

with the organization’s safety and soundness. There is no change in the substance or methodology of this information collection. The change in burden is due to a decrease in the estimated number of respondents. The burden hours decreased by 358 from 4,368 to 4,010.

Request for Comment

Comments are invited on: (a) Whether the collection of information is necessary for the proper performance of the FDIC’s functions, including whether the information has practical utility; (b) the accuracy of the estimates of the burden of the information collection, including the validity of the methodology and assumptions used; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques

or other forms of information technology. All comments will become a matter of public record.

Federal Deposit Insurance Corporation.
Dated at Washington, DC, on April 28, 2023.

James P. Sheesley,
Assistant Executive Secretary.
[FR Doc. 2023–09529 Filed 5–3–23; 8:45 am]
BILLING CODE 6714–01–P

FEDERAL DEPOSIT INSURANCE CORPORATION

Update to Notice of Financial Institutions for Which the Federal Deposit Insurance Corporation Has Been Appointed Either Receiver, Liquidator, or Manager

AGENCY: Federal Deposit Insurance Corporation.
ACTION: Update listing of financial institutions in liquidation.

SUMMARY: Notice is hereby given that the Federal Deposit Insurance Corporation (Corporation) has been appointed the sole receiver for the following financial institution effective as of the Date Closed as indicated in the listing.

SUPPLEMENTARY INFORMATION: This list (as updated from time to time in the **Federal Register**) may be relied upon as “of record” notice that the Corporation has been appointed receiver for purposes of the statement of policy published in the July 2, 1992, issue of the **Federal Register** (57 FR 29491). For further information concerning the identification of any institutions which have been placed in liquidation, please visit the Corporation website at www.fdic.gov/bank/individual/failed/banklist.html, or contact the Chief, Receivership Oversight at RO@fdic.gov or at Division of Resolutions and Receiverships, FDIC, 600 North Pearl Street, Suite 700, Dallas, TX 75201.

INSTITUTIONS IN LIQUIDATION
[In alphabetical order]

FDIC ref. No.	Bank name	City	State	Date closed
10543	First Republic Bank	San Francisco	CA	05/01/2023

Federal Deposit Insurance Corporation.
Dated at Washington, DC, on May 1, 2023.
James P. Sheesley,
Assistant Executive Secretary.
[FR Doc. 2023–09528 Filed 5–3–23; 8:45 am]
BILLING CODE 6714–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–23–1027]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled “Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery” to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on February 6, 2023, to obtain comments from the public and affected agencies. CDC received one comment related to the

previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.
CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:
(a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
(c) Enhance the quality, utility, and clarity of the information to be collected;
(d) Minimize the burden of the collection of information on those who are to respond, including, through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and
(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570. Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.

Proposed Project

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (OMB Control No. 0920–1027, Exp. 8/31/2023)—Extension—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC is requesting a three-year Extension for the Generic ICR titled Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery. During the past three-year approval period, CDC has submitted eight GenICs consisting of 750 responses. The collections included web-based surveys, focus groups, and assessments. The information collection activities conducted under this extension will continue to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration’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. 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 CDC and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

Feedback collected under this Generic Clearance will continue to provide useful information, but it will not yield data that can be generalized to the overall population. This type of Generic Clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: (1) the target population to which generalizations will be made; (2) the

sampling frame; (3) the sample design (including stratification and clustering); (4) the precision requirements or power calculations that justify the proposed sample size; (5) the expected response rate; (6) methods for assessing potential non-response bias; (7) the protocols for data collection; and 8) any testing procedures that were or will be undertaken prior fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other Generic mechanisms that are designed to yield quantitative results.

Respondents will be screened and selected from Individuals and Households, Businesses, Organizations, and/or State, Local or Tribal Government. Below we provide CDC’s projected annualized estimate for the next three years. There is no cost to respondents other than their time. The estimated annualized burden hours for this data collection activity are 9,690.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Type of collection	Number of respondents	Annual frequency per response	Hours per response
Individuals and Households, Businesses, Organizations, and/or State, Local or Tribal Government.	Online surveys	10,500	1	30/60
	Discussion Groups	280	1	120/60
	Focus groups	640	1	120/60
	Website/app usability testing	2,000	1	30/60
	Interviews	800	1	120/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2023–09518 Filed 5–3–23; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–23–1359; Docket No. CDC–2023–0034]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of

government information, invites the general public and other federal agencies the opportunity to comment on a continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled National Survey of Syringe Services Programs (NSSSP). This data collection which proposes assess and monitor SSP operational characteristics and services, funding resources, community relations, and key operational and programmatic successes and challenges, and support timely analysis and dissemination of national program evaluation survey findings.

DATES: CDC must receive written comments on or before July 3, 2023.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2021–0034 by either of the following methods:

- *Federal eRulemaking Portal:* www.regulations.gov. Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600

Clifton Road NE, MS H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to www.regulations.gov.

Please note: Submit all comments through the Federal eRulemaking portal (www.regulations.gov) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329; Telephone: 404–639–7570; Email: omb@cdc.gov.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also