

environmental chemicals, physical substances, or mixtures (collectively referred to as "substances") cause adverse effects on reproduction and/or development and provide opinion on whether these substances are hazardous for humans. Information about CERHR can be obtained from its homepage (<http://cerhr.niehs.nih.gov>).

Dated: April 7, 2011.

**John R. Bucher,**

*Associate Director, National Toxicology Program.*

[FR Doc. 2011-9182 Filed 4-14-11; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Public Meeting of the Presidential Commission for the Study of Bioethical Issues

**AGENCY:** Presidential Commission for the Study of Bioethical Issues, Office of the Assistant Secretary for Health, Department of Health and Human Services.

**ACTION:** Notice of meeting.

**SUMMARY:** The Presidential Commission for the Study of Bioethical Issues will conduct its fifth meeting in May. At this meeting, the Commission will discuss the topic of Federal standards regarding human subjects protection in Federally funded scientific studies.

**DATES:** The meeting will take place Wednesday, May 18, 2011, from 9 a.m. to approximately 4:45 p.m., and Thursday, May 19, 2011, from 9 a.m. to approximately 1:15 p.m.

**ADDRESSES:** The Warwick New York Hotel, 65 West 54th Street, New York, NY 10019. Phone 212-247-2700.

**FOR FURTHER INFORMATION CONTACT:** Hillary Wicai Viers, Communications Director, Presidential Commission for the Study of Bioethical Issues, 1425 New York Avenue, NW., Suite C-100, Washington, DC 20005. Telephone: 202-233-3963. E-mail: [Hillary.Viers@bioethics.gov](mailto:Hillary.Viers@bioethics.gov). Additional information may be obtained at <http://www.bioethics.gov>.

**SUPPLEMENTARY INFORMATION:** Pursuant to the Federal Advisory Committee Act of 1972, Public Law 92-463, 5 U.S.C. app. 2, notice is hereby given of the fifth meeting of the Presidential Commission for the Study of Bioethical Issues (PCSBI). The meeting will be held from 9 a.m. to approximately 4:45 p.m. on Wednesday, May 18, 2011, and from 9 a.m. to approximately 1:15 p.m. on Thursday, May 19, 2011, at the Warwick New York Hotel, New York, NY. The

meeting will be open to the public with attendance limited to space available. The meeting will also be webcast at <http://www.bioethics.gov>.

Under authority of Executive Order 13521, dated November 24, 2009, the President established PCSBI to serve as a public forum and advise him on bioethical issues generated by novel and emerging research in biomedicine and related areas of science and technology. The Commission is charged to identify and promote policies and practices that assure ethically responsible conduct of scientific research, healthcare delivery, and technological innovation. In undertaking these duties, the Commission will examine specific bioethical, legal, and social issues related to potential scientific and technological advances; examine diverse perspectives and possibilities for useful international collaboration on these issues; and recommend legal, regulatory, or policy actions as appropriate.

The main agenda item for this fifth meeting is to review Federal as well as transnational standards of human subjects protections in scientific studies supported by the Federal government as requested by President Obama on November 24, 2010.

The draft meeting agenda and other information about PCSBI, including information about access to the webcast, will be available at <http://www.bioethics.gov>.

The Commission welcomes input from anyone wishing to provide public comment on any issue before it. There will be a public comment session in the afternoon on May 18, 2011. Individuals who would like to provide public comment at that time should notify Esther Yoo by telephone at 202-233-3960, or e-mail at [Esther.Yoo@bioethics.gov](mailto:Esther.Yoo@bioethics.gov) before May 9, 2011. To accommodate as many speakers as possible the time for each individual to speak may be limited. If the number of individuals wishing to speak is greater than can reasonably be accommodated during the scheduled meeting, the Commission may randomly select speakers from among those who register to speak.

Anyone planning to attend the meeting who needs special assistance, such as sign language interpretation or other reasonable accommodations, should also notify Esther Yoo (contact information above) in advance of the meeting. The Commission will make every effort to accommodate persons who need special assistance.

Written comments will also be accepted and are especially welcome. Please address written comments by e-

mail to [info@bioethics.gov](mailto:info@bioethics.gov), or by mail to the following address: Public Commentary, Presidential Commission for the Study of Bioethical Issues, 1425 New York Ave., NW, Suite C-100, Washington, DC 20005. Comments will be publicly available, including any personally identifiable or confidential business information that they contain. Trade secrets should not be submitted.

Dated: April 6, 2011.

**Valerie H. Bonham,**

*Executive Director, Presidential Commission for the Study of Bioethical Issues.*

[FR Doc. 2011-9123 Filed 4-14-11; 8:45 am]

**BILLING CODE 4154-06-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier CMS-10369]

#### 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 of Information Collection:* Solicitation for Applications: Community-based Care Transitions Program; *Use:* The Community-based Care Transitions Program (CCTP) described in Section 3026 of the Affordable Care Act will run for 5 years with a mandated start date of January 1, 2011. This program provides funding to community-based organizations in partnership with acute care hospitals for the provision of care transition services delivered to high risk

Medicare beneficiaries. The legislation provides \$500,000,000 for the program. The goals of the CCTP are to improve transitions of beneficiaries from the inpatient hospital setting to other care settings, to improve quality of care, to reduce readmissions for high risk beneficiaries, and to document measureable savings to the Medicare program. *Form Number:* CMS-10369 (OMB#: 0938-NEW); *Frequency:* Once; *Affected Public:* State, Local, or Tribal Governments; Private Sector—Business or other for-profits and not-for-profit institutions; *Number of Respondents:* 1,000; *Total Annual Responses:* 1,000; *Total Annual Hours:* 80,000. (For policy questions regarding this collection contact Juliana Tiongson at 410-786-0342. 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.gov/PaperworkReductionActof1995/PRAL/list.asp#TopOfPage> 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 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 *June 14, 2011*:

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.

Dated: April 6, 2011.

**Martique Jones,**

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

[FR Doc. 2011-9125 Filed 4-12-11; 11:15 am]

**BILLING CODE 4120-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Medicare & Medicaid Services**

[Document Identifier: CMS-216-94 and CMS-10112]

**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:* Extension of a currently approved collection; *Title of Information Collection:* Organ Procurement Organization/Histocompatibility Laboratory Statement of Reimbursable Costs, manual instructions and supporting regulations contained in 42 CFR 413.20 and 413.24; *Use:* This form is required by statute and regulation for participation in the Medicare program. The information is used to determine payment for Medicare. Organ Procurement Organizations and Histocompatibility Laboratories are the users. *Form Number:* CMS-216-94 (OMB# 0938-0102); *Frequency:* Yearly; *Affected Public:* Business or other for-profit, Not-for-profit institutions; *Number of Respondents:* 115; *Total Annual Responses:* 115; *Total Annual Hours:* 5175. (For policy questions regarding this collection contact Angela Havrilla at 410-786-4516 or Amelia Citerone at 410-786-3901. For all other issues call 410-786-1326.)

2. *Type of Information Collection Request:* Extension without change of a currently approved collection; *Title of Information Collection:* Phone Surveys

of Products and Services for Medicare Payment Validation and Supporting Regulations in 42 CFR 405.502. *Use:* The phone surveys of products and services for Medicare payment validation and supporting regulations in 42 CFR 405.502 will be used to identify specific products/services provided to Medicare beneficiaries and the costs associated with the provision of those products/services. The information collected will be used to validate the Medicare payment amounts for those products/services and institute revisions of payment amounts where necessary. The respondents will be the companies that have provided the product/service under review to Medicare beneficiaries. *Form Number:* CMS-10112 (OMB# 0938-0939); *Frequency:* Occasionally; *Affected Public:* Private sector—Business or other for-profit; *Number of Respondents:* 4,000; *Total Annual Responses:* 4,000; *Total Annual Hours:* 16,000. (For policy questions regarding this collection contact Michael Rich at 410-786-6856. For all other issues call 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 *May 16, 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: April 8, 2011.

**Michelle Shortt,**

*Director, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.*

[FR Doc. 2011-9024 Filed 4-14-11; 8:45 am]

**BILLING CODE 4120-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Medicare & Medicaid Services**

[Document Identifier: CMS-304 and CMS-304a; and CMS-368 and CMS-R-144]

**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.