



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

*Print Date: 11/5/24*

<b>Title:</b>	NCEHATSDR Service Delivery (Generic Clearance)
<b>Project Id:</b>	0900f3eb81cc43d8
<b>Accession #:</b>	-NCEH-3/10/21-c43d8
<b>Project Contact:</b>	Yan S Ding
<b>Organization:</b>	NCEH/ATSDR
<b>Status:</b>	<b>Project In Progress</b>
<b>Intended Use:</b>	<b>Project Determination</b>
<b>Estimated Start Date:</b>	02/01/2022
<b>Estimated Completion Date:</b>	01/31/2025
<b>CDC/ATSDR HRPO/IRB Protocol #:</b>	NA
<b>OMB Control #:</b>	0923-0047

## **Determinations**

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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	8/5/24	Dignam_Timothy A. (ted9) CIO HSC
PRA: PRA Applies		8/5/24	Dignam_Timothy A. (ted9) CIO OMB / PRA
ICRO: PRA Applies	OMB Approval date: 2/17/22 OMB Expiration date: 2/28/25	8/5/24	Zirger_Jeffrey (wtj5) ICRO Reviewer

## Description & Funding

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### Description

Priority: Standard

Priority Justification:

CDC Priority Area for this Project: Not selected

Determination Start Date: 03/10/21

Description:

This is a request to publish the 60-day Federal Register Notice for ATSDR #Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery# (OMB Control No. 0923-0047; expiration date 01/31/2022). The information collection activity provides a means to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Federal government's commitment to improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study.

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**

This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

**Objective:**

The solicitation of feedback will target areas such as: timeliness, appropriateness, accuracy of information, courtesy, efficiency of service delivery, and resolution of issues with service delivery. Responses will be assessed to plan and inform efforts to improve or maintain the quality of service offered to the public. If this information is not collected, vital feedback from customers and stakeholders on the Agency's services will be unavailable.

**Does your project measure health disparities among populations/groups experiencing social, economic, geographic, and/or environmental disadvantages?:** Not Selected

**Does your project investigate underlying contributors to health inequities among populations /groups experiencing social, economic, geographic, and/or environmental disadvantages?:** Not Selected

**Does your project propose, implement, or evaluate an action to move towards eliminating health inequities?:** Not Selected

**Activities or Tasks:** New Collection of Information, Data, or Biospecimens

**Target Populations to be Included/Represented:** General US Population

**Tags/Keywords:** Feedback

**CDC's Role:**

Activity originated and designed by CDC staff, or conducted at the specific request of CDC, or CDC staff will approve study design and data collection as a condition of any funding provided ; CDC employees or agents will obtain data by intervening or interacting with participants ; CDC employees or agents will obtain or use anonymous or unlinked data or biological specimens ; CDC employees or agents will obtain or use identifiable (including coded) private data or biological specimens ; CDC employees will participate as co-authors in presentation(s) or publication(s) ; CDC employees will provide substantial technical assistance or oversight

**Method Categories:**

Discussion Group; Focus Group; Individual Interviews (Qualitative); Needs Assessment; QA/QI

ATSDR will only submit a collection for approval under this generic clearance if it meets the following conditions: # The collections are voluntary; # The collections are low-burden for respondents (based on considerations of total burden hours, total number of respondents, or burden-hours per respondent) and are low-cost for both the respondents and the Federal Government; # The collections are noncontroversial and do not raise issues of concern to other Federal agencies; # Any collection is targeted to the solicitation of opinions from respondents who have experience with the program or may have experience with the program in the

**Methods:** near future; # Personally identifiable information (PII) is collected only to the extent necessary and is not retained; # Information gathered is intended to be used only internally for general service improvement and program management purposes and is not intended for release outside of the agency (if released, the agency must indicate the qualitative nature of the information); # Information gathered will not be used for the purpose of substantially informing influential policy decisions; and # Information gathered will yield qualitative information; the collections will not be designed or expected to yield statistically reliable results or used as though the results are generalizable to the population of study.

**Collection of Info, Data or Biospecimen:** The types of collections that this generic clearance covers include, but are not limited to: # Customer comment cards/complaint forms # Small discussion groups # Focus Groups of customers, potential customers, delivery partners, or other stakeholders # Cognitive laboratory studies, such as those used to refine questions or assess usability of a website; # Qualitative customer satisfaction surveys (e.g., post-transaction surveys; opt-out web surveys) # In-person observation testing (e.g., website or software usability tests)

**Expected Use of Findings/Results and their impact:** The solicitation of feedback will target areas such as: timeliness, appropriateness, accuracy of information, courtesy, efficiency of service delivery, and resolution of issues with service delivery. Responses will be assessed to plan and inform efforts to improve or maintain the quality of service offered to the public. If this information is not collected, vital feedback from customers and stakeholders on the Agency's services will be unavailable.

**Could Individuals potentially be identified based on Information Collected?** Yes

**Will PII be captured (including coded data)?** Yes

**Does CDC have access to the identifiers (including coded data)?:** Yes

**Is this project covered by an Assurance of Confidentiality?** No

**Does this activity meet the criteria for a Certificate of Confidentiality (CoC)?** No

**Is there a formal written agreement prohibiting the release of identifiers?** No

## Funding

Funding Type	Funding Title	Funding #	Original Budget Yr	# Years Award	Budget Amount
Other-Staff time only - \$0	Staff time only - \$0				

## HSC Review

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## HSC Attributes

Program Evaluation Yes  
Quality Assurance / Improvement Yes  
Other - Service Delivery and Program Improvement Yes

## Additional Ethical Considerations

Resulting data collected are not published but will be used to improve agency products.

## Regulation and Policy

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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 wavier

Informed consent for adults No Selection

<b>Children capable of providing assent</b>	No Selection
<b>Parental permission</b>	No Selection
<b>Alteration of authorization under HIPAA Privacy Rule</b>	No Selection

### **Requested Waivers of Documentation of Informed Consent**

<b>Informed consent for adults</b>	No Selection
<b>Children capable of providing assent</b>	No Selection
<b>Parental permission</b>	No Selection

### **Consent process shown in an understandable language**

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

### **Clinical Trial**

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

### **Other Considerations**

<b>Exception is requested to PHS informing those tested about HIV serostatus</b>	No Selection
<b>Human genetic testing is planned now or in the</b>	No Selection

future

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

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### Institutions

Will you be working with an outside Organization or Institution? Yes

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
Sharon Flores	09/01/2026	06/07/2027				Co-Investigator	fil4@cdc.gov	770-488-4182	NATIONAL CENTER FOR ENVIRONMENTAL HEALTH
Yan Ding	06/26/2026	06/26/2027		12/01/2026		Principal Investigator	yad6@cdc.gov	770-488-7934	NATIONAL CENTER FOR ENVIRONMENTAL HEALTH

## Data

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### DMP

Proposed Data Collection Start Date: 2/1/22

Proposed Data Collection End Date: 1/31/25

Proposed Public Access Level: Non-Public

Non-Public Details:

**Reason For Not Releasing Data:** Other - Quality improvement for ATSDR services

**Public Access Justification:** This is a generic IC request to allow gathering of feedback on services provided by ATSDR for the purposes of quality improvement.

**How Access Will Be Provided for Data:** Data will be used by ATSDR to evaluate and improve services.

**Plans for Archival and Long Term Preservation:**

**Spatiality**

Spatiality (Geographic Locations) yet to be added .....

**Dataset**

Dataset Title	Dataset Description	Data Publisher /Owner	Public Access Level	Public Access Justification	External Access URL	Download URL	Type of Data Released	Collection Start Date	Collection End Date
Dataset yet to be added...									

**Supporting Info**



Current	CDC Staff Member and Role	Date Added	Description	Supporting Info Type	Supporting Info
	Zirger_Jeffrey (wtj5) ICRO Reviewer	08/05/2024	NOA 0923-0047 (2022)	Notice of Action	NOA 0923-0047_2022.pdf
Current	Ding_Yan (Shirley) (yad6) Project Contact	08/05/2024	60D FRN ATSDR	Paperwork Reduction Act Form	0 - 60-day_FRN_ATSDR_0923-0047 Generic Service Delivery.docx
Current	Ding_Yan (Shirley) (yad6) Project Contact	08/05/2024	60D FRN request	Paperwork Reduction Act Form	0 - 60-day FRN Request 0923-0047 Generic Service Delivery.docx
	Davis_Stephanie I. (sgd8) CIO OMB / PRA	03/11/2021	60D Federal Register Notice package for OMB Control No. 0923-0047; expiration date 01/31 /2022	Other	0923-0047 Final 60 day 2021 Extension to ICRO.zip
	Abel_Jason A. (jza5) CIO HSC	03/11/2021	Updated 60 day package	Other	0923-0047 60 day 2021 Extension Updated per OS review. zip
	Davis_Stephanie I. (sgd8) CIO HSC	03/11/2021	Zipfile of 60-day Files with comments and edits	Other	0923-0047 60D 2021 Extnsn rev OS.zip
	Abel_Jason A. (jza5) Project Contact	03/10/2021	60 day PRA package	Other-60 day PRA package	0923-0047 60 day 2021 Extension.zip



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