

**DEPARTMENT OF DEFENSE****GENERAL SERVICES  
ADMINISTRATION****NATIONAL AERONAUTICS AND  
SPACE ADMINISTRATION**

[OMB Control No. 9000–0113; Docket 2016–0053; Sequence 43]

**Information Collection; Acquisition of Helium**

**AGENCY:** Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

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

**SUMMARY:** Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division (MVCB) will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement concerning acquisition of helium.

**DATES:** Submit comments on or before January 23, 2017.

**ADDRESSES:** Submit comments identified by Information Collection 9000–0113, Acquisition of Helium, 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 9000–0113. Select the link “Comment Now” that corresponds with “Information Collection 9000–0113, Acquisition of Helium”, Follow the instructions provided on the screen. Please include your name, company name (if any), and “Information Collection 9000–0113, Acquisition of Helium”, on your attached document.

- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0113, Acquisition of Helium.

**Instructions:** Please submit comments only and cite Information Collection 9000–0113, Acquisition of Helium, 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](http://www.regulations.gov), approximately two to three days after submission to verify posting (except

allow 30 days for posting of comments submitted by mail).

**FOR FURTHER INFORMATION CONTACT:** Ms. Mahruba Uddowla, Procurement Analyst, Office of Governmentwide Acquisition Policy, GSA, via telephone 703–605–2868 or via email to [mahruba.uddowla@gsa.gov](mailto:mahruba.uddowla@gsa.gov).

**SUPPLEMENTARY INFORMATION:****A. Purpose**

The Helium Act (Pub. L. 86–777) (50 U.S.C. 167a, *et seq.*) and the Department of the Interior’s implementing regulations (30 CFR parts 601 and 602) require Federal agencies to procure all major helium requirements from the Bureau of Land Management, Department of the Interior.

FAR 8.5, Acquisition of Helium, and the clause 52.208–8 Required Sources for Helium and Helium Usage Data, requires that the Contractor provide to the Contracting Officer the following data within 10 days after the Contractor or subcontractor receives a delivery of helium from a Federal helium supplier; (i) The name of the supplier; (ii) The amount of helium purchased; (iii) The delivery date(s); and (iv) the location where the helium was used. Such information will facilitate enforcement of the requirements of the Helium Act and the contractual provisions requiring the use of Government helium by agency contractors.

The information is used in administration of certain Federal contracts to ensure contractor compliance with contract clauses. Without the information, the required use of Government helium cannot be monitored and enforced effectively. The FAR requires that the contractor provide helium purchase information 10 days after delivery from a federal helium supplier, not for the contractor to forecast what they are going to purchase.

**B. Annual Reporting Burden**

In consultation with subject matter experts at the Department of the Interior, Bureau of Land Management, Helium Operations, the number of estimated responses per year was verified as being within an acceptable range, as was the average time required to read and prepare information which was estimated at 1 hour per response.

*Respondents:* 26.  
*Responses per Respondent:* 1.  
*Total Responses:* 26.  
*Hours per Response:* 1.  
*Total Burden Hours:* 26.

**C. Public Comments**

*Public comments are particularly invited on:* Whether this collection of

information is necessary for the proper performance of functions of the FAR, 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; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

**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. 9000–0113, Acquisition of Helium, in all correspondence.

Dated: November 17, 2016.

**Lorin S. Curit,**

*Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.*

[FR Doc. 2016–28062 Filed 11–21–16; 8:45 am]

**BILLING CODE 6820–EP–P**

**GENERAL SERVICES  
ADMINISTRATION**

[Notice–ISP–2016–03; Docket No. 2016–0002; Sequence No. 23]

**Privacy Act of 1974: Revised Privacy Act System of Records; Withdrawal**

**AGENCY:** General Services Administration (GSA).

**ACTION:** Notice of a Privacy Act system of records; withdrawal.

**SUMMARY:** Access Certificates for Electronic Services GSA/GOVT–5 original intent was to facilitate secure, on-line communication between federal automated information systems and the public, using digital signature technologies to authenticate and verify identity. The reason for the cancellation notice is the system is no longer in use as a government system and has been updated as a commercial affiliate program of the Federal PKI.

**DATES:** *Effective:* November 22, 2016.

**FOR FURTHER INFORMATION CONTACT:** Call or email the GSA Privacy Act Officer:

telephone 571-388-6570; or email [gsa.privacyact@gsa.gov](mailto:gsa.privacyact@gsa.gov).

**Pranjali Desai**,  
Director, Office of Information Management,  
General Services Administration.

[FR Doc. 2016-28084 Filed 11-21-16; 8:45 am]

BILLING CODE 6820-34-P

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[60Day-17-17CQ]

**Proposed Data Collection Submitted for Public Comment and Recommendations—Zika Virus Associated Neurologic Illness Case Control Study; Correction**

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

**ACTION:** Notice; correction.

**SUMMARY:** The Centers for Disease Control and Prevention (CDC) published a document in the **Federal Register** of November 17, 2016, concerning request

for comments on *Proposed Data Collection Submitted for Public Comment and Recommendations—Zika Virus Associated Neurologic Illness Case Control Study*. The document provided the incorrect agency identification number (60Day-17-17ZQ).

**FOR FURTHER INFORMATION CONTACT:** Leroy Richardson, 1600 Clifton Road, MS D-74, Atlanta, GA 30333; telephone (404) 639-4965; email: [omb@cdc.gov](mailto:omb@cdc.gov).

**Correction**

In the **Federal Register** of November 17, 2016, in FR Doc. 2016-27692, on page 81143, in the first column (first heading), correct the agency identification number to read:

[60Day-17-17CQ; Docket No. CDC-2016-0107]

Dated: November 17, 2016.

**Leroy A. Richardson**,  
Chief, Information Collection Review Office,  
Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2016-28072 Filed 11-21-16; 8:45 am]

BILLING CODE 4163-18-P

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Proposed Information Collection Activity; Comment Request**

**Proposed Projects**

*Title:* State Plan Child Support Collection and Establishment of Paternity Title IV-D, OCSE-100.

*OMB No.:* 0970-0017.

*Description:* The Office of Child Support Enforcement has approved a IV-D state plan for each state. Federal regulations require states to amend their state plans only when necessary to reflect new or revised federal statutes or regulations or material change in any state laws, regulations, policies, or IV-D agency procedures. The requirement for submission of a state plan and plan amendments for the Child Support Enforcement program is found in sections 452, 454, and 466 of the Social Security Act.

*Respondents:* State IV-D Agencies.

**ANNUAL BURDEN ESTIMATES**

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
State Plan (OCSE-100)	54	5	.5	135
State Plan Transmittal (OCSE-21-U4)	54	5	.25	67.5

*Estimated Total Annual Burden Hours:* 202.5.

In compliance with the requirements of the Paperwork Reduction Act of 1995 (Pub. L. 104-13, 44 U.S.C. Chap 35), the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington DC 20201. Attn: ACF Reports Clearance Officer. Email address: [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov). All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the

agency's estimate of the burden of the proposed collection of information; (c) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

**Robert Sargis**,  
Reports Clearance Officer.

[FR Doc. 2016-28107 Filed 11-21-16; 8:45 am]

BILLING CODE 4184-01-P

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2015-E-4875]

**Determination of Regulatory Review Period for Purposes of Patent Extension; ANAVIP**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) has determined the regulatory review period for ANAVIP and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of an application to the Director of the U.S. Patent and Trademark Office (USPTO), Department of Commerce, for the extension of a patent which claims that human biological product.