



**Tracking NO. No Funding**

- 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

Name	Role (project officer, investigator, consultant, etc.)	Scientific ethics number Prin
Rachel Kaufmann		7214

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

**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.**

Goal of the project: To provide Cooperative Agreement applicants and awardees with templates for the creation of Data Management Plans (DMP).  
 Intended use of the resulting data: DMPs are required of entities using federal funds to collect or generate data. DMPs will be submitted to CDC by grant and cooperative agreement awardees for assessment to verify that they are concordant with CDC's data sharing policy.  
 Methods to be used to collect: DMPs will be submitted as standalone sections of the NOFO and annual continuation applications; revisions can also be submitted by the awardees whenever needed.  
 How data will be analyzed: Once DMPs are received, the DMP is then assessed for completeness and adherence to CDC policy and guidelines.

**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
Rachel Kaufmann - Assoc Director Science       staff member completing this form	06/12/2019	<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> please approve

<p>Rachel Kaufmann - Assoc Director Science</p> <p>Team Lead</p>	<p>06/13/2019</p>	<p><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</p> <p>(check if applicable)  <input type="checkbox"/> Local IRB  <input type="checkbox"/> CDC Exemption  <input type="checkbox"/> CDC IRB</p> <p><u>Comments:</u></p>
<p>-</p> <p>Division ADS</p>	<p>MM/dd/yyyy</p>	<p><input 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</p> <p>(check if applicable)  <input type="checkbox"/> Local IRB  <input type="checkbox"/> CDC Exemption  <input type="checkbox"/> CDC IRB</p> <p><u>Comments:</u></p>
<p>Joan Redmond Leonard - PUBLIC HEALTH ANALYST</p> <p>CUC ADS, Deputy ADS, or Human Subjects Contact</p>	<p>06/27/2019</p>	<p><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</p> <p>(check if applicable)  <input type="checkbox"/> Local IRB  <input type="checkbox"/> CDC Exemption  <input type="checkbox"/> CDC IRB</p> <p><u>Comments:</u></p>