

more rigorous designs that address: The target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods of assessing potential nonresponse bias, the protocols for data collection, and 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.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary.

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FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060-0550; FR ID 337624]

Information Collection Being Reviewed by the Federal Communications Commission Under Delegated

AGENCY: Federal Communications Commission.

ACTION: Notice; request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act (PRA) of 1995, the Federal Communications Commission (FCC or Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collections. Comments are requested concerning: whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection burden on small business concerns with fewer than 25 employees. The FCC may not conduct or sponsor a collection of information unless it displays a currently valid OMB control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the

PRA that does not display a valid OMB control number.

DATES: Written PRA comments should be submitted on or before June 1, 2026. If you anticipate that you will be submitting comments but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: Direct all PRA comments to Cathy Williams, FCC, via email *PRA@fcc.gov* and to *Cathy.Williams@fcc.gov*.

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Cathy Williams at (202) 418-2918.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060-0550.

Title: Local Franchising Authority Certification, FCC Form 328; Section 76.910, Franchising Authority Certification.

Form No.: FCC Form 328.

Type of Review: Extension of a currently approved collection.

Respondents: State, local or tribal governments; Businesses or other for-profit entities.

Number of Respondents and Responses: 7 respondents; 13 responses.
Estimated Time per Response: 2 hours.

Frequency of Response: One-time reporting requirement; Third party disclosure requirement.

Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this collection of information is contained in section 3 of the Cable Television Consumer Protection and Competition Act of 1992 (47 U.S.C. 543), as well as sections 4(i), 4(j), and 623 of the Communications Act of 1934, as amended, and section 111 of the STELA Reauthorization Act of 2014.

Total Annual Burden: 26 hours.

Total Annual Cost: None.

Needs and Uses: On June 3, 2015, the Commission released a Report and Order, MB Docket No. 15-53; FCC 15-62. The Report and Order adopted a rebuttable presumption that cable operators are subject to competing provider effective competition. The information collection requirements have not changed since they were last approved by the Office of Management and Budget (OMB). The information collection requirements consist of:

FCC Form 328. Pursuant to section 76.910, a franchising authority must be certified by the Commission to regulate the basic service tier and associated equipment of a cable system within its jurisdiction. To obtain this certification, the franchising authority must prepare and submit FCC Form 328. The Report

and Order revised § 76.910 to require a franchising authority filing Form 328 to submit specific evidence demonstrating its rebuttal of the presumption in § 76.906 that the cable system is subject to competing provider effective competition pursuant to § 76.905(b)(2). The franchising authority bears the burden of submitting evidence rebutting the presumption that competing provider effective competition, as defined in § 76.905(b)(2), exists in the franchise area. Unless a franchising authority has actual knowledge to the contrary, it may rely on the presumption in section 76.906 that the cable system is not subject to one of the other three types of effective competition.

Evidence establishing lack of effective competition. If the evidence establishing the lack of effective competition is not otherwise available, § 76.910(b)(4) provides that franchising authorities may request from a multichannel video programming distributor (MVPD) information regarding the MVPD's reach and number of subscribers. An MVPD must respond to such request within 15 days. Such responses may be limited to numerical totals.

Franchising authority's obligations if certified. Section 76.910(e) of the Commission's rules currently provides that, unless the Commission notifies the franchising authority otherwise, the certification will become effective 30 days after the date filed, provided, however, that the franchising authority may not regulate the rates of a cable system unless it: (1) Adopts regulations (i) consistent with the Commission's regulations governing the basic tier and (ii) providing a reasonable opportunity for consideration of the views of interested parties, within 120 days of the effective date of the certification; and (2) notifies the cable operator that the franchising authority has been certified and has adopted the required regulations.

Federal Communications Commission.

Aleta Bowers,

Federal Register Liaison Officer, Office of the Secretary.

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GOVERNMENT ACCOUNTABILITY OFFICE

Request for Nominations for the Board of Governors of the Patient-Centered Outcomes Research Institute (PCORI)

AGENCY: Government Accountability Office.

ACTION: Request for letters of nomination and resumes.

SUMMARY: The Patient Protection and Affordable Care Act gave the Comptroller General of the United States responsibility for appointing up to 21 members to the Board of Governors of the Patient-Centered Outcomes Research Institute (PCORI). In addition, the Directors of the Agency for Healthcare Research and Quality and the National Institutes of Health, or their designees, are members of the Board. As the result of terms ending in September 2026, the Government Accountability Office (GAO) is accepting nominations in the following categories: a physician, a quality improvement or health services researcher, and a representative of private payers who represents employers who self-insure employee benefits. Nominations should be sent to the email address listed below. Acknowledgement of receipt will be provided within a week of submission.

DATES: Letters of nomination and resumes should be submitted no later than May 1, 2026, to ensure adequate opportunity for review and consideration of nominees prior to appointment.

ADDRESSES: Submit letters of nomination and resumes to PCORI@gao.gov. Include PCORI nominations in the subject line of the email.

FOR FURTHER INFORMATION CONTACT: Ray Sendejas at SendejasR@gao.gov or (202) 512-7114 if you do not receive an acknowledgement or need additional information. For general information, contact GAO's Office of Public Affairs at PublicAffairs@gao.gov.

Authority: 42 U.S.C. 1320e; 26 U.S.C. 9511.

Orice Williams Brown,

Acting Comptroller General of the United States.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-26-0147]

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 "National Blood Collection and Utilization Survey (NBCUS)" 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 December 5, 2025 to obtain comments from the public and affected agencies. CDC received 10 comments 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

National Blood Collection and Utilization Survey (NBCUS)—New—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The National Blood Collection and Utilization Survey (NBCUS) is a biennial survey of the blood collection and utilization community designed to produce reliable and accurate estimates of national and regional collections, utilization, and safety of all blood products. The survey includes a core of standard questions on blood collection, processing, and utilization practices. Proposed changes from the 2023 survey include adjustments to answer options to make them more straightforward, removal of policy questions that were required of blood centers by the end of 2023, defining a blood shortage, and addition of several new questions. New questions included information about bacterial transfusion-transmitted infections found in blood, length of time any blood shortage lasted, cold-stored platelets, and pathogen-reduced cryoprecipitated units. The rapidly changing environment in blood supply and demand makes it important to have regular, periodic data describing the state of U.S. blood collections and transfusions for understanding the dynamics of blood safety and availability.

Survey respondents will consist of community-based blood collection centers, hospital-based blood collection centers, and transfusing hospitals, except those reporting fewer than 100 inpatient surgeries per year. For the purposes of this ICR, federal burden is only being placed on facilities located within the 50 states and the District of Columbia.

CDC will take over the NBCUS data collection activities from HHS/OASH and requests OMB approval for an estimated 4,612 annual burden hours. There is no cost to respondents other than their time to participate.