

the approximate time requested for the presentation. Oral comments made at the public meeting must also be submitted to the NIOSH Docket Office in writing in order to be considered by the Agency.

*Request for Information:* NIOSH seeks to obtain materials, including published and unpublished reports and research findings, to evaluate the possible health risks of occupational exposure to carbon nanotubes and nanofibers. Examples of requested information include, but are not to be limited to:

(1) Identification of industries or occupations in which exposures to carbon nanotubes and nanofibers can occur;

(2) Trends in the production and use of carbon nanotubes and nanofibers;

(3) Exposure measurement data;

(4) Case reports or other health information demonstrating possible health effects in workers exposed to carbon nanotubes or nanofibers;

(5) Reports of experimental in vivo and in vitro studies that provide evidence of a dose-relationship between exposure to carbon nanotubes and nanofibers and biological activity;

(6) Reports of experimental data on the airborne characteristics of carbon nanotubes or nanofibers, including information on the amounts that are inhalable and respirable;

(7) Criteria and rationale for including workers in a medical surveillance and screening program;

(8) Description of work practices and engineering controls used to reduce or prevent workplace exposure to carbon nanotubes and nanofibers; and

(9) Educational materials for worker safety and training on the safe handling of carbon nanotubes and nanofibers.

**ADDRESSES:** Written comments or requests to attend or present at the meeting, identified by docket number NIOSH-161-A, may be submitted by any of the following ways:

- *Mail:* NIOSH Docket Office, Robert A. Taft Laboratories, MS-C34, 4676 Columbia Parkway, Cincinnati, OH 45226.

- *Facsimile:* (513) 533-8285.

- *E-mail:* [nioshdocket@cdc.gov](mailto:nioshdocket@cdc.gov).

All information received in response to this notice will be available for public examination and copying at the NIOSH Docket Office, 4676 Columbia Parkway, Room 111, Cincinnati, OH 45226. A complete electronic docket containing all comments submitted will be available on the NIOSH Web page at <http://www.cdc.gov/niosh/docket>, and comments will be available in writing by request. NIOSH includes all comments received without change in

the docket, including any personal information provided. All electronic comments should be formatted as Microsoft Word. Please make reference to docket number NIOSH 161-A.

**FOR FURTHER INFORMATION CONTACT:**

Ralph D. Zumwalde, NIOSH, Robert A. Taft Laboratories MS-C32, 4676 Columbia Parkway, Cincinnati, OH 45226, telephone (513) 533-8320.

Dated: December 13, 2010.

**Tanja Popovic,**

*Deputy Associate Director for Science, Centers for Disease Control and Prevention.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Medicare & Medicaid Services**

[Document Identifier: CMS-R-153]

**Agency Information Collection Activities: Submission for OMB Review; 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), Department of Health and Human Services, 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 function; (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:* Reinstatement with change of a previously approved collection; *Title of Information Collection:* Medicaid Drug Utilization Review (DUR) Annual Report; *Use:* The DUR program is required to assure that prescriptions are appropriate, medically necessary and are not likely to result in adverse medical results. Each State DUR program must consist of prospective drug use review, retrospective drug use review, data assessment of drug use

against predetermined standards, and ongoing educational outreach activities. In addition, States are required to submit an annual DUR program report that includes a description of the nature and scope of State DUR activities. Over the years, technology has changed as has the practice of the pharmacy. Therefore, CMS has revised the old survey vehicle to more fully address the current practices and areas of concern with the Medicaid Pharmacy Programs. *Form Number:* CMS-R-153 (OMB#: 0938-0659); *Frequency:* Annually; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 51; *Total Annual Responses:* 51; *Total Annual Hours:* 20,298. (For policy questions regarding this collection contact Madlyn Kruh at 410-786-3239. 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 address at <http://www.cms.hhs.gov/PaperworkReductionActof1995>, or e-mail your request, including your address, phone number, OMB number, and CMS document identifier, to [Paperwork@cms.hhs.gov](mailto:Paperwork@cms.hhs.gov), or call the Reports Clearance Office on (410) 786-1326.

To be assured consideration, comments and recommendations for the proposed information collections must be received by the OMB desk officer at the address below, no later than 5 p.m. on *January 24, 2011*: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer. *Fax Number:* (202) 395-6974. *E-mail:* [OIRA\\_submission@omb.eop.gov](mailto:OIRA_submission@omb.eop.gov).

Dated: December 17, 2010.

**Martique Jones,**

*Director, Regulations Development Division-B, Office of Strategic Operations and Regulatory Affairs.*

[FR Doc. 2010-32196 Filed 12-22-10; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Medicare & Medicaid Services**

[Document Identifier CMS-10367]

**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