research efforts, leading to lower participation rates and possibly different duration of intervention sessions than might be anticipated in a live roll-out.
- Time spent locating prisoners for post-liberation intervention sessions increased due to additional complexities of doing so remotely that may not reflect future methods.

Activity-based costing exercise

Row headings on [Tables 35–37](#) display the items identified in Stage 1 of the ABC process via day consultation. Tables were then populated in Steps 2–4 detailing all activities, staff costs and the unit cost sources. Separate estimates are provided for the Scotland and England study sites to account for differences in services and local salaries of staff providing services.

[Table 35](#) breaks down staff costs associated with delivering the intervention. The intervention service providers (CGL) at each study site were asked to provide a typical salary cost for the different roles delivering the APPRAISE intervention so that the intervention staff salary cost could be estimated.

TABLE 35 APPRAISE intervention staff delivery costs

APPRAISE activities	England cost	Scotland cost
Session 1 Approximately 45 minutes with a co-ordinator plus an additional participant-locating cost	Co-ordinator annual salary cost = £21,630–£21,812 45 minutes	Co-ordinator annual salary cost = £23,218.59–£26,983.44 45 minutes only
Session 2 Approximately 20 minutes with an intervention support worker plus an additional participant-locating cost	Support worker annual salary cost = £18,213–£19,847 20 minutes	Support worker annual salary cost = £18,895.07–£19,744.94 20 minutes
Session 3 Approximately 20 minutes with an intervention support worker plus an additional participant-locating cost	Support worker annual salary cost = £18,213–£19,847 20 minutes	Support worker annual salary cost = £18,895.07–£19,744.94 20 minutes only
Session 4 Approximately 20 minutes with an intervention support worker plus an additional participant-locating cost	Support worker annual salary cost = £18,213–£19,847 20 minutes	Support worker annual salary cost = £18,895.07–£19,744.94 20 minutes

Values obtained for these, correct at the time of the pilot study undertaking (2020), are included in [Table 35](#). Participant locating time proved more difficult to determine and hence it was more difficult to provide a meaningful estimation of the staff cost associated with it. In the pilot study these tended to be lower-paid staff, such as support workers or administrative staff, attempting to make contact with a participant. The time taken to schedule a follow-up intervention session or to successfully make contact to start the intervention session remotely (by telephone) often varied significantly. Some individuals could be reached quickly (within 10–15 minutes) or it could be ascertained quickly that the listed provided contact methods could no longer be used, for example their phone number was no longer in use or e-mails to their address were bouncing back etc. This was in direct contrast to those who were unable to be reached after frequent attempts, by different methods of contact, or those who would reply once and then would stop replying. Successful completion of follow-up interventions was low, even with large amounts of time spent attempting to make contact. However, it was difficult to establish how much of an effect the COVID-19 pandemic had in this lack of successful contact; thus it needs to be an area to explore further in a future cost analysis, rather than attaching a cost estimate to the participant-contacting time with limited insight.

[Table 36](#) documents the one-off and the ongoing staff training costs that would be associated with implementing the APPRAISE intervention in practice. Initial APPRAISE interventionist staff training consisted of a one-time in-depth training session lasting approximately 2 hours. This was followed by refresher courses every 6 months typically lasting 20–30 minutes, and weekly 5–10-minute debriefs to allow the intervention staff to catch up with the research staff around their progress and to provide updates and to discuss any concerns or queries. However, disentangling research-only costs from those that would be present in a real-world setting may require separate sampling, and/or may be better handled by attributing varying proportions of activities to each in sensitivity analysis in a next-stage RCT. These staff costs, associated with the intervention delivery training, would vary dependent on the numbers of the different staff roles that were required in each location, and across the two study sites.

[Table 37](#) presents other resource-utilisation costs required when delivering the APPRAISE intervention. As several of the questionnaires form part of the screening and recruitment process, and prison security processes and procedures prevented all non-paper-based data-collection options, printing costs totalling £6.45 per participant were unavoidable.

Depending on which session was being undertaken a plethora of different methods may be used to contact participants for follow-up intervention sessions. These included the cost of using a telephone or text messages, sending written letters or using the internet for social media or WhatsApp. In the pilot study, all of the required printing was conducted and costed by the research staff at the University

TABLE 36 APPRAISE ABI staff training costs

Training	England cost	Scotland cost
Initial training Generally, 2 hours and dependent on number of staff trained	Co-ordinator annual salary cost = £21,630–£21,812 Support worker annual salary cost = £18,213–£19,847	Co-ordinator annual salary cost = £23,218.59–£26,983.44 Support worker annual salary cost = £18,895.07–£19,744.94
Refresher training Generally, 30 minutes and dependent on number of staff trained	Co-ordinator annual salary cost = £21,630–£21,812 Support worker annual salary cost = £18,213–£19,847	Co-ordinator annual salary cost = £23,218.59–£26,983.44 Support worker annual salary cost = £18,895.07–£19,744.94
Ongoing weekly debrief in prison Generally, 10 minutes and dependent on number of staff	Co-ordinator annual salary cost = £21,630 - £21,812 Support worker annual salary cost = £18,213 - £19,847	Co-ordinator annual salary cost = £23,218.59–£26,983.44 Support worker annual salary cost = £18,895.07–£19,744.94

TABLE 37 Other resource costs associated with the APPRAISE ABI

Resource	Resource breakdown	Price per item
Printing	APPRAISE information booklet	£0.42
	Consent form	£0.57
	Participant contact details form	£0.12
	Baseline questionnaire (with AUDIT-C)	£2.43
	APPRAISE alcohol unit postcard	£0.19
	APPRAISE follow-up intervention booklet	£2.60
Contacting participants	Letters	Printing letter – £0.12
		Envelope – £0.11
		2nd-class postage – £0.66
	Telephone calls	£0.25 per minute
	Text messages	£0.10 per message
	Internet (WhatsApp or social media)	£0.10/MB

of Edinburgh, which included the follow-up letters. Any further attempts to contact participants or to conduct phone-based follow-up sessions were conducted using a researcher's phone, but with a new pay-as-you-go sim card. The sim card was free to obtain, but the phone costs have been quoted from the listed provider's pay-as-you-go costs at the time of the study. Costs of these in future studies or eventual roll-outs may vary, and users of these results may wish to substitute their own local estimates.

Factors, such as rooms in prisons needed to undertake sessions were provided free and so were not included in the pilot ABC exercise. While these are anticipated to have opportunity costs associated with them, costs at a given prison will vary substantially and arguably only incur an opportunity cost if the room was needed for another purpose. Financial values for these are challenging to estimate reliably and may be misleading if they appear overly specific. Nevertheless, their need is noted in a more qualitative sense, with recommendation that future work or roll-outs include these costs and use local estimates for decision purposes where possible.

Adaptation of Economic Form 90

The adapted version of the Economic Form 90 survey developed as part of this study can be found in [Appendix 9](#). While the COVID pandemic prevented data collection, it was still possible to identify candidate sources of price weights for items included in the survey. These are listed in [Table 38](#).

Note that this table provides potential price weight sources as opposed to specific price weights and values for a given base year. This is to provide a guide for future survey users to source weights appropriate to their analysis which may not have been published at time of writing.

While the specific content of each may change in a given year of publication, standard UK price weight sources, such as the Personal Social Services Research Unit (PSSRU) unit costs of health and social care series⁷⁸ and the NHS reference cost report series⁷⁹ are likely to be suitable for the majority if not all of the health and social care aspects of the survey. PSSRU additionally has some content for substance misuse including alcohol services, albeit with more limited content.

A small number of items were identified as being likely to require bespoke or local pricing to best reflect the differences in regional availability of services and prices, namely those relating to housing costs and

TABLE 38 Candidate unit cost sources for adapted Economic Form 90 questionnaire

Item	Unit	Unit cost source	Notes
Healthcare costs			
GP consultations	Per consultation	PSSRU unit costs ⁷⁸	
Practice nurse consultations	Per consultation	PSSRU unit costs	
Psychiatrist	Per consultation	PSSRU unit costs	
Psychiatric nurse	Per consultation	PSSRU unit costs	
Inpatient admissions	Per day	NHS reference costs ⁷⁹ /ISD cost book ⁸⁰	
Emergency dept. visits	Per visit	NHS reference costs/ISD cost book	
Social care costs			
Social worker in person	Per consultation	PSSRU unit costs	
Social worker phone	Per consultation	PSSRU unit costs	
Addiction services			
Alcohol services	Per consultation?	PSSRU unit costs	
Drug services	Per consultation?	PSSRU unit costs	
Third sector	Per consultation?	Recommend local pricing	Costs will depend on organisation in question. Recommend bespoke costing, such as private communications.
Criminal justice system costs			
Arrests	Per event	See comments in text below	
Custody	Per night	See comments in text below	
Prison	Per night	See comments in text below	
Hospital security detail	Per night	See comments in text below	
Welfare			
Stayed at hostel	Per night	Recommend local pricing	
Stayed at B&B	Per night	Recommend local pricing	
Residential care/supported accommodation/shelter	Per night	Recommend local pricing	
Housing benefit	Total GBP paid	N/A	Recorded in GBP. No price weight required.
Universal credit	Total GBP paid	N/A	Recorded in GBP. No price weight required.
ISD, Information Services Division.			

third-sector alcohol services. While a similar argument for local costing could be made for health and social care aspects, housing and third-sector alcohol services are likely to be part of costing perspectives falling out with a national programme or organisation (such as the NHS), but rather on the individual or a programme regional budget.

Price weights for criminal justice aspects may warrant some minor rewording of the survey to better match the granularity of the available sources in the literature. For example, Ramponi *et al.* break down

costs by form of crime (e.g. assault/robbery/criminal damage) rather than as components of system use (e.g. arrest/custody/prison).⁸¹ In the absence of a recurring national publication for such items, researchers may wish to identify price weights prior to survey issue and adapt the questionnaire to suit.

Response rates of Economic Form 90

While the adapted Economic Form 90 was issued at baseline, its follow-up issues were hampered by the pandemic, as shown by [Table 39](#).

The baseline data completeness and self-reported responses are shown in [Table 40](#) (health and social care), [Table 41](#) (living arrangements), [Table 42](#) (employment status) and [Table 43](#) (criminal justice service utilisation).

TABLE 39 Number of adapted Economic Form 90 issued by time point

Site	Baseline			6 months			12 months		
	Intervention	Control	Total	Intervention	Control	Total	Intervention	Control	Total
England	46	44	90	0	0	0	2	2	4
Scotland	22	20	42	2	1	3	7	7	14
Total	68	64	132	2	1	3	9	9	18

TABLE 40 Completeness and self-reported responses to health and social care aspects of adapted Economic Form 90

Item		Intervention	Control	Total		
Total number of prisoners		68	100.0%	64	100.0%	132 100.0%
Since (date), have you been to hospital?	Missing/unknown: N, %	0	0.0%	0	0.0%	0 0.0%
	No: N, %	47	69.1%	44	68.8%	91 68.9%
	Yes: N, %	21	30.9%	20	31.3%	41 31.1%
If Yes, how many times?	Missing/unknown: N, %	0	0.0%	0	0.0%	0 0.0%
	Answered: N, %	21	100.0%	20	100.0%	41 100.0%
	Visits: mean, SD	2.4	1.70	1.7	0.90	2.1 1.40
If Yes, did you stay overnight?	Missing/unknown: N, %	0	0.0%	0	0.0%	0 0.0%
	No: N, %	11	16.2%	8	12.5%	19 14.4%
	Yes: N, %	10	14.7%	12	18.8%	22 16.7%
If Yes, how many nights in total?	Missing/unknown: N, %	1	4.8%	0	0.0%	1 2.4%
	Answered: N, %	9	42.9%	12	60.0%	21 51.2%
	Nights: mean, SD	10.7	9.80	4.3	7.70	7 9.00
Are you registered with a GP?	Missing/unknown: N, %	1	1.5%	1	1.6%	2 1.5%
	No: N, %	11	16.2%	9	14.1%	20 15.2%
	Yes: N, %	56	82.4%	54	84.4%	110 83.3%
If registered, have you visited a GP?	Missing/unknown: N, %	1	1.8%	0	0.0%	1 0.9%

continued

TABLE 40 Completeness and self-reported responses to health and social care aspects of adapted Economic Form 90 (continued)

Item		Intervention		Control		Total	
	No: N, %	19	33.9%	23	42.6%	42	38.2%
	Yes: N, %	36	64.3%	31	57.4%	67	60.9%
If Yes, how many times?	Missing/unknown: N, %	1	2.8%	0	0.0%	1	-1.5%
	Answered: N, %	35	97.2%	31	100%	66	98.5%
	Visits: mean, SD	3.1	3.00	2.9	2.60	3	2.80
If registered, have you visited a practice nurse?	Missing/unknown: N, %	1	1.8%	0	0.0%	1	0.9%
	No: N, %	40	71.4%	45	83.3%	85	77.3%
	Yes: N, %	15	26.8%	9	16.7%	24	21.8%
If Yes, how many times?	missing/unknown: n, %	3	20.0%	1	11.1%	4	16.7%
	Answered: N, %	12	80.0%	8	88.9%	20	83.3%
	Visits: mean, SD	2.3	1.60	2.4	1.80	2.3	1.60
	Missing/unknown: N, %	1	1.5%	1	1.6%	2	1.5%
Since (date), have you been visited by a community psychiatric nurse?	Missing/unknown: N, %	0	0.0%	1	1.6%	1	0.8%
	No: N, %	65	95.6%	56	87.5%	121	91.7%
	Yes: N, %	3	4.4%	7	10.9%	10	7.6%
If Yes, how many times?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	Answered: N, %	3	100.0%	7	100.0%	10	100.0%
	Visits: mean, SD	1.7	0.60	2.4	2.20	2.2	1.90
Since (date), have you been to any addiction services?	Missing/unknown: N, %	2	2.9%	1	1.6%	3	2.3%
	No: N, %	48	70.6%	45	70.3%	93	70.5%
	Yes: N, %	18	26.5%	18	28.1%	36	27.3%
If Yes, did you stay overnight for any visit?	Missing/unknown: N, %	2	11.1%	4	22.2%	6	16.7%
	No: N, %	16	88.9%	14	77.8%	30	83.3%
	Yes: N, %	0	0.0%	0	0.0%	0	0.0%
Since (date), have you been to A&E?	Missing/unknown: N, %	2	2.9%	0	0.0%	2	1.5%
	No: N, %	31	45.6%	35	54.7%	66	50.0%
	Yes: N, %	35	51.5%	29	45.3%	64	48.5%
If Yes, how many times?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	Answered: N, %	35	100.0%	29	100.0%	64	100.0%
	Visits: mean, SD	1.7	1.4	1.5	0.9	1.6	1.2
If Yes, were any of these times when you were in prison?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	No: N, %	27	77.1%	20	69.0%	47	73.4%

TABLE 40 Completeness and self-reported responses to health and social care aspects of adapted Economic Form 90 (continued)

Item		Intervention		Control		Total	
If Yes, how many times?	Yes: N, %	8	22.9%	9	31.0%	17	26.6%
	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	Answered: N, %	8	100.0%	9	100.0%	17	100.0%
If Yes, did you ever use an emergency ambulance?	Visits: mean, SD	2	1.8	1.1	0.3	1.5	1.3
	Missing/unknown: N, %	0	0.0%	1	3.4%	1	1.6%
	No: N, %	16	45.7%	12	41.4%	28	43.8%
If Yes, how many times?	Yes: N, %	19	54.3%	16	55.2%	35	54.7%
	Missing/unknown: N, %	1	5.3%	0	0.0%	1	2.9%
	Answered: N, %	18	94.7%	16	100%	34	97.1%
Since (date), have you been working with a social worker?	Trips: mean, SD	1.6	1.1	1.2	0.4	1.4	0.9
	Missing/unknown: N, %	0	0.0%	2	3.1%	2	1.5%
	No: N, %	51	75.0%	47	73.4%	98	74.2%
If Yes, how many times have you been in contact?	Yes: N, %	17	25.0%	15	23.4%	32	24.2%
	Missing/unknown: N, %	2	11.8%	0	0.0%	2	6.3%
	Answered: N, %	15	88.2%	15	100.0%	30	93.8%
If Yes, how many times were by phone?	Contacts: mean, SD	14.7	12.9	10.5	8	12.6	10.8
	Missing/unknown: N, %	4	24%	1	7%	5	16%
	Answered: N, %	13	76%	14	93%	27	84%
If Yes, how many times were in person?	Contacts: mean, SD	3.1	4	4.8	8	4	6.3
	Missing/unknown: N, %	0	0%	0	0%	0	0%
	Answered: N, %	17	100%	15	100%	32	100%
If Yes, how many times were while you were in your home?	Contacts: mean, SD	11.9	12.7	7.4	7.1	9.8	10.5
	Missing/unknown: N, %	4	24%	1	7%	5	16%
	Answered: N, %	13	76%	14	93%	27	84%
Since (date), have you tried to access help from any other support services?	Contacts: mean, SD	10.1	15.1	0.9	1.1	5.3	11.3
	Missing/unknown: N, %	0	0.0%	2	3.1%	2	1.5%
	No: N, %	52	76.5%	45	70.3%	97	73.5%
If Yes, how many times?	Yes: N, %	16	23.5%	17	26.6%	33	25.0%
	Missing/unknown: N, %	3	18.8%	6	35.3%	9	27.3%
	Answered: N, %	13	81.3%	11	64.7%	24	72.7%
	Contacts: mean, SD	4.8	5.7	9.7	8.7	7.1	7.5

TABLE 41 Completeness and self-reported responses to living arrangements aspects of adapted Economic Form 90

Item		Intervention		Control		Total	
What are your current living arrangements?							
Living alone	Unchecked	50	73.5%	44	68.8%	94	71.2%
	Checked	18	26.5%	20	31.3%	38	28.8%
Living with a partner	Unchecked	55	80.9%	51	79.7%	106	80.3%
	Checked	13	19.1%	13	20.3%	26	19.7%
Living with other family members	Unchecked	47	69.1%	42	65.6%	89	67.4%
	Checked	21	30.9%	22	34.4%	43	32.6%
Living with friends	Unchecked	66	97.1%	63	98.4%	129	97.7%
	Checked	2	2.9%	1	1.6%	3	2.3%
Owned or rented property (you must be the tenant/joint tenant)	Unchecked	64	94.1%	61	95.3%	125	94.7%
	Checked	4	5.9%	3	4.7%	7	5.3%
Council property/housing association (you must be the tenant/joint tenant)	Unchecked	61	89.7%	58	90.6%	119	90.2%
	Checked	7	10.3%	6	9.4%	13	9.8%
Hostel	Unchecked	68	100.0%	63	98.4%	131	99.2%
	Checked	0	0.0%	1	1.6%	1	0.8%
Bed and breakfast	Unchecked	66	97.1%	63	98.4%	129	97.7%
	Checked	2	2.9%	1	1.6%	3	2.3%
Residential care/shelter/supported accommodation	Unchecked	61	89.7%	64	100.0%	125	94.7%
	Checked	7	10.3%	0	0.0%	7	5.3%
Other (including rough sleeping, sofa surfing)	Unchecked	59	86.8%	54	84.4%	113	85.6%
	Checked	9	13.2%	10	15.6%	19	14.4%
Since (date), have you been homeless at any point?	Missing/unknown: N, %	3	4.4%	0	0.0%	3	2.3%
	No: N, %	44	64.7%	44	68.8%	88	66.7%
	Yes: N, %	21	30.9%	20	31.3%	41	31.1%
If Yes, how many nights?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	Answered: N, %	21	100.0%	20	100.0%	41	100.0%
	Nights: mean, SD	123.4	109	97.5	77	110.8	94.5
Since (date), have you stayed in any of the following?							
Hostel	Unchecked	64	94.1%	62	96.9%	126	95.5%
	Checked	4	5.9%	2	3.1%	6	4.5%
If Yes, how many times?	Missing/unknown: N, %	0	0.0%	1	50.0%	1	16.7%

TABLE 41 Completeness and self-reported responses to living arrangements aspects of adapted Economic Form 90 (continued)

Item		Intervention		Control		Total	
	Answered: N, %	4	100.0%	1	50.0%	5	83.3%
	Nights: mean, SD	28.3	14.8	3	--	23.2	17.1
Bed and breakfast	Unchecked	62	91.2%	61	95.3%	123	93.2%
	Checked	6	8.8%	3	4.7%	9	6.8%
If Yes, how many times?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	Answered: N, %	6	100.0%	3	100.0%	9	100.0%
	Nights: mean, SD	40	70.2	58	68.6	46	65.8
Residential care/shelter/ supported accommodation	Unchecked	59	86.8%	62	96.9%	121	91.7%
	Checked	9	13.2%	2	3.1%	11	8.3%
If Yes, how many times?	Missing/unknown: N, %	0	0.0%	1	50.0%	1	9.1%
	Answered: N, %	9	100.0%	1	50.0%	10	90.9%
	Nights: mean, SD	81.3	73.1	60.0	--	79.2	69.2

TABLE 42 Completeness and self-reported responses to employment and benefits aspects of adapted Economic Form 90

Item		Intervention		Control		Total	
Are you currently employed?	Missing/unknown: N, %	0	0.0%	1	1.6%	1	0.8%
	No: N, %	52	76.5%	52	81.3%	104	78.8%
	Yes: N, %	16	23.5%	11	17.2%	27	20.5%
If No, have you been employed at any point since (date)?	Missing/unknown: N, %	0	0.0%	1	1.9%	1	1.0%
	No: N, %	43	82.7%	43	82.7%	86	82.7%
	Yes: N, %	9	17.3%	8	15.4%	17	16.3%
If Yes, how many jobs have you had during this time?	Missing/unknown: N, %	3	18.8%	1	9.1%	4	14.8%
	One	13	81.3%	10	90.9%	23	85.2%
If Yes, how many weeks have you been employed between (date) and now?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	Answered: N, %	16	100.0%	11	100.0%	27	100.0%
	Weeks: mean, SD	22.5	7.2	31.4	50.1	26.1	31.9
Do you currently receive any money from a public or government source?	Missing/unknown: N, %	1	1.5%	1	1.6%	2	1.5%
	No: N, %	17	25.0%	23	35.9%	40	30.3%
	Yes: N, %	50	73.5%	40	62.5%	90	68.2%

continued

TABLE 42 Completeness and self-reported responses to employment and benefits aspects of adapted Economic Form 90 (continued)

Item		Intervention		Control		Total	
If Yes, approximately how much do you receive in total per week?	Missing/unknown: N, %	5	10.0%	1	2.5%	6	6.7%
	Answered: N, %	45	90.0%	39	97.5%	84	93.3%
	GBP: mean, SD	114.7	93.2	117.1	75.7	115.8	85.0
If No, have you been employed at any point since (date)?	Missing/unknown: N, %	0	0.0%	1	1.9%	1	1.0%
	No: N, %	43	82.7%	43	82.7%	86	82.7%
	Yes: N, %	9	17.3%	8	15.4%	17	16.3%
If Yes, how many jobs have you had during this time?	Missing/unknown: N, %	3	18.8%	1	9.1%	4	14.8%
	One	13	81.3%	10	90.9%	23	85.2%
If Yes, how many weeks have you been employed between (date) and now?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	Answered: N, %	16	100.0%	11	100.0%	27	100.0%
	Weeks: mean, SD	22.5	7.2	31.4	50.1	26.1	31.9

TABLE 43 Completeness and self-reported responses to criminal justice aspects of adapted Economic Form 90

Item		Intervention		Control		Total	
Since (date), have you been arrested?	Missing/unknown: N, %	0	0	0	0	0	0
	No: N, %	0	0	1	1.6%	1	0.8%
	Yes: N, %	68	1	63	98.4%	131	99.2%
If Yes, how many times?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	Answered: N, %	68	100%	63	100.0%	131	100.0%
	Times: mean, SD	2.4	3.6	2.7	2.4	2.5	2.1
Since (date), have you spent at least one night in custody?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	No: N, %	1	1.5%	0	0.0%	1	0.8%
	Yes: N, %	67	98.5%	64	100.0%	131	99.2%
If Yes, was this police custody, prison, or both?	Missing/unknown: N, %	1	1.5%	0	0.0%	1	0.8%
	Police custody: N, %	0	0.0%	0	0.0%	0	0.0%
	Prison: N, %	2	2.9%	2	3.1%	4	3.0%
	Both: N, %	65	95.6%	62	96.9%	127	96.2%
If Yes, how many nights spent on remand?	Missing/unknown: N, %	1	1.5%	1	1.6%	2	1.5%

TABLE 43 Completeness and self-reported responses to criminal justice aspects of adapted Economic Form 90 (*continued*)

Item	Intervention	Control	Total				
	Answered: N, %	66	98.5%	63	98.4%	129	98.5%
	Nights: mean, SD	15.3	24.2	24	42	19.5	34.2
If Yes, how many nights spent in police custody?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	Answered: N, %	67	100.0%	64	100.0%	131	100.0%
	Nights: mean, SD	18.6	24.2	27	42.2	22.7	34.4
Since (date), have you been in court?	Missing/unknown: N, %	0	0.0%	0	0.0%	0	0.0%
	No: N, %	1	1.5%	0	0.0%	1	0.8%
	Yes: N, %	67	98.5%	64	100.0%	131	99.2%
If Yes, how many times?	Missing/unknown: N, %	0	0.0%	1	1.6%	1	0.8%
	Answered: N, %	67	100.0%	63	98.4%	130	99.2%
	Times: mean, SD	2.3	2.2	2.6	2.6	2.5	2.4

Completeness of the survey at baseline was generally very good, with only three questions having over five missing entries.

Feasibility of accessing Police National Computer and National Health Service service use data

Although it is possible that COVID-19 played a role in our inability to obtain these data, we do not think that the pandemic was solely responsible. Two key issues are at play: prisoner confidentiality and data systems. We found that gaining approval to obtain the PNC or NHS identifiers was difficult to impossible due to a combination of prison and data-protection concerns. Although these concerns may fade as more evidence of the benefits of the APPRAISE intervention mounts, we conclude that it is very likely that these concerns will still exist for a definitive trial, and possibly be even more salient for some stakeholders in that context. Exacerbating the confidentiality issues were data-system issues that prevented data access on a large scale. Although this issue depends on the jurisdiction, in each of our two jurisdictions (Scotland and England) at least one of the data sources required case-by-case data extraction at some step of the process. We could not extract either PNC or NHS data on a large scale in either England or Scotland, making the use of such data prohibitively costly for a definitive RCT.

Discussion

We aimed to assess the feasibility of collecting economic data to inform the design of a potential future RCT of APPRAISE in a UK prison setting. We did not seek to provide a definitive measurement of cost, but rather inform methods by which they could be measured in a future trial. This has been achieved with mixed results. While many of the issues faced were a result of the COVID-19 pandemic, it is important to separate ambiguity of findings due to extenuating circumstances from findings of issues in data collection. Data obtained despite such circumstances are still likely to be viable without them, if not more so, and we were able to document the stages of an ABC exercise and provide discussion around how to assess the costs associated with implementing the APPRAISE intervention. We believe this to be one of the first pieces of work in this area and the methods that we have presented can be applied more broadly to determine the costs associated with implementing a similar intervention in practice.

While not the aim of the study, we have also provided early estimates of some of the costs associated with the implementation of the APPRAISE intervention, in two different prison systems. There were important differences between the two different sites, primarily around the differing roles and numbers of members of prison and intervention staff involved, the location of delivery and availability of rooms and the level of access to prisoner data at the time of data collection. Such differences are inevitable given the heterogeneity of prison regimes in general in terms of local service provision, facilities, staff and the management of each prison site. Nevertheless, the ABC process itself as described could be reapplied to new contexts, and elements of the activities observed, such as necessary paperwork and durations of phone sessions would reasonably be expected to be common to other settings.

There were challenges in separating research-specific costs from those which might reasonably be expected in a practice. For example, in the England site the researcher recruiting the participants to the study was unable to access the prison database to identify further potential participants and their current location. This meant that the process of identification was more cumbersome at times and it may have taken longer to locate eligible participants, which would likely increase the staff cost associated with the intervention. Assuming that, in a real-world setting, intervention staff or prison staff would have access to such databases and would not need to spend the extra time locating individuals for their APPRAISE intervention, the time taken may not accurately reflect that needed in a live roll-out. In relation to this, prison staff costs were intentionally excluded in the staff costs listed in [Table 35](#) because, while they were involved and helped with the process of locating participants within the research trial setting, it is currently unclear what role they would play in practice. Such details will need to reflect the final design of an intervention or will need to be explored in future work, though we have no reason to believe the ABC approach could not be used to do so.

The COVID-19 pandemic is also likely to have influenced the follow-up and the retention of study participants. This issue is common to the study as a whole rather than to economic assessment in particular, and is discussed in broader terms in other chapters of this report. Specific to the economic evaluation pilot, substantial time was spent by intervention staff trying to locate participants for their follow-up interventions and often being unable to access the information they required or finding that participant court dates had been moved or postponed due to COVID-19. More time spent locating participants for the post-liberation intervention sessions has direct implications on the cost of the APPRAISE intervention, as it obviously increases the staff costs associated with the intervention. Disentangling the direct impact of the COVID-19 pandemic on locating participants remotely from the normal challenges of this population was not possible in this pilot study. Follow-up in the CJS is often very complicated and it is frequently challenging to retain participants within a study due to a number of logistical reasons, mostly being unable to re-establish contact with them.⁸² Consequently, this is something to consider when planning for a future RCT with the APPRAISE intervention and ensuring that the lower-paid staff, such as administrative or support workers, could be responsible for locating participants to reduce overall costs, if deemed necessary.

The timing of the first COVID lockdowns in 2020 had a significant effect on the participant recruitment and the study's data collection after both prison sites went into an immediate lockdown. Hence limited data could be collected for the remaining duration of the pilot, post April 2020. As a result, it also became difficult to work with prison staff due to their sharply increased workloads and new unforeseen pressures. The majority of any further interactions were conducted virtually (via telephone or e-mail), rather than the face-to-face meetings and follow-ups as planned. This is extremely likely to have had an effect on the information we were able to collect around the cost estimates. For some specific items, such as the cost of space in the prison, we were unable to obtain meaningful estimates. Context is critical for this factor, as costs of this are expected to vary substantially between prisons. It may be sufficient to note the duration for which a private room is required, and allow users of this information to apply their own local estimates where necessary.

We were able to adapt the Economic Form 90 to a UK context and provide a list of candidate price weight sources for future use. We were also able to issue the survey to 132 prisoners at baseline with very high completeness, though follow-up time points were heavily disrupted by the pandemic. Of note, the baseline survey was researcher-administrated. It is not possible to know how well the survey would have been completed in follow-up time points without the confounding of the pandemic; however, the completeness at baseline at least implies the form was well accepted and understood. Completeness included questions relating to criminal justice activity, suggesting that any adaptation of the questionnaire to allow for more bespoke or specific price weights to be applied is likely to be viable following a similar format.

Disruptions to data collection prohibited our ability to validate the survey against routine data sources. The matter may be rendered a moot point as PNC and NHS data proved too difficult to obtain at large scale for this population, thus suggesting they should not be included as part of a definitive RCT. Some element of prisoner self-report was always anticipated to be unavoidable; however, the loss of routine data as a potential alternative source places increased importance on validation of the revised Economic Form 90. While the original Economic Form 90 has been validated for a US alcohol treatment population,⁶⁰ larger-scale future trials would be advised to build in a pilot stage to validate and potentially refine our adapted UK version prior to full-scale administration.

Chapter 8 Prison survey

In this chapter, we present the methods undertaken and the results obtained from conducting a survey to further understand the alcohol services provided in male remand prisons across England, Wales and Scotland and how the COVID-19 pandemic has affected them. A short online survey was distributed to all prison governors of male remand prisons in England, Wales and Scotland. The survey questions were largely closed questions with simple 'yes' or 'no' responses, but some open-ended questions were included to obtain a more in-depth understanding in specific areas. Out of the 55 prison governors invited to take part in the survey by e-mail, 17 completed the survey.

Background

During 2020 and the subsequent years that followed; the COVID-19 pandemic was seen to have a monumental impact across the world.⁸³ It affected every aspect of society from the increased demands on the healthcare system, the negative effects on employment and businesses and the adverse effects on mental health, with individuals being left isolated from friends and family.^{83,84} The prison population was not spared from the impact of the pandemic, and both positive and negative implications associated with the prisoner and prison staff mental health outcomes have been reported.^{85,86} Prisons across the world immediately went into lockdown.⁸⁷ This left prisoners being unable to maintain contact with their family and friends and they often found themselves isolated and confined to their cells for extended periods of time, in order to minimise the risk of a COVID-19 outbreak within the prison.^{85,88} Although in one study this was argued to have led to a reduction in the cases of self-harm being reported,⁸⁶ it was also said to contribute to the deepening of the pains of imprisonment in another.⁸⁹

As expected there has been an outpouring of research projects focusing on the different impacts of the pandemic, across different populations. There have been examples of research focusing on the impact of COVID-19 within the prison setting.⁸⁵⁻⁸⁹ However, to date there has been no research which has specifically focused on the impact to the alcohol treatment services, which were routinely available to prisoners pre pandemic. Although the APPRAISE feasibility study commenced in 2019, the data collection and the follow-ups spanned across the pandemic years of 2020 and 2021. The restrictions imposed caused a significant impact on the research study. The study's recruitment process was significantly affected by the prisons going into lockdown, with Scotland's recruitment having to be halted before reaching the desired target sample. In addition, the follow-ups were hampered by being unable to be undertaken face to face and often it proved difficult to confirm the location of prisoners, with court dates and release dates being postponed as a result of the pandemic.

As the world started opening up again and restrictions began to ease, it felt salient to conduct a survey, within the APPRAISE study, that explored the impact of the COVID-19 pandemic on prisons and the availability of the alcohol treatment services. The purpose of the survey was to give more context to the delays experienced in the APPRAISE study, but also to establish the wider implications of the lockdown on services for alcohol within the prison. As the APPRAISE study focused on male remand prisoners, the survey sought to include all prisons within England, Wales and Scotland that held male remand prisoners. The alcohol treatment services that worked within the prison and were the intervention workers used to deliver the APPRAISE intervention (CGL) were largely unable to complete the data collection due to the pandemic. This impacted hugely on the data collected and hence the findings of the study. Therefore, we anticipated by undertaking this survey it would help us to understand what the prisons did in relation to alcohol services throughout the pandemic, and to fill in the gaps in our data that the research team were not able to complete.

Aims and objectives

The aim of conducting the survey, as a component of the APPRAISE study, was to further understand the alcohol services that are currently provided in male remand prisons across England, Wales and Scotland and how the COVID-19 pandemic has affected their delivery.

The specific objectives to achieve the above aim were:

- to establish what alcohol services are currently provided within male remand prisons
- to explore the prison governors' understanding of alcohol brief interventions
- to understand how the COVID-19 pandemic has impacted the services available in male remand prisons
- to identify whether the impact of the COVID-19 pandemic upon services could be avoided in the future.

Methods

Survey design

The survey was developed to ask questions related to the delivery of alcohol brief interventions and other alcohol treatment services, within the prison setting, in order to ascertain what services the prison currently provided. Questions were largely closed, with 'yes' or 'no' responses, but did include some open-ended questions to obtain a more in-depth understanding in specific areas. The latter part of the survey asked participants whether services were affected by the COVID-19 pandemic and, if so, what could be done to ameliorate this in the future. The survey also provided participants with the opportunity to say whether there was anything that was not asked that they wished to express around the impact of the pandemic. The survey finished by asking whether any participants would be willing to engage with the research team in the context of a future study in remand prisons. It was made clear that this response was in no way binding, but simply a way of gauging an initial expression of interest for planned future work.

The survey was distributed online using the JISC tool (www.onlinesurveys.ac.uk/), which was used to both develop and distribute the survey. By using an online platform it ensured that the survey was accessible across a wide geographical reach and that participants did not need to return hard copies, avoiding a pitfall for completion. In addition, by using the online platform it also allowed the survey data to be transferred to SPSS for analysis, which avoided security being compromised by surveys being returned and entered by hand. JISC also allowed the scheduling of e-mail reminders to be sent to prospective participants reminding them about completing the survey.

Setting

The primary setting of the APPRAISE study was two specific remand prison sites in England, Wales and Scotland. However, to get a broader range of responses across different remand prisons the survey component was extended to invite all prison governors, across England, Wales and Scotland, working within prisons with men on remand to take part in the survey.

Sampling and recruitment

As the contact details for prison governors are freely available online, online searches were conducted to obtain the e-mail addresses of all of the governors, across the UK. E-mails were then sent to each prison governor which included a link to the online survey, alongside an information leaflet. The information leaflet was used to explain the purpose of the survey and what the participants' responses would be used for.

Consent

The survey asked all participants to provide their informed consent, before completion, via a consent form built into the start of the survey. It stated that participation was voluntary and that participants did not have to answer any questions that made them feel uncomfortable. It also reassured participants that all personal details, such as names and places, would be removed, in order to ensure they were not identifiable from their responses. In addition, participants were informed that they could withdraw their consent at any point during the survey and could even withdraw their responses following its completion. It was stated that participants could withdraw their responses, for up to 2 weeks, following the completion of the survey, by contacting the lead researcher and hence the respective contact details were supplied.

Data analysis

Analysis of the survey data took place once all surveys were anonymised. SPSS was employed to analyse the survey data obtained. As the survey primarily consisted of yes or no responses the data analysis mainly consisted of percentage calculations. For the more open-ended questions, simple thematic analysis was undertaken to ascertain qualitative statements, which could be used to draw conclusions and answer the survey's objectives.

Data management

All contact with prospective participants was undertaken electronically, along with the creation of all necessary research materials. Any data collected was stored in accordance with GDPR 2016/679. The exported data were stored in a shared folder on a secure network drive which was only accessible by the research team. All identifiable information was removed from the survey results, in order to protect the identity of participants. A password-protected database was stored in a secure network folder at Teesside University, which contained the prison names and the survey IDs, in order to assist the removal of a participant's responses, if they requested to remove themselves from the study. This file was securely deleted once the survey had concluded and the survey withdrawal period had closed. Teesside University acted as the data processor and data owner. Data and research materials were also stored in a Microsoft Teams group which was only accessible to members of the research team and was used to upload, store and circulate all relevant study files and documents securely.

Ethics

Although the APPRAISE study obtained ethical clearance from the University of Edinburgh, a separate ethics application was submitted to Teesside University ethics within the school of Social Sciences, Humanities and Law, for this survey component – 7 October 2021 (2021-6427). Ethical clearance was also obtained from HMPPS on 21 December 2021 (HMPPS 2019-2240) via an amendment to the original approval of the overall APPRAISE study. Participants were notified that their survey responses would be confidential, and only if they disclosed something which indicated a risk to them or someone else would it potentially be necessary to breach confidentiality. In this case the procedure involved the project supervisor being informed, and if deemed appropriate the project supervisor would take further action based on the safeguarding protocol. However, confidentiality did not need to be broken across any of the included participants.

Results

Obtained sample

Out of the 59 prisons in the UK, successful e-mail invitations were sent to 55 governors inviting them to take part in the survey by e-mail; 17 completed the survey. Difficulties were encountered during this part of the research process due to the apparent turnover in staff undertaking the prison governor role. Many e-mail invitations were returned as not being able to be delivered and more investigating had to be carried out.

Governors were asked to disclose the capacity of their prison and, from those who answered, the smallest prison held 577, and the largest 1600.

Findings

The findings of the survey are presented below in relation to the objectives discussed above. The response rate of 17 out of 55 completing the survey needs to be borne in mind when considering the study findings. It may be that those prisons that were more engaged in alcohol services responded and as such self-selected.

Current alcohol services within male remand prisons

The survey showed a wealth of knowledge on the different services offered within prisons with men on remand. The survey gathered information on both the service providers and the services offered. These services included:

- Alcoholics Anonymous (AA) sessions: 58.8% ($n = 10$).
- NHS services: 35.3% ($n = 6$).
- Psychotherapy/Counselling/11 support: 29.4% ($n = 5$).
- In-cell work packs: 23.5% ($n = 4$).
- Smart recovery: 17.6% ($n = 3$).
- Support services via community partners: 11.8% ($n = 2$).
- Group work: 11.8% ($n = 2$).
- Detox services: 11.8% ($n = 2$).
- Substance misuse services: 5.8% ($n = 1$).
- CJSW Addictions team: 5.8% ($n = 1$).

The above services are provided by a broad range of providers. These included: NHS services, AA sessions, Change Grow Live, DART (drug and recovery team), Inclusion team, criminal justice social worker (CJSW) Addictions team, Chaplaincy support, Scotland and Midlothian Offender Recovery Service (EMORS), Dfodol, trained prison staff, SPS, Forward Trust, Drugs Action, Turning Point and third-sector services. The most frequently mentioned provider was NHS services ($n = 10$, 41.2%).

Governors' understanding of alcohol brief interventions

All governors were asked about their knowledge of alcohol brief interventions. When initially asked, 82.4% ($n = 14$) stated that they did know what the intervention was.

The survey then asked all governors if to the best of their knowledge their prison delivered alcohol brief interventions. Ten (58.8%) said that they thought their prison did deliver the intervention. Within the survey they were then given some information on what the intervention entails and asked again if they knew if they were delivering them. This time, more governors declared that they think they are delivering the intervention ($n = 11$, 64.7%). This shows that the knowledge of the intervention helped with making a decision about whether alcohol brief interventions are being delivered.

One of the questions at the end of the survey asked those who wished to, to leave an e-mail address if they were interested in taking part in any future trial around alcohol brief interventions. For this question, 64.7% ($n = 11$) did express this interest, showing an interest in developing their alcohol provisions further.

The impact of the coronavirus disease discovered in 2019 pandemic upon services available in male remand prisons

The pandemic undoubtedly affected the alcohol services within the prisons with men on remand in England, Scotland and Wales and that were included in the overall APPRAISE study (HMP England and

HMP Scotland). The survey also established this, with 15 of the 17 prisons (88.2%) answering yes to 'did the pandemic impact alcohol services within your prison?'

The impact on services was answered in an open question, with governors giving detailed responses to the question of how services were impacted. A summary of some of the service impact is as follows:

- group sessions being suspended
- face-to-face sessions being suspended
- services suspended – being unable to attend the prison
- staffing issues
- interventions being reduced
- a lack of resources.

The majority of governors mentioned the services that entailed group sessions being affected ($n = 8$, 47.1%). This and all of the barriers were due to not being able to see the men face to face and the social distancing measures in place. The conditions imposed on the prison by government to stop the spread of the COVID-19 virus and keep the men and staff safe meant staff and organisations other than prison officers and healthcare staff were unable to enter the prison. There were some prison governors who mentioned an attempt at using telephone support but due to staffing pressures this was unsuccessful too. Staff numbers were limited due to isolation rules or being vulnerable or (potentially) contagious.

This has shown that although there are some services being carried out with in-cell packs, most alcohol services rely on face-to-face intervention work, mostly in groups, and carried out by outside organisations.

Interestingly, we asked the prison governors about 'other services' delivered in the prison and whether or not these were impacted by the pandemic. In this instance, 100% ($n = 17$) answered 'yes'. The reasons for impact were all the same as those mentioned above in relation to the disruption to alcohol services. While the majority of services mentioned were related to drugs and smoking, there were also other services disrupted, such as health care, therapeutic services, gambling support, education, religious services, the gym and, importantly, the overall regime of the prison.

Best practice to avoid any future disruption to services

When asked whether the disruptions due to the pandemic would have a long-term effect on the prison, there was a divide between responses, with 41.2% ($n = 7$) stating they felt it would, and 58.5% ($n = 10$) stating no, they would not. Those that felt the affect would be felt for a long time had different reasons for this. However, most responses discussed that this was from the fact that there was a long gap in services, and the impact of that having a long-term effect, for example, not having treatment and falling more unwell. One governor discussed the DART team not having access to the prisoners, which ultimately led to fewer services now being delivered, and these have not yet been fully increased back to pre-pandemic levels. All of those who responded stated they were unable to say when they thought this long-term affect would end, but one governor predicted 'years to come'.

Every governor who answered the survey had an opinion on how best to avoid a future disruption. First, we gave options as to what they thought would lead to less disruption should a pandemic arise in the future. The most favoured appeared to be something tangible for the prisoners to work on in their cells:

- online sessions: 76.5% ($n = 13$)
- workbook/paper sessions: 82.4% ($n = 14$)
- staff continuing interventions but wearing full personal protective equipment (PPE): 70.6% ($n = 12$).

PPE featured heavily in some of the open-ended questions and was considered a barrier to overcoming further disruption if not dealt with more quickly in a future pandemic.

Other suggestions given by those who returned the survey included comments about better provisions for in-cell phones, or in some circumstances the praising of purple visits which allowed video calls. This highlighted some good that came from the learning of the COVID-19 pandemic and new innovations. Aside from comments regarding a better, more appropriate use of technology should this ever happen again, one governor commented on the services being seen as 'essential' and therefore not something that is withdrawn in the same way it was in 2020.

Chapter 9 Discussion and conclusions

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For individuals in contact with the criminal justice system, the prevalence of at-risk drinking is far higher than in the general population. There is very little evidence of efficacy or effectiveness of alcohol interventions in reducing risky drinking amongst those in the criminal justice system and in particular men on remand in prison. This is compounded by the limited evidence for the optimum timing of delivery, recommended length, content, implementation and economic benefit of an extended alcohol intervention in the prison setting. APPRAISE aimed to provide vital evidence to inform a future definitive RCT of an extended alcohol intervention for men on remand in prison.

APPRAISE specifically aimed to undertake a two-arm, parallel-group, individually randomised pilot study of a self-efficacy-enhancing psychosocial alcohol intervention for men on remand in prison.

Results relating to objectives are detailed in the individual chapters and summarised below.

Objective 1

To pilot the study measures and evaluation methods to assess the feasibility of conducting a future definitive multicentre, pragmatic, parallel-group RCT.

- 1a. Is it feasible to conduct a future multicentre RCT of a self-efficacy-enhancing psychosocial alcohol intervention for men on remand?

We cannot say that it is feasible to conduct a future definitive trial as we were unable to follow up participants – of course COVID-19 played a part in the low response rates but, at this present stage, with the results we have we are not yet in a position to move this work forward to a definitive study.

- 1b. Can we obtain reasonable estimates of the parameters necessary to inform the design and sample size calculation for a future definitive multicentre RCT? This includes standard deviations of potential continuous primary outcomes and estimates of recruitment, retention and follow-up rates.

Yes, but only partially. We can obtain reasonable estimates of standard deviations of outcomes based on the baseline data collected. However, the amount of valid data collected at follow-up was too little to be able to provide us with reliable estimates of parameters for outcomes at follow-up.

The pre-pandemic recruitment data give us reasonable estimates of recruitment rates in a future trial. However, the retention and follow-up rates we observed in the APPRAISE trial were likely to have been heavily influenced by the COVID-19 pandemic and associated lockdown measures, and therefore these are not reliable indicators of what they might be in a future trial.

- 1c. How well do participants complete the questionnaires necessary for a future definitive RCT?

At baseline there was a very small amount of missing data for a few questionnaires/outcomes, for example, 'Readiness to Change Ruler: The people who are important to me support me in reducing

my drinking', and for the 'Drinking Refusal Self-Efficacy Questionnaire' and for the 'Negative Alcohol Expectancy Questionnaire'. The amount missing was very similar between sites at baseline. At follow-up, the amount of missing data was very substantial in both sites. We were able to do slightly better at the Scotland site because 'we were able to have some access to the prison site and identify where participants were located'. In addition, the same researcher conducted the majority of the follow-ups in Scotland at 12 months as conducted the recruitment and baseline measures, which may have had an effect on participants' willingness to respond. However, numbers are small so we must be cautious with this assertion. In terms of data completeness of the questionnaires at each site, this was similar between the sites ([Report Supplementary Material 1 S1.1](#) and [S1.2](#)). It is, however, really difficult to read too much into the differences because the numbers completing data at each site were so small. Similarly, the different questionnaires had similar amounts of missing data at follow-up (see [Table 28](#)) and we cannot really say too much about differences of ± 1 or ± 2 . In summary, we did not find any notable differences across the different sites/instruments. However, it could be that self-completion was a factor; at baseline the questionnaire was not self-completed, but RA-delivered. In addition, we also observed and reflected on the suitability of certain questions, particularly if the questionnaire was to be self-completed (which was reflected in later amendments) and this would warrant further consideration in future research for this population.

1d. Can we collect economic data needed for a future definitive RCT?

We were able to design and implement an ABC study that provided detailed estimates of the implementation costs of the APPRIASE intervention. That we could do this despite COVID-19 complications suggests that our methodology will be appropriate for a definitive RCT. We nonetheless note that additional attention needs to be devoted to some aspects of our ABC study. In particular, more attention should be devoted to assessing the prison space needed to deliver the intervention and more attention should be given to separating research costs from true intervention costs when it comes to scheduling follow-up intervention visits. We were also able to modify the Economic Form 90 for use with a UK prison population and found no evidence that it was difficult for study subjects to complete. However, we were unable to validate the data-collection instrument and recommend that such validation be part of a definitive trial.

1e. Can we access recidivism data from the PNC databases for trial participants?

1f. Can we access health data from routine NHS data sources for trial participants?

Although it is possible that COVID-19 played a role in our inability to obtain these data, we do not think that it was solely responsible. Two key issues are at play: prisoner confidentiality and data systems. We found that gaining approval to obtain the PNC or NHS identifiers was difficult to impossible due to a combination of prison and ethics board concerns. Although these concerns may fade as more evidence of the benefits of the APPRAISE intervention mounts, we conclude that it is very likely that these concerns will still exist for a definitive trial, and possibly be even more salient for some stakeholders in that context. Exacerbating the confidentiality issues were data-system issues that prevented data access on a large scale. Although this issue depends on the jurisdiction, in each of our two jurisdictions at least one of the data sources required case-by-case data extraction at some step of the process. We could not extract either PNC or NHS data on a large scale in either England or Scotland, making the use of such data prohibitively costly for a definitive RCT.

Objective 2

To assess intervention fidelity

2a. What proportion of the interventions are delivered as per protocol?

We recorded three of the in-prison interventions. They all scored highly and had maximum scores on the BECCI Scale.

2b. Is there any evidence of contamination between the two conditions and/or between those workers delivering the intervention?

We did not find any evidence of contamination between workers delivering the intervention. At the Scotland study site, the intervention was only delivered by one interventionist. In England two interventionists were delivering the intervention. There was mention of discussion of the study between one of the remand participants and another (see sub-section 'Coherence: acceptability of the APPRAISE intervention' in [Remand participants](#)) but the participant stated that he didn't know if the other participant got the intervention or not. No other evidence was obtained of contamination; however, it would be prudent to explore this again in any future work in the absence of a global pandemic, lockdown and restrictions across the prison estate.

2c. To what extent was the intervention changing process variables consistent with the underpinning theory?

[Table 28](#) shows the results for changes in process variables (secondary outcomes). The available data do not allow answering the question if the change in process variables from baseline to 6 and 12 months was greater in the intervention group than in the control group in central process variables (goals, self-efficacy, outcome expectations, according to social cognitive theory), as we were able to obtain only limited data for change in mediators (especially at 6 months) and only a small number of participants were released, which precludes interpretation of change in mediators after release from prison. We found small and inconsistent changes in process variables with considerable variation. Therefore, we cannot answer this research question.

Objective 3

To qualitatively explore the feasibility and acceptability of a self-efficacy-enhancing psychosocial alcohol intervention and study measures to staff and for men on remand and on liberation.

3a. How acceptable are the trial and intervention procedures (including context and any barriers and facilitators) to the following key stakeholders: men on remand in prison and on liberation; prison staff (including healthcare staff); commissioners; policy-makers and third-sector partners?

The qualitative process evaluation showed that the trial and intervention procedures were acceptable to staff and men on remand. We found that it was important for an individual to feel comfortable and trusting of an interventionist before sharing their personal experiences and consumption. Staff did report that the prison setting and resources could affect the work due to the lack of available space and the limited staff capacity. However, in general participants believed that delivering the APPRAISE intervention did not significantly increase their workload. The intervention training and content appeared to be largely acceptable, although the length and content could have been condensed. The APPRAISE intervention delivery benefited from buying into the intervention and being motivated and engaged with the content.

Similar findings were found around the prison's logistical issues affecting the APPRAISE intervention delivery feasibility, such as space and staff capacity, the importance of engagement with prisoners and the lack of currently available support for prisoners on remand from the stakeholder interviews. Additional themes and subthemes that emerged were around the intricacies of the prison environment and how it affected the study and intervention delivery, the individual differences across the setting and across the participants and also further barriers and facilitators to delivery of the ABI.

A key finding, observed in all three participant groups, was that the in-prison APPRAISE sessions were deemed to be more acceptable and feasible than the post-liberation intervention sessions. This was observed due to a plethora of different factors, such as being unable to re-establish contact with males upon their release and individuals relapsing to previous risky alcohol consumption when alcohol becomes more freely available.

Objective 4

To assess whether operational progression criteria for conducting a future definitive RCT are met across trial arms and study sites and, if so, to develop a protocol for a future definitive trial. Operational progression criteria are based on previous research results.⁶

A large proportion of this study was carried out during the COVID-19 pandemic and as such we cannot demonstrate the full extent to how this affected the study. However, we did not meet the criteria for developing a protocol for a future definitive study.

4a. Do the two prisons invited to the study agree to take part?

Yes. Both Scotland and England prisons agreed to take part in the study and stayed involved until COVID-19 restrictions closed access to the project down.

4b. Based on knowledge from previous data, do at least 90 eligible participants consent to take part and be randomised across the trial arms?

No. Although in the England site, where recruitment began earlier than the Scotland site, we recruited 90 participants, the COVID-19 pandemic and associated lockdown restrictions meant that we were not able to enter the prison and therefore could not achieve the target of 90 participants in the Scotland site ($n = 42$). However, in a future study without lockdown restrictions we are confident that the required number of participants can be recruited.

4c. Do at least 70% of participants who consent to the trial receive the intervention?

Yes, partially. Of the 68 consented and randomised to the trial 53/68 (78% of participants) received the in-prison intervention, although much lower numbers received the post-prison component of the intervention. Only one participant received the day-3 post-liberation intervention, none received the day-7 post-liberation intervention, and one person received the 21-day post-liberation intervention, at least in part because of the COVID-19 context. A post-prison-liberation component of the intervention may not be viable in a future trial.

4d. Are at least 60% of those who received the intervention followed up at 12 months across trial arms and study sites?

No. Only 13% (18/132) of those who received the intervention were followed up at 12 months. Although the COVID-19 pandemic was undoubtedly a factor in the low percentage followed up, we observed high percentages of participants who were not released at 12 months; this was worsened by COVID-19 because of the number of those on remand awaiting trial due to suspended and closed courts; or of those who were released, many could not be contacted or could not be located. We consider this to be the greatest barrier in designing a future trial that will achieve satisfactory follow-up rates.

Objective 5

To ascertain what alcohol services are available in male remand prisons and how COVID-19 has affected services.

From 59 prisons in Scotland and England, successful e-mails were sent to 55 prison governors. Seventeen (31%) were completed. A limitation of the study was the barriers to freely accessing governor e-mails. While this was handled by a gatekeeper who worked for SPS for the Scottish prisons, no such facilitation took place in England, and while most e-mails were freely available, many were not. Two-thirds of the sample were interested in hearing more about ABIs.

5a. What alcohol services are currently provided within male remand prisons?

There are a wide range of services from a wide variety of providers. The most common were AA sessions (59%), NHS services (35%) and psychotherapy, counselling or 1--1 support (29%).

5b. To explore the prison governors' understanding of brief interventions.

Two-thirds of the governors understood what ABIs are and that they were being delivered.

5c. To understand how the COVID-19 pandemic has impacted the services available in male remand prisons.

We found that the COVID-19 pandemic had a devastating affect across all prisons in relation to the services they provide. It was also useful to establish what the prison governors deemed important learning from the pandemic, in this instance mostly around PPE and a better use of technology.

5d. To identify whether the impact of the COVID-19 pandemic upon services could be avoided in the future.

There were major impacts in relation to services during COVID-19 which included all sessions being suspended and there being staffing issues and a lack of resources.

Patient and public involvement

Patient and public involvement (PPI) has informed and influenced the development of the study through various stages. PPI representatives involved in the research design include: those with experience of being in prison and the criminal justice system, a social enterprise group, prison employees and NHS staff. At the time of the study conception and development, our PPI co-applicant, Sharon Mercado (SM), was a community justice mentor (CJM) at the Wise Group. As part of our PPI work, she brought together service user representatives and held a participatory workshop with men who had experience of being in prison on remand. The aim of the workshop was to discuss and provide suggestions and feedback regarding development of the study design. Their views informed and influenced the study design: need for pictorial information for study material to address literacy/readability; strategies to maximise follow-up by obtaining contact details of family/significant others, homeless shelters and community justice support workers. SM was a member of the PRISM-A study Advisory Board and has continued to work with us as a co-applicant on the APPRAISE study, where she informed the research design and follow-up strategy using experience of her own follow-up work as a CJM with prisoners on liberation. SM continued to provide input throughout the duration of the study as a member of the PMG and in the

recommendations for a future definitive trial. Wider PPI was present through TSC membership where we also had wider representation.

The PPI was an extremely positive experience for the team and the study benefited hugely from all their input. This was particularly the case when we were reviewing the impact of COVID on follow-up and considering the mitigations we could take moving to self-completion questionnaires, change of primary outcome measure and adaptations to some of the secondary outcome measures. The input from SM was key. The impact of COVID and the subsequent restrictions on face-to-face meetings did have a negative effect on PPI engagement. The face-to-face meetings were key for the relationships we had been building and meeting face-to-face with PPI colleagues was always a positive experience for everyone.

Reflections and lessons learned from patient and public involvement involvement

Our reflections and learning of our PPI engagement in the APPRAISE study was that we should in future look to strengthen the PPI presence within any future studies. It would have been useful to have created a PPI-led space for the PPI members for them to have the opportunity to meet away from the PMG and TSC. On reflection the TSC and PMG meetings could be quite technical at times, with language used that PPI colleagues felt they were less familiar with. As SM was the sole PPI member of the PMG, in future it would be useful to have a minimum of two members to have a more balanced group. If we created a separate PPI space where PPI members could meet outside of formal meetings to set the agenda, it could be used as a safe space for them to discuss and provide regular feedback to the TSC and PGM. It may be possible to also run workshops linked to the study but also to have informal activity too. With SM we would generally meet ahead of the PMG and have refreshments together; this worked well and has informed the idea of a separate PPI space for PPI colleagues.

SM has provided her own insightful reflections on the research study in relation to what went well and what could have been done differently to improve the PPI representation and involvement. These reflections and recommendations, in her own words, are presented here.

Engaging with offenders is not easy. It takes time and patience and recognising that sometimes, despite the willingness to engage, it doesn't always happen the way you anticipated.

Holding a focus group needs careful planning especially as you need to make sure that those attending will feel safe, and anything said is confidential. Offenders' lives, especially those that are involved with substance misuse/alcohol, are chaotic. Therefore, making sure those attending are not going to meet anyone else that they have issues with (enemies, debts etc.) requires input. Despite all the support to attend (incentives were included, food vouchers, travel expenses), the focus group was not attended by all those invited. However, even with a small number, input was good. Those still on remand were also on an individual basis asked for input regarding the information leaflets, and what works best in terms of follow-up appointments after liberation.

To summarise, the whole experience was challenging. Regardless of the research, working with offenders/ex-offenders can be difficult, especially with those who due to their alcoholism can have chaotic lives. This makes it hard to continue engagement with follow-up appointments. There are things that can be done which may assist in getting increased involvement.

- *Building up more trust and spending more time with the offender can help. Rather than just a few engagements, have a few more or enlist support from other agencies working with the offender just to build up the momentum and remind the offender of the research project. This could prevent the offender from disengaging.*
- *Better offender tracking after liberation. Some people may have a mentor or a support worker: find out (hopefully before liberation) who is currently working and keep up communication with them, they may be able to inform you of a person's whereabouts.*

- *Offering incentives may help. Whilst offering financial incentives (£20 per community engagement), it can help. It is recommended that any financial rewards are given in food/shop vouchers or pre-payment energy support.*
- *Invite those who are now in recovery and have recent relevant experience to come and talk to the research team at the start of the project. It can be a good opportunity to find out more about the lives of those affected by alcohol and why engagement or non-engagement happens.*

Equality, diversity and inclusion

Equality, diversity and inclusion (EDI) was a core value of the purpose of our research by the very nature of the setting of the research, the focus on alcohol-related harm and the population we were focusing on. Conducting the study required EDI to be at the forefront of our work from inception with our awareness of the lack of evidence, to meeting ethical approval requirements and working within the criminal justice system. An overview of our experience and reflections are provided in relation to participants, the evidence base, the study design, the research team and PPI representation.

Population characteristics and optimising participation

We are aware that undertaking research within the prison setting is challenging and not commonplace and for trials research it is even less common. The challenge is increased even further when there is a focus on the remand population. The study was developed and conceptualised by drawing on the collective experiences of the research team, our previous research and work with those incarcerated and from the published literature. From this we had a unique understanding of our population and who should be in the study. In our previous research we had undertaken interviews with men in prison who were on remand and asked them specifically how we could optimise participation in a future trial. We had also undertaken interviews with prison service colleagues providing care in the prison setting. Their reflections and insights directly informed the study design and how we approached the research as described in [Chapter 3](#) to be as inclusive as possible. We also purposely undertook the study in two different prison estates to explore how the impact of the variations in category of the prison population and the estate regimes impacted participation. This was a factor from our previous work we had become aware of and remains a key factor for researchers to consider in any future studies along with the use of digital technology from the use of digital recorders to use of mobile phones.

Gaps in the evidence base

To the best of our knowledge we are not aware of any other trials for men on remand focusing on the delivery of a self-efficacy-enhancing extended alcohol brief intervention, at least in the UK, hence the need to undertake the pilot trial. The key gap that remains for this particular population is how best to design a study that provides an optimum follow-up process. We have provided recommendations in [Chapter 9](#) for future studies.

Encouraging under-represented groups

Our PPI engagement with service users during the conception and design phase of the study supported the development of the study material through use of images and the material layout. This was specifically with the aim of optimising accessibility and inclusion.

Research team inclusivity and development opportunities

Our research team comprised a range of under-represented stakeholder groups and individuals largely seen in our TSC membership. The research team also had a good gender balance with a larger percentage of women and early-career research representation evident as co-authors. The ethos and values of the team ensured development-supported opportunities were available to take on key elements of the study, for example, process evaluation interviews and data analyses and COVID survey. Similarly, we provided development opportunities for those coming from a non-research background, for example, our co-applicant service user (SM) and our researcher who collected the baseline data at the England study site.

Patient and public involvement representation

Our PPI representation is outlined in the section [Patient and public involvement](#). The study participants were representative of those from our previous work with this population. The impact of COVID has been described elsewhere in the report. Our additional COVID impact survey provided some further data on how this population were affected in relation to the delivery of alcohol services during the pandemic.

Strengths and limitations

Strengths of the study

- The APPRAISE study, to the best of our knowledge, is the first pilot trial of an alcohol-focused self-efficacy-enhancing intervention for men on remand.
- The study provides significant insight into the feasibility and acceptability of pilot trials for this particular population.
- The ethical approval process for the prison setting across devolved countries provided useful insights into how to navigate the submission and approvals process across a range of ethics committees.
- The study provided good evidence of the feasibility of recruitment, training interventionists and subsequent in-prison delivery of the APPRAISE intervention to 73% of men on remand, randomised to the intervention, within a highly complex prison setting.
- The economic findings provide valuable insights, as we believe this to be the first of its kind for this type of intervention in this setting and that the methods discussed will likely be relevant and transferable across other settings.
- Useful insight regarding the access to PNC and NHS service use data was obtained.
- The process evaluation provided useful insights into the perspectives of the remand participants, intervention delivery team and wider stakeholders on the acceptability and feasibility of implementation of the intervention.

Limitations of the study

- Protracted multiple ethical approvals across devolved countries and processes meant a significant delay in recruitment commencing, particularly for the Scotland site.
- We were unable to deliver the post-liberation elements of the intervention.
- The onset of the COVID-19 pandemic resulted in prison restrictions and no access to the prison site, meaning we were unable to identify if participants had been released or not at 6 months, with no access still at the England site at 12 months and very limited access at the Scotland site at 12 months.
- Although the COVID-19 pandemic was undoubtedly a factor in the low percentage followed up, only 13% (18/132) of those who received the intervention were followed up at 12 months.
- Changes to the study protocol to include self-completion of follow-up data may have made it difficult for participants to complete the survey and resulted in larger proportions of missing data as opposed to RA survey completion at baseline, where missing data were minimal.
- We did not include probation services in the trial, which on reflection would have strengthened our post-liberation follow-up process.

Recommendations regarding a definitive randomised trial

From the study progression criteria for a full RCT, we have identified the following recommendations:

1. Buy-in for a research trial of this nature in prison requires significant pre-study trust and relationship development, buy-in from the prison estate, governor, prison officers, peer mentors and embedded third-sector services.

2. Recruitment and randomisation of men on remand to a future APPRAISE RCT are possible, with trust in the research team an important factor.
3. Training team members of existing alcohol services to deliver APPRAISE as per protocol is possible.
4. Delivery of the in-prison APPRAISE intervention is possible and would require appropriate space, time and team member capacity.
5. Economic evaluation is possible.
6. Post-liberation intervention delivery would only be possible if there was a robust follow-up process identified and in place.
7. Six-month and 12-month follow-up would only be possible if there was a robust follow-up process identified and in place. What that might look like needs to be carefully considered, explored and understood in an already complex liberation and rehabilitation statutory and non-statutory agency landscape. This may be achieved through the use of in-depth individual case studies or a realist data-capture approach to cover this period.
8. Further exploration of the inclusion of and collaboration with the probation service in the service delivery and implementation of the APPRAISE intervention may offer higher success rates of post-liberation intervention delivery and a more robust follow-up process.
9. Further exploration of the feasibility of utilising routinely collected data to evaluate outcomes post prison release is also warranted, for example, presentation at community alcohol services, presentation to GPs, and/or liver function test readings on routinely collected blood-test investigations.
10. Future research funding amongst prisoner populations must be made available to address the feasibility issues identified in this pilot trial so that a best-practice framework can be developed and utilised by research teams seeking to conduct much-needed research amongst this under-served group.

Conclusion

Addressing alcohol harm in prisons, at what can be considered a 'teachable moment', where there is an opportunity for reflection on their risky drinking and their current position, could potentially reduce the risk of re-offending and costs to society, while helping address health inequalities. For those on remand in prison, the prevalence of at-risk drinking is between 62% and 68%.⁷

Many men on remand do not have the opportunity to access mainstream prison health or public health services. An intervention such as APPRAISE offers an opportunity to provide an extended behavioural-based alcohol intervention to men on remand. The APPRAISE study has identified that despite the complexities of ethical approval and the time taken to build relationships and trust, it is possible to undertake key elements of a future RCT, but not all, namely follow-up. Further focused research needs to be undertaken to explore, identify and develop a robust process and system to optimise follow-up post liberation. A recent pilot trial in prisons has shown promise in this area, having identified satisfactory follow-up at 6 months, but not at 12 months.⁹⁰

The evidence base to meet the needs of men on remand in relation to risky drinking remains weak. However, there are opportunities to build on the work of APPRAISE to ensure that those in the prison setting have equal access to interventions that have the potential to positively impact their relationship with and use of alcohol.

Additional information

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All authors agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

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Patient data statement

This work uses data provided by patients and collected by the NHS as part of their care and support. Using patient data is vital to improve health and care for everyone. There is huge potential to make better use of information from people's patient records, to understand more about disease, develop new treatments, monitor safety and plan NHS services. Patient data should be kept safe and secure, to protect everyone's privacy, and it's important that there are safeguards to make sure that they are stored and used responsibly. Everyone should be able to find out about how patient data are used. #datasaveslives You can find out more about the background to this citation here: <https://understandingpatientdata.org.uk/data-citation>.

Data-sharing statement

All requests for data should be sent to the corresponding author. Access to available anonymised data may be granted following review.

Ethics statement

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Appendix 1 GRIPP2 short form

Section and topic	Item	Reported on page no.
1: Aim	Report the aim of PPI in the study	<i>Structure of the report.</i> p. 5
2: Methods	Provide a clear description of the methods used for PPI in the study	<i>Patient and public involvement.</i> p. 109–10
3: Study results	Outcomes – report the results of PPI in the study, including both positive and negative outcomes	<i>Patient and public involvement.</i> p. 109–10
4: Discussion and conclusions	Outcomes – comment on the extent to which PPI influenced the study overall. Describe positive and negative effects	<i>Patient and public involvement.</i> p. 109–10
5: Reflections/ critical perspective	Comment critically on the study, reflecting on the things that went well and those that did not, so others can learn from this experience	<i>Patient and public involvement.</i> p. 109–10

Appendix 2 CONSERVE-CONSORT

Use CONSERVE-CONSORT for completed trial reports and CONSERVE-SPIRIT for trial protocols.

CONSERVE-CONSORT Extension: [DATE]					
Item	Item title	Description	Page no.		
I.	Extenuating circumstances	Describe the circumstances and how they constitute extenuating circumstances.	Overall study recruitment . p. 31		
II.	Important modifications	<p>a. Describe how the modifications are important modifications.</p> <p>b. Describe the impacts and mitigating strategies, including their rationale and implications for the trial.</p> <p>c. Provide a modification timeline.</p>	<p>Changes to the original study protocol. p. 28</p> <p>Changes to the original study protocol, Table 6. pp. 28–29</p> <p>Changes to the original study protocol. p. 28</p>		
III.	Responsible parties	State who planned, reviewed and approved the modifications.	<p>Changes to the original study protocol. p. 28</p> <p>CI, PMG, TSC, study sponsor</p>		
IV.	Interim data	<p>If modifications were informed by trial data, describe how the interim data were used, including whether they were examined by study group, and whether the individuals reviewing the data were blinded to the treatment allocation.</p> <p>For each row, if important modifications occurred check 'direct impact' and/or 'mitigating strategy' and describe the changes in the trial manuscript or supplement. Check 'no change' for items that are unaffected in the extenuating circumstance.</p>	N/A		
CONSORT number and item					
		No change	Impact ^a	Mitigating strategy ^b	Page no.
1	Title and abstract	No change			
2	Introduction	No change			
3	Methods: trial design	No change			
4	Methods: participants	No change			
5	Methods: interventions	No change			
6	Methods: outcomes		Yes	Change of primary outcome measure	Follow-up data collection pp. 44–6
7	Methods: sample size		Yes	None	Changes to the original study protocol . p. 51
8–10	Methods: randomisation	No change			
11	Methods: blinding	No change			
12	Methods: statistical methods		Yes	Focus was on descriptive analyses and baseline characteristics given the low numbers providing follow-up data	Statistical methods/analysis plan . p. 46

continued

CONSORT number and item	No change	Impact ^a	Mitigating strategy ^b	Page no.
13	Results: participant flow	No change		
14	Results: recruitment	No change		
15	Results: baseline data	No change		
16	Results: numbers analysed	Yes	Reduced sample size at follow-up and so focus was on descriptive analyses and baseline characteristics	Statistical methods/analysis plan . p. 46
17	Results: outcomes and estimation	Yes	Change of primary outcome measure	Follow up data collection
18	Results: ancillary analyses	Yes	Could not do originally planned exploratory analyses comparing agreement and full logic model analyses	Statistical methods/analysis plan . p. 49
19	Results: harms	No change		
20	Discussion: limitations	Yes	None	Equality, diversity and inclusion . pp. 154–5
21	Discussion: generalisability	Yes	None	Strengths and limitations . p. 155
23	Other information: registration	No change		
24	Other information: protocol	Yes	Protocol was updated to reflect changes	Changes to the original study protocol, Table 6
25	Other information: funding	Yes	Study extension secured and additional survey conducted focusing on impact of alcohol services for men on remand during COVID-19	Survey Chapter 8

a Aspects of the trial that are directly affected or changed by the extenuating circumstance and are not under the control of investigators, sponsor or funder.

b Aspects of the trial that are modified by the study investigators, sponsor or funder to respond to the extenuating circumstance or manage the direct impacts on the trial.

Appendix 3 Template for Intervention Description and Replication Checklist

The TIDieR (Template for Intervention Description and Replication) Checklist:

Information to include when describing an intervention and the location of the information

Item number ^a	Item	Where located ^b	
		Primary paper (page or appendix number)	Other ^c (details)
	BRIEF NAME		
1.	Provide the name or a phrase that describes the intervention.	Origins and development of the APPRAISE intervention , p. 7	
	WHY		
2.	Describe any rationale, theory, or goal of the elements essential to the intervention.	APPRAISE intervention: theory, elements and logic model , pp. 7–10	
	WHAT		
3.	Materials: describe any physical or informational materials used in the intervention, including those provided to participants or used in intervention delivery or in training of intervention providers. Provide information on where the materials can be accessed (e.g. online appendix, URL).	APPRAISE intervention materials , p. 10	
4.	Procedures: describe each of the procedures, activities, and/or processes used in the intervention, including any enabling or support activities.	APPRAISE intervention procedures , pp. 10–11	
	WHO PROVIDED		
5.	For each category of intervention provider (e.g. psychologist, nursing assistant), describe their expertise, background and any specific training given.	APPRAISE intervention training , p. 11	
	HOW		
6.	Describe the modes of delivery (e.g. face-to-face or by some other mechanism, such as internet or telephone) of the intervention and whether it was provided individually or in a group.	APPRAISE Intervention delivery mode and location; APPRAISE post-liberation intervention modifications due to COVID-19 , pp. 11–14	
	WHERE		
7.	Describe the type(s) of location(s) where the intervention occurred, including any necessary infrastructure or relevant features.	APPRAISE Intervention delivery mode and location; APPRAISE post-liberation intervention modifications due to COVID-19 , pp. 11–14	
	WHEN and HOW MUCH		
8.	Describe the number of times the intervention was delivered and over what period of time including the number of sessions, their schedule, and their duration, intensity or dose.	APPRAISE Intervention delivery mode and location; APPRAISE post-liberation intervention modifications due to COVID-19 , pp. 11–14 Table 3	

continued

Item number ^a	Item	Where located ^b	
		Primary paper (page or appendix number)	Other ^c (details)
TAILORING			
9.	If the intervention was planned to be personalised, titrated or adapted, then describe what, why, when and how.	APPRAISE intervention procedures , pp. 10–11 APPRAISE post-liberation intervention modifications due to COVID-19 , pp. 13–14	
MODIFICATIONS			
10. ^d	If the intervention was modified during the course of the study, describe the changes (what, why, when, and how).	APPRAISE post liberation intervention modifications due to COVID-19 , pp. 13–14, Randomisation and blinding , p. 19	
HOW WELL			
11.	Planned: if intervention adherence or fidelity was assessed, describe how and by whom, and if any strategies were used to maintain or improve fidelity, describe them.	<i>Process evaluation aims and objectives – context</i> , pp. 97–100 Table 30	
12. ^d	Actual: if intervention adherence or fidelity was assessed, describe the extent to which the intervention was delivered as planned.	Chapter 6 , pp. 67–82	

a We strongly recommend using this checklist in conjunction with the TIDieR guide (see [BMJ 2014;348:g1687](#)), which contains an explanation and elaboration for each item. The focus of TIDieR is on reporting details of the intervention elements (and where relevant, comparison elements) of a study. Other elements and methodological features of studies are covered by other reporting statements and checklists and have not been duplicated as part of the TIDieR checklist. When a *randomised trial* is being reported, the TIDieR checklist should be used in conjunction with the CONSORT statement (see [www.consort-statement.org](#)) as an extension of *Item 5 of the CONSORT 2010 Statement*. When a *clinical trial protocol* is being reported, the TIDieR checklist should be used in conjunction with the SPIRIT statement as an extension of *Item 11 of the SPIRIT 2013 Statement* (see [www.spirit-statement.org](#)). For alternative study designs, TIDieR can be used in conjunction with the appropriate checklist for that study design (see [www.equator-network.org](#)).

b *Authors* – use N/A if an item is not applicable for the intervention being described. *Reviewers* – use '?' if information about the element is not reported or not sufficiently reported.

c If the information is not provided in the primary paper, give details of where this information is available. This may include locations such as a published protocol or other published papers (provide citation details) or a website (provide the URL).

d If completing the TIDieR checklist for a protocol, these items are not relevant to the protocol and cannot be described until the study is complete.

Appendix 4 APPRAISE intervention tool (V3_20October)



The image shows a green rectangular form for the APPRAISE intervention tool. At the top left is a logo consisting of three concentric white circles. To the right of the logo, the word "APPRAISE" is written in large, bold, white capital letters, with the tagline "Rethink your drink." below it in a smaller white font. The form contains several input fields for data collection:

- AUDIT score**: A single white square input field.
- Time when started**: A row of four white square input fields.
- Time when complete**: A row of four white square input fields.
- Participant ID**: A row of five white square input fields.

This resource has been produced as part of a research project in to extended Alcohol Brief Interventions in the prison setting.

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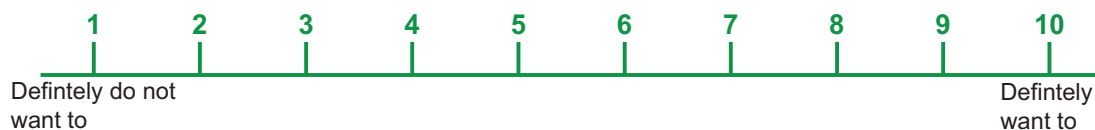
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Teviot Place
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All rights reserved. Material contained in this publication may not be reproduced in whole or part without prior permission of the APPRAISE project or other copyright holders. While every effort is made to ensure that the information given here is accurate, no legal responsibility is accepted for any errors, omissions or misleading statements.

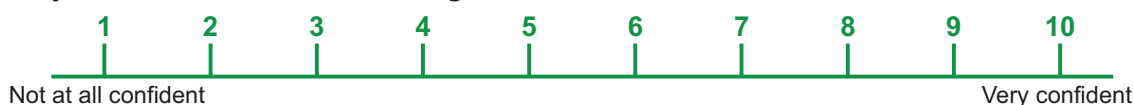
Element 1: Preliminary discussion

- Affirm verbal consent, confidentiality, engagement rules and trust

How much do you want to reduce (or stop) drinking when you are released?



If you decided to reduce (or stop) upon release, how confident do you feel about succeeding with this?



How ready are you to reduce (or stop) drinking when you are released?



I have someone I can talk to about reducing my drinking



The people who are important to me support me in reducing my drinking



I have made a detailed plan regarding when, where and how to reduce my drinking



Things to remember

Discuss answers to readiness to change scales, e.g. Why did you score 7 and not an 8?



Element 1: Preliminary discussion

Below is a list of situations in which people find themselves drinking alcohol. For each situation, please tell us, on a scale of 0–5, whether you think it is a high-risk situation? (0 is no risk and 5 is high risk.)

<i>Situation</i>	<i>Score (0 to 5)</i>
When I am eating dinner	
When someone offers me a drink	
When my spouse or partner is drinking	
When my friends are drinking	
When I am at a pub or club	
When I am angry	
When I feel frustrated	
When I am worried	
When I feel upset	
When I feel nervous	
When I feel sad	
When I am watching TV	
When I am at lunch	
When I am on the way home from work	
When I am listening to music or reading	
When I am by myself	
When I have finished playing sport	
When I first arrive home	
When I feel like celebrating	



Element 2: Acquiring and providing information

ELEMENT TWO

- Provide AUDIT score
- Acquire participants perception of impact:

Have you thought about the impact of alcohol on you and your life?

Impacts of alcohol

Job

- Absent from work
- Loss of earnings
- Poor performance

Family and relationships

- Family breakdown
- Strained relationships
- Unprotected sex
- Domestic abuse
- Child abuse and neglect
- Financial problems

Community

- Accidents, falls and injury
- Anti-social behaviour
- Crime and violence
- Road traffic accidents

Health

- Heart disease
- Cancer
- Stroke
- High blood pressure
- Liver damage
- Infertility
- Pancreatitis, gastritis, indigestion
- Depression
- Dementia
- Anxiety, stress and panic disorders
- Amnesia/memory loss
- Suicide

Present Unit Card: Provide brief information about recommended drinking levels and units.



ELEMENT FOUR

Element 4: Increasing awareness

- Discuss pros and cons of drinking including physiological sensations
- Discuss alternative solutions to dealing with anxiety and stress

Are there any positive things you associate with your use of alcohol?

What are the costs of drinking?

People have lots of reasons why they choose to change or maintain their drinking behaviours.
Let's make a note of yours.

Write down the benefit of reducing or stopping drinking:

7

Element 8: Self-evaluation and self-reinforcement

- Discuss if participant has used self-monitoring or rewarded their successes
- Reiterate the importance of these tools
- Praise attempts and successes
- Relapse should not be attributed to internal stable causes such as willpower or ability
- Tools such as the diary card might help them keep track
- Encourage them to reward themselves for achieving their goal.

Element 9: Culmination

- Summarise your conversation
- Remind the participants of the continuing follow-up.

ELEMENT FIVE & SIX

Write down an example of a strategy you might use in one of your high-risk situations:

Element 6: Goal setting

- Set specific and realistic goals for alcohol reduction with participants

Things to remember

- Set **realistic** and **short-term** sub-goals
- Ensure these are achievable and provide confirmation that participants are able to achieve them



Element 7: Relapse

- Discuss likelihood of relapse
- Discuss general reasons for relapse
- Encourage participant to verbalise specific situations which may result in relapse
- Identify appropriate strategies for relapse prevention

Remember if relapse occurs it is possible to find a more successful way of responding to that situation next time.

Things to remember

- Reiterate that relapse can happen
- Reiterate that relapse occurs typically as a response to stressful situations, social pressure and/or interpersonal conflict
- Remind them of their skills to address relapse and how they can learn to cope with difficult situations

Element 8: Self-evaluation and self-reinforcement

- Reiterate the importance of self-monitoring and identifying successes
- Relapse should not be attributed to internal stable causes such as willpower or ability.
- Tools such as the diary card might help them keep track
- Encourage them to reward themselves for achieving their goal.

Tear along the dotted line and take away with you

Why do you want to reduce or stop drinking?

What is your specific goal upon release?

How will you achieve your goal?

What will make it difficult to achieve this goal?

How will you overcome that?

Who do you call for support? Identify a specific person.

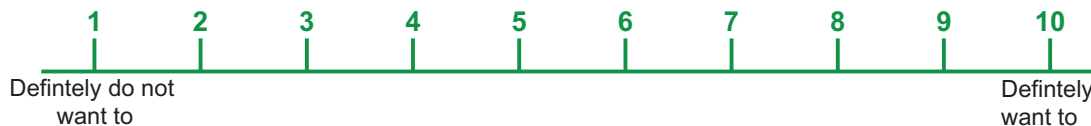
www.changegrowlive.org

Change, Grow, Live - Edinburgh 0131 557 5273

Change, Grow, Live - Durham 0191 4477994



How much do you want to reduce (or stop) drinking when you are released?



If you decided to reduce (or stop) upon release, how confident do you feel about succeeding with this?



How ready are you to reduce (or stop) drinking when you are released?



I have someone I can talk to about reducing my drinking



The people who are important to me support me in reducing my drinking



I have made a detailed plan regarding when, where and how to reduce my drinking



Below is a list of situations in which people find themselves drinking alcohol. For each situation, please tell us, on a scale of 0–5, whether you think it is a high-risk situation? (0 is no risk and 5 is high risk.)

<i>Situation</i>	<i>Score (0 to 5)</i>
When I am eating dinner	
When someone offers me a drink	
When my spouse or partner is drinking	
When my friends are drinking	
When I am at a pub or club	
When I am angry	
When I feel frustrated	
When I am worried	
When I feel upset	
When I feel nervous	
When I feel sad	
When I am watching TV	
When I am at lunch	
When I am on the way home from work	
When I am listening to music or reading	
When I am by myself	
When I have finished playing sport	
When I first arrive home	
When I feel like celebrating	

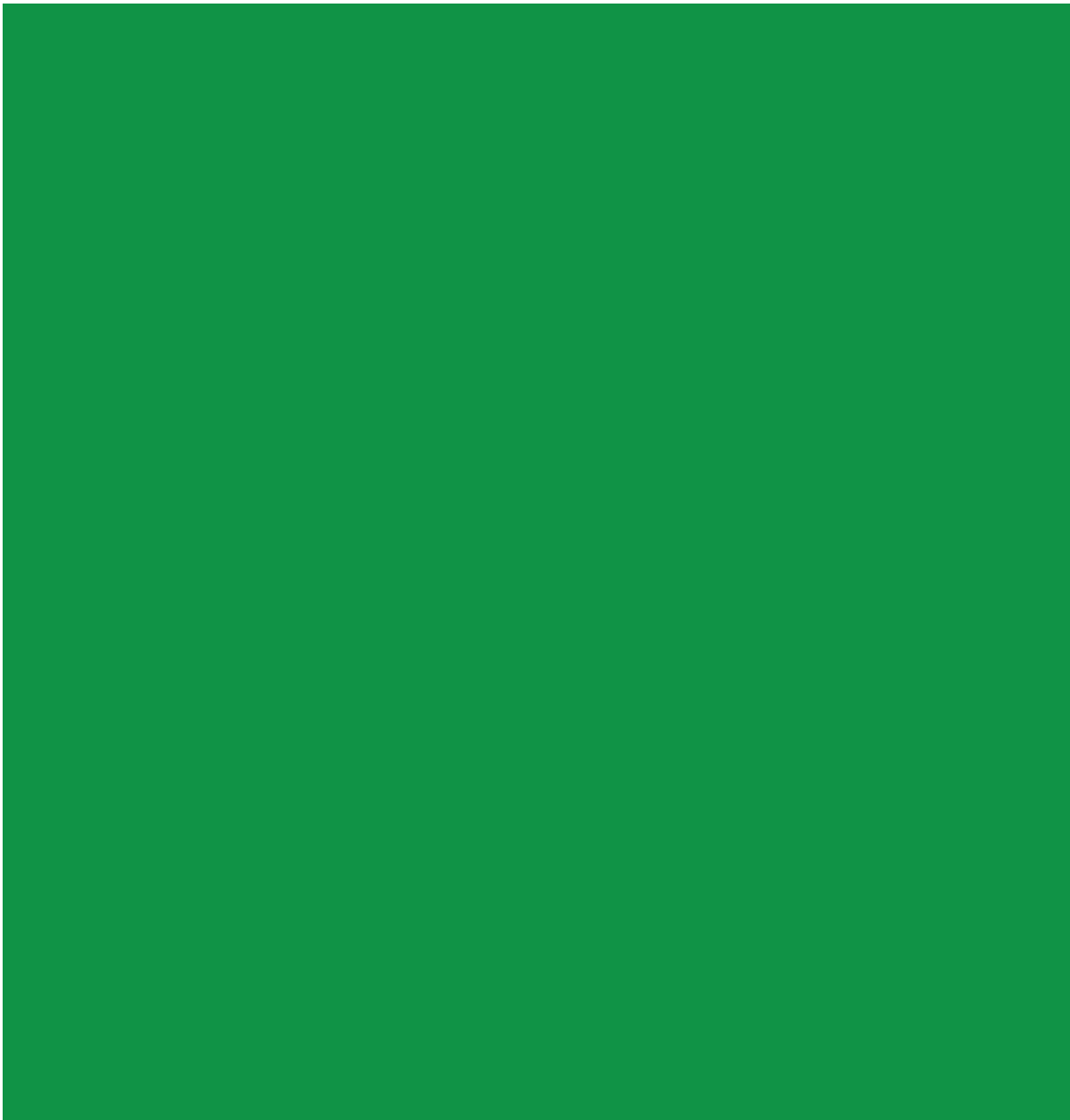
Next steps and final checks

Remind participants about the follow up on day 3, 7 and 21 after they are released.


Confirm preferred contact details for follow-up:

**Remember to give:
Unit card and participant generated plan.**






Appendix 5 Intervention tool postcard (Unit card) (V2_25Sept19)




One pint of beer/lager/cider = **between 2 & 3 units***

One 25 ml measure of spirit (whisky/brandy/dark rum) = **1 unit**




UK Chief Medical Office recommend that adults should not drink more than 14 units a week. Units should be spread evenly over 3 or more days.




175 ml glass of wine = **between 2 & 3 units***

250 ml glass of wine = **between 3 & 4 units***

750 ml bottle of wine = **between 9 & 10 units***



^aExact units will depend on the specific brand and corresponding ABV.



APPRASE

How many units are in your drink?

**If you would like more information or to
speak with someone please contact:**

www.changegrowlive.com

Drinkline

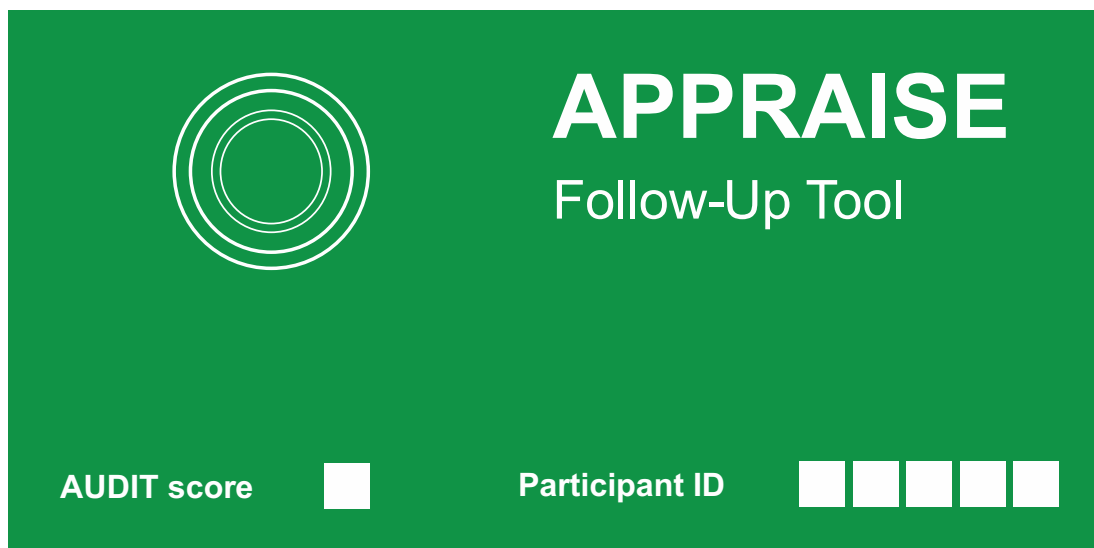
Helpline: 0300 123 1110

Offers advice and information for people worried
about their own drinking, and support to the family
and friends of people who are drinking.



APPRAISE

Appendix 6 Follow-up manual (V1_25 Sept19)



This resource has been produced as part of a research project in to extended Alcohol Brief Interventions in the prison setting.

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Element 1: Preliminary discussion

- Re-establish dialogue
- Affirm verbal consent, confidentiality, engagement rules and trust

Brief background

Participant's high-risk situations

Participant's goals

Participant's strategies

3

Intervention Follow-up 1: Day 3 post-liberation

- Check-in with participant
- Reconfirm previous goals (use previous page 3 as an aide)
- Work through elements 5 to 9 of the APPRAISE intervention

Date delivered

Time started

Time finished

Intervention Follow-up 2: Day 7 post-liberation

- Check-in with participant
- Reconfirm previous goals (use previous page 3 as an aide)
- Work through elements 5 to 9 of the APPRAISE intervention

Date delivered

Time started

Time finished

Intervention Follow-up 3: Day 21 post-liberation

- Check-in with participant
- Reconfirm previous goals (use previous page 3 as an aide)
- Work through elements 5 to 9 of the APPRAISE intervention

Date delivered

Time started

Time finished

Element 5: Situational appraisal and coping strategies

- Discuss strategies used to reduce consumption

**What has worked?
What can they build upon to maintain their goals?**

Examples discussed in session one of intervention:

- Switch type of drink to a lower strength
 - Smaller measures
 - Pace yourself
 - Only buy alcohol when needed
 - Have drink free days
 - Occupy yourself
 - Buy yourself something with the money saved.
- Discuss high-risk situations

**Have you been in any high-risk situations since
being released?
Can you tell me what strategies?
How did that go?
And what strategies will you use next time?**

Things to remember

- Ensure participant verbalises strategies
- Provide encouragement and praise
- Develop and model strategies with participants
- Allow participants to talk through the strategy and visualise themselves carrying it out successfully
- Encourage **rehearsal** of strategies that end in success

6

Element 6: Goal-setting

- Review goals set in session one of the intervention to make sure these are still relevant, specific and realistic.

Things to remember

- Set **realistic** and **short-term** sub-goals building on what they have already achieved
- Ensure these are achievable and provide confirmation they are able to achieve them

Element 7: Relapse

- Identify any relapse which has happened
- Discuss general reasons for relapse
- Encourage participants to verbalise specific situations which have or may resulted in relapse
- Identify appropriate strategies for relapse prevention

Remember if relapse occurs it is possible to find a more successful way of responding to that situation next time.

Things to remember

- Reiterate that relapse can happen
- Reiterate that relapse occurs typically as a response to stressful situations, social pressure and/or interpersonal conflict
- Remind them of their skills to address relapse and how they can learn to cope with difficult situations

Element 8: Self-evaluation and self-reinforcement

- Discuss if participant has used self-monitoring or rewarded their successes
- Reiterate the importance of these tools
- Praise attempts and successes
- Relapse should not be attributed to internal stable causes such as willpower or ability
- Tools such as the diary card might help them keep track
- Encourage them to reward themselves for achieving their goal.

Element 9: Culmination

- Summarise your conversation
- Remind the participants of the continuing follow-up.

Next steps and final checks

Remind participants about the continuing follow-up on day 3, 7 and 21 after they are released.

Is this still your preferred mode of contact? Confirm preferred contact details for follow-up (if different):





THE UNIVERSITY
of EDINBURGH



Appendix 7 APPRAISE training part 1

Course: <u>APPRAISE Training Part 1</u>	Duration: 2 hours
Aim: To enable staff to deliver brief interventions as part of the APPRAISE Research	
Objectives: The learners will: <ul style="list-style-type: none"> • Have an awareness and understanding of the APPRAISE RESEARCH • Understand AUDIT Scores and what they Mean • Have an awareness of the Readiness to Change Rulers & Risky Situation Assessments and why we use them • Be confident working with the APPRAISE booklet • Be able to deliver session one of the APPRAISE intervention 	

Timing	Content	Tutor Activity	Learner Activity	Resources
	Arrival and registration	Meet and greet	Arrive, sign in	Signing in sheet
5 mins	Introductions	Introduce the trainer(s) to the group). Ask the participants to briefly introduce themselves including their names, where they work	Listening, asking questions	
5 mins	Aims and Objectives	Read through aims and objectives	Listening	
5 mins	Introduction to APPRAISE	Play you tube clip	Listening, questions	https://www.youtube.com/watch?v=SQALC5MDsQs

5 mins	Introduction to AUDIT	Give AUDIT Questionnaire, Explain Scoring, & Remind them of the APPRAISE threshold of ≥ 8	Listening	https://uoe-my.sharepoint.com/:w/g/personal/apprstud_ed_ac_uk/ETpR_IKyQ25HlvxGXfmi8kBnBFsi_qnw54QtahvQBrbHw?e=cO4rYc
30 mins	APPRAISE booklet	Play you tube clip and hand out paper copy of booklet and APPRAISE Elements Facilitate discussion and feedback	Listening, asking questions	APPRAISE Booklet https://uoe-my.sharepoint.com/:b/g/personal/apprstud_ed_ac_uk/EQQkdsb4ovlNsnG5x49SdMYBjlqIpueKJcX5oLkF5TfZA?e=1bkyn4
				APPRAISE Elements sheet https://uoe-my.sharepoint.com/:w/g/personal/apprstud_ed_ac_uk/ET61B33L4bNPiNY1pSIcH2sBz9OoOSVGZwFIODEugKyLhQ?e=cJubew APPRAISE Booklet Video https://www.youtube.com/watch?v=Jc_1fZk4jBM

40 mins	Intervention example	Facilitate discussion and feedback	Listening, discussion, active participation	<p>APPRAISE Role Play Video</p> <p>https://www.youtube.com/watch?v=EMh1xXsTWno</p> <p>OR</p> <p>Facilitator role plays intervention delivery (this was done in Edinburgh)</p>
30 mins	Workshops	<p>Split the participants into pairs.</p> <p>Each pair to practice using APPRAISE booklet</p> <p>This last section of the training is to enable the participant to use the booklet effectively</p> <p>Staff will be at hand to answer any questions throughout the exercises</p> <p>Each participant is allocated 15</p>	Discussion, listening, asking, roleplay	<p>APPRAISE Booklet</p> <p>https://uoe-my.sharepoint.com/:b:/g/personal/apprstud_ed_ac_uk/EQQkdsb4ovlNsnG5x49SdMYBjlqIpueKJcX5oLkF5TfZA?e=1bkyn4</p>

		minutes then will be asked to swap places with partner		
5 mins	Evaluation and Close	Recap on aims and objectives. Answer any questions		Confirm Date, time for part 2

Appendix 8 Six-question Likert scale survey

The APPRAISE study: Training Evaluation Form

1. Following the training, my understanding of the APPRAISE study has improved significantly:	Strongly disagree	Disagree	Neither/Nor Agree	Agree	Strongly Agree
	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
2. Following the training, my understanding of the APPRAISE brief intervention has improved significantly:	Strongly disagree	Disagree	Neither/Nor Agree	Agree	Strongly Agree
	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
3. The training was what I expected:	Strongly disagree	Disagree	Neither/Nor Agree	Agree	Strongly Agree
	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
4. Following the training, I feel confident that I will be able to deliver the intervention using the APPRAISE booklet:	Strongly disagree	Disagree	Neither/Nor Agree	Agree	Strongly Agree
	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
5. The videos used during the training were helpful:	Strongly disagree	Disagree	Neither/Nor Agree	Agree	Strongly Agree
	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
6. Overall how would you rate the training?	Strongly disagree	Disagree	Neither/Nor Agree	Agree	Strongly Agree
	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Other comments:

.....

.....

.....

Date:

Site: HMP Edinburgh

HMP Durham

Appendix 9 Adapted Economic Form 90

Adapted from the Econ Form 90⁶⁰

HEALTH ECONOMICS QUESTIONNAIRE

We are asking for details of the last 6 months. Ensure that the date is calculated for each participant.

	Yes	No	Don't Know	No Answer
Since (date), have you been to Hospital?				
IF YES , how many times have you been to hospital during this time?				
	Yes	No	Don't Know	No Answer
Did you stay overnight for any of them?				
How many nights did you stay in total?				
Why were you admitted to hospital				
Notes:				

	Yes	No	Don't Know	No Answer
Are you registered with a GP?				
<u>IF YES</u> , since (date), have you been seen by a GP?				
<u>IF YES</u> , how many times have you been seen by a GP?				
	Yes	No	Don't Know	No Answer
<u>IF YES</u> , since (date), have you been seen by a practice nurse?				
<u>IF YES</u> , how many times have you been seen by a practice nurse?				
Notes:				

	Yes	No	Don't Know	No Answer
Since (date), have you been visited by a Community Psychiatric Nurse?				
<u>IF YES</u> , how many times have you been visited by a Community Psychiatric Nurse?				

Notes:

	Yes	No	Don't Know	No Answer
Since (date), have you been to any addiction services?				
<u>If YES</u>, what treatment have you received and who from?				
<u>If YES</u>, did you stay overnight for any visit?				
<u>If YES</u>, how many nights did you stay in total over all visits?				
Notes:				

	Yes	No	Don't Know	No Answer
Since (date), have you been to A&E?				
<u>If YES</u>, how many times have you gone to A&E?				
	Yes	No	Don't Know	No Answer

IF YES, were any of your visits to A&E while you were in prison?				
IF YES, how many times did you visit A&E while in prison?				
	Yes	No	Don't Know	No Answer
Did you ever use an emergency ambulance?				
IF YES, how many times have you used an emergency ambulance?				
Notes:				

	Yes	No	Don't Know	No Answer
Since (date), have you been working with a social worker?				
IF YES, how many times in total have you been in contact with them?				
IF YES, how many times has the contact been over the phone?				
IF YES, how many times has the contact been in person?				

IF YES, were any of your visits to A&E while you were in prison?				
IF YES, how many times did you visit A&E while in prison?				
	Yes	No	Don't Know	No Answer
Did you ever use an emergency ambulance?				
IF YES, how many times have you used an emergency ambulance?				
Notes:				

	Yes	No	Don't Know	No Answer
Since (date), have you been working with a social worker?				
IF YES, how many times in total have you been in contact with them?				
IF YES, how many times has the contact been over the phone?				
IF YES, how many times has the contact been in person?				

<p><u>IF YES,</u> how many times have they visited you in your home?</p>	
<p>Notes:</p>	

	Yes	No	Don't Know	No Answer
<p>Since (date), have you tried to access help from any other support services (e.g. any third-sector organisations, AA)?</p>				
<p><u>IF YES,</u> what other agencies have you been to for support?</p>				
<p><u>If agencies are identified,</u> how many times have you been to each?</p>				
<p>Notes:</p>				

<p>What are your current living arrangements?</p>	<p><input type="checkbox"/> Living alone</p> <p><input type="checkbox"/> Living with partner</p> <p><input type="checkbox"/> Living with other family members</p> <p><input type="checkbox"/> Living with friends</p> <p><input type="checkbox"/> Owned or rented property (you must be the tenant/joint tenant)</p> <p><input type="checkbox"/> Council property/Housing Association (you must be the tenant/joint tenant)</p> <p><input type="checkbox"/> Hostel</p> <p><input type="checkbox"/> Bed & Breakfast</p> <p><input type="checkbox"/> Residential Care/Shelter/Supported Accommodation</p> <p><input type="checkbox"/> Other (including rough sleeping, sofa surfing):</p> <p>Other: _____</p>			
<p>Since (date), have you been homeless at any point?</p> <p>N.b You are homeless if you have nowhere to stay and are living on the streets, but you can be homeless even if you have a roof over your head. You count as homeless if you are: staying with friends or family. Staying in a hostel, night shelter or B&B.²</p>	<p>Yes</p>	<p>No</p>	<p>Don't Know</p>	<p>No Answer</p>
<p>IF YES, how many nights were you homeless for?</p>				

<p><u>IF YES,</u> where did you sleep during this time?</p>	
<p>Since (date), have you stayed in any of the following?</p>	<p>Hostel <input type="checkbox"/></p> <p>B & B <input type="checkbox"/></p> <p>Residential Care/Shelter/Supported Accommodation <input type="checkbox"/></p>
<p><u>IF YES,</u> how many nights did you stay in each?</p>	<p>Hostel: _____</p> <p>B & B: _____</p> <p>Residential Care/Shelter/Supported Accommodation: _____</p>
<p>Notes:</p>	

	Yes	No	Don't Know	No Answer
<p>Are you currently employed? (If it is cash in hand work please put this in the notes section)</p>				
<p><u>IF NO,</u> have you been employed at any point since (date)?</p>				
<p><u>IF YES,</u> how many weeks have you been employed for between (date) and now?</p>				

IF YES, how many jobs have you had during this time?	
Notes:	

	Yes	No	Don't Know	No Answer
Do you currently receive any money from a public or government source (eg. Housing benefit, universal credit)				
IF YES, approximately how much do you receive in total per week?				
Notes:				

	Yes	No	Don't Know	No Answer
Since (date), have you been arrested?				
IF YES, how many times have you been arrested?				
	Yes	No	Don't Know	No Answer
Since (date), have you spent at least one night in custody?				

<p>IF YES, how many times have you spent at least one night in custody?</p>				
<p>If yes, was this police custody, prison, or both?</p>	<p><input type="checkbox"/> Police Custody</p> <p><input type="checkbox"/> Prison</p> <p><input type="checkbox"/> Both</p>			
<p>If yes, how many nights have you spent in custody?</p>	<p>Police Custody: _____</p> <p>Remand: _____</p> <p>Total: _____</p>			
	<p>Yes</p>	<p>No</p>	<p>Don't Know</p>	<p>No Answer</p>
<p>Since (date), have you been in court?</p>				
<p>If yes, how many times have you been in court?</p>				





Appendix 10 Participant Locator Form

A pilot study of an Alcohol Brief Intervention for male remand prisoners: The APPRAISE Study







Participant Locator Form - A pilot study of an Alcohol Brief Intervention for male remand prisoners: The APPRAISE

We aim use multiple methods to contact you for follow up. If you consent to us contacting you by the method, please tick yes and provide details.

HOW	YES	NO	DETAILS
Mobile 			
Text message 			
Voicemail 			
Email 			

A pilot study of an Alcohol Brief Intervention for male remand prisoners: The APPRAISE Study

HOW	YES	NO	DETAILS
Social Media 			
Home Address 			
Significant Other 			
Others involved in your care 			

Please sign that you agree to us contacting you through the details you have provided:

Appendix 11 Trial Steering Committee and Subcommittee terms of reference



A two-arm parallel-group individually randomised Prison Pilot study of a male Remand Alcohol Intervention for Self-efficacy Enhancement: the APPRAISE study

Terms of reference for the Trial Steering Committee (TSC)

These ToR will guide the scientific, administrative and operational direction of the APPRAISE Study.

Chief investigator (CI): Professor Aisha Holloway, School of Health and Social Science, the University of Edinburgh

Aims and objectives

The TSC has the primary aims of monitoring implementation of the APPRAISE feasibility trial, providing an independent assessment of the data analysis and determining if a future trial is merited.

The TSC has the following objectives:

- Provide overall supervision of the trial on behalf of the trial sponsor and funder and ensure it is conducted to rigorous standards.
- Comment on the progress of the trial and adherence to protocol.
- Consider new information of relevance to the research question.
- Provide advice, through the chair, to the chief investigator and trial funder on all appropriate aspects of the trial.
- Provide evidence to support any requests for extensions.

In addition to the TSC, the chair plus three members of the TSC (who specialise in health economics, statistics, and qualitative methodology and analysis) will form a subcommittee responsible for data monitoring and ethics.

The subcommittee has the following objectives:

- To monitor the data and make recommendation to the TSC on whether there are any ethical or safety reasons why the trial should not continue.
- To ensure that the safety, rights and well-being of the trial participants remain paramount.
- The subcommittee will consider the need for any interim analysis.
- The subcommittee is the only body who, if necessary, will have access to the unblinded data.

Meeting

The TSC will meet bi-annually. Members are able to join the meeting by teleconferencing. A meeting will be considered quorate when at least seven members are in attendance. Aisha Holloway and Victoria Guthrie will be responsible for calling, organising and minuting the meeting.

The subcommittee will meet prior to the TSC meeting. Members are able to join the meeting by teleconferencing. There is no quorate to this meeting. Jamie Smith will be the administrator of this group and responsible for minuting the meeting.

Duration: the group will function for the entire duration of the APPRAISE study.

Membership

Current membership is as follows:

Name	Position
Professor Frank Sullivan (Chair)	Professor of Primary Care Medicine, Director of Research
Professor Olivia Wu	Professor of Health Technology Assessment (Health Economics and Health Technology Assessment), Director of the HEHTA Research Unit
Dr Fergus Daly	Statistician at Frontier Science
Hon Professor Alex McMahon	Chair of the National Prisoner Health Care Network, Executive Director, Nursing, Midwifery, and Allied Health Professionals, Executive Lead, REAS and Prison Healthcare, NHS Lothian
Dr Justina Murray	CEO Scottish Families Affected by Drugs and Alcohol
Ms Joanne O'Connor	Director at Junction 42
Ms Jessica Davidson	Senior Clinical Forensic Charge Nurse
Dr James Carnie	Director of Research and Evaluation at the Scottish Prison Service
Mr Frank Reilly	Director at the Scottish Recovery Network
Ms Tracey Stewart	Community Alcohol-Related Damage Service (CARDS) Volunteer Co-ordinator at Rowan Alba
Dr Anne Whittaker	Associate Professor and Clinical Academic
Ms Alison Douglas	Chief Executive at Alcohol Focus Scotland
Mr Pete White	Chief Executive at Positive Prison? Positive Future
Mr Nigel Johnson	
Membership of the PMG: Aisha Holloway (CI); Dorothy Newbury-Birch (English Study Site Lead); Victoria Guthrie (Senior Research Fellow [SRF] and Project Manager); Natalie Connor (Research Associate, English Study Site); Jamie Smith (Research Assistant). Other members of the PMG as necessary.	
Note Membership of the group will be reviewed as appropriate and as required.	

Appendix 12 Consolidated Health Economic Evaluation Reporting Standards 2022 checklist

Topic	No.	Item	Location where item is reported
Title			
	1	Identify the study as an economic evaluation and specify the interventions being compared.	Chapter 7 Title 3.12 & 3.13
Abstract			
	2	Provide a structured summary that highlights context, key methods, results, and alternative analyses.	N/A as economics part of a larger study
Introduction			
Background and objectives	3	Give the context for the study, the study question, and its practical relevance for decision-making in policy or practice.	Chapter 7 Introductory Paragraphs
Methods			
Health economic analysis plan	4	Indicate whether a health economic analysis plan was developed and where available.	Section 7.2
Study population	5	Describe characteristics of the study population (such as age range, demographics, socioeconomic, or clinical characteristics).	Section 3.2
Setting and location	6	Provide relevant contextual information that may influence findings.	Section 3.1
Comparators	7	Describe the interventions or strategies being compared and why chosen.	3.12 & 3.13
Perspective	8	State the perspective(s) adopted by the study and why chosen.	Chapter 7 Introductory Paragraphs

Topic	No.	Item	Location where item is reported
Time horizon	9	State the time horizon for the study and why appropriate.	Chapter 7 Introductory Paragraphs
Discount rate	10	Report the discount rate(s) and reason chosen.	Chapter 7 Introductory Paragraphs
Selection of outcomes	11	Describe what outcomes were used as the measure(s) of benefit(s) and harm(s).	Not Applicable. Feasibility of cost estimation study.
Measurement of outcomes	12	Describe how outcomes used to capture benefit(s) and harm(s) were measured.	Not Applicable. Feasibility of cost estimation study.
Valuation of outcomes	13	Describe the population and methods used to measure and value outcomes.	Not Applicable. Feasibility of cost estimation study.
Measurement and valuation of resources and costs	14	Describe how costs were valued.	Tables 35–38
Currency, price date, and conversion	15	Report the dates of the estimated resource quantities and unit costs, plus the currency and year of conversion.	Chapter 7 Introductory Paragraphs
Rationale and description of model	16	If modelling is used, describe in detail and why used. Report if the model is publicly available and where it can be accessed.	Not Applicable. Feasibility of data collection study
Analytics and assumptions	17	Describe any methods for analysing or statistically transforming data, any extrapolation methods, and approaches for validating any model used.	Not Applicable. Feasibility of data collection study
Characterising heterogeneity	18	Describe any methods used for estimating how the results of the study vary for subgroups.	Not Applicable. Feasibility of data collection study

Topic	No.	Item	Location where item is reported
Characterising distributional effects	19	Describe how impacts are distributed across different individuals or adjustments made to reflect priority populations.	Not Applicable. Feasibility of data collection study
Characterising uncertainty	20	Describe methods to characterise any sources of uncertainty in the analysis.	Not Applicable. Feasibility of data collection study
Approach to engagement with patients and others affected by the study	21	Describe any approaches to engage patients or service recipients, the general public, communities, or stakeholders (such as clinicians or payers) in the design of the study.	Section 9.6
Results			
Study parameters	22	Report all analytic inputs (such as values, ranges, references) including uncertainty or distributional assumptions.	Not Applicable. Feasibility of data collection study
Summary of main results	23	Report the mean values for the main categories of costs and outcomes of interest and summarise them in the most appropriate overall measure.	Tables 40–43
Effect of uncertainty	24	Describe how uncertainty about analytic judgments, inputs, or projections affect findings. Report the effect of choice of discount rate and time horizon, if applicable.	Not Applicable. Feasibility of data collection study
Effect of engagement with patients and others affected by the study	25	Report on any difference patient/service recipient, general public, community, or stakeholder involvement made to the approach or findings of the study.	Section 7.4
Discussion			

Topic	No.	Item	Location where item is reported
Study findings, limitations, generalisability and current knowledge	26	Report key findings, limitations, ethical or equity considerations not captured, and how these could affect patients, policy, or practice.	Section 7.4
Other relevant information			
Source of funding	27	Describe how the study was funded and any role of the funder in the identification, design, conduct, and reporting of the analysis.	Funded by NIHR-PH
Conflicts of interest	28	Report authors conflicts of interest according to journal or International Committee of Medical Journal Editors requirements.	Submitted to funder

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