Advisory Group at prevention.council@hhs.gov.


SUPPLEMENTARY INFORMATION: On June 10, 2010, the President issued Executive Order 13544 to comply with the statutes under Section 4001 of the Patient Protection and Affordable Care Act, Public Law 111–148. This legislation mandated that the Advisory Group was to be established within the Department of Health and Human Services. The charter for the Advisory Group was approved by the Secretary of Health and Human Services on June 23, 2010; the charter was filed with the appropriate Congressional committees and Library of Congress on June 24, 2010. The Advisory Group has been established as a non-discretionary Federal advisory committee.

The Advisory Group has been established to provide recommendations and advice to the National Prevention, Health Promotion and Public Health Council (the “Council”). The Advisory Group shall provide assistance to the Council in carrying out its mission.

The Advisory Group membership shall consist of not more than 25 non-Federal members to be appointed by the President. The membership shall include a diverse group of licensed health professionals, including integrative health practitioners who have expertise in (1) worksite health promotion; (2) community services, including community health centers; (3) preventive medicine; (4) health coaching; (5) public health education; (6) geriatrics; and (7) rehabilitation medicine. There are currently 22 members of the Advisory Group. This will be the sixth meeting of the Advisory Group.

Public attendance at the web meeting is limited. Members of the public who wish to attend the web meeting must register by 12:00 p.m. EST June 19, 2012. Individuals should notify the designated contact to register for public attendance at prevention.council@hhs.gov.

Individuals who plan to attend the web meeting and need special assistance and/or accommodations should notify the designated contact for the Advisory Group. The public will have opportunity to provide electronic written comments to the Advisory Group on the June 25, 2012 during the web meeting. Any member of the public who wishes to have printed material distributed to the Advisory Group for this scheduled web meeting should submit material to the designated point of contact for the Advisory Group no later than 12:00 p.m. EST June 14, 2012.


BILLING CODE 4150–28–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Medicare & Medicaid Services

[Document Identifier CMS–10320]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS) is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

This information collection is mandated by Sections 1103 and 10102 of The Patient Protection and Affordability Care Act. Public Law 111–148 (ACA). Once all of the information is collected from insurance issuers of major medical health insurance (hereon referred to as issuers) and other affected parties, it will be displayed at http://www.healthcare.gov. Issuers are required to provide information quarterly, and healthcare.gov will be updated on a periodic schedule during each quarter. The information provided will help the general public make educated decisions about organizations providing private health care insurance.

In accordance with the provisions of the ACA referenced above, the U.S. Department of Health and Human Services created a Web site called healthcare.gov to meet these and other provisions of the law, and data collection was conducted for six months based upon an emergency information collection request. The interim final rule published on May 5, 2010 served as the emergency Federal Register Notice for the prior Information Collection Request (ICR). The Office of Management and Budget (OMB) reviewed this ICR under emergency processing and approved the ICR on April 30, 2010. The CCIIO is currently submitting a new ICR to OMB for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection is published to obtain comments from the public and affected agencies.

CCII0 is currently updating a system (hereon referred to as web portal) where State Departments of Insurance and issuers may log in using a custom user ID and password validation. The States may be asked to provide information on issuers in their State and various Web sites maintained for consumers. The issuers will be tasked with providing information on their major medical insurance products and plans. They will ultimately be given the choice to download a basic information template to enter data then upload into the web portal itself; or to submit .xml files containing their information. Once the
States and issuers submit their data, they will receive an email notifying them of any errors, and that their submission was received.

CCIIO is mandating the issuers verify and update their information on a quarterly basis and is requesting that States verify State-submitted information on an annual basis. In the event that an issuer enhances its existing plans, proposes new plans, or deactivates plans, the organization would be required to update the information in the web portal. Changes occurring during the three month quarterly periods will be allowed utilizing effective dates for both the plans and rates associated with the plans.

Information that is to be collected from State high risk pools will be collected from The National Association of State Comprehensive Health Insurance Plans (NASCHIP) at this time. Updates to this information may be submitted voluntarily. The estimated hour burden on issuers for the Plan Finder data collection in the first year is estimated as 90,400 total burden hours, or 113 hours per organization. This estimate is based on an assumed average of 450 individual plan issuers and 700 small group plan issuers per each of the four quarterly collections. It includes 30 hours per organization for training and communication, and 700 small group plan issuers per each of the four quarterly collections. It includes 30 hours per organization for training and communication. Additionally, for each of the issuers it includes 11 hours of preparation time, one hour of login and upload time, two hours of troubleshooting and data review and one half hour for attestation per organization per quarterly refresh. The estimated hour burden on the States is informed by the fact that they have already submitted the data once and are informed by the fact that they have already submitted the data once and only need to update. The overall hours estimate is 575, or 11.5 per Department of Insurance. This is premised on two hours of training and communication, 8 hours for data collection, and one half hour of submission.

Form Number: CMS–10320 (OMB#: 0938–1086); Frequency: Reporting—Annually/Quarterly; Affected Public: Health insurance issuers in the individual and small group markets; Number of Respondents: 801; Total Annual Responses: 3,051; Total Annual Hours: 90,400. (For policy questions regarding this collection contact Joe Mercer at 301–492–4265. For all other issues call 410–786–1326.)

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS' Web site at http://www.cms.hhs.gov/PaperworkReductionActof1995, or email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov, or call the Reports Clearance Office at 410–786–1326.

In commenting on the proposed information collections please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in one of the following ways by August 6, 2012.

1. Electronically. You may submit your comments electronically to http://www.regulations.gov. Follow the instructions for “Comment or Submission” or “More Search Options” to find the information collection document(s) accepting comments.

2. By regular mail. You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.


Martique Jones,
Director, Regulations Development Group, Division B, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2012–13480 Filed 6–1–12; 11:15 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Study of Coordination of Tribal TANF and Child Welfare Services.
OMB No.: New Collection.
Description: Study of Coordination of Tribal TANF and Child Welfare Services is sponsored by the Office of Planning, Research and Evaluation (OPRE), Administration for Children and Families of the U.S. Department of Health and Human Services. The study examines the approaches and strategies utilized by tribes and tribal organizations that were awarded the grants for Coordination of Tribal TANF and Child Welfare Services to Tribal Families at Risk of Child Abuse or Neglect.

The descriptive study of these programs that serve tribal communities will document the way in which the tribal grantees are creating and adapting culturally relevant and appropriate approaches, systems, and programs to increase coordination and enhance service delivery to address child abuse and neglect. The study will also document challenges faced and lessons learned to inform the field of practice as well as policymakers and funders at various levels.

The proposed information collection activities consist of semi-structured interviews, conducted at each of the 14 tribal communities, and a grantee feedback survey on the usefulness of periodically held cross-grantee learning events.

Respondents: Program director(s), tribal TANF and child welfare staff and supervisors, program partners, and tribal leaders or elders. The information collection does not include direct interaction with individuals or families that receive the services.

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Annual number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total Annual burden hours</th>
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<tr>
<td>Interview Protocol for Program Staff</td>
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<td>3</td>
<td>1.5</td>
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<tr>
<td>Interview Protocol for TANF and CW Staff</td>
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<td>Interview Protocol for Tribal or Community Partners</td>
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<td>Interview Protocol for Tribal Leaders or Elders</td>
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<td>Feedback Form for Community of Learning Events</td>
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<td>5</td>
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</table>

Estimated Total Annual Burden Hours: ........................................ .......................... 153