

return it, using the instructions provided on the survey form.

### C. Reporting Requirements

#### 1. Progress and Financial Reports

- Grantees must provide annual and final progress reports. The final progress report must summarize information from the annual reports, describe the accomplishments of the project, and describe next steps for implementing plans developed during the grant period.

- Grantees must provide annual and final financial status reports. These reports may be included as separate sections of annual and final progress reports or can be separate documents. Because SAMHSA is extremely interested in ensuring that its best practices efforts can be sustained, your financial reports must explain plans to ensure the sustainability (see Glossary) of efforts initiated under this grant. Initial plans for sustainability should be described in year 1 of the grant. In each subsequent year, you should describe the status of the project, successes achieved and obstacles encountered in that year.

- SAMHSA will provide guidelines and requirements for these reports to grantees 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 the grantee's progress toward meeting its goals.

#### 2. Government Performance and Results Act

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. These requirements will be specified in the NOFA for each funding opportunity.

#### 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 (301-443-8596) of any materials based on the SAMHSA-funded 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 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

The NOFAs provide contact information for questions about program issues.

For questions on grants management issues, contact: Stephen Hudak, Office of Program Services, Division of Grants Management, Substance Abuse and Mental Health Services Administration, 5600 Fishers Lane, Rockwall II 6th Floor, Rockville, MD 20857, (301) 443-9666, [shudak@samhsa.gov](mailto:shudak@samhsa.gov).

### VIII. Other Information

#### A. SAMHSA Confidentiality and Participant Protection Requirements and Protection of Human Subjects Regulations

You must describe your procedures relating to Confidentiality, Participant Protection and the Protection of Human Subjects Regulations in Section I of your application, using the guidelines provided below. Problems with confidentiality, participant protection, and protection of human subjects identified during peer review of your application may result in the delay of funding.

#### Confidentiality and Participant Protection

All applicants must address each of the following elements relating to confidentiality and participant protection. You must describe how you will address these requirements.

##### 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 target 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.).

- 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.