

burden for the focus group is estimated to be ~55 hours (3,280 minutes) total. Including both telephone interviews and focus group sessions, the total new

burden for this revision request will be an additional ~68 hours (321 individuals) at \$4,421 total, compared with the original OMB approved burden

of 1,570 hours (4,435 individuals) at \$97,460 total. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Pathologist	IHC telephone interview	20	1	20/60	7
	IHC telephone interview—contacted	27	1	1/60	0.45
	IHC focus group	12	1	1.50	18
	IHC focus group—invitation	100	1	5/60	8
	IHC focus group—consent form	12	1	5/60	1
Laboratory Directors	IHC telephone interview	10	1	20/60	3
	IHC telephone interview—contacted	27	1	1/60	0.45
	IHC focus group	6	1	1.50	9
	IHC focus group—invitation	50	1	5/60	4
Laboratory Managers	IHC focus group—consent form	6	1	5/60	0.50
	IHC telephone interview	10	1	20/60	3
	IHC telephone interview—contacted	27	1	1/60	0.45
	IHC focus group	6	1	1.50	9
	IHC focus group—invitation	50	1	5/60	4
Total	IHC focus group—consent form	6	1	5/60	0.50
	68.00

Leroy A. Richardson,
 Chief, Information Collection Review Office,
 Office of Scientific Integrity, Office of the
 Associate Director for Science, Office of the
 Director, Centers for Disease Control and
 Prevention.

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**DEPARTMENT OF HEALTH AND
 HUMAN SERVICES**

**Centers for Disease Control and
 Prevention**

[60 Day-16-0968; Docket No. CDC-2015-0104]

**Proposed Data Collection Submitted
 for Public Comment and
 Recommendations**

AGENCY: Centers for Disease Control and
 Prevention (CDC), Department of Health
 and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease
 Control and Prevention (CDC), as part of
 its continuing efforts to reduce public
 burden and maximize the utility of
 government information, invites the
 general public and other Federal
 agencies to take this opportunity to
 comment on proposed and/or
 continuing information collections, as
 required by the Paperwork Reduction
 Act of 1995. This notice invites
 comment on a proposed information
 collection entitled “Monitoring and
 Reporting System for DELTA FOCUS

Awardees”. CDC will use the
 information collected to monitor
 cooperative agreement awardees and to
 identify challenges to program
 implementation and achievement of
 outcomes.

DATES: Written comments must be
 received on or before January 25, 2016.

ADDRESSES: You may submit comments,
 identified by Docket No. CDC-2015-
 0104 by any of the following methods:

Federal eRulemaking Portal:
 Regulation.gov. Follow the instructions
 for submitting comments.

Mail: Leroy A. Richardson,
 Information Collection Review Office,
 Centers for Disease Control and
 Prevention, 1600 Clifton Road NE., MS-
 D74, Atlanta, Georgia 30329.

Instructions: All submissions received
 must include the agency name and
 Docket Number. All relevant comments
 received will be posted without change
 to Regulations.gov, including any
 personal information provided. For
 access to the docket to read background
 documents or comments received, go to
 Regulations.gov.

Please note: All public comment should be
 submitted through the Federal eRulemaking
 portal (Regulations.gov) or by U.S. mail to the
 address listed above.

FOR FURTHER INFORMATION CONTACT: To
 request more information on the
 proposed project or to obtain a copy of
 the information collection plan and
 instruments, contact the Information
 Collection Review Office, Centers for
 Disease Control and Prevention, 1600

Clifton Road NE., MS-D74, Atlanta,
 Georgia 30329; phone: 404-639-7570;
 Email: omb@cdc.gov.

SUPPLEMENTARY INFORMATION: Under the
 Paperwork Reduction Act of 1995 (PRA)
 (44 U.S.C. 3501-3520), Federal agencies
 must obtain approval from the Office of
 Management and Budget (OMB) for each
 collection of information they conduct
 or sponsor. In addition, the PRA also
 requires Federal agencies to provide a
 60-day notice in the **Federal Register**
 concerning each proposed collection of
 information, including each new
 proposed collection, each proposed
 extension of existing collection of
 information, and each reinstatement of
 previously approved information
 collection before submitting the
 collection to OMB for approval. To
 comply with this requirement, we are
 publishing this notice of a proposed
 data collection as described below.

Comments are invited on: (a) Whether
 the proposed collection of information
 is necessary for the proper performance
 of the functions of the agency, including
 whether the information shall have
 practical utility; (b) the accuracy of the
 agency’s estimate of the burden of the
 proposed collection of information; (c)
 ways to enhance the quality, utility, and
 clarity of the information to be
 collected; (d) ways to minimize the
 burden of the collection of information
 on respondents, including through the
 use of automated collection techniques
 or other forms of information
 technology; and (e) estimates of capital
 or start-up costs and costs of operation,

maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information.

Proposed Project

Monitoring and Reporting System for DELTA FOCUS Awardees, (OMB Control No. 0920–0968, expiration 5/31/2016)—Extension—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Intimate Partner Violence (IPV) is a serious, preventable public health problem that affects millions of Americans and results in serious consequences for victims, families, and communities. IPV occurs between two people in a close relationship. The term

“intimate partner” describes physical, sexual, or psychological harm by a current or former partner or spouse. IPV can impact health in many ways, including long-term health problems, emotional impacts, and links to negative health behaviors. Given these factors, the Family Violence Prevention and Services Act (42 U.S.C. 10401) provides an important opportunity for the advancement of public health and reduction of IPV. Support and guidance for programs addressing IPV have been provided through cooperative agreement funding and technical assistance administered by CDC’s National Center for Injury Prevention and Control (NCIPC). CDC seeks to continue collecting information needed to monitor cooperative agreement programs funded under Domestic Violence Prevention Enhancement and Leadership through Alliances, Focusing on Outcomes for Communities United with States DELTA FOCUS (FOA CDC–RFA–CE13–130).

Information to be collected will provide crucial data for program performance monitoring and provide CDC with the capacity to respond in a timely manner to requests for information about the program from the Department of Health and Human Services (HHS), the White House, Congress, and other sources. Awardees will report progress and activity information to CDC on an annual

schedule using the Program Management Information System (PMIS) consisting of fillable electronic templates and submitted via Grant Solutions.

CDC will use the information collected to monitor each awardee’s progress and to identify facilitators and challenges to program implementation and achievement of outcomes. Monitoring allows CDC to determine whether an awardee is meeting performance goals and to make adjustments in the type and level of technical assistance provided to them, as needed, to support attainment of their objectives. CDC’s monitoring and evaluation activities also allow CDC to provide oversight of the use of federal funds, and to identify and disseminate information about successful prevention and control strategies implemented by awardees. These functions are central to the NCIPC’s broad mission of reducing the burden of injury and violence. Finally, the information collection allows CDC to monitor the increased emphasis on partnerships and programmatic collaboration, and is expected to reduce duplication of effort, enhance program impact and maximize the use of federal funds.

This is an extension request for three years. Participation in the information collection is required as a condition of funding. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
State Domestic Violence Coalitions ..	DELTA FOCUS PMIS: Semi-annual reporting.	10	2	3	60
Total	60

Leroy A. Richardson,
Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Request for Nominations of Candidates To Serve as Members of the Community Preventive Services Task Force (CPSTF); Reopening of Nomination Period

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Disease Control and Prevention (CDC) within

the Department of Health and Human Services (HHS) announces the reopening of the nomination period for individuals qualified to serve as members of the Community Preventive Services Task Force (CPSTF). The nomination period originally closed on November 9, 2015.

DATES: Nomination packages must be received by December 8, 2015. Complete nomination packages must be submitted by the deadline in order to be considered. Individuals who submitted a nomination package during the original nomination period do not need to re-submit their nomination package to be considered.