Total Burden Hours: 41,298.
Total Annual Burden Hours: 148,912.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 3090–0292, FFATA Subaward and Executive Compensation Reporting Requirements, in all correspondence.

Dated: April 18, 2017.

Steve Grewal,

Deputy Chief Information Officer, General Services Administration.

[FR Doc. 2017–08600 Filed 4–27–17; 8:45 am]

BILLING CODE 6820–WY–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0291; Docket No. 2017–0001; Sequence 3]

Information Collection; FSRS Registration Requirements for Prime Grant Awardees

AGENCY: Office of the Integrated Award Environment, General Services Administration (GSA).

ACTION: Notice of request for public comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve a renewal of the currently approved information collection requirement regarding FSRS Registration Requirements for Prime Grant Awardees.

DATES: Submit comments on or before June 27, 2017.

ADDRESSES: Submit comments identified by Information Collection 3090–0291, FSRS Registration Requirements for Prime Grant Awardees by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by searching OMB control number 3090–0291. Select the link ''Comment Now'' that corresponds with ''Information Collection 3090–0291, FSRS Registration Requirements for Prime Grant Awardees.'' Follow the instructions provided on the screen. Please include your name, company name (if any), and ''Information Collection 3090–0291, FSRS Registration Requirements for Prime Grant Awardees on your attached document.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: IC 3090–0291, FSRS Registration Requirements for Prime Grant Awardees.

Instructions: Please submit comments only and cite Information Collection 3090–0291, FSRS Registration Requirements for Prime Grant Awardees, in all correspondence related to this collection. All comments received will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided.

FOR FURTHER INFORMATION CONTACT: John Corro, Procurement Analyst, Office of the Integrated Award Environment, GSA, at telephone number 202–215–9767; or via email at *john.corro@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

The Federal Funding Accountability and Transparency Act (P.L. 109–282, as amended by section 6202(a) of P.L. 110–252), known as FFATA or the Transparency Act, requires information disclosure of entities receiving Federal financial assistance through Federal awards such as Federal contracts, sub-contracts, grants and sub-grants, FFATA 2(a),(2),(i),(ii). The system that collects this information is called the FFATA Sub-award Reporting System (FSRS, *www.fsrs.gov*). This information collection requires information necessary for prime awardee registration in FSRS to create a user log-in and enable sub-award reporting for their entity. To register in FSRS for a user log-in, an entity is required to provide their Data Universal Numbering System (DUNS) number. FSRS then pulls core data about the entity from their System for Award Management (SAM) registration to include the legal business name, physical address, mailing address and Commercial and Government Entity (CAGE) code. The entity completes the FSRS registration by providing contact information within the entity for approval.

If a prime awardee has already registered in FSRS to report contracts-related Transparency Act financial data, a new log-in will not be required. In addition, if a prime awardee had a user account in the Electronic Subcontract Reporting System (eSRS), a new log-in will not be required.

B. Annual Reporting Burden

Respondents: 5,678.
Responses Per Respondent: 1.
Total Annual Responses: 5,678.
Hours Per Response: .5.
Total Burden Hours: 2,839.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755.

Please cite OMB Control No. 3090–0291, FSRS Registration Requirements for Prime Grant Awardees, in all correspondence.

Dated: April 18, 2017.

Steve Grewal,

Deputy Chief Information Officer, General Services Administration.

[FR Doc. 2017–08602 Filed 4–27–17; 8:45 am]

BILLING CODE 6820–XY–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0118; Docket 2017–0001; Sequence 2]

Information Collection; Statement of Witness, Standard Form 94

AGENCY: Federal Vehicle Policy Division, General Services Administration (GSA).

ACTION: Notice of a request for comments regarding a reinstatement, with change, to an OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995, GSA has submitted to the Office of Management and Budget (OMB) a request to review and approve a reinstatement, with change, to an

information collection requirement concerning Standard Form 94, Statement of Witness.

DATES: Submit comments on or before June 27, 2017.

FOR FURTHER INFORMATION CONTACT: Ray Wynter, Federal Vehicle Policy Division, 202–501–3802, or via email at *ray.wynter@gsa.gov.*

ADDRESSES: Submit comments identified by Information Collection 3090–0118, Statement of Witness, SF 94, by any of the following methods: Regulations.gov: *http://www.regulations.gov.*

Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link ''Submit a Comment'' that corresponds with ''Information Collection 3090–0118, Statement of Witness, SF 94.'' Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 3090–0118, Statement of Witness, SF 94'' on your attached document.

- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Sosa/IC 3090–0118, Statement of Witness, SF 94.

Instructions: Please submit comments only and cite Information Collection 3090–0118, Statement of Witness, SF 94, in all correspondence related to this collection. Comments received generally will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two to three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

SUPPLEMENTARY INFORMATION:

A. Purpose

GSA is requesting the Office of Management and Budget (OMB) to review and approve information collection, 3090–0118, Statement of Witness, SF 94. This form is used by all Federal agencies to report accident information involving U.S. Government motor vehicles.

B. Annual Reporting Burden

Respondents: 874.
Responses per Respondent: 1.
Total Annual Responses: 874.
Hours per Response: .333.
Total Burden Hours: 291.

C. Public Comment

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 3090–0118, Statement of Witness, SF 94, in all correspondence.

Dated: April 18, 2017.

Steve Grewal,

Deputy Chief Information Officer, General Services Administration.

[FR Doc. 2017–08603 Filed 4–27–17; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed changes to the currently approved information collection project: ''*Developing a Registry of Registries.*''

DATES: Comments on this notice must be received by June 27, 2017.

ADDRESSES: Written comments should be submitted to: Doris Lefkowitz, Reports Clearance Officer, AHRQ, by email at *doris.lefkowitz@AHRQ.hhs.gov.*

Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427–1477, or by email at *doris.lefkowitz@AHRQ.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Proposed Revision of a Currently Approved Collection Project: ''Developing a Registry of Registries.'' OMB Control Number 0935–0203

In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501–3521, AHRQ invites the public to comment on this proposed information collection on the development of a registry of patient registries. Patient registries have received significant attention and funding in recent years. Similar to controlled studies, patient registries represent some burden to patients (*e.g.,* time to complete patient reported outcome measures, risk of loss of privacy), who often participate voluntarily in hopes of improving knowledge about a disease or condition. Patient registries also represent a substantial investment of health research resources. Despite these factors, patient registries are not required to be registered in *ClinicalTrials.gov*, presenting the potential for duplication of efforts and insufficient dissemination of findings that are not published in the peer-reviewed literature. To fulfill the obligation to patients and to ensure that resources are used in the most efficient manner, registries need to be listed in a manner similar to that of trials in *ClinicalTrials.gov*.

By providing a centralized point of collection for information about all patient registries in the United States, the Registry of Patient Registries (RoPR) enhances patient registry information, extracted from *ClinicalTrials.gov*, building on AHRQ's efforts to describe the quality, appropriateness, and effectiveness of health services (and patient registries in particular) in a more readily available, central location.

The RoPR database system aims to achieve the following objectives:

(1) Provide a searchable database of patient registries in the United States (to promote collaboration, reduce redundancy, and improve transparency);

(2) Facilitate the use of common data fields and definitions in similar health conditions (to improve opportunities for sharing, comparing, and linkage);

(3) Provide a public repository of searchable summary results (including results from registries that have not yet been published in the peer-reviewed literature);

(4) Offer a search tool to locate existing data that researchers can request for use in new studies; and

(5) Serve as a recruitment tool for researchers and patients interested in participating in patient registries.