

## Attachment D

### Rapid Response Suicide Investigation Data Collection: Generic Information Collection (GenIC) Investigation Protocol Template

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#### **TITLE**

Provide a title for the investigation.

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#### **CDC STAFF LEADING INVESTIGATION**

List the CDC staff leading the investigation.

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#### **INTRODUCTION**

Describe the circumstances of the investigation, including the geographic location, any vulnerable population, and initiating event (e.g., partner requesting CDC assistance, CDC's suicide monitoring that detected area or population with possible cluster or elevated suicide rates).

Specify which circumstances justify the Rapid Response Suicide Investigation Data Collection. Examples of circumstances, include, but are not limited to:

- Increased rate in suicidal behavior or suicide
  - Specific description of vulnerable subpopulation(s)
  - Possible or known suicide cluster or contagion
  - Need for new information about trends, risk/protective factors, vulnerable populations (e.g., adolescents), and other topics (e.g., common methods/location) in suicidal behavior or suicides to inform prevention strategies
  - Public or political concern (e.g., possible suicide cluster or increasing trends in a state, county, community or vulnerable population)
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#### **PURPOSE**

Describe the objectives of the investigation. Specify the external partners (e.g., local, state, territory, and tribal health authorities; other federal agencies; local and state leaders; schools; or other partner organizations) involved with the urgent response. Describe the type of CDC technical assistance requested/provided. Describe the purpose of the data collection activities.

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**METHODS**

Describe the proposed data collection methods.

Investigation and analytic objectives:

Investigation design:

Case definition(s):

Investigation population:

Sample design (if appropriate):

Data collection approach:

Power/sample size calculations/minimum detectable differences (if appropriate):

Respondents:

Anticipated burden hours:

**BURDEN ESTIMATE**

Data Collection Instrument Name	Type of Respondent	Data Collection Mode	No. Respondents (A)	No. Responses per Respondent (B)	Burden per Response in Minutes (C)	Total Burden in Hours (A x B x C)/60*

Data analysis plan:

Intended application of findings:

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**RESULTS AND DISSEMINATION**

Describe how results will be synthesized and reported to the external partner.

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## **INVESTIGATION CONSIDERATIONS**

Explain why the Privacy Act does or does not apply. If applicable, describe steps being taken ensure privacy of respondent data?:

Describe whether or not the investigation requires Human Subjects Protections. Attach NCIPC Determination of Human Subjects review?:

Are incentives provided? If yes, provide a justification for incentive plan.:

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## **INVESTIGATIVE TEAM**

List full investigative team, including CDC staff and external partner staff.

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## **CITATIONS**

Provide references for works cited.

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## **ATTACHMENTS**

Provide the draft data collection forms to be used in the investigation; specify respondents for each form.