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.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Dr. Sarah L. Glavin, Deputy Director, Office of Science Policy, Analysis and Communication, National Institute of Child Health and Human Development, 31 Center Drive Room 2A18, Bethesda, Maryland 20892, or call non-toll free number (301) 496–1877 or E-mail your request, including your address, to glavins@mail.nih.gov.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Dated: November 5, 2010.

Sarah L. Glavin,
Deputy Director, Office of Science Policy, Analysis and Communications, National Institute of Child Health and Human Development, National Institutes of Health.

[FR Doc. 2010–28701 Filed 11–12–10; 8:45 am]
BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health

Proposed Collection; Comment Request; California Health Interview Survey Cancer Control Module (CHIS–CCM) 2011 (NCI)

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection: Title: California Health Interview Survey Cancer Control Module (CHIS–CCM) 2010. Type of Information Collection Request: Revision. Need and Use of Information Collection: The NCI has sponsored five Cancer Control Modules in the California Health Interview Survey (CHIS), and will be sponsoring a sixth this year. CHIS is a telephone survey that collects population-based, standardized health-related data to assess California’s progress in meeting Healthy People 2010 objectives for the nation and the State. The CHIS sample is designed to provide statistically reliable estimates statewide, for California counties, and for California’s ethnically and racially diverse population. Initiated by the UCLA Center for Health Policy Research, the California Department of Health Services, and the California Public Health Institute, the survey is funded by a number of public and private sources. It was first administered in 2001 to 55,428 adults and subsequently in 2003 to 42,043 adults, in 2005 to 43,020 adults, and in 2007 to 48,150 adults. These adults are a representative sample of California’s non-institutionalized population living in households. CHIS 2011 is planned for continual administration to 48,150 adults and 3,316 adolescent Californians. This study will allow NCI to examine patterns and trends in cancer screening and follow-up, as well as to study other cancer-related topics such as tobacco control, diet, physical activity, obesity, and human papillomavirus. Additionally, CHIS is designed to be comparable to the National Health Interview Survey (NHIS) data in order to conduct comparative analyses. CHIS provides enhanced estimates for cancer risk factors and screening among racial/ethnic minority populations. Frequency of Response: Once. Affected public: Individuals. Types of Respondents: U.S. adults and adolescents (persons 12 years of age and older). The total annual burden hours requested are 2,177 (see Table 1). There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

TABLE 1—ESTIMATES OF ANNUALIZED HOUR BURDEN

<table>
<thead>
<tr>
<th>Type of respondent</th>
<th>Form type</th>
<th>Number of respondents</th>
<th>Frequency of response</th>
<th>Average time per response—minutes/hours</th>
<th>Annual hour burden</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adults</td>
<td>Adult Pilot</td>
<td>50</td>
<td>1</td>
<td>8/60 (0.133)</td>
<td>6.67</td>
</tr>
<tr>
<td></td>
<td>Adult Survey</td>
<td>16,000</td>
<td>1</td>
<td>8/60 (0.133)</td>
<td>2,133.33</td>
</tr>
<tr>
<td>Adolescents</td>
<td>Adolescent Pilot</td>
<td>6</td>
<td>1</td>
<td>2/60 (0.033)</td>
<td>0.20</td>
</tr>
<tr>
<td></td>
<td>Adolescent Survey</td>
<td>1,100</td>
<td>1</td>
<td>2/60 (0.033)</td>
<td>36.67</td>
</tr>
<tr>
<td>Total</td>
<td></td>
<td>17,156</td>
<td></td>
<td></td>
<td>2,176.87</td>
</tr>
</tbody>
</table>

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proposed performance of the function of the agency, including whether the information will have practical utility; (2) 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) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to 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.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Nancy Breen, PhD, Project Officer, National Cancer Institute, EPN 4094, 6130 Executive Boulevard MSC 7344, Bethesda, Maryland 20852–7344, or call non-toll
free number 301–496–4675 or e-mail your request, including your address to: breen@mail.nih.gov.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of this publication.

Dated: November 9, 2010.

Vivian Horovitch-Kelley, NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2010–28648 Filed 11–12–10; 8:45 am]

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Income Withholding for Support (IWO).

OMB No.: 0970–0154.

ANNUAL BURDEN ESTIMATES

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Income Withholding for Support (Form)</td>
<td>58</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>e-IWO Record Layouts</td>
<td>58</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

Estimated Total Annual Burden Hours: 0.

Additional Information

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L’Enfant Promenade, SW., Washington, DC 20447. Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: infocollection@acf.hhs.gov.

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the Federal Register.

Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project. Fax: 202–395–7285.

E-mail: OIRA SUBMISSION@OMB.EOP.GOV.

Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–28615 Filed 11–12–10; 8:45 am]

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–2336–FN]

Medicare and Medicaid Programs; Approval of Det Norske Veritas Healthcare for Deeming Authority for Critical Access Hospitals

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Final notice.

SUMMARY: This final notice announces our decision to approve Det Norske Veritas Healthcare (DNVHC) for recognition as a national accreditation program for critical access hospitals seeking to participate in the Medicare or Medicaid programs.

DATES: Effective Date: This final notice of approval is effective December 23, 2010, through December 23, 2014.

FOR FURTHER INFORMATION CONTACT: Lillian Williams, (410) 786–8636. Patricia Chmielewski, (410) 786–6899.

SUPPLEMENTARY INFORMATION:

I. Background

Under the Medicare program, eligible beneficiaries may receive covered services in a critical access hospitals (CAHs) provided certain requirements are met. Sections 1820(c)(2)(B) and 1861(mm) of the Social Security Act (the Act) establish distinct criteria for facilities seeking designation as a CAH. The minimum requirements that a CAH must meet to participate in Medicare are set forth in regulation at 42 CFR part 485, subpart F. Conditions for Medicare payment for CAHs are set forth at § 413.70. Applicable regulations concerning provider agreements are located in 42 CFR part 489, and those pertaining to facility survey and