

Title: ACF Program Instruction: Children's Justice Act.
OMB No.: 0980-0196.

Description: The Program Instruction, prepared in response to the enactment of the Children's Justice Act (CJA), as set forth in Title II of Public Law 108-36, Child Abuse Prevention and Treatment Act Amendments of 2003, provides direction to the States and Territories to accomplish the purposes of assisting States in developing, establishing and operating programs designed to improve: (1) The handling of child abuse and neglect cases, particularly

child sexual abuse and exploitation, in a manner that limits additional trauma to the child victim; (2) the handling of cases of suspected child abuse or neglect-related fatalities; (3) the investigation and prosecution of cases of child abuse and neglect, particularly child sexual abuse and exploitation; and (4) the handling of cases involving children with disabilities or serious health-related problems who are victims of abuse and neglect. This Program Instruction contains information collection requirements that are found

in Public Law 108-36 at Sections 107(b) and 107(d), and pursuant to receiving a grant award. The information being collected is required by statute to be submitted pursuant to receiving a grant award. The information submitted will be used by the agency to ensure compliance with the statute; to monitor, evaluate and measure grantee achievements in addressing the investigation and prosecution of child abuse and neglect; and to report to Congress.

Respondents: State Governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Application & Annual Report	52	1	60	3,120

Estimated Total Annual Burden Hours: 3,120.

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: infocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments 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) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: November 16, 2010.

Robert Sargis,
Reports Clearance Officer.
[FR Doc. 2010-29189 Filed 11-18-10; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Office of the Secretary

Office of the Commissioner of Food and Drugs; Delegation of Authority

Notice is hereby given that I have delegated to the Commissioner of Food and Drugs the authorities vested in the Secretary of Health and Human Services under section 4 of the Federal Cigarette Labeling and Advertising Act (FCLAA)(15 U.S.C. 1333), as amended by the Family Smoking Prevention and Tobacco Control Act (Tobacco Control Act).

These authorities may be redelegated. These authorities shall be exercised under the Department's policy on regulations and the existing delegation of authority to approve and issue regulations.

I hereby ratify and affirm any actions taken by the Commissioner of Food and Drugs, or other FDA officials, which involved the exercise of the authorities delegated herein prior to the effective date of this delegation. This delegation is effective upon date of signature.

Dated: November 9, 2010.

Kathleen Sebelius,
Secretary.
[FR Doc. 2010-29150 Filed 11-18-10; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2010-N-0001]

Joint Meeting of the Anesthetic and Life Support Drugs Advisory Committee and the Drug Safety and Risk Management Advisory Committee; Cancellation

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Joint Meeting of the Anesthetic and Life Support Drugs Advisory Committee and the Drug Safety and Risk Management Advisory Committee scheduled for December 2, 2010, is cancelled. This meeting was announced in the **Federal Register** of November 1, 2010 (75 FR 67093). This meeting has been cancelled because the Agency believes the information received from previous advisory committee meetings is adequate to allow the Agency to address the specific concerns in the application that were delineated in the **Federal Register** notice of November 1, 2010.

FOR FURTHER INFORMATION CONTACT: Kalyani Bhatt, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, rm. 2417,

Silver Spring, MD 20993-0002, 301-796-9001, FAX: 301-847-8533, e-mail: kalyani.bhatt@fda.hhs.gov, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), codes 3014512529 or 3014512535. Please call the Information Line for up-to-date information on this meeting.

Dated: November 15, 2010.

Joanne Less,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2010-29280 Filed 11-18-10; 8:45 am]

BILLING CODE 4160-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2010-N-0001]

Cardiovascular and Renal Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Cardiovascular and Renal Drugs Advisory Committee.

General Function of the Committees: To provide advice and recommendations to the Agency on FDA's regulatory issues.

Date and Time: The meeting will be held on December 8, 2010, from 8 a.m. to 3:30 p.m.

Location: FDA White Oak Campus, Building 31, the Great Room, White Oak Conference Center, Rm. 1503, 10903 New Hampshire Ave., Silver Spring, MD 20993-0002. Information regarding special accommodations due to a disability, visitor parking and transportation may be accessed at: <http://www.fda.gov/AdvisoryCommittees/default.htm>; under the heading "Resources for You", click on "White Oak Conference Center Parking and Transportation Information for FDA Advisory Committee Meetings". Please note that visitors to the White Oak Campus must have a valid driver's license or other picture ID, and must enter through Building 1.

Contact Person: Elaine Ferguson, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993-0002, 301-796-9001, FAX: 301-847-8533, e-mail:

elaine.ferguson@fda.hhs.gov, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 3014512533. Please call the Information Line for up-to-date information on this meeting. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site and call the appropriate advisory committee hot line/phone line to learn about possible modifications before coming to the meeting.

Agenda: On December 8, 2010, from 8 a.m. to 11:30 a.m., the committee will discuss and provide general advice on the appropriate clinical study design for thromboxane receptor antagonists for prevention of cardiovascular events (such as heart attacks) in patients with aspirin intolerance due to immunologically-based adverse reactions (adverse events related to immune system function), specifically in the setting of coronary artery bypass grafting (also referred to as "heart bypass surgery").

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at <http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm>. Scroll down to the appropriate advisory committee link.

Procedure: On December 8, 2010, from 8 a.m. to 11:30 a.m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before December 6, 2010. Oral presentations from the public will be scheduled between approximately 10 a.m. and 11 a.m. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before November 29, 2010. Time allotted for each presentation may be limited. If the

number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by November 30, 2010.

Closed Presentation of Data: On December 8, 2010, from 12:30 p.m. to 3:30 p.m., the meeting will be closed to permit discussion and review of trade secret and/or confidential commercial information (5 U.S.C. 552b(c)(4)).

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Elaine Ferguson at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at <http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm> for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: November 15, 2010.

Joanne Less,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2010-29278 Filed 11-18-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant