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## Trial study design to test a bilingual digital health tool for alcohol use disorders among Latino emergency department patients

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### Abstract

We describe an emergency department (ED)-based, Latino patient focused, unblinded, randomized controlled trial to empirically test if automated bilingual computerized alcohol screening and brief intervention (AB-CASI), a digital health tool, is superior to standard care (SC) on measures of alcohol consumption, alcohol-related negative behaviors and consequences, and 30-day treatment engagement. The trial design addresses the full spectrum of unhealthy drinking from high-risk drinking to severe alcohol use disorder (AUD). In an effort to surmount known ED-based alcohol screening, brief intervention, and referral to treatment process barriers, while addressing racial/ethnic alcohol-related health disparities among Latino groups, this trial will purposively use a digital health tool and seek enrollment of English and/or Spanish speaking self-identified adult Latino ED patients. Participants will be randomized (1:1) to AB-CASI or SC, stratified by AUD severity and preferred language (English vs. Spanish). The primary outcome will be the number of binge drinking days assessed using the 28-day timeline followback method at 12 months post-randomization. Secondary outcomes will include mean number of drinks/week and number of episodes of driving impaired, riding with an impaired driver, injuries, arrests, and tardiness and days absent from work/school. A sample size of 820 is necessary to provide 80% power to detect a 1.14 difference between AB-CASI and SC in the primary outcome. Showing efficacy of this promising bilingual ED-based brief intervention tool in Latino patients has the potential to widely and efficiently expand prevention efforts and facilitate meaningful contact with specialized treatment services.

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## Keywords

Alcohol screening; Behavioral intervention; Emergency department; Digital health; Health disparities; Latino

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## 1. Introduction

### 1.1. U.S. Latinos and alcohol-related disease burden

Disparities in drinking-consequences carry heavy disease burden for U.S. Latinos. This is expected to worsen as the Latino population doubles by 2060 becoming nearly 1/3rd of the U.S. population [1–3]. Studies document greater burden of disease from both social and health perspectives (e.g., impaired driving/crashes; cirrhosis morbidity/mortality) among Latino drinkers [4–10]. The first national alcohol survey to emphasize race/ethnicity was conducted in 1984 [11]. Subsequent studies show complexity and importance of racial/ethnic variations in drinking and consequences [12–15]. Latino men have high prevalence of daily heavy drinking (more likely to binge drink) [6,16–21]. Moreover, while Non-Latino Whites are more likely to become dependent, once alcohol dependent, Latinos have higher prevalence of recurrent/persistent dependence [7,17,22,23]. Latinos also face higher rates of negative consequences (e.g., impaired driving, impaired driving-related arrests) [24,25]. While research on distinctions between/within Latino population subgroups and AUD prevalence has increased, it remains very limited.

### 1.2. U.S. emergency departments and alcohol use disorders

National Hospital Ambulatory Medical Care Survey data show that alcohol remains a major contributor to ED visits [26]. In 2017, U.S. EDs had more than 138 million visits (43.3/100 persons; over 22 million visits by Latinos); nearly 40 million injury-related. For the chronic alcohol-related condition category, there were over 4.2 million alcohol-misuse/abuse/dependence visits. Between 2006 and 2014, U.S. ED alcohol-related visits increased by over 61% [27]. Acute and chronic alcohol-related visits also increased by over 51% and 75% respectively. Another study showed that hours spent in care for alcohol-related ED visits nearly doubled from 5.6 million (2001) to 11.6 million (2011), a 108.5% increase while overall ED hours only increased 54.0% with corresponding 232.2% increase in ED resource utilization (e.g., lab tests, CTs/MRIs) [28]. A more recent study revealed that between 1999 and 2017, among those 16 y/o, annual alcohol-related deaths doubled (35,914 to 72,558) and alcohol-related death rate increased by 50% [29].

### 1.3. EDs offer opportunity to address alcohol-related disparities

ED visits coupled with alcohol screening, brief intervention, and referral to treatment (SBIRT), provide unique opportunity to intervene and address alcohol-related health disparities, particularly among vulnerable populations [30]. Nearly 20 years ago, ED-SBIRT was conceptualized. Since then, it has been rigorously tested in cohort and large RCTs [31–33]. Moreover, for approximately 15 years, the American College of Surgeons mandated SBIRT use in verified trauma centers [34]. However, past research identifies barriers (e.g., time burden, language, intervention fidelity) to consistently applying ED-SBIRT [35–37].

Use of automated bilingual (English and Spanish) computerized alcohol screening and intervention (AB-CASI) has shown it to be compelling in reducing time and resource burden while providing patient anonymity/confidentiality in self-disclosure of sensitive [38,39]. AB-CASI's automated-bilingual-scripted intervention facilitates intervention consistency. So, it can bolster intervention fidelity and integrity by reducing individual variability found when care providers use face-to-face brief interventions. This is particularly important as variations exist in accuracy and reliability of measures across different racial/ethnic groups with tailored intervention feed-back [40–43]. We describe the construction of our AB-CASI clinical trial, highlighting design decisions and considerations.

## 2. Methods

With a health disparity focus, this randomized controlled trial will accomplish three aims: 1) To compare the efficacy of AB-CASI to Standard Care (SC) in the reduction of alcohol consumption in unhealthy (i.e., high-risk) drinkers [44]; 2) To compare the efficacy of AB-CASI to SC in the reduction of alcohol-related negative health behaviors and consequences; 3) To compare the efficacy of AB-CASI to SC in 30-day treatment engagement. Further, given the paucity of ED-based alcohol SBIRT research conducted in Latino subgroups, the trial will explore variation of AB-CASI on alcohol consumption, alcohol-related negative health behaviors and consequences, and 30-day treatment engagement across Latino subpopulations (Puerto-Rican, Mexican-American, Cuban-American, South/Central American) as well as other potential modifiers (age, birthplace, gender, preferred language, dependence, reason for ED visit, and smoking status). Participants will be consented and enrolled in English or Spanish according to their language preference.

At the time of development of this trial, the DSM-IV was widely in use. From the outset, this investigation encompassed the inclusion of the whole spectrum of unhealthy drinking which at that time, definitionally, included those drinking above the low-risk limits (i.e., women and all those older than 65 y/o who have > 3 drinks/occasion or > 7 drinks/week; men who have > 4 drinks/occasion or > 14 drinks/week) through dependence [44]. In May of 2013, the DSM-5 was released and what was previously categorized in the DSM-IV as two different alcohol disorders (i.e., alcohol abuse and alcohol dependence) became one disorder (i.e., AUD) with three levels of severity; mild, moderate, and severe. Currently, the NIAAA points to the U.S. Department of Health and Human Services and U.S. Department of Agriculture's 2015–2020 Dietary Guidelines for Americans to reference high-risk drinking (i.e., for women 4 drinks on any day or 8 drinks per week; for men 5 drinks on any day or 15 drinks per week; binge drinking - drinking 4 drinks for women and 5 drinks for men within about 2 h) [45,46]. Here forward, we use the most current alcohol disorder nomenclature to minimize confusion.

Data are being collected using electronic case report forms in the Oncore Clinical Trials Management System (Forte Research, Madison, WI). The protocol for this study was reviewed and approved by Human Research Protection Program and Institutional Review Board at Yale University and the trial was registered with [Clinicaltrials.gov](https://clinicaltrials.gov) with identifier [NCT02247388](https://clinicaltrials.gov/ct2/show/study/NCT02247388).

## 2.1. Design

This study is an unblinded, parallel group RCT designed to evaluate the efficacy of AB-CASI in reducing alcohol use when compared to SC among adult Latino ED drinkers. Table 1 lists key RCT planning design decision and respective rationale.

## 2.2. Participants and setting

The study is being conducted in a large tertiary care center urban ED, American College of Surgeons verified Level 2 trauma center in the northeastern U.S. At the time when trial was started, the city in which the ED is located was known to have 44% of households speaking another language other than English at home and Latinos made up 38% of the more than 144,000 total residents. The population of the hospital's primary catchment area was 400,000 and included a diverse ethnic and cultural mix. The annual census of the ED was over 77,000 visits of which 35% were Latino, 32% White, 31% Black, and Asian/American Indian/Hawaiian Pacific Islander/Other 2%. To be included in the study, the participant must be an ED patient, self-identify as Latino, and be found to be a high-risk drinker [44,46]. Further details of enrollment criteria are shown in Table 2.

## 2.3. Measures

The study assessments and their respective timing (Fig. 1.) were organized in a manner that would allow for rigorous testing of efficacy of the AB-CASI intervention compared to a SC condition with relation to alcohol consumption, negative health behaviors and consequences, and 30-day treatment engagement. Although participants are not blinded to the intervention, interviewers conducting the assessments are blinded to intervention assignment.

## 2.4. Screening health quiz (pre-enrollment screening)

Administered in English or Spanish, the Health Quiz contains questions regarding alcohol, tobacco, exercise and seatbelt use [47,48]. The embedded questionnaire (i.e., brief alcohol use pattern screening embedded within a broader personal injury prevention survey) approach has been noted by the World Health Organization to improve the reliability of self-reported behavior. The alcohol questions embedded in the questionnaire ask three standard quantity and frequency questions [44]; 1) On average, how many days per week do you drink alcohol?; 2) On a typical day when you drink, how many drinks do you have?; 3) How many times in the past month have you had "X" or more drinks on any occasion?, where "X" is 5 for men and 4 for women. Patients who admit to a high-risk pattern drinking with consistency (e.g., frequent binge drinking) are identified and approached for consent and inclusion in the trial [44,46].

## 2.5. Alcohol use disorders identification test ((AUDIT) baseline)

The AUDIT is used at baseline to identify patients with alcohol dependence (AUDIT score 20) for the purpose of study sample stratification [49]. This metric has good operating characteristics in an emergency department setting [41]. The 10 AUDIT questions cover drinking behavior, adverse psychological reactions, alcohol-related problems, quantity and frequency of consumption. Examples of the AUDIT questions include, "On a typical week, how often do you have a drink containing alcohol- that is beer, wine, liquor, or distilled

spirits?” with response options as never; monthly or less; two to four times a month; two to three times a week; four or more times a week; “How often during the last year have you been unable to remember what happened the night before because you had been drinking?” with response options of never, less than monthly, monthly, weekly, daily or almost daily.

## **2.6. Timeline Followback ((TLFB) Baseline, 1-mo, 6-mo, 12-mo)**

The validity of self-report data on alcohol consumption has been documented previously and has been used ubiquitously for decades in the alcohol use disorder literature. This overwhelming evidence supports our use of the 28-day TLFB by telephone interview as a good method for capturing quantity and frequency of alcohol consumption and frequency of bingeing episodes [50–52]. Typically, the interviewer uses a calendar to facilitate questioning of the study participant’s retrospective daily alcohol use (i.e., the previous month). By this method, the interviewer uses “anchors” (e.g., special days of the week for the participant, weekends, and/or holidays in a particular month) to enhance the study participant’s recall of their alcohol consumption both in quantity and frequency.

## **2.7. Revised injury behavior checklist ((RIBC) baseline, 12-mo)**

The RIBC will facilitate assessment of the patient’s injury history, if they needed medical treatment for an injury, or were drinking within hours of an injury event. Originally developed for an adolescent population by Starfield [53], it was revised for use with an injured adult population by Longabaugh [54]. Its construct validity was established by relating it to the AUDIT variables in both college and ED populations. Examples of questions asked in the RIBC include, “During the past 6 months, were you injured while driving a car, truck, or bus?,” with response options as, no, yes, how many times were you injured, for how many of these injuries were you treated by a doctor, for how many of these injuries had you been drinking alcohol within 2 h of the injury; “During the past 6 months, were you injured by being physically attacked?,” with response options as, no, yes, how many times were you injured, for how many of these injuries were you treated by a doctor, for how many of these injuries had you been drinking alcohol within 2 h of the injury.

## **2.8. Short inventory of problems ((SIP) baseline, 1-mo, 6-mo, 12-mo)**

As a validated shortened version of the Drinker Inventory of Consequences (DrInC) [55,56]. The SIP contains 15-items and measures several domains (i.e., physical, social, intra/interpersonal) of negative consequences due to drinking. Examples of questions asked in the SIP include, “In the past 6 months, how often have you done impulsive things that you regretted later when you have been drinking?,” with response options as, never, once or a few times, once or twice a week, daily or almost daily; “During the past 6 months, how much have you spent too much or lost a lot of money because of your drinking?,” with response options as, not at all, a little, somewhat, very much.

## **2.9. Brief event data ((BED) baseline, 1-mo, 6-mo, 12-mo)**

The BED contains questions related to motor vehicle events, legal problems, and employment-related events. It provides information that will enable evaluation of alcohol-related negative consequences (driving impaired, riding with impaired driver, injuries,

arrests, tardiness, days absent from school or work). Items from this assessment are from the Non-Study Medical Services Form [57,58]. Examples of questions asked in the BED include, “Have you been a driver of a car involved in a crash after drinking or being intoxicated (past 6 months)?,” with response options as, no, almost, yes—once, yes—more than once, and if yes is initially answered, additional questions about who were the passengers in the car (family members, close friends, co-workers, some other person, a minor) is asked; “In the past 30 days, how many full or part days have you missed work because of your own health problems or illness, a family member’s health problem or illness, because of a legal problem, or other problem?,” with response options as, how many were full-days, and how many were part-days.

#### **2.10. Treatment services review ((TSR) Baseline, 1-mo, 6-mo, 12-mo)**

The TSR is a brief structured interview that will be administered to collect information on the type and amount of services received by participants [59]. This includes ED visits, hospitalizations, primary medical care visits and self-help sources of support (e.g. Alcoholics Anonymous). All patients who self-report treatment at a specialized treatment facility will have their data verified by the agency. Consent to contact the treatment agency is part of the original consent. Examples of questions asked as part of the TSR include, “Were you hospitalized [INPATIENT] for at least one night for any of the following reasons during the past 6 months?,” with response options as yes or now for, medical problem, surgical problem, psychiatric disorder, alcohol use disorder, substance use disorder; “Have you been to see a doctor, dentist, nurse, nurse practitioner, physician’s assistant, counselor, or chiropractor for medical care in an OUTPATIENT setting (include visits related to your participation in this study, or hospitalizations) for any of the following reasons during the past 6 months?,” with response options as yes or now for medical problem, problem related to STI, testing for HIV, surgical problem, psychiatric disorder, alcohol use disorder, substance use disorder.

#### **2.11. AB-CASI intervention**

The theoretical underpinnings of the AB-CASI intervention are rooted in development and use of the Brief Negotiation Interview (BNI). Over the last 20 years, members of our study team and colleagues, have successfully developed, refined, and empirically tested the BNI in large clinical trials [31–33,60–63]. Originally developed with colleagues at Boston University in conjunction with Rollnick [64], the BNI has been improved over time and enhanced operationally to include 4 key components that include: 1) Raise the Subject of alcohol consumption; 2) Provide Feedback on the patient’s drinking levels and effects; 3) Enhance Motivation to reduce drinking; 4) Negotiate and Advise a plan of action [63]. Moreover, the psychometric properties of the BNI have been previously tested showing good to excellent results [65]. The purpose of the BNI is to assist patients to reduce or abstain from unhealthy alcohol use, or to engage in formal treatment. Combining techniques founded in motivational interviewing as well as the stages of change model, this alcohol SBIRT intervention takes approximately 10-min to complete and has been effectively used in other clinical intervention trials focused on alcohol and drugs [31,33,66,67].



AB-CASI is a bilingual (English or Spanish) digital health tool that was developed for automated bilingual ED-SBIRT [39,68]. The version of the AB-CASI tool used runs on iPads® and is taken to the patient's bedside by a trained bilingual research assistant. Questions and messages are displayed on the iPad® screen and spoken in English or Spanish through headphones for patient privacy. Recognizing that some Latino patients may be conversationally competent in English but prefer to consume more personal health information in Spanish, patients are able to select the tool interface language based upon their comfort and preference (i.e., in both text and audio) [68]. Demographic information is collected followed by automated administration of the AUDIT. With pre-programmed logic branching, patients who screen as high-risk drinkers receive an automated BNI, including automated personalized feedback, readiness to change evaluation, reasons for cutting down, goal setting, a printed personalized alcohol use reduction plan, and counseling referral information (Fig. 2.). Patients found to be dependent by the AUDIT also receive an automated BNI that includes respectful and reflective questioning, personalized non-judgmental feedback, and supportive communication intentionally focused on treatment engagement. AB-CASI is able to reliably identify high-risk drinkers and AUD in the ED, requires little time, and is highly accepted by patients [38,39].

### 2.12. Standard care (SC) condition

Patients randomized to SC do not receive the AB-CASI intervention. However, they receive SC as provided by the treating emergency care provider. All SC patients receive an informational sheet with primary care follow-up recommended. Consultation with social workers are at the discretion of the treating emergency care provider. All requirements for alcohol screening and treatment referral are performed according to the American College of Surgeons (ACS) Level 2 trauma designation [34]. Further, in order to assess the nature of the care provided by the treating emergency care provider, review of the ED record of each enrolled study patient assigned to standard condition will be coded for emergency care provider/physician-initiated assessment (alcohol-related), any intervention and/or referral to treatment services (i.e. any documented discussion about alcohol use or referral to treatment facility in the ED treatment record or discharge instructions).

### 2.13. Follow-up telephone assessments

Formal telephone assessments by blinded, bilingual research assistants are planned at 1-, 6-, and 12-months. The 1-month follow-up will focus on collecting the extent and frequency of early treatment engagement defined as patient's report of receiving care in a treatment program that addresses their AUD (e.g. outpatient or inpatient detoxification, therapeutic community) and/or participation in a self-help program (e.g. A.A.). During this assessment, participants are asked if they received any physician-initiated advice regarding their use of alcohol during their visit to the ED. The 6-month follow-up is designed to collect early effects of the intervention on alcohol consumption and to allow enough time to collect sufficient number of negative behaviors and consequences (e.g., impaired driving motor vehicle crash, missing full or partial days of work, contact with the court/criminal justice system) by use of the BED measure, as well as all treatment engagement episodes. The 12-month follow-up assessment is designed to detect long-term effects on alcohol consumption, alcohol-related negative behaviors and consequences, and delayed referral to treatment

engagement. Several successful brief intervention studies have used these time intervals and we have chosen the same intervals to facilitate later comparison of previously published studies [54,66,69]. Because of the possibility of a “sleeper effect” or delayed emergence of treatment efficacy [70] [71], it is imperative to conduct the assessments to the 12-month interval and evaluate the effect of the intervention at each follow-up. The use of ancillary services, the occurrence of other injuries and/or illnesses, or advice from other persons may reinforce the BNI. Each of these facets, like the BNI, have the potential to help facilitate readiness for change and movement within the stages of behavior change as described in Prochaska’s transtheoretical model for behavior change [72].

#### **2.14. Data analytic plan**

This study was designed as a single-site, randomized parallel group design to test the efficacy of AB-CASI compared to SC in reducing alcohol consumption, alcohol related negative health behaviors and consequences and increase 30-day treatment engagement in Latino unhealthy drinkers, from high-risk drinking to severe AUD. All analyses will consider participants according to their randomized assignment regardless of adherence to protocol (i.e., intention to treat analysis will be conducted).

#### **2.15. Analysis of the primary outcome**

The primary objective of the analysis is to test whether AB-CASI will reduce the number of binge episodes more than SC at 12 months. A generalized linear mixed model (GLMM) with a Negative Binomial distribution will be used to estimate differences in the number of binge drinking episodes in the past 28 days at 12 months. More specifically the mixed model will include fixed effects for intervention (AB-CASI vs. SC), time (1, 6, 12 months), and the interaction of intervention with time. Additional fixed effects will be included for baseline covariates (baseline number of drinks per week, baseline number of binge episodes, gender, English vs. Spanish preferred language and dependent status). This analysis will assume that missing data occurs at random (i.e. MAR, not informative). The inclusion of baseline, 1–6- and 12-months outcome data in the model will assist in meeting this assumption. Furthermore, we will evaluate patterns of missing data as well as determine baseline characteristics that are predictive of dropout. If identified, these characteristics will be included in the model to meet the MAR assumption. Modification of the intervention effect by preferred language will be evaluated at the 0.10 significance level by including two and three-way interactions of language with intervention and time. If not significant, these interactions will be excluded and intervention effects pooled across preferred language strata. Similar procedures will be used to assess modification by dependence status. Linear contrasts (at the 0.05 two-sided significance level) will be used to estimate intervention group differences and 95% confidence intervals at the 1-, 6- and 12-month time points.

#### **2.16. Analysis of secondary outcomes**

We will test whether number of drinks per week, negative behaviors and consequences (episodes of impaired driving, riding with an impaired driver, injuries, arrests, tardiness, days absent from work/school and SIP) during the 12-month follow-up will be improved in subjects receiving AB-CASI compared to those receiving SC. Similar repeated measures mixed model analysis as that specified for the primary outcome will be implemented for



each of the secondary outcomes. Comparison of all secondary outcomes between study groups will be evaluated at the two-sided 0.01 significance level to control inflated type I error from multiple significance testing.

### 2.17. Analysis of tertiary outcome

We will test the effect of the AB-CASI compared to SC on 30-day treatment engagement. Mantel-Haenszel chi-square analysis will be used to compare the likelihood of 30-day treatment engagement in AB-CASI to SC while adjusting for preferred language and dependent status. Significance will be judged at the two-sided 0.05 significance level. Heterogeneity of treatment effect will be evaluated by the Breslow-Day test. Participants dropping out or lost-to-follow-up will be considered to be not engaged in treatment for the primary analysis.

### 2.18. Heterogeneity of treatment effects (HTE)

In addition to the stratification factors (AUD severity, preferred language), HTE on the primary outcome will be assessed for subgroups based on factors assessed at baseline (Latino ancestry, age, birthplace, gender, reason for ED visit and smoking status). These subgroup analyses will be conducted within the Generalized Linear Mixed Model framework in an evaluation similar to that proposed for investigating modification by the stratification factors of dependence status and preferred language as described above. Significant interactions will be followed by the estimation and summarization of intervention effects within subgroups at both 1-, 6- and 12-month time points.

### 2.19. Sample size

Estimation of sample size is based on randomizing and following a sufficient number of unhealthy drinkers to evaluate the primary hypothesis that AB-CASI will result in greater 12-month reductions in the primary outcome, the number of episodes of binge drinking over the past 28-days, compared to SC. Fleming et al. [69]. demonstrated that the number of binge episodes in the past 30-days was reduced by 1.14 in the intervention compared to control conditions. D'Onofrio et al. [73]. reported similar findings in an RCT conducted in hazardous and harmful drinkers. Given the following: 1) power of 80%, 2) a two-sided 0.05 significance level, 3) a standard deviation for number of binge episodes in the past 28 days of 5.2, and 4) a 1:1 intervention allocation, a sample size of 327 subjects per group will be required to detect a 1.14 difference between AB-CASI and SC in the number of binge episodes in the past 28 days at 12 months. A total of 820 unhealthy drinkers will be enrolled and randomized to accommodate up to 20% dropout. To maximize the ability to explore modification by preferred language, we will enroll an equal number of preferred English and Spanish speaking participants.

## 3. Summary

The described first-of-a-kind ED-RCT, has been intentionally designed to address alcohol-related health disparities in adult U.S. Latino ED patients using AB-CASI. In this trial, we will test the efficacy of the AB-CASI intervention against a SC condition and compare alcohol consumption, negative health behaviors and consequences, and 30-day treatment

engagement. Moreover, we will explore variations in intervention outcomes between Latino subpopulations. This study harnesses bilingual digital health tool providing a tablet-delivered brief negotiation interview (BNI). The intervention is conducted in a busy clinical setting that offers unique and important access to this vulnerable population that can benefit from directed disease prevention and health promotion efforts to close alcohol-related disparity gaps. Of particular note, this trial provides the opportunity to expand the evidence that well-known ED-SBIRT barriers (e.g., practitioner time burden, cost of intervention personnel, maintaining intervention fidelity, providing intervention in other languages) can be effectively surmounted. This could potentially reinvigorate and bolster prevention efforts to further advance national ED activities and programs addressing AUDs and ED-SBIRT practice known to currently lag behind national guidelines [74].

The design and efforts in this clinical trial are particularly unique for five reasons. First, the study is closely aligned with the NIAAA's Strategic Plan to Address Health Disparities and its most recent overall Strategic Plan (2017–2021) committing to advance the science in health disparities and developing interventions that benefit the health of at-risk populations [75,76]. Second, the design of the AB-CASI intervention has been recognized and singled out as a promising approach to address alcohol-related health disparities among racial/ethnic minorities specifically in the area of screening and brief intervention in unique settings that call for the use of innovative methods [77]. Third, with few exceptions [78], nearly all U.S. ED-SBIRT studies have not enrolled Spanish speaking participants. Fourth, none have used an automated bilingual intervention approach that facilitates disclosure of sensitive information. As a result, opportunities have been missed to capitalize on such trials in order to advance the knowledge of AUD and more specifically alcohol-related health disparities in patient-oriented outcomes among the largest minority population in the U.S. Fifth, the described trial, unlike many preceding ED-SBIRT RCTs, sets out to address and enroll participant from the full spectrum of unhealthy drinkers from high-risk drinking to severe AUD. Every day emergency care providers treat patients that present to the ED as a result of unhealthy drinking. As such, an evaluation of a broader ED-SBIRT intervention approach, that is, enrolling the full spectrum of AUD, is more congruent with the what emergency care providers routinely encounter; AUD patients with severity that results in major adverse events, such as acute and chronic physical and/or psychological harm.

We recognize that while the preliminary study of AB-CASI, prior to its proposed scientific testing described here, has shown it to be promising [38,39], it is reasonable to consider that in some U.S. ED settings, the deployment of AB-CASI may still be limited in its full and consistent implementation. In this context, limitations may arise related to the training and accountability of the specific ED personnel/staff that would be responsible for deploying the AB-CASI intervention iPads®. Further, given the ever-rapid-advancing software and hardware technologies, it is possible that the ongoing cost of keeping the AB-CASI intervention technologically current could ultimately blunt or eliminate any significant financial advantage and cost-savings in administering and sustaining meaningful ED-SBIRT efforts. Finally, with the anticipated extensive individual variability of ED workflow in U.S. EDs, it is reasonable to consider that some barriers to optimal integration of the overall AB-CASI intervention process will arise. However, without a large pragmatic study of AB-

CASI, it's difficult to say with any level of certainty that the noted limitations would be insurmountable.

If found to be effective, the AB-CASI intervention could provide more definitive evidence that not only can many previously identified ED-SBIRT barriers be overcome, but also that ED-SBIRT in the described manner could be scaled up and pragmatically implemented to significantly enhance and improve current alcohol brief intervention efforts in the ED. Moreover, because the ED is healthcare safety net for more vulnerable populations that are at greater risk for the development of alcohol-related injury, AUD, and requiring referral for specialized treatment services, an AB-CASI approach could improve the early identification AUD patient and help facilitate their referral to treatment services. On an even broader scale, if AB-CASI is found to be effective, its logical to consider that this approach could afford valuable opportunities for intervention adaptation. That is, it may also lend itself to systematic implementation of AB-CASI in a number of other languages as well as use in primary care where more recent literature suggests the need for greater scope of not only intentional screening but also monitoring of reductions in alcohol use as well as medical treatment of AUDs, both to the benefit of patients [79,80].

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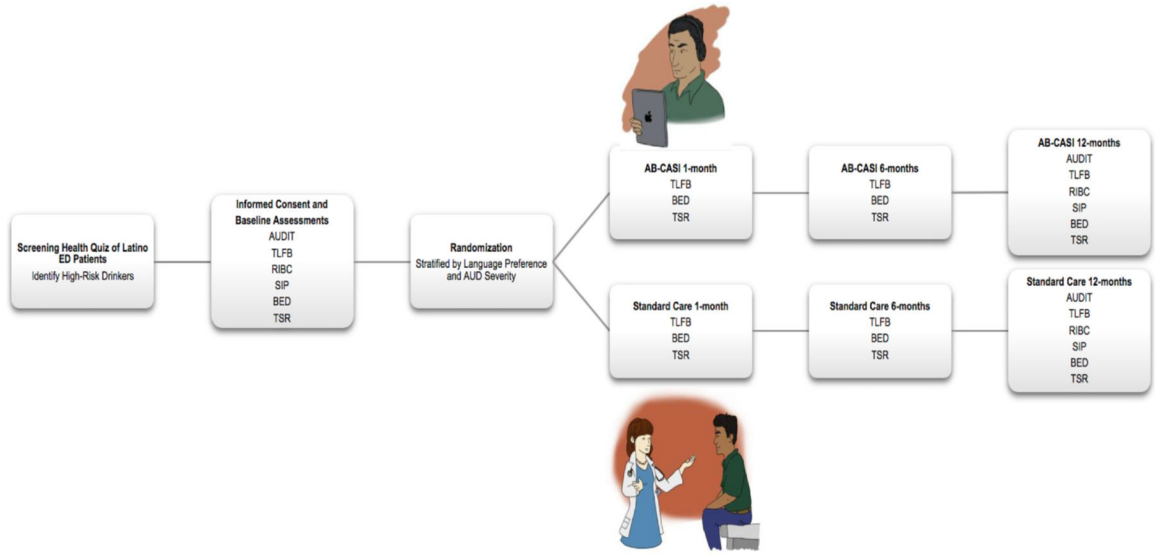
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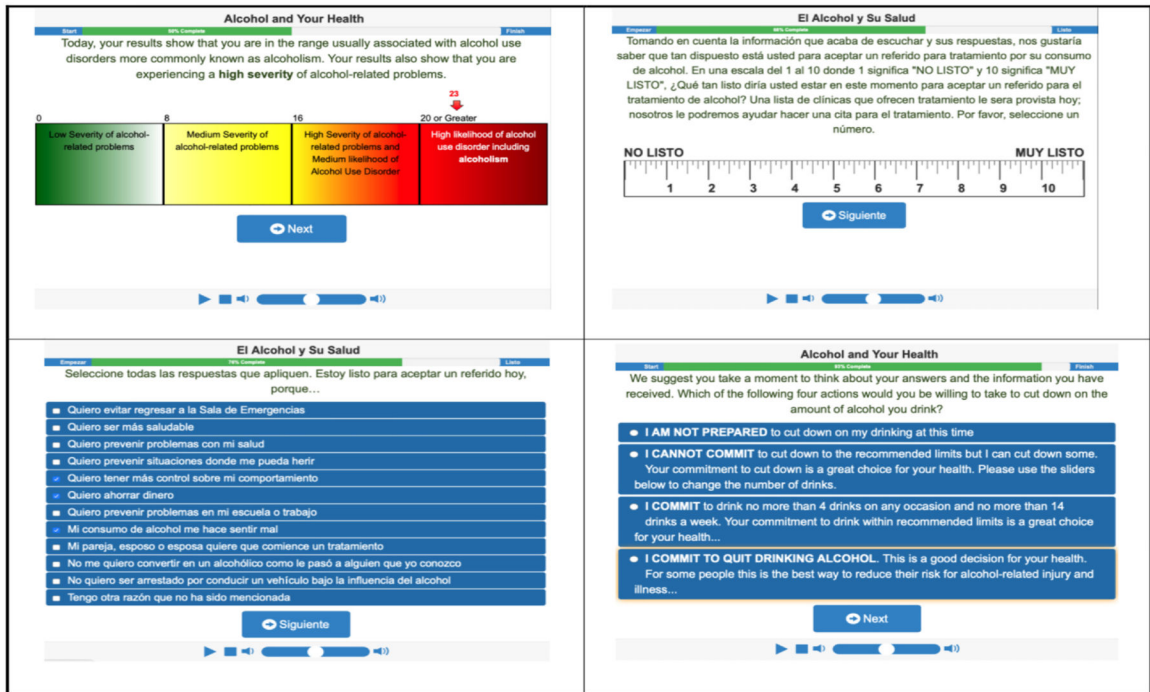
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**Fig. 1.** Schedule of assessments. Alcohol Use Disorders Identification Test ((AUDIT); Timeline Followback (TLFB); Revised Injury Behavior Checklist (RIBC); Short Inventory of Problems (SIP); Brief Event Data (BED); Treatment Services Review (TSR).



**Fig. 2.**  
Automated Bilingual Computerized Alcohol Screening and Intervention (AB-CASI):  
Example screen views in English and Spanish.

**Table 1**

Design decisions and rationale.

<b>Decision</b>	<b>Rationale</b>
Enrollment of self-identified Latino ED patients	Address alcohol-related health disparities in a highly vulnerable group in a ED-SBIRT context
Bilingual (English and Spanish) ED-SBIRT	Address known major language barrier limitation in ED-SBIRT
Prescreen [Health Screen]	Ensure enrollment of consistent above low-risk limit drinkers
5 baseline assessments	Capture alcohol use quantity and frequency, severity of AUD, negative and injury-related outcomes, type and amount of treatment services received
Randomization	Facilitate testing efficacy of AB-CASI intervention
Stratification [dependence and language]	Explore treatment modification
1-mo, 6-mo, 12-mo follow-up assessment	Longitudinal assessment testing efficacy in outcomes of interest

**Table 2**

Inclusion and exclusion criteria.

Inclusion	Exclusion
Self-identified adult Latino ED patients	Current enrollment in alcohol or substance abuse treatment program
English and/or Spanish speaking	At the time of enrollment known to be pregnant
High-risk drinking [44,46]	Current ED visit for acute psychosis (i.e. suicidal or homicidal ideation)
	Condition that precludes interview or AB-CASI use i.e., life threatening injury/illness including sexual assault, poor decisional capacity due to cognitive impairment
	Police custody
	Inability to provide two contact numbers for follow-up