disclosing or otherwise making available to Magellan, any non-public information relating to KMI; and (3) institute procedures and requirements throughout the various entities of the proposed respondents to ensure that non-public information is protected as required by the proposed Order.

Paragraph II.D. provides that, for the time period that Carlyle or Riverstone holds, directly or indirectly, any interest in Magellan, Carlyle and Riverstone shall not, without providing thirty days advance written notification, acquire any stock, share capital, equity or other interest in KMI other than the interest acquired through the Acquisition. This prior notice gives the Commission the opportunity to analyze additional purchases of KMI by the proposed Respondents that may change the economic incentives of the proposed Respondents. Advance notice is not required in certain limited situations where investments are effectively passive or where the Respondents' relative ownership interests would not change. In such situations, the Respondents must provide notification under Paragraph II.E. within ten days after such acquisitions.

C. Implementation Monitor

To assure that the firewall provisions of Paragraphs II.B. and II.C. of the Order are properly implemented and enforced, the Order requires an Implementation Monitor to monitor these obligations. Pursuant to Paragraph IV, Mr. Kevin Sudy, an Associate Director at Navigant Consulting, will be appointed as the Implementation Monitor and shall serve until such time as he reports to the Commission that the parties have established adequate procedures under the terms of the proposed Order and the Commission notifies the parties that such procedures are acceptable. The Commission reserves the right subsequently to reinstate the monitor as necessary and appropriate to ensure compliance by Respondents with the terms of the proposed Order. The Implementation Monitor is important to assuring compliance with the firewall provisions of the Order.

D. Notice Provisions

Paragraph II.E. requires the proposed Respondents to provide the Commission with written notice within ten days if they (1) no longer hold any interest in Magellan, other than a wholly passive investment, (2) no longer hold any interest in Magellan, (3) no longer hold any interest in KMI or no longer have the ability to influence or have representation at KMI, (4) acquire interest in interest in KMI through a passive investment fund, or (5) acquire any interest in Magellan.

Paragraph III of the proposed Order requires the proposed Respondents to send notice of the Order, Complaint, and Analysis to Aid Public Comment in this matter to certain persons likely to have competitively sensitive information subject to this Order or likely to be impacted by the firewall provisions of the Order, including persons on the Magellan and KMI Boards of Directors, and other persons involved in the Acquisition of KMI.

Paragraph V.A. requires periodic reports until the Implementation Monitor and the Commission are satisfied that the firewalls are properly established and adequately protect the flow of non-public information as required by the Order. Paragraph V.B. requires annual reports until the Order terminates in ten years.

Paragraph VI requires the proposed Respondents to give the Commission prior notice of certain events that may change their obligations under the Order.

E. Additional Provisions

Paragraph VII allows the Commission to have access to personnel and documents at the offices of the proposed Respondents with proper notice for purposes of determining or securing compliance with this Order.

Paragraph VIII provides that the Order shall terminate after ten years.

V. The Order to Maintain Assets

The Commission has also issued an Order to Maintain Assets in this proceeding, which effectively requires the proposed Respondents to adhere to the terms of the proposed Order during the time period leading up to their proposed Acquisition of equity interests in KMI.

VI. Opportunity for Public Comment

The proposed Consent Agreement has been placed on the public record for thirty (30) days for receipt of comments by interested persons. The Commission has also issued its Complaint in this matter. Comments received during this comment period will become part of the public record. After thirty days, the Commission will again review the proposed Consent Agreement and the comments received and will decide whether it should withdraw from the Agreement or make final the Agreement's proposed Order.

By accepting the proposed Consent Agreement subject to final approval, the Commission anticipates that the competitive problems alleged in the Complaint will be resolved. The purpose of this analysis is to invite public comment on the proposed Order to aid the Commission in its determination of whether it should make final the proposed Order contained in the Agreement. This analysis is not intended to constitute an official interpretation of the proposed Order, nor is it intended to modify the terms of the proposed Order in any way.

By direction of the Commission, Commissioner Leibowitz dissenting and Commissioner Rosch recused.

Donald S. Clark,

Secretary.

[FR Doc. E7–1479 Filed 1–30–07; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: OS–0990–0220; 60-Day Notice]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Office of the Secretary, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection 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.

Type of Information Collection Request: Extension.

Title of Information Collection: Voluntary Academic and Industry Partner Surveys to Implement Executive Order 12862 and 5 U.S.C. 305 for the Dept. of Health and Human Services.

Form/OMB No.: 0990–0220.

Use: The Office of Acquisition Management Policy (OAMP) under the Assistant Secretary for Administration and Management (ASAM) and the Office of Grants (OG) under the Assistant Secretary for Resources and

Technology (ASRT), Office of the Secretary, Department of Health and Human Services (HHS) request that the Office of Management and Budget (OMB) extend its existing approval under Clearance No. 0990–0220 for HHS to undertake voluntary surveys of HHS' partners in academia and industry (e.g., Principal Investigators, business offices, and vendors) through January 31, 2010. To comply with Executive Order 12862, Setting Customer Service Standards (the EO), HHS again plans to systematically survey its grant recipients and contractors to compile their evaluations of the Department's grants and procurement processes, and to improve the way we conduct business with them.

These voluntary surveys will continue to be a collaborative effort, with OAMP and OG providing leadership, oversight, and a methodology; and the HHS Operating Divisions (OPDIVs) conducting the surveys for their own operations. Each OPDIV will conduct web-based surveys of its partners to obtain feedback for improving business processes. The grant recipients and contractors to be surveyed are sufficiently familiar with the Department and its OPDIVs to make this feedback extremely useful. These surveys will give OAMP, OG, and each of the OPDIVs an opportunity to understand and evaluate grant and procurement quality standards, as well as to incorporate best industry or public sector standards into OPDIV practices.

Frequency: Reporting every 3 years.

Affected Public: Business or other for-profit, Not-for-profit institutions, Federal Government.

Annual Number of Respondents: 2133.

Total Annual Responses: 2133.

Average Burden per Response: 10.75 minutes.

Total Annual Hours: 382.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, e-mail your request, including your address, phone number, OMB number, and OS document identifier, to *Sherette.funncoleman@hhs.gov*, or call the Reports Clearance Office on (202) 690–6162. Written comments and recommendations for the proposed information collections must be received within 60 days, and directed to the OS Paperwork Clearance Officer at the following address: Department of Health and Human Services, Office of the Secretary, Assistant Secretary for Resources and Technology, Office of Resources Management, *Attention:* Sherrette Funn-Coleman (0990–0220), Room 537–H, 200 Independence Avenue, SW., Washington, DC 20201.

Dated: January 23, 2007.

Alice Bettencourt,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

[FR Doc. E7–1464 Filed 1–30–07; 8:45 am]

BILLING CODE 4151–17–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–07–07AL]

Proposed Data Collections Submitted for Public Comment and Recommendations

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 Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–5960 and send comments to Joan Karr, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an e-mail to *omb@cdc.gov*.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

Proposed Project

Evaluation of the Successful Business Strategies to Prevent Heart Disease and Stroke Toolkit—NEW—Division for Heart Disease and Stroke Prevention (DHDSP), National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Under Part C (Centers for Disease Control and Prevention) of the Statement of Organization Functions, and Delegations of Authority of the Department of Health and Human Services (45 FR 67772–76, dated October 14, 1980, and corrected at 45 FR 69296, October 20, 1980, as amended most recently at 70 FR 72842–72843, dated December 7, 2005), the Division for Heart Disease and Stroke Prevention, National Center for Chronic Disease Prevention and Health Promotion, Centers for Disease Control and Prevention was established. This Division plans, directs, and coordinates programs to reduce morbidity, risk factors, costs, disability, mortality, and disparities associated with heart disease, stroke, and other cardiovascular disease outcomes. Under this Division, formative research was conducted to identify effective interventions and promising practices for preventing heart disease and stroke at the work site. In 2005, this research resulted in the development of a Successful Business Strategies to Prevent Heart Disease and Stroke Toolkit. The toolkit provides state programs with suggestions about which health benefits, services, and interventions can improve employee cardiovascular health, prevent heart disease and stroke, and reduce related costs. The second phase of this project focuses on disseminating and evaluating the Successful Business Strategies to Prevent Heart Disease and Stroke Toolkit.

As part of the Toolkit evaluation, the CDC has employed contractor support to design and conduct a Web-based survey of State Health Departments to gather information on their experiences with the Toolkit. The contractor will collect and analyze all data from this survey. The CDC has also contracted to make revisions to the Toolkit based on results of this survey, ongoing feedback from the States, and feedback from employers through interviews.

There are no costs to respondents except their time to complete the survey.