service on the HSRB. EPA will evaluate each candidate to assess whether there is any conflict of financial interest, appearance of a lack of impartiality, or prior involvement with matters likely to be reviewed by the Board.

Nominations will be evaluated on the basis of several criteria, including: Their professional background, expertise and experience that would contribute to the diversity of perspectives of the committee; interpersonal, verbal and written communication skills and other attributes that would contribute to the HSRB’s collaborative process; consensus building skills; absence of any financial conflicts of interest or the appearance of a lack of impartiality, or lack of independence, or bias; and the availability to attend meetings and administrative sessions, participate in teleconferences, develop policy recommendations to the Administrator, and prepare recommendations and advice in reports.

Nominations should include a resume or C.V. providing the nominee’s educational background, qualifications, leadership positions in national associations or professional societies, relevant research experience and publications along with a short (one page) biography describing how the nominee meets the above criteria and other information that may be helpful in evaluating the nomination, as well as the nominee’s current business address, e-mail address, and daytime telephone number. Interested candidates may self-nominate.

To help the Agency in evaluating the effectiveness of its outreach efforts, nominees are requested to inform the Agency of how you learned of this opportunity.

Final selection of HSRB members is a discretionary function of the Agency and will be announced on the HSRB Web site at [http://www.epa.gov/osa/hsrb/index.htm](http://www.epa.gov/osa/hsrb/index.htm) as soon as selections are made.

**ADDRESSES:** Submit your nominations by July 6, 2011, identified by Docket ID No. EPA–HQ–ORD–2011–0503, by any of the following methods:

- Internet: [http://www.regulations.gov](http://www.regulations.gov);
- Follow the on-line instructions for submitting comments.
- E-mail: ORD.Docket@epa.gov.
- Hand or Courier Delivery: EPA Docket Center (EPA/DC), Room 3304, EPA West Building, 1200 Constitution Ave., NW., Washington, DC 20460, Attention Docket ID No. EPA–HQ–ORD–2011–0503. Deliveries are accepted from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. Special arrangements should be made for deliveries of boxed information.


Dated: June 8, 2011.

Paul T. Anastas,
EPA Science Advisor.

[FR Doc. 2011–14681 Filed 6–13–11; 8:45 am]

**BILLING CODE 6560–50–P**

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**FARM CREDIT ADMINISTRATION**

**FARM CREDIT ADMINISTRATION Board; Sunshine Act; Special Meeting**

**AGENCY:** Farm Credit Administration.

**SUMMARY:** Notice is hereby given, pursuant to the Government in the Sunshine Act (5 U.S.C. 552b(e)(3)), of the special meeting of the Farm Credit Administration Board (Board).

**DATE AND TIME:** The special meeting of the Board will be held at the offices of the Farm Credit Administration in McLean, Virginia, on June 22, 2011, from 10 a.m. until such time as the Board concludes its business.

**FOR FURTHER INFORMATION CONTACT:** Dale L. Aultman, Secretary to the Farm Credit Administration Board, (703) 883–4009, TTY (703) 883–4036.

**ADDRESSES:** Farm Credit Administration, 1501 Farm Credit Drive, McLean, Virginia 22102–5900.

**SUPPLEMENTARY INFORMATION:** This meeting of the Board will be open to the public (limited space available). In order to increase the accessibility to Board meetings, persons requiring assistance should make arrangements in advance. The matter to be considered at the meeting is:

- Open Session
  - Request to Merge U.S. AgBank FCB with CoBank ACB

_Dated:_ June 10, 2011.

**FARM CREDIT ADMINISTRATION Board; Sunshine Act; Special Meeting**

**FARM CREDIT ADMINISTRATION**

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**FEDERAL COMMUNICATIONS COMMISSION**

**Information Collection Being Reviewed by the Federal Communications Commission Under Delegated Authority**

**AGENCY:** Federal Communications Commission.

**ACTION:** Notice and request for comments.

**SUMMARY:** As part of its continuing effort to reduce paperwork burden and as required by the Paperwork Reduction Act (PRA) of 1995 (44 U.S.C. 3501–3520), the Federal Communications Commission invites the general public and other Federal agencies to take this opportunity to comment on the following information collection(s). Comments are requested concerning: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; (b) the accuracy of the Commission’s burden estimate; (c) ways to enhance the quality, utility, and clarity of the information collected; (d) ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and (e) ways to further reduce the information burden for small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid OMB control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act (PRA) that does not display a valid OMB control number.

**DATES:** Written Paperwork Reduction Act (PRA) comments should be submitted on or before August 15, 2011. If you anticipate that you will be submitting PRA comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the FCC contact listed below as soon as possible.

**ADDRESSES:** Submit your PRA comments to Nicholas A. Fraser, Office of Management and Budget, via fax at 202–395–5167 or via Internet at Nicholas.A.Fraser@omb.eop.gov and to Benish Shah, Federal Communications Commission, via the Internet at Benish.Shah@fcc.gov. To submit your PRA comments by e-mail send them to: PRA@fcc.gov.
FOR FURTHER INFORMATION CONTACT: Benish Shah, Office of Managing Director, (202) 418–7866.

SUPPLEMENTARY INFORMATION:
OMB Control No.: 3060–0532.
Title: Sections 2.1033 and 15.121, Scanning Receiver Compliance Exhibits.
Form No.: N/A.
Type of Review: Extension of a currently approved collection.
Respondents: Business or other for-profit.
Number of Respondents: 25 respondents; 25 responses.
Estimated Time per Response: 1 hour.
Frequency of Response: One time reporting requirement and third party disclosure requirement.
Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this information collection is contained in 47 U.S.C. Sections 154(i), 301, 302, 303(e), 303(f), 303(g), 303(k), 304 and 307.
Total Annual Burden: 25 hours.
Annual Cost Burden: $1,250.
Privacy Act Impact Assessment: N/A.
Nature and Extent of Confidentiality: The Commission’s rules require that certain portions of scanning receiver applications for certification will remain confidential after the effective date of the grant of the application. No other assurances of confidentiality are provided to respondents.
Needs and Uses: This collection will be submitted as an extension (no change in reporting and/or third party disclosure requirements) after this 60-day comment period to the Office of Management and Budget (OMB) in order to obtain the full three year clearance.
The FCC rules under 47 CFR 2.1033 and 15.121 require manufacturers of scanning receivers to design their equipment so that it has 38 dB of image rejection for Cellular Service frequencies, tuning, control and filtering circuitry are inaccessible and any attempt to modify the scanning receiver to receive Cellular Service transmissions will likely render the scanning receiver inoperable. The Commission’s rules also require manufacturers to submit information with any application for certification that describes the testing method used to determine compliance with the 38 dB image rejection ratio, the design features that prevent modification of the scanning receiver to receive Cellular Service transmissions, and the design steps taken to make tuning, control, and filtering circuitry inaccessible. Furthermore, the FCC requires equipment to carry a statement assessing the vulnerability of the scanning receiver to modification and to have a label affixed to the scanning receiver, similar to the following as described in section 15.121:
Warning: Modification of this device to receive cellular radiotelephone service signals is prohibited under FCC Rules and Federal Law.
The Commission uses the information required in this equipment authorization process to determine whether the equipment that is being marketed complies with the Congressional mandate in the Telephone Disclosure and Dispute Resolution Act of 1992 (TDDRA) and applicable Commission rules.
Federal Communications Commission.
Marlene H. Dortch,
Secretary, Office of the Secretary, Office of Managing Director.
[FR Doc. 2011–14642 Filed 6–13–11; 8:45 am]
BILLING CODE 6712–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–11–11DD]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–3960 or send an e-mail to ombr/cdc.gov. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project


Background and Brief Description

The Division of Blood Disorders, located within the National Center on Birth Defects and Developmental Disabilities, implements health promotion and wellness programs designed to prevent secondary conditions in people with bleeding and clotting disorders. There are few public health problems as serious as deep vein thrombosis (DVT) and pulmonary embolism (PE), yet these conditions receive little attention. DVT/PE is an underdiagnosed, serious, preventable medical condition that occurs when a blood clot forms in a deep vein. These clots usually develop in the lower leg, thigh, or pelvis, but they can also occur in the arm. In more than one third of people affected by DVT, clots can travel to the lungs and cause PE, a potentially fatal condition.

The precise number of people affected by DVT/PE is unknown, but estimates range from 300,000 to 600,000 annually in the United States. DVT/PE is associated with substantial morbidity and mortality: One third of people with DVT/PE will have a recurrence within 10 years and one third of people die within 1 month of diagnosis. Among people who have had a DVT, one third will have long-term complications (post-thrombotic syndrome), such as swelling, pain, discoloration, and scaling in the affected limb. In some cases, the symptoms can be so severe that a person can become disabled. More troubling, sudden death is the first symptom in about one quarter of people who have a PE.

The Division of Blood Disorders submitted questions to the 2007 HealthStyles survey to determine the public’s knowledge of DVT, its common symptoms, and risk factors. Although over 60% of respondents identified pain and swelling as symptoms, 60% did not identify tenderness (often the first sign of DVT) as a symptom. Only 36% of respondents knew that a DVT was a blood clot in a vein, and most could not identify common risk factors for DVT such as sitting for a long period of time (e.g., during air travel); having a leg or foot injury; having a family member who has had a DVT; taking birth control pills; or getting older; and certain groups could not identify risk factors that specifically applied to their risk. The results of this survey demonstrate the need for greater awareness of DVT and its risk factors and the data show that there are many opportunities to develop audience specific messages that are age specific and culturally appropriate.

Much of the morbidity and mortality associated with DVT/PE could be prevented with early and accurate diagnosis and management. DVT/PE is preventable. It is important for people to be able to recognize the signs and symptoms and know when to seek care and available treatment. Individuals, families, and their support communities can reduce their risk by understanding DVT/PE and its risk factors. DVT/PE affects people of all races and ages.