of the information to be collected; and (d) ways to minimize the burden of the information collection on respondents, including through the use of automated collection techniques or other forms of information technology.

DATES: Consideration will be given to all comments received by July 17, 2015.

ADDRESSES: You may submit comments, identified by docket number and title, by any of the following methods:

- Federal eRulemaking Portal: *http://www.regulations.gov.* Follow the instructions for submitting comments.
- Mail: Department of Defense, Office of the Deputy Chief Management Officer, Directorate of Oversight and Compliance, Regulatory and Audit Matters Office, 9010 Defense Pentagon, Washington, DC 20301–9010.

Instructions: All submissions received must include the agency name, docket number and title for this **Federal Register** document. The general policy for comments and other submissions from members of the public is to make these submissions available for public viewing on the Internet at *http://www.regulations.gov* as they are received without change, including any personal identifiers or contact information.

Any associated form(s) for this collection may be located within this same electronic docket and downloaded for review/testing. Follow the instructions at *http://www.regulations.gov* for submitting comments. Please submit comments on any given form identified by docket number, form number, and title.

FOR FURTHER INFORMATION CONTACT: To request more information on this proposed information collection or to obtain a copy of the proposal and associated collection instruments, please write to the Defense Health Services Systems (DHSS) Program Executive Office (PEO), ATTN: CDR Patrick Amersbach, Defense Health Headquarters (DHHQ) 7700 Arlington Boulevard, Falls Church ,VA 22042–2902, or call 703–681–0845.

SUPPLEMENTARY INFORMATION:

Title; Associated Form; and OMB Number: Centralized Credentials Quality Assurance System (CCQAS); OMB Control Number 0720–TBD.

Needs and Uses: CCQAS v2.9.11 is an automated Tri-Service, Web-based database containing credentialing, privileging, risk management, and adverse actions information on direct healthcare providers in the MHS. CCQAS also allows providers to apply for privileges online. This latter capability allows for a privileging workflow for new providers, for transfers (TDY and PCS), for modification of privileges, and for renewal of privileges and staff reappointment within the system. CCQAS was CAC enforced December 2009 and as part of the Federal Health Care Center, North Chicago, VA PIV users gained access in October 2010. In November 2011, CCQAS was PKI/SSO integrated.

Affected Public: Individuals or Households.

Annual Burden Hours: 80,000.

Number of Respondents: 40,000.

Responses per Respondent: 1.

Average Burden per Response: 2 hours.

Frequency: On occasion.

Currently, CCQAS provides credentialing, privileging, risk-management and adverse actions capabilities which support medical quality assurance activities in the direct care system. CCQAS is fully deployed world-wide and is used by all Services (Army, Navy, Air Force) and Components (Guard, Reserve). CCQAS serves users functioning at the facility (defined by an individual UIC), Service, and DoD levels. Access to CCQAS modules and capabilities within each module is permissions-based, so that users have access tailored to the functions they perform and sensitive information receives maximal protection. Within each module, access control is available to the screen level.

Dated: May 13, 2015.

Aaron Siegel,

Alternate OSD Federal Register, Liaison Officer, Department of Defense.

[FR Doc. 2015–11971 Filed 5–15–15; 8:45 am]

BILLING CODE 5001–06–P

DEPARTMENT OF DEFENSE

Office of the Secretary

Uniform Formulary Beneficiary Advisory Panel; Notice of Federal Advisory Committee Meeting

AGENCY: Assistant Secretary of Defense (Health Affairs), DoD.

ACTION: Notice of meeting.

SUMMARY: The Department of Defense is publishing this notice to announce a Federal Advisory Committee meeting of the Uniform Formulary Beneficiary Advisory Panel (hereafter referred to as the Panel).

DATES: Thursday, June 11, 2015, from 9:00 a.m. to 1:00 p.m.

ADDRESSES: Naval Heritage Center Theater, 701 Pennsylvania Avenue NW., Washington, DC 20004.

FOR FURTHER INFORMATION CONTACT: Mr. William H. Blanche, Alternate DFO, Uniform Formulary Beneficiary Advisory Panel, 7700 Arlington Boulevard, Suite 5101, Falls Church, VA 22042–5101. Telephone: (703) 681–2890. Fax: (703) 681–1940. Email Address: *dha.ncr.health-it.mbx.baprequests@mail.mil.*

SUPPLEMENTARY INFORMATION:

This meeting is being held under the provisions of the Federal Advisory Committee Act of 1972 (Title 5, United States Code (U.S.C.), Appendix, as amended) and the Government in the Sunshine Act of 1976 (5 U.S.C. 552b, as amended).

Purpose of Meeting: The Panel will review and comment on recommendations made to the Director of Defense Health Agency, by the Pharmacy and Therapeutics Committee, regarding the Uniform Formulary.

Meeting Agenda

1. Sign-In
2. Welcome and Opening Remarks
3. Public Citizen Comments
4. Scheduled Therapeutic Class Reviews (Comments will follow each agenda item)
   a. Newer Oral Anticoagulants and Warfarin
   b. Hepatitis Antivirals (Hepatitis C)
5. Designated Newly Approved Drugs in Already-Reviewed Classes
6. Pertinent Utilization Management Issues
7. Panel Discussions and Vote

Meeting Accessibility: Pursuant to 5 U.S.C. 552b, as amended, and 41 Code of Federal Regulations (CFR) 102–3.140 through 102–3.165, and the availability of space, this meeting is open to the public. Seating is limited and will be provided only to the first 220 people signing-in. All persons must sign-in legibly.

Administrative Work Meeting: Prior to the public meeting, the Panel will conduct an Administrative Work Meeting from 8:00 a.m. to 9:00 a.m. to discuss administrative matters of the Panel. The Administrative Work Meeting will be held at the Naval Heritage Center, 701 Pennsylvania Avenue NW., Washington, DC 20004. Pursuant to 41 CFR 102–3.160, the Administrative Work Meeting will be closed to the public.

Written Statements: Pursuant to 41 CFR 102–3.140, the public or interested organizations may submit written statements to the membership of the Panel at any time or in response to the stated agenda of a planned meeting. Written statements should be submitted to the Panel's Designated Federal Officer

(DFO). The DFO's contact information can be obtained from the General Services Administration's Federal Advisory Committee Act Database at *http://facadatabase.gov/.*

Written statements that do not pertain to the scheduled meeting of the Panel may be submitted at any time. However, if individual comments pertain to a specific topic being discussed at a planned meeting, then these statements must be submitted no later than 5 business days prior to the meeting in question. The DFO will review all submitted written statements and provide copies to all the committee members.

Public Comments: In addition to written statements, the Panel will set aside 1 hour for individuals or interested groups to address the Panel. To ensure consideration of their comments, individuals and interested groups should submit written statements as outlined in this notice; but if they still want to address the Panel, then they will be afforded the opportunity to register to address the Panel. The Panel's DFO will have a ''Sign-Up Roster'' available at the Panel meeting for registration on a first-come, first-serve basis. Those wishing to address the Panel will be given no more than 5 minutes to present their comments, and at the end of the 1 hour time period, no further public comments will be accepted. Anyone who signs-up to address the Panel, but is unable to do so due to the time limitation, may submit their comments in writing; however, they must understand that their written comments may not be reviewed prior to the Panel's deliberation.

To ensure timeliness of comments for the official record, the Panel encourages that individuals and interested groups consider submitting written statements instead of addressing the Panel.

Dated: May 13, 2015.

Aaron Siegel,

Alternate OSD Federal Register Liaison Officer, Department of Defense.

[FR Doc. 2015–11977 Filed 5–15–15; 8:45 am]

BILLING CODE 5001–06–P

DEPARTMENT OF DEFENSE

Department of the Army

[Docket ID: USA–2015–0017]

Proposed Collection; Comment Request

AGENCY: Armed Forces Medical Examiner (AFMES), DoD.

ACTION: Notice.

SUMMARY: In compliance with the *Paperwork Reduction Act of 1995,* the Office of the Armed Forces Repository Specimen Samples for the Identification or Remains (AFRSSIR), a part of the Armed Forces Medical Examiner System (AFMES), announces a proposed public information collection and seeks public comment on the provisions thereof. Comments are invited 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 information collection; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the information collection on respondents, including through the use of automated collection techniques or other forms of information technology.

DATES: Consideration will be given to all comments received by July 17, 2015.

ADDRESSES: You may submit comments, identified by docket number and title, by any of the following methods:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Department of Defense, Office of the Deputy Chief Management Officer, Directorate of Oversight and Compliance, Regulatory and Audit Matters Office, 9010 Defense Pentagon, Washington, DC 20301–9010.

Instructions: All submissions received must include the agency name, docket number and title for this **Federal Register** document. The general policy for comments and other submissions from members of the public is to make these submissions available for public viewing on the Internet at *http://www.regulations.gov* as they are received without change, including any personal identifiers or contact information. Any associated form(s) for this collection may be located within this same electronic docket and downloaded for review/testing. Follow the instructions at *http://www.regulations.gov* for submitting comments. Please submit comments on any given form identified by docket number, form number, and title.

FOR FURTHER INFORMATION CONTACT: To request more information on this proposed information collection or to obtain a copy of the proposal and associated collection instruments, please write to the Office of the Armed Forces Repository of Specimen Samples for the Identification of Remains, Armed Forces Medical Examiner System (AFMES), 115 Purple Heart Drive, Dover AFB, DE, 19902–5051, ATTN: Mr. John Martin, Legal Counsel, AFMES at (302) 346–8634.

SUPPLEMENTARY INFORMATION:

Title; Associated Form; and OMB Number: Donor Specimen Card, OMB Control Number: 0702–XXXX.

Needs and Uses: The information collected will be used for the identification of human remains. The principal purpose of the information is to identify reference specimen samples that will routinely be stored and not analyzed until needed for remains identification program purposes.

Affected Public: Individuals or Households and Federal Government.

Annual Burden Hours: 62,500.

Number of Respondents: 250,000.

Responses per Respondent: 1.

Average Burden per Response: 15 minutes.

Frequency: On Occasion.

Respondents are deploying civilian or contractors and military personnel family members. The principal purpose of the collection is identify reference specimen samples that will be stored and not analyzed until needed for remain identification purposes. The donors at various military collection points and other federal agencies provide a blood sample which is stained on laboratory grade blood stain card (BSC). The identifying information on the blood stain card provided the donor reflects the individual's full name, signature, social security number (SSN), date of birth collection date and branch of service. The BSC is air dried and vacuumed sealed in a poly foil pouch. An adhesive label reflecting the donor information and redacted (SSN) is printed on the label, along with the unique accession number. In the event of the donor's death, the blood sample is scientifically analyzed and a DNA profile is created. This profile is then compared with the post-mortem sample obtained at the autopsy for positive identification.

Dated: May 12, 2015.

Aaron Siegel,

Alternate OSD Federal Register Liaison Officer, Department of Defense.

[FR Doc. 2015–11857 Filed 5–15–15; 8:45 am]

BILLING CODE 5001–06–P

DEPARTMENT OF EDUCATION

Applications for New Awards; Predominantly Black Institutions Competitive Grant Program

AGENCY: Office of Postsecondary Education, Department of Education