

Department of Health and Human Services

Substance Abuse and Mental Health Services Administration

Cooperative Agreements for Screening, Brief Intervention, Referral and Treatment (SBIRT) (Initial Announcement)

Request for Applications (RFA) No. TI-08-001

Catalogue of Federal Domestic Assistance (CFDA) No.: 93.243

Key Dates:

Application Deadline	Applications are due by January 31, 2008.
Intergovernmental Review (E.O. 12372)	Applicants must comply with E.O. 12372 if their State(s) participates. Review process recommendations from the State Single Point of Contact (SPOC) are due no later than 60 days after application deadline.
Public Health System Impact Statement (PHSIS)/Single State Agency Coordination	Applicants must send the PHSIS to appropriate State and local health agencies by application deadline. Comments from Single State Agency are due no later than 60 days after application deadline.

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Executive Summary:

The Substance Abuse and Mental Health Services Administration, Center for Substance Abuse Treatment is accepting applications for fiscal year (FY) 2008 for Cooperative Agreements for Screening, Brief Intervention, Referral and Treatment (SBIRT). The purpose of this program is to expand and enhance State/Tribal substance abuse treatment service systems.

Funding Opportunity Title:	Cooperative Agreement for Screening, Brief Intervention, Referral and Treatment (SBIRT)
Funding Opportunity Number:	TI-08-001
Due Date for Applications:	January 31, 2008
Anticipated Total Available Funding:	Up to \$10.044 million
Estimated Number of Awards:	4
Estimated Award Amount:	Up to \$2.52 million per year
Length of Project Period:	Up to 5 years
Eligible Applicants:	States, Territories, federally recognized tribes, and tribal organizations are eligible to apply. [See Section III-1 of this RFA for complete eligibility information.]

I. FUNDING OPPORTUNITY DESCRIPTION

1. INTRODUCTION

The Substance Abuse and Mental Health Services Administration, Center for Substance Abuse Treatment is accepting applications for fiscal year (FY) 2008 for Cooperative Agreements for Screening, Brief Intervention, Referral and Treatment (SBIRT). The purpose of this program is to expand and enhance State/Tribal substance abuse treatment service systems by:

- Expanding the State's/Tribe's continuum of care to include screening, brief intervention, referral, and brief treatment (SBIRT) in general medical and other community settings (e.g., community health centers, nursing homes, schools and student assistance programs, occupational health clinics, hospitals, emergency departments);
- Supporting clinically appropriate services for persons at risk for, or diagnosed with, a Substance Use Disorder (i.e., Substance Abuse or Dependence). (Note: for the purpose of this RFA, "at risk" is defined as persons who are using substances but who do not yet meet the criteria for a diagnosis of Substance Use Disorders¹); and
- Identifying systems and policy changes to increase access to treatment in generalist and specialist settings.

Screening, Brief Intervention, Referral and Treatment (SBIRT) is one of SAMHSA's services grant programs. SAMHSA's services grants are designed to address gaps in substance abuse treatment services and/or to increase the ability of States, units of local government, American Indian/Alaska Native tribes and tribal organizations, and community- and faith-based organizations to help specific populations or geographic areas with serious, emerging substance abuse problems. SAMHSA intends that its services grants result in the delivery of services as soon as possible after award. Substantial service delivery should begin by the 6th month of the project at the latest.

SBIRT grants are authorized under Section 509 of the Public Health Service Act, as amended. This announcement addresses Healthy People 2010 focus area 26 (Substance Abuse).

2. EXPECTATIONS

Background

The National Drug Control Strategy (NDCS) emphasizes: (1) preventing drug use before it starts; (2) intervening and healing those who already use drugs; and (3) disrupting the market for illicit substances (ONDCP, 2007). SBIRT's focus on early intervention and treatment is a central component of NDCS.

¹ For purposes of this announcement, the need for treatment is discussed in terms of the categories of substance use disorders (substance dependence and substance abuse) used in the American Psychiatric Association Diagnostic and Statistical Manual of Mental Disorders (DSM-IV; 1994).

Federal programs, including those operated by SAMHSA/CSAT, have tended to emphasize either universal prevention strategies aimed at those who have never initiated use (Mrazek and Haggerty, 1994) or specialist treatment for those who are dependent (Gerstein and Harwood, 1990). Little attention has been paid to the large group of individuals who use drugs but are not yet dependent who could successfully reduce drug use through “early intervention” (Fleming, 2002; Klitzner et al., 1992). There is an emerging body of research and clinical experience that supports use of the SBIRT approach as providing effective early intervention for persons at risk for, or diagnosed with, a Substance Use Disorder (i.e., Substance Abuse or Dependence).²

The specialist treatment system is often not appropriate for persons at risk for a Substance Use Disorder, nor can that system alone address the needs of all those persons diagnosed with either a Substance Abuse or a Substance Dependence Disorder. Moreover, most people with or at risk for a Substance Use Disorder are unlikely to seek help from the specialty treatment system. According to SAMHSA’s National Survey on Drug Use and Health (NSDUH) in 2006, 23.6 million persons age 12 or older needed treatment for a Substance Use Disorder involving alcohol or an illicit drug. Of these, only 2.5 million (10.6 percent) received treatment at a specialty facility. Of the 21.1 million persons who needed but did not get treatment, only 940,000 (4.5 percent) recognized that they needed help for their problem and only one third of these individuals (314,000 persons) made an effort to get treatment.³

Consequently, new program efforts are needed to provide funding to introduce or expand screening and brief intervention and brief treatment for persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence). These new program efforts need to be initiated in general medical and other community settings (e.g., community health centers, nursing homes, schools and student assistance programs, occupational health clinics, hospitals, emergency departments).

SBIRT is valuable not just for individuals who use alcohol and illicit drugs but also for those who use prescription drugs non-medically. One area of concern highlighted by the 2006 NSDUH is the growing misuse of prescription drugs. For example, non-medical use of prescription drugs among young adults increased from 5.4 percent in 2002 to 6.4 percent in 2006, due largely to an increase in the non-medical use of pain relievers.⁴

² E.g., see Babor, 2004; Babor et al. 2002; Baker et al. 2001; Barry, 1999; Bernstein et al., 1997; Blow, 1999; Conrod et al., 2000; Dennis et al., 2002a and b; Fleming, 2002; Gil et al., 2004; Gray et al., 2005; Grenard et al., 2007; Humeniuk, 2006; Kelso, 2002; Rohsenow et al., 2004; Samet et al., 1996; Stephens et al., 1994; Stephens, et al., 2000; Stephens, et al., 2004; Sullivan et al., 1997; WHO ASSIST Working Group, 2002; Zweben and Fleming, 1999). A complete listing of the references and resources (including evidence-based practices/services) cited in this document can be found at http://www.mayatech.com/sbirt/tools_resources/references.htm. Copies of the listing are available in the application kits distributed by SAMHSA’s National Clearinghouse for Alcohol and Drug Information.

³ SAMHSA, 2006 National Survey on Drug Use and Health, September 2006

⁴ Ibid.

Services Design

Grantees will be required to adopt and implement a treatment system that includes all of the following components:

- *Screening, Identification, Brief Intervention, Referral, and Treatment.* This involves the implementation of a system within community and specialist settings that screens for and identifies persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence) including such disorders involving use of prescription medications. SAMHSA is committed to: (1) standardizing the screening process so that screening produces consistent results across sites, in terms of individuals identified as requiring brief intervention or referral to treatment; and (2) detecting and intervening with individuals who have a Substance Use Disorder, but whose problems are not so severe that they require specialty treatment. Grantees will be required to use the Alcohol, Smoking, and Substance Involvement Screening Test (ASSIST) tool for adults and the Car, Relax, Alone, Forget, Friends, Trouble (CRAFFT) for screening adolescents. Depending on the level of problems identified, the system either provides for a brief intervention within the generalist setting, when appropriate, or motivates and refers persons with a probable diagnosis of substance abuse or substance dependence to the specialist setting for assessment, diagnosis, and either brief or long-term treatment. This includes training in self-management and involvement in mutual help groups, as appropriate (Workgroup on Substance Abuse Self-Help Organizations, 2003).⁵ The evidence-based approaches and tools utilized for screening, brief intervention, referral, and treatment may vary; however, the core components of SBIRT remain (see Appendix I). As part of brief intervention, applicants must screen and assess clients for the presence of co-occurring substance use (abuse and dependence) and mental disorders and use the information obtained from screening and assessment to develop appropriate treatment approaches for persons identified as having such co-occurring disorders. For more information on the process of selecting screening instruments to identify co-occurring substance use and mental disorders, go to http://www.cocce.samhsa.gov/cod_resources/cod_presentations.aspx.
- *Sequential Assessment and Diagnosis.* This involves having a system in place to ensure that individuals who screen positive for substance use disorders are appropriately assessed for the presence of co-morbid physical and mental disorders so that a diagnosis is made, an initial treatment plan is developed, and a referral is made to the clinically appropriate community or specialist treatment setting as dictated by the person's clinical status. Mental health services are not part of this cooperative agreement.
- *Treatment.* This involves having a system in place to ensure that individuals who are diagnosed with a substance use disorder are provided an opportunity to undergo an integrated pharmacological and psychosocial treatment regimen in order to reduce or eliminate their harmful consumption and its adverse effects in the clinically appropriate

⁵ References and resources that support this approach are available at www.mayatech.com/sbirt/tools_resources/references.htm.

community or specialist treatment setting. This includes training in self-management and involvement in mutual help groups, as appropriate. Mental health services are not part of this cooperative agreement.

- *Continued Management Support.* This involves having a system in place to ensure that individuals who complete their formal treatment episode will receive long term management support (care management and recovery support services), as appropriate for their level of disability and relapse potential in the clinically appropriate community or specialist treatment setting. This includes training in self-management and involvement in mutual help groups.⁶

Patients manifesting signs of intoxication, withdrawal symptoms, and other physical problems that require emergency care or urgent action would be managed in other components of the generalist or specialist treatment systems. While stabilization and detoxification may be required for some persons presenting to community agencies, the availability of treatment resources, financing mechanisms, and other access barriers vary from those encountered in treating individuals who do not require withdrawal and stabilization. This variation is recognized in the differentiation of the levels and settings of services for detoxification and rehabilitation in the latest version of the American Society of Addiction Medicine (ASAM) Patient Placement Criteria (ASAM, 2001); Gastfriend et al., 2000) and the guidelines developed by the Evidence-Based Clinical Practice Guideline Working Group for the Veterans Health Administration and Department of Defense (2001).

The applicant can propose to implement the project in as many sub-recipient communities⁷ as it wishes. For each sub-recipient community chosen, the applicant must demonstrate need and potential for systems change to rapidly initiate the SBIRT approach. Each sub-recipient community must receive sufficient funds to enable the grantee to document an impact using the SBIRT performance targets.

Policy and Systems Change

Eliminating barriers through policy and systems change is a major emphasis of this program (e.g., Pauly, 1991; Libertoff, 1999; Zarkin et al., 1995). Integrating SBIRT in community settings requires an analysis of the policies, procedures and practices that inhibit and facilitate implementation. You must include in your application a change plan that reflects an understanding of the general reasons why people do not seek services as well as how barriers which prevent individuals from successfully accessing the clinically appropriate level of care may apply to your system of care. These barriers may include laws, regulations, eligibility requirements for service receipt, facility and provider eligibility requirements, varied funding streams, coverage limitations, a lack of patient placement criteria or standardized screening and

⁶ References and resources that support this model include CSAT's Treatment Improvement Protocols (TIPs) and Technical Assistance Publications (TAPs).

⁷ For purposes of this announcement, a community may be a geopolitical unit (city, county), a health district or human services region, or a substate planning area as defined for purpose of allocating Substance Abuse Prevention and Treatment Block Grant (SAPTBG) funds.

sequential assessment protocols, etc. In addition, your change plan must describe the policy and systems changes you will undertake to eliminate existing barriers that impede people from accessing the treatment they need. For example, enhanced collaboration with federally Qualified Health Centers (FQHCs) may be an appropriate application for SBIRT within the context of this RFA. See Appendix J for further discussion of policy and systems change.

Project Phases

Phase I: Project Planning and Start-Up. This phase is expected to last approximately 6 months. During this time, CSAT will work collaboratively with the grantee, project staff, and Policy Steering Committee members (see *Phase II: Operations*). The start-up tasks to be completed in this phase are, at minimum:

- Selecting the members of the Policy Steering Committee (see *Phase II: Operations* for membership).
- Developing a solid organizational structure that involves or enlists the participation of an appropriate array of service providers and funders representing the full spectrum of community and specialist services required to serve the needs of persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence) in the sub-recipient communities.
- Refining the project management, reporting, quality improvement, and cost control mechanisms.
- Refining the needs assessment and survey of existing system gaps and precisely identifying the target populations and communities to be served.
- Refining the plan to provide training and technical assistance, including information about SBIRT methods, training for staff in the community and specialist settings in carrying out SBIRT, and technical assistance to the overall project and its sub-recipients.
- Finalizing all necessary interagency agreements, contracts, subcontracts, billing procedures and fiscal controls, and reporting and monitoring procedures with the agency or agencies in the communities that will deliver services.
- Introducing reporting instruments and obtaining baseline data covering existing levels of service, patient/client needs, program performance characteristics, and training and technical assistance.
- Developing a plan for garnering and sustaining necessary policy changes and resources required to continue the project following the period of Federal support.
- Demonstrating that required resources not included in the Federal budget request are adequate and readily accessible.

- Initiating service delivery in the expanded continuum of care in each sub-recipient target community, if required.
- Establishing the mechanism for monitoring performance against targets for: (1) reducing drug use by patients receiving services through the SBIRT project; (2) increasing the number of persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence) who receive treatment in each sub-recipient community; (3) increasing the number of community settings where SBIRT services are provided; and (4) providing treatment services within approved cost parameters for each treatment modality.
- Submitting an acceptable final Project Implementation Plan that includes specific objectives and milestones, implementation timeframes and designation of staff responsible for accomplishing individual program objectives.

Release of funds for project implementation will be contingent on CSAT approval of the Project Implementation Plan finalized during the initial phase and submitted for approval by the end of the third month following award. At the conclusion of Phase I, every component of the project should be fully operational.

Phase II: Operations. This phase is expected to last approximately 4 years and 3 months. During this time, CSAT will work collaboratively with the grantee, project staff, State Substance Abuse Authority (SSA), and other relevant agencies, Policy Steering Committee members, and sub-recipients to implement project management, monitoring and reporting, training, technical assistance to sub-recipients, and service delivery. In Phase II, the grantee will be responsible for these activities:

- Operation of the Policy Steering Committee (and its subcommittees, if appropriate), including regular meetings; monitoring project activities and achievements with regard to the specific objectives and milestones, implementation timeframes, and designation of staff according to the Project Implementation Plan; and communications with the sub-recipients and CSAT Project Officers.
- Determining the need for and providing the requisite training and technical assistance needed to achieve project goals.
- Project management, reporting, quality improvement, and cost control.
- Managing the continuation award process to the sub-recipients.
- Accomplishing and tracking systems change (i.e., overcoming funding and other resource barriers, policy changes, improving linkages among specialist and community agencies, providing training and technical assistance, carrying out service delivery in the expanded continuum of care in each sub-recipient target community) and achieving the targets for: (1) reducing drug use by patients receiving treatment through the SBIRT project; (2) increasing the number of persons at risk for, or diagnosed with, a Substance Use Disorder

(Substance Abuse or Dependence) who receive treatment in each sub-recipient community; (3) increasing the number of community settings where SBIRT services are provided; and (4) providing treatment services within approved cost parameters for a given treatment modality.

- Refining operations as barriers are encountered and lessons are learned through feedback from the monitoring and reporting systems.

Phase III: Phase Out. During the final 3 months of the cooperative agreement award, CSAT will work cooperatively with the grantee, project staff, Policy Steering Committee members, and sub-recipients to make the transition from the cooperative agreement to State/Tribe and local control and to sustain the system changes achieved by the project.

2.1 Using Evidence-Based Practices

SAMHSA's services grants are intended to fund services or practices that have a demonstrated evidence base and that are appropriate for the target population. An evidence-based practice, also called EBP, refers to approaches to prevention or treatment that are validated by some form of documented scientific evidence. In your application, you will need to:

- Identify the evidence-based practice you propose to implement.
- Identify and discuss the evidence that shows that the practice is effective. [See note below.]
- Discuss the population(s) for which this practice has been shown to be effective and show that it is appropriate for your target population(s). [See note below.]

Note: SAMHSA recognizes that EBPs have not been developed for all populations and/or service settings. For example, certain interventions for American Indians/Alaska Natives, rural or isolated communities, or recent immigrant communities may not have been formally evaluated and, therefore, have a limited or nonexistent evidence base. In addition, other interventions that have an established evidence base for certain populations or in certain settings may not have been formally evaluated with other subpopulations or within other settings. Applicants proposing to serve a population with an intervention that has not been formally evaluated with that population are encouraged to provide other forms of evidence that the practice(s) they propose is appropriate for the target population. Evidence may include unpublished studies, preliminary evaluation results, clinical (or other professional association) guidelines, findings from focus groups with community members, etc. You may describe your experience either with the target population or in managing similar programs. Information in support of your proposed practice needs to be sufficient to demonstrate the appropriateness of your practice to the people reviewing your application.

- Document the evidence that the practice you have chosen is appropriate for the outcomes you want to achieve.

- Explain how the practice you have chosen meets SAMHSA’s goals for this grant program.
- Describe any modifications/adaptations you will need to make to this practice to meet the goals of your project and why you believe the changes will improve the outcomes. We expect that you will implement your evidence-based service/practice in a way that is as close as possible to the original service/practice. However, SAMHSA understands that you may need to make minor changes to the service/practice to meet the needs of your target population or your program, or to allow you to use resources more efficiently. You must describe any changes to your proposed service/practice that you believe are necessary for these purposes. You may describe your own experience either with the target population or in managing similar programs. However, you will need to convince the people reviewing your application that the changes you propose are justified.
- Explain why you chose this evidence-based practice over other evidence-based practices.

Resources for Evidence-Based Practices:

You will find information on evidence-based practices in SAMHSA’s *Guide to Evidence-Based Practices on the Web* at www.samhsa.gov/ebpwebguide. SAMHSA has developed this Web site to provide a simple and direct connection to Web sites with information about evidence-based interventions to prevent and/or treat mental and substance use disorders. The *Guide* provides a short description and a link to dozens of Web sites with relevant evidence-based practices information – either specific interventions or comprehensive reviews of research findings.

Please note that SAMHSA’s *Guide to Evidence-Based Practices* also references another SAMHSA Web site, the National Registry of Evidence-Based Programs and Practices (NREPP). NREPP is a searchable database of interventions for the prevention and treatment of mental and substance use disorders. NREPP is intended to serve as a decision support tool, not as an authoritative list of effective interventions. *Being included in NREPP, or in any other resource listed in the Guide, does not mean an intervention is “recommended” or that it has been demonstrated to achieve positive results in all circumstances.* You must document that the selected practice is appropriate for the specific target population and purposes of your project.

In addition to the Web site noted above, you may provide information on research studies to show that the services/practices you plan to implement are evidence-based. This information is usually published in research journals, including those that focus on minority populations. If this type of information is not available, you may provide information from other sources, such as unpublished studies or documents describing formal consensus among recognized experts.

2.2 Services Delivery

SBIRT Services: Grantees must develop a systematic approach and must devote not less than **65 percent (65%)** of their award to expand and enhance their service system to carry out the following services, face-to-face, in community agencies, including establishing referral linkages to specialist treatment agencies/providers:

- **Screening** for the presence of Substance Use Disorders

- **Brief Interventions** (1 to 5 sessions)
- **Brief Treatment** (up to 12 sessions) and monitoring of individuals who use drugs but are not yet dependent
- **Referral to Treatment** (when indicated) for those who have a Substance Use Disorder. Persons who qualify for a diagnosis of drug abuse or dependence and who are non-responsive to an initial brief intervention or brief treatment must be referred for specialty treatment.

Specialty Treatment Services: While the focus of this initiative is on screening, brief intervention, and brief treatment, it is critical to ensure that appropriate services are available to treat persons for whom such services in community settings are not appropriate. Therefore, a portion of the award may be used for referral to specialty treatment. Grantees may use up to **15 percent (15%)** of the award to expand specialty modalities (outreach/pretreatment services, methadone and non-methadone outpatient services, and residential services) for persons screened in this program who require more intensive and prolonged specialty treatments who lack other treatment payment recourse.

Service delivery should begin by the 6th month of the project at the latest.

2.3 Infrastructure Development (maximum 15% of total grant award)

Although services grant funds must be used primarily for face-to-face direct services, SAMHSA recognizes that infrastructure changes may be needed to implement the services or improve their effectiveness. You may use up to 15% of the total services grant award for the following types of infrastructure development, if necessary to support the direct service expansion of the grant project, such as:

- Developing partnerships with other service providers for service delivery.
- Enhancing your computer system, management information system (MIS), electronic health records, etc.
- Training/workforce development to help your staff or other providers in the community identify mental health or substance abuse issues or provide effective services consistent with the purpose of the grant program.
- Applying for the approval of State Medicaid funding for the appropriate reimbursement codes expeditiously. In addition, providers are encouraged to bill reimbursing entities for screening and brief intervention services. Case reimbursement through Medicaid will allow a portion of funds to be used as the State's Medicaid matching fund requirement.

2.4 Data Collection and Performance Measurement

All SAMHSA grantees are required to collect and report certain data so that SAMHSA can meet its obligations under the Government Performance and Results Act (GPRA). CSAT's GPRA requirements for this program include data collection and real time reporting about cooperative

agreement-supported service recipients at baseline/intake, discharge and 6 months after intake. Grantees are also expected to obtain a minimum of an 80 percent (80%) follow-up rate on those clients selected as part of the follow-up sample. Data must be entered into the CSAT GPRA Web-based data entry and reporting system on a real-time basis. Grantees also are required to submit specified aggregate data in semi-annual reports. These requirements should be considered when preparing the data collection, monitoring, and reporting budget section of the application. This information will be gathered using the data collection tool referenced below. The collection of these data will enable CSAT to report on the National Outcome Measures (NOMs), which have been defined by SAMHSA as key priority areas relating to substance abuse. You must document your ability to collect and report the required data in “Section E: Performance Assessment and Data” of your application. Grantees must collect and report data using the Discretionary Services Client Level GPRA Tool, which can be found at www.samhsa-gpra.samhsa.gov. This Website must be used to enter full GPRA baseline/intake, discharge and 6 months after intake data. Hard copies are available in the application kits available by calling the SAMHSA Information Line at 1-877-SAMHSA7 [TDD: 1-800-487-4889].

Many grantees have found that an automated system to create a database and upload GPRA information has been quite helpful. As such, applicants are expected to include an automated system for these activities as part of their proposal.

Applicants should carefully note that there are three categories of services or combinations of services to be supported by these cooperative agreement funds and each category has specific reporting requirements with regards to GPRA.

The three categories of services or combinations of services to be provided to individuals include:

- Screening Only;
- Screening and Brief Intervention (BI); and
- Screening and Brief Treatment (BT) or Screening and Referral to Other Types of Treatment for Substance Use Disorders (RT).

Varying levels of data are required on clients in each category of care. Drug use, employment status, housing status, criminal justice status, social connectedness, access and retention will all be measured using specific sections of the GPRA tool. There are also follow-up data specifications. Follow-up data will be required on 10 percent (10%) of the clients served in **each** category of care requiring intervention/treatment (BI, BT, and RT). Please see Appendix L for the reporting requirements for each category of services.

Grantees will be required use the ASSIST to screen adults and the CRAFFT to screen adolescents. The screening and collection of the GPRA information must be face-to-face. Additional screening instruments/tools may be used with the agreement of the SAMHSA Project Officer.

Grantees must comply with GPRA data collection and reporting requirements, including continuous reporting⁸ of progress in meeting the targets proposed (in the application) for the number of persons to be served and the collection of the specified CSAT GPRA Core Client Outcomes at specified time points. Grantees are required to collect and report client level data for the overall project and for each sub-recipient using the CSAT GPRA data entry and reporting system.

Other Reporting Requirements

CSAT's GPRA Core Client Outcome domains are:

- Number of individuals served;
- Percent of service recipients who: have no past month substance use; have no or reduced alcohol or illegal drug consequences; are permanently housed in the community/living in a stable housing environment; are employed/in school; have no or reduced involvement with the criminal/juvenile justice system; have increased social connectedness; and, have good or improved health and mental health status.

Applicants must clearly state which GPRA service population(s) they propose to address as target populations. For more information, as well as the electronic versions of the CSAT GPRA materials, go to www.samhsa-gpra.samhsa.gov.

Performance data will be reported to the public, the Office of Management and Budget (OMB) and Congress as part of SAMHSA's budget request.

2.5 Performance Assessment

Grantees must assess their projects, addressing the performance measures described in Section I-2.4. The assessment should be designed to help you determine whether you are achieving the goals, objectives and outcomes you intend to achieve and whether adjustments need to be made to your project. You will be required to report on your progress achieved, barriers encountered, and efforts to overcome these barriers in a performance assessment report to be submitted at least annually.

In addition to assessing progress against the performance measures required for this program, your performance assessment must also consider outcome and process questions, such as the following:

Outcome Questions:

- What was the effect of the intervention on participants?
- What program/contextual factors were associated with outcomes?
- What individual factors were associated with outcomes?

⁸ Continuous reporting is defined as entering client level data into the GPRA Web-based data system within 7 business days of collection.

- How durable were the effects?

Process Questions:

- How closely did implementation match the plan?
- What types of deviation from the plan occurred?
- What led to the deviations?
- What effect did the deviations have on the planned intervention and performance assessment?
- Who provided (program staff) what services (modality, type, intensity, duration), to whom (individual characteristics), in what context (system, community), and at what cost (facilities, personnel, dollars)?

SAMHSA may conduct a cross-site evaluation of all SBIRT projects funded under its SBIRT funding announcements. Projects funded under this announcement will be included in the cross-site evaluation and all grantees are required to participate in this evaluation.

Many State Medicaid agencies, as part of their approval for funding, require a third party evaluation to certify the fidelity and quality of services delivery. Applicants are encouraged to include this third party evaluation of the fidelity and quality of SBIRT services in the design of their proposal.

No more than 20% of the total grant award may be used for data collection, performance measurement, and performance assessment, e.g., activities required in Sections I-2.4 and 2.5 above.

2.6 Grantee Meetings

Grantees must plan to send a minimum of two people (including the Project Director) to at least one joint grantee meeting in each year of the grant, and you must include a detailed budget and narrative for this travel in your budget. At these meetings, grantees will present the results of their projects and Federal staff will provide technical assistance. Each meeting will be 3 days. These meetings are usually held in the Washington, D.C., area and attendance is mandatory.

II. AWARD INFORMATION

Funding Mechanism:	Cooperative Agreement
Anticipated Total Available Funding:	\$10.044 million
Estimated Number of Awards:	4
Estimated Award Amount:	Up to \$2.52 million per year
Length of Project Period:	Up to 5 years

This program is being announced prior to the appropriation for FY 2008 for SAMHSA's programs, with funding estimates based on the President's budget request for FY 2008. Applications are invited based on the assumption that sufficient funds will be appropriated for FY 2008 to permit funding of a reasonable number of applications solicited. All applicants are reminded, however, that we cannot guarantee that sufficient funds will be appropriated to permit SAMHSA to fund any applications.

Proposed budgets cannot exceed \$2.52 million in total costs (direct and indirect) in any year of the proposed project. Annual continuation awards will depend on the availability of funds, grantee progress in meeting project goals and objectives, timely submission of required data and reports, and compliance with all terms and conditions of award.

Applicants should be aware that the amount to be awarded for continuation awards in year 5 is expected to be 95% of the amount awarded for continuation awards in year 4. This is being done to create a pool of funds for supplemental performance based awards (described below). [Note: Applicants should not reduce their requested fifth year amounts relative to year 4; this adjustment will be made by SAMHSA at the time the year 5 continuation awards are negotiated.]

Supplemental Awards Based on Performance: Section VI-2, Administrative and National Policy Requirements, of this RFA discusses a grantee's proposed performance targets and explains that failure to meet stated goals and objectives may result in suspension or termination of the grant award, or in the reduction or withholding of continuation awards. Conversely, an SBIRT grantee that exceeds its performance targets or demonstrates efficiencies may receive a supplemental award based on performance to maintain its high level of performance.

For year 5 of the SBIRT Cooperative Agreements, CSAT will review each grantee's Government Performance and Results Act (GPRA) data submissions and assess whether a grantee has: 1) met or exceeded its target for the number of clients served by 25 percent or more; 2) met or exceeded its target for 6-month follow-ups⁹; and 3) provided services within approved cost-bands. Any grantee that has demonstrated appropriate financial management of the grant and has exceeded its targets for the number of clients served by 25 percent or more, exceeded its target for 6-month follow-ups, and provided services within allowable cost bands, may receive a supplemental award of up to 5 percent of the fifth year requested amount based on performance. Supplemental award amounts will be determined on a sliding scale based on availability of funds and the grantee's achievement of performance goals and demonstration of sound fiscal management. **Applicants should be aware that SAMHSA/CSAT does not plan to make supplemental awards to all grantees, and that it is possible that no grantees will receive supplemental awards based on performance.**

Eligible grantees will be asked to submit a narrative and budget justification for the supplemental award that maintains the increase in its targets during the final year of the project. The supplemental award based on performance is for the purpose of the grantee maintaining, at a minimum, the additional number of clients for the remainder the project. A grantee receiving a supplemental award based on performance may be subject to additional site visits and/or audits to verify the accuracy of the client data reported.

⁹ The follow-up rate must be at least 80 percent of the number of clients selected as part of the follow-up sample.

Cooperative Agreement

These awards are being made as cooperative agreements because they require substantial post-award Federal programmatic participation in the conduct of the project. Under this cooperative agreement, the roles and responsibilities of grantees and SAMHSA staff are:

Role of Grantee:

- Comply with the terms and conditions of the cooperative agreement award.
- Agree to provide SAMHSA with data required for GPRA.
- Collaborate with CSAT staff in project implementation and monitoring.
- Select members and organize/conduct regular meetings of the project's Policy Steering Committee (PSC).
- Implement and monitor activities of the cooperative agreement project, including accountability for sub-recipients' service delivery.
- Collect, evaluate, and report Statewide treatment project and GPRA data.
- Respond to requests for program-related data.
- Document intended and actual systemic changes resulting from the project's activities.
- Submit the final Project Implementation Plan by the end of the third month following the award.
- Participate in the CSAT cross-site SBIRT evaluation.

Role of SAMHSA Staff:

- Collaborate in selection of PSC members; review and approve final membership.
- Work collaboratively with the grantee, project staff, and PSC members to finalize the Project Implementation Plan; review and approve the plan to be submitted by the end of the third month for release of funds for implementation (i.e., Phases II and III).
- Provide best practice program information, resource materials, and technical assistance, e.g., examples of model programs, financing strategies and benefit designs, and screening and assessment tools/protocols to help grantees identify, select, and replicate evidence-based practices for implementing SBIRT.
- Provide guidance on how to assess resource allocation strategies in order to re-direct treatment resources toward an emphasis on persons at risk, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence).
- Review and approve sub-recipient contracts and awards.
- Actively participate in PSC discussions.
- Work cooperatively with the grantee to make the transition from the cooperative agreement to State/Tribal and local control and to sustain the system changes achieved by the project.

Policy Steering Committee will:

- Provide a freestanding policy steering committee to provide strategic policy and operational advice on the SBIRT project to the grantee as well as provide advice on integrating SBIRT into the existing system of care and on policies, as appropriate.

- Consist of 15 to 20 members and a chair, to be appointed by the grantee, with CSAT's approval.
- Represent the Office of the Governor/highest ranking official and diverse stakeholders in the State/Tribe, including, for example, representatives from:
 - Relevant State agencies (including the SSA)
 - Member of the State Medicaid Agency
 - Community specialist treatment organizations
 - General and specialist healthcare organizations (e.g., Federally Qualified Community Health Centers, hospitals, family practice clinics, emergency departments, ob-gyn clinics)
 - Occupational health clinics and employee assistance programs or human resources departments
 - Student health centers and student assistance programs
 - Unions and member assistance programs
 - Financial organizations (e.g., insurers, fiscal intermediaries, employers)
 - Professional and trade associations
 - Recovery community organizations
 - Community coalitions
 - Training agencies and universities
 - Employers and business coalitions
 - Insurers and managed care organizations
- Hold the initial meeting within 60 days of award and continue to meet once a month for the first year and quarterly in subsequent years.
- Coordinate with other State/Tribal agencies, commissions, and offices (including the SSA) as appropriate.

III. ELIGIBILITY INFORMATION

1. ELIGIBLE APPLICANTS

Eligible applicants are the immediate office of the Chief Executive (e.g., Governor) in the States, Territories, and the District of Columbia; or the highest ranking official and/or the duly authorized official of a federally recognized American Indian/Alaska Native tribe or tribal organization. Tribal organization means the recognized body of any AI/AN tribe; any legally established organization of American Indians/Alaska Natives which is controlled, sanctioned, or chartered by such governing body or which is democratically elected by the adult members of the Indian community to be served by such organization and which includes the maximum participation of American Indians/Alaska Natives in all phases of its activities. **Eligible applicants in the same geographic region may apply as a consortium and must designate a single fiscal agent.**

The Governor/highest ranking official will designate a lead official to be Project Director for the cooperative agreement. That individual may be, but is not required to be, part of the SSA. However, the services to be provided through this cooperative agreement program are to be

integrated into the current system of care. Therefore, SAMHSA expects that the SSA will be involved in the project.

The Chief Executive of the State, Territory, or District of Columbia, or the highest ranking official and/or the duly authorized official of the Tribal Organization must sign the application. Following the initial award, the Chief Executive or highest ranking official may delegate responsibility for the grant, including signatory authority for continuation applications, to a State Agency, State Official, or duly authorized official.

Updated on Nov. 29, 2007: States/tribes that have been recipients of SBIRT cooperative agreements are not eligible to apply.

2. COST SHARING and MATCH REQUIREMENTS

Cost sharing/match are not required in this program.

3. OTHER

3.1 Additional Eligibility Requirements

You must comply with the following requirements, or your application will be screened out and will not be reviewed: use of the PHS 5161-1 application form; application submission requirements in Section IV-3 of this document; and formatting requirements provided in Appendix A of this document.

3.2 Evidence of Experience and Credentials

SAMHSA believes that only existing, experienced, and appropriately credentialed organizations with demonstrated infrastructure and expertise will be able to provide required services quickly and effectively. You must meet three additional requirements related to the provision of services.

The three requirements are:

- A provider organization for direct client services (e.g., substance abuse treatment) appropriate to the grant must be involved in the proposed project. The provider may be the applicant or another organization committed to the project. More than one provider organization may be involved;
- Each direct service provider organization must have at least 2 years experience (as of the due date of the application) providing relevant services in the geographic area(s) in which services are to be provided (official documents must establish that the organization has provided relevant services for the last 2 years); and
- Each direct service provider organization must comply with all applicable local (city, county) and State/tribal licensing, accreditation, and certification requirements, as of the due date of the application.

[Note: The above requirements apply to all service provider organizations. A license from an individual clinician will not be accepted in lieu of a provider organization's license.]

In **Appendix 1** of your application, you must: (1) identify at least one experienced, licensed service provider organization; (2) include a list of all direct service provider organizations that have agreed to participate in the proposed project, including the applicant agency if the applicant is a treatment or prevention service provider organization; and (3) include the Statement of Assurance (provided in Appendix C of this announcement), signed by the authorized representative of the applicant organization identified on the face-page (SF 424 v2) of the application, attesting that all participating service provider organizations:

- meet the 2-year experience requirement;
- meet applicable licensing, accreditation, and certification requirements; and
- if the application is within the funding range for grant award, the applicant will provide the Government Project Officer (GPO) with the required documentation within the time specified.

In addition, if, following application review, your application's score is within the funding range, the GPO will call you and request that the following documentation be sent by overnight mail:

- a letter of commitment that specifies the nature of the participation and what service(s) will be provided from every service provider organization that has agreed to participate in the project;
- official documentation that all participating organizations have been providing relevant services for a minimum of 2 years before the date of the application in the area(s) in which the services are to be provided; and
- official documentation that all participating service provider organizations comply with all applicable local (city, county) and State/tribal requirements for licensing, accreditation, and certification or official documentation from the appropriate agency of the applicable State/tribal, county, or other governmental unit that licensing, accreditation, and certification requirements do not exist.

If the GPO does not receive this documentation within the time specified, the application will not be considered for an award.

IV. APPLICATION AND SUBMISSION INFORMATION

1. ADDRESS TO REQUEST APPLICATION PACKAGE

You may request a complete application kit from the SAMHSA Information Line at 1-877-SAMHSA7 [TDD: 1-800-487-4889].

You also may download the required documents from the SAMHSA Web site at www.samhsa.gov/grants/apply.aspx

Additional materials available on this Web site include:

- a grant writing technical assistance manual for potential applicants;
- standard terms and conditions for SAMHSA grants;
- guidelines and policies that relate to SAMHSA grants (e.g., guidelines on cultural competence, consumer and family participation, and evaluation); and
- a list of certifications and assurances referenced in item 21 of the SF 424 v2.

2. CONTENT AND FORM OF APPLICATION SUBMISSION

2.1 Application Kit

SAMHSA application kits include the following documents:

- PHS 5161-1 (revised July 2000) – Includes the face page (SF 424 v2), budget forms, assurances, certification, and checklist. You must use the PHS 5161-1. **Applications that are not submitted on the required application form will be screened out and will not be reviewed.**
- Request for Applications (RFA) – Provides a description of the program, specific information about the availability of funds, and instructions for completing the grant application. This document is the RFA. The RFA will be available on the SAMHSA Web site (www.samhsa.gov/grants/index.aspx) and a synopsis of the RFA is available on the Federal grants Web site (www.Grants.gov).

You must use all of the above documents in completing your application.

2.2 Required Application Components

Applications must include the required ten application components (Face Page, Abstract, Table of Contents, Budget Form, Project Narrative and Supporting Documentation, Appendices, Assurances, Certifications, Disclosure of Lobbying Activities, and Checklist).

- **Face Page** – SF 424 v2 is the face page. This form is part of the PHS 5161-1. [Note: Applicants must provide a Dun and Bradstreet (DUNS) number to apply for a grant or cooperative agreement from the Federal Government. SAMHSA applicants are required to provide their DUNS number on the face page of the application. Obtaining a DUNS number is easy and there is no charge. To obtain a DUNS number, access the Dun and Bradstreet Web site at www.dunandbradstreet.com or call 1-866-705-5711. To expedite the process, let Dun and Bradstreet know that you are a public/private nonprofit organization getting ready to submit a Federal grant application.]

- ❑ **Abstract** – Your total abstract should not be longer than 35 lines. It should include the project name, population to be served (demographics and clinical characteristics), strategies/interventions, project goals and measurable objectives, including the number of people to be served annually and throughout the lifetime of the project, etc. In the first five lines or less of your abstract, write a summary of your project that can be used, if your project is funded, in publications, reporting to Congress, or press releases.
- ❑ **Table of Contents** – Include page numbers for each of the major sections of your application and for each appendix.
- ❑ **Budget Form** – Use SF 424A, which is part of the PHS 5161-1. Fill out Sections B, C, and E of the SF 424A. A sample budget and justification is included in Appendix H of this document.
- ❑ **Project Narrative and Supporting Documentation** – The Project Narrative describes your project. It consists of Sections A through E. Sections A-E together may not be longer than 30 pages. (Remember that if your Project Narrative starts on page 5 and ends on page 35, it is 31 pages long, not 30 pages.) More detailed instructions for completing each section of the Project Narrative are provided in “Section V – Application Review Information” of this document.

The Supporting Documentation provides additional information necessary for the review of your application. This supporting documentation should be provided immediately following your Project Narrative in Sections F through I. There are no page limits for these sections, except for Section H, Biographical Sketches/Job Descriptions. Additional instructions for completing these sections are included in Section V under “Supporting Documentation.” Supporting documentation should be submitted in black and white (no color).

- ❑ **Appendices 1 through 5** – Use only the appendices listed below. If your application includes any appendices not required in this document, they will be disregarded. Do not use more than a total of 30 pages for Appendices 1, 3 and 4 combined. There are no page limitations for Appendices 2 and 4. Do not use appendices to extend or replace any of the sections of the Project Narrative. Reviewers will not consider them if you do.
 - *Appendix 1:* (1) Identification of at least one experienced, licensed service provider organization; (2) a list of all direct service provider organizations that have agreed to participate in the proposed project, including the applicant agency, if it is a treatment or prevention service provider organization; (3) the Statement of Assurance (provided in Appendix C of this announcement) signed by the authorized representative of the applicant organization identified on the face page of the application, that assures SAMHSA that all listed providers meet the 2-year experience requirement, are appropriately licensed, accredited, and certified, and that if the application is within the funding range for an award, the applicant will send the GPO the required documentation within the specified time; (4) letters of commitment/support.
 - *Appendix 2:* Data Collection Instruments/Interview Protocols

- *Appendix 3: Sample Consent Forms*
 - *Appendix 4: Letter to the SSA (if applicable; see Section IV-4 of this document)*
 - *Appendix 5: Training and Technical Assistance Plan and Signed Agreement*
- ❑ **Assurances** – Non-Construction Programs. You must read the list of assurances provided on the SAMHSA Web site or in the application kit before signing the face page (SF 424 v2) of the application. You are also required to complete the Assurance of Compliance with SAMHSA Charitable Choice Statutes and Regulations Form SMA 170. This form will be posted on SAMHSA’s Web site with the RFA and provided in the application kits.
 - ❑ **Certifications** – You must read the list of certifications provided on the SAMHSA Web site or in the application kit before signing the face page (SF 424 v2) of the application.
 - ❑ **Disclosure of Lobbying Activities** – You must submit Standard Form LLL found in the PHS 5161-1. Federal law prohibits the use of appropriated funds for publicity or propaganda purposes, or for the preparation, distribution, or use of the information designed to support or defeat legislation pending before the Congress or State legislatures. This includes “grass roots” lobbying, which consists of appeals to members of the public suggesting that they contact their elected representatives to indicate their support for or opposition to pending legislation or to urge those representatives to vote in a particular way. If no lobbying is to be disclosed, mark N/A on the form.
 - ❑ **Checklist** – Use the Checklist found in PHS 5161-1. The Checklist ensures that you have obtained the proper signatures, assurances and certifications. If you are submitting a paper application, the Checklist should be the last page.

2.3 Application Formatting Requirements

Please refer to *Appendix A, Checklist for Formatting Requirements and Screenout Criteria for SAMHSA Grant Applications*, for SAMHSA’s basic application formatting requirements. Applications that do not comply with these requirements will be screened out and will not be reviewed.

3. SUBMISSION DATES AND TIMES

Applications are due by close of business on **January 31, 2008**. Hard copy applications are due by 5:00 PM (EST). Electronic applications are due by 12:00 midnight (EST). **Hand carried applications will not be accepted. Applications may be shipped using only DHL, Federal Express (FedEx), United Parcel Service (UPS), or the United States Postal Service (USPS).**

You will be notified by postal mail that your application has been received.

Your application must be received by the application deadline or it will not be considered for review. Please remember that mail sent to Federal facilities undergoes a security screening

prior to delivery. You are responsible for ensuring that you submit your application so that it will arrive by the application due date and time.

If an application is mailed to a location or office (including room number) that is not designated for receipt of the application and, as a result, the designated office does not receive your application by the deadline, your application will be considered late and ineligible for review.

SAMHSA will not accept or consider any applications sent by facsimile.

SAMHSA accepts electronic submission of applications through www.Grants.gov. Please refer to Appendix B for “Guidance for Electronic Submission of Applications.”

4. INTERGOVERNMENTAL REVIEW (E.O. 12372) REQUIREMENTS

This grant program is covered under Executive Order (EO) 12372, as implemented through Department of Health and Human Services (DHHS) regulation at 45 CFR Part 100. Under this Order, States may design their own processes for reviewing and commenting on proposed Federal assistance under covered programs. Certain jurisdictions have elected to participate in the EO process and have established State Single Points of Contact (SPOCs). A current listing of SPOCs is included in the application kit and can be downloaded from the Office of Management and Budget (OMB) Web site at www.whitehouse.gov/omb/grants/spoc.html.

- Check the list to determine whether your State participates in this program. You **do not** need to do this if you are an American Indian/Alaska Native tribe or tribal organization.
- If your State participates, contact your SPOC as early as possible to alert him/her to the prospective application(s) and to receive any necessary instructions on the State’s review process.
- For proposed projects serving more than one State, you are advised to contact the SPOC of each affiliated State.
- The SPOC should send any State review process recommendations to the following address within 60 days of the application deadline. **For United States Postal Service:** Crystal Saunders, Director of Grant Review, Office of Program Services, Substance Abuse and Mental Health Services Administration, Room 3-1044, 1 Choke Cherry Road, Rockville, MD **20857**. ATTN: SPOC – Funding Announcement No. **TI-08-001**. Change the zip code to **20850** if you are using another delivery service.

In addition, if you are a community-based, non-governmental service provider and you are not transmitting your application through the State, you must submit a Public Health System Impact Statement (PHSIS)¹⁰ to the head(s) of appropriate State or local health agencies in the area(s) to

¹⁰ Approved by OMB under control no. 0920-0428; Public reporting burden for the Public Health System Reporting Requirement is estimated to average 10 minutes per response, including the time for copying the face page of SF 424 v2 and the abstract and preparing the letter for mailing. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The

be affected no later than the application deadline. The PHSIS is intended to keep State and local health officials informed of proposed health services grant applications submitted by community-based, non-governmental organizations within their jurisdictions. If you are a State or local government or American Indian/Alaska Native tribe or tribal organization, you are not subject to these requirements.

The PHSIS consists of the following information:

- a copy of the face page of the application (SF 424 v2); and
- a summary of the project, no longer than one page in length, that provides: 1) a description of the population to be served; 2) a summary of the services to be provided; and 3) a description of the coordination planned with appropriate State or local health agencies.

For SAMHSA grants, the appropriate State agencies are the Single State Agencies (SSAs) for substance abuse and mental health. A listing of the SSAs can be found on SAMHSA's Web site at www.samhsa.gov. If the proposed project falls within the jurisdiction of more than one State, you should notify all representative SSAs.

If applicable, you must include a copy of a letter transmitting the PHSIS to the SSA in **Appendix 4, "Letter to the SSA."** The letter must notify the State that, if it wishes to comment on the proposal, its comments should be sent not later than 60 days after the application deadline to the following address. **For United States Postal Service:** Crystal Saunders, Director of Grant Review, Office of Program Services, Substance Abuse and Mental Health Services Administration, Room 3-1044, 1 Choke Cherry Road, Rockville, MD **20857**. ATTN: SSA – Funding Announcement No. **TI-08-001**. Change the zip code to **20850** if you are using another delivery service.

In addition:

- Applicants may request that the SSA send them a copy of any State comments.
- The applicant must notify the SSA within 30 days of receipt of an award.

5. FUNDING LIMITATIONS/RESTRICTIONS

Cost principles describing allowable and unallowable expenditures for Federal grantees, including SAMHSA grantees, are provided in the following documents, which are available at www.samhsa.gov/grants/management.aspx:

- Institutions of Higher Education: OMB Circular A-21

OMB control number for this project is 0920-0428. Send comments regarding this burden to CDC Clearance Officer, 1600 Clifton Road, MS D-24, Atlanta, GA 30333, ATTN: PRA (0920-0428).

- State and Local Governments and Federally Recognized Indian Tribal Governments: OMB Circular A-87
- Nonprofit Organizations: OMB Circular A-122
- Hospitals: 45 CFR Part 74, Appendix E

In addition, SAMHSA's SBIRT grant recipients must comply with the following funding restrictions:

- No more than 15% of the total grant award may be used for developing the infrastructure necessary for expansion of services.
- No more than 20% of the total grant award may be used for data collection and performance assessment, including incentives for participating in the required data collection follow-up.

SAMHSA grantees must also comply with SAMHSA's standard funding restrictions, which are included in Appendix G.

6. OTHER SUBMISSION REQUIREMENTS

You may submit your application in either electronic or paper format:

Submission of Electronic Applications

SAMHSA accepts electronic submission of applications through www.Grants.gov. Electronic submission is voluntary. No review points will be added or deducted, regardless of whether you use the electronic or paper format.

To submit an application electronically, you must use the www.Grants.gov apply site. You will be able to download a copy of the application package from www.Grants.gov, complete it off-line, and then upload and submit the application via the Grants.gov site. E-mail submissions will not be accepted.

Please refer to Appendix B for detailed instructions on submitting your application electronically.

Submission of Paper Applications

You must submit an original application and 2 copies (including appendices). The original and copies must not be bound. Do not use staples, paper clips, or fasteners. Nothing should be attached, stapled, folded, or pasted.

Send applications to the address below:

For United States Postal Service:

Crystal Saunders, Director of Grant Review
Office of Program Services
Substance Abuse and Mental Health Services Administration
Room 3-1044
1 Choke Cherry Road
Rockville, MD **20857**

Change the zip code to **20850** if you are using another delivery service.

Do not send applications to other agency contacts, as this could delay receipt. Be sure to include “**SBIRT - TI-08-001**” in item number 12 on the face page (SF 424 v2) of any paper applications. If you require a phone number for delivery, you may use (240) 276-1199.

Hand carried applications will not be accepted. Applications may be shipped using only DHL, Federal Express (FedEx), United Parcel Service (UPS), or the United States Postal Service (USPS).

SAMHSA will not accept or consider any applications sent by facsimile.

V. APPLICATION REVIEW INFORMATION

1. EVALUATION CRITERIA

The Project Narrative describes what you intend to do with your project and includes the Evaluation Criteria in Sections A-E below. Your application will be reviewed and scored according to the quality of your response to the requirements in Sections A-E.

- In developing the Project Narrative section of your application, use these instructions, which have been tailored to this program. **These are to be used instead of the “Program Narrative” instructions found in the PHS 5161-1.**
- The Project Narrative (Sections A-E) together may be no longer than 30 pages.
- You must use the five sections/headings listed below in developing your Project Narrative. Be sure to place the required information in the correct section, **or it will not be considered**. Your application will be scored according to how well you address the requirements for each section of the Project Narrative.
- Reviewers will be looking for evidence of cultural competence in each section of the Project Narrative, and will consider how well you address the cultural competence aspects of the evaluation criteria when scoring your application. SAMHSA’s guidelines for cultural competence can be found on the SAMHSA Web site at www.samhsa.gov. Click on “Grants/Applying for a New SAMHSA Grant/Guidelines for Assessing Cultural Competence.”

- The Supporting Documentation you provide in Sections F-I and Appendices 1-4 will be considered by reviewers in assessing your response, along with the material in the Project Narrative.
- The number of points after each heading is the maximum number of points a review committee may assign to that section of your Project Narrative. Although scoring weights are not assigned to individual bullets, applicants are encouraged to respond to each bulleted statement.

Section A: Statement of Need (10 points)

- Describe the need for treatment in the State/Tribe and for each community in which SBIRT will be implemented. Include as much documentation as possible, with the focus on differentiating clinically appropriate treatment for persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence).
- Describe for the State/Tribe and for each community in which SBIRT will be implemented the current resources and continuum of care for persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence), including the provider and practitioner resources, and the funding streams available for intervention and treatment services in the generalist and specialist systems. Explain why the existing services are insufficient or inappropriate to respond to the demand for services and the treatment needs of the target population chosen for this application.
- Describe how the applicant currently plans for, funds, and provides intervention and treatment services within its continuum of care (including SBIRT if it is part of the current continuum of care) and how SBIRT (or SBIRT expansion) can be integrated into the financing and provider systems. Include a discussion of the use of patient placement criteria and standardized screening and sequential assessment protocols, if these are used, and the modalities in which persons are placed. If the modalities that the applicant funds do not match those for which CSAT calculates program costs, provide a crosswalk that aligns your modalities and costs for each with those that CSAT tracks for GPRA.
- Describe for the State/Tribe and for each community in which SBIRT will be implemented the potential for policy and systems change and the strategies to rapidly initiate SBIRT.

Section B: Proposed Evidence-Based Service/Practice (30 points)

- Clearly state the purpose, goals and objectives of your proposed project. Describe how achievement of the goals will produce meaningful and relevant results (e.g., increase access, availability, prevention, outreach, pre-services, treatment, and/or intervention) and support SAMHSA's goals for the program.
- Identify the evidence-based service/practice that you propose to implement for each treatment modality (BI, BT, RT) and the source of your information. (See Section I-2.1,

Using Evidence-Based Practices.) Discuss the evidence that shows that this practice is effective with the target population. If the evidence is limited or non-existent for the target population, provide other information to support your selection of the intervention for the target population.

- Discuss any screening tools you plan to use other than the ASSIST and CRAFFT tools. Provide a justification for the use of these additional tools, including the evidence that supports their use with the target population.
- Document the evidence that the practice you have chosen is appropriate for the outcomes you want to achieve.
- Identify and justify any modifications or adaptations you will need to make to the proposed practice to meet the goals of your project and why you believe the changes will improve the outcomes.
- Explain why you chose this evidence-based practice over other evidence-based practices. If this is not an evidence-based practice, explain why you chose this intervention over other interventions.
- Describe how the proposed project will address issues of age, race, ethnicity, culture, language, sexual orientation, disability, literacy, and gender in the target population, while retaining fidelity to the chosen practice.
- Demonstrate how the proposed service/practice will meet your goals and objectives. Provide a logic model that links need, the services or practice to be implemented, and outcomes. (See Appendix D for a sample logic model.)

Section C: Proposed Implementation Approach (35 points)

- Provide a detailed Project Implementation Plan that explains how you propose to use project funds in conjunction with other available funding sources to provide SBIRT services. All funding sources that are or could be used to pay for screening and treatment of Substance Use Disorders (e.g., State General Fund, Medicaid, Medicare, Preventive Health and Health Services Block Grant, Community Health Center grants, commercial insurance, Substance Abuse Prevention and Treatment Block Grant, Temporary Assistance for Needy Families (TANF) Block Grant, Child Care and Development (CCDF) Block Grant, Maternal and Child Health Block Grant) should be addressed, but the focus should be on the three major funding streams that you will use to increase support and decrease barriers.
- Provide a realistic time line for the entire project period (chart or graph) showing key activities, milestones, and responsible staff. [Note: The time line should be part of the Project Narrative. It should not be placed in an appendix.]

- Describe how you will increase the number of generalist settings that provide SBIRT in each sub-recipient community as a result of the award and redirection of other funding sources.
- Describe how you will provide SBIRT within its continuum of care, within the geographic areas proposed, including a description of the modalities and services to be provided, the protocols that will be used for standardizing screening, assessment, determining the level of service required, referral, brief intervention, brief treatment, and follow up. Provide a justification for the procedures to be used, including a discussion of the evidence-based services/practices that you propose to implement.
- Describe how you will screen and assess clients for Substance Use Disorders involving prescription medications and for the presence of co-occurring substance use (abuse and dependence) and mental disorders and use the information obtained from the screening and assessment to develop appropriate treatment approaches for the persons identified as having such co-occurring disorders.
- Discuss the target population's language, beliefs, norms and values, as well as socioeconomic factors that must be considered in delivering programs to this population, and how the proposed approach addresses these issues.
- If you choose to expend funds for other treatment modalities within the continuum of care, describe how these services will be implemented. Applicants that do not seek to fund specific components of their continuum of care through the SBIRT cooperative agreement must provide evidence that there is a sufficient amount of services in those elements of the continuum (modalities) for each community to be included.
- Provide a plan describing existing barriers to treatment (e.g., laws, regulations, eligibility, requirements for service receipt, facility and provider eligibility requirements, varied funding streams, coverage limitations, a lack of patient placement criteria or standardized screening and sequential assessment protocols, etc.) in the State/Tribe and in each community in which SBIRT will be implemented. Include in the plan a discussion of how you will overcome barriers to accessing clinically appropriate care, using the SBIRT approach. Whenever possible, apply findings from recent literature and other information that demonstrates a thorough understanding of the issues faced in introducing SBIRT into the applicant's continuum of care. Include literature citations in Section F, Supporting Documentation, of your application.
- Describe the linkages to be developed between the participating specialist and community agencies for referrals, cooperation in case management, and information sharing.
- Provide a plan to make available training and technical assistance to sub-recipient communities, including information about SBIRT methods, training for staff in the community and specialist settings in carrying out SBIRT, and technical assistance to the

overall project and its sub-recipients.¹¹ Include the plan and signed agreement in **Appendix 5** of the application. You must also include this activity in your budget and budget justification in **Section G** of your application.

- Describe how you will increase access and availability of services to a larger number of persons at risk, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence) as a result of the award. State clearly the number of additional persons to be served (annually and over the entire project period) through the proposed cooperative agreement in each element of SBIRT (i.e., number of persons projected to receive Screening Only, number of persons projected to receive Screening and Brief Intervention [BI], and number of persons projected to receive Screening and Brief Treatment [BT] or Screening and Referral to Other Types of Treatment for Substance Use Disorders [RT]) and the number of persons to receive clinically appropriate treatment in all other modalities within the system. Show that the targets are feasible and reasonable. Describe how the target population will be identified, recruited, and retained.
- Describe the expected outcomes of treatment (e.g., decreased drug use in those patients receiving services through SBIRT) and the means by which you determined these targets. Show that the targets are feasible and reasonable.
- Describe your plan to continue the project after the funding period ends. Also describe how program continuity will be maintained when there is a change in the operational environment (e.g., staff turnover, change in project leadership) to ensure stability over time.

Section D: Staff and Organizational Experience (10 points)

- Discuss the capability and experience of the applicant organization and other participating organizations with similar projects and populations. Demonstrate that the applicant organization and other participating organizations have linkages to the target population and ties to grassroots/community-based organizations that are rooted in the culture and language of the target population.
- Provide a complete list of staff positions for the project, showing the role of each and their level of effort and qualifications. Include the Project Director and other key personnel, such as treatment/prevention personnel.
- Discuss how key staff have demonstrated experience in serving the target population and are familiar with the culture and language of the target population. If the target population is multicultural and multilingual, describe how the staff are qualified to serve this population.

¹¹ The State may wish to consider subcontracting with the SAMHSA/CSAT funded Addiction Technology Training Centers (ATTCs) already working with its SSA or an in-State resource. A list of ATTCs, the States covered by each, and contact information is provided in Appendix I. If a subcontract with the ATTC, another academic institution, or a vendor is used, the plan should include the cost for providing these activities as a separate budget component.

- Describe the resources available for the proposed project (e.g., facilities, equipment), and provide evidence that services will be provided in a location that is adequate, accessible, compliant with the Americans with Disabilities Act (ADA), and amenable to the target population. If the ADA does not apply to your organization, please explain why.

Section E: Performance Assessment and Data (15 points)

- Document your ability to collect and report on the required performance measures as specified in Section I-2.4 of this RFA. Describe your plan for data collection, management, analysis and reporting. Specify and justify any additional measures or instruments you plan to use for your grant project.
- Describe how data will be used to manage the project and assure continuous quality improvement.
- Provide a per-person or unit cost of the project to be implemented. You can calculate this figure by: 1) taking the total cost of the project over the lifetime of the grant and subtracting 20% for data and performance assessment; 2) dividing this number by the total unduplicated (i.e., each client can be counted only one time regardless of the number of visits) number of persons to be served. The applicant will be required to project costs by year of the grant to show progress on cost effectiveness.

Program Costs. The following are considered reasonable ranges by treatment modality:

- Residential: \$3,000 to \$10,000
- Outpatient (Non-Methadone): \$1,000 to \$5,000
- Outpatient (Methadone): \$1,500 to \$8,000
- Intensive Outpatient: \$1,000 to \$7,500
- Screening/Brief Intervention/Brief Treatment/Outreach/Pretreatment Services: \$200 to \$1,200
- Drug Court Programs (regardless of client treatment modality): \$3,000 to \$5,000
- Peer Recovery Support Services: \$1,000 to \$2,500

The outreach and pretreatment services cost band applies only to outreach and pretreatment programs that do not offer treatment services but operate with a network of substance abuse treatment facilities. Treatment programs that add outreach and pretreatment services to a treatment modality or modalities are expected to fall within the cost band for that treatment modality.

- Describe your plan for conducting the performance assessment as specified in Section I-2.5 of this RFA and document your ability to conduct the assessment.

NOTE: Although the budget for the proposed project is not a scored review criterion, the Review Group will be asked to comment on the appropriateness of the budget after the merits of the application have been considered.

SUPPORTING DOCUMENTATION

Section F: Literature Citations. This section must contain complete citations, including titles and all authors, for any literature you cite in your application.

Section G: Budget Justification, Existing Resources, Other Support. You must provide a narrative justification of the items included in your proposed budget, as well as a description of existing resources and other support you expect to receive for the proposed project. Be sure to show that no more than 15% of the total grant award will be used for infrastructure development, if necessary, and that no more than 20% of the total grant award will be used for data collection and performance assessment. An illustration of a budget and narrative justification is included in Appendix H of this document.

Section H: Biographical Sketches and Job Descriptions.

- Include a biographical sketch for the Project Director and other key positions. Each sketch should be 2 pages or less. If the person has not been hired, include a position description and/or a letter of commitment with a current biographical sketch from the individual.
- Include job descriptions for key personnel. Job descriptions should be no longer than 1 page each. Level of effort by percentage of FTE should be included.
- Information on what should be included in biographical sketches and job descriptions can be found on page 22, Item 6, in the Program Narrative section of the PHS 5161-1 instruction page, available on the SAMHSA Web site.

Section I: Confidentiality and SAMHSA Participant Protection/Human Subjects: You must describe procedures relating to Confidentiality, Participant Protection and the Protection of Human Subjects Regulations in Section I of your application, using the guidelines provided below.

Confidentiality and Participant Protection:

Because of the confidential nature of the work in which many SAMHSA grantees are involved, it is important to have safeguards protecting individuals from risks associated with their participation in SAMHSA projects. All applicants must address the seven bullets below. Appendix F of this RFA provides a more detailed discussion of issues applicants should consider in addressing these seven bullets. If some are not applicable or relevant to the proposed project, simply state that they are not applicable and indicate why. In addition to addressing these seven bullets, read the section that follows entitled Protection of Human Subjects Regulations to determine if the regulations may apply to your project. If so, you are required to describe the process you will follow for obtaining Institutional Review Board (IRB) approval. While we encourage you to keep your responses brief, there are no page limits for this section and no points will be assigned by the Review Committee. Problems with confidentiality, participant protection, and the protection of human subjects identified during peer review of the application must be resolved prior to funding.

- ❑ Identify foreseeable risks or adverse effects due to participation in the project and/or in the data collection (performance assessment) activities (including physical, medical, psychological, social, legal, and confidentiality) and provide your procedures for minimizing or protecting participants from these risks. Identify plans to provide guidance and assistance in the event there are adverse effects to participants.
- ❑ Describe the target population and explain why you are including or excluding certain subgroups. Explain how and who will recruit and select participants.
- ❑ State whether participation in the project is voluntary or required. If you plan to provide incentives/compensate participants, specify the type (e.g., money, gifts, coupons), and the value of any such incentives. Provide justification that the use of incentives is appropriate, judicious, and conservative and that incentives do not provide an “undue inducement” which removes the voluntary nature of participation. Incentives should be the minimum amount necessary to meet the programmatic and performance assessment goals of the grant. Applicants should determine the minimum amount that is proven to be effective by consulting with existing local programs and reviewing the relevant literature. In no case may the value of an incentive paid for with SAMHSA discretionary grant funds exceed \$20. (See Appendix F: Confidentiality and Participant Protection.) For the purpose of this project, incentives may be provided for follow-up only.
- ❑ Describe data collection procedures, including sources (e.g., participants, school records) and the data collecting setting (e.g., clinic, school). Provide copies of proposed data collection instruments and interview protocols in **Appendix 2** of your application, “Data Collection Instruments/Interview Protocols.” State whether specimens such as urine and/or blood will be obtained and the purpose for collecting the specimens. If applicable, describe how the specimens and process will be monitored to ensure both the safety of participants and the integrity of the specimens.
- ❑ Explain how you will ensure privacy and confidentiality of participants’ records, data collected, interviews, and group discussions. Describe where the data will be stored, safeguards (e.g., locked, coding systems, storing identifiers separate from data), and who will have access to the information.
- ❑ Describe the process for obtaining and documenting consent from adult participants and assent from minors along with consent from their parents or legal guardians. Provide copies of all consent forms in **Appendix 3** of your application, “Sample Consent Forms.” If needed, give English translations.
- ❑ Discuss why the risks are reasonable compared to expected benefits from the project.

Protection of Human Subjects Regulations

SAMHSA expects that most grantees funded under this announcement will not have to comply with the Protection of Human Subjects Regulations (45 CFR 46), which requires Institutional Review Board (IRB) approval. However, in some instances, the applicant’s proposed

performance assessment design may meet the regulation’s criteria of research involving human subjects. For assistance in determining if your proposed performance assessment meets the criteria in 45 CFR 46, Protection of Human Subjects Regulations, refer to the SAMHSA decision tree on the SAMHSA Web site, under “Applying for a New SAMHSA Grant,” <http://www.samhsa.gov/grants/apply.aspx>.

Applicants whose projects must comply with the Human Subjects Regulations must, in addition to the bullets above, fully describe the process for obtaining IRB approval. While IRB approval is not required at the time of grant award, these grantees will be required, as a condition of award, to provide documentation that an Assurance of Compliance is on file with the Office for Human Research Protections (OHRP). IRB approval must be received in these cases prior to enrolling clients in the project. General information about Human Subjects Regulations can be obtained through OHRP at <http://www.hhs.gov/ohrp>, or ohrp@osophs.dhhs.gov, or (240) 453-6900. SAMHSA–specific questions should be directed to the program contact listed in Section VII of this announcement.

2. REVIEW AND SELECTION PROCESS

SAMHSA applications are peer-reviewed according to the evaluation criteria listed above. For those programs where the individual award is over \$100,000, applications also must be reviewed by the appropriate National Advisory Council.

Decisions to fund a grant are based on:

- the strengths and weaknesses of the application as identified by peer reviewers and, when applicable, approved by the Center for Substance Abuse Treatment’s National Advisory Council;
- availability of funds; and
- equitable distribution of awards in terms of geography (including urban, rural and remote settings) and balance among target populations and program size.

VI. ADMINISTRATION INFORMATION

1. AWARD NOTICES

After your application has been reviewed, you will receive a letter from SAMHSA through postal mail that describes the general results of the review, including the score that your application received.

If you are approved for funding, you will receive an **additional** notice through postal mail, the Notice of Grant Award, signed by SAMHSA’s Grants Management Officer. The Notice of Grant Award is the sole obligating document that allows you to receive Federal funding for work on the grant project.

If you are not funded, you may re-apply if there is another receipt date for the program.

2. ADMINISTRATIVE AND NATIONAL POLICY REQUIREMENTS

- If your application is funded, you must comply with all terms and conditions of the grant award. SAMHSA’s standard terms and conditions are available on the SAMHSA Web site at <http://www.samhsa.gov/grants/management.aspx>.
- If your application is funded, you must also comply with the administrative requirements outlined in 45 CFR Part 74 or 45 CFR Part 92, as appropriate. For more information see the SAMHSA Web site (<http://www.samhsa.gov/grants/management.aspx>).
- Depending on the nature of the specific funding opportunity and/or your proposed project as identified during review, SAMHSA may negotiate additional terms and conditions with you prior to grant award. These may include, for example:
 - actions required to be in compliance with confidentiality and participant protection/human subjects requirements;
 - requirements relating to additional data collection and reporting;
 - requirements relating to participation in a cross-site evaluation; or
 - requirements to address problems identified in review of the application.
- If your application is funded, you will be held accountable for the information provided in the application relating to performance targets. SAMHSA program officials will consider your progress in meeting goals and objectives, as well as your failures and strategies for overcoming them, when making an annual recommendation to continue the grant and the amount of any continuation award. Failure to meet stated goals and objectives may result in suspension or termination of the grant award, or in reduction or withholding of continuation awards.
- Grant funds cannot be used to supplant current funding of existing activities. “Supplant” is defined as replacing funding of a recipient’s existing program with funds from a Federal grant.
- In an effort to improve access to funding opportunities for applicants, SAMHSA is participating in the U.S. Department of Health and Human Services “Survey on Ensuring Equal Opportunity for Applicants.” This survey is included in the application kit for SAMHSA grants and is posted on the SAMHSA Web site. You are encouraged to complete the survey and return it, using the instructions provided on the survey form.

3. REPORTING REQUIREMENTS

In addition to the data reporting requirements listed in Section I-2.4, you must comply with the following reporting requirements:

3.1 Progress and Financial Reports

- You will be required to submit annual and final progress reports, as well as annual and final financial status reports.
- Because SAMHSA is extremely interested in ensuring that treatment and prevention services can be sustained, your progress reports should explain plans to ensure the sustainability of efforts initiated under this grant.
- If your application is funded, SAMHSA will provide you with guidelines and requirements for these reports at the time of award and at the initial grantee orientation meeting after award. SAMHSA staff will use the information contained in the reports to determine your progress toward meeting its goals.

3.2 Government Performance and Results Act (GPRA)

The Government Performance and Results Act (GPRA) mandates accountability and performance-based management by Federal agencies. To meet the GPRA requirements, SAMHSA must collect performance data (i.e., “GPRA data”) from grantees. The performance requirements for SAMHSA’s SBIRT grant program are described in Section I-2.4 of this document under “Data Collection and Performance Measurement.”

3.3 Publications

If you are funded under this grant program, you are required to notify the Government Project Officer (GPO) and SAMHSA’s Publications Clearance Officer (240-276-2130) of any materials based on the SAMHSA-funded grant project that are accepted for publication.

In addition, SAMHSA requests that grantees:

- Provide the GPO and SAMHSA Publications Clearance Officer with advance copies of publications.
- Include acknowledgment of the SAMHSA grant program as the source of funding for the project.
- Include a disclaimer stating that the views and opinions contained in the publication do not necessarily reflect those of SAMHSA or the U.S. Department of Health and Human Services, and should not be construed as such.

SAMHSA encourages successful applicants to publish related to their SBIRT grant. Publications related to the effectiveness of screening and drugs are especially encouraged.

SAMHSA reserves the right to issue a press release about any publication deemed by SAMHSA to contain information of program or policy significance to the substance abuse treatment/substance abuse prevention/mental health services community.

VII. AGENCY CONTACTS

For questions about program issues contact:

Tom Stegbauer
Center for Substance Abuse Treatment
Substance Abuse and Mental Health Services Administration
1 Choke Cherry Lane
Room 5-1099
Rockville, Maryland 20850
(240) 276-2965
tom.stegbauer@samhsa.hhs.gov

For questions on grants management issues contact:

Kathleen Sample
Office of Program Services, Division of Grants Management
Substance Abuse and Mental Health Services Administration
1 Choke Cherry Road
Room 7-1089
Rockville, Maryland 20857
(240) 276-1407
kathleen.sample@samhsa.hhs.gov

Appendix A – Checklist for Formatting Requirements and Screenout Criteria for SAMHSA Grant Applications

SAMHSA’s goal is to review all applications submitted for grant funding. However, this goal must be balanced against SAMHSA’s obligation to ensure equitable treatment of applications. For this reason, SAMHSA has established certain formatting requirements for its applications. If you do not adhere to these requirements, your application will be screened out and returned to you without review.

- Use the PHS 5161-1 application form.
- Applications must be received by the application due date and time, as detailed in Section IV-3 of this grant announcement.
- Information provided must be sufficient for review.
- Text must be legible. Pages must be typed in black ink, single-spaced, using a font of Times New Roman 12, with all margins (left, right, top, bottom) at least one inch each. (For Project Narratives submitted electronically, see separate requirements in Section IV-6 of this announcement under “Submission of Electronic Applications.”)
- To ensure equity among applications, page limits for the Project Narrative cannot be exceeded.
- Paper must be white paper and 8.5 inches by 11.0 inches in size.

To facilitate review of your application, follow these additional guidelines. Failure to adhere to the following guidelines will not, in itself, result in your application being screened out and returned without review. However, the information provided in your application must be sufficient for review. Following these guidelines will help ensure your application is complete, and will help reviewers to consider your application.

- The 10 application components required for SAMHSA applications should be included and submitted in the following order:
 - § Face Page (Standard Form 424 v2, which is in PHS 5161-1)
 - § Abstract
 - § Table of Contents
 - § Budget Form (Standard Form 424A, which is in PHS 5161-1)
 - § Project Narrative and Supporting Documentation
 - § Appendices
 - § Assurances (Standard Form 424B, which is in PHS 5161-1)
 - § Certifications
 - § Disclosure of Lobbying Activities (Standard Form LLL, which is in PHS 5161-1)
 - § Checklist (a form in PHS 5161-1)

- Applications should comply with the following requirements:
 - § Provisions relating to confidentiality and participant protection specified in Section V-1 of this announcement.
 - § Budgetary limitations as specified in Sections I, II, and IV-5 of this announcement.
 - § Documentation of nonprofit status as required in the PHS 5161-1.
- Pages should be typed single-spaced in black ink with one column per page. Pages should not have printing on both sides.
- Pages should be numbered consecutively from beginning to end so that information can be located easily during review of the application. The abstract page should be page 1, the table of contents should be page 2, etc. The four pages of Standard form 424 v2 are not to be numbered. Appendices should be labeled and separated from the Project Narrative and budget section, and the pages should be numbered to continue the sequence.
- The page limits for Appendices stated in Section IV-2.2 of this announcement should not be exceeded.
- Send the original application and two copies to the mailing address in Section IV-6 of this document. Please do not use staples, paper clips, and fasteners. Nothing should be attached, stapled, folded, or pasted. Do not use heavy or lightweight paper or any material that cannot be copied using automatic copying machines. Odd-sized and oversized attachments such as posters will not be copied or sent to reviewers. Do not include videotapes, audiotapes, or CD-ROMs.

Appendix B – Guidance for Electronic Submission of Applications

If you would like to submit your application electronically, you may search www.Grants.gov for the downloadable application package by the funding announcement number (called the opportunity number) or by the Catalogue of Federal Domestic Assistance (CFDA) number. You can find the CFDA number on the first page of the funding announcement.

You must follow the instructions in the User Guide available at the www.Grants.gov apply site, on the Help page. In addition to the User Guide, you may wish to use the following sources for help:

- By e-mail: support@Grants.gov
- By phone: 1-800-518-4726 (1-800-518-GRANTS). The Customer Support Center is open from 7:00 a.m. to 9:00 p.m. Eastern Time, Monday through Friday, excluding Federal holidays.

If this is the first time you have submitted an application through Grants.gov, you must complete four separate registration processes before you can submit your application. Allow at least two weeks (10 business days) for these registration processes, prior to submitting your application. The processes are: 1) DUNS Number registration; 2) Central Contractor Registry (CCR) registration; 3) Credential Provider registration; and 4) Grants.gov registration.

It is strongly recommended that you submit your grant application using Microsoft Office products (e.g., Microsoft Word, Microsoft Excel, etc.). If you do not have access to Microsoft Office products, you may submit PDF files. Directions for creating PDF files can be found on the Grants.gov Web site. Use of file formats other than Microsoft Office or PDF may result in your file being unreadable by our staff.

The Project Narrative must be a separate document in the electronic submission. Formatting requirements for SAMHSA grant applications are described in Appendix A of this announcement. These requirements also apply to applications submitted electronically, with the following exceptions only for Project Narratives submitted electronically in Microsoft Word. These requirements help ensure the accurate transmission and equitable treatment of applications.

- *Text legibility:* Use a font of Times New Roman 12, line spacing of single space, and all margins (left, right, top, bottom) of at least one inch each. Adhering to these standards will help to ensure the accurate transmission of your document.
- *Amount of space allowed for Project Narrative:* The Project Narrative for an electronic submission may not exceed **15,450** words. **If the Project Narrative for an electronic submission exceeds the word limit, the application will be screened out and will not be reviewed.** To determine the number of words in your Project Narrative document in Microsoft Word, select file/properties/statistics.

Keep the Project Narrative as a separate document. Please consolidate all other materials in your application to ensure the fewest possible number of attachments. Be sure to label each file according to its contents, e.g., “Appendices 1-3”, “Appendices 4-5.”

Ensure all pages in your application are numbered consecutively, with the exception of the standard forms in the PHS-5161 application package. **Documents containing scanned images must also contain page numbers to continue the sequence.** Failure to comply with these requirements may affect the successful transmission and consideration of your application.

Applicants are strongly encouraged to submit their applications to Grants.gov early enough to resolve any unanticipated difficulties prior to the deadline. You may also submit a back-up paper submission of your application. Any such paper submission must be received in accordance with the requirements for timely submission detailed in Section IV-3 of this announcement. The paper submission must be clearly marked: **“Back-up for electronic submission.”** The paper submission must conform with all requirements for non-electronic submissions. If both electronic and back-up paper submissions are received by the deadline, the electronic version will be considered the official submission.

After you electronically submit your application, you will receive an automatic acknowledgement from Grants.gov that contains a Grants.gov tracking number. It is important that you retain this number. **Include the Grants.gov tracking number in the top right corner of the face page (SF 424 v2) for any paper submission. Receipt of the tracking number is the only indication that Grants.gov has successfully received and validated your application. If you do not receive a Grants.gov tracking number, you may want to contact the Grants.gov help desk for assistance.**

The Grants.gov Web site does not accept electronic signatures at this time. Therefore, you must submit a signed paper original of the face page (SF 424 v2), the assurances (SF 424B), and hard copy of any other required documentation that cannot be submitted electronically. **You must include the Grants.gov tracking number for your application on these documents with original signatures, on the top right corner of the face page, and send the documents to the following address. The documents must be received at the following address within 5 business days after your electronic submission.** Delays in receipt of these documents may impact the score your application receives or the ability of your application to be funded.

For United States Postal Service:

Crystal Saunders, Director of Grant Review
Office of Program Services
Substance Abuse and Mental Health Services Administration
Room 3-1044
1 Choke Cherry Road
Rockville, MD **20857**
ATTN: Electronic Applications

For other delivery services, change the zip code to 20850.

If you require a phone number for delivery, you may use (240) 276-1199.

Appendix C - Statement of Assurance

As the authorized representative of [*insert name of applicant organization*]

_____, I assure SAMHSA that all participating service provider organizations listed in this application meet the two-year experience requirement and applicable licensing, accreditation, and certification requirements. If this application is within the funding range for a grant award, we will provide the SAMHSA Government Project Officer (GPO) with the following documents. I understand that if this documentation is not received by the GPO within the specified timeframe, the application will be removed from consideration for an award and the funds will be provided to another applicant meeting these requirements.

- a letter of commitment that specifies the nature of the participation and what service(s) will be provided from every service provider organization listed in Appendix 1 of the application, that has agreed to participate in the project;
- official documentation that all service provider organizations participating in the project have been providing relevant services for a minimum of 2 years prior to the date of the application in the area(s) in which services are to be provided. Official documents must definitively establish that the organization has provided relevant services for the last 2 years; and
- official documentation that all participating service provider organizations are in compliance with all local (city, county) and State/tribal requirements for licensing, accreditation, and certification or official documentation from the appropriate agency of the applicable State/tribal, county, or other governmental unit that licensing, accreditation, and certification requirements do not exist. (Official documentation is a copy of each service provider organization's license, accreditation, and certification. Documentation of accreditation will not be accepted in lieu of an organization's license. A statement by, or letter from, the applicant organization or from a provider organization attesting to compliance with licensing, accreditation and certification or that no licensing, accreditation, certification requirements exist does not constitute adequate documentation.)

Signature of Authorized Representative

Date

Appendix D – Sample Logic Model

A Logic Model is a tool to show how your proposed project links the purpose, goals, objectives, and tasks stated with the activities and expected outcomes or “change” and can help to plan, implement, and assess your project. The model also links the purpose, goals, objectives, and activities back into planning and evaluation. A Logic Model is a *picture* of your project. It graphically shows the activities and progression of the project. It should also describe the relationships among what resources you put in (inputs), what you do (outputs), and what happens or results (outcomes). Based on both your planning and evaluating activities, you can then make a “logical” chain of “if-then” relationships.

Look at the graphic on the following page to see the chain of events that links the inputs to program components, the program components to outputs, and the outputs to outcomes (goals).

The framework you set up to build your model is based on a review of your Statement of Need, in which you state the conditions that gave rise to the project with your target group. Then you look at the **Inputs**, which are the resources, contributions, time, staff, materials, and equipment you will invest to change these conditions. These inputs then are organized into the **Program Components**, which are the activities, services, interventions and tasks that will reach the target population. These outputs then are intended to create **Outputs** such as changes or benefits for the consumer, families, groups, communities, organizations and SAMHSA. The understanding and further evidence of what works and what does not work will be shown in the **Outcomes**, which include achievements that occur along the path of project operation.

*The logic model presented is not a required format and SAMHSA does not expect strict adherence to this format. It is presented only as a sample of how you can present a logic model in your application.

Sample Logic Model*

Resources (Inputs)	Program Components (Activities)	Outputs (Objectives)	Outcomes (Goals)
Examples	Examples	Examples	Examples
<p>People Staff – hours Volunteer – hours</p> <p>Funds</p> <p>Other resources Facilities Equipment Community services</p>	<p>Outreach Intake/Assessment Client Interview</p> <p>Treatment Planning Treatment by type: Methadone maintenance Weekly 12-step meetings Detoxification Counseling sessions Relapse prevention Crisis intervention</p> <p>Special Training Vocational skills Social skills Nutrition Child care Literacy Tutoring Safer sex practices</p> <p>Other Services Placement in employment Prenatal care Child care Aftercare</p> <p>Program Support Fundraising Long-range planning Administration Public Relations</p>	<p>Waiting list length Waiting list change Client attendance Client participation</p> <p>Number of Clients: Admitted Terminated Inprogram Graduated Placed</p> <p>Number of Sessions: Per month Per client/month</p> <p>Funds raised Number of volunteer hours/month</p> <p>Other resources required</p>	<p>Inprogram: Client satisfaction Client retention</p> <p>In or postprogram: Reduced drug use – self reports, urine, hair Employment/school progress Psychological status Vocational skills Social skills Safer sexual practices Nutritional practices Child care practices Reduced delinquency/crime</p>

Appendix E – Logic Model Resources

Chen, W.W., Cato, B.M., & Rainford, N. (1998-9). Using a logic model to plan and evaluate a community intervention program: A case study. *International Quarterly of Community Health Education*, 18(4), 449-458.

Edwards, E.D., Seaman, J.R., Drews, J., & Edwards, M.E. (1995). A community approach for Native American drug and alcohol prevention programs: A logic model framework. *Alcoholism Treatment Quarterly*, 13(2), 43-62.

Hernandez, M. & Hodges, S. (2003). *Crafting Logic Models for Systems of Care: Ideas into Action*. [Making children's mental health services successful series, volume 1]. Tampa, FL: University of South Florida, The Louis de la Parte Florida Mental Health Institute, Department of Child & Family Studies. <http://cfs.fmhi.usf.edu> or phone (813) 974-4651

Hernandez, M. & Hodges, S. (2001). Theory-based accountability. In M. Hernandez & S. Hodges (Eds.), *Developing Outcome Strategies in Children's Mental Health*, pp. 21-40. Baltimore: Brookes.

Julian, D.A. (1997). Utilization of the logic model as a system level planning and evaluation device. *Evaluation and Planning*, 20(3), 251-257.

Julian, D.A., Jones, A., & Deyo, D. (1995). Open systems evaluation and the logic model: Program planning and evaluation tools. *Evaluation and Program Planning*, 18(4), 333-341.

Patton, M.Q. (1997). *Utilization-Focused Evaluation* (3rd Ed.), pp. 19, 22, 241. Thousand Oaks, CA: Sage.

Wholey, J.S., Hatry, H.P., Newcome, K.E. (Eds.) (1994). *Handbook of Practical Program Evaluation*. San Francisco, CA: Jossey-Bass Inc.

Appendix F – Confidentiality and Participant Protection

1. Protect Clients and Staff from Potential Risks

- Identify and describe any foreseeable physical, medical, psychological, social, and legal risks or potential adverse effects as a result of the project itself or any data collection activity.
- Describe the procedures you will follow to minimize or protect participants against potential risks, **including risks to confidentiality**.
- Identify plans to provide guidance and assistance in the event there are adverse effects to participants.
- Where appropriate, describe alternative treatments and procedures that may be beneficial to the participants. If you choose not to use these other beneficial treatments, provide the reasons for not using them.

2. Fair Selection of Participants

- Describe the target population(s) for the proposed project. Include age, gender, and racial/ethnic background and note if the population includes homeless youth, foster children, children of substance abusers, pregnant women, or other targeted groups.
- Explain the reasons for including groups of pregnant women, children, people with mental disabilities, people in institutions, prisoners, and individuals who are likely to be particularly vulnerable to HIV/AIDS.
- Explain the reasons for including or excluding participants.
- Explain how you will recruit and select participants. Identify who will select participants.

3. Absence of Coercion

- Explain if participation in the project is voluntary or required. Identify possible reasons why participation is required, for example, court orders requiring people to participate in a program.
- If you plan to compensate participants, state how participants will be awarded incentives (e.g., money, gifts, etc.). Provide justification that the use of incentives is appropriate, judicious, and conservative and that incentives do not provide an “undue inducement” which removes the voluntary nature of participation. Incentives should be the minimum amount necessary to meet the programmatic and performance assessment goals of the grant. Applicants should determine the minimum amount that is proven effective by consulting with existing local programs and reviewing the relevant literature. In no case

may the value if an incentive paid for with SAMHSA discretionary grant funds exceed \$20. For the purpose of this project, incentives may be provided for follow-up only.

- State how volunteer participants will be told that they may receive services intervention even if they do not participate in or complete the data collection component of the project.

4. Data Collection

- Identify from whom you will collect data (e.g., from participants themselves, family members, teachers, others). Describe the data collection procedures and specify the sources for obtaining data (e.g., school records, interviews, psychological assessments, questionnaires, observation, or other sources). Where data are to be collected through observational techniques, questionnaires, interviews, or other direct means, describe the data collection setting.
- Identify what type of specimens (e.g., urine, blood) will be used, if any. State if the material will be used just for evaluation or if other use(s) will be made. Also, if needed, describe how the material will be monitored to ensure the safety of participants.
- Provide in **Appendix 2, “Data Collection Instruments/Interview Protocols,”** copies of all available data collection instruments and interview protocols that you plan to use.

5. Privacy and Confidentiality

- Explain how you will ensure privacy and confidentiality. Include who will collect data and how it will be collected.
- Describe:
 - How you will use data collection instruments.
 - Where data will be stored.
 - Who will or will not have access to information.
 - How the identity of participants will be kept private, for example, through the use of a coding system on data records, limiting access to records, or storing identifiers separately from data.

NOTE: If applicable, grantees must agree to maintain the confidentiality of alcohol and drug abuse client records according to the provisions of **Title 42 of the Code of Federal Regulations, Part II.**

6. Adequate Consent Procedures

- List what information will be given to people who participate in the project. Include the type and purpose of their participation. Identify the data that will be collected, how the data will be used and how you will keep the data private.
- State:
 - Whether or not their participation is voluntary.
 - Their right to leave the project at any time without problems.
 - Possible risks from participation in the project.
 - Plans to protect clients from these risks.
- Explain how you will get consent for youth, the elderly, people with limited reading skills, and people who do not use English as their first language.

NOTE: If the project poses potential physical, medical, psychological, legal, social or other risks, you **must** obtain written informed consent.

- Indicate if you will obtain informed consent from participants or assent from minors along with consent from their parents or legal guardians. Describe how the consent will be documented. For example: Will you read the consent forms? Will you ask prospective participants questions to be sure they understand the forms? Will you give them copies of what they sign?
- Include, as appropriate, sample consent forms that provide for: (1) informed consent for participation in service intervention; (2) informed consent for participation in the data collection component of the project; and (3) informed consent for the exchange (releasing or requesting) of confidential information. The sample forms must be included in **Appendix 3, “Sample Consent Forms”**, of your application. If needed, give English translations.

NOTE: Never imply that the participant waives or appears to waive any legal rights, may not end involvement with the project, or releases your project or its agents from liability for negligence.

- Describe if separate consents will be obtained for different stages or parts of the project. For example, will they be needed for both participant protection in treatment intervention and for the collection and use of data?
- Additionally, if other consents (e.g., consents to release information to others or gather information from others) will be used in your project, provide a description of the consents. Will individuals who do not consent to having individually identifiable data collected for evaluation purposes be allowed to participate in the project?

7. Risk/Benefit Discussion

Discuss why the risks are reasonable compared to expected benefits and importance of the knowledge from the project.

Protection of Human Subjects Regulations

Applicants may also have to comply with the Protection of Human Subjects Regulations (45 CFR 46), depending on the evaluation and data collection procedures proposed and the population to be served.

Applicants must be aware that even if the Protection of Human Subjects Regulations do not apply to all projects funded, the specific performance assessment design proposed by the applicant may require compliance with these regulations. For assistance in determining if your proposed performance assessment meets the criteria in 45 CFR 46, Protection of Human Subjects Regulations, refer to the SAMHSA decision tree on the SAMHSA Web site, under “Applying for a New SAMHSA Grant,” <http://www.samhsa.gov/grants/apply.aspx>.

Applicants whose projects must comply with the Protection of Human Subjects Regulations must describe the process for obtaining Institutional Review Board (IRB) approval fully in their applications. While IRB approval is not required at the time of grant award, these applicants will be required, as a condition of award, to provide the documentation that an Assurance of Compliance is on file with the Office for Human Research Protections (OHRP) and that IRB approval has been received prior to enrolling any clients in the proposed project.

General information about Protection of Human Subjects Regulations can be obtained on the Web at <http://www.hhs.gov/ohrp>. You may also contact OHRP by e-mail (ohrp@osophs.dhhs.gov) or by phone (240/453-6900). SAMHSA-specific questions related to Protection of Human Subjects Regulations should be directed to the program contact listed in Section VII of this RFA.

Appendix G – Funding Restrictions

SAMHSA grant funds must be used for purposes supported by the program and may not be used to:

- Pay for any lease beyond the project period.
- Provide services to incarcerated populations (defined as those persons in jail, prison, detention facilities, or in custody where they are not free to move about in the community).
- Pay for the purchase or construction of any building or structure to house any part of the program. (Applicants may request up to \$75,000 for renovations and alterations of existing facilities, if necessary and appropriate to the project.)
- Provide residential or outpatient treatment services when the facility has not yet been acquired, sited, approved, and met all requirements for human habitation and services provision. (Expansion or enhancement of existing residential services is permissible.)
- Pay for housing other than residential mental health and/or substance abuse treatment.
- Provide inpatient treatment or hospital-based detoxification services. Residential services are not considered to be inpatient or hospital-based services.
- Make direct payments to individuals to induce them to enter prevention or treatment services. However, SAMHSA discretionary grant funds may be used for non-clinical support services (e.g., bus tokens, child care) designed to improve access to and retention in prevention and treatment programs.
- Make direct payments to individuals to encourage attendance and/or attainment of prevention or treatment goals. However, SAMHSA discretionary grant funds may be used for non-cash incentives of up to \$20 to encourage attendance and/or attainment of prevention or treatment goals when the incentives are built into the program design and when the incentives are the minimum amount that is deemed necessary to meet program goals. SAMHSA policy allows an individual participant to receive more than one incentive over the course of the program. However, non-cash incentives should be limited to the minimum number of times deemed necessary to achieve program outcomes. A grantee or treatment or prevention provider may also provide up to \$20 cash or equivalent (coupons, bus tokens, gifts, child care, and vouchers) to individuals as incentives to participate in required data collection follow up. This amount may be paid for participation in each required interview.
- Food is generally unallowable unless it's an integral part of a conference grant or program specific, e.g., children's program, residential.

- Implement syringe exchange programs, such as the purchase and distribution of syringes and/or needles.
- Pay for pharmacologies for HIV antiretroviral therapy, sexually transmitted diseases (STD)/sexually transmitted illnesses (STI), TB, and hepatitis B and C, or for psychotropic drugs.

SAMHSA will not accept a “research” indirect cost rate. The grantee must use the “other sponsored program rate” or the lowest rate available.

Appendix H – Sample Budget and Justification

ILLUSTRATION OF A SAMPLE DETAILED BUDGET AND NARRATIVE JUSTIFICATION TO ACCOMPANY SF 424A: SECTION B FOR 01 BUDGET PERIOD

OBJECT CLASS CATEGORIES

Personnel

Job Title	Name	Annual Salary	Level of Effort	SAMHSA Funded	Non-Federal Sources	TOTAL
Project Director	J. Doe	\$30,000	1.0	\$30,000	\$-0-	
Clinical Director	J. Doe			\$-0-	In-Kind	
Secretary	Unnamed	\$18,000	0.5	\$-0-	\$ 9,000	
Counselor	R. Down	\$25,000	1.0	\$25,000	\$-0-	
SUBTOTAL				\$55,000	\$9,000	
Enter Personnel subtotal on 424A, Section B, 6.a.						\$64,000
<u>Fringe Benefits</u> (24%)				\$15,360	\$-0-	
SUBTOTAL				\$15,360	\$-0-	
Enter Fringe Benefits subtotal on 424A, Section B, 6.b.						\$15,360
<u>Travel</u>						
2 trips for SAMHSA Meetings for 2 Attendees (Airfare @ \$600 x 4 = \$2,400) + (per diem @ \$120 x 4 x 6 days = \$2,880)						
				\$5,280	\$-0-	
Local Travel (500 miles x .24 per mile)				\$-0-	\$120	
[Note: Current Federal Government per diem rates are available at www.gsa.gov .]						
SUBTOTAL				\$5,280	\$120	
Enter Travel subtotal on 424A, Section B, 6.c.						\$ 5,400
<u>Equipment</u> (List Individually)						
"Equipment" means an article of nonexpendable, tangible personal property having a useful life of more than one year and an acquisition cost which equals the lesser of (a) the capitalization level established by the governmental unit or nongovernmental applicant for financial statement purposes, or (b) \$5000.						
SUBTOTAL				\$-0-	\$-0-	
Enter Equipment subtotal on 424A, Section B, 6.d.						\$-0-

ILLUSTRATION OF DETAILED BUDGET AND NARRATIVE JUSTIFICATION (cont'd.)

Supplies

Office Supplies	\$500	\$-0-
Computer Software – Microsoft Word	\$-0-	500

Enter Supplies subtotal on 424A, Section B, 6.e. \$1,000

CONTRACTUAL COSTS

Evaluation

Job Title	Name	Annual Salary	Level of Effort	SAMHSA Funded	Non-Federal Sources	TOTAL
Evaluator	J. Wilson	\$48,000	.05	\$24,000	\$-0-	
Other Staff		\$18,000	1.0	\$18,000	\$-0-	
Fringe Benefits (25%)				\$10,500	\$-0-	
Travel						
2 trips x 1 Evaluator (\$600 x 2)				\$ 1,200	\$-0-	
Per Diem @ \$120 x 6				720	\$-0-	
Supplies (General Office)				500	\$-0-	
Evaluation Contractual Direct Costs				\$54,920	\$-0-	
Evaluation Contractual Indirect Costs (19%)				\$10,435	\$-0-	
Evaluation Contract Subtotal				\$65,355		
SUBTOTAL				\$65,355	\$-0-	\$65,355

Training

Job Title	Name	Annual Salary	Level of Effort	SAMHSA Funded	Non-Federal Sources	TOTAL
Coordinator	M. Smith	\$ 12,000	0.5	\$12,000	\$-0-	
Admin. Asst.	N. Jones	9,000	0.5	9,000	\$-0-	
Fringe Benefits (25%)				5,250	\$-0-	
Travel						
2 Trips for Training						
Airfare @ \$600 x 2				\$1,200	\$-0-	
Per Diem \$120 x 2 x 2 days				480	\$-0-	
Local (500 miles x .24/mile)				120	\$-0-	
Supplies						
Office Supplies				\$500	\$-0-	
Software (Microsoft Word)				\$500	\$-0-	
Training Contractual Direct Costs Subtotal				\$40,025	\$-0-	\$40,025
Training Contractual Indirect Costs Subtotal				\$-0-	\$-0-	\$-0-

ILLUSTRATION OF DETAILED BUDGET AND NARRATIVE JUSTIFICATION (cont'd.)

SUBTOTAL	\$105,380	\$-0-	\$105,380
Enter Contractual subtotal on 424A, Section B, 6.f.			\$105,380
	SAMHSA Funded	Non-Federal Sources	TOTAL
<u>OTHER</u>			
Rent (500 Sq. Ft. x \$9.95)	\$ 4,975	\$-0-	
Telephone	\$ 500	\$-0-	
Maintenance (e.g., van)	\$-0-	\$ 2,500	
Audit	\$-0-	\$ 3,000	
Consultants = Expert @ \$250/day X 6 day (If expert is known, should list by name)	\$ 1,500	\$-0-	
SUBTOTAL	\$6,957	\$5,500	
Enter Other subtotal on 424A, Section B, 6.h.			\$12,475
<u>TOTAL DIRECT CHARGES</u> (sum of 6.a-6.h)			
Enter Total Direct on 424A, Section B, 6.i.			\$192,640
<u>INDIRECT CHARGES</u>			
15% of Salary and Wages (copy of negotiated Indirect Cost Rate Agreement attached) [\$64,000 X 15% = \$9,600]			
Enter Indirect Costs subtotal of 424A, Section B, 6.j.			\$9,600
Enter TOTALS on 424A, Section B, 6.k. (sum of 6i and 6j)			\$202,240

JUSTIFICATION

PERSONNEL - Describe the role and responsibilities of each position.

FRINGE BENEFITS - List all components of the fringe benefit rate.

EQUIPMENT - List equipment and describe the need and the purpose of the equipment in relation to the proposed project.

SUPPLIES - Generally self-explanatory; however, if not, describe need. Include explanation of how the cost has been estimated.

TRAVEL - Explain need for all travel other than that required by SAMHSA.

CONTRACTUAL COSTS - Explain the need for each contractual arrangement and how these components relate to the overall project.

OTHER - Generally self-explanatory. If consultants are included in this category, explain the need and how the consultant's rate has been determined. If rent is requested, provide the name of the owner of the building/facility. If anyone related to the project owns the building which is a less than arms length arrangement, provide cost of ownership/use allowance.

INDIRECT COST RATE - If your organization has no indirect cost rate, please indicate whether your organization plans to: a) waive indirect costs if an award is issued; or b) negotiate and establish an indirect cost rate with DHHS within 90 days of award issuance.

OTHER SOURCES – If other non-Federal sources of funding, including match or cost sharing as a total operating budget is included, provide the name of the source, e.g., in-kind, foundation, program income, Medicaid, State funds, applicant organization, etc., and explain its use.

CALCULATION OF FUTURE BUDGET PERIODS
(based on first 12-month budget period)

Review and verify the accuracy of future year budget estimates. Increases or decreases in the future years must be explained and justified. (NOTE: salary cap of \$186,600 is effective for all FY 2008 awards.)

	First 12-month Period	Second 12-month Period	Third 12-month Period
Personnel			
Project Director	30,000	30,000	30,000
Secretary*	9,000	18,000	18,000
Counselor	25,000	25,000	25,000
TOTAL PERSONNEL	64,000	73,000	73,000

*Increased from 50% to 100% effort in 02 through 03 budget periods.

Fringe Benefits (24%)	15,360	17,520	17,520
Travel	5,400	5,400	5,400
Equipment	-0-	-0-	-0-
Supplies**	1,000	520	520

**Increased amount in 01 year represents costs for software.

Contractual			
Evaluation***	65,355	67,969	70,688
Training	40,025	40,025	40,025

***Increased amounts in 02 and 03 years reflect the increase in client data collection.

Other	1,500	1,500	1,500
Total Direct Costs	192,640	205,934	208,653
Indirect Costs (15% S&W)	9,600	9,600	9,600
TOTAL COSTS	202,240	216,884	219,603

The Federal dollars requested for all object class categories for the first 12-month budget period are entered on Form 424A, Section B, Column (1), lines 6a-6i. The total Federal dollars requested for the second through the fifth 12-month budget periods are entered on Form 424A, Section E, Columns (b) – (e), line 20. The RFA will specify the maximum number of years of support that may be requested.

Appendix I- Core Components of SBIRT

Screening – brief screening incorporated into the normal routine in medical and other community settings that provides identification of individuals at risk for Substance Use Disorders.

SAMHSA is committed to: (1) standardizing the screening process so that screening produces consistent results across sites, in terms of individuals identified as requiring brief intervention or referral to treatment; and (2) detecting and intervening with individuals who have a Substance Use Disorder, but whose problems are not so severe that they require specialty treatment.

Therefore, grantees will be required to screen adults using the Alcohol, Smoking, and Substance Involvement Screening Test (ASSIST) tool for adults and the Car, Relax, Alone, Forget, Friends, Trouble (CRAFFT) tool for screening adolescents. Additional screening tools may be used with the agreement of the SAMHSA Project Officer.

Brief Intervention – face-to-face discussion that is focused on raising an individual’s awareness of his/her substance use and motivating the individual toward behavioral change. Brief interventions are considered to be 1 to 5 sessions in length.

Brief Treatment – a distinct level of face-to-face care that consists of a limited course of highly focused cognitive behavioral clinical sessions. Brief treatment is considered to be 6 to 12 sessions in length. It may occur in the same session as the initial screening or in follow-up sessions. It is conducted by licensed clinical practitioners.

Referral – a proactive process that facilitates access to care for individuals who are assessed to have a Substance Use Disorder requiring a more intensive treatment specialty.

Appendix J – Policy and Systems Change

A substantial body of research is related to barriers to access to health care, in general, as well as barriers to treatment for Substance Use Disorders, specifically. Various approaches exist to identify and classify barriers (Barry et al., 2004; Fiorentine, 1993; Institute of Medicine, 1990; PLNP, 2000; Schermer et al., 2003; Weisner and Schmidt, 1999). Less is known about those enabling factors that increase help-seeking and access (Fortney and Booth, 2001; Grant, 1997; Weisner and Schmidt, 2001).

Of major concern are the numerous studies documenting the failure of primary care physicians to identify persons at high risk of or already experiencing a Substance Use Disorder and initiating the appropriate referral for evaluation and treatment (Degutis, 2003; Hack and Adger, 2002; NCASA, 2002; Saitz et al., 1997). Such identification in mainstream medical care settings is necessary because perceived illness severity and stigma also may act as barriers to treatment.

Addiction policy and service provision in States/Tribes occurs within the context of both general health systems and financing arrangements and carved-out specialty prevention and treatment systems (Denmeade and Rouse, 1994; Jainchill, N., 2004). The implications of these arrangements for the diffusion of SBIRT must be considered in project formulation and implementation. Financial accessibility implies that the cost of the service is reasonable and there is no disincentive to use needed services because of their costs or the method of reimbursement. However, many studies identify barriers due to the manner in which substance abuse treatment is financed, such as a lack of parity with physical illness in commercial and public insurance leading to high co-pays and restrictions of payment for diagnostic assessments, and lack of coverage or payment for nontraditional specialist treatment modalities (e.g., residential therapeutic communities) or for screening services in emergency departments and primary care settings (Buck and Umland, 1997; Reader and Sullivan, 1992; Rivara et al., 2000; Rockett et al., 2005; Schmidt and Weisner, 2005; Sing et al., 1998). For example, several States still have Uniform Accident and Sickness Policy Provision Law (UPPL) provisions in their insurance policies that permit insurers to deny coverage for injuries sustained due to intoxication or because of the influence of other drugs. Although these policies are not always enforced, they may serve as disincentives to the provision of SBIRT-related services (Rivara, et al., 2000).

Another often cited barrier is the multiple, separate, and fragmented Federal, State, local, and private funding streams accessed by frequently uncoordinated agencies¹² that have different coverage policies, codes, and procedures for treatment modalities and ancillary services, different eligibility criteria for providers and patients, different reporting requirements, different placement criteria, and inconsistent benefit designs (Garnick et al, 2002; Gerstein and Schlessinger et al., 1991; Harwood, 1990; Horgan and Merrick, 2001; Hodgkin et al., 2000; Hodgkin et al., 2004; Johnson, 1999; Moss, 1998; Rockett et al., 2005; Schmidt and Weisner, 2005).

¹²The complexity of these multiple treatment subsystems at the State level has been described in a report submitted to SAMHSA/CSAT by the National Association of State and Drug Abuse Directors (NASADAD, 2002).

Appendix K - Resources for Implementing Screening, Brief Intervention, Referral, and Treatment

Background

For purposes of this cooperative agreement, CSAT will not require grantees to use a specific methodology for determining need, implementing systems change, or introducing SBIRT within their continuum of care. CSAT is not recommending a specific approach for developing collaboration among participating generalist and specialist providers. CSAT is requiring specific protocols for carrying out the individual activities involved (the screening instrument will be the ASSIST (Alcohol, Smoking and Substance Involvement Screening Test; World Health Organization (WHO) ASSIST Working Group, 2002, which provides the basis for brief intervention, brief treatment, and referral to specialty treatment). Other screening tools may be used with the agreement of the CSAT Project Officer. The applicant is required to describe and justify the strategies that will be implemented under the proposed cooperative agreement project and to describe the methods that will be used to assess need, eliminate barriers to access, and to carry out each of these activities. Wherever possible, the applicant should provide a description of any prior services or research projects on which their proposed approach is based.

In order to introduce some commonality in responses, CSAT will present an overview of terminology and anticipated issues and provide illustrative references that can serve as resources for proposal development and project implementation. The resources and references provided are not presented as an inclusive listing that must be used in proposal preparation.

Terminology

The financing of treatment for substance use problems has differed from the rest of health care financing in part because the public sector through block and categorical grants has been the major payer for services (e.g., Alemi and Sullivan, 2007; Beaston-Blaakman et al., 2007; Bray et al., 2007; Horgan and Merrick, 2001; Horgan et al., 2001). For example, Medicaid and other forms of health insurance require a clinical diagnosis and a determination of medical necessity for admission to treatment, while SAPT does not. The lack of common terminology has created problems in understanding who receives what services for treatment of Substance Use Disorders with what outcomes (Coffey et al., 2001).

Developing the policies and data for studying utilization and designing policies to increase access to clinically appropriate treatment requires use of common terms with clear definitions, starting with identifying the conditions for which treatment is needed. Diagnosis is the process of identifying and labeling specific diseases; diagnostic criteria for Substance Abuse and Dependence Disorders reflect the consensus of researchers and clinicians as to precisely which patterns of behavior or physiological characteristics constitute symptoms of these conditions. (Babor et al., 2005; Babor et al., 2001; National Institute on Alcohol Abuse and Alcoholism (NIAAA), 2003; National Institute on Drug Abuse (NIDA), 1997; 1999). Agreement on diagnosis in this field is relatively new, and the definitions and techniques for establishing diagnoses are evolving. Having a consistent set of diagnostic criteria allows clinicians to plan treatment and monitor treatment progress; enables policymakers, and planners to ensure the availability of needed treatment resources in each community; helps health care insurers and

other funders to decide whether treatment will be reimbursed; and allows patients access to medical insurance coverage.

As noted in the RFA, the need for treatment is discussed in terms of the categories used in the American Psychiatric Association Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV; 1994).¹³ DSM-IV includes a category called "Substance Related Disorders" that is divided into two major subcategories, Substance Induced Disorders and Substance Use Disorders. The focus of this program is on that part of the continuum of care that addresses treatment of Substance Use Disorders and not the treatment of Substance Induced Disorders, namely Substance Intoxication and Substance Withdrawal. Patients manifesting signs of intoxication, withdrawal symptoms, and other physical problems that require emergency care or urgent action would be managed in other components of the generalist or specialist treatment systems, stabilized and medically cleared before being screened for presence of a Substance Use Disorder (Veterans Health Administration and Department of Defense (VHA/DoD), 2001; Room et. al., 2005; Babor et al., 2005).

The DSM-IV identifies two Substance Use Disorders, Substance Abuse Disorder and Substance Dependence Disorder. Substance Use Disorders are further differentiated by type of drug primarily involved (e.g., amphetamine, alcohol, cocaine, marijuana/cannabis). DSM-IV is the diagnostic approach primarily used in this country for determining treatment eligibility, developing substance-specific treatments, and conducting epidemiological and clinical research.

Based on the DSM-IV, *Substance Abuse Disorder* is characterized by the presence of social or health-related problems related to the person's consistent pattern of substance use. *Substance Dependence Disorder* is characterized by a cluster of recognizable symptoms, including physical withdrawal, loss of control over use episodes, and continued use of substance despite knowledge of having a physical or psychological problem that is likely caused by substance.

The World Health Organization has also developed diagnostic criteria for the purpose of compiling statistics on all causes of death and illness, including those related to substance abuse or dependence. These criteria are published as the *International Classification of Diseases (ICD)*. In the current revision, ICD-10, substance dependence is defined in a way that is similar to the DSM-IV. The diagnosis focuses on an interrelated cluster of psychological symptoms, such as craving; physiological signs, such as tolerance and withdrawal; and behavioral indicators, such as the use of alcohol to relieve withdrawal discomfort. However, in a departure from the DSM-IV, rather than include the category "abuse," ICD-10 includes the concept of "harmful use." This category was created so that health problems related to alcohol and other drug use would not be underreported. Harmful use implies alcohol or drug use that causes either physical or mental damage in the absence of dependence (Babor, 2002; Babor et al, 2005). The ICD classification approach has served as the basis for much of the research underlying the use of brief interventions.

¹³ For a discussion of the methodology change, see Epstein, 2002, Substance Dependence, Abuse, and Treatment: Findings from the 2000 National Household Survey on Drug Abuse, Appendix C: Measurement of Dependence, Abuse, Treatment, and Treatment Need.

Review of the literature and discussions with practitioners and State Substance Abuse Authorities (SSAs) established that, while most of the research establishing the effectiveness of this approach has focused on alcohol use problems and disorders and has used the **problems** approach rather than the clinical approach, there is an emerging body of research and clinical experience that supports use of the SBIRT approach for persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence) who are experiencing problems related to the use of illicit drugs, particularly for marijuana use disorders (e.g., Babor, 2004; Babor et al. 2002; Baker et al. 2001; Barry, 1999; Bernstein et al., 1997; Blow, 1999; Conrod et al., 2000; Dennis et al., 2002a and b; Fleming, 2002; Gil et al., 2004; Gray et al., 2005; Grenard et al., 2007; Humeniuk, 2006; Kelso, 2002; Rohsenow et al., 2004; Samet et al., 1996; Stephens et al., 1994; Stephens, et al., 2000; Stephens, et al., 2004; Sullivan et al., 1997; WHO ASSIST Working Group, 2002; Zweben and Fleming, 1999).

While the effort to develop brief interventions for persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence) has not been as extensive as that for persons with alcohol problems, there have been several precedents. A number of States have already begun to introduce protocols for screening and brief intervention for both alcohol and drug use problems and disorders into their continuum of care (e.g., Harrison et al., 1996; Hartwell et al., 1996; Kroutil et al., 1997; New York State Office of Alcoholism and Substance Abuse Services (New York OASAS), 1996).¹⁴ Yet, in contrast to more traditional treatment services, early intervention services are often not specifically defined or regulated (IOM, 1990; Klitzner, et al., 1992). For purposes of this announcement, early intervention services (brief interventions) are those treatment procedures designed for persons who are exhibiting some problems associated with alcohol or other drug use but whose problems are not deemed serious enough to warrant treatment within a specialist setting. This would include those persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence). Early intervention services are sometimes identified as pre-treatment interventions (Blow, 1998) or clinical preventive services (U.S. Preventive Services Task Force (USPSTF), 1998) or indicated preventive interventions (Mrazek and Haggerty, 1994). The goal of early intervention is to prevent the problems from becoming more serious, and to promote total abstinence from alcohol and other illegal drugs. Early intervention could include an assessment of substance use and related problems, individual counseling provided by a health care practitioner, or participation in school-based or community-based educational or counseling programs designed to deter further substance use and promote healthier alternatives.

Rather than negating the public health approach to defining primary, secondary, and tertiary prevention as some have held, the IOM model can be seen as complementary, expanding the

¹⁴ A number of other States have included similar characterizations for differentiating intervention and treatment in their rules or planning efforts (e.g., Louisiana, Minnesota, Florida, North Carolina, Connecticut, Vermont, and Washington). For example, South Dakota has defined its approach as part of the regulations governing licensure of treatment facilities: "A facility that provides Early Intervention and Outpatient Services is a nonresidential facility that provides direct supportive client contact, indirect or collateral client contact, community information and liaison services. The program also provides formally planned counseling services to those persons harmfully affected by alcohol or drugs and who have been determined not to be in need of or accepting of structured outpatient or residential services." <http://legis.state.sd.us/rules/rules/6716A.htm#67:16:11:03.04>. Apparently some States (e.g., Florida) define intervention as both a treatment and non-treatment activity.

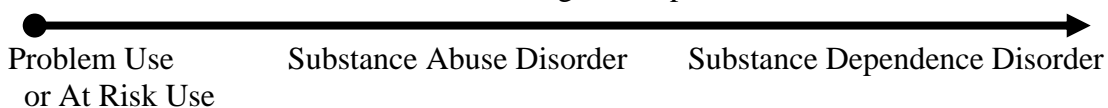
public health approach. The newer IOM model can be seen as actually further differentiating the public health construct of primary prevention into the categories of universal, selected, and indicative interventions, and the public health constructs of secondary and tertiary prevention into the categories of treatment and maintenance, respectively. The early intervention activities overlap the boundaries between primary prevention (indicated prevention) and secondary prevention (case finding).

In filling out the treatment portion of a State's continuum of care, the purpose of screening for substance use problems is to identify those persons who should receive either a brief intervention or referral for additional screening and assessment to establish whether more intensive treatment for a Substance Use Disorder (SUD) is needed. The persons screened *may* or *may not* meet the DSM-IV criteria for a Substance Abuse or Dependence Disorder (American Psychiatric Association, 1994). If they do not, but are deemed to be at risk users, then the same technology is employed as a clinical preventive service (or indicated preventive intervention). In practice, the activities are the same. However, the distinction is important for developing financing policies, for conducting epidemiological research and for tracking treatment access, appropriateness, utilization, and effectiveness.

Since diagnosis has not always been used as a criterion for admission to treatment in publicly funded treatment programs, States and service providers will need to introduce and agree upon a uniform approach to diagnosis as part of their implementation of this program and efforts to provide sustained funding for SBIRT, particularly through public and private health insurance mechanisms.

Integrating the Diagnostic and Problems Approaches

As noted, the DSM-IV term Substance Use Disorders can be used to refer to a range of substance-related problems that require treatment. A spectrum of Substance Use Disorders, from least to most serious, which encompasses the problems approach used in developing screening protocols for the use of brief interventions might be represented as follows:



In general, *problem or at-risk use* means use that exceeds an established threshold. The threshold may be defined in different ways depending on the source, the population, and other local conditions. The majority of work for developing such classifications in order to identify persons who could benefit from a brief intervention has been carried out for alcohol use problems and disorders. For example, the WHO manuals for introducing screening and brief intervention into primary care present general guidelines for assigning “risk levels” based upon AUDIT scores that conform to the spectrum above and lay out a spectrum of intervention and treatment responses.

Table 1: AUDIT Guidelines for Determining Intervention Strategy¹⁵

Risk Level	Intervention	Audit Score
I	Education	0-7
II	Simple Advice	8-15
III	Simple Advice plus Brief Counseling and Continued Monitoring	16-19
IV	Referral to Specialist for Diagnostic Evaluation and Treatment	20-40

The risk levels are used as a basis for making clinical judgments to tailor interventions to the particular conditions of individual patients, assuming that higher AUDIT scores are generally indicative of more severe levels of risk and problems or dependence. The guidelines are to serve as a starting point for an appropriate intervention. If a patient is not successful at the initial level of intervention, then the protocol calls for follow-up to develop a plan to step the patient up to the next level of intervention (Hornig and Chueh, 2004; Babor and Higgins-Biddle, 2001; Babor et al., 2001).

This approach is similar to that used for other screening tests, such as the Drug Abuse Screening Test (DAST) and the ASSIST (see Tables 2 and 3, respectively).

Table 2: DAST Guidelines for Determining Intervention Strategy¹⁶

<i>Score</i>	<i>Degree of Problems Related to Drug Abuse</i>	<i>Suggested Action</i>
0	No Problems Reported	None At This Time
1-2	Low Level	Monitor, Reassess At A Later Date
3-5	Moderate Level	Further Investigation
6-8	Substantial Level	Intensive Assessment

¹⁵ Based on Babor and Higgins-Biddle (2001) Brief Intervention For Hazardous And Harmful Drinking: A Manual for Use in Primary Care, Box 2, p.12

¹⁶ Based on Skinner HA (1982)

Table 3: ASSIST Guidelines for Determining Intervention Strategy¹⁷

<i>Alcohol</i>		<i>All Other Substances</i>	
Score	Risk Level/Intervention	Score	Risk Level/Intervention
0-10	Low Education	0-3	Low Education
11-19	Low-Moderate Brief Intervention	4-19	Low-Moderate Brief Intervention
20-26	Moderate-High Brief Intervention and Brief Treatment	20-26	Moderate-High Brief Intervention and Brief Treatment
27+	High Brief Intervention and Referral	27+	High Brief Intervention and Referral

These classification systems reflect the different patterns of drug use consumption and problems that call for differential societal responses that reflect differences in the drug (substance) used, the history, frequency, and amount used, as well as the existence and severity of associated physical, emotional, and social consequences of use. The Institute of Medicine committee that carried out a Congressionally mandated study of the evolution, effectiveness and financing of public and private drug treatment systems (Gerstein and Harwood, 1990) described a four level classification system reflecting these patterns that was a starting point in developing their initial estimates of the need for treatment, a model that was adapted for creating national estimates of the treatment gap. Table 4 depicts individual drug use patterns and interventions associated with each pattern of use. Each stage of use elicits a different type of societal response. The definitions for the categories are:

Use: Low or infrequent doses: experimental, occasional, “social.” Damaging consequences are rare or minor.

Abuse: Higher doses and/or frequencies: sporadically heavy, intensive. Effects are unpredictable, sometimes severe.

Dependence: High, frequent doses: compulsion, craving, withdrawal. Severe consequences are very likely.

¹⁷ Based on Henry-Edwards S, Humeniuk R, Ali R, Poznyak V, Monteiro (2003)

Table 4: Individual Drug Use Patterns and Intervention Strategies¹⁸

Stage	Category of Use	Use Pattern	Reason	Consequences	Societal Responses	
	Abstinence				Prevention Programs	
Early/light	Use	Low or infrequent doses	Experimental, occasional, “social”	Minor	Prevention Programs	Mild sanctions
Late/heavy	Abuse	Higher doses and/or frequencies	Sporadically heavy	Unpredictable, sometimes severe		
Late/heavy	Dependence	High, frequent doses	Compulsion, craving, withdrawal	Severe	Treatment programs	Severe sanctions

In the SBIRT approach, all persons are first screened and referred to the appropriate sector (community generalist, non-specialty or specialty) for intervention or treatment. Persons with a mild or moderate level of substance use problems would most often be offered a brief intervention in the non-specialty primary health care, criminal justice, educational, employment, or social service setting. Referral to intensive treatment in the specialty sector would be made only for those whose life situation is so unstable that prognosis is poor without specialty treatment or for those who fail to respond to an initial brief intervention--the stepped care approach (Sobell and Sobell, 1999, 2000).

Persons with substantial or severe problems would be referred from screening to specialty sequential assessment and treatment where problem and personal assessment would lead to assignment to more differentiated types of treatment modalities and levels of care, using a formal set of patient placement criteria.

Recent efforts have attempted to integrate the problem and diagnostic approaches, using both the research literature and clinical experience to refine the methods for screening, referring, and treating person’s based on these concepts (e.g., American Society of Addiction Medicine (ASAM), 2001; American Psychiatric Association, 1994; Beich et al., 2003; VHA/DoD, 2001). A possible model for this integration is presented in Table 5. The model also attempts to integrate the public health and IOM models for defining the continuum of care.

¹⁸ Based on Figure 3-1. A model of individual drug history, Gerstein and Harwood (1990:61)

Table 5: Integrating the Problem and Diagnostic Perspectives--A Possible Model

Problems	Risk Category or Diagnosis	Intervention Strategy	Exposures\ Sessions	Follow-up Suggested: Track: use, risk factors, and problems
No problems	No risk Or low risk	Universal prevention	Variable	Periodic re-screen: every year
Mild problems	At low risk	Clinical preventive service Selective prevention-brief advice	1-2	Periodic re-screen: every year
Moderate problems	At high risk	Clinical preventive service Indicated prevention Brief advice Brief intervention	1-2	Periodic re-screen every 6 months for 3 years, every year if no relapse
	Substance Abuse Disorder (DSM-IV, Axis I)	Brief advice Brief intervention Brief treatment	1-2 1-5 6-20	Periodic re-screen and booster session: every 3 months for 2 years, every 6 months for 2 years, every year if no relapse
Substantial problems	Substance Dependence Disorder (DSM-IV, Axis I)	Sequential assessment; match to clinically appropriate consumption and quality of life treatment strategies	21-60+	Periodic re-screen: every 3 months for 2 years, every 6 months for 2 years, every year if no relapse
Severe problems	Substance Dependence Disorder (DSM-IV, Axis I)	Sequential assessment; match to clinically appropriate consumption and quality of life treatment strategies	Variable; Based on individual response to treatment	Periodic re-screen: every 3 months for 2 years, every 6 months for 2 years, every year if no relapse

Using either the problems approach or the clinical approach, it is well recognized that within each community there is a spectrum of persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence). In keeping with recent summaries of the international research literature, it is estimated that the majority of adults (approximately 75%) are either abstainers or light or moderate users including some persons with Substance Use Disorders (who qualify for a diagnosis of abuse or dependence on alcohol or illicit drugs).

Persons in this group experience either no problems or mild or moderate substance use-related problems. There is a small but often highly visible minority of heavy, dependent users with major substance-related problems (estimated at approximately 5%). In between these extremes, there is a sizeable group of persons (20%) who may be drinking or using illicit drugs substantially or heavily and who have encountered substantial or severe problems related to their substance use. The concepts have been more difficult to address for illicit drugs, since any use could be seen as “abuse” because of potential legal consequences. As will be noted below, treatment is not necessarily the best societal response for persons at risk for, or diagnosed with, a Substance Use Disorder, (Substance Abuse or Dependence), but a brief intervention, early in their use career may well be.

These findings suggest that the continuum of care in each community must include a spectrum of primary, secondary, and tertiary prevention responses that parallels the spectrum of problems associated with use and that the diagnostic and problem approaches must be reconciled to ensure introduction of evidence based clinical protocols (NIDA, 1999). Research on effectiveness of specific approaches continues, but there is sufficient evidence available to lead to the policy conclusion that more widespread SBIRT efforts will decrease the medical and social costs of illicit drug use.

Using a method similar to that employed by Skinner and his colleagues in the original development of screening for establishing brief interventions as a valid technology, persons can be classified into four graded categories of drug and alcohol use problems, each of which should lead to a different treatment or intervention strategy being employed and to a different set of resource requirements (See Tables 3 and 5.):

Mild level of substance use problems. Use is light or moderate; symptoms are rated as mild or moderate; dependence is probably not present or, if present, is psychological rather than physical; life problems related to use are rated as absent, mild, or moderate.

Moderate level of substance use problems. Use is medium, substantial, or heavy; symptoms are rated as moderate; psychosocial problems related to use are likely and rated as moderate; psychological dependence may still be characteristic, but there are increasing signs of physical dependence, such as withdrawal symptoms; related life problems are rated as mild and/or moderate.

Substantial level of substance use problems. Use is substantial or heavy; symptoms are rated as substantial; physical dependence is likely; physical disorders, mental disorders, and psychosocial problems related to substance use are rated as moderate and/or substantial.

Severe level of substance use problems. Use is heavy; symptoms are rated as substantial and/or severe; physical dependence is highly pronounced; life problems are rated as substantial and/or severe; serious physical disorders and mental disorders related to use, such as liver disease, are likely.

As presented in Table 5, persons can also be classified as either persons with some level of risk for, or diagnosed with, a Substance Use Disorder, either Abuse or Dependence (those with no problems, mild or moderate problems), or persons with Substance Use Disorders (those who qualify for a diagnosis of drug abuse or dependence, either with moderate problems or with substantial and severe problems). The act of diagnosis shifts the nature of the services from prevention to treatment.

In developing strategies to increase access to clinically appropriate treatment, SAMHSA wants States to focus on the resources needed for improved screening, intervention, referral and treatment for Substance Use Disorders in order to increase the resources devoted to identifying and intervening with persons at risk for, or diagnosed with, a Substance Use Disorder (Substance Abuse or Dependence), as part of the generalist health care system. States should be able to provide for a similar linkage between whatever classification system the State is using and the DSM-IV categories in the protocol.

Resources for Implementing Screening

In health care, screening refers to a process designed to identify people who have, or who are at risk of having, an illness or disorder. The purpose of screening is to target persons for treatment, so as to reduce the long-term morbidity and mortality related to the condition. In addition, by intervening early and raising the individual's level of concern about the risk factors and substance-related problems, it is expected that screening for drug and alcohol problems in community settings can itself reduce subsequent use.

Two types of screening procedures are typically used. The first type includes self-report questionnaires and structured interviews; the second, clinical laboratory tests that can detect biochemical changes associated with excessive alcohol consumption (e.g., Peterson, 2004) or illicit drug use.

Commonly used and scientifically-validated instruments include the ASSIST (Newcombe et al., 2005; WHO ASSIST Working Group, 2002), AUDIT (Donovan et al., 2006; Reinert and Allen, 2007), DAST (McCabe et al., 2006; Yudko et al., 2007), and CRAFFT (Cummins et al., 2003; Knight et al., 2002; Knight et al., 2003). Several new instruments have been developed, but not yet rigorously tested to assess harmful use of either alcohol or drugs (e.g., the CAGE-D, the TCUDS, the GAIN-QS, the PDES). Several can be administered in 15-20 minutes or less. In some instances, two or three items have shown promise as screening tools (Brown and Rounds, 1995; Brown et al., 1997; 2001).

A bibliography containing descriptions and evaluations of various interview, questionnaire, and laboratory test screening approaches is available from Project Cork:

Project Cork. 2007. *CORK Bibliography: Screening Tests*. January 2005 to present, prepared : June 2006, 65 Citations.

www.projectcork.org/bibliographies/data/Bibliography_Screening_Tests.html.

Screening instruments have been developed or modified for use with different target populations, notably adolescents, offenders within the criminal justice system, and welfare recipients, women,

and the elderly. Several have been translated into other languages and have been evaluated for cultural sensitivity.

It is well recognized that screening instruments used with adolescents must be developmentally appropriate, valid and reliable, and practical for use in busy medical settings. One example of a brief substance abuse screening instrument recently developed specifically for use with adolescents is the CRAFFT test.

Cummins L, Huang C, Karen K, Burns KM, Blume AW, Larimer M, Marlatt GA. 2003. Validity of the CRAFFT in American-Indian and Alaska-Native adolescents: Screening for drug and alcohol risk. *Journal of Studies on Alcohol* ;64: 727-732.

Jull, A. 2003. "The CRAFFT Test Was Accurate for Screening for Substance Abuse Among Adolescent Clinic Patients." *Evid. Based. Nurs.* 6(1): 23.

Levy, S., Sherritt, L., Harris, S. K., Gates, E. C., Holder, D. W., Kulig, J. W., and Knight, J. R. 2004. "Test-Retest Reliability of Adolescents' Self-Report of Substance Use." *Alcoholism: Clinical & Experimental Research* 28(8): 1236-41.

Knight JR, Sherritt L, Shrier LA, Harris SK, Chang G. 2002. Validity of the CRAFFT substance abuse screening test among adolescent clinic patients. *Arch Pediatr Adolesc Med.* 156(6): 607-14.

Knight JR; Sherritt L; Shrier LA; Harris SK; Gates EC; Chang G. 2003. Validity of a brief alcohol screening test among adolescents: A comparison of the AUDIT, POSIT, CAGE, and the CRAFFT. *Alcoholism: Clinical and Experimental Research* 27: 67-73.

Additional screening tests and procedures targeted at adolescents, including the PDES and the GAIN-QS, are described in these publications:

Winters KC. 1992. Development of an adolescent alcohol and other drug abuse screening scale: Personal Experience Screening Questionnaire. *Addict Behav.* 17(5): 479-90.

Winters KC. 1999. **Screening and Assessing Adolescents For Substance Use Conditions.** Treatment Improvement Protocol (TIP) Series 31 DHHS Publication No. (SMA) 99-3282.

Winters KC. 1999. **Treatment of Adolescents With Substance Use Conditions.** Treatment Improvement Protocol (TIP) Series 32. DHHS Publication No. (SMA) 99-3283.

Winters KC. 2001. Assessing adolescent substance use problems and other areas of functioning: State of the art. In: PM Monti, SM. Colby, and TA. O'Leary (eds). **Adolescents, Alcohol, and Substance Abuse: Reaching Teens Through Brief Interventions.** New York, Guilford Publications, Inc., pp. 80-108.

Dennis ML 1998. **Global Appraisal of Individual Needs (GAIN) manual: Administration, Scoring and Interpretation**, (Prepared with funds from CSAT TI 11320). Bloomington IL: Lighthouse Publications.
www.chestnut.org/LI/GAIN/GAIN_QS/index.html

Martino S, Grilo CM, and Fehon DC 2000. Development of the drug abuse screening test for adolescents (DAST-A). *Addictive Behaviors* 25(1): 57-70.

Screening tests and procedures targeted at the elderly are described in these publications:

Beullens, J. and Aertgeerts, B. 2004. "Screening for Alcohol Abuse and Dependence in Older People Using DSM Criteria: a Review." *Aging Ment.Health* 8(1): 76-82.

Blow, F.C. Consensus Panel Chair. 1998. **Substance Abuse Among Older Adults**. Treatment Improvement Protocol (TIP) Series 26. DHHS Publication No. (SMA) 98-3179.

Blow FC and Barry KL. 1999-2000. Advances in alcohol screening and brief intervention with older adults. *Advances in Medical Psychotherapy*. 10:107-124

Screening tests and procedures targeted at persons in the criminal justice system are described in these publications:

Inciardi JA Consensus Panel Chair 1994. **Screening and Assessment for Alcohol and Other Drug Abuse Among Adults in the Criminal Justice System**. Treatment Improvement Protocol (TIP) Series 7. DHHS Publication No. (SMA) 94B2076

Peters, RH, Greenbaum, PE, Steinberg, ML, Carter, CR, Ortiz, MM, Fry, BC, Valle, SK. 2000. Effectiveness of screening instruments in detecting substance use conditions among prisoners. *Journal Substance Abuse Treatment*: 18(4): 349-58.

Simpson DD. 2001. Core set of TCU forms. Fort Worth: Texas Christian University, Institute of Behavioral Research. www.ibr.tcu.edu.

Efforts are also ongoing to develop methods for screening within the dual diagnosis population:

Maisto SA, Carey MP, Carey KB, Gordon CM, and Gleason JR. 2000. Use of the AUDIT and the DAST-10 to identify alcohol and drug use disorders among adults with a severe and persistent mental illness. *Psychological Assessment* 12(2): 186-192.

Rosenberg SD, Drake RE, Wolford GL, Mueser KT, Oxman TE, Vidaver RM, Carrieri KL, Luckoor R. 1998. Dartmouth Assessment of Lifestyle Instrument (DALI): a substance use condition screen for people with severe mental illness. *Am J Psychiatry* 155(2): 232-8.

Resources for Implementing Brief Interventions and Brief Treatments

There are now a variety of approaches that have been labeled as Brief Interventions (BI) and Brief Treatments (BT). Examples of approaches that address specific drugs are the Cannabis Youth Treatment protocol and the Adult Marijuana Treatment protocol, developed through CSAT funded testing of models originally developed through NIDA and NIAAA research.

Brief intervention and brief treatment strategies range from relatively unstructured advice-giving, to counseling and formalized feedback, to formal structured manuals for the number, duration, frequency, and content of sessions. Many of the protocols are based on behavioral self-control training, motivational interviewing, and cognitive-behavioral psychotherapy.

One of the most extensive efforts to attempt to conceptualize and differentiate Brief Interventions and Brief Treatments (and Long Term Treatments) is CSAT's TIP 34: **Brief Interventions and Brief Therapies for Substance Abuse**, published in 1999. The Consensus Panel for CSAT TIP #34 describes the two activities as follows:

Brief Intervention

Brief interventions are those practices that aim to investigate a potential problem and motivate an individual to begin to do something about his/her substance abuse, either by natural, client-directed means or by seeking additional substance abuse treatment.

Brief Treatment (Therapy)

Brief treatment (therapy) is a systematic, focused process that relies on assessment, client engagement, and rapid implementation of change strategies. Brief therapies usually consist of more (as well as longer) sessions than brief interventions. The duration of brief therapies is reported to be anywhere from 1 session (Bloom, 1997) to 40 sessions (Sifneos, 1987), with the typical therapy lasting between 6 and 20 sessions. Twenty sessions are usually the maximum because of limitations placed by many managed care organizations. Any therapy may be brief by accident or circumstance, but the focus of this TIP 34 is on *planned* brief therapy. The therapies described here may involve a set number of sessions or a set range (e.g., from 6 to 10 sessions), but they always work within a time limitation that is clear to both therapist and client.

In distinguishing between Brief Intervention and Brief Treatments, Zweben and Fleming (1999) characterize Brief Interventions as a low-cost, effective treatment alternative for alcohol and drug problems that use time-limited, self-help and preventive strategies to promote reductions in the case of nondependent clients, and in the case of dependent clients to facilitate their referral to specialized treatment programs. The primary goal in all cases is to increase motivation for behavior change. Brief interventions do not teach specific cognitive or behavioral skills, nor do they attempt to change a client's social environment.

Some researchers, practitioners, and policy analysts have suggested that the differentiation should be made on the basis of the number of sessions, with Brief Intervention typically lasting 1-3 sessions, not more than 5 sessions, and Brief Treatment typically consisting of 6 or more

sessions but not more than 20 sessions. Others have limited Brief Interventions to only 1 or 2 sessions and Brief Treatments to no more than 6 sessions.

Brief interventions and brief therapies may be thought of as elements on a continuum of care, but they can be distinguished from each other according to differences in outcome goals. Interventions are generally aimed at motivating a client to perform a particular action (e.g., to enter treatment, change a behavior, think differently about a situation), whereas therapies are used to address larger concerns (such as altering personality, maintaining abstinence, or addressing long-standing problems that exacerbate substance abuse).

A bibliography containing descriptions and evaluations of various brief intervention and brief treatment approaches is available from Project Cork.

Project Cork. 2007. *CORK Bibliography: Brief Treatment in Substance Abuse*. January 2005 to present, prepared : June 2006, 71 Citations.
www.projectcork.org/bibliographies/data/Bibliography_Brief_Treatment.html

Resources for Protocol Development

Treatment Improvement Protocols (TIPs) are best practice guidelines for the treatment of substance abuse. CSAT draws on the experience and knowledge of clinical, research, and administrative experts to produce the TIPs, which are distributed to a growing number of facilities and individuals across the country. Examples of protocols, screening instruments, and methods for carrying out activities required to implement the SBIRT program can also be found in several Treatment Improvement Protocols (TIPS) published by CSAT. TIPS can be accessed on the internet through the Treatment Improvement Exchange at:
www.treatment.org/Externals/tips.html

Barry KL Consensus Panel Chair. 1999. **Brief Interventions And Brief Therapies for Substance Abuse**. Treatment Improvement Protocol (TIP) Series 34. DHHS Publication No. (SMA) 99-3353.

Blow FC. Consensus Panel Chair. 1998. **Substance Abuse Among Older Adults**. Treatment Improvement Protocol (TIP) Series 26. DHHS Publication No. (SMA) 98-3179.

Miller WR. Consensus Panel Chair. 1999. **Enhancing Motivation for Change in Substance Abuse Treatment**. Treatment Improvement Protocol (TIP) Series 35. DHHS Publication No. (SMA) 99-3354.

Rostenberg PO. Consensus Panel Chair. 1995. **Alcohol and Other Drug Screening of Hospitalized Trauma Patients**. Treatment Improvement Protocol (TIP) Series 16. DHHS Publication No. (SMA) 95-3039.

Sacks S, Ries RK. Consensus Panel Co-Chairs. 2005. **A Guide to Substance Abuse Services for Primary Care Clinicians**. Treatment Improvement Protocol (TIP) Series 24. DHHS Publication No. (SMA) 05-3922.

Siegal H.A Consensus Panel Chair. 1998. **Comprehensive Case Management for Substance Abuse Treatment**. Treatment Improvement Protocol (TIP) Series 27. DHHS Publication No. (SMA) 98-3222.

Sullivan E., Fleming, M. Consensus Panel Co-Chairs. 1997. **A Guide to Substance Abuse Services for Primary Care Clinicians**. Treatment Improvement Protocol (TIP) Series 24. DHHS Publication No. (SMA) 97-3139.

Winters KC. Consensus Panel Chair. 1999. Treatment of Adolescents With Substance Use Conditions. Treatment Improvement Protocol. **(TIP) Series 32. DHHS Publication No. (SMA) 99-3283.**

An excellent example of a protocol that can guide implementation of a systematic approach to expanding the continuum of care is that developed by the VA/DoD Evidence-Based Clinical Practice Guideline Working Group, Veterans Health Administration, Department of Veterans Affairs, and Health Affairs, Department of Defense (2001). Electronic copies of the guideline are available from: Office of Quality and Performance web site: www.oqp.med.va.gov/cpg/SUD/SUD_Base.htm.

The VA/DoD guideline consists of five modules that address inter-related aspects of care for patients with Substance Use Disorders. Module A, Assessment and Management in Primary Care, provides a summary of the evidence base for the use of screening and brief interventions and outlines pathways for referral to specialty treatment.

Module A:	Assessment and Management in Primary Care includes screening, brief intervention, and specialty referral considerations.
Module C:	Care Management emphasizes chronic disease management for patients unwilling or unable to pursue rehabilitation goals.
Module P:	Addiction-Focused Pharmacotherapy addresses use of currently approved medications as part of treatment for alcohol and opioid dependence.
Module R:	Assessment and Management in Specialty Care focuses on patients in need of further assessment or motivational enhancement or who endorse rehabilitation goals.
Module S:	Stabilization addresses detoxification and pharmacological management of withdrawal symptoms.

The VA/DOD Guidelines and the TIPS are presented here as examples that may or may not fit a particular State's definition of its continuum of care. New York State has developed its own procedures, as may have other States:

New York State Office of Alcoholism and Substance Abuse Services (New York OASAS). 1996. **Changing Directions: Reference Manual for Early Intervention Services**. Albany NY: New York OASAS.

Brief Intervention Manuals

As noted in the RFA, CSAT has recently supported development and evaluation of manualized brief intervention and brief treatment strategies for adolescents and adults with marijuana use disorders that can be utilized.

Manuals in the Cannabis Youth Treatment (CYT) Series include:

Sample S., and Kadden R. 2002. **Motivational Enhancement Therapy and Cognitive Behavioral Therapy for Adolescent Cannabis Users: 5 Sessions**. Cannabis Youth Treatment (CYT) Series, Volume 1. <http://ncadi.samhsa.gov/govpubs/bkd384/>

Webb C, Scudder M, Kaminer Y, and Kadden R 2002. **The Motivational Enhancement Therapy and Cognitive Behavioral Therapy Supplement: 7 Sessions of Cognitive Behavioral Therapy for Adolescent Cannabis Users**. Cannabis Youth Treatment (CYT) Series, Volume 2. <http://ncadi.samhsa.gov/govpubs/bkd385>

Hamilton NL., Brantley LB, Tims FM, Angelovich N., and McDougall B. 2002. **Family Support Network for Adolescent Cannabis Users**. Cannabis Youth Treatment (CYT) Series, Volume 3. <http://ncadi.samhsa.gov/govpubs/bkd386/cyt3.pdf>

Godley SH., Meyers RJ, Smith JE, Karvinen T, Titus JC, Godley MD., Dent G, Passetti L, and Kelberg P. 2002. **The Adolescent Community Reinforcement Approach for Adolescent Cannabis Users**. Cannabis Youth Treatment (CYT) Series, Volume 4.

Little, HA. 2002. **Multidimensional Family Therapy for Adolescent Cannabis Users**, Cannabis Youth Treatment (CYT) Series, Volume 5.

These efforts build on prior research done under the auspices of the National Institute on Drug Abuse (NIDA), the National Institute on Alcoholism and Alcohol Abuse (NIAAA) and the World Health Organization (WHO), which have also issued several manuals that can also serve as resources in project development:

Babor TF and Higgins-Biddle JF. 2001. **Brief Intervention For Hazardous And Harmful Drinking: A Manual for Use in Primary Care**. Geneva: World Health Organization. WHO/MSD/MSB/01.6b.

Babor TF, Higgins-Biddle JC, Saunders JB, and Monteiro, MG. 2001. **AUDIT: The Alcohol Use Disorders Identification Test: Guidelines for Use in Primary Care. Second Edition**. Geneva: World Health Organization. WHO/MSD/MSB/01.6a.

Carroll KM 1998. **A Cognitive-Behavioral Approach: Treating Cocaine Addiction.** National Institute on Drug Abuse Therapy Manuals for Drug Addiction, Manual 1, NIH Publication 98-4308.

Miller WR, Zweben A, DiClemente CC, et al. 1992. **Motivational Enhancement Therapy Manual: A Clinical Research Guide for Therapists Treating Individuals with Alcohol Abuse and Dependence.** NIAAA Project MATCH Monograph Series Vol. 2. DHHS Publication No. (ADM) 92-1894.

National Institute on Alcohol Abuse and Alcoholism (NIAAA) **1995. The Physicians' Guide to Helping Patients With Alcohol Problems.** NIH Publication No. 95-3769.

National Institute on Alcohol Abuse and Alcoholism (NIAAA) 2003. **Helping Patients with Alcohol Problems: A Health Practitioner's Guide.** NIH Publication No. 03-3769. Bethesda, MD: U.S. Department of Health and Human Services, Public Health Service, National Institutes of Health.

National Institute on Drug Abuse (NIDA). 1999. **Principles of Drug Addiction Treatment: A Research Based Guide.** NIH Publication No. 00-4180 (reprinted 2000). Rockville MD: NIH/NIDA.

National Institute on Drug Abuse (NIDA). 2006. **Principles of Drug Abuse Treatment for Criminal Justice Populations.** NIH Publication No. 06-5316. Rockville MD: NIH/NIDA.

Roberts LJ and McCrady BS 2002. **Alcohol Problems in Intimate Relationships: Identification and Intervention - A Guide for Marriage and Family Therapists.** Rockville MD: National Institute on Alcohol Abuse and Alcoholism.

Resources for Analyzing Barriers and Implementing Systems Change

Additional resources for analyzing barriers to access and linkage between the generalist and specialist agencies and devising policy changes are provided by CSAT Technical Assistance Publications (TAPs). TAPs are publications, manuals, and guides developed by CSAT to offer practical responses to emerging issues and concerns in the substance abuse treatment field. Each TAP is developed by an expert who has had firsthand experience with the topic. TAPs can be accessed on the internet through the Treatment Improvement Exchange at:

www.treatment.org/Taps/

TAPs that may be useful resources include:

Crowe AH. and R Reeves. 1994. **Treatment for Alcohol and Other Drug Abuse: Opportunities for Coordination.** Technical Assistance Publication (TAP) Series 11. DHHS Publication No. (SMA) 94-2075.

Hansen C. 1995. **Forecasting the Cost of Chemical Dependency Treatment Under Managed Care: The Washington State Study.** Technical Assistance Publication (TAP) Series 15. DHHS Publication No. (SMA) 95-3045).

Moss S. 1998. **Contracting for Managed Substance Abuse and Mental Health Services: A Guide for Public Purchasers.** CSAT Technical Assistance Publication Series, Number 22. www.treatment.org/taps/tap22/TAP22TOC.htm

Samuels PN, Anita R. Marton AR. 1999. **Navigating the Pathways: Lessons and Promising Practices in Linking Alcohol and Drug Services with Child Welfare.** CSAT Technical Assistance Publication Series, Number 24. www.treatment.org/Taps/Tab24.pdf

Other publications that can be used to understand development of cost estimates, financing analyses, and systems change strategies are.

Alemi F and Sullivan T. 2007. "An Example of Activity Based Costing of Treatment Programs." *Am J Drug Alcohol Abuse* 33(1): 89-99.

Beaston-Blaakman A, Shepard D, Horgan C, Ritter G. 2007. "Organizational and Client Determinants of Cost in Outpatient Substance Abuse Treatment." *J Ment Health Policy Econ* 10(1): 3-13.

Bray JW, Zarkin GA, Davis KL, Mitra D, Higgins-Biddle JC, Babor TF. 2007. The effect of screening and brief intervention for risky drinking on health care utilization in managed care organizations. *Med Care.* 45(2): 177-82.

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The emphasis in this RFA is on expanding the State's continuum of care to include screening, brief intervention, referral, and treatment (SBIRT) in general medical and other community settings (e.g. community health centers, nursing homes, schools and student assistant programs, occupational health clinics, hospitals, and emergency departments). It is recognized that SBIRT activities are being, or could be, carried out in non-medical community settings (viz., student assistance programs, employee assistance programs, and welfare offices, drug courts, senior citizen centers).

While most of the research has been focused on screening in primary care medical settings, the approach can be effectively applied in many other contexts as well. In many cases, procedures have already been developed and used in these community settings for specific instruments, such as the AUDIT. To provide an example, Table 6 summarizes information about the settings, screening personnel, and target groups considered appropriate for a screening program using the screening instrument.

Table 6: Personnel, Settings and Groups Considered Appropriate for a Screening Program Using Screening Instruments¹⁹

Setting	Target Group	Screening Personnel
Primary care clinic	Medical patients	Nurse, social worker
Emergency room	Accident victims, Intoxicated patients, trauma victims	Physicians, nurses, or staff, health educators
Physician's office Surgery Prenatal and perinatal clinics	Medical patients	General practitioners, family physicians, physician extenders, nurses, or staff
General Hospital wards Outpatient clinic	Patients with hypertension, heart disease, gastrointestinal or neurological disorders	Internists, physician extenders, nurses, staff
Psychiatric hospital	Psychiatric patients, particularly those who are suicidal	Psychiatrists, psychologists, counselors, staff
Court, jail, prison	DWI offenders, violent criminals	Officers, counselors, probation officers
Other health-related facilities	Persons demonstrating impaired social or occupational functioning (e.g. marital discord, child neglect, etc.)	Health and human service workers
Military Services	Enlisted men and officers	Medics
Welfare Offices	Applicants and clients	Social Workers, case aides
Workplace Employee Assistance Program	Workers, especially those having problems with productivity, absenteeism or accidents	Employee assistance staff

A State that includes such efforts in their proposal must recognize that these efforts must comport to the diagnostic considerations outlined here. Examples of such activities can be found in these and other publications:

Peters RH and Wexler HK, Consensus Panel Co-Chairs 2005. **“Substance Abuse Treatment for Adults in the Criminal Justice System.” Treatment Improvement Protocol (TIP) series number 44.** DHHS Publication No. (SMA) 05-4056

White WL and Dennis M. 2002. **The cannabis youth treatment experiment: Key lessons for student assistance programs.** *Student Assistance Journal*, 14: 16-19.

¹⁹ Modified from Box 1, Personnel, Settings and Groups Considered Appropriate for a Screening Programme Using the AUDIT (Babor et al., 2001)

Young, N. K. 1996. **Alcohol and Other Drug Treatment: Policy Choices in Welfare Reform**. Washington DC: National Association of State Alcohol and Drug Abuse Directors.

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Young NK and Gardner SL. 2002. **Navigating the Pathways: Lessons and Promising Practices in Linking Alcohol and Drug Services with Child Welfare**. . Technical Assistance Publication (TAP) Series 27. SAMHSA Publication No. (SMA) 02-3639.

Resources for Developing Need Estimates

Resources that can be referred to for developing estimates of need for treatment and resource availability are:

DeWit DJ and Rush B 1996. Assessing the Need for Substance Abuse Services: A Critical Review of Needs Assessment Models. *Evaluation and Program Planning*. 19(1): 41-64.

Epstein JF 2002. **Substance Dependence, Abuse, and Treatment: Findings from the 2000 National Household Survey on Drug Abuse**. (DHHS Publication No. SMA 02-3642, NHSDA Series A-16). Rockville MD: Substance Abuse and Mental Health Services Administration, Office of Applied Studies

Gerstein DR and Green LW (eds). 1993. **Preventing Drug Abuse: What Do We Know?** Washington DC: National Academy Press. (Chapter 2 and Appendix)

Gerstein DR and Harwood HJ (eds). 1990. **Treating Drug Problems**, Vol. I. Washington DC: National Academy Press. (Chapter 3)

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Maxwell JC (ed). 2001. **Multiple Indicator Analysis: Using Secondary Data to Analyze Illicit Drug Use**. (DHHS Publication No. SMA 01-3539. Rockville, MD: Center for Substance Abuse Treatment and Center for Mental Health Services). Substance Abuse and Mental Health Services Administration.

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Office of Applied Studies. 2002. **National and State Estimates of the Drug Abuse Treatment Gap: 2000 National Household Survey on Drug Abuse (NHSDA Series H-**

14, DHHS Publication No. SMA 02-3640). Rockville, MD: Substance Abuse and Mental Health Services Administration. www.samhsa.gov/oas/TXgap/toc.htm

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Collaboration with Addiction Technology Training Centers as a Training Resource

SAMHSA/CSAT funds a network of 14 independent regional Addiction Technology Transfer Centers (ATTCs) and a National Office (www.nattc.org). The ATTCs constitute a nationwide, multi-disciplinary resource that draws upon the knowledge, experience and latest work of recognized experts in the field of addictions. A list of ATTCs, the States covered, and contact information is provided in Table 7. Each ATTC serves as a resource to 2 or more States, having memoranda of understanding with the State Substance Abuse Authorities (SSAs). For additional information related to ATTC's, please visit www.nattc.org.

Table 7: Addiction Technology Transfer Center Contacts

<p>Maine, New Hampshire, Vermont, Massachusetts, Connecticut, Rhode Island ATTC of New England Center for Alcohol and Addiction Studies Brown University Providence, Rhode Island 02912 (401) 863-6486 www.attc-ne.org Director: Susan Storti, PhD, RN</p>	<p>Georgia, South Carolina, North Carolina Southeast ATTC Morehouse School of Medicine CORK Institute Atlanta, Georgia 30310 (404) 752-1016 www.sattc.org Director: Kay Gresham Morrison, LCSW, ACSW</p>
<p>New York, Pennsylvania Northeast ATTC Institute for Research, Education and Training in Addictions Pittsburgh, Pennsylvania 15219 (866) 246-5344 www.ireta.org/attc Director: Michael Flaherty, PhD</p>	<p>Virginia, Kentucky, Tennessee, West Virginia Mid-Atlantic ATTC Virginia Commonwealth University Richmond, Virginia 23298-0469 (804) 828-9910 www.mid-attc.org Director: Paula Horvatich, PhD</p>
<p>District of Columbia, Delaware, Maryland, New Jersey Central East ATTC DANYA Institute Silver Spring, Maryland 20910 (240) 645-1145 www.ceattc.org Director: Thomas Durham, PhD</p>	<p>Illinois, Ohio, Indiana, Michigan Great Lakes ATTC Jane Addams College of Social Work University of Illinois-Chicago Chicago, Illinois 60612 (312) 996-1373 www.glattc.org Director: Lonnetta Albright</p>

**Iowa, North Dakota, South Dakota,
Minnesota, Wisconsin**

Prairielands ATTC
University of Iowa
Iowa City, Iowa 52242
(319) 335-5368

www.pattc.org

Director: Anne Helene Skinstad, PhD

**Nebraska, Missouri, Kansas, Oklahoma
Arkansas**

Mid-America ATTC
University of Missouri-Kansas City
Kansas City, Missouri 64110
(816) 482-1100

www.mattc.org

Director: Pat Stilen, LCSW, CADAC

**Nevada, Montana, Wyoming, Utah,
Colorado, Idaho**

Mountain West ATTC
University of Nevada, Reno
Reno, Nevada 89557
(775) 784-6265

www.mwattc.org

Principal Investigator: Nancy Roget, MS
Co-PI: Gary L. Fisher, PhD

**Alaska, Washington, Oregon, Hawaii,
Pacific Islands**

Northwest Frontier ATTC
Salem, Oregon 97303
(503) 373-1322

www.nfattc.org

Director: Steve Gallon, PhD

Texas, Louisiana, New Mexico

Gulf Coast ATTC
University of Texas
Center for Social Work Research
Austin, Texas 78703
(512) 232-0608

www.utattc.net

Director: Richard Spence, PhD
Director: Philip Orrick, BS

California, Arizona

Pacific Southwest ATTC
UCLA Integrated Substance Abuse
Programs
Los Angeles California 90025
(310) 267-5408

www.psattc.org/

Director: Thomas Freese, PhD
Co-Director: Michael Shafer, PhD

Puerto Rico, US Virgin Islands

Caribbean Basin and Hispanic ATTC
Centro de Estudios en Adiccion
Universidad Central del Caribe
Call Box 60-327

Bayamon, Puerto Rico 00960-6032
(787) 785-4211

web <http://cbattc.uccaribe.edu/>

Director: Rafaela Robles, EdD

Alabama, Florida, Mississippi

Southern Coast ATTC
Florida Certification Board
Tallahassee Florida 32301
(850) 222-6731

www.scattc.org

Director: Pam Waters, MEd

National Office

University of Missouri - Kansas City
Kansas City, MO 64110-2499
(816) 235-6888

www.nattc.org/

Director: Mary Beth Johnson, MSW

Appendix L- Reporting Requirements for SBIRT

The following are the reporting requirements for each category of services or combination of services to be provided to individuals:

Screening Only

For clients who are screened and who, based on the results of the screen, should not require any level of substance abuse intervention or treatment services, the following will be required for each grantee and/or each community, if applicable:

Baseline Client Level Data

Baseline (at screening) CSAT-GPRA data elements limited to the demographics must be collected on all clients in this category. (See Sections A and B of the GPRA tool.) This individual client level data will be used to count unduplicated clients served. No further data collection will be required on these clients.

Screening and Brief Intervention (BI)

For all clients who are screened and who, based on the results of the screen, should or do receive brief intervention, the following must be collected and reported:

Baseline Client Level Data

Baseline (at screening) CSAT-GPRA data elements limited to the demographic, and substance use domains must be collected on all clients in this category of service. (See Sections A and B of the GPRA tool.) This individual client level data will be used to count unduplicated clients served. It is important that all clients complete a tracking information sheet in the event they are selected for follow-up.

Discharge Client Level Data

For all clients in this category, discharge data must be submitted to CSAT. If a Brief Intervention is completed more than 7 days from the time of intake, Sections A, B, J and K of the GPRA tool must be completed on the client. If the intervention is 7 days or less from the time of intake, Sections A, J and K should be completed.

Follow-up Client Level Data

For a representative 10 percent (10%) sample of clients in this category who should have or did receive brief intervention, the follow-up GPRA items asked are limited to the substance use domain and follow-up sections of the tool. (See Sections A, B and I of the GPRA tool.) Data must be collected at 6 months after baseline and entered into the CSAT web-based GPRA data entry and reporting system. CSAT will provide grantees the sampling method to obtain the representative sample of 10 percent (10%). Grantees will be notified which clients have been selected as part of the representative sample and need to be located for follow-up via a Web-based notification report. Grantees are expected to achieve a follow-up rate of at least 80 percent (80%) of those selected for the follow-up sample.

For example, if 100 patients are screened and should receive Brief Intervention, 10 clients will be in the CSAT selected sample to be followed up. Grantees will be required to attempt to locate all 10 clients. It is required that at a minimum eight of these clients complete a follow-up interview.

Aggregated Data

In the semi-annual report, the grantee must also provide data about the costs for the delivery of screening and brief intervention, including the mean, median, and range of costs overall, by facility type, and region and sub-recipient, if applicable. The grantee must also discuss how such costs compare to the CSAT approved cost parameters for screening and brief intervention and what efforts they are undertaking to bring costs into line with those expected.

Screening and Brief Treatment (BT) or Screening and Referral to Other Types of Treatment for Substance Use Disorders (RT)

For all clients that are screened and require either brief treatment or other treatment, the following must be collected and reported:

Baseline Client Level Data

Baseline (at screening) using all of the CSAT GPRA data elements must be collected on all clients in this category of service. (See Sections A through G of the GPRA tool.) It is important that all clients complete a tracking information sheet in the event they are selected for follow-up.

Discharge Client Level Data

For all clients in this category, discharge data must be submitted to CSAT. If a Brief Treatment is completed more than 7 days from the time of intake, Sections A through G, J and K of the GPRA tool must be completed on the client. If the treatment is 7 days or less from the time of intake, Sections A, J and K of the GPRA tool must be completed.

Follow-up Client Level Data

For a representative 10 percent (10%) sample of clients in this category who, based on the results of their screening, should have or did receive services beyond brief intervention, follow-up data (all domains, see Sections A through I of the GPRA tool) are to be collected at 6 months after the initiation of substance abuse treatment services and entered into the GPRA web-based data entry and reporting system. CSAT will provide grantees the sampling method to obtain the representative sample of 10 percent (10%). Grantees will be notified which clients have been selected as part of the representative sample and need to be located for follow-up via a web based notification report. Grantees are expected to achieve a follow-up rate of at least 80 percent (80%) of those selected.

Note that two levels of intervention (BT, RT) are being described here. A 10 percent (10%) sample is required for each of the two levels (BT, RT).

For example, if 100 patients are screened and should receive Brief Treatment, 10 clients will be in the sample to be followed up. Grantees will be required to attempt to locate all 10 clients. It is required that at a minimum eight of these clients complete a follow-up interview.

In addition, if 100 patients were screened and should receive a Referral to Treatment, 10 clients will be in the sample to be followed up. Grantees will be required to attempt to locate all 10 clients. It is required that at a minimum 8 of these clients complete a follow-up interview.

Aggregated Data

In the semi-annual report, the grantee must also provide data about the costs for the delivery of screening and brief treatment as well as all other treatment modalities supported by this cooperative agreement including the mean, median, and range of costs overall, by modality, facility type and region, and sub-recipient, if applicable. The grantee must also discuss how such costs compare to the CSAT approved cost parameters for screening and brief intervention and what efforts they are undertaking to bring costs into line with those expected.