Respondents: Submitters to the GTR are expected to include clinical laboratories, researchers, and entities that report and interpret tests performed elsewhere. The GTR is not limited to U.S. respondents; it will also include submissions from outside the United States. Information will be collected and managed using an online submission system.

Estimate of Burden: Although participation in the GTR is voluntary, in order to participate, respondents must provide information for a certain subset of fields, identified as the “minimal fields.” GTR includes 31 minimal fields and 85 optional fields. Sixteen of the 31 minimal fields refer to contact data and other information about the laboratory, which the respondent completes only once. These data will autopopulate new test records, leaving 15 minimal fields that require completion. The GTR will also support bulk submission as an XML file or uploading subsets of information from spreadsheets, which will significantly reduce the burden for laboratories that want to provide information on multiple genetic tests. The annualized cost to respondents is estimated at $1,103.

### Estimates of Hour Burden

<table>
<thead>
<tr>
<th>Type of respondents</th>
<th>Estimated number of respondents</th>
<th>Estimated number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Estimated total annual burden hours requested</th>
</tr>
</thead>
<tbody>
<tr>
<td>Laboratory</td>
<td>770</td>
<td>12</td>
<td>Minimal Fields: 0.5</td>
<td>Minimal Fields: 4,620.</td>
</tr>
<tr>
<td>Personnel</td>
<td></td>
<td></td>
<td>Optional Fields: 2.5</td>
<td>Optional Fields: 23,100.</td>
</tr>
<tr>
<td>Total</td>
<td>770</td>
<td>3.0</td>
<td></td>
<td>27,720.</td>
</tr>
</tbody>
</table>

Request For Comments: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to: Office of Management and Budget, Office of Regulatory Affairs, OIRA_submission@omb.eop.gov or by fax to [202] 395–6974. Attention: Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instrument, contact: Amy P. Patterson, M.D., Associate Director for Science Policy, NIH, by mail to the Office of Biotechnology Activities, 6705 Rockledge Dr., Suite 750, Bethesda, MD 20892; telephone (301) 496–9838; fax (301) 496–9839; or email gtr@od.nih.gov; or refer to the GTR Web site at http://oba.od.nih.gov/gtr/gtr.html.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

Dated: November 16, 2011.

Amy P. Patterson, Associate Director for Science Policy, NIH.

[FR Doc. 2011–30286 Filed 11–22–11; 8:45 am] BILLY CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentablematerial, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; Hemodialysis and Markers of Heart Failure.

Date: December 5, 2011.
Time: 2 p.m. to 3 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Barbara A WoynarowskA, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 754, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 402–7172, woynarowska@niddk.nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; Ancillary Study to the Hispanic Community Health Study.

Date: December 7, 2011.
Time: 3 p.m. to 4 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Carol J. Goter-Robinson, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 748, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–7791, goterrobinsnc@extra.niddk.nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; Ancillary Study to the Hispanic Community Health Study.

Date: December 8, 2011.
Time: 4:30 p.m. to 6 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy
DEPARTMENT OF HOMELAND SECURITY

[Docket No. DHS–2011–0098]

Agency Information Collection Activities: Submission for Review; Information Collection Request for the Department of Homeland Security (DHS), Science and Technology, Protected Repository for the Defense of Infrastructure Against Cyber Threats (PREDICT) Program

AGENCY: Science and Technology Directorate, DHS.

ACTION: 60-day Notice and request for comment.

SUMMARY: The Department of Homeland Security (DHS), Science & Technology (S&T) Directorate invites the general public to comment on data collection forms for the Protected Repository for the Defense of Infrastructure against Cyber Threats (PREDICT) program, and is a revision of a previously approved collection. The PREDICT program facilitates the accessibility of computer and network operational data for use in cyber security research and development through the establishment of distributed repositories of security-relevant network operations data, and the application procedures, protection policies, and review processes necessary to make this data available to the cyber defense research community. The forms allow the PREDICT initiative to provide a central repository, accessible through a Web-based portal (https://www.predict.org/) that catalogs current computer network operational data, provide secure access to multiple sources of data collected as a result of use and traffic on the Internet, and facilitate data flow among PREDICT participants for the purpose of developing new models, technologies and products that support effective threat assessment and increase cyber security capabilities. The PREDICT Coordinating Center (PCC) has established application procedures, protection policies, and review processes necessary to make this data available to the cyber defense research community, and PREDICT has been operational since Fall 2008. In order for a user to access PREDICT data, s/he must complete a registration form to establish a user account. The information collected is used by the DHS S&T PREDICT program to determine the authenticity and validate the requestor’s stated research against the data requested.

The DHS invites interested persons to comment on the following form and instructions (hereinafter "Forms Package") for the S&T PREDICT program. Interested persons may receive a copy of the Forms Package by contacting the DHS S&T PRA Coordinator. This notice and request for comments is required by the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. chapter 35).

DATES: Comments are encouraged and will be accepted until January 23, 2012.

ADDRESSES: Interested persons are invited to submit comments, identified by docket number DHS–2011–0098, by one of the following methods:

• E-mail: Millie.Ives@dhs.gov. Please include docket number DHS–2011–0098 in the subject line of the message.
• Fax: (202) 254–6171. (Not a toll-free number.)
• Mail: Science and Technology Directorate, ATTN: Chief Information Office—Millie Ives, 245 Murray Drive, Mail Stop 0202, Washington, DC 20528.

FOR FURTHER INFORMATION CONTACT: DHS S&T PRA Coordinator Millie Ives (202) 254–6828. (Not a toll free number.)

SUPPLEMENTARY INFORMATION: The information will be collected via the DHS S&T secure Web site at http://www.predict.org/. The PREDICT Web site employs only secure web-based technology to collect information from users to both reduce the burden and increase the efficiency of this collection.

The Department is committed to improving its information collection and urges all interested parties to suggest how these materials can further reduce burden while seeking necessary information under the Act. DHS is particularly interested in comments that:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Suggest ways to enhance the quality, utility, and clarity of the information to be collected; and

(4) Suggest ways to minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submissions of responses.

Overview of This Information Collection

(1) Type of Information Collection: Revision of a currently approved collection.

(2) Title of the Form/Collection: Science and Technology, Protected Repository for the Defense of Infrastructure against Cyber Threats (PREDICT) program.


(4) Affected public who will be asked or required to respond, as well as a brief