the Internet. A search capability for all Center for Devices and Radiological Health guidance documents is available at *http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/GuidanceDocuments/default.htm.* Guidance documents are also available at *http://www.regulations.gov.* Persons unable to download an electronic copy of ''Display Devices for Diagnostic Radiology'' may send an email request to *CDRH-Guidance@fda.hhs.gov* to receive an electronic copy of the document. Please use the document number 1500022 to identify the guidance you are requesting.

IV. Paperwork Reduction Act of 1995

The draft guidance refers to previously approved collections of information found in FDA regulations. These collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in 21 CFR part 807 have been approved under OMB control number 0910–0120 and the collections of information in 21 CFR part 801 have been approved under OMB control number 0910–0485.

Dated: February 2, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016–02521 Filed 2–8–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: HHS–OS–4040–New–30D]

Agency Information Collection Activities; Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, has submitted an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB) for review and approval. The ICR is for a new collection. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public on this ICR during the review and approval period.

DATES: Comments on the ICR must be received on or before March 10, 2016.

ADDRESSES: Submit your comments to *OIRA_submission@omb.eop.gov* or via facsimile to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, *Information.CollectionClearance@hhs.gov* or (202) 690–6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the Information Collection Request Title and document identifier HHS–OS–4040–New–30D for reference.

Information Collection Request Title: DATA Act Sec. 5. ''Simplifying Federal Award Reporting'' Grants Pilot

Abstract: Public Law 113–101, The Digital Accountability and Transparency Act of 2014 (DATA Act) expands the Federal Funding Accountability and Transparency Act of 2006 by increasing accountability and transparency in Federal spending. Section 5 of the DATA Act (''Sec. 5. Simplifying Federal Award Reporting'') tasks the Director of the Office of Management and Budget (OMB) to establish a pilot program (Sec. 5 (b)).

OMB has designated the Department of Health and Human Services (HHS) as the executing agent of the pilot program. Within HHS, the DATA Act Program Management Office (PMO) (DAP) has been established under the Office of the Assistant Secretary for Financial Resources (ASFR) in order to implement this pilot program. ASFR/DAP, in coordination with Grants.gov, is requesting a generic clearance for the purpose of conducting tests under the pilot program to obtain qualitative and quantitative data and gain an understanding of the burden imposed on Federal recipients.

The DAP has designed several test models to evaluate recipient burden and assess quality of data. The goal of these test models is to determine whether new technology, data standards, processes, and forms aid in reducing recipient burden and increase the accuracy and quality of the data submitted. Under this clearance, a variety of methods (surveys, focus groups, etc.) could be used to collect data, with the exact nature of the questions currently undetermined. DAP expects these questions to include, but not be limited to, topics pertaining to the Standard Form (SF) 424, the Consolidated Federal Financial Reports, and the expanded Single Audit form (SF–SAC). If this data is not collected, the requirements of the DATA Act Section 5 pilot will not be met. The types of collections that this generic clearance covers include, but are not limited to:

- Surveys,
- Focus Groups,
- Other qualitative methods such as interviews, small discussion groups, and case studies.

Likely Respondents: Recipients of Federal contracts, grants, and sub-awards.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Estimated annual reporting burden				
Type of collection	Number of respondents	Annual frequency per response	Hours per response	Total hours
Surveys, Focus Groups, and other qualitative methods	300	1	56.25	16,875
Total	300			16,875

Darius Taylor,

Information Collection Clearance Officer.

[FR Doc. 2016–02472 Filed 2–8–16; 8:45 am]

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