



**U.S. Department of
Health and Human Services**
Centers for Disease
Control and Prevention

Print Date: 5/21/26

Title: Performance Measures Project

Project Id: 0900f3eb826b2025

Accession #: OPPE-EEU-5/18/26-b2025

Project Contact: Clay Cooksey

Organization: OPPE/PEO/PEOB/EEU

Status: **Project In Progress**

Intended Use: **Project Determination**

Estimated Start Date: 07/01/2026

Estimated Completion Date: 06/29/2029

CDC/ATSDR HRPO/IRB Protocol #:

OMB Control #: 0920-1282

OMB Discontinuation Date:

Determinations

Determination	Justification	Completed	Entered By & Role
HSC: Does NOT Require HRPO Review	Not Research / Other <i>45 CFR 46.102(l)</i> Program Evaluation Quality Assurance / Improvement	5/20/26	Chirila_Robert (uno4) CIO HSC
PRA: PRA Applies		5/20/26	Zirger_Jeffrey (wtj5) OMB / PRA
ICRO: PRA Applies	OMB Approval date: 6/13/23 OMB Expiration date: 6/30/26	5/20/26	Zirger_Jeffrey (wtj5) ICRO Reviewer

Description & Funding

Description

Priority:	Higher Priority
Date Needed:	05/27/2026
Priority Justification:	Current GenIC expires June 30,2026. ICRO needs research determination for processing new GenIC package
CDC Priority Area for this Project:	Other CDC Priority - Performance and Evaluation
Determination Start Date:	05/19/26
Description:	Generic Information Collection for NOFO program performance measures
IMS/CIO/Epi-Aid/Lab-Aid/Chemical Exposure Submission:	No
IMS Activation Name:	Not selected
Submitted through IMS Clearance Matrix:	Not selected
Primary Scientific Priority:	Not selected
Secondary Scientific Priority (s):	Not selected
Task Force Responsible:	Not selected
CIO Emergency Response Name:	Not selected
Epi-Aid Name:	Not selected
Lab-Aid Name:	Not selected

Assessment of Chemical Exposure Name:	Not selected
Goals/Purpose	To provide a flexible and timely approval framework that facilitates the collection of information needed to measure and improve the performance of recipients of CDC/ATSDR funding. Through participation in the Performance Measures Project (PMP), CDC/ATSDR programs and recipients will: 1) Develop strong performance measurement tools and practices; 2) define and operationalize priority performance measures tailored to a specific public health initiative; and 3) establish common data collection and reporting expectations across all recipients.
Objective:	CDC#s Performance Measures Project (PMP) will work with up to 40 new CDC programs. The number of recipients per CDC /ATSDR program will vary dependent on program activity, and the expected average is approximately 35 recipients per program.
Activities or Tasks:	New Collection of Information, Data, or Biospecimens
Target Populations to be Included/Represented:	General US Population
Tags/Keywords:	Public Health
CDC's Role:	CDC employees will provide substantial technical assistance or oversight
Method Categories:	Technical Assistance
Methods:	<p>Respondents will be recipients of CDC/ATSDR funding. CDC#s Performance Measures Project (PMP) will work with up to 40 new CDC programs. The number of recipients per CDC/ATSDR program will vary dependent on program activity, and the expected average is approximately 35 recipients per program. Data will be analyzed by the CDC/ATSDR program to assess recipient progress toward the key outcomes defined by the public health initiative. The information collected is meant primarily for program improvement and accountability; it is not intended to be used as the principal basis for policy decisions.</p> <p>Participating CDC/ATSDR programs will provide the program-specific information collection forms to each recipient to facilitate standardized performance data collection at the local level. Recipients will report these standardized performance data directly to CDC/ATSDR programs on a periodic basis to assess progress toward achieving desired program outcomes as defined by the program. The default periodic basis is annual reporting, but programs may propose alternate reporting schedules if needed.</p>
Collection of Info, Data or Biospecimen:	
Expected Use of Findings/Results and their impact:	The intended use of the resulting data is to generate general purpose statistics, to assess program progress (at both the local recipient-specific and CDC/ATSDR program-wide levels), to provide data that can inform program evaluation, to assist with program planning/management, and to improve the accountability of federal funds provided to recipients.
Could Individuals potentially be identified based on Information Collected?	No
Dual Use Research potential?:	No

Funding

Funding yet to be added

HSC Review

HSC Attributes

Program Evaluation	Yes
Quality Assurance / Improvement	Yes

Regulation and Policy

Do you anticipate this project will require review by a CDC IRB or HRPO? No

Estimated number of study participants

Population - Children Protocol Page #:

Population - Minors Protocol Page #:

Population - Prisoners Protocol Page #:

Population - Pregnant Women Protocol Page #:

Population - Emancipated Minors Protocol Page #:

Suggested level of risk to subjects

Do you anticipate this project will be exempt research or non-exempt research

Requested consent process wavers

Informed consent for adults No Selection

Children capable of providing assent No Selection

Parental permission No Selection

Alteration of authorization under HIPAA Privacy Rule No Selection

Requested Waivers of Documentation of Informed Consent

Informed consent for adults	No Selection
Children capable of providing assent	No Selection
Parental permission	No Selection

Consent process shown in an understandable language

Reading level has been estimated	No Selection
Comprehension tool is provided	No Selection
Short form is provided	No Selection
Translation planned or performed	No Selection
Certified translation / translator	No Selection
Translation and back-translation to/from target language(s)	No Selection
Other method	No Selection

Clinical Trial

Involves human participants	No Selection
Assigned to an intervention	No Selection
Evaluate the effect of the intervention	No Selection
Evaluation of a health related biomedical or behavioral outcome	No Selection
Registerable clinical trial	No Selection

Other Considerations

Exception is requested to PHS informing those bested about HIV serostatus	No Selection
Human genetic testing is planned now or in the future	No Selection
Involves long-term storage of identifiable biological specimens	No Selection
Involves a drug, biologic, or device	No Selection

Conducted under an Investigational New Drug exemption or Investigational Device Exemption

No Selection

Institutions & Staff

Institutions

Will you be working with an outside Organization or Institution? No

Institutions yet to be added

Staff

Staff Member	SIQT Exp. Date	CITI Biomedical Exp. Date	CITI Social & Behavioral Exp. Date	CITI Good Clinical Practice Exp. Date	CITI Good Laboratory Practice Exp. Date	Staff Role	Email	Phone	Organization
Destiny Bruce	07/10 /2026					Project Coordinator	rln0@cdc.gov	404-639-0764	PERFORMANCE MONITORING AND ACCOUNTABILITY UNIT

Data

DMP

Proposed Data Collection Start Date: 7/1/26

Proposed Data Collection End Date: 6/29/29

Proposed Public Access Level: Non-Public

Non-Public Details:

Reason For Not Releasing Data: CIO conducting this project does not fund or own the data and is not responsible for making it available

Public Access Justification: The data is managed by the CIO, not OPPE

How Access Will Be Provided for Data: This is controlled by the CIO

Plans for Archival and Long Term Preservation: This is managed by the CIO

Spatiality

Country	State/Province	County/Region
United States		

Supporting Info

Current	CDC Staff Member and Role	Date Added	Description	Supporting Info Type	Supporting Info
	Zirger_Jeffrey (wtj5) ICRO Reviewer	05/20/2026	NOA 0920-1282 (2023)	Notice of Action	NOA 0920-1282_2023.pdf
Current	Cooksey_Clay (zwx2) Project Contact	05/19/2026	Current PMP Guidance that demonstrates what is required of participating programs	Other-Enter new type	PMP GenIC Package Guidance.docx



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