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NCIPC Determination

General Information

Project Title

National Intimate Partner and Sexual Violence Survey - 5 (NISVS-5): Data Collection

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Project Officer(s) Asad Patwary Division: DVP Telephone: (404) 718-5190

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Proposed Project Dates:

Start: 09/26/2022 Ending: 05/31/2027

Ex: MM/DD/YYYY

Ex: MM/DD/YYYY

Funding Mechanism

- Cooperative Agreement #: _____ Funding FOA#: _____
- Grant #: _____ Funding FOA#: _____
- Contract#: 75D30122F15662
- No funding (Specify): _____

Describe the purpose, methods, and outcomes of the project (Use space provided - Abstract of purpose, methods and outcomes)

The National Intimate Partner and Sexual Violence Survey (NISVS) is an ongoing, nationally representative public health surveillance survey that collects information about experiences of sexual violence (SV), stalking, and intimate partner violence (IPV) among non-institutionalized English and Spanish speaking women and men aged 18 years or older in the United States.

A contractor (RTI) will administer the NISVS survey on behalf of CDC. This survey will be conducted over time to provide population-based national prevalence estimates of sexual violence, stalking, and intimate partner violence. The purpose of this project is to conduct a population-based survey that will provide accurate and reliable estimates of the prevalence of SV, stalking, and IPV at the national and state levels.

Outcomes of the project are to provide detailed information regarding the magnitude and characteristics of sexual violence, stalking, and intimate partner violence at the national and state levels. Data points will include lifetime and 12-month prevalence Respondent characteristics will include sex, gender identity, sexual orientation, and race/ethnicity. Data will provide critical information about experiences with SV, stalking, and IPV prevalence characteristics, including circumstances and context in which the violence occurred, the victim-perpetrator relationship, and the respondent's age when the violence first occurred. Data from this project will provide ongoing public health surveillance and monitoring of SV, stalking, and IPV and inform efforts to prevent these forms of violence.

Describe the roles and responsibilities of CDC and any partner organizations (e.g., grantee, contractor).

CDC will oversee the project and review and approve: (1) materials used in data collection; (2) data analyses; (3) sampling procedures; (4) training procedures. CDC will not have access to personally-identifiable information.

RTI (contractor) will carry out the sampling and recruitment, design the web survey program, conduct data collection, train phone interviewers, clean and de-identify the data, conduct data, and analyses.

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National Intimate Partner and Sexual Violence Survey - 5 (NISVS-5)

Proposed Project Dates:

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Applicability of Human Subjects Regulations

Please check appropriate category:

I. Activity is not research. Primary intent is public health practice: disease/injury control, surveillance, improvement of programs or services. Objectives focused on a specific population.

- A.** Epidemic/endemic **disease/injury control** activity; collected data directly relate to *immediate* disease control needs (e.g., epi-aid).
- B.** Routine **disease/injury surveillance** activity; data used for disease control program or policy purposes for a specific health condition/disease in a specific population and setting. (Includes disease reporting)
- C.** **Program evaluation/monitoring** activity; data are used primarily for assessing, monitoring or improving a program, policy, or a communications activity (e.g., message testing) in a specific population/setting.
- D.** Purchase orders or contracts for services or equipment.

-OR-

II. Activity is research but does NOT involve human subjects. Primary intent is to develop or contribute to generalizable knowledge, but data is obtained solely from non-human sources or not living individuals, or anonymous existing data collected for another purpose are being analyzed:

- A.** Activity is research involving collection/analysis of data about health facilities or other organizations or units, which are not individual persons
- B.** Activity is research using existing unlinked or anonymous data previously collected for another purpose.
- C.** Activity is research involving data and/or specimens from *deceased persons*.

-OR-

III. Activity is research involving human subjects but CDC is not engaged. CDC employees including visiting scientists, fellows, and on-site contractors (but not off-site contractors or other collaborators)
*Will NOT obtain consent or data by intervening or interacting with participants
*Will NOT have access to identifiable (including coded) private data or biological specimens

NOTE: Once local IRB approval has been obtained please forward a copy (electronic preferred) to the NCIPC Human Subjects Coordinator for records keeping purposes.

-OR-

IV. Activity is research involving human subjects and could be exempt according to the categories specified in the regulations 45 CFR 46.101(b). Educational practices, Educational tests, surveys, interviews, or observation of public behavior. Existing data, documents, records (e.g., not identifiable, publicly available). Demonstration projects. This will require final determination from CDC-HRPO.

-OR-

V. Activity is research involving human subjects, CDC is engaged, and CDC IRB approval will be sought.

Required Signatures

Kristin

Holland -S

Digitally signed by
Kristin Holland -S
Date: 2022.10.31
14:29:23 -04'00'

10/31/2022

Branch/Team Official (e.g., Branch chief or Team Lead)

Date

Kathleen C.

Basile -S

Digitally signed by
Kathleen C. Basile -S
Date: 2022.10.31
14:41:08 -04'00'

10/31/2022

Division Official (e.g., ADS, Director)

Date

Karen C.

Angel -S

Digitally signed by
Karen C. Angel -S
Date: 2022.11.01
08:31:18 -04'00'

11/01/2022

Human Subjects Coordinator

Date

Office Use Only

NCIPC Determination

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National Intimate Partner and Sexual Violence Survey - 5 (NISVS-5)

Proposed Project Dates:

Start: 09/26/2022 Ending: 05/31/2027

Applicability of OMB-PRA Regulations

Please check appropriate category:

I. Does the activity involve collecting identical information from 10 or more respondents within a one year period

Including evaluation/monitoring (Examples of collections: Surveys & Interviews (Phone & On-line) - Focus groups – Surveillance – Program evaluation- Program monitoring).

No

Yes - Type of collection:

<input checked="" type="checkbox"/> Mail-backquestionnaire	<input type="checkbox"/> On-site questionnaire	<input type="checkbox"/> Personal interview	<input checked="" type="checkbox"/> Telephone survey
<input type="checkbox"/> Testing/assessment form	<input checked="" type="checkbox"/> Web-based survey	<input type="checkbox"/> Focus groups	Record abstractions
<input type="checkbox"/> Evaluation	<input type="checkbox"/> Observation	<input type="checkbox"/> Workshop	Discussion group
<input type="checkbox"/> Comment card			
<input type="checkbox"/> Other (Explain) _____			

II. Is NCIPC Sponsoring the data collection? Check all that apply.

- A.** NCIPC will initiate or request a data collection.
- B.** NCIPC will develop or design the data collection.
- C.** NCIPC will manage or own the data collection.
- D.** NCIPC will be directing the data collection.
- E.** NCIPC staff will interact/intervene participants.
- F.** NCIPC is requesting specific data reports.
- G.** NCIPC will disseminate the data as an official report or study.
- H.** NCIPC is not conducting any of the above activities and therefore is not sponsoring the data collection.