The Variability of Outcomes Used in Efficacy and Effectiveness Trials of Alcohol Brief Interventions: A Systematic Review

Shorter, G. W., Bray, J. W., Giles, E. L., O'Donnell, A. J., Berman, A. H., Holloway, A., Heather, N., Barbosa, C., Stockdale, K. J., Scott, S. J., Clarke, M., & Newbury-Birch, D. (2019). The Variability of Outcomes Used in Efficacy and Effectiveness Trials of Alcohol Brief Interventions: A Systematic Review. Journal of studies on alcohol and drugs, 80(3), 286-298. https://doi.org/10.15288/jsad.2019.80.286

Published in:
Journal of studies on alcohol and drugs

Document Version:
Peer reviewed version

Queen's University Belfast - Research Portal:
Link to publication record in Queen's University Belfast Research Portal

Publisher rights
Copyright 2019 Alcohol Research Documentation. This work is made available online in accordance with the publisher's policies. Please refer to any applicable terms of use of the publisher.

General rights
Copyright for the publications made accessible via the Queen's University Belfast Research Portal is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy
The Research Portal is Queen's institutional repository that provides access to Queen's research output. Every effort has been made to ensure that content in the Research Portal does not infringe any person's rights, or applicable UK laws. If you discover content in the Research Portal that you believe breaches copyright or violates any law, please contact openaccess@qub.ac.uk.
Shorter, G., Bray, J., Giles, E., O'Donnell, A., Berman, A., Holloway, A., Heather, N., Barbosa, C., Stockdale, K., Scott, S., Clarke, M., & Newbury-Birch, D. (2019). The variability of outcomes used in efficacy and effectiveness trials of alcohol brief interventions: A systematic review. *Journal of Studies on Alcohol and Drugs, 80*, 286-298. 
https://doi.org/https://doi.org/10.15288/jsad.2019.80.286

Number of tables 3
Number of figures 2

Mapping the variability of outcomes used in efficacy and effectiveness trials of alcohol brief interventions: A systematic review

Gillian W. Shorter BSc PhD 1,2*, Jeremy W. Bray BA MA PhD 3, Emma L. Giles BSc PhD 2, Amy J. O'Donnell BSc PhD 4, Anne H. Berman BSc MSc PhD 5,6, Aisha Holloway BSc PhD 7, Nick Heather BA MSc PhD 8, Carolina Barbosa PharmD MSc PhD 9, Kelly J. Stockdale BA MA PhD 10, Stephanie J. Scott BSc MA PhD 2, Mike Clarke BA DPhil 11, Dorothy Newbury-Birch BSc PhD 2

1 Institute of Mental Health Sciences, School of Psychology, Ulster University, Coleraine, UK
2 School of Health and Social Care, Teesside University, Middlesbrough, UK
3 Department of Economics, Bryan School of Business and Economics, University of North Carolina at Greensboro, Greensboro, NC, USA.
4 Institute of Health and Society, Newcastle University, Newcastle Upon Tyne, UK
5 Stockholm Health Care Services, Stockholm County Council, Stockholm, Sweden
6 Centre for Psychiatry Research, Department of Clinical Neuroscience, Karolinska Institutet, Stockholm, Sweden.
7 Nursing Studies, School of Health in Social Science, University of Edinburgh, Edinburgh, UK.
8 Faculty of Health and Life Sciences, Northumbria University, Newcastle upon Tyne, UK
9 Behavioral Health Economics Program, RTI International, Chicago, IL, USA.
10 School of Psychological and Social Sciences, York St. John University, York, UK
11 Northern Ireland Methodology Hub, Queen’s University Belfast, Belfast, UK

* Corresponding Author

Dr Gillian W Shorter
Address: Institute for Mental Health Sciences, School of Psychology, Ulster University, Cromore Road, Coleraine, BT52 1SA
Email: g.shorter@qub.ac.uk [amended]
Telephone: +4428 70 124025

Funding: We acknowledge the funding from Alcohol Research UK (Research Innovation Grant Number: R2016/04). The funding body had no role in the design of the study and the writing of this manuscript.

Registration: PROSPERO review registration (CRD42016047185) on 20 September 2016 prior to review start (Shorter et al., 2016a). This work is part of and informs the Outcome Reporting in Brief Intervention Trials: Alcohol project (ORBITAL) which aims to create a core outcome set or minimum data standard for alcohol brief intervention trials. This ORBITAL project arose from collaborations in the International Network on Brief Interventions for Alcohol and Other Drugs [INEBRIA] Research Measurement Standardization Special Interest Group. This was registered at the start of the project at the
COMET Initiative (Shorter et al., 2016b) and the protocol has been published (Shorter et al., 2017).
Abstract

Objective: To characterize recent alcohol brief intervention (ABI) efficacy and effectiveness trials; summarize outcomes; and show how variability in outcomes and reporting compromises the evidence base.

Method: A systematic review and narrative synthesis of articles from 10 databases were undertaken (Jan 2000-Nov 2017); study selection represented recent, readily available publications. Alcohol brief intervention definitions were informed by National Institute of Clinical Excellence (NICE) Public Health Guideline 24: Alcohol use disorders: prevention. The review was conducted using Centre for Reviews and Dissemination (CRD) guidance and pre-registered on PROSPERO (CRD42016047185). Seven a priori specified domains were used to classify outcomes: biomarkers, alcohol related outcomes, economic factors/resource use, health measures, life impact, intervention factors, and psychological/behavioral factors.

Results: The search identified 405 trials from 401 eligible papers. In 405 trials, 2641 separate outcomes were measured in approximately 1560 different ways. The most common outcomes used were number of drinks consumed in a week and frequency of heavy episodic drinking. Biomarkers were least frequently used. The most common primary outcome was weekly drinks. By trial type, the most frequent outcome in efficacy and effectiveness trials was frequency of heavy drinking.

Conclusions: Consumption outcomes predominated; however, no single outcome was found in all trials. This comprehensive outcome map for ABI effectiveness and efficacy trials can aid decision making in future trials. There was diversity of instruments, time points, and outcome descriptions in methods and results sections. Compliance with reporting guidance
would support data synthesis and improve trial quality. This review establishes need for a core outcome set/minimum data standard (COS) and supports the Outcome Reporting in Brief Interventions: Alcohol initiative (ORBITAL) to improve standards in the ABI field through a COS for effectiveness and efficacy randomized trials.

**Keywords:** Alcohol drinking; alcohol brief interventions, randomized controlled trials, outcome assessment, core outcome set, systematic review
Introduction

Alcohol brief interventions (ABIs) are key strategies to address problematic alcohol use worldwide (Coffield et al., 2001; National Institute for Health and Clinical Excellence, 2010; US Preventive Services Task Force, 2004; World Health Organisation [WHO], 2016). Numerous systematic reviews (Ballesteros et al., 2003; Ballesteros et al., 2004a; Ballesteros et al., 2004b; Beich et al., 2003; Bertholet et al., 2005; Kaner et al., 2018; O'Donnell et al., 2014) suggest ABIs in primary health care are effective, but these reviews report substantial outcome heterogeneity, limiting the strength of conclusions. Commentators have urged caution in making broad clinical practice recommendations as a result (Bernstein et al., 2009; Bernstein et al., 2010; Field et al., 2010; Heather, 2016; McCambridge & Saitz, 2017; Saitz et al., 2006; Saitz, 2010).

An avoidable problem is the diversity in definition and measurement of outcomes used. This reduces the ability to meaningfully synthesize available information. For example, in a recent and comprehensive review (Kaner et al., 2018), authors excluded 22 of 69 otherwise eligible studies due to outcome reporting issues. Differing outcomes across studies weakens meta-analyses of the efficacy and effectiveness of ABIs and contributes to research waste as not all articles can be used for the evidence base (Glasziou, 2014). Given the number of reviews mentioning outcome heterogeneity across all populations in which ABIs are now employed, it is no longer appropriate to dismiss this heterogeneity as a limitation, when it can and should be addressed.

To address outcome heterogeneity in ABI trials, future ABI studies should use a coherent, consistent set of outcomes, known as a core outcome set (COS). A COS is a feature of a mature research base, and many healthcare areas have developed, or are developing,
COS to support advances in their field (COMET Initiative, 2017). A COS reduces selective and inconsistent reporting in research trials, improves the quality of treatment guidance for a condition, and increases the number of studies synthesized in systematic reviews. Both the Consolidated Standards of Reporting Trials (CONSORT) (Moher et al., 2010) and the Standard Protocol Items: Recommendations for Intervventional Trials (SPIRIT) statements recommend COS use, and a formal process for COS development has been established by the Core Outcome Measures in Effectiveness Trials (COMET) Initiative (Williamson et al., 2017; Williamson et al., 2012). A COS is a minimum reporting standard, and does not restrict the measurement of additional outcomes. A comprehensive map of outcomes can support decision making on other outcomes to be measured alongside the COS; reducing a potential source of conflict in trial planning (Daykin et al., 2016; Daykin et al., 2017).

Recognizing the benefits an ABI COS could provide, the International Network on Brief Interventions for Alcohol and Other Drugs (INEBRIA) Research Measurement Standardization-Special Interest Group (IRMS-SIG) established the Outcome Reporting in Brief Intervention Trials: Alcohol (ORBITAL) project to derive a COS using COMET guidelines. This systematic review is a component of ABI COS development and follows the ORBITAL protocol (Shorter et al., 2017). Although numerous systematic reviews on ABI have been conducted, most have aimed to establish efficacy, effectiveness, and/or cost-effectiveness, and their included studies meet a restrictive set of eligibility criteria, including their pre-specified outcome of interest. No study to date has compiled all outcomes used across ABI studies. This paper fills this gap through a definitive catalogue of outcomes used in recent ABI trial literature. Such a catalogue is needed to a) map outcomes used to demonstrate efficacy and effectiveness in peer-reviewed, published ABI trials, b) demonstrate the variability in outcome type and measurement, c) highlight methodological issues in the
ABI field around outcomes and reporting, d) inform COS development, including identifying outcomes for a Delphi prioritization exercise (see Shorter et al., In Press), and e) support ABI trial protocol decision making on outcomes by trial area.

**Methods**

A review protocol was registered in advance on PROSPERO (CRD42016047185) (Shorter, et al., 2016a). Medline [OVID], EMBASE, PsycINFO [OVID], Health Management Information Consortium [HMIC], Cumulative Index to Nursing and Allied Health Literature [CINAHL], Allied and Complementary Medicine Database [AMED], Cochrane Library, ERIC [EBSCO], Web of Science, Google Scholar, Clinicaltrials.gov, and WHO International Clinical Trials Registry Platform [ICTRP] databases were searched to identify trials published between January 2000-November 2017 in peer-reviewed journals. This date range provided balance between capturing an extensive evidence base and reflecting the current state of ABI research. We focus on peer-reviewed publications as readily available in the public domain and reflect the work of the COS target audience (policy, practice, and research). Core search concepts related to three domains: alcohol use; brief interventions; and randomized trials. Terms were coupled with relevant MeSH/thesaurus terms, truncated as appropriate, and variant spellings were used to identify useful records (Supplementary Material A contains the OVID search which was adapted for other databases).

Eligible studies were individual or cluster randomized trials focused on efficacy or effectiveness of ABIs designed to reduce alcohol consumption published in peer-reviewed journals. Trials that did not analyze outcomes by randomized arm were excluded (e.g. subsample analysis only). Papers with the same trial registration number were included if they assessed different outcomes in each. Specific search parameters are described below.
Population: Current drinkers (at least one drink in the past year) who were aged 16 years or above. Trials of drinkers aged 15 years or below were excluded, as were trials including individuals seeking treatment for alcohol problems, following related UK NICE guidance (National Institute for Health and Clinical Excellence, 2010).

Intervention: ABIs were defined as those suitable for drinkers not seeking treatment for an alcohol problem but who are identified by screening as having, or being at risk of, problems from their alcohol use (National Institute for Health and Clinical Excellence, 2010). This definition covers brief advice and extended brief interventions, delivered once or more frequently. An ABI should assess an individual’s alcohol use and provide feedback on their alcohol assessment. Trials including a multicomponent intervention arm or where one or more intervention components addressed non-alcohol related health behaviors (e.g., smoking cessation) were included if alcohol intervention components and outcomes could be clearly distinguished.

Comparator: Comparators could be any active or control intervention.

Outcomes: All outcomes analyzed by randomized arm were extracted including detail of how the outcome was defined and measured if possible. This was used to estimate the variability in outcome measurement, i.e. to what extent an outcome in one paper was exactly the same as in another paper (what the outcome represents, how it was measured and scored, and time period referred to). Other extracted information included: number and nature of sample randomized (sex, age, and population), trial details, including region, number of trial arms, trial arm composition, trial type (efficacy/effectiveness/not reported), and details of follow up timing. These were summarized either as number (%) or mean (SD) of trials.
included with indication of missing data in the total number. Broad indicators of trial reporting quality were included: stating ‘trial’ in the title or including a participant flow chart in line with early CONSORT guidance (Begg et al., 1996). Where study information was not provided this was stated. Effectiveness and efficacy ABI reviews often contact authors for missing data, we did not do so as our aim was to highlight where reporting falls short to improve standards in the field like other, similar high quality methodological reviews (Harman et al., 2017; Riddle et al., 2008; Thornley & Adams, 1998).

A taxonomy was created to map outcomes under seven domains: alcohol related outcomes; biomarkers; health measures; economic factors/social impacts; psychological/behavioral factors; life impact; and intervention factors. This was influenced by a range of sources. The first draft was informed by a presentation at the COMET V meeting in Amsterdam (attended by GWS in September 2016), since published in Dodd et al. (2018). However, given the ABI topic area is not directly concerned with physical pathology, many clinical factors in this taxonomy were irrelevant (e.g. musculoskeletal outcomes), whereas other outcomes were not specific enough (e.g. emotional functioning/wellbeing). Other sources included the Outcome Measurement Sets for Clinical Trials [OMERACT] filter (Boers et al., 2014); this was helpful to derive core areas such as death, life impact, resource use, or pathophysiological manifestations, but was too broad to capture outcomes relevant to ABIs. The Patient-Reported Outcomes Measurement Information System [PROMIS] (Cella et al., 2007) provided elaboration to describe some outcomes in ABI trials (anxiety, depression, or sleep disturbance) but there were classification limitations, and some outcomes (e.g. PROMIS alcohol use questionnaire) were absent from ABI papers. We drew upon health economic reviews to inform the economic outcomes domain (Barbosa et al., 2015; Barbosa et al., 2010; Bray et al., 2011). Outcome data extracted from ABI trials were
used to refine the taxonomy further. GWS created the taxonomy, this was then refined by others (NH, DNB, JWB, AHB, CB, and ELG).

Search results were downloaded to EndNote version X7 and de-duplicated. GWS screened all titles and abstracts of papers and excluded those that did not meet the inclusion criteria. DNB checked 28% of these for accuracy; discrepancies were resolved by discussion. All full text versions of potentially eligible papers were reviewed by GWS, and all double-screened by one of ELG, DNB, JWB, AJOD, AHB, and AH; discrepancies were resolved by discussion. Extraction forms were piloted by GWS and KJS. All data were extracted by GWS, and all extracted data cross-checked for accuracy by at least one of ELG, DNB, SJS, KJS, JWB, AJOD and AHB. Data were presented from all trials, and split by population (‘primary care’, ‘emergency department’, ‘University/College’, ‘general population’ (i.e. a general adult sample not selected as having specific characteristics), ‘other healthcare’, and ‘other’ populations (including workplaces or job related populations (n=14), veteran populations (n=19), community sample of persons with an intellectual disability (n=1), homeless population (n=1), criminal justice populations (n=14), licensed premises (n=1), sports clubs (n=1), and young people aged 16+ years (n=6)). A PRISMA checklist is in Supplementary Material B.

Results

Searches identified 33,134 papers after de-duplication to be screened by title and abstract for eligibility. Exclusion at title and abstract stage reflected unambiguous violation of the above PICO (population, intervention, comparator, outcomes) on the basis of topic area (i.e. not alcohol) or a known alcohol treatment sample (such as Project MATCH). Any unclear matches were referred to full text assessment for closer inspection; 1,612 papers were retrieved for full text evaluation against PICO criteria, and 401 were deemed eligible (Figure
1). The 401 included papers covered 405 individual trials (some papers reported two trials), representing 182,272 randomized participants in total (see Supplementary Material C for included papers)

<<<Figure 1>>> The mean trial size was 450 individuals (range=12-7,935). Typically, higher numbers were randomized in ‘primary care’, ‘emergency department’, and ‘general population’ samples compared to the remaining populations (Table 1). There were slightly more males than females on average (mean % male=56.2; SD=28.1); highest in the ‘other’ population. Most trials were conducted in North America (60.7%); this was particularly evident in the ‘University/College’ population with 81.1% of trials from this region. Two-arm trial designs predominated, with trials in the ‘emergency department’ and ‘University/College’ populations more likely to have more than two arms. Around 83% of trials had a non-ABI control group. ‘Other’ populations were most likely to have a non-ABI control (91.2%) and ‘General population’ were least likely (75.0%). More trials were declared by their authors as efficacy trials (52.8%) compared to effectiveness trials (42.0%). Twenty-one trials did not state their type. Only ‘University/College’ populations had more effectiveness trials (56.1%) than efficacy trials (40.2%).

<<<Table 1>>> Just over half the trials indicated they were a trial in the paper’s title (52.1%); 63.7% included a flow chart of participants through the trial. ‘University/College’ populations were least likely to report these elements, with ‘general population’ trials more likely to state they were a trial (67.2%), and ‘other healthcare’ populations more likely to include a flow chart (78.2%). Broadly similar percentages had two or three data collection waves (42% and 38%...
respectively). Longer-term follow-up of two or more years was more likely in ‘primary care’ (n=7; 14%). Short-term follow-up was more likely in ‘University/College’ samples. Overall, trialists most often selected three-month intervals for follow-up (e.g. three or six months). Over time, there was a general increase in the number of ABI trials published per year. The largest number were published in 2014. The number of trials per year is given in Figure 2.

Outcomes

Overall, 2,641 outcomes were extracted from 405 trials. Only 285 trials stated if their outcomes were primary or secondary. The mean number of outcomes per trial was 6.5 (ranging from 1-56); highest in ‘primary care’, and lowest in ‘University/College’ samples. On average, there were two primary, and four secondary outcomes reported in the included trials. Most trials had at least one alcohol related outcome measure. The highest percentage of trials with at least one health outcome was in the ‘primary care’ or ‘other healthcare’ population, least likely in the ‘University/College’ population. Economic factors or social impacts were most likely in the ‘primary care’ or ‘emergency department’ population. Psychological factors were found in around 28% of trials, most commonly in ‘other healthcare’ populations. Life impact outcomes were present in 56 trials. Less than 10% of trials looked at intervention factors and ‘University/College’ samples were more likely to have one outcome of this type. Biomarkers were infrequently used: only 13 trials had measured at least one biomarker; more likely in ‘primary care’ or ‘other’ populations.

Alcohol related outcomes

Alcohol related outcomes include those connected to the amount or pattern of alcohol consumption, those related to the comorbid use of other substances and those reflecting
substance use disorder symptomology. As such, it is broader than just alcohol consumption measures but we have retained the term “alcohol related outcomes” for ease of exposition and to maintain consistency with our protocol (Shorter et al., 2017) and Delphi study (Shorter et al., In Press). In the 405 trials, there were 1,456 alcohol related outcomes measured in 744 different ways (Table 2). The most commonly reported alcohol related outcome variables were frequency of heavy drinking (n=213), weekly drinks (n=205), alcohol related problems or consequences (n=190), typical quantity (n=137), typical frequency (n=117), and hazardous or harmful drinking (n=111). Many of the infrequently-measured outcomes were also the most diversely measured. An exception included ‘at risk drinking’ (which measures risk derived from publicly-available recommendations such as weekly or single episode limits).

By population, ‘primary care’ trials were most likely to report weekly drinks, frequency of heavy drinking, and at-risk drinking. This was somewhat similar to the ‘general population’, ‘emergency department’ and ‘University/College’ populations, which often measured weekly drinks, frequency of heavy drinking, and alcohol-related problems or consequences. The majority of trials that measured blood alcohol concentration were in the ‘University/College’ population. ‘Other healthcare’ populations often measured typical and heavy drinking frequencies, and hazardous and harmful drinking. Frequency of heavy drinking was the most commonly reported outcome in both efficacy and effectiveness trials, with the number of drinks consumed in a week the most frequent primary outcome.

<<<Table 2>>>

Other outcomes

In total, 32 biomarker outcomes were reported across the 405 trials (Table 3). Of these, the most commonly reported was gamma-glutamyltransferase (GGT). Biomarkers were only found in ‘primary care’, ‘other healthcare’, and ‘other’ populations. The most frequent
biomarker in efficacy trials was GGT, in effectiveness trials it was Carbohydrate-deficient transferrin (CDT). GGT and CDT tied as the most common primary outcome.

In the economic factors/social impacts domain, the most commonly reported outcomes were driving related offences and hospitalizations. This domain includes some overlap with measures of alcohol related consequences in the alcohol related outcome domain, but measures in this domain are intended to assess social costs and impacts, not to assess the possibility of a diagnosable alcohol disorder. In ‘primary care’, the most commonly reported economic factors/social impacts outcomes were driving-related offences, hospitalizations, other criminal justice use, or other healthcare use. For ABI trials set in the ‘emergency department’, the most common were seeking alcohol treatment, driving-related offences, and emergency healthcare use. In ‘other healthcare’ populations, the most commonly-assessed economic variable was that of provider intervention costs. In ‘other’ populations, given the composition of this group, other criminal justice use was most common. Economic factors/social impacts measures were not commonly reported by ‘general population’ or ‘University/College’ ABI trials. The intervention cost to the provider was the most common economic factors/social impacts measure for efficacy trials. Driving related offences was the most common measure in effectiveness trials, and the most reported primary outcome.

Health outcomes most commonly reported were alcohol-exposed pregnancy factors, psychological health measures, sexual violence or coercion, and severity of depression symptoms. In ‘primary care’, cardiac factors, psychological health, and physical health were most commonly reported. In ‘general population’ samples, alcohol-exposed pregnancy factors or severity of depression were more commonly reported. Sleep disruption was only measured in ‘University/College’ ABI trials. ‘Other healthcare’ populations most commonly
reported alcohol-exposed pregnancy factors. The most frequent efficacy outcome in this
domain was psychological health; the most common outcome in effectiveness trials was
alcohol-exposed pregnancy factors. The most commonly reported outcome from the
intervention factors domain was intervention satisfaction; true for both effectiveness and
efficacy trials. ABI trials in ‘University/College’ and ‘general population’ samples were more
likely to ask participants about this outcome.

In the domain of psychological and behavioral factors, the most commonly reported
outcomes across all trials were drinking refusal self-efficacy, alcohol outcome expectancies,
risky behaviors, and readiness to change. ABI trials in the ‘primary care’ and ‘emergency
department’ populations were least likely to measure these outcomes. By contrast,
‘University/College’ samples were particularly likely to measure the perception of others’
drinking, for example, the typical quantity drunk by a student at their institution. For ‘general
population’ samples, drinking refusal self-efficacy and readiness to change were most
common. In ‘other healthcare’ populations, risky behaviors were the most commonly
reported; these include aspects such as sex without effective contraception. Finally, in ‘other’
populations, anger and aggression, drinking refusal self-efficacy, other psychological factors,
and readiness to change were the most commonly reported outcomes. The most frequent
measure in efficacy trials was readiness to change; for effectiveness trials it was perception of
others’ drinking. The most common primary outcome for both was engagement in risky
behaviors. Life impact measures were most commonly role functioning or relationship factors
or quality of life. The former was most common in effectiveness trials (and as a primary
outcome), the latter the most common for efficacy trials.

<<<Table 3>>>
Discussion

This review is the first to go beyond stating outcome heterogeneity as a weakness in ABI systematic reviews; it quantifies the heterogeneity and inconsistency in outcomes reported in effectiveness and efficacy trials of ABIs. Overall, there were 2,641 outcomes measured in approximately 1,560 different ways, truly a “Tower of Babel”. The estimated 1,560 different ways outcomes were measured may be a conservative guess of the true variability given the lack of precision on how outcomes were measured. The variation in the outcomes used and reported across ABI trials reflects similar reviews conducted in different research areas (Harman et al., 2017). For the ABI field, the substantial heterogeneity represents an important challenge. Meta-analyses will continue to be compromised as they cannot draw on all evidence to decide whether ABIs work as intended. Just over half (53%) measured the most common consumption measure frequency of heavy drinking; this creates a considerable conflict between the drive for inclusion of all studies meeting criteria in high quality systematic reviews, and the ability to include all studies in the meta-analysis.

Determining efficacy or effectiveness depends on outcomes measured, and therefore all ABI trial papers should contain sufficient detail on outcome measurement. One way this may affect meta-analyses is through the combination of an outcome (e.g. weekly drinks) which hides considerable variability. For example, “weekly drinks” may refer to an average week, a typical week, or the last week. It may refer to a typical week in the past month, 28 days, 90 days, six months, or since last measurement. The definition of drink may be specified or left to the respondent. Weekly drinks may be reported directly or calculated based on other information in a range of different questionnaires. We can calculate some differences to be equivalent, but some measure genuine differences and their combination compromises the validity of estimates. At a minimum trials should report a) what the outcome is, b) the
question or questionnaire used to measure and how this is used (e.g. scale score, or the binary above and below a cut-off point), c) measure of aggregation such as mean value or mean individual difference, and d) time point (e.g. 1, 4, and 8 weeks post intervention).

Some trials did not specify whether their outcomes were primary or secondary outcomes. This could be because the trial was a pilot study and specification may not be required (Eldridge et al., 2016), or it might be stated in a trial registry. However, excluding this from reporting is problematic (Begg, et al., 1996; Moher, et al., 2010). In addition, although one might expect trials to have only one primary outcome, we found, of those who specified, the average was two primary outcomes. This was an under-estimate of the average because some papers only reported secondary outcomes; their primary outcome(s) were in other papers with the same trial registration number. The correct interpretation of secondary outcomes is ‘through’ the primary analysis on the premise that, if the primary outcome is positive, then secondary outcomes can help to understand how the ABI worked. The secondary designation may also be useful for outcomes more distal on the causal pathway that reduced drinking would be expected to change. If the primary outcome is neutral, the secondary outcomes are hypothesis-generating. If the primary outcome is ‘negative’, the secondary outcomes provide insight into how the treatment caused harm (Freemantle, 2001). If change is shown in some primary outcomes but not others, interpretation can become difficult and it may be a challenge to state the ABI brought about change. To improve the aggregation of trials into the evidence base, outcomes (from a COS or otherwise) should be detailed, identified as primary or secondary with a clear statistical analysis plan, well reported in results sections which include point estimates and variability around estimates, and follow reporting guidance.
Alcohol related outcomes, particularly consumption outcomes, were the predominant outcomes measured in ABI trials. Although some have called for an increase in biomarkers in ABI trials (Kypri, 2007) this call has not been heeded; most outcomes were self-reported. ABI effectiveness or efficacy meta-analyses rely on the outcomes reported without validating them against objective measures (Moyer et al., 2002), exacerbating the problem of outcome heterogeneity in ABI trials. Our review provides the first systematic and quantifiable evidence to support previous calls for standard definitions of ABI outcomes to compare across studies (Bernstein et al., 2010).

Despite efforts to identify literature from across the globe, most trials were from North American or European countries. This may reflect the predominance of publishing or funding opportunities available to those researchers, the high levels of hazardous and harmful use of alcohol in these countries (Rehm et al., 2009), or be a consequence of the pre-specified databases searched. We attempted to minimize English language bias and improve the quality of the review by including studies reported in languages other than English (Moher et al., 2003). The searching was largely conducted in English, and our ability to extract data from articles in languages other than English was limited, as shown in the CONSORT flowchart (Figure 1). Although focusing on peer-reviewed literature may have also limited the number of non-English articles included, it is in keeping with our intention to focus on those articles that are likely to be most accessible and influential for many decision makers. Our searches of the grey literature, which constitute a separate part of our PROSPERO-registered systematic review not reported here, will be one opportunity to explore how improving access to a wider range of literature from low-resource settings or from reports in languages other than English, may influence the evidence base. This limitation is likely to have shown additional heterogeneity in findings, as the number of valid trials increased.
There was also a predominance of efficacy trials in the included studies, and attention should turn towards effectiveness trials within the different populations. Efficacious interventions may not be effective in routine practice (McCambridge & Saitz, 2017). Some trials did not specify their trial approach as either efficacy or effectiveness; although this may be a consequence of challenges of specification across the efficacy to effectiveness continuum (Heather, 2014). Short-term follow up was common, as reported by other systematic reviews (Moyer, et al., 2002). This is perhaps expected given effect sizes tend to be larger at early follow up, and there are concerns about the longitudinal effects of ABI (Donoghue et al., 2014). The predominant follow-up interval was around three months between data collection points. With around 20% of studies with four or more follow-up points, there is a balance between minimizing loss to follow-up, timely collection of only important information, and respondent burden (Lin et al., 2012).

By synthesizing outcome selection, this review offers the opportunity to consider outcome choice and the implications for the ABI field; other healthcare areas have noted the importance of design in attrition (Kilburn et al., 2014). Others have considered respondent burden (Cunningham et al., 1999; Kypri, 2007); as the number of outcomes reported was 56 in one trial, this may need careful consideration. Decision making around which outcomes to use for particular trials can be assisted by this outcome map, broken down by research area, effectiveness/efficacy, and primary/secondary/other outcomes. The structure of this outcome map was the process of discussion between co-authors, and we recognize that other structures of categorization may also exist.
This review highlights the importance of a COS for efficacy or effectiveness ABI randomized trials. This review will contribute to the efforts to establish a COS using high quality, established methodologies (Williamson, et al., 2017), which will improve and standardize reporting in the ABI field. This review also informs a preliminary list of outcomes for a related e-Delphi prioritization exercise (Shorter et al., In Press) and for discussion at the consensus meeting as outlined in the ORBITAL protocol (Shorter et al., 2017). We aimed to better understand how the extent of variability and reporting of outcomes compromises the evidence base and have conclusively shown this variability is considerable, and reporting is incomplete. The ability of users of ABI research to compare and understand findings is restricted because we do not know what exactly was measured and how, nor can we confidently compare seemingly alike outcomes. We did not seek to improve the completeness of the data by contacting the original authors, but used the incompleteness (contrary to usual systematic review practice) as a tool to highlight shortcomings in the field. We must improve issues of reporting and methodological quality to advance the field; the ABI evidence base cannot move from middle age to more established without it (Babor et al., 2007).

Acknowledgements

We would like to thank the INEBRIA Measurement Standardization SIG and participants at the INEBRIA 2016 conference workshop on the topic for their useful commentary/feedback.
References

Babor, T. F., McRee, B. G., Kassebaum, P. A., Grimaldi, P. L., Ahmed, K., & Bray, J. (2007). Screening, Brief Intervention, and Referral to Treatment (SBIRT): toward a public health approach to the management of substance abuse. Substance Abuse, 28(3), 7–30. doi:10.1300/J465v28n03_03. Medline

Ballesteros, J., Ariño, J., González-Pinto, A., & Querejeta, I. (2003). [Effectiveness of medical advice for reducing excessive alcohol consumption. Meta-analysis of Spanish studies in primary care]. Gaceta Sanitaria, 17(2), 116–122. doi:10.1016/S0213-9111(03)71708-7. Medline

Ballesteros, J., Duffy, J. C., Querejeta, I., Ariño, J., & González-Pinto, A. (2004a). Efficacy of brief interventions for hazardous drinkers in primary care: systematic review and meta-analyses. Alcoholism, Clinical and Experimental Research, 28(4), 608–618. doi:10.1097/01.ALC.0000122106.84718.67. Medline

Ballesteros, J., González-Pinto, A., Querejeta, I., & Ariño, J. (2004b). Brief interventions for hazardous drinkers delivered in primary care are equally effective in men and women. Addiction (Abingdon, England), 99(1), 103–108. doi:10.1111/j.1360-0443.2004.00499.x. Medline

Barbosa, C., Cowell, A., Bray, J., & Aldridge, A. (2015). The cost-effectiveness of alcohol screening, brief intervention, and referral to treatment (SBIRT) in emergency and outpatient medical settings. Journal of Substance Abuse Treatment, 53, 1–8. doi:10.1016/j.jsat.2015.01.003. Medline

Barbosa, C., Godfrey, C., & Parrott, S. (2010). Methodological assessment of economic evaluations of alcohol treatment: what is missing? Alcohol and Alcoholism (Oxford, Oxfordshire), 45(1), 53–63. doi:10.1093/alcalc/agg067. Medline

Begg, C., Cho, M., Eastwood, S., Horton, R., Moher, D., Olkin, I., et al. (1996). Improving the quality of reporting of randomized controlled trials. The CONSORT statement. JAMA, 276(8), 637–639. doi:10.1001/jama.1996.03540080059030. Medline

Beich, A., Thorsen, T., & Rollnick, S. (2003). Screening in brief intervention trials targeting excessive drinkers in general practice: systematic review and meta-analysis. BMJ (Clinical Research Ed.), 327(7414), 536–542. doi:10.1136/bmj.327.7414.536. Medline

Bernstein, E., Bernstein, J. A., Stein, J. B., & Saizt, R. (2009). SBIRT in emergency care settings: are we ready to take it to scale? Academic Emergency Medicine, 16(11), 1072–1077. doi:10.1111/j.1553-2712.2009.00549.x. Medline

Bernstein, J. A., Bernstein, E., & Heeren, T. C. (2010). Mechanisms of change in control group drinking in clinical trials of brief alcohol intervention: implications for bias toward the null. Drug and Alcohol Review, 29(5), 498–507. doi:10.1111/j.1465-3362.2010.00174.x. Medline

Bertholet, N., Daeppen, J.-B., Wietlisbach, V., Fleming, M., & Burnand, B. (2005). Reduction of alcohol consumption by brief alcohol intervention in primary care: systematic review and meta-analysis. Archives of Internal Medicine, 165(9), 986–995. doi:10.1001/archinte.165.9.986. Medline

Boers, M., Kirwan, J. R., Wells, G., Beaton, D., Gossec, L., d’Agostino, M. A., et al. (2014). Developing core outcome measurement sets for clinical trials: OMERACT filter 2.0.
Bray, J. W., Cowell, A. J., & Hinde, J. M. (2011). A systematic review and meta-analysis of health care utilization outcomes in alcohol screening and brief intervention trials. Medical Care, 49(3), 287–294. doi:10.1097/MLR.0b013e318203624f. Medline

Cella, D., Yount, S., Rothrock, N., Gershon, R., Cook, K., Reeve, B., et al., & the PROMIS Cooperative Group. (2007). The Patient-Reported Outcomes Measurement Information System (PROMIS): progress of an NIH Roadmap cooperative group during its first two years. Medical Care, 45(5, Suppl 1), S3–S11. doi:10.1097/01.mlr.0000258615.42478.55. Medline

Coffield, A. B., Maciosek, M. V., McGinnis, J. M., Harris, J. R., Caldwell, M. B., Teutsch, S. M., et al. (2001). Priorities among recommended clinical preventive services. American Journal of Preventive Medicine, 21(1), 1–9. doi:10.1016/S0749-3797(01)00308-7. Medline

COMET Initiative. (2017). How to search the COMET Initiative database. Retrieved from https://stream.liv.ac.uk/eqyg4t36

Cunningham, J. A., Ansara, D., Wild, T. C., Toneatto, T., & Koski-Jännes, A. (1999). What is the price of perfection? The hidden costs of using detailed assessment instruments to measure alcohol consumption. Journal of Studies on Alcohol, 60(6), 756–758. doi:10.15288/jsa.1999.60.756. Medline

Daykin, A., Selman, L. E., Cramer, H., McCann, S., Shorter, G. W., Sydes, M. R., et al. (2016). What are the roles and valued attributes of a Trial Steering Committee? Ethnographic study of eight clinical trials facing challenges. Trials, 17(1), 307. doi:10.1186/s13063-016-1425-y. Medline

Daykin, A., Selman, L. E., Cramer, H., McCann, S., Shorter, G. W., Sydes, M. R., et al. (2017). ‘We all want to succeed, but we’ve also got to be realistic about what is happening’: an ethnographic study of relationships in trial oversight and their impact. Trials, 18(1), 612. doi:10.1186/s13063-017-2305-9. Medline

Dodd, S., Clarke, M., Becker, L., Mavergamas, C., Fish, R., & Williamson, P. R. (2018). A taxonomy has been developed for outcomes in medical research to help improve knowledge discovery. Journal of Clinical Epidemiology, 96, 84–92. doi:10.1016/j.jclinepi.2017.12.020. Medline

Donoghue, K., Patton, R., Phillips, T., Deluca, P., & Drummond, C. (2014). The effectiveness of electronic screening and brief intervention for reducing levels of alcohol consumption: a systematic review and meta-analysis. Journal of Medical Internet Research, 16(6), e142. doi:10.2196/jmir.3193. Medline

Eldridge, S. M., Chan, C. L., Campbell, M. J., Bond, C. M., Hopewell, S., Thabane, L., & Lancaster, G. A., & the PAFS consensus group. (2016). CONSORT 2010 statement: extension to randomised pilot and feasibility trials. Pilot and Feasibility Studies, 2(1), 64. doi:10.1186/s40814-016-0105-8. Medline

Field, C. A., Baird, J., Saizt, R., Caetano, R., & Monti, P. M. (2010). The mixed evidence for brief intervention in emergency departments, trauma care centers, and inpatient hospital settings: what should we do? Alcoholism, Clinical and Experimental Research, 34(12), 2004–2010. doi:10.1111/j.1530-0277.2010.01297.x. Medline
Freemantle, N. (2001). Interpreting the results of secondary end points and subgroup analyses in clinical trials: should we lock the crazy aunt in the attic? BMJ (Clinical Research Ed.), 322(7292), 989–991. doi:10.1136/bmj.322.7292.989, Medline

Glasziou, P., Altman, D. G., Bossuyt, P., Boutron, I., Clarke, M., Julious, S., et al. (2014). Reducing waste from incomplete or unusable reports of biomedical research. Lancet, 383(9913), 267–276. doi:10.1016/S0140-6736(13)62228-X, Medline

Harman, N. L., James, R., Wilding, J., & Williamson, P. R., & the SCORE-IT study team. (2017). SCORE-IT (Selecting Core Outcomes for Randomised Effectiveness trials In Type 2 diabetes): a systematic review of registered trials. Trials, 18(1), 597. doi:10.1186/s13063-017-2317-5, Medline

Heather, N. (2014). The efficacy-effectiveness distinction in trials of alcohol brief intervention. Addiction Science & Clinical Practice, 9(1), 13. doi:10.1186/1940-0640-9-13, Medline

Heather, N. (2016). Spreading alcohol brief interventions from health care to non-health care settings: Is it justified? Drugs Education Prevention & Policy, 23(5), 359–364. doi:10.1080/09687637.2016.1187113

Kaner, E. F. S., Beyer, F. R., Muirhead, C., Campbell, F., Pienaar, E. D., Bertholet, N., et al. (2018). Effectiveness of brief alcohol interventions in primary care populations. Cochrane Database of Systematic Reviews, 2(2), CD004148. doi:10.1002/14651858.CD004148.pub4, Medline

Kilburn, L. S., Banerji, J., & Bliss, J. M., & the NCRI Breast Clinical Studies Group. (2014). The challenges of long-term follow-up data collection in non-commercial, academically-led breast cancer clinical trials: the UK perspective. Trials, 15(1), 379. doi:10.1186/1745-6215-15-379, Medline

Kypri, K. (2007). Methodological issues in alcohol screening and brief intervention research. Substance Abuse, 28(3), 31–42. doi:10.1300/J465v28n03_04, Medline

Lin, J. Y., Lu, Y., & Tu, X. (2012, June). How to avoid missing data and the problems they pose: design considerations. Shanghai Jingshen Yixue, 24(3), 181–184 Medline

McCormadige, J., & Saitz, R. (2017). Rethinking brief interventions for alcohol in general practice. BMJ (Clinical Research Ed.), 356, j116. doi:10.1136/bmj.j116, Medline

Moher, D., Hopewell, S., Schulz, K. F., Montori, V., Gøtzsche, P. C., Devereaux, P. J., et al. (2010). CONSORT 2010 explanation and elaboration: updated guidelines for reporting parallel group randomised trials. BMJ (Clinical Research Ed.), 340(mar23 1), c869. doi:10.1136/bmj.c869, Medline

Moher, D., Pham, B., Lawson, M. L., & Klassen, T. P. (2003). The inclusion of reports of randomised trials published in languages other than English in systematic reviews. Health Technology Assessment, 7(41), 1–90. doi:10.3310/hta7410, Medline

Moyer, A., Finney, J. W., Swearingen, C. E., & Vergun, P. (2002). Brief interventions for alcohol problems: a meta-analytic review of controlled investigations in treatment-seeking and non-treatment-seeking populations. Addiction (Abingdon, England), 97(3), 279–292. doi:10.1046/j.1360-0443.2002.00018.x, Medline

National Institute for Health and Clinical Excellence. (2010). NICE Public Health (PH) Guideline 24: Alcohol use disorders – preventing harmful drinking. Retrieved from: http://www.nice.org.uk/PH24
O'Donnell, A., Anderson, P., Newbury-Birch, D., Schulte, B., Schmidt, C., Reimer, J., & Kaner, E. (2014). The impact of brief alcohol interventions in primary healthcare: a systematic review of reviews. Alcohol and Alcoholism (Oxford, Oxfordshire), 49(1), 66–78. doi:10.1093/alcalc/agt170. Medline

Rehm, J., Mathers, C., Popova, S., Thavorncharoensap, M., Teerawattananon, Y., & Patra, J. (2009). Global burden of disease and injury and economic cost attributable to alcohol use and alcohol-use disorders. Lancet, 373(9682), 2223–2233. doi:10.1016/S0140-6736(09)60746-7. Medline

Riddle, D. L., Stratford, P. W., & Bowman, D. H. (2008). Findings of extensive variation in the types of outcome measures used in hip and knee replacement clinical trials: a systematic review. Arthritis Care and Research, 59(6), 876–883. doi:10.1002/art.23706. Medline

Saitz, R., Svikis, D., D’Onofrio, G., Kraemer, K. L., Perl, H., & Research, E. (2006). Challenges applying alcohol brief intervention in diverse practice settings: populations, outcomes, and costs. Alcoholism, Clinical and Experimental Research, 30(2), 332–338. doi:10.1111/j.1530-0277.2006.00038.x. Medline

Saitz, R. (2010). Candidate performance measures for screening for, assessing, and treating unhealthy substance use in hospitals: advocacy or evidence-based practice? Annals of Internal Medicine, 153(1), 40–43. doi:10.7326/0003-4819-153-1-201007060-00008. Medline

Shorter, G. W., Heather, N., Bray, J. W., Berman, A. H., Giles, E. L., O’Donnell, A. J., et al. (in press). Prioritization of outcomes in efficacy and effectiveness alcohol brief intervention trials: International Multi-stakeholder e-Delphi Consensus Study to inform a core outcome set.[Pagination TBC]. Journal of Studies on Alcohol and Drugs.

Shorter, G. W., Heather, N., Bray, J. W., Giles, E. L., Holloway, A., Barbosa, C., et al. (2017). The ‘Outcome Reporting in Brief Intervention Trials: Alcohol’ (ORBITAL) framework: protocol to determine a core outcome set for efficacy and effectiveness trials of alcohol screening and brief intervention. Trials, 18(1), 611. doi:10.1186/s13063-017-2335-3. Medline

Shorter, G. W., & Heather, N. Newbury- Birch, D., Giles, E. L., Holloway, A., Stockdale, K. J., . . . O’Donnell, A. J. (2016b). Outcome Reporting for Brief Intervention Trials (ORBIT): COMET Initiative Protocol. Retrieved from http://cometinitiative.org/studies/details/957

Shorter, G. W. Newbury- Birch, D., Heather, N., Giles, E. L., Holloway, A., Bray, J. W., . . . O’Donnell, A. J. (2016a). Systematic review to identify and appraise outcome measures and domains used in trials evaluating alcohol screening and brief interventions: the Outcome Reporting in Brief Intervention Trials (ORBIT) project review. Retrieved from https://www.crd.york.ac.uk/PROSPERO/display_record.php?RecordID=47185

Thornley, B., & Adams, C. (1998). Content and quality of 2000 controlled trials in schizophrenia over 50 years. BMJ (Clinical Research Ed.), 317(7167), 1181–1184. doi:10.1136/bmj.317.7167.1181. Medline

US Preventive Services Task Force. (2004). Screening and Behavioral Counseling Interventions in Primary Care to Reduce Alcohol Misuse: Recommendation Statement. Retrieved from http://www.uspreventiveservicestaskforce.org/3rduspstf/alcohol/alcomisrs.htm

Linking powered by eXtyles
Williamson, P. R., Altman, D. G., Bagley, H., Barnes, K. L., Blazeby, J. M., Brookes, S. T., et al. (2017). The COMET Handbook: version 1.0. Trials, 18(S3, Suppl 3), 280. doi:10.1186/s13063-017-1978-4. Medline

Williamson, P. R., Altman, D. G., Blazeby, J. M., Clarke, M., Devane, D., Gargon, E., & Tugwell, P. (2012). Developing core outcome sets for clinical trials: issues to consider. Trials, 13(1), 132. doi:10.1186/1745-6215-13-132. Medline

World Health Organisation [WHO]. (2016). Management of Substance Abuse. Retrieved from http://www.who.int/substance_abuse/activities/sbi/en/
Figure 1: PRISMA Flow Diagram for the systematic review of outcomes in efficacy and effectiveness trials of alcohol brief interventions

*Other sources were the cited and citing readily available peer-reviewed papers of those included in the review which met the inclusion criteria
Table 1: Features of readily available peer-reviewed randomized trials of alcohol brief interventions (published 01/01/00-31/10/17)

|                                | Overall N=405 | Primary care N=50 | Emergency Department N=47 | University/College N=132 | General population N=64 | Other healthcare N=55 | Other N=57 |
|--------------------------------|---------------|-------------------|---------------------------|--------------------------|--------------------------|-----------------------|------------|
| **Trial size Mean (SD)**       | 450.1 (730.1) | 540.0 (967.6)     | 628.3 (692.9)             | 414.1 (662.3)            | 549.5 (1074.3)           | 277.3 (246.5)        | 362.4 (404.7) |
| **Range of those randomized**  | 12-7935       | 29-6897           | 45-4476                   | 18-5227                  | 29-7935                  | 29-975               | 12-1449    |
| **Sex of those randomized**    |               |                   |                           |                          |                          |                      |            |
| **Male**                       | 56.2 (28.1)   | 65.4 (27.0)       | 67.6 (17.2)               | 45.9 (17.6)              | 56.1 (28.4)              | 46.4 (39.1)          | 76.1 (27.3) |
| **Not reported/split by arm**  | 65            | 9                 | 17                        | 15                       | 6                        | 8                    | 10         |
| **Age of those randomized**    | 31.6 (13.3)   | 48.2 (14.8)       | 30.9 (6.5)                | 20.4 (2.3)               | 39.1 (10.4)              | 37.3 (10.5)          | 34.1 (14.1) |
| **Trial region**               |               |                   |                           |                          |                          |                      |            |
| **North America**              | 246 (60.7%)   | 23 (46.0%)        | 32 (68.1%)                | 107 (81.1%)              | 32 (50.0%)               | 22 (40.0%)           | 30 (52.6%)  |
| **Australia & NZ**             | 20 (4.9%)     | 1 (2.0%)          | 1 (2.1%)                  | 6 (4.5%)                 | 4 (6.3%)                 | 3 (5.5%)             | 5 (8.8%)    |
| **Europe**                     | 110 (27.2%)   | 18 (36.0%)        | 13 (27.7%)                | 16 (12.1%)               | 25 (39.1%)               | 19 (34.5%)           | 19 (33.3%)  |
| **South America**              | 4 (1.0%)      | 0                 | 1 (2.1%)                  | 2 (1.5%)                 | 0                        | 0                    | 1 (1.8%)    |
| **Africa**                     | 10 (2.5%)     | 4 (8.0%)          | 1 (0.8%)                  | 0                        | 5 (9.1%)                 | 0                    | 0           |
| **Asia**                       | 15 (3.7%)     | 4 (8.0%)          | 0                         | 3 (4.7%)                 | 6 (10.9%)                | 2 (3.5%)             |            |
| **# of data collection waves** |               |                   |                           |                          |                          |                      |            |
| **2 waves**                    | 168 (41.8%)   | 20 (40.8%)        | 16 (34.0%)                | 63 (48.1%)               | 27 (42.2%)               | 19 (34.5%)           | 23 (41.1%)  |
| **3 waves**                    | 153 (38.1%)   | 18 (36.7%)        | 24 (51.1%)                | 44 (33.6%)               | 24 (37.5%)               | 29 (52.7%)           | 14 (25.0%)  |
| **4+ waves**                   | 81 (20.1%)    | 11 (22.4%)        | 7 (14.9%)                 | 24 (18.3%)               | 13 (20.3%)               | 7 (12.8%)            | 19 (33.9%)  |
| **Follow up wave timing**      |               |                   |                           |                          |                          |                      |            |
| **0 – 2 weeks**                | 10 (2.5%)     | 0                 | 0                         | 8 (6.1%)                 | 0                        | 1 (1.8%)             | 1 (1.8%)    |
| **>2w – 1 month**              | 91 (22.5%)    | 5 (10.0%)         | 2 (4.3%)                  | 54 (40.9%)               | 13 (20.1%)               | 8 (14.5%)            | 9 (15.8%)   |
| **>1m - 2m**                   | 56 (13.8%)    | 3 (6.0%)          | 0                         | 27 (20.5%)               | 9 (14.1%)                | 7 (12.7%)            | 10 (17.5%)  |
| **>2m - 3m**                   | 168 (41.5%)   | 19 (38.0%)        | 25 (53.2%)                | 44 (33.3%)               | 30 (46.9%)               | 20 (36.4%)           | 30 (52.6%)  |
| N (%) >3m - 4m | 14 (3.5%) | 1 (2.0%) | 1 (2.1%) | 5 (3.8%) | 1 (1.6%) | 2 (3.6%) | 4 (7.0%) |
| N (%) >4m - 6m | 202 (49.9%) | 30 (60.0%) | 23 (48.9%) | 51 (38.6%) | 36 (56.3%) | 33 (60.0%) | 29 (50.9%) |
| N (%) >6m – 9m | 32 (7.9%) | 3 (6.0%) | 3 (6.4%) | 10 (7.6%) | 3 (4.7%) | 8 (14.5%) | 5 (8.8%) |
| N (%) >9m – 12m | 120 (29.6%) | 24 (48.0%) | 30 (63.8%) | 20 (15.2%) | 14 (21.9%) | 20 (36.4%) | 12 (21.1%) |
| N (%) >12m – 18m | 14 (3.5%) | 0 | 2 (4.3%) | 2 (1.5%) | 0 | 4 (7.3%) | 0 | 6 (10.5%) |
| N (%) >18m – 24m | 17 (4.2%) | 6 (12.0%) | 0 | 3 (2.3%) | 4 (6.3%) | 3 (5.5%) | 1 (1.8%) |
| N (%) >24m | 13 (1.7%) | 7 (14.0%) | 1 (2.1%) | 3 (2.3%) | 2 (3.1%) | 0 | 0 | 0 |

| # and type of outcomes Mean (SD) # primary outcomes (n=285) | 2.4 (2.0) | 2.7 (3.0) | 2.3 (1.3) | 2.9 (2.4) | 2.3 (1.3) | 1.9 (1.3) | 2.0 (1.6) |
| Mean (SD) # secondary (n=285) | 4.1 (6.0) | 5.3 (6.8) | 5.1 (6.4) | 2.6 (3.6) | 4.3 (8.7) | 4.5 (4.1) | 3.9 (5.0) |
| Mean (SD) # not specified (n=120) | 6.4 (4.5) | 11.2 (8.8) | 5.9 (2.8) | 6.0 (4.1) | 4.8 (1.7) | 5.6 (4.1) | 8.1 (5.6) |
| Mean (SD) # outcomes per trial (n=405) | 6.5 (5.8) | 8.4 (7.3) | 7.0 (5.8) | 5.7 (3.9) | 6.3 (8.0) | 6.1 (4.1) | 6.5 (5.0) |

| Trials with at least one Alcohol related outcomes | 388 (95.8%) | 46 (92.0%) | 45 (95.7%) | 132 (100%) | 63 (98.4%) | 52 (94.5%) | 50 (87.7%) |
| N (%) Health | 80 (19.8%) | 16 (32.0%) | 6 (12.8%) | 10 (7.6%) | 16 (25.0%) | 19 (34.5%) | 13 (22.8%) |
| N (%) Economic factors/social impacts | 87 (21.5%) | 20 (40.0%) | 23 (48.9%) | 13 (9.8%) | 5 (7.8%) | 8 (14.5%) | 18 (31.6%) |
| each of the Psychological/Behavioral Factors | 114 (28.1%) | 5 (10.0%) | 11 (23.4%) | 47 (35.6%) | 14 (21.9%) | 21 (38.2%) | 16 (28.1%) |
| following Life Impact | 57 (14.1%) | 9 (18.0%) | 8 (17.0%) | 13 (9.8%) | 11 (17.2%) | 6 (10.9%) | 10 (17.5%) |
| domains Biomarkers | 13 (3.2%) | 6 (12.0%) | 0 (0.0%) | 0 (0.0%) | 0 (0.0%) | 2 (3.6%) | 5 (8.8%) |
| N (%) Intervention factors | 38 (9.4%) | 1 (2.0%) | 3 (6.4%) | 14 (10.6%) | 14 (21.9%) | 2 (3.6%) | 4 (7.0%) |

* Other includes workplaces or job related populations (n=14), veteran populations (n=19), community sample of persons with an intellectual disability (n=1), homeless population (n=1), criminal justice populations (n=14), licensed premises (n=1), sports clubs (n=1), and young people (n=6).
Figure 2: Number of alcohol brief intervention papers published per year by population
Table 2: Frequency and variability in reporting of alcohol related outcome variables overall and by population

| Outcome                                                                 | N times measured | N ways measured | Ratio of variability* | Trial type*** | Outcome type**** | Primary/ Secondary/ Not specified | Primary care N=50 | Emergency Department N=47 | University/ College N=132 | General population N=63 | Other healthcare N=55 | Other N=57 |
|------------------------------------------------------------------------|------------------|-----------------|------------------------|--------------|------------------|-----------------------------------|--------------------|---------------------------|---------------------------|--------------------------|------------------------|-----------|
| Frequency of heavy drinking                                           | 213              | 128             | 0.6                    | 105/103/5    | 90/65/58         | 30                                | 32                 | 72                        | 28                        | 24                       | 18                     | 18        |
| Number of drinks consumed in a week                                    | 205              | 63              | 0.3                    | 99/97/9      | 121/27/57       | 28                                | 19                 | 85                        | 37                        | 18                       | 18                     | 18        |
| Alcohol-related problems or consequences                               | 190              | 73              | 0.4                    | 84/98/8      | 47/62/81        | 9                                 | 28                 | 97                        | 26                        | 8                        | 22                     | 18        |
| Typical quantity                                                      | 137              | 63              | 0.5                    | 81/49/7      | 53/38/46        | 17                                | 16                 | 45                        | 22                        | 19                       | 18                     | 18        |
| Typical frequency                                                      | 117              | 73              | 0.6                    | 58/50/9      | 47/25/45        | 5                                 | 12                 | 43                        | 16                        | 22                       | 19                     | 19        |
| Hazardous or harmful drinking                                         | 111              | 27              | 0.2                    | 83/20/8      | 34/49/28        | 21                                | 15                 | 14                        | 15                        | 25                       | 21                     | 18        |
| Blood alcohol concentration                                            | 76               | 39              | 0.5                    | 41/33/2      | 28/18/30        | 0                                 | 1                  | 62                        | 7                         | 3                        | 3                      | 3         |
| At risk drinking                                                      | 72               | 63              | 0.9                    | 48/23/1      | 34/29/9         | 9                                 | 8                  | 15                        | 15                        | 6                        | 5                      | 5         |
| Largest number of drinks on occasion                                  | 57               | 36              | 0.6                    | 22/31/4      | 23/17/17        | 0                                 | 10                 | 30                        | 9                         | 5                        | 3                      | 3         |
| Days abstinent                                                        | 44               | 28              | 0.6                    | 28/13/3      | 16/22/6         | 8                                 | 4                  | 0                         | 7                         | 14                       | 11                     | 11        |
| Combined consumption measure                                          | 42               | 17              | 0.4                    | 21/9/2       | 20/9/13         | 5                                 | 3                  | 10                        | 12                        | 7                        | 5                      | 5         |
| Tobacco                                                               | 29               | 17              | 0.6                    | 5/24/0       | 7/17/5          | 2                                 | 0                  | 13                        | 9                         | 3                        | 2                      | 2         |
| Cannabis/marijuana use                                                | 26               | 19              | 0.7                    | 7/18/1       | 6/13/7          | 2                                 | 2                  | 16                        | 2                         | 2                        | 2                      | 2         |
| Number of drinks in a month                                            | 22               | 12              | 0.5                    | 15/7/0       | 13/3/6          | 5                                 | 1                  | 12                        | 3                         | 1                        | 0                      | 0         |
| Dependence symptomatology                                             | 19               | 10              | 0.5                    | 14/4/1       | 3/10/6          | 3                                 | 5                  | 3                         | 3                         | 3                        | 3                      | 3         |
| Polydrug use (alcohol +)                                               | 17               | 12              | 0.7                    | 9/7/1        | 6/8/3           | 2                                 | 1                  | 2                         | 4                         | 6                        | 2                      | 2         |
| Frequency of intoxication                                             | 15               | 9               | 0.6                    | 9/5/1        | 1/4/10          | 2                                 | 3                  | 6                         | 0                         | 2                        | 2                      | 2         |
| Other substance use                                                    | 13               | 11              | 0.8                    | 10/3/0       | 2/7/4           | 5                                 | 0                  | 2                         | 0                         | 2                        | 4                      | 2         |
| Problems with other substances                                        | 13               | 13              | 1                      | 4/8/1        | 44/5            | 0                                 | 3                  | 7                         | 1                         | 2                        | 0                      | 0         |
| Drinks on a specific occasion                                          | 9                | 8               | 0.9                    | 5/4/0        | 2/1/6           | 0                                 | 1                  | 7                         | 0                         | 0                        | 1                      | 0         |
| Number of drinks in other period                                       | 8                | 6               | 0.8                    | 8/0/0        | 6/0/2           | 1                                 | 2                  | 1                         | 0                         | 3                        | 1                      | 0         |
| Number of drinks consumed                                             | 6                | 4               | 0.7                    | 4/2/0        | 0/2/4           | 0                                 | 0                  | 4                         | 0                         | 2                        | 0                      | 0         |
in two weeks

| Measure                          | 1/3/1 | 0/4/1 | 0/0 | 0/4/0 | 0/0/4 | 0/0 | 4/0 | 0/0 | 1/1/2 | 1/1/2 | 2/0/0 | 2/0/0 | 0/0 | 1/1/0 | 1/1/0 |
|--------------------------------|-------|-------|-----|-------|-------|-----|-----|-----|-------|-------|-------|-------|-----|-------|-------|
| Other consumption measure**    | 5      | 5     | 1   | 1/3/1 | 0/4/1 | 0   | 0   | 1   | 2     | 1     | 1     |       |     |       |       |
| Abuse symptomatology           | 4      | 4     | 1   | 2/2/0 | 1/1/2 | 2   | 0   | 0   | 0     | 0     | 1     | 1     | 1   |       |       |
| Drinking game participation    | 4      | 2     | 0.5 | 0/4/0 | 0/0/4 | 0   | 0   | 4   | 0     | 0     |       |       |     |       |       |
| Preloading alcohol             | 2      | 2     | 1   | 2/0/0 | 2/0/0 | 0   | 0   | 1   | 1     | 1     | 0     | 0     |     |       |       |
| Totals                         | 1456   | 744   |     | 176   | 166   | 546 | 219 | 179 | 170   |       |       |       |     |       |       |

* Ratio of variability is the calculation of the number of variables by the approximate number of ways measured; a higher number suggests greater variability. ** Includes the following measures (times measured) drinking non-beverage alcohol (1), average time spent drinking (1), substance use successfully verified by a significant other (1), drinking the number of drinks planned to consume that night/meeting personal drinking goal (1), whether the participant thought their drinking decreased, increased or stayed the same (1). *** Refers to the number of times an outcome appeared in an effectiveness or efficacy trial, or a trial not specified as either. **** Refers to the number of times that an outcome appeared as first, second, or not specified as either.
Table 3: Frequency and variability in reporting of non-consumption variables overall and by population

| Domain and Outcome                        | N times measured | N ways measured | Ratio of variability* | Trial type** | Outcome type*** |
|-------------------------------------------|------------------|-----------------|-----------------------|--------------|-----------------|
| **Biomarkers**                            |                  |                 |                       | Efficacy/     | Primary/        |
| Gamma-glutamyltransferase                 | 10               | 7               | 0.7                   | 9/1/0        | 3/4/3          |
| Carbohydrate-deficient transferrin        | 7                | 4               | 0.6                   | 4/3/0        | 3/4/0          |
| Mean corpuscular volume                   | 6                | 4               | 0.7                   | 5/1/0        | 2/3/1          |
| Alanine aminotransferase                 | 4                | 3               | 0.8                   | 3/1/0        | 2/2/0          |
| Aspartate aminotransferase               | 4                | 3               | 0.8                   | 3/1/0        | 2/0/2          |
| Ethyl Glucuronide/ethyl sulfate           | 1                | 1               | 1                     | 0/1/0        | 0/1/0          |
| **Economic factors/social impacts**       |                  |                 |                       |              |                 |
| Driving related offences                  | 60               | 36              | 0.6                   | 17/41/2      | 17/36/7        |
| Hospitalizations                          | 36               | 11              | 0.3                   | 26/9/1       | 5/29/2         |
| Use of/seeking alcohol treatment          | 35               | 18              | 0.5                   | 24/9/2       | 5/25/5         |
| Other criminal justice use                | 35               | 26              | 0.7                   | 17/18/0      | 10/19/6        |
| Emergency healthcare use                  | 30               | 8               | 0.3                   | 18/11/1      | 7/21/2         |
| General or other healthcare use           | 29               | 23              | 0.8                   | 23/5/1       | 2/23/4         |
| Intervention cost provider                | 28               | 5               | 0.2                   | 28/0/0       | 0/28/0         |
| GP/primary care use                       | 24               | 13              | 0.5                   | 23/1/0       | 0/24/0         |
| Alcohol related injuries                  | 18               | 10              | 0.6                   | 11/6/1       | 2/6/10         |
| Outpatient healthcare                     | 14               | 8               | 0.6                   | 12/2/0       | 0/12/2         |
| Social care use                           | 13               | 10              | 0.8                   | 13/0/0       | 1/13/0         |
| Over the counter/prescribed medication use| 11               | 5               | 0.5                   | 10/1/0       | 1/8/2          |
| Alcohol related offences                  | 11               | 8               | 0.7                   | 1/10/0       | 6/3/2          |
| Use of self-help for alcohol              | 9                | 8               | 0.9                   | 6/2/1        | 0/7/2          |
| Quality adjusted life years               | 9                | 1               | 0.1                   | 9/0/0        | 0/9/0          |
| Other service use                         | 7                | 7               | 1                     | 7/0/0        | 0/7/0          |
| Intervention cost client                  | 6                | 5               | 0.8                   | 6/0/0        | 0/6/0          |

Linking powered by eXtyles
| Category                                               | Count | Frequency | Effectiveness | Mean    | Standard Deviation | Median | Mode | Total |
|--------------------------------------------------------|-------|-----------|---------------|---------|--------------------|--------|------|-------|
| **Injuries (general)**                                 | 4     | 3         | 0.8           | 2/2/0   | 1/2/1              | 2      | 2    |       |
| **Incremental cost-effectiveness ratio**               | 3     | 1         | 0.3           | 3/0/0   | 0/3/0              | 2      | 1    |       |
| **Intervention cost overall/not specified**            | 3     | 2         | 0.7           | 3/0/0   | 0/3/0              | 3      |      |       |
| **Other health economic measures**                     | 3     | 3         | 1             | 0/2/1   | 0/1/2              |        |      | 3     |
| **Productivity losses**                               | 2     | 1         | 0.5           | 2/0/0   | 0/2/0              | 2      |      |       |
| **Societal perspectives**                             | 2     | 1         | 0.5           | 2/0/0   | 0/2/0              | 2      |      |       |
| **Substance free reinforcement**                       | 2     | 1         | 0.5           | 0/2/0   | 0/2/0              |        |      |       |
| **Totals**                                             | 401   | 220       | 0.5           |         |                    | 149    | 102  | 23    | 19    | 41    | 67    |
| **Health**                                             |       |           |               |         |                    |        |      |       |
| **Alcohol-exposed pregnancy factors**                  | 31    | 16        | 0.5           | 12/19/0 | 14/13/4            | 4      | 7    | 18    | 2     |
| **Psychological health**                              | 26    | 15        | 0.6           | 20/4/2  | 0/23/3            | 7      | 3    | 1     | 2     | 6     | 7     |
| **Sexual violence or coercion**                        | 25    | 21        | 0.8           | 17/8/0  | 4/16/5            | 2      | 4    | 5     | 6     | 8     |
| **Severity of depression symptoms**                    | 24    | 13        | 0.5           | 13/10/1 | 6/17/1            | 4      | 4    | 7     | 4     | 5     |
| **Physical health**                                    | 13    | 6         | 0.5           | 13/0/0  | 0/11/2           | 7      | 2    | 1     | 1     | 2     |
| **General health**                                     | 10    | 9         | 0.9           | 9/1/0   | 1/5/4             | 3      | 1    | 2     | 3     | 1     |
| **Cardiac outcomes**                                   | 8     | 2         | 0.3           | 8/0/0   | 0/8/0             | 6      |      |       |       |       |
| **Other health factors**                               | 7     | 7         | 1             | 7/0/0   | 0/5/2            | 2      | 2    | 3     |       |       |
| **Severity of anxiety symptoms**                       | 6     | 6         | 1             | 2/4/0   | 1/5/0            | 2      | 2    | 2     | 1     |       |
| **Severity of PTSD symptoms**                          | 6     | 4         | 0.7           | 3/3/0   | 1/5/0            | 1      | 2    | 2     | 1     |       |
| **Weight/obesity**                                     | 6     | 4         | 0.7           | 6/0/0   | 0/6/0            | 2      |      |       | 4     |       |
| **Sleep disturbance**                                  | 6     | 6         | 1             | 6/0/0   | 0/6/0            |       |      |       | 6     |       |
| **Mortality/Death**                                    | 5     | 5         | 1             | 0/5/0   | 0/0/5            |       |      |       |       | 5     |
| **Suicidality**                                        | 3     | 2         | 0.7           | 2/0/1   | 0/3/0            | 2      |      | 1     |       |       |
| **Totals**                                             | 176   | 116       | 0.7           |         |                    | 35     | 12   | 23    | 30    | 43    | 33    |
| **Intervention factors**                              |       |           |               |         |                    |        |      |       |
| **Intervention satisfaction**                          | 73    | 54        | 0.7           | 29/27/17| 1/47/25          | 3      | 2    | 31    | 26    | 6     | 5     |
| **Intervention delivered/used as expected**            | 15    | 13        | 0.9           | 12/3/0  | 1/9/5            | 4      | 4    | 4     | 6     | 1     |
| **Perceived change in alcohol use**                    | 9     | 9         | 1.0           | 3/6/0   | 0/6/3            | 3      | 5    |       |       | 1     |
| **Other intervention factors**                         | 2     | 2         | 1.0           | 1/1/0   | 0/2/0            |       |      |       |       | 2     |
| **Totals**                                             | 99    | 78        | 0.8           |         |                    | 3      | 6    | 38    | 39    | 7     | 6     |
| **Psychological/behavioral factors**                   |       |           |               |         |                    |        |      |       |
| **Readiness to change**                                | 80    | 50        | 0.6           | 60/17/3 | 4/49/27          | 4      | 14   | 18    | 17    | 12    | 15    |
| Category                                      | Count | Variability | Min | Median | Max | 
|----------------------------------------------|-------|-------------|-----|--------|-----| 
| Risky behaviors                               | 49    | 0.9         | 36/13/0 | 18/22/9 | 9   | 
| Drinking refusal self-efficacy               | 43    | 0.7         | 32/8/3  | 1/25/17 | 1   | 2  | 8  | 17  | 2  | 13  | 2  | 12  |
| Alcohol outcome expectancies                  | 42    | 1           | 34/8/0  | 2/28/12 | 2   | 11 | 15 | 2   | 12  | 13  | 11  | 2  |
| Perception of others’ drinking               | 40    | 1           | 4/36/0  | 1/10/29 | 37  | 1  | 2  | 13  | 2  | 2  | 1  | 12  |
| Protective behavioral strategy use            | 17    | 0.9         | 2/15/0  | 9/4/4   | 17  | 13  | 11  | 7   | 1   | 13  | 11  | 2  |
| Anger or aggression                           | 14    | 1           | 13/1/0  | 1/1/12  | 13  | 13  | 11  | 7   | 1   | 13  | 11  | 2  |
| Other psychological factors                   | 14    | 0.9         | 9/5/0   | 0/6/8   | 1   | 2  | 6  | 13  | 11  | 7   | 1   | 1  |
| Sexual factors                                | 11    | 0.7         | 10/1/0  | 2/6/3   | 3   | 7  | 1  | 13  | 11  | 7   | 1   | 1  |
| Knowledge of alcohol                          | 10    | 1           | 9/1/0   | 1/9/0   | 8   | 1  | 1  | 13  | 11  | 7   | 1   | 1  |
| Negative/positive views of alcohol            | 9     | 0.9         | 9/0/0   | 0/6/3   | 1   | 2  | 6  | 13  | 11  | 7   | 1   | 1  |
| Alcohol demand curve measures                 | 7     | 0.9         | 0/7/0   | 0/3/4   | 7   | 1  | 1  | 13  | 11  | 7   | 1   | 1  |
| Others’ concern about drinking                | 7     | 0.6         | 5/1/1   | 0/4/3   | 1   | 2  | 1  | 13  | 11  | 7   | 1   | 1  |
| Drinking to cope                              | 6     | 1           | 4/1/1   | 1/1/4   | 4   | 1  | 1  | 13  | 11  | 7   | 1   | 1  |
| Alcohol-induced memory loss                   | 5     | 0.8         | 4/0/1   | 0/3/2   | 1   | 1  | 1  | 13  | 11  | 7   | 1   | 1  |
| Readiness to receive help                     | 5     | 1           | 4/1/0   | 1/2/2   | 1   | 1  | 1  | 13  | 11  | 7   | 1   | 1  |
| Guilt after drinking                          | 4     | 0.8         | 3/0/1   | 0/2/2   | 1   | 1  | 1  | 13  | 11  | 7   | 1   | 1  |
| Drinking in the morning                       | 3     | 0.7         | 2/0/1   | 0/2/1   | 1   | 1  | 1  | 13  | 11  | 7   | 1   | 1  |
| Impulsivity                                   | 2     | 1           | 1/1/0   | 0/1/1   | 1   | 1  | 1  | 13  | 11  | 7   | 1   | 1  |
| Goals and goal striving                       | 2     | 1           | 1/0/1   | 0/1/1   | 1   | 1  | 1  | 13  | 11  | 7   | 1   | 1  |
| Totals                                       | 363   | 0.8         | 20      | 23      | 110 | 68 | 58 | 84  | 13  | 11  | 7   | 1   | 1  |

**Life impact**

| Category                                      | Count | Variability | Min | Median | Max | 
|----------------------------------------------|-------|-------------|-----|--------|-----| 
| Role functioning/relationship factors         | 66    | 0.7         | 26/26/14 | 5/44/17 | 4   | 12 | 16 | 5   | 14 | 15 |
| Quality of life                               | 48    | 0.6         | 44/1/3  | 1/44/3  | 9   | 5  | 22 | 3   | 9  | 1 |
| Totals                                       | 114   | 0.6         | 13      | 17      | 16  | 27 | 17 | 24  | 13 | 11  | 7   | 1   | 1  |

* Ratio of variability is the calculation of the number of variables by the approximate number of ways measured; a higher number suggests greater variability. ** Refers to the number of times an outcome appeared in an effectiveness or efficacy trial, or a trial not specified as either. *** Refers to the number of times that an outcome appeared as first, second, or not specified as either.