

(2) The indispensability of the information to the success of an investigation or establishing a violation; and

(3) The need to protect against the destruction of records or information or to preserve testimony that may be necessary to investigate and prosecute violations of the rules of the Authority.

(g) Failure to comply with a subpoena or with the other provisions of this Rule may be penalized by the imposition of one or more penalties set forth in Rule 8200.

(h) The Authority may administer oaths to witnesses and require witnesses to testify under oath in matters within the jurisdiction of the Authority.

#### 8410. Review of Alleged Rule Violations

(a) The Authority may on its own initiative review any evidence or information submitted from any source which establishes a reasonable probability that one or more rules of the Authority in the Rule 2200 Series or the Rule 8100 Series have been violated or have not been enforced. If the evidence or information pertains to the running of a Covered Horserace, the evidence or information should be submitted to the Authority within 72 hours after the Covered Horserace. If the Authority deems the evidence or information submitted to be lacking in credibility, the Authority is not obligated to take further action. The Authority's decision whether to take further action is not subject to appeal.

(b) If the Authority determines there is a reasonable probability that the rules of the Authority have been violated or have not been enforced, the Authority shall conduct an investigation of the matter. The Authority may in its discretion:

(1) Assign the matter for a hearing pursuant to Rule 8320(b) or Rule 8330; or

(2) Issue a Notice of Suspected or Actual Violation pursuant to Rule 8200(d).

#### 8420. Civil Actions

The Authority shall request and obtain the approval of the Commission before initiating any civil action pursuant to 15 U.S.C. 3054(j). If the Commission does not act upon the request within 20 days of receipt, the Authority's request to initiate a civil action pursuant to 15 U.S.C. 3054(j) shall be deemed approved by the Commission.

By direction of the Commission.

**April J. Tabor,**

*Secretary.*

[FR Doc. 2025-17296 Filed 9-8-25; 8:45 am]

**BILLING CODE 6750-01-P**

### OFFICE OF GOVERNMENT ETHICS

#### **Agency Information Collection Activities; Submission for OMB Review; Proposed Collection; Comment Request for a Modified OGE Form 201 Request an Individual's Ethics Documents**

**AGENCY:** Office of Government Ethics (OGE).

**ACTION:** Notice of request for agency and public comments.

**SUMMARY:** After this first round notice and public comment period, the U.S. Office of Government Ethics (OGE) plans to submit a proposed modified OGE Form 201, entitled "Request an Individual's Ethics Documents," to the Office of Management and Budget (OMB) for review and approval of a three-year extension under the Paperwork Reduction Act of 1995. The OGE Form 201 is used by persons requesting access to executive branch public financial disclosure reports and other covered records.

**DATES:** Written comments by the public and agencies on this proposed extension are invited and must be received by November 10, 2025. Information on how to submit a written comment can be found in the **ADDRESSES** section of this notice.

Additionally, a virtual public meeting will be held on October 6, 2025 from 2:00-3:00 p.m. Eastern Time. Information on how to register for the meeting can be found in the **SUPPLEMENTARY INFORMATION** section of this notice.

**ADDRESSES:** Comments may be submitted to OGE, by any of the following methods:

*Email:* [usoge@oge.gov](mailto:usoge@oge.gov). (Include reference to "OGE Form 201 Paperwork Comment" in the subject line of the message.)

*Mail, Hand Delivery/Courier:* Office of Government Ethics, Suite 750, 250 E Street SW, Washington, DC 20024, Attention: McEvan Baum, Assistant Counsel.

*Instructions:* Comments may be posted on OGE's website, [www.oge.gov](http://www.oge.gov). Sensitive personal information, such as account numbers or Social Security numbers, should not be included. Comments generally will not be edited to remove any identifying or contact information.

#### **FOR FURTHER INFORMATION CONTACT:**

McEvan Baum at the U.S. Office of Government Ethics; telephone: 202-482-9287; TTY: 800-877-8339; Email: [usoge@oge.gov](mailto:usoge@oge.gov). An electronic copy of the OGE Form 201 version used to manually submit access requests to OGE or other executive branch agencies by mail or FAX is available in the Forms Library section of OGE's website at <http://www.oge.gov>. A paper copy may also be obtained, without charge, by contacting Mr. Baum. An automated version of the OGE Form 201, also available on OGE's website, enables the applicant to electronically fill out, submit and receive access to copies of the public financial disclosure reports certified by the U.S. Office of Government Ethics.

#### **SUPPLEMENTARY INFORMATION:**

*Title:* OGE Form 201 Request an Individual's Ethics Documents.

*Agency Form Number:* OGE Form 201.

*OMB Contro Number:* 3209-0002.

*Type of Information Collection:* Extension with modifications of a currently approved collection.

*Type of Review Request:* Regular.

*Respondents:* Individuals requesting access to executive branch public financial disclosure reports and other covered records.

*Estimated Annual Number of Respondents:* 12,461.

*Estimated Time per Response:* 10 minutes.

*Estimated Total Annual Burden:* 2,077 hours.

*Abstract:* The OGE Form 201 collects information from, and provides certain information to, persons who seek access to OGE Form 278 Public Financial Disclosure Reports, including OGE Form 278-T Periodic Transaction Reports, and other covered records. The form reflects the requirements of the Ethics in Government Act, subsequent amendments pursuant to the STOCK Act, and OGE's implementing regulations that must be met by a person before access can be granted. These requirements include the address of the requester, as well as any other person on whose behalf a record is sought, and acknowledgement that the applicant is aware of the prohibited uses of executive branch public disclosure financial reports. See 5 U.S.C. 13107(b) and (c) and 13122(b)(1) and 5 CFR 2634.603(c) and (f). Executive branch departments and agencies are encouraged to utilize the OGE Form 201 for individuals seeking access to public financial disclosure reports and other covered documents. OGE permits departments and agencies to use or

develop their own forms as long as the forms collect and provide all of the required information.

OGE currently has OMB approval for six versions of the form: OGE's online application (English, Arabic, Chinese, French, and Spanish), and a PDF version in English.

OGE is proposing two changes to both English language versions of the form. First, OGE proposes updating the language of its Privacy Act Statement. Second, OGE proposes replacing the outdated Ethics in Government Act citation with the current citation.

OGE is also proposing additional changes only to the English language PDF version of the form, as summarized below:

- Adding text to the Supplemental Information section, page 3 (section 5a), and page 5 (above the large text box) to inform requesters of general retention periods that may impact availability of documents they are requesting.
- Modifying the text in several places to improve accuracy, clarity, consistency, and align with plain language principles.

To view a pdf draft containing the proposed changes, visit [https://www.oge.gov/web/OGEnsf/0/7CB/E89A1496B40E785258CEC00555DC2/\\$FILE/OGEnsf/20Form%20201%20Renewal\\_Draft\\_website.pdf](https://www.oge.gov/web/OGEnsf/0/7CB/E89A1496B40E785258CEC00555DC2/$FILE/OGEnsf/20Form%20201%20Renewal_Draft_website.pdf).

To register to attend the virtual public meeting on October 6, 2025 from 2:00–3:00 p.m. Eastern Time, visit <https://events.gcc.teams.microsoft.com/event/2a12717b-0338-4df7-af40-342375b8dd51@c0abca44-0182-40a9-8010-01ec94254f77>.

**Request for Comments:** Agency and public comment is invited specifically on the need for and practical utility of this information collection, the accuracy of OGE's burden estimate, the enhancement of quality, utility and clarity of the information collected, and the minimization of burden (including the use of information technology). Comments received in response to this notice will be summarized for and included with the OGE request for extension of OMB paperwork approval. The comments will also become a matter of public record.

Specifically, OGE seeks public comment on the following:

- What problems do you have using the form?
- Are there sections of the form or instructions that are unclear?
- Is there information provided that is confusing?
- What additional information would be helpful?

**Authority:** 44 U.S.C. 3506(c)(2)(A); 5 U.S.C. 13107(b) and (c); 5 U.S.C. 13122(b)(1); 5 CFR 2634.603(c) and (f).

Approved: September 4, 2025.

**Shelley K. Finlayson,**

*Chief of Staff and Program Counsel, Office of Government Ethics.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[30Day–25–0600]

#### 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 “CDC Model Performance Evaluation Program (MPEP) for *Mycobacterium tuberculosis* Drug Susceptibility Testing” 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 June 16, 2025, to obtain comments from the public and affected agencies. CDC received no public 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:

- 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;
- 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;
- Enhance the quality, utility, and clarity of the information to be collected;
- 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

- 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](http://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

CDC Model Performance Evaluation Program (MPEP) for *Mycobacterium tuberculosis* Drug Susceptibility Testing (OMB Control No. 0920–0600, Exp. 9/30/2025)—Extension—National Center for HIV, Viral Hepatitis, STD, and Tuberculosis Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

#### Background and Brief Description

The CDC is requesting an Extension of a currently approved information collection from participants in the CDC Model Performance Evaluation Program (MPEP) for *Mycobacterium tuberculosis* Drug Susceptibility Testing for a period of three years. Extension of this information will not require changes in the scope of the project. As part of the Extension, CDC is requesting a non-substantive change to the title of the data collection to “CDC Model Performance Evaluation (MPEP) for *Mycobacterium tuberculosis* Drug Susceptibility Testing” from “CDC Model Performance Evaluation Program (MPEP) for *Mycobacterium tuberculosis* and Nontuberculous Mycobacteria Drug Susceptibility Testing” to reflect that nontuberculous mycobacteria are no longer included in the program.

Tuberculosis (TB) is a continuing public health problem. The overall number of cases, and associated incidence rate, in the United States has increased over the past several years. High rates are reported among non-U.S.-born persons, correctional populations, people experiencing homelessness, and individuals reported to have diabetes. To reach the goal of eliminating TB, this