

1. DATE ISSUED MM/DD/YYYY | 2. CFDA NO. | 3. ASSISTANCE TYPE  
 07/30/2016 | 93.073 | Cooperative Agreement

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**  
**Centers for Disease Control and Prevention**  
**CDC Office of Financial Resources**  
 2920 Brandywine Road  
 Atlanta, GA 30341

1a. SUPERSEDES AWARD NOTICE dated  
 except that any additions or restrictions previously imposed remain  
 in effect unless specifically rescinded

4. GRANT NO. | 5. ACTION TYPE  
 1 NU50DD000001-01-00 | New  
 Formerly

6. PROJECT PERIOD MM/DD/YYYY | MM/DD/YYYY  
 From 08/01/2016 | Through 07/31/2021

7. BUDGET PERIOD MM/DD/YYYY | MM/DD/YYYY  
 From 08/01/2016 | Through 07/31/2017

**NOTICE OF AWARD**  
 AUTHORIZATION (Legislation/Regulations)  
 [AWARD AUTHORITY NOT DEFINED FOR DD16-1603]

8. TITLE OF PROJECT (OR PROGRAM)  
 National Center on Birth Defects and Developmental Disabilities (NCBDDD)

9a. GRANTEE NAME AND ADDRESS  
 State of Missouri  
 920 Wildwood Dr  
 Jefferson City, MO 65109-5796

9b. GRANTEE PROJECT DIRECTOR  
 Loise Wambugh  
 920 Wildwood Drive  
 Jefferson City, MO 65102-0570  
 Phone: 573-751-6343

10a. GRANTEE AUTHORIZING OFFICIAL  
 Mr. Bret Fischer  
 920 WILDWOOD DR  
 JEFFERSON CITY, MO 65109-5796  
 Phone: 573-751-6014

10b. FEDERAL PROJECT OFFICER  
 William Paradies  
 Centers for Disease Control and Prevention  
 CCHP/NCBDDD/DBDDD/BDB  
 Atlanta, GA 30333  
 Phone: 404-498-3919

**ALL AMOUNTS ARE SHOWN IN USD**

<b>11. APPROVED BUDGET (Excludes Direct Assistance)</b>	
I Financial Assistance from the Federal Awarding Agency Only	<b>I</b>
II Total project costs including grant funds and all other financial participation	
a. Salaries and Wages .....	124,759.00
b. Fringe Benefits .....	62,380.00
c. Total Personnel Costs .....	187,139.00
d. Equipment .....	0.00
e. Supplies .....	30,218.00
f. Travel .....	87,489.00
g. Construction .....	0.00
h. Other .....	5,989.00
i. Contractual .....	45,000.00
j. TOTAL DIRECT COSTS →	355,835.00
k. INDIRECT COSTS	44,165.00
<b>l. TOTAL APPROVED BUDGET</b>	<b>400,000.00</b>
m. Federal Share	400,000.00
n. Non-Federal Share	0.00

<b>12. AWARD COMPUTATION</b>	
a. Amount of Federal Financial Assistance (from item 11m)	400,000.00
b. Less Unobligated Balance From Prior Budget Periods	0.00
c. Less Cumulative Prior Award(s) This Budget Period	0.00
<b>d. AMOUNT OF FINANCIAL ASSISTANCE THIS ACTION</b>	<b>400,000.00</b>
<b>13. Total Federal Funds Awarded to Date for Project Period</b>	<b>400,000.00</b>

<b>14. RECOMMENDED FUTURE SUPPORT</b> (Subject to the availability of funds and satisfactory progress of the project):			
YEAR	TOTAL DIRECT COSTS	YEAR	TOTAL DIRECT COSTS
a. 2		d. 5	
b. 3		e. 6	
c. 4		f. 7	

**15. PROGRAM INCOME SHALL BE USED IN ACCORD WITH ONE OF THE FOLLOWING ALTERNATIVES:**

a. DEDUCTION b. ADDITIONAL COSTS c. MATCHING d. OTHER RESEARCH (Add / Deduct Option) e. OTHER (See REMARKS)	<b>b</b>
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**16. THIS AWARD IS BASED ON AN APPLICATION SUBMITTED TO, AND AS APPROVED BY, THE FEDERAL AWARDING AGENCY ON THE ABOVE TITLED PROJECT AND IS SUBJECT TO THE TERMS AND CONDITIONS INCORPORATED EITHER DIRECTLY OR BY REFERENCE IN THE FOLLOWING:**

a. The grant program legislation  
 b. The grant program regulations.  
 c. This award notice including terms and conditions, if any, noted below under REMARKS.  
 d. Federal administrative requirements, cost principles and audit requirements applicable to this grant.

In the event there are conflicting or otherwise inconsistent policies applicable to the grant, the above order of precedence shall prevail. Acceptance of the grant terms and conditions is acknowledged by the grantee when funds are drawn or otherwise obtained from the grant payment system.

REMARKS (Other Terms and Conditions Attached -  Yes  No)

GRANTS MANAGEMENT OFFICIAL: Patricia French, Grants Management Officer

17. OBJ CLASS	41.51	18a. VENDOR CODE		18b. EIN		19. DUNS	878092600	20. CONG. DIST.	03
FY-ACCOUNT NO.	DOCUMENT NO.	CFDA	ADMINISTRATIVE CODE	AMT ACTION FIN ASST	APPROPRIATION				
21. a.	6-939061A	b. 000001DD16	c. 93.073	d. DD	e. \$200,000.00	f. 75-16-0956			
22. a.	6-93906FV	b. 000001DD16	c. 93.073	d. DD	e. \$200,000.00	f. 75-1519-0943			
23. a.		b.	c.	d.	e.	f.			

NOTICE OF AWARD (Continuation Sheet)

PAGE 2 of 2	DATE ISSUED 07/30/2016
GRANT NO. 1 NU50DD000001-01-00	

**Direct Assistance**

BUDGET CATEGORIES	PREVIOUS AMOUNT (A)	AMOUNT THIS ACTION (B)	TOTAL (A + B)
Personnel	\$0.00	\$0.00	\$0.00
Fringe Benefits	\$0.00	\$0.00	\$0.00
Travel	\$0.00	\$0.00	\$0.00
Equipment	\$0.00	\$0.00	\$0.00
Supplies	\$0.00	\$0.00	\$0.00
Contractual	\$0.00	\$0.00	\$0.00
Construction	\$0.00	\$0.00	\$0.00
Other	\$0.00	\$0.00	\$0.00
<b>Total</b>	<b>\$0.00</b>	<b>\$0.00</b>	<b>\$0.00</b>

# AWARD ATTACHMENTS

State of Missouri

1 NU50DD000001-01-00

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1. Terms and Conditions
2. Summary Statement

Funding Opportunity Announcement (FOA) Number: DD16-1605

Award Number: **1 NU50 DD 000001-01**

Award Type: Cooperative Agreement

Applicable Regulations: 45 Code of Federal Regulations (CFR) Part 75, Uniform Administrative Requirements, Cost Principles, and Audit Requirements for HHS Awards

## **45 CFR Part 75 supersedes regulations at 45 CFR Part 74 and Part 92**

### **AWARD INFORMATION**

**Incorporation:** The Centers for Disease Control and Prevention (CDC) hereby incorporates Funding Opportunity Announcement number DD16-1605: Surveillance, intervention, and referral to services activities for infants with microcephaly or other adverse outcomes linked with the Zika virus and application dated June 28, 2016, as may be amended, which are hereby made a part of this Research award hereinafter referred to as the Notice of Award (NoA). The Department of Health and Human Services (HHS) grant recipients must comply with all terms and conditions outlined in their NoA, including grants policy terms and conditions contained in applicable HHS Grants Policy Statements, and requirements imposed by program statutes and regulations and HHS grant administration regulations, as applicable; as well as any requirements or limitations in any applicable appropriations acts. The term grant is used throughout this notice and includes cooperative agreements.

**Note:** In the event that any requirement in this Notice of Award, the Funding Opportunity Announcement, the HHS GPS, 45 CFR Part 75, or applicable statutes/appropriations acts conflict, then statutes and regulations take precedence.

**Approved Funding:** Funding in the amount of **\$400,000** is approved for the Year **01** budget period, which is **August 01, 2016** through **July 31, 2017**. All future year funding will be based on satisfactory programmatic progress and the availability of funds.

**Budget Revision Requirement:** By **September 01, 2016** the grantee must submit the following to the GMS via email:

- 1) **Supplies:** For miscellaneous office supplies requested, include the estimated amount per month and number of months supplies are requested in this budget category.
- 2) **Contractual:** Include the following information for each Contractor: Name of Contractor, Method of Selection, Period of Performance, Scope of Work, Method of Accountability, and an Itemized Budget and Justification.
- 3) **Travel:** Provide a narrative justification describing what the travel staff members will perform. List where travel will be undertaken, number of trips planned, who will be making the trips, and approximate dates.
- 4) **Other:** Provide a narrative justification related to program objectives

Failure to submit the required information in a timely manner may adversely affect the future funding of this project. If the information cannot be provided by the due date, you are required to contact the GMS/GMO identified in the **Staff Contacts section of this notice before the due date.**

**Note:** Refer to the Payment Information section for draw down and Payment Management System (PMS) subaccount information.

**Objective/Technical Review Statement Response Requirement:** The review comments on the strengths and weaknesses of the proposal are provided as part of this award. A response to the weaknesses in these statements must be submitted to and approved, in writing, by the Grants Management Specialist/Grants Management Officer (GMS/GMO) noted in the Staff Contacts section of this NoA, no later than 30 days from the budget period start date. Failure to submit the required information by the due date, **September 01, 2016**, will cause delay in programmatic progress and will adversely affect the future funding of this project.

**Award Funding:** Not funded by the Prevention and Public Health Fund

**Program Income:** Any program income generated under this grant or cooperative agreement will be used in

accordance with the Addition alternative.

**Addition alternative:** Under this alternative, program income is added to the funds committed to the project/program and is used to further eligible project/program objectives.

**Note:** The disposition of program income must have written prior approval from the GMO.

## FUNDING RESTRICTIONS AND LIMITATIONS

**Indirect Costs:** Indirect costs are approved based on the Indirect Cost Rate Agreement dated February 4, 2016, which calculates indirect costs as follows, a Fixed is approved at a rate of 23.6% of the base, which includes, direct salaries and wages, including all fringe benefits. The effective dates of this indirect cost rate are from July 1, 2016 to June 30, 2017.

A. Cap on Salaries (Sec. 203): None of the funds appropriated in this title shall be used to pay the salary of an individual, through a grant or other extramural mechanism, at a rate in excess of Executive Level II.

Note: The salary rate limitation does not restrict the salary that an organization may pay an individual working under an HHS contract or order; it merely limits the portion of that salary that may be paid with Federal funds.

B. Gun Control Prohibition (Div. H, Title II, Sec. 217): None of the funds made available in this title may be used, in whole or in part, to advocate or promote gun control.

C. Lobbying Restrictions (Sec. 503):

- 503(a): No part of any appropriation contained in this Act or transferred pursuant to section 4002 of Public Law 111-148 shall be used, other than for normal and recognized executive-legislative relationships, for publicity or propaganda purposes, for the preparation, distribution, or use of any kit, pamphlet, booklet, publication, electronic communication, radio, television, or video presentation designed to support or defeat the enactment of legislation before the Congress or any State or local legislature or legislative body, except in presentation of the Congress or any State or local legislature itself, or designed to support or defeat any proposed or pending regulation, administrative action, or order issued by the executive branch of any State or local government itself.
- 503 (b): No part of any appropriation contained in this Act or transferred pursuant to section 4002 of Public Law 111-148 shall be used to pay the salary or expenses of any grant or contract recipient, or agent acting for such recipient, related to any activity designed to influence the enactment of legislation, appropriations, regulation, administrative action, or Executive order proposed or pending before the Congress or any State government, State legislature or local legislature or legislative body, other than normal and recognized executive legislative relationships or participation by an agency or officer of an State, local or tribal government in policymaking and administrative processes within the executive branch of that government.
- 503(c): The prohibitions in subsections (a) and (b) shall include any activity to advocate or promote any proposed, pending or future Federal, State or local tax increase, or any proposed, pending, or future requirement or restriction on any legal consumer product, including its sale or marketing, including but not limited to the advocacy or promotion of gun control.

For additional information, see Additional Requirement 12 at

<http://www.cdc.gov/grants/additionalrequirements/index.html> and Anti Lobbying Restrictions for CDC Grantees at [http://www.cdc.gov/grants/documents/Anti-Lobbying\\_Restrictions\\_for\\_CDC\\_Grantees\\_July\\_2012.pdf](http://www.cdc.gov/grants/documents/Anti-Lobbying_Restrictions_for_CDC_Grantees_July_2012.pdf)

D. Needle Exchange (Sec. 521): Notwithstanding any other provision of this Act, no funds appropriated in this Act shall be used to carry out any program of distributing sterile needles or syringes for the hypodermic injection of any illegal drug.

E. Blocking access to pornography (Sec. 526): (a) None of the funds made available in this Act may be used to maintain or establish a computer network unless such network blocks the viewing, downloading, and exchanging of pornography; (b) Nothing in subsection (a) shall limit the use of funds necessary for any Federal, State, tribal, or local law enforcement agency or any other entity carrying out criminal investigations, prosecution, or adjudication activities.

**Rent or Space Costs:** Grantees are responsible for ensuring that all costs included in this proposal to establish billing or final indirect cost rates are allowable in accordance with the requirements of the Federal award(s) to which they apply, including 45 CFR Part 75, Uniform Administrative Requirements, Cost Principles, and Audit Requirements for HHS Awards. The grantee also has a responsibility to ensure sub-recipients expend funds in compliance with applicable federal laws and regulations. Furthermore, it is the responsibility of the grantee to ensure rent is a legitimate direct cost line item, which the grantee has supported in current and/or prior projects and these same costs have been treated as indirect costs that have not been claimed as direct costs. If rent is claimed as direct cost, the grantee must provide a narrative justification, which describes their prescribed policy to include the effective date to the assigned Grants Management Specialist (GMS) identified in the CDC Contacts for this award.

**Trafficking In Persons:** This award is subject to the requirements of the Trafficking Victims Protection Act of 2000, as amended (22 U.S.C. Part 7104(g)).

**Cancel Year:** 31 U.S.C. Part 1552(a) Procedure for Appropriation Accounts Available for Definite Periods states the following, On September 30<sup>th</sup> of the 5<sup>th</sup> fiscal year after the period of availability for obligation of a fixed appropriation account ends, the account shall be closed and any remaining balances (whether obligated or unobligated) in the account shall be canceled and thereafter shall not be available for obligation or expenditure for any purpose. An example is provided below:

Fiscal Year (FY) 2016 funds will expire September 30, 2021. All FY 2016 funds should be drawn down and reported to Payment Management Services (PMS) prior to September 30, 2021. After this date, corrections or cash requests will not be permitted.

## REPORTING REQUIREMENTS

**Annual Federal Financial Report (FFR, SF-425):** The Annual Federal Financial Report (FFR) SF-425 is required and must be to your grants management specialist (GMS) no later than 90 days after the end of budget period. The FFR for this budget period is due to the GMS/GMO by **October 30, 2017**. Reporting timeframe is **August 01, 2016** through **July 31, 2017**.

The FFR may be downloaded from the following website below and submitted to the GMS via email.  
[https://www.whitehouse.gov/sites/default/files/omb/grants/approved\\_forms/SF-425.pdf](https://www.whitehouse.gov/sites/default/files/omb/grants/approved_forms/SF-425.pdf)

The FFR should only include those funds authorized and disbursed during the timeframe covered by the report. The final FFR must indicate the exact balance of unobligated funds and may not reflect any unliquidated obligations. There must be no discrepancies between the final FFR expenditure data and the Payment Management System's (PMS) cash transaction data. All Federal reporting in PMS is unchanged

Failure to submit the required information in a timely manner may adversely affect the future funding of this project. If the information cannot be provided by the due date, the grantee is required to contact the Grants Officer listed in the contacts section of this notice before the due date.

FFR (SF-425) instructions for CDC Grantees are available at <http://grants.nih.gov/grants/forms.htm>. For further information, contact GrantsInfo@nih.gov. Additional resources concerning the eFSR/FFR system, including a User Guide and an on-line demonstration, can be found on the eRA Commons Support Page: <http://www.cdc.gov/od/pgo/funding/grants/eramain.shtm>.

**Performance Reporting:** The annual Performance Report is due 90 days after the end of the final budget period. Please reference the closeout requirements which follows in this NOA.

**Audit Requirement:** Domestic Organizations: An organization that expends \$750,000 or more in a fiscal year in

Federal awards shall have a single or program-specific audit conducted for that year in accordance with the provisions of 45 CFR Part 75. The audit period is an organization's fiscal year. The audit must be completed along with a data collection form (SF-SAC), and the reporting package shall be submitted within the earlier of 30 days after receipt of the auditor's report(s), or nine (9) months after the end of the audit period. The audit report must be sent to

Federal Audit Clearing House Internet Data Entry System

Electronic Submission: [https://harvester.census.gov/facides/\(S\(0vkw1zaelyzjibnahocga5i0\)\)/account/login.aspx](https://harvester.census.gov/facides/(S(0vkw1zaelyzjibnahocga5i0))/account/login.aspx)

AND

Office of Grants Services, Financial Assessment and Audit Resolution Unit

Electronic Copy to:

[OGS.Audit.Resolution@cdc.gov](mailto:OGS.Audit.Resolution@cdc.gov) (CDC Office of Grants Services)

After receipt of the audit report, CDC will resolve findings by issuing Final Determination Letters

Audit requirements for Subrecipients to whom 45 CFR 75 Subpart F applies: The grantee must ensure that the subrecipients receiving CDC funds also meet these requirements. The grantee must also ensure to take appropriate corrective action within six months after receipt of the subrecipient audit report in instances of non-compliance with applicable Federal law and regulations (45 CFR 75 Subpart F and HHS Grants Policy Statement). The grantee may consider whether subrecipient audits necessitate adjustment of the grantee's own accounting records. If a subrecipient is not required to have a program-specific audit, the grantee is still required to perform adequate monitoring of subrecipient activities. The grantee shall require each subrecipient to permit the independent auditor access to the subrecipient's records and financial statements. The grantee must include this requirement in all subrecipient contracts.

Note: The standards set forth in 45 CFR Part 75 Subpart F will apply to audits of fiscal years beginning on or after December 26, 2014.

**Federal Funding Accountability and Transparency Act (FFATA):**

In accordance with 2 CFR Chapter 1, Part 170 Reporting Sub-Award And Executive Compensation Information, Prime Awardees awarded a federal grant are required to file a FFATA sub-award report by the end of the month following the month in which the prime awardee awards any sub-grant equal to or greater than \$25,000.

Pursuant to 45 CFR Part 75, §75.502, a grant sub-award includes the provision of any commodities (food and non-food) to the sub-recipient where the sub-recipient is required to abide by terms and conditions regarding the use or future administration of those goods. If the sub-awardee merely consumes or utilizes the goods, the commodities are not in and of themselves considered sub-awards.

2 CFR Part 170: [http://www.ecfr.gov/cgi-bin/text-idx?tpl=/ecfrbrowse/Title02/2cfr170\\_main\\_02.tpl](http://www.ecfr.gov/cgi-bin/text-idx?tpl=/ecfrbrowse/Title02/2cfr170_main_02.tpl)

FFATA: [www.fsrs.gov](http://www.fsrs.gov).

Reporting of First-Tier Sub-awards

Applicability: Unless you are exempt (gross income from all sources reported in last tax return is under \$300,000), you must report each action that obligates \$25,000 or more in Federal funds that does not include Recovery funds (as defined in section 1512(a)(2) of the American Recovery and Reinvestment Act of 2009, Pub. L. 111-5) for a sub-award to an entity.

Reporting: Report each obligating action of this award term to [www.fsrs.gov](http://www.fsrs.gov). For sub-award information, report no later than the end of the month following the month in which the obligation was made. (For example, if the obligation was made on November 7, 2010, the obligation must be reported by no later than December 31, 2010). You must report the information about each obligating action that the submission instructions posted at [www.fsrs.gov](http://www.fsrs.gov) specify.

Total Compensation of Recipient Executives: You must report total compensation for each of your five most highly compensated executives for the preceding completed fiscal year, if:

- The total Federal funding authorized to date under this award is \$25,000 or more;
- In the preceding fiscal year, you received—
  - 80 percent or more of your annual gross revenues from Federal procurement contracts (and subcontracts) and Federal financial assistance subject to the Transparency Act, as defined at 2 CFR Part 170.320 (and sub-awards); and
  - \$25,000,000 or more in annual gross revenues from Federal procurement contracts (and subcontracts) and Federal financial assistance subject to the Transparency Act, as defined at 2 CFR Part 170.320 (and sub-awards); and
  - The public does not have access to information about the compensation of the executives through periodic reports filed under section 13(a) or 15(d) of the Securities Exchange Act of 1934 (15 U.S.C. Part 78m(a), 78o(d)) or section 6104 of the Internal Revenue Code of 1986. (To determine if the public has access to the compensation information, see the U.S. Security and Exchange Commission total compensation filings at <http://www.sec.gov/answers/execomp.htm>).

Report executive total compensation as part of your registration profile at <http://www.sam.gov>. Reports should be made at the end of the month following the month in which this award is made and annually thereafter.

Total Compensation of Sub-recipient Executives: Unless you are exempt (gross income from all sources reported in last tax return is under \$300,000), for each first-tier sub-recipient under this award, you must report the names and total compensation of each of the sub-recipient's five most highly compensated executives for the sub-recipient's preceding completed fiscal year, if:

- In the sub-recipient's preceding fiscal year, the sub-recipient received—
  - 80 percent or more of its annual gross revenues from Federal procurement contracts (and subcontracts) and Federal financial assistance subject to the Transparency Act, as defined at 2 CFR Part 170.320 (and sub-awards); and
  - \$25,000,000 or more in annual gross revenues from Federal procurement contracts (and subcontracts), and Federal financial assistance subject to the Transparency Act (and sub-awards); and
  - The public does not have access to information about the compensation of the executives through periodic reports filed under section 13(a) or 15(d) of the Securities Exchange Act of 1934 (15 U.S.C. Part 78m(a), 78o(d)) or section 6104 of the Internal Revenue Code of 1986. (To determine if the public has access to the compensation information, see the U.S. Security and Exchange Commission total compensation filings at <http://www.sec.gov/answers/execomp.htm>).

You must report sub-recipient executive total compensation to the grantee by the end of the month following the month during which you make the sub-award. For example, if a sub-award is obligated on any date during the month of October of a given year (i.e., between October 1st and 31st), you must report any required compensation information of the sub-recipient by November 30th of that year.

Definitions:

- Entity means all of the following, as defined in 2 CFR Part 25 (Appendix A, Paragraph(C)(3)):
  - Governmental organization, which is a State, local government, or Indian tribe;
  - Foreign public entity;



- Domestic or foreign non-profit organization;
- Domestic or foreign for-profit organization;
- Federal agency, but only as a sub-recipient under an award or sub-award to a non-Federal entity.
- Executive means officers, managing partners, or any other employees in management positions.
- Sub-award: a legal instrument to provide support to an eligible sub-recipient for the performance of any portion of the substantive project or program for which the grantee received this award. The term does not include the grantees procurement of property and services needed to carry out the project or program (for further explanation, see 45 CFR Part 75). A sub-award may be provided through any legal agreement, including an agreement that the grantee or a sub-recipient considers a contract.
- Sub-recipient means an entity that receives a sub-award from you (the grantee) under this award; and is accountable to the grantee for the use of the Federal funds provided by the sub-award.
- Total compensation means the cash and non-cash dollar value earned by the executive during the grantee's or sub-recipient's preceding fiscal year and includes the following (for more information see 17 CFR Part 229.402(c)(2)):
  - Salary and bonus
  - Awards of stock, stock options, and stock appreciation rights. Use the dollar amount recognized for financial statement reporting purposes with respect to the fiscal year in accordance with the Statement of Financial Accounting Standards No. 123 (Revised 2004) (FAS 123R), Shared Based Payments.
  - Earnings for services under non-equity incentive plans. This does not include group life, health, hospitalization or medical reimbursement plans that do not discriminate in favor of executives, and are available generally to all salaried employees.
  - Change in pension value. This is the change in present value of defined benefit and actuarial pension plans.
  - Above-market earnings on deferred compensation which is not tax-qualified.
  - Other compensation, if the aggregate value of all such other compensation (e.g. severance, termination payments, value of life insurance paid on behalf of the employee, perquisites or property) for the executive exceeds \$10,000.

## GENERAL REQUIREMENTS

**Travel Cost:** In accordance with HHS Grants Policy Statement, travel costs are only allowable where such travel will provide direct benefit to the project or program. There must be a direct benefit imparted on behalf of the traveler as it applies to the approved activities of the NoA. To prevent disallowance of cost, the grantee is responsible for ensuring that only allowable travel reimbursements are applied in accordance with their organization's established travel policies and procedures. Grantees approved policies must meet the requirements of 45 CFR Part 75, as applicable.

**Food and Meals:** Costs associated with food or meals are allowable when consistent with applicable federal regulations and HHS policies and guidance, which can be found at [http://www.hhs.gov/asfr/ogapa/acquisition/effspendpol\\_memo.html](http://www.hhs.gov/asfr/ogapa/acquisition/effspendpol_memo.html). In addition, costs must be proposed in accordance with grantee approved policies and a determination of reasonableness has been performed by the

grantees. Grantee approved policies must meet the requirements of 45 CFR Part 75, as applicable.

**Prior Approval:** All requests, which require prior approval, must bear the signature of an authorized official of the business office of the grantee organization as well as the principal investigator or program or project director named on this NoA. The grantee must submit these requests by **March 30, 2017** or no later than 120 days prior to this budget period's end date. Any requests received that reflect only one signature will be returned to the grantee unprocessed. Additionally, any requests involving funding issues must include an itemized budget and a narrative justification of the request.

The following types of requests require prior approval unless waived by approved expanded authorities (see Award Information section).

- Lift funding restriction, withholding, or disallowance
- Redirection of funds
- Change in scope
- Implement a new activity or enter into a sub-award that is not specified in the most recently approved budget
- Apply for supplemental funds
- Response to the Objective/Technical Review Statement
- Change in key personnel
- Extensions
- Conferences or meetings that exceed cost threshold

Templates for prior approval requests can be found at:

<http://www.cdc.gov/grants/alreadyhavegrant/priorapprovalrequests.html>

**Key Personnel:** In accordance with 2 CFR Parts 200.308 and 215.25(c)(2) & (3), CDC grantees must obtain prior approval from CDC for (1) change in the project director/principal investigator, business official, authorized organizational representative or other key persons specified in the FOA, application or award document; and (2) the disengagement from the project for more than three months, or a 25 percent reduction in time devoted to the project, by the approved project director or principal investigator.

**Inventions:** Acceptance of grant funds obligates grantees to comply with the standard patent rights clause in 37 CFR Part 401.14.

**Publications:** Publications, journal articles, etc. produced under a CDC grant support project must bear an acknowledgment and disclaimer, as appropriate, for example:

This publication (journal article, etc.) was supported by the Grant or Cooperative Agreement Number, **1 NU50 DD00001-01**, funded by the Centers for Disease Control and Prevention. Its contents are solely the responsibility of the authors and do not necessarily represent the official views of the Centers for Disease Control and Prevention or the Department of Health and Human Services.

**Acknowledgment Of Federal Support:** When issuing statements, press releases, requests for proposals, bid solicitations and other documents describing projects or programs funded in whole or in part with Federal money, all awardees receiving Federal funds, including and not limited to State and local governments and grantees of Federal research grants, shall clearly state:

- percentage of the total costs of the program or project which will be financed with Federal money
- dollar amount of Federal funds for the project or program, and
- percentage and dollar amount of the total costs of the project or program that will be financed by non-governmental sources.

**Copyright Interests Provision:** This provision is intended to ensure that the public has access to the results and accomplishments of public health activities funded by CDC. Pursuant to applicable grant regulations and CDC's Public Access Policy, Recipient agrees to submit into the National Institutes of Health (NIH) Manuscript Submission (NIHMS) system an electronic version of the final, peer-reviewed manuscript of any such work developed under this award upon acceptance for publication, to be made publicly available no later than 12 months after the official date of publication. Also at the time of submission, Recipient and/or the Recipient's

submitting author must specify the date the final manuscript will be publicly accessible through PubMed Central (PMC). Recipient and/or Recipient's submitting author must also post the manuscript through PMC within twelve (12) months of the publisher's official date of final publication; however the author is strongly encouraged to make the subject manuscript available as soon as possible. The recipient must obtain prior approval from the CDC for any exception to this provision.

The author's final, peer-reviewed manuscript is defined as the final version accepted for journal publication, and includes all modifications from the publishing peer review process, and all graphics and supplemental material associated with the article. Recipient and its submitting authors working under this award are responsible for ensuring that any publishing or copyright agreements concerning submitted articles reserve adequate right to fully comply with this provision and the license reserved by CDC. The manuscript will be hosted in both PMC and the CDC Stacks institutional repository system. In progress reports for this award, recipient must identify publications subject to the CDC Public Access Policy by using the applicable NIHMS identification number for up to three (3) months after the publication date and the PubMed Central identification number (PMCID) thereafter.

**Disclaimer for Conference/Meeting/Seminar Materials:** Disclaimers for conferences/meetings, etc. and/or publications: If a conference/meeting/seminar is funded by a grant, cooperative agreement, sub-grant and/or a contract the grantee must include the following statement on conference materials, including promotional materials, agenda, and internet sites:

Funding for this conference was made possible (in part) by the Centers for Disease Control and Prevention. The views expressed in written conference materials or publications and by speakers and moderators do not necessarily reflect the official policies of the Department of Health and Human Services, nor does the mention of trade names, commercial practices, or organizations imply endorsement by the U.S. Government.

**Logo Use for Conference and Other Materials:** Neither the Department of Health and Human Services (HHS) nor the CDC logo may be displayed if such display would cause confusion as to the funding source or give false appearance of Government endorsement. Use of the HHS name or logo is governed by U.S.C. Part 1320b-10, which prohibits misuse of the HHS name and emblem in written communication. A non-federal entity is unauthorized to use the HHS name or logo governed by U.S.C. Part 1320b-10. The appropriate use of the HHS logo is subject to review and approval of the HHS Office of the Assistant Secretary for Public Affairs (OASPA). Moreover, the HHS Office of the Inspector General has authority to impose civil monetary penalties for violations (42 CFR Part 1003). Accordingly, neither the HHS nor the CDC logo can be used by the grantee without the express, written consent of either the CDC Project Officer or the CDC Grants Management Officer. It is the responsibility of the grantee to request consent for use of the logo in sufficient detail to ensure a complete depiction and disclosure of all uses of the Government logos. In all cases for utilization of Government logos, the grantee must ensure written consent is received from the Project Officer and/or the Grants Management Officer.

**Equipment and Products:** To the greatest extent practicable, all equipment and products purchased with CDC funds should be American-made. CDC defines equipment as tangible non-expendable personal property (including exempt property) charged directly to an award having a useful life of more than one year AND an acquisition cost of \$5,000 or more per unit. However, consistent with grantee policy, a lower threshold may be established. Please provide the information to the Grants Management Officer to establish a lower equipment threshold to reflect your organization's policy.

The grantee may use its own property management standards and procedures, provided it observes provisions in applicable grant regulations found at 45 CFR Part 75.

**Federal Information Security Management Act (FISMA):** All information systems, electronic or hard copy, that contain federal data must be protected from unauthorized access. This standard also applies to information associated with CDC grants. Congress and the OMB have instituted laws, policies and directives that govern the creation and implementation of federal information security practices that pertain specifically to grants and contracts. The current regulations are pursuant to the Federal Information Security Management Act (FISMA), Title III of the E-Government Act of 2002, PL 107-347.

FISMA applies to CDC grantees only when grantees collect, store, process, transmit or use information on behalf of HHS or any of its component organizations. In all other cases, FISMA is not applicable to recipients of grants, including cooperative agreements. Under FISMA, the grantee retains the original data and intellectual property,

and is responsible for the security of these data, subject to all applicable laws protecting security, privacy, and research. If/When information collected by a grantee is provided to HHS, responsibility for the protection of the HHS copy of the information is transferred to HHS and it becomes the agency's responsibility to protect that information and any derivative copies as required by FISMA. For the full text of the requirements under Federal Information Security Management Act (FISMA), Title III of the E-Government Act of 2002 Pub. L. No. 107-347, please review the following website:

[http://frwebgate.access.gpo.gov/cgi-bin/getdoc.cgi?dbname=107\\_cong\\_public\\_laws&docid=f:publ347.107.pdf](http://frwebgate.access.gpo.gov/cgi-bin/getdoc.cgi?dbname=107_cong_public_laws&docid=f:publ347.107.pdf)

**Pilot Program for Enhancement of Contractor Employee Whistleblower Protections:** Grantees are hereby given notice that the 48 CFR section 3.908, implementing section 828, entitled "Pilot Program for Enhancement of Contractor Employee Whistleblower Protections," of the National Defense Authorization Act (NDAA) for Fiscal Year (FY) 2013 (Pub. L. 112-239, enacted January 2, 2013), applies to this award.

#### Federal Acquisition Regulations

As promulgated in the Federal Register, the relevant portions of 48 CFR section 3.908 read as follows (note that use of the term "contract," "contractor," "subcontract," or "subcontractor" for the purpose of this term and condition, should be read as "grant," "grantee," "subgrant," or "subgrantee"):

3.908 Pilot program for enhancement of contractor employee whistleblower protections.

3.908-1 Scope of section.

(a) This section implements [41 U.S.C. 4712](#).

(b) This section does not apply to-

(1) DoD, NASA, and the Coast Guard; or

(2) Any element of the intelligence community, as defined in section 3(4) of the National Security Act of 1947 (50 U.S.C. 3003(4)). This section does not apply to any disclosure made by an employee of a contractor or subcontractor of an element of the intelligence community if such disclosure-

(i) Relates to an activity of an element of the intelligence community; or

(ii) Was discovered during contract or subcontract services provided to an element of the intelligence community.

3.908-2 Definitions.

As used in this section-

"Abuse of authority" means an arbitrary and capricious exercise of authority that is inconsistent with the mission of the executive agency concerned or the successful performance of a contract of such agency.

"Inspector General" means an Inspector General appointed under the Inspector General Act of 1978 and any Inspector General that receives funding from, or has oversight over contracts awarded for, or on behalf of, the executive agency concerned.

3.908-3 Policy.

(a) Contractors and subcontractors are prohibited from discharging, demoting, or otherwise discriminating against an employee as a reprisal for disclosing, to any of the entities listed at paragraph (b) of this subsection, information that the employee reasonably believes is evidence of gross mismanagement of a Federal contract, a gross waste of Federal funds, an abuse of authority relating to a Federal contract, a substantial and specific danger to public health or safety, or a violation of law, rule, or regulation related to a Federal contract (including the competition for or negotiation of a contract). A reprisal is prohibited even if it is undertaken at the request of an executive branch official, unless the request takes the form of a non-discretionary directive and is within the authority of the executive branch official making the request.

(b) Entities to whom disclosure may be made.

(1) A Member of Congress or a representative of a committee of Congress.

(2) An Inspector General.

(3) The Government Accountability Office.

(4) A Federal employee responsible for contract oversight or management at the relevant agency.

(5) An authorized official of the Department of Justice or other law enforcement agency.

(6) A court or grand jury.

(7) A management official or other employee of the contractor or subcontractor who has the responsibility to investigate, discover, or address misconduct.

(c) An employee who initiates or provides evidence of contractor or subcontractor misconduct in any judicial or administrative proceeding relating to waste, fraud, or abuse on a Federal contract shall be deemed to have made a disclosure.

3.908-9 Contract clause.

Contractor Employee Whistleblower Rights and Requirement to Inform Employees of Whistleblower Rights (Sept. 2013)

(a) This contract and employees working on this contract will be subject to the whistleblower rights and remedies in the pilot program on Contractor employee whistleblower protections established at [41 U.S.C. 4712](#) by section 828 of the National Defense Authorization Act for Fiscal Year 2013 (Pub. L. 112-239) and FAR [3.908](#).

(b) The Contractor shall inform its employees in writing, in the predominant language of the workforce, of employee whistleblower rights and protections under [41 U.S.C. 4712](#), as described in section [3.908](#) of the Federal Acquisition Regulation.

(c) The Contractor shall insert the substance of this clause, including this paragraph (c), in all subcontracts over the simplified acquisition threshold.

## PAYMENT INFORMATION

**Automatic Drawdown (Direct/Advance Payments):** Payment under this award will be made available through the Department of Health and Human Services (HHS) Payment Management System (PMS). PMS will forward instructions for obtaining payments.

PMS correspondence, mailed through the U.S. Postal Service, should be addressed as follows:

Director, Payment Management Services  
P.O. Box 6021  
Rockville, MD 20852  
Phone Number: (877) 614-5533  
Email: [PMSSupport@psc.gov](mailto:PMSSupport@psc.gov)  
Website: <http://www.dpm.psc.gov/help/help.aspx?explorer.event=true>

**Note:** To obtain the contact information of DPM staff within respective Payment Branches refer to the links listed below:

- University and Non-Profit Payment Branch:  
[http://www.dpm.psc.gov/contacts/dpm\\_contact\\_list/univ\\_nonprofit.aspx?explorer.event=true](http://www.dpm.psc.gov/contacts/dpm_contact_list/univ_nonprofit.aspx?explorer.event=true)
- Governmental and Tribal Payment Branch:  
[http://www.dpm.psc.gov/contacts/dpm\\_contact\\_list/gov\\_tribal.aspx?explorer.event=true](http://www.dpm.psc.gov/contacts/dpm_contact_list/gov_tribal.aspx?explorer.event=true)
- Cross Servicing Payment Branch:  
[http://www.dpm.psc.gov/contacts/dpm\\_contact\\_list/cross\\_servicing.aspx](http://www.dpm.psc.gov/contacts/dpm_contact_list/cross_servicing.aspx)
- International Payment Branch:  
Bhavin Patel (301) 492-4918  
Email: [Bhavin.patel@psc.hhs.gov](mailto:Bhavin.patel@psc.hhs.gov)

If a carrier other than the U.S. Postal Service is used, such as United Parcel Service, Federal Express, or other commercial service, the correspondence should be addressed as follows:

U.S. Department of Health and Human Services  
Division of Payment Management  
7700 Wisconsin Avenue, Suite 920  
Bethesda, MD 20814

To expedite your first payment from this award, attach a copy of the Notice of Grant/Cooperative Agreement to your payment request form.

**Payment Management System Subaccount:** Effective October 1, 2013, a new HHS policy on subaccounts requires the CDC setup payment subaccounts within the Payment Management System (PMS) for all grant awards. Funds awarded in support of approved activities have been obligated in a newly established subaccount in the PMS, herein identified as the "P" Account". A "P" Account is a subaccount created specifically for the purpose of tracking designated types of funding in the PMS.

Funds must be used in support of approved activities in the FOA and the approved application. All award funds must be tracked and reported separately.

The grant document number and subaccount title (below) must be known in order to draw down funds from this P Account.

Grant Document Number: **000001DD16**  
Subaccount Title **DD16-1605-ZIKACOAG16**

**Acceptance of the Terms of an Award:** By drawing or otherwise obtaining funds from the grant Payment Management Services, the grantee acknowledges acceptance of the terms and conditions of the award and is obligated to perform in accordance with the requirements of the award. If the recipient cannot accept the terms, the recipient should notify the Grants Management Officer within thirty (30) days of receipt of this award notice.

**Certification Statement:** By drawing down funds, the grantee certifies that proper financial management controls and accounting systems, to include personnel policies and procedures, have been established to adequately administer Federal awards and funds drawn down. Recipients must comply with all terms and conditions outlined in their NoA, including grant policy terms and conditions contained in applicable HHS Grant Policy Statements, and requirements imposed by program statutes and regulations and HHS grants administration regulations, as applicable; as well as any regulations or limitations in any applicable appropriations acts.

## **CLOSEOUT REQUIREMENTS**

Grantees must submit closeout reports in a timely manner. Unless the Grants Management Specialist/Grants Management Officer (GMS/GMO) approves a deadline extension the grantee must submit all closeout reports within 90 days after the last day of the final budget period. Reporting timeframe is **August 01, 2016** through **July 31, 2021**. Failure to submit timely and accurate final reports may affect future funding to the organization or awards under the direction of the same Project Director/Principal Investigator (PD/PI).

All manuscripts published as a result of the work supported in part or whole by the cooperative grant must be submitted with the progress reports.

An original plus two copies of the reports must be mailed to the GMS for approval by the GMO by the due date noted. Ensure the Award and Program Announcement numbers shown above are on the reports.

The final and other programmatic reports required by the terms and conditions of the NoA are the following.

**Final Performance Report:** An original and two copies are required. At a minimum, the report should include the following:

- Statement of progress made toward the achievement of originally stated aims.
- Description of results (positive or negative) considered significant.
- List of publications resulting from the project, with plans, if any, for further publication.



**Final Federal Financial Report (FFR, SF-425):** The FFR should only include those funds authorized and actually expended during the timeframe covered by the report. The Final FFR, SF-425 is required and must be submitted to your grants management specialist (GMS) no later than 90 days after the end of budget period. This report must indicate the exact balance of unobligated funds and may not reflect any unliquidated obligations. Should the amount not match with the final expenditures reported to the Department of Health and Human Services' Payment Management Services (PMS), you will be required to update your reports to PMS accordingly. Remaining unobligated funds will be de-obligated and returned to the U.S. Treasury.

If the final reports (FFR and Final Progress Report) cannot be submitted within 90 days after the end of the project period, in accordance with 45 CFR Part 75.381 (Closeout), the grantee must submit a letter requesting an extension that includes the justification for the delay and state the expected date the CDC Procurement and Grants Office will receive the reports. All required documents must be mailed to the business contact identified in Staff Contacts.

**Equipment Inventory Report:** An original and two copies of a complete inventory must be submitted for all major equipment acquired or furnished under this project with a unit acquisition cost of \$5,000 or more. The inventory list must include the description of the item, manufacturer serial and/or identification number, acquisition date and cost, percentage of Federal funds used in the acquisition of the item. The grantee should also identify each item of equipment that it wishes to retain for continued use in accordance with 45 CFR Part 75. These requirements do apply to equipment purchased with non-federal funds for this program. The awarding agency may exercise its rights to require the transfer of equipment purchased under the assistance award referenced in the cover letter. CDC will notify the grantee if transfer to title will be required and provide disposition instruction on all major equipment. Equipment with a unit acquisition cost of less than \$5,000 that is no longer to be used in projects or programs currently or previously sponsored by the Federal Government may be retained, sold, or otherwise disposed of, with no further obligation to the Federal Government. If no equipment was acquired under this award, a negative report is required.

**Final Invention Statement:** An original and two copies of a Final Invention Statement are required. Electronic versions of the form can be downloaded by visiting <http://grants1.nih.gov/grants/hhs568.pdf>. If no inventions were conceived under this assistance award, a negative report is required. This statement may be included in a cover letter.

## **CDC ROLES AND RESPONSIBILITIES**

**Roles and Responsibilities:** Grants Management Specialists/Officers (GMO/GMS) and Program/Project Officers (PO) work together to award and manage CDC grants and cooperative agreements. From the pre-planning stage to closeout of an award, grants management and program staff have specific roles and responsibilities for each phase of the grant cycle. The GMS/GMO is responsible for the business management and administrative functions. The PO is responsible for the programmatic, scientific, and/or technical aspects. The purpose of this factsheet is to distinguish between the roles and responsibilities of the GMO/GMS and the PO to provide a description of their respective duties.

**Grants Management Officer:** The GMO is the federal official responsible for the business and other non-programmatic aspects of grant awards including:

- Determining the appropriate award instrument, i.e.; grant or cooperative agreement
- Determining if an application meets the requirements of the FOA
- Ensuring objective reviews are conducted in an above-the-board manner and according to guidelines set forth in grants policy
- Ensuring grantee compliance with applicable laws, regulations, and policies
- Negotiating awards, including budgets
- Responding to grantee inquiries regarding the business and administrative aspects of an award
- Providing grantees with guidance on the closeout process and administering the closeout of grants
- Receiving and processing reports and prior approval requests such as changes in funding, carryover, budget redirection, or changes to the terms and conditions of an award
- Maintaining the official grant file and program book

The GMO is the only official authorized to obligate federal funds and is responsible for signing the NoA, including revisions to the NoA that change the terms and conditions. The GMO serves as the counterpart to the business officer of the recipient organization.

**GMO Contact:** See Staff Contacts below for the assigned GMO

**Grants Management Specialist:** The GMS is the federal staff member responsible for the day-to-day management of grants and cooperative agreements. The GMS is the primary contact of recipients for business and administrative matters pertinent to grant awards. Many of the functions described above are performed by the GMS on behalf of the GMO.

**GMS Contact:** See Staff Contacts below for the assigned GMS

**Program/Project Officer:** The PO is the federal official responsible for the programmatic, scientific, and/or technical aspects of grants and cooperative agreements including:

- The development of programs and FOAs to meet the CDC's mission
- Providing technical assistance to applicants in developing their applications e.g. explanation of programmatic requirements, regulations, evaluation criteria, and guidance to applicants on possible linkages with other resources
- Providing technical assistance to grantees in the performance of their project
- Post-award monitoring of grantee performance such as review of progress reports, review of prior approval requests, conducting site visits, and other activities complementary to those of the GMO/GMS

### **Programmatic Contact**

William Paradies

#### **Project Officer**

Centers for Disease Control and Prevention  
National Center on Birth Defects and Developmental Disabilities (NCBDDD)  
4770 Buford Highway, NE Mailstop E-86  
Atlanta, GA 30341  
Telephone: (404) 498-3919  
Email: [wep2@cdc.gov](mailto:wep2@cdc.gov)

### **Business and Grants Policy Contact**

Patricia French

#### **Grants Management Officer (GMO)**

Centers for Disease Control  
2920 Brandywine Rd.  
Atlanta, GA 30341  
Telephone: 770-488-2849  
Email: [pff6@cdc.gov](mailto:pff6@cdc.gov)

Natasha Grant

#### **Grants Management Specialist (GMS)**

Centers for Disease Control  
2920 Brandywine Rd.  
Atlanta, GA 30341  
Telephone: 770-488-1649  
Email: [ngrant@cdc.gov](mailto:ngrant@cdc.gov)



**TECHNICAL REVIEW/SUMMARY STATEMENT  
FOA #CDC-RFA-DD16-1605**

**Surveillance, intervention, and referral to services activities for infants with microcephaly  
or other adverse outcomes linked with the Zika virus**

<b><u>APPLICANT:</u></b>	Missouri Department of Health and Senior Services
<b><u>APPLICATION NO.:</u></b>	NU50DD2016000032
<b><u>APPLICATION DATED:</u></b>	6/28/2016
<b><u>RECOMMENDATION:</u></b>	Approved
<b><u>YR 1 FUNDING REQUESTED:</u></b>	\$400,000
<b><u>SCORE:</u></b>	67

**Jorge Rosenthal / Cara Mai**  
Technical Reviewer

**7/27/2016**

**SUMMARY OF PROJECT:**

Currently the state of Missouri has a passive birth defects surveillance system. This results in a 2-year lag time for identifying birth defect cases. The Missouri Zika Surveillance, Intervention, and Referral to Services (MOZSIRS) project aims at establishing an active, real-time population-based surveillance system that collects and disseminates valid and reliable data on microcephaly and adverse pregnancy outcomes with a possible link to maternal Zika virus infection in pregnancy. The surveillance and data utilization activities will focus on women of childbearing age, pregnant women, and infants with possible or diagnosed microcephaly or other selected CNS defects associated with the Zika virus in the state.

The MOZSIRS will build collaborations among Missouri public health agencies and other relevant stakeholders particularly with regard to dissemination of information in a timely manner, and in accordance with CDC guidelines. The surveillance system will collect standardized data from all Missouri hospitals and the State Public Health Laboratory (SPHL) and will link and evaluate health and developmental outcomes for children with microcephaly and select CNS conditions with a possible link to maternal Zika virus. Collected information will be shared with all the 115 local public health agencies (LPHAs) in Missouri and collaborating public health agencies and programs as well with CDC.

**OBJECTIVES:**

- Establish and implement a rapid (real-time) population-based surveillance of microcephaly and select CNS defects (live births, stillbirths and terminations).
- To build surveillance capacity through collaboration with internal and external partners
- To provide referral services to those children diagnosed with microcephaly and other CNS defects to health programs and social services
- To establish a centralized pooled clinical case and surveillance system meeting deadlines specified by CDC

- To examine health and development outcomes of selected children born to women with positive or inconclusive Zika virus test

### **METHODS:**

- The applicant proposed to establish a rapid population-based surveillance of microcephaly and selected CNS defects using an active case-finding methodology and the standard case definition
- Target population: Women of childbearing-age (15-44 yrs), pregnant women and infants with possible or diagnosed microcephaly or other selected CNS defects associated with the Zika virus in the state of Missouri
- Case ascertainment of any potential/possible cases of microcephaly and selected CNS defects for “all” pregnancy outcomes (live births, stillbirths and terminations).
- Positive or inconclusive maternal Zika virus tests will be linked with the records of these pregnant women
- Bureau of Vital Statistics will “finalize” agreements with hospitals, prenatal care clinics and the laboratory to obtain data
- Collaboration with CDC funded and non-funded projects:
  - BVS and Zika Pregnancy Registry (“creates” active surveillance) through electronic records sharing
  - Bureau of Health care Analysis and Data Dissemination to link patient abstracted data to birth certificate data
  - Bureau of Vital Records will modify the birth certificate to capture microcephaly and Zika infection (addition of check boxes for both)

### **1. Evaluation Criteria: Approach (50 points)**

#### **STRENGTHS:**

- Applicant proposes to move from a passive birth defects surveillance system to an active, “real-time” population-based surveillance system that will ascertain defects among all pregnancy outcomes
  - Baseline data on the prevalence of microcephaly in the state is reported
- Applicant will reach agreements and coordinate activities with different CDC and non-CDC funded projects to achieve real-time data exchange
- Applicant will use data to provide information and awareness to the target population and health services providers
- Applicant will refer infants diagnosed with microcephaly and other CNS birth defects to relevant health, developmental and social services.

#### **WEAKNESSES/RECOMMENDATIONS:**

- Applicant appears to have a primary focus on Zika positive cases of women and their affected infants, as opposed to cases of infants with birth defects, e.g., microcephaly and other CNS conditions, of Zika response interest. It is important for the applicant to be sure to include rapid, real-time response of infants with birth defects that might be associated with

Zika virus.

- Case ascertainment: It is not clear from the proposal how the applicant will assure valid and complete case identification.
- Active surveillance: Applicant does not provide or explain the mechanisms by which the surveillance system will be active. It appears that the use of the checkbox for microcephaly will be how the applicant identifies infants (from the birth certificate).
- Applicant provides no indication of how non-live births will be ascertained.
- Case type and reporting: Differentiation between Zika virus linked and non-linked cases. The applicant does not differentiate how to discern between these cases and how it will count and report each of them on a timely basis. Their proposed use of the birth certificate as one way to obtain the data is not optimal.
- Linkage to patients’ records, and provider training are not clearly explained.

**Recommended Score for #1 – Approach:        \_30\_**

<b>Poor</b>	<b>Fair</b>	<b>Good</b>	<b>Very Good</b>	<b>Outstanding</b>
<b>0-24</b>	<b>25-34</b>	<b>35-39</b>	<b>40-44</b>	<b>45-50</b>

**2. Evaluation Criteria: Evaluation and Performance Measurement (25 points)**

**STRENGTHS:**

- Applicant presents a detailed work plan with monthly targets
- Applicant presents short, medium and long term outcomes and activities
- Applicant proposes a set of questions to address the evaluation of the performance measurement – both activity evaluation and outcome evaluation are proposed

**WEAKNESSES/RECOMMENDATIONS:**

- Applicant’s year 1 work plan does not include many measureable performance measures and some key terms are left undefined (e.g. “timely data for Zika virus infection” – this is not defined)

**Recommended Score for #2 – Evaluation and Performance Measurement:        \_20\_**

<b>Poor</b>	<b>Fair</b>	<b>Good</b>	<b>Very Good</b>	<b>Outstanding</b>
<b>0-11</b>	<b>12-16</b>	<b>17-19</b>	<b>20-22</b>	<b>23-25</b>

**3. Evaluation Criteria: Applicant's Organizational Capacity to Implement the Approach (25 points)**

**STRENGTHS:**

- Current staff have experience with passive birth defects surveillance
- Applicant included the hiring of additional staff for data analysis and management and to train providers and others
- Applicant provided letters of support

**WEAKNESSES/RECOMMENDATIONS**

- No clear evidence of current or proposed staff capacity to conduct active birth defects surveillance
- 2.5 FTEs are currently to be determined

**Recommended Score for #3 – Applicant's Organizational Capacity to Implement the Approach:     \_\_17\_\_**

<b>Poor</b>	<b>Fair</b>	<b>Good</b>	<b>Very Good</b>	<b>Outstanding</b>
<b>0-11</b>	<b>12-16</b>	<b>17-19</b>	<b>20-22</b>	<b>23-25</b>

**4. Evaluation Criteria: Budget Justification and Adequacy of Facilities (not scored)**

The budget will be evaluated for the extent to which it is reasonable, clearly justified, and consistent with the intended use of the cooperative agreement funds. The applicant shall describe and indicate the availability of facilities and equipment necessary to carry out this project. Award recipients agree to use cooperative agreement funds for travel by project staff agreed-upon by CDC to participate in CDC-sponsored workshops or other called meetings.

**COMMENTS:**

- We verified the accuracy, allowability, and reasonableness of the proposed cost.
- The grantee will need to submit a revised budget with their response to the Summary Statement weaknesses and recommendations.
- Personnel: Applicant has a vacant Health Program Representative position proposed at 100% for 12 months. If the position is not anticipated to be filled by start date 8/1/2016, the applicant should submit a revised budget reflecting the anticipated start date after award.
- Supplies: Breakdown of supplies, estimated quantities and item costs.
- Contractual: Applicant should provide the following Contractual elements: 1) Name of Contractor; 2) Method of Selection; 3) Period of Performance; 4) Scope of Work; 5) Method of Accountability and 6) Itemized Budget and Justification.