

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Family History and Diamond Blackfan Anemia, DD11-010, Initial Review

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC) announces the aforementioned meeting:

Time and Date

11 a.m.–5 p.m., April 19, 2011 (Closed).

Place: Teleconference.

Status: The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92-463.

Matters To Be Discussed: The meeting will include the initial review, discussion, and evaluation of “Family History and Diamond Blackfan Anemia, DD11-010, initial review.”

Contact Person for More Information: Michael Dalmat, DrPH, Scientific Review Officer, CDC, National Center for Chronic Disease Prevention and Health Promotion, Office of the Director, Extramural Research Program Office, 4770 Buford Highway, NE., Mailstop K-92, Atlanta, GA 30341. Telephone: (770) 488-6423, E-mail: MED1@CDC.GOV.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: January 7, 2011.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2011-1233 Filed 1-20-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention (CDC)—Ethics Subcommittee (ES)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC)

announces the following meeting of the aforementioned subcommittee:

Times and Dates:

1 p.m.–5 p.m., February 17, 2011; 8:30 a.m.–12:30 p.m., February 18, 2011.

Place: CDC, Thomas R. Harkin Global Communications Center, Distance Learning Auditorium, 1600 Clifton Road, NE., Atlanta, GA 30333. This meeting is also available by teleconference. Please dial (877) 928-1204 and enter code 4305992.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 60 people. To accommodate public participation in the meeting, a conference telephone line will be available. The public is welcome to participate during the public comment. The public comment periods are tentatively scheduled for 4 p.m.–4:15 p.m. on February 17, 2011 and from 12 p.m.–12:15 p.m. on February 18, 2011.

Purpose: The ES will provide counsel to the ACD, CDC, regarding a broad range of public health ethics questions and issues arising from programs, scientists and practitioners.

Matter To Be Discussed: Agenda items will include the following: A review of public comments submitted on the ethical considerations document for the allocation of ventilators during a severe pandemic; finalizing the presentation for the ACD on ethical issues related to non-communicable disease prevention and control; and a review of the outcome of discussions with state, tribal, local, and territorial health officers on public health ethics challenges, including an assessment of available options towards the development of practical tools to assist health departments in their efforts to address these challenges.

The agenda is subject to change as priorities dictate.

Contact Person for More Information: For security reasons, members of the public interested in attending the meeting should contact Drue Barrett, PhD, Designated Federal Officer, ACD, CDC-ES, 1600 Clifton Road, NE., M/S D-50, Atlanta, Georgia 30333. Telephone (404) 639-4690. E-mail: d Barrett@cdc.gov. The deadline for notification of attendance is February 11, 2011.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: January 14, 2011.

Lorenzo J. Falgiano,

Acting Director, Management Analysis and Service Office, Centers for Disease Control and Prevention.

[FR Doc. 2011-1232 Filed 1-20-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-R-148, CMS-R-266, and CMS-216]

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:* Extension without change of a currently approved collection; *Title of Information Collection:* Limitations on Provider Related Donations and Health Care Related Taxes; Limitation on Payments for Disproportionate Share Hospitals and Supporting Regulations in 42 CFR 433.68, 433.74 and 447.272; *Use:* This information collection is necessary to ensure compliance with Sections 1903 and 1923 of the Social Security Act for the purpose of preventing payments of Federal financial participation on amounts prohibited by statute; *Form Number:* CMS-R-148 (OMB#: 0938-0618); *Frequency:* Quarterly and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 50; *Total Annual Responses:* 40; *Total Annual Hours:* 3,200. (For policy questions regarding this collection

contact Rory Howe at 410-786-4878. For all other issues call 410-786-1326.)

2. Type of Information Collection

Request: Extension of a currently approved collection; **Title of Information Collection:** Medicaid Disproportionate Share Hospital Annual Reporting; **Use:** Section 1923(j)(i) of the Social Security Act requires States to submit an annual report that identifies each disproportionate share hospital (DSH) that received a DSH payment under the State's Medicaid program in the preceding fiscal year and the amount of DSH payments paid to that hospital in the same year and such other information as the Secretary determines necessary to ensure the appropriateness of DSH payments. The information supplied will also satisfy the requirements under section 1923(a)(2)(D) of the Act; **Form Number:** CMS-R-266 (OMB#: 0938-0746); **Frequency:** Yearly; **Affected Public:** State, Local, or Tribal Governments; **Number of Respondents:** 52; **Total Annual Responses:** 52; **Total Annual Hours:** 1,976. (For policy questions regarding this collection contact Rory Howe at 410-786-4878. For all other issues call 410-786-1326.)

3. 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 the 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:** 5,175.

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 e-mail your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov, or call the Reports Clearance Office on (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 *March 22, 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.

Martique Jones,

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

[FR Doc. 2011-1265 Filed 1-20-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2011-N-0015]

Agency Information Collection Activities; Proposed Collection; Comment Request; Orphan Drugs; Common European Medicines Agency/ Food and Drug Administration Application Form for Orphan Medicinal Product Designation (Form FDA 3671)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the procedures by which sponsors of orphan drugs may request eligibility for the incentives by implementing a program as outlined in the Orphan Drug Act and the joint adoption by FDA and the European Medicines Agency (EMA) of the Common EMA/FDA Application Form for Orphan Medicinal Product Designation (form FDA 3671).

DATES: Submit either electronic or written comments on the collection of information by March 22, 2011.

ADDRESSES: Submit electronic comments on the collection of information to <http://www.regulations.gov>. Submit written comments on the collection of information to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT:

Jonna Capezzuto, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50-400B, Rockville, MD 20850. 301-796-3794. Jonnalynn.Capezzuto@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501-3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.