

**THE PERFORMANCE MEASURES PROJECT: IMPROVING PERFORMANCE MEASUREMENT
AND MONITORING BY CDC PROGRAMS**

PART B: STATISTICAL METHODS

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List of Attachments

1. Sample Recipient Technical Specifications Codebook and Data Reporting Guide

B. Collections of Information Employing Statistical Methods

1. Respondent Universe and Sampling Methods

CDC/ATSDR programs are eligible to request OMB approval through the Performance Measures Project (“PMP”) generic if they have collaborated with CDC’s Performance and Evaluation Office (PEO) to develop their Notice of Funding Opportunity (NOFO) or other funding announcement and have developed a core set of priority performance measures and data reporting plans for recipients. Collaboration with PEO ensures that appropriate performance and evaluation principles are incorporated into the activities and reporting plans for these awards. PEO anticipates that the majority of programmatic activities will be funded through the cooperative agreement mechanism but eligibility may also include activities funded through grants or contracts. Eligibility is limited to non-research activities but includes qualifying activities conducted in domestic or international settings.

Information will be collected from the recipients of qualifying awards. Recipients (“respondents”) will typically be units of state or local government (e.g., state or local health departments) but may include other public health partners (e.g., private sector organizations or ministries of health).

Performance monitoring is required. No statistical sampling method will be used.

Sixteen GENICs were approved in the previous approval period (June 2023 – June 2026) and CDC/ATSDR requests OMB approval to continue these information collections. In addition, CDC/ATSDR anticipates that up to 40 new programs will be phased in to the PMP over the next three years.

2. Procedures for the Collection of Information

The CDC/ATSDR program funding a particular public health initiative will collect information from each recipient on the schedule required by the funding mechanism and documented in the Program’s Data Management Plan. Recipients will report progress on their performance measures using a cooperative agreement-specific information collection adapted from a Sample Recipient Technical Specifications Codebook and a Recipient Data Reporting Guide (**Attachment 2**). Instructions will be provided to recipients for completing the templates.

3. Methods to Maximize Response Rates and Deal with Nonresponse

Periodic reporting based on requirements stated in the funding mechanism is required of all recipients. Response rates are expected to be 100%.

4. Test of Procedures or Methods to be Undertaken

No testing of procedures or methods will be undertaken for this Revision ICR. The request is based on CDC’s experience with the PMP generic. All recipients receiving funding from a specific NOFO will use the NOFO-specific tool.

5. Individuals Consulted on Statistical Aspects and Individuals Collecting and/or Analyzing Data

The individuals responsible for design and management of the information collection include:

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