

requested. This includes the time needed to review instructions, to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and

maintaining information, and disclosing and providing information, to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information, and to

transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
MRC Unit Leader	1,000	6	1.0	6,000

Keith A. Tucker,

*Information Collection Clearance Officer,
Department of Health and Human Services.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Chronic Fatigue Syndrome Advisory Committee

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of the Assistant Secretary for Health.

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Services is hereby giving notice that the Chronic Fatigue Syndrome Advisory Committee (CFSAC) will hold a meeting. The meeting will be open to the public.

DATES: The meeting will be held on Wednesday, October 3, 2012 and Thursday, October 4, 2012 from 9 a.m. until 5 p.m.

ADDRESSES: Department of Health and Human Services; Hubert H. Humphrey Building; 200 Independence Avenue SW., Room 800, Washington, DC 20201. For a map and directions to the Hubert H. Humphrey building, please visit <http://www.hhs.gov/about/hhhmap.html>.

FOR FURTHER INFORMATION CONTACT: Nancy C. Lee, M.D., Designated Federal Officer, Chronic Fatigue Syndrome Advisory Committee, Department of Health and Human Services, 200 Independence Avenue SW., Room 712E, Washington, DC 20201. Any questions about meeting registration or public comment sign-up should be directed to CFSACOctober2012@seamoncorporation.com. Please direct other inquiries to cfsac@hhs.gov.

SUPPLEMENTARY INFORMATION: CFSAC was established on September 5, 2002 to

advise, consult with, and make recommendations to the Secretary through the Assistant Secretary for Health, on a broad range of topics including: (1) The current state of knowledge and research on the epidemiology, etiologies, biomarkers, treatment, and risk factors relating to chronic fatigue syndrome (CFS), to identify potential opportunities in these areas; (2) the impact and implications of current and proposed diagnosis and treatment methods for CFS; (3) development and implementation of programs to inform the public, health care professionals, and the biomedical research communities about CFS; and (4) strategies to improve the quality of life for CFS patients.

The agenda for this meeting is being developed and will be posted on the CFSAC Web site, <http://www.hhs.gov/advcomcfs> when finalized. The meeting will be live-video streamed at www.HHS.gov/Live and archived through the CFSAC Web site: www.hhs.gov/advocomcfs. Listening-only audio via telephone will be available on both days. Call-in information will be posted on the CFSAC Web site.

Individuals who plan to attend should register at the following link by September 28, 2012: <http://www.blsmmeetings.net/CFSACOctober2012>. Attendance by visitors who are not U.S. citizens is welcome, but prior approval is required by sending a request to CFSACOctober2012@seamoncorporation.com. Members of the media will also need to register. All attendees will be required to show government-issued picture identification (state or federal) for entry into the federal building. Attendees will receive a wrist band that must be worn the entire time. Security requires all non-federal employees to be escorted the entire time they are in the building. Upon leaving the building for any reason individuals will be required to

follow the security steps mentioned above and receive a new wrist band.

Members of the public will have the opportunity to provide public comments at the meeting or via telephone. International calls cannot be accommodated. A separate sign-up process for requesting time for public comment must be completed by September 24, 2012 at the following link: <http://www.blsmmeetings.net/CFSACPublicCommentOctober2012>. It is requested that individuals wishing to provide public comment submit a copy of their testimony (5 pages or less) in advance. It is preferred that individuals email their testimony (in MS WORD format, single spaced, using a 12 point font) to CFSACOctober2012@seamoncorporation.com by Monday, September 24, 2012. Testimony may also be mailed to the following address: Chronic Fatigue Syndrome Advisory Committee, Department of Health and Human Services, 200 Independence Avenue SW. Room 712E, Washington, DC, 20201. Mailed testimony must be received no later than Monday, September 24, 2012. Note: PDF files, hand-written notes and photographs will not be accepted. Requests for public comment and written testimony will not be accepted through the CFSAC mailbox. Also, the CFSAC mailbox will not respond to questions about specific public comment requests.

All public comment becomes part of the public record, available for viewing and posted on the CFSAC Web site. All testimony and printed material submitted for the meeting are part of the official meeting record and will be uploaded to the CFSAC Web site and made available for public inspection. Testimony and materials submitted should not include sensitive personal information, such as social security number, birthdates, driver's license number, state identification or foreign country equivalent, passport number, financial account number, or credit or debit card number. Sensitive health information, or non-public corporate or

trade association information, such as trade secrets or other proprietary information should be excluded from any materials submitted. If you wish to remain anonymous the document must specify this.

We will confirm your time for public comment via email by September 28, 2012. Each speaker will be limited to five minutes per speaker; no exceptions will be made. We will give priority to individuals who have not provided public comment within the previous year.

Persons who wish to distribute printed materials to CFSAC members should submit one copy to Designated Federal Officer at cfsac@hhs.gov, prior to Friday, September 28, 2012. Submissions are limited to five typewritten pages.

Dated: September 4, 2012

Nancy C. Lee,

Designated Federal Officer, Chronic Fatigue Syndrome Advisory Committee.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10445, CMS-10164, CMS-10143 and CMS-838]

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.

1. *Type of Information Collection*
Request: New collection; *Title:* Medicare

Advantage Quality Bonus Payment Demonstration; *Use:* In response to the provision of the Affordable Care Act, beginning in 2012, quality bonus payments (QBP) are given to all plans earning four or five stars in Medicare's Star Rating program. As an extension of this legislation, CMS launched the Medicare Advantage Quality Bonus Payment Demonstration, which accelerates the phase-in of QBPs by extending bonus payments to three-star plans and eliminating the cap on blended county benchmarks that would otherwise limit QBPs. Through this demonstration, CMS seeks to understand how incentive payments impact plan quality across a broader spectrum of plans.

The data collection effort will be conducted in the form of a survey of Medicare Advantage Organizations (MAOs) and up to 10 case studies with MAOs in order to supplement what can be learned from the analyses of administrative and financial data for MAOs, and from an environmental and literature scan. The data collected is needed to evaluate the QBP demonstration to better understand what impact the demonstration has had on MAO operations and their efforts to improve quality. The data collection instrument is a survey questionnaire designed to capture information on how MAOs perceive the demonstration and are planning for or implementing changes in quality initiatives and to identify factors that help or hinder the capacity to achieve quality improvement and that influence the decision calculus to make changes. Specifically, the information is expected to provide a detailed picture to CMS of the kinds of quality initiatives utilized by MAOs and some preliminary information on how they assess the effectiveness of these programs. The survey is designed to provide an overall picture of the QBP that can be used for national comparisons across plans as part of the larger evaluation of the QBP demonstration.

The case studies will be conducted as a series of open-ended discussions with MAO staff that will be guided by a discussion protocol. The case studies will supplement the information gathered from the survey and data analysis, providing valuable context and details about successful quality improvement activities. The case studies are particularly well suited to exploring the detailed characteristics of the plans' quality improvement activities, emphasizing the decision-making and thought processes underlying the structure and direction of their efforts and capturing the

contextual factors that impact the nature, structure, and scope of the programs. *Form Number:* CMS-10445 (OCN: 0938-New); *Frequency:* Annual; *Affected Public:* Private Sector—Business or other for-profits; *Number of Respondents:* 730; *Total Annual Responses:* 1,280; *Total Annual Hours:* 683. (For policy questions regarding this collection contact Gerald Riley at 410-786-6699. For all other issues call 410-786-1326.)

2. *Type of Information Collection*
Request: Reinstatement with a change of a previously approved collection; *Title:* Medicare Electronic Data Interchange (EDI) Registration and Electronic Data Interchange (EDI) Enrollment Form; *Use:* The purpose of this collection to obtain information that will be subsequently used during transaction exchange for identification of Medicare providers/suppliers and authorization of requested Electronic Data Interface (EDI) functions. The EDI Enrollment and the Medicare Registration Forms are completed by Medicare providers/suppliers and submitted to Medicare contractors. Authorization is needed for providers and suppliers to send and receive HIPAA standard transactions directly (or through a designated 3rd party) to and from Medicare contractors. Medicare contractors would use the information for initial set-up and maintenance of the access privileges. The use of the standard form provides an efficient uniform means by which Medicare captures information necessary to drive Medicare EDI security and EDI access privileges. All EDI providers will complete and sign the EDI Enrollment Form along with the Medicare EDI Registration Form. They will also reconfirm their access privileges annually.

The information collected will be uploaded into Medicare contractor computer systems. Medicare contractors will store this information in a database accessed at the time of provider connection to the Medicare Data Contractor Network (MDCN). When authentication is successful and connectivity is established, transactions may be exchanged. The information will be stored in a computer data base and used to authenticate the user on day-to-day electronic commerce, support the submitter and password administration function, and validate access relationships between providers/suppliers and their designated EDI submitter/receiver on a per transaction basis. *Form Number:* CMS-10164 (OCN: 0938-0983); *Frequency:* Once; *Affected Public:* Private Sector—Business or other for-profits, Not for-profit institutions; *Number of Respondents:*