



# REQUEST FOR DETERMINATION OF RESEARCH STATUS

To be completed by the staff member with lead responsibility for the project and approved by branch chief (if applicable) and Division ADS. A separate PGO funding memo is required if project is research and involves human subjects regardless of the CDC staff role.

- Instructions:**
- (1) Use this form to declare: (a) the research status of any project, (b) role or roles of CDC staff
  - (2) A short summary should be attached offering specific details about the project and the role of staff.
  - (3) Be sure to complete all applicable items, obtain appropriate signatures and submit this form for approval.

**Tracking Number:** To Be Determined

(Use PGO number if cooperative agreement, grant, etc.)

**Date submitted:** 08/13/2018

**Title of Project:** DP18-1811 Partner Actions to Improve Oral Health Outcomes

**Dates for project period:**

**Dates for funding (if applicable):**

**Beginning:** 09/01/2018

**Beginning:** 09/01/2018

**Ending:** 08/31/2023

**Ending:** 08/31/2019

**Project is (choose one):**

*NOTE: Revision, as used below, refers to any substantive change made to the project including scope of project, funding restrictions, personnel, role of CDC staff member, determination of research status, etc.*

**New**

**Revision**

**Continuation, without revision(s)**

**Continuation, with revision(s)**

**Lead staff member:**

**Contact information:**

**Please indicate your role(s) in this project:**

**Name:** Marcia Parker

**Division:** DOH

**Project officer**  **Technical monitor**

**User ID:** KUV7

**Telephone:** 770-488-6075

**Principal investigator**  **Investigator**

**Scientific Ethics number:** \_\_\_\_\_

**Mailstop:** F-80

**Consultant**  **Other (please explain)**

**1. Are any or all of the activities within this project DESIGNED to contribute to generalizable knowledge (i.e., research)?**

**YES**  **NO**

**If YES, list those activities which are research:**

**2. Is this CDC project research or public health practice (check all that apply)?**

**Research**

**Public health practice**

*Check one:*

*Check all that apply:*

**Human subjects involved**

**Emergency Response**  **Surveillance**

**Human subjects not involved**

**Program evaluation**  **Other (please explain)**

**3. If RESEARCH involving human subjects, has the project or research activities been reviewed by the CDC IRB for human subjects protection?**

a.  **NO, New project, not yet reviewed**

d.  **YES, Reviewed and approved by CDC**

b.  **NO, Existing project, not ready to submit**

**If YES, please list protocol number and**

c.  **NO, Submitted for approval**

**expiration date** \_\_\_\_\_

e.  **NO, RESEARCH, no CDC investigators (CDC IRB not required)**

f.  **N/A (Not Applicable)**

**If RESEARCH, list any other CDC staff involved in this project, please include the name, role, and scientific ethics number**

## Tracking NO. To Be Determined

Name	Role (project officer, investigator, consultant, etc.)	Scientific ethics number Prin
Marcia Parker		

**IF YOU THINK THE RESEARCH PROJECT MIGHT QUALIFY AS EXEMPT RESEARCH (as identified in 45CFR46.101), PLEASE ANSWER questions 4-6, OTHERWISE SKIP TO question 7.**

4. Does the proposed research involve prisoners?  
 YES If YES, this research cannot be exempted and must be reviewed by an IRB (skip to question 7).  
 NO
5. Does the proposed research involve fetuses, pregnant women, or human in vitro fertilization as targets (such that Subpart B would apply)?  
 YES If YES, this research cannot be exempted and must be reviewed by an IRB (skip to question 7).  
 NO

### Educational Research

- 6.1 Is this research conducted in established or commonly accepted educational settings, AND does the research involve normal educational practices (e.g., research on regular and special education strategies or research on the effectiveness of, or comparison among instructional techniques, curricula or classroom management methods)?  
 YES  NO

### Research Involving Surveys, Interview Procedures (including Focus groups), Observation of Public Behavior, or Educational Tests

- 6.2 Will this research use educational tests (cognitive, diagnostic, aptitude, achievement), survey procedures, interview procedures or observation of public behavior?  
 YES  NO If NO skip 6.3

Will children (<18 years of age) be research subjects?

- YES If YES, this research cannot be exempted and must be reviewed by an IRB (skip to item 7)  
 NO

- 6.2.1 Is the information obtained recorded in such a manner that human subjects can be identified directly or indirectly through identifiers (such as a code) linked to the subjects;  
 YES  NO

- 6.2.2 Will any disclosure of the human subjects' responses outside of the research setting have the potential to place the subjects at risk of criminal or civil liability, or be damaging to the subjects' financial standing, employability or reputation? (Examples here may include: the collection of sensitive data regarding the subjects' (or relatives' or associates') possible substance abuse, sexuality, criminal history or intent, medical or psychological condition, financial status, or similarly compromising information).  
 YES  NO

- 6.3 Will this research use educational tests (cognitive, diagnostic, aptitude, achievement), survey procedures, interview procedures, or observation of public behavior but the research is not exempt under paragraph 6.2 of this section:

YES  NO If NO skip to 6.4

- 6.3.1 Will this research involve human subjects that are elected or appointed public officials or candidates for public office?  
 YES  NO

- 6.3.2 Does federal statute(s) require(s) without exception that confidentiality of the personally identifiable information will be maintained throughout the research and thereafter? (Note: CDC can use this exemption criterion only in the case where a 308(d) Assurance of Confidentiality has been obtained to cover the research).  
 YES  NO

### Existing Data Which Is Publicly Available or Unidentifiable

- 6.4 Does this research involve only the collection or study of existing\* data, documents, records, pathological or diagnostic specimens? (\* 'existing' means existing before the study begins)?

YES  NO If NO skip to 7

- 6.4.1 Is this material or information publicly available?  
 YES  NO

**Tracking NO. To Be Determined**

**6.4.2 Is this material or information recorded in such a manner by the investigator that the subjects cannot be identified directly or indirectly through identifiers linked to the subjects?**

(Note: If a link is created by an investigator even temporarily, for research purposes, this criterion is not met. If a temporary link is created by clinical staff who already have access to the data, this criterion is met).

- YES (there are no identifying information and no unique identifiers or codes)YES
- NO (there are identifiers (including codes))

**7. Please prepare and attach a short summary paragraph (<1 page); if this is new:**

- a. **Be sure to include the purpose of the project, specific details about the project and the role of the CDC staff member (s) in the project. In explaining one's role as a consultant be particularly careful to identify involvement in things like: study design decisions, oversight of protocol development, participation in review of data collection procedures, and participation in data analysis and/or manuscript preparation, as well as whether there will be access to identifiable or personal data.**
- b. **Explain your project status selection (research--non-exempt, exempt, no CDC investigator or not involving human subjects; public health practice). If you selected research not involving human subjects be sure to indicate if the data includes any personal information (e.g., name, SSN), linkable study identification numbers or codes, or geographical information.**

The DP18-1811 Partner Actions to Improve Oral Health Outcomes program is a five-year cooperative agreement to continue CDC investment in and support of oral health promotion and disease prevention programs. This program does not include research. The purpose of this Notice of Funding Opportunity (NOFO) is to build the strength and effectiveness of state and territorial oral health programs to prevent and control oral diseases and related conditions. Partner Actions to Improve Oral Health Outcomes will have two components. Applicants may choose to apply for component 1 only, component 2 only, or both. Under component 1, the recipient will implement priority strategies such as providing technical assistance and capacity building resources for states, conducting state oral health program assessments, and conducting assessments and providing technical assistance to territorial oral health programs. Note: Funds for the assessment and technical assistance strategy for territorial oral health programs are available ONLY for year 1 of the period of performance. Funds for State oral health infrastructure and capacity building are available for each year of the period of performance. Under component 2, the recipient will work with five programs selected under NOFO DP18-1810 to implement medical-dental integration activities that integrate oral health with other chronic disease programs. The recipient will provide technical assistance for medical-dental integration programs, and collect and information supporting the effectiveness of medical-dental integration programs and strategies. The proposed program will replace and build upon NOFO DP13-1313 [FY2013-FY2018]. Successful implementation and execution of the NOFO strategies will result in decreases in dental caries, oral health disparities, and co-morbid chronic diseases.

CDC's role is to provide programmatic, evaluation, epidemiologic, and technical assistance for recipients and partner organizations through programmatic and one-on-one technical assistance and consultation, national training, workshops, web conferences, and other forms of guidance. CDC will also facilitate technical assistance between recipients and other CDC partners as needed.

**8. Please list the primary project site and all collaborating site(s).**

**Explanation of project components:**

**9. If project involves research that is funded extramurally, list amount of award that should be restricted pending IRB approval and describe which project components will be affected, if known:**

Approvals (signature and position title)	Date	Research Determination / Remarks
Casey Hannan - Supervisory Health Scientist       staff member completing this form	08/13/2018	<input checked="" type="checkbox"/> Public health practice <input type="checkbox"/> Research not involving human subjects <input type="checkbox"/> Research involving human subjects, no CDC investigators <input type="checkbox"/> Research involving human subjects, CDC investigators, exempt <input type="checkbox"/> Research involving human subjects, CDC investigators, not exempt  (check if applicable) <input type="checkbox"/> Local IRB <input type="checkbox"/> CDC Exemption <input type="checkbox"/> CDC IRB  <u>Comments:</u> Approved.

